

# Human Papillomavirus and Related Diseases Report

## UNITED KINGDOM OF GREAT BRITAIN AND NORTHERN IRELAND

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## **Executive summary**

Human papillomavirus (HPV) infection is now a well-established cause of cervical cancer and there is growing evidence of HPV being a relevant factor in other anogenital cancers (anus, vulva, vagina and penis) and head and neck cancers. HPV types 16 and 18 are responsible for about 70% of all cervical cancer cases worldwide. HPV vaccines that prevent against HPV 16 and 18 infection are now available and have the potential to reduce the incidence of cervical and other anogenital cancers.

This report provides key information for United Kingdom of Great Britain and Northern Ireland on cervical cancer, other anogenital cancers and head and neck cancers, HPV-related statistics, factors contributing to cervical cancer, cervical cancer screening practices, and HPV vaccine introduction. The report is intended to strengthen the guidance for health policy implementation of primary and secondary cervical cancer prevention strategies in the country.

Table 1: Key Statistics

Population						
	r (Female population aged >=15 yrs)		28.3 million			
Burden of cervical cancer an						
Annual number of cervical cance	er cases		3791			
Annual number of cervical cance			1121			
Crude incidence rates per 100,0	00 population:	Male	Female			
	Cervical cancer	-	11.0			
	Anal cancer	1.62	3.32			
	Vulva cancer	-	4.43			
	Vaginal cancer -					
	Penile cancer 2.27					
	Oropharyngeal cancer	6.29	2.04			
	Oral cavity cancer	11.7	6.95			
	Laryngeal cancer	6.31	1.46			
Burden of cervical HPV infe	etion					
Prevalence (%) of HPV 16 and/or	HPV 18 among women with:					
		Normal cytology	3.2			
		lesions (LSIL/CIN-1)	29.6			
	High-grade cervical lesions (HS	IL/CIN-2/CIN-3/CIS)	58.6			
	79.0					
Other factors contributing to						
Smoking prevalence (%) [95% U			18.5 [14.6-22.4]			
Total fertility rate (live births pe	er women)		1.9			
Oral contraceptive use (%)			28			
HIV prevalence (%) [95% UI], w	omen (15-49 years)		0.1 [0.1-0.1]			
Sexual behaviour			2.4.0/2.2			
	ave had sexual intercourse (men/women)		24.0/32.0			
Range of median age at first sex			16.0-19.1/16.0-20.9			
Cervical screening practices Existence of official national rec			V			
			Yes			
Starting year of current recomm	endations		- 37			
Active invitation to screening		<u> </u>	Yes			
	screening test used, and screening interval or	frequency of screen-	Varies by region			
ings HPV vaccine in females						
HPV vaccination programme			Introduced			
Year of introduction			2008			
Year of estimation of HPV vaccin	nation coverage		2003			
HPV coverage – first dose (%)	ianion coverage		77			
HPV coverage – last dose (%)			59			
Discount the mariful action for more information			อฮ			

<sup>\*</sup> Please see the specific sections for more information.

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## 1 Introduction



Figure 1: United Kingdom of Great Britain and Northern Ireland and Northern Europe

Information Centre aims to compile and centralise updated data and statistics on human papillomavirus (HPV) and related cancers. This report aims to summarise the data available to fully evaluate the burden of disease in United Kingdom of Great Britain and Northern Ireland and to facilitate stakeholders and relevant bodies of decision makers to formulate recommendations on the prevention of cervical cancer and other HPV-related cancers. Data include relevant cancer statistic estimates, epidemiological determinants of cervical cancer such as demographics, socioeconomic factors, risk factors, burden of HPV infection in women and men, cervical screening and immunization practices. The report is structured into the following sections:

Section 2, Demographic and socioeconomic factors. This section summarises the socio-demographic profile of United Kingdom of Great Britain and Northern Ireland. For analytical purposes, United Kingdom of Great Britain and Northern Ireland is classified in the geographical region of Northern Europe (Figure 1, lighter blue), which is composed of the following countries: Åland Islands, Channel Islands, Denmark, Estonia, Finland, Faeroe Islands, Guernsey, Isle of Man, Ireland, Iceland, Jersey, Lithuania, Latvia, Norway, Svalbard and Jan Mayen Islands, and Sweden. Throughout the report, United Kingdom of Great Britain and Northern Ireland estimates will be complemented with corresponding regional estimates.

**Section 3, Burden of HPV related cancers.** This section describes the current burden of invasive cervical cancer and other HPV-related cancers in United Kingdom of Great Britain and Northern Ireland ith estimates of prevalence, incidence, and mortality rates. Information in other HPV-related cancers includes other anogenital cancers (anus, vulva, vagina, and penis) and head and neck cancers (oral cavity, oropharyngeal, and larynx).

**Section 4, HPV related statistics**. This section reports on prevalence of HPV and HPV type-specific distribution in United Kingdom of Great Britain and Northern Ireland, in women with normal cytology, precancerous lesions and invasive cervical cancer. In addition, the burden of HPV in other anogenital cancers (anus, vulva, vagina, and penis), head and neck cancers (oral cavity, oropharynx, and larynx)

1 INTRODUCTION -3-

and men are presented.

**Section 5, Factors contributing to cervical cancer**. This section describes factors that can modify the natural history of HPV and cervical carcinogenesis such as smoking, parity, oral contraceptive use, and co-infection with HIV.

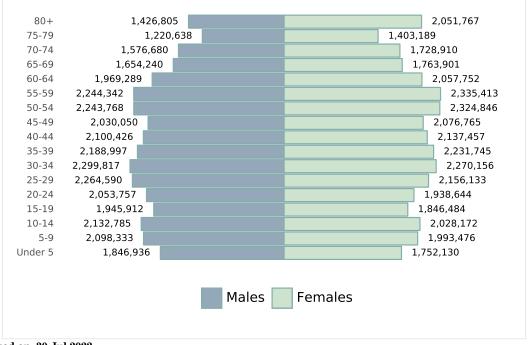
**Section 6, Sexual and reproductive health behaviour indicators**. This section presents sexual and reproductive behaviour indicators that may be used as proxy measures of risk for HPV infection and anogenital cancers, such as age at first sexual intercourse, average number of sexual partners, and anal intercourse among others.

**Section 7, HPV preventive strategies**. This section presents preventive strategies that include basic characteristics and performance of cervical cancer screening status, status of HPV vaccine licensure introduction, and recommendations in national immunisation programmes.

**Section 8, Protective factors for cervical cancer**. This section presents male circumcision and the use of condoms.

#### 2 Demographic and socioeconomic factors

Figure 2: Population pyramid of United Kingdom of Great Britain and Northern Ireland for 2022



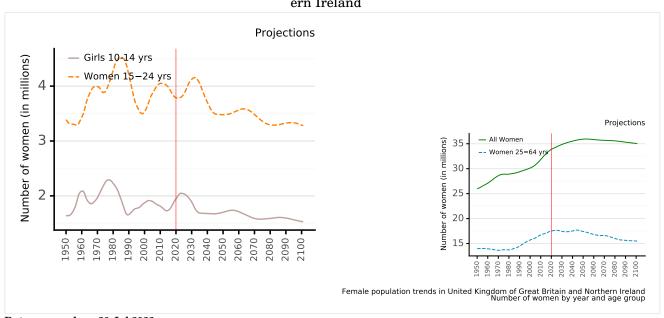
Data accessed on 30 Jul 2022

Refers to the United Kingdom of Great Britain and Northern Ireland. For statistical purposes, the data for United Kingdom do not include Anguilla, Bermuda, British Virgin Islands, Cayman Islands, Falkland Islands (Malvinas), Gibraltar, Guernsey, Isle of Man, Jersey, Montserrat, Saint Helena, Turks and Caicos Islands. Please refer to original source for methods of estimation

Year of estimate: 2022

Data Sources:
United Nations, Department of Economic and Social Affairs, Population Division (2022). World Population Prospects 2022, Online Edition. [Accessed on July 30, 2022].

Figure 3: Population trends in four selected age groups in United Kingdom of Great Britain and Northern Ireland



Data accessed on 30 Jul 2022

Refers to the United Kingdom of Great Britain and Northern Ireland. For statistical purposes, the data for United Kingdom do not include Anguilla, Bermuda, British Virgin Islands, Cayman Islands, Falkland Islands (Malvinas), Gibraltar, Guernsey, Isle of Man, Jersey, Montserrat, Saint Helena, Turks and Caicos Islands Please refer to original source for methods of estimation.

Year of estimate: 2022

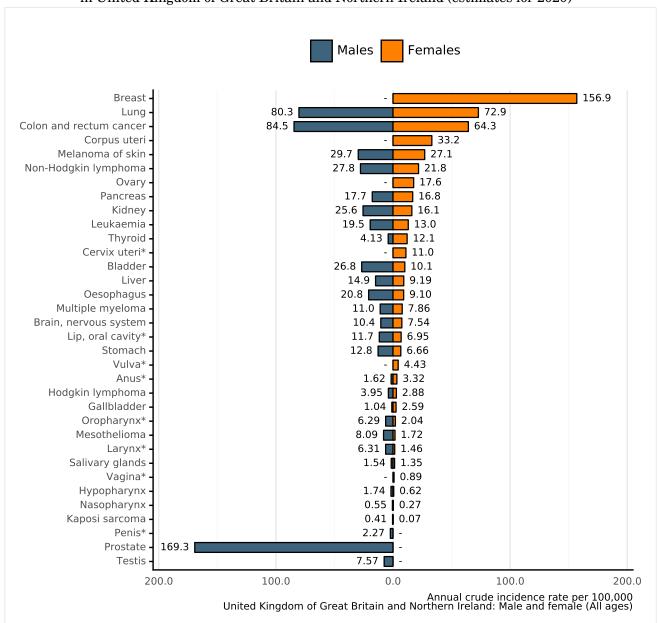
United Nations, Department of Economic and Social Affairs, Population Division (2022). World Population Prospects 2022, Online Edition. [Accessed on July 30, 2022].

## 3 Burden of HPV related cancers

HPV is the cause of almost all cervical cancer cases and is responsible for an important fraction of other anogenital and head and neck cancer. Here, we present the most recent estimations on the burden of HPV-associated cancer.

#### 3.1 HPV related cancers incidence

Figure 4: Comparison of HPV related cancers incidence to other cancers in men and women of all ages in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

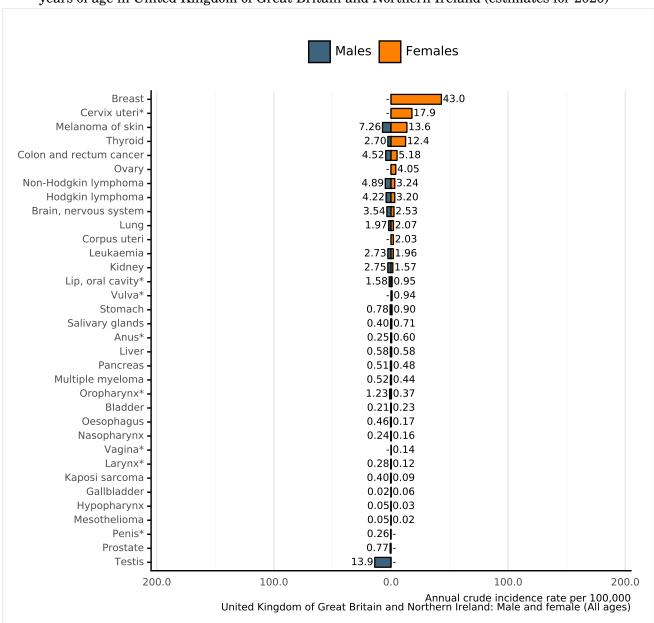
For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

Non-melanoma skin cancer is not included Rates per 100,000 men per year.

Rates per 100,000 men per year. Rates per 100,000 women per year.

Data Sources

Figure 5: Comparison of HPV related cancers incidence to other cancers among men and women 15-44 years of age in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



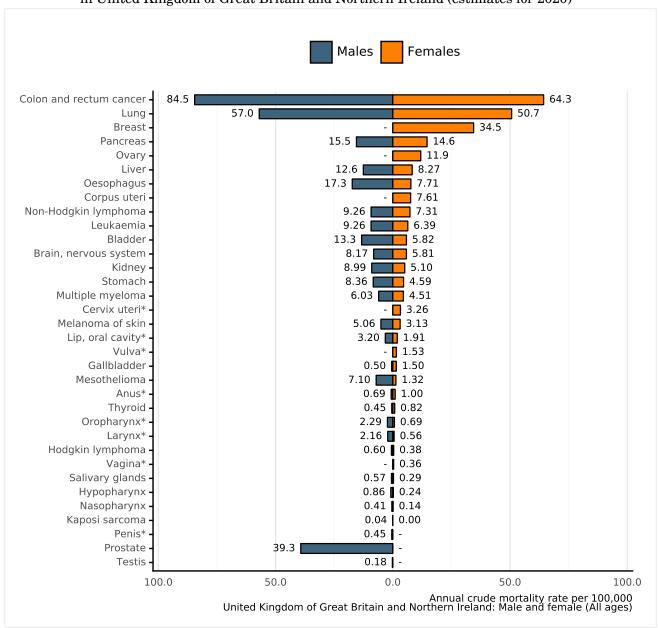
Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods Non-melanoma skin cancer is not included

Rates per 100,000 men per year. Rates per 100,000 women per year

## 3.2 HPV related cancers mortality

Figure 6: Comparison of HPV related cancers mortality to other cancers in men and women of all ages in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

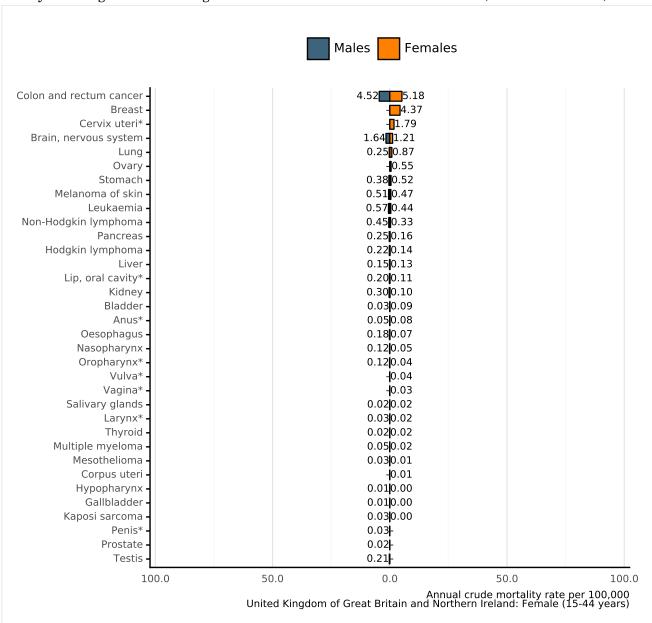
For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods Non-melanoma skin cancer is not included

Rates per 100,000 men per year.

Rates per 100,000 women per year.

Data Sources

Figure 7: Comparison of HPV related cancers mortality to other cancers among men and women 15-44 years of age in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



#### Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods Non-melanoma skin cancer is not included

Rates per 100,000 men per year. Rates per 100,000 women per year

#### 3.3 Cervical cancer

Cancer of the cervix uteri is the  $4^{th}$  most common cancer among women worldwide, with an estimated 604,127 new cases and 341,831 deaths in 2020. Worldwide, mortality rates of cervical cancer are substantially lower than incidence with a ratio of mortality to incidence to 57% (GLOBOCAN 2020). The majority of cases are squamous cell carcinoma followed by adenocarcinomas. (Vaccine 2006, Vol. 24, Suppl 3; Vaccine 2008, Vol. 26, Suppl 10; Vaccine 2012, Vol. 30, Suppl 5; IARC Monographs 2007, Vol. 90)

This section describes the current burden of invasive cervical cancer in United Kingdom of Great Britain and Northern Ireland and in comparison to geographic region, including estimates of the annual number of new cases, deaths, incidence, and mortality rates.

### 3.3.1 Cervical cancer incidence in United Kingdom of Great Britain and Northern Ireland

**Key Stats.** 

About 3,791 new cervical cancer cases are diagnosed annually in United Kingdom of Great Britain and Northern Ireland (estimations for 2020).

Cervical cancer ranks\* as the 12th leading cause of female cancer in United Kingdom of Great Britain and Northern Ireland.

Cervical cancer is the 2<sup>nd</sup> most common female cancer in women aged 15 to 44 years in United Kingdom of Great Britain and Northern Ireland.

Table 2: Cervical cancer incidence in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

Indicator	United Kingdom of Great Britain and Northern Ireland	Northern Europe	World
Annual number of new cancer cases	3,791	6,666	604,127
Uncertainty intervals of new cancer cases [95% UI]	[3,563-4,034]	[6,414-6,927]	[582,031-627,062]
Crude incidence rate <sup>b</sup>	11.0	12.4	15.6
Age-standardized incidence rate <sup>b</sup>	9.91	10.4	13.3
Cumulative risk (%) at 75 years old <sup>a</sup>	0.81	0.90	1.39

Data accessed on 27 Jan 2021

<sup>\*</sup> Ranking of cervical cancer incidence to other cancers among all women according to highest incidence rates (ranking 1st) excluding non-melanoma skin cancer. Ranking is based on crude incidence rates (actual number of cervical cancer cases). Ranking using age-standardized rate (ASR) may differ

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

<sup>a</sup> Cumulative risk (incidence) is the probability or risk of individuals getting from the disease during ages 0-74 years. For cancer, it is expressed as the % of new born children who would be expected to develop from a particular cancer before the age of 75 if they had the rates of cancer observed in the period in the absence of competing causes.  $^b$  Rates per 100,000 women per year.

Table 3: Cervical cancer incidence in United Kingdom of Great Britain and Northern Ireland by cancer

	registry			
Cancer registry	Period	N cases <sup>a</sup>	Crude rate <sup>b</sup>	ASR <sup>b</sup>
East Anglia <sup>1</sup>	1993-1997	547	10.2	7.2
East of England Region <sup>2</sup>	1998-2002	536	7.7	5.4
${ m England}^3$	2003-2007	11638	9.1	6.8
England and Wales <sup>4</sup>	1988-1990	13000	16.7	12.5
England: Mersey <sup>1</sup>	1993-1997	863	14	10.3
England: Merseyside and Cheshire <sup>2</sup>	1998-2002	728	11.9	8.3
England: North Western <sup>2</sup>	1998-2002	1289	12.2	8.8
England, North Western <sup>3</sup>	2003-2007	1654	9.9	7.4
England: North Western Region <sup>1</sup>	1993-1997	1611	15.7	11.7
England: Northern and Yorkshire <sup>2</sup>	1998-2002	2081	12.3	9.2
England, Northern and Yorkshire <sup>3</sup>	2003-2007	1941	11.3	8.8
England: Oxford Region <sup>2</sup>	1998-2002	407	5.9	4.2
England, Oxford Region <sup>3</sup>	2003-2007	569	8.1	6.3
England: South and Western Regions <sup>2</sup>	1998-2002	1809	10.6	7.5
England, South and Western Regions <sup>3</sup>	2003-2007	1634	9.3	7
England: South Thames Region <sup>1</sup>	1993-1997	1709	9.8	6.9
England: South Western Region <sup>4</sup>	1988-1992	1200	14.2	10.3
England: Thames <sup>2</sup>	1998-2002	2842	7.9	5.6
England, Thames <sup>3</sup>	2003-2007	2198	7.4	5.3
England: Trent <sup>2</sup>	1998-2002	1257	10.3	7.5
England, Trent <sup>3</sup>	2005-2007	841	11.1	8.9
England: Wessex <sup>4</sup>	1988-1992	1197	15.8	11.6
England: West Midlands <sup>2</sup>	1998-2002	1448	10.8	8.1
England, West Midlands <sup>3</sup>	2003-2007	1308	9.6	7.5
England: West Midlands Region <sup>1</sup>	1993-1997	1776	13.2	10.1
England: Yorkshire Region <sup>1</sup>	1993-1997	1387	14.6	10.8
Northern Ireland <sup>3</sup>	2003-2007	430	9.8	8
$Scotland^3$	2003-2007	1432	10.8	8.2
Scotland: Ayrshire <sup>5</sup>	1970-1972	77	13.6	10.1
Scotland: East <sup>6</sup>	1983-1987	158	15.4	11.3
Scotland: North <sup>6</sup>	1983-1987	111	19	15.1
Scotland: North-East <sup>6</sup>	1983-1987	238	17.2	13.7
Scotland: South-East <sup>6</sup>	1983-1987	602	19.5	15.2
Scotland: West <sup>4</sup>	1988-1992	1177	16.7	12.7
England, East of England Region <sup>3</sup>	2003-2007	982	6.9	5.1
$ m Wales^3$	2003-2007	842	11.1	8.4
National <sup>7</sup>	2008-2012	15526	9.7	7.7
England <sup>7</sup>	2008-2012	12606	9.4	7.5
England, East <sup>7</sup>	2008-2012	1229	8.3	6.6
England, East Midlands <sup>7</sup>	2008-2012	1218	10.7	8.8
England, London <sup>7</sup>	2008-2012	$-\frac{1470}{1470}$	7.2	5.4
England, North East <sup>7</sup>	2008-2012	810	12.2	10.4
England, North West <sup>7</sup>	2008-2012	1842	10.3	8.3
England, South West <sup>7</sup>	2008-2012	1392	10.4	8.7
England, South East <sup>7</sup>	2008-2012	1733	7.9	6.3
6, ·				

Continued on next page

Table 3 - continued from previous page

Cancer registry	Period	N cases <sup>a</sup>	Crude rate <sup>b</sup>	ASR <sup>b</sup>
England, West Midlands <sup>7</sup>	2008-2012	1460	10.3	8.5
England, Yorkshire and The Humber <sup>7</sup>	2008-2012	1452	10.9	9.1
Northern Ireland <sup>7</sup>	2008-2012	535	11.6	9.8
$Scotland^7$	2008-2012	1596	11.8	9.1
Wales <sup>7</sup>	2008-2012	789	10.1	8

#### Data accessed on 5 Oct 2018

Please refer to original source (available at http://ci5.iarc.fr/CI5-XI/Default.aspx)

ASR: Age-standardized rate, Standardized rates have been estimated using the direct method and the World population as the reference.

<sup>a</sup> Accumulated number of cases during the period in the population covered by the corresponding registry.

b Rates per 100,000 women per year.

Parkin, D.M., Whelan, S.L., Ferlay, J., Teppo, L., and Thomas, D.B., eds (2002). Cancer Incidence in Five Continents, Vol. VIII. IARC Scientific Publications No. 155, Lyon, IARC.

<sup>2</sup> Curado, M. P., Edwards, B., Shin, H.R., Storm, H., Ferlay, J., Heanue, M. and Boyle, P., eds (2007). Cancer Incidence in Five Continents, Vol. IX. IARC Scientific Publications No. 160, Lyon, IARC.

3 Forman D, Bray F, Brewster DH, Gombe Mbalawa C, Kohler B, Piñeros M, Steliarova-Foucher E, Swaminathan R and Ferlay J eds (2013). Cancer Incidence in Five Continents, Vol. X

 $<sup>(</sup>electronic\ version)\ Lyon,\ IARC.\ http://ci5.iarc.fr$ 

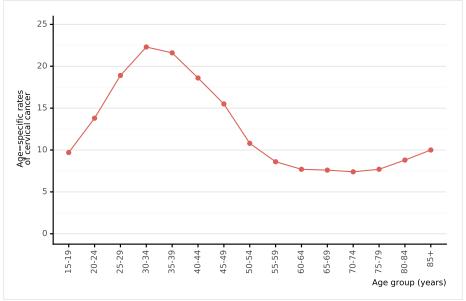
<sup>&</sup>lt;sup>4</sup> Parkin, D.M., Whelan, S.L., Ferlay, J., Raymond, L., and Young, J., eds (1997). Cancer Incidence in Five Continents, Vol. VII. IARC Scientific Publications No. 143, Lyon, IARC.

<sup>&</sup>lt;sup>5</sup> Waterhouse, J., Muir, C.S., Correa, P., Powell, J., eds (1976). Cancer Incidence in Five Continents, Vol. III. IARC Scientific Publications No. 15, Lyon, IARC.

<sup>6</sup> Parkin, D.M., Muir, C.S., Whelan, S.L., Gao, Y.-T., Ferlay, J., Powell, J., eds (1992). Cancer Incidence in Five Continents, Vol. VI. IARC Scientific Publications No. 120, Lyon, IARC.

<sup>&</sup>lt;sup>7</sup> Bray F, Colombet M, Mery L, Piñeros M, Znaor A, Zanetti R and Ferlay J, editors (2017). Cancer Incidence in Five Continents, Vol. XI (electronic version). Lyon: International Agency for Research on Cancer. Available from:  $\verb|http://ci5.iarc.fr|, accessed [05 October 2018].$ 

Figure 8: Age-specific incidence rates of cervical cancer in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



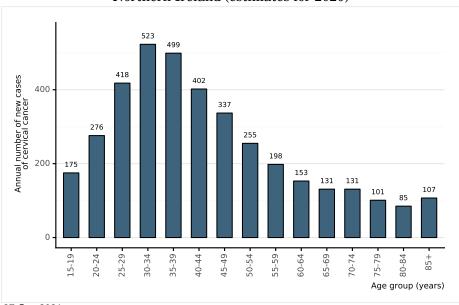
#### Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

Rates per 100,000 women per year.

Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

Figure 9: Annual number of new cases of cervical cancer in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



### Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

- · For age-standardised incidence rates of cervical cancer of United Kingdom of Great Britain and Northern Ireland (estimates for 2020) please refer to Figure 73
- · For annual number of new cases of cervical cancer by age group in United Kingdom of Great Britain and Northern Ireland (estimates for 2020) please refer to Figure 74
- For comparison of age-specific cervical cancer incidence rates in United Kingdom of Great Britain and Northern Ireland, within the region, and the rest of world please refer to Figure 75

## 3.3.2 Cervical cancer incidence by histology in United Kingdom of Great Britain and Northern Ireland

Table 4: Age-standardised incidence rates of cervical cancer in United Kingdom of Great Britain and Northern Ireland by histological type and cancer registry

Cancer registry <sup>1</sup>	Period	Squamo	Adeno	Other	Unspec.
National	2008-2012	5.2	1.5	0.4	0.1
England	2008-2012	5.3	1.5	0.4	0.1
England, East	2008-2012	4.6	1.5	0.3	0.1
England, East Midlands	2008-2012	6	2.2	0.4	0.1
England, London	2008-2012	3.7	1.1	0.2	0.1
England, North East	2008-2012	8	1.6	0.6	0.1
England, North West	2008-2012	6	1.5	0.4	0.1
England, South West	2008-2012	6.2	1.8	0.4	0.1
England, South East	2008-2012	4.3	1.4	0.3	0.1
England, West Midlands	2008-2012	5.8	1.7	0.6	0.2
England, Yorkshire and The Humber	2008-2012	6.9	1.5	0.4	0.2
Northern Ireland	2008-2012	7.2	1.8	0.4	0.2
Scotland	2008-2012	6.7	1.8	0.4	0.1
Wales	2008-2012	-	-	-	-

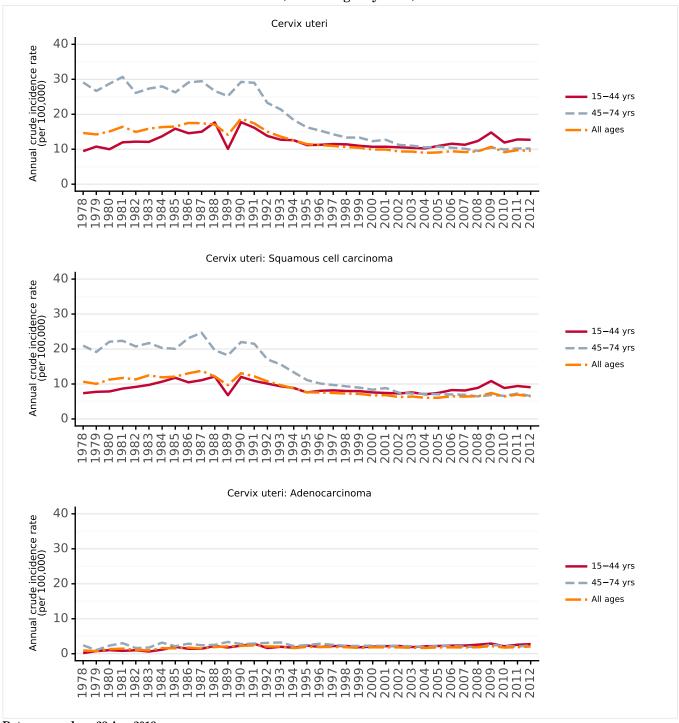
#### Data accessed on 5 Oct 2018

Rates per 100,000 women per year.
Standarized rates have been estimated using the direct method and the World population as the references.
Adeno: adenocarcinoma; Other: Other carcinoma; Squamous: Squamous cell carcinoma; Unspec: Unspecified carcinoma;

Data Sources:

1 Bray F, Colombet M, Mery L, Piñeros M, Znaor A, Zanetti R and Ferlay J, editors (2017). Cancer Incidence in Five Continents, Vol. XI (electronic version). Lyon: International Agency for Research on Cancer. Available from: http://ci5.iarc.fr, accessed [05 October 2018].

Figure 10: Time trends in cervical cancer incidence in United Kingdom of Great Britain and Northern Ireland (cancer registry data)



Data accessed on 28 Aug 2018

The following regional cancer registries provided data and contributed to their national estimate: Scotland, Northern Ireland, England: East, North East, North West, South East, South West, Yorkshire and The Humber, East Midlands, West Midlands, London

Estimated annual percentage change based on the trend variable from the net drift for 25 years, from 1983-2007.

Data Sources:
Ferlay J, Colombet M and Bray F. Cancer Incidence in Five Continents, CI5plus: IARC CancerBase No. 9 [Internet]. Lyon, France: International Agency for Research on Cancer; 2018. Available from: http://ci5.iarc.fr
Vaccarella S, Lortet-Tieulent J, Plummer M, Franceschi S, Bray F. Worldwide trends in cervical cancer incidence: Impact of screening against changes in disease risk factors. eur J Cancer 2013;49:3262-73.

## 3.3.3 Cervical cancer mortality in United Kingdom of Great Britain and Northern Ireland

### **Key Stats.**

About 1,121 cervical cancer deaths occur annually in United Kingdom of Great Britain and Northern Ireland are diagnosed annually (estimations for 2020).

Cervical cancer ranks\* as the 16th leading cause of cancer deaths of female cancer deaths in United Kingdom of Great Britain and Northern Ireland.

Cervical cancer is the 3<sup>rd</sup> leading cause of cancer deaths in women aged 15 to 44 years in United Kingdom of Great Britain and Northern Ireland.

Table 5: Cervical cancer mortality in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

Indicator	United Kingdom of Great Britain and Northern Ireland	Northern Europe	World
Annual number of deaths	1,121	2,134	341,831
Uncertainty intervals of mortality cancer cases [95% UI]	[1,036-1,213]	[1,984-2,296]	[324,231-360,386]
Crude mortality rate <sup>b</sup>	3.26	3.97	8.84
Age-standardized mortality rate <sup>b</sup>	1.92	2.18	7.25
Cumulative risk (%) at 75 years old <sup>a</sup>	0.19	0.22	0.82

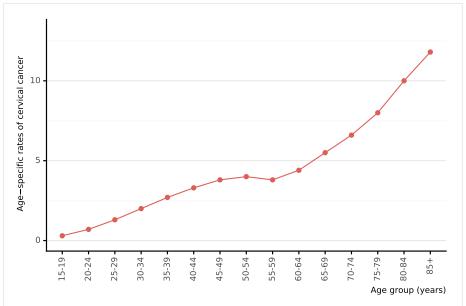
#### Data accessed on 27 Jan 2021

<sup>\*</sup> Ranking of cervical cancer incidence to other cancers among all women according to highest incidence rates (ranking 1st) excluding non-melanoma skin cancer. Ranking is based on crude incidence rates (actual number of cervical cancer cases). Ranking using age-standardized rate (ASR) may differ

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a Cumulative risk (mortality) is the probability or risk of individuals dying from the disease during ages 0-74 years. For cancer, it is expressed as the % of new born children who would be expected to die from a particular cancer before the age of 75 if they had the rates of cancer observed in the period in the absence of competing causes b Rates per 100,000 women per year.

Figure 11: Age-specific mortality rates of cervical cancer in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

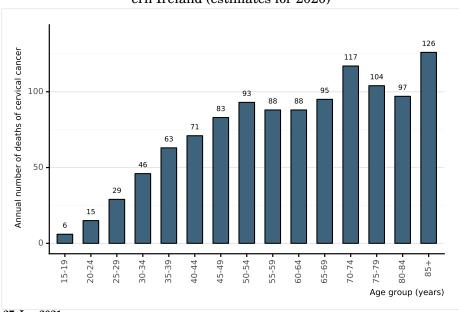


Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods a Rates per 100,000 women per year.

Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

Figure 12: Annual number of deaths of cervical cancer in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



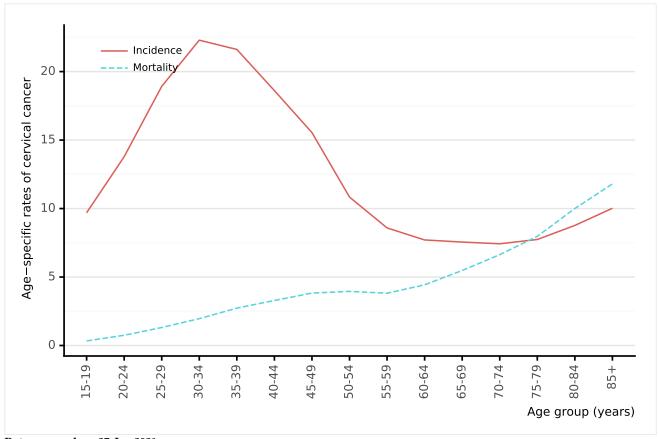
Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods Data Sources:

- · For age-standardised mortality rates of cervical cancer of United Kingdom of Great Britain and Northern Ireland (estimates for 2020) please refer to Figure 105
- For annual number of deaths of cervical cancer by age group in United Kingdom of Great Britain and Northern Ireland (estimates for 2020) please refer to Figure 106
- For comparison of age-specific cervical cancer mortality rates in United Kingdom of Great Britain and Northern Ireland, within the region, and the rest of world please refer to Figure 107

## 3.3.4 Cervical cancer incidence and mortality comparison in United Kingdom of Great Britain and Northern Ireland

Figure 13: Comparison of age-specific cervical cancer incidence and mortality rates in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to  $\frac{\text{http://gco.iarc.fr/today/data-sources-methods}}{a} \text{ Rates per } 100,000 \text{ women per year.}$ 

Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

Table 6: Premature deaths and disability from cervical cancer in United Kingdom of Great Britain and Northern Ireland, Europe and the rest of the world (estimates for 2019)

	United Kingdom	of Great Br	itain and Northern Ireland	Europe	е	World
Indicator	Number		Rate	Number	Rate	Number
DALYs (95% UI) <sup>a</sup>	43,304		128	824,336	189	8,955,013
DALIS (95% UI)	(40,596-55,323)		(120-163)	(726, 198-913, 992)	(166-209)	(7,547,733-9,978,462)
YLLs (95% UI) <sup>b</sup>	41,651		123	793,756	182	8,712,962
ILLS (95% UI)	(39,197-53,322)		(115-157)	(703,004-877,841)	(161-201)	(7,365,279-9,728,886)
YLDs (95% UI) <sup>c</sup> (	1,653		5 (3-7)	30,580	7 (5-10)	242,051
	(1,059-2,429)			(21,266-42,064)		(171,644-326,024)

#### Data accessed on 29 Apr 2021

Rate per 100,000 women

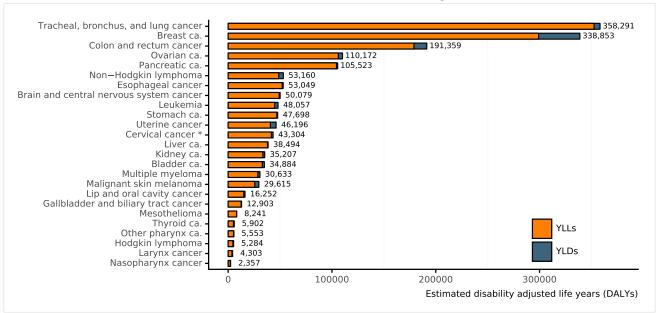
<sup>a</sup> DALYs (95% UI): estimated disability adjusted life years (95% uncertainty interval)

 $^b$  YLLs (95% UI): years of life lost (95% uncertainty interval)

 $^c$  YLDs (95% UI): estimated years lived with disability (95% uncertainty interval)

GBD 2019 Diseases and Injuries Collaborators. Global burden of 369 diseases and injuries in 204 countries and territories, 1990-2019: a systematic analysis for the Global Burden of Disease Study 2019. Lancet. 2020 Oct 17;396(10258):1204-1222

Figure 14: Comparison of annual premature deaths and disability from cervical cancer in United Kingdom of Great Britain and Northern Ireland to other cancers among women (estimates for 2019)



Data accessed on 29 Apr 2021

YLLs: years of life lost YLDs: years lived with disability

<u>Data Sources:</u>
GBD 2019 Diseases and Injuries Collaborators. Global burden of 369 diseases and injuries in 204 countries and territories, 1990-2019: a systematic analysis for the Global Burden of Disease Study 2019. Lancet. 2020 Oct 17;396(10258):1204-1222

### Anogenital cancers other than the cervix

Data on HPV role in anogenital cancers other than cervix are limited, but there is an increasing body of evidence strongly linking HPV DNA with cancers of anus, vulva, vagina, and penis. Although these cancers are much less frequent compared to cervical cancer, their association with HPV make them potentially preventable and subject to similar preventative strategies as those for cervical cancer. (Vaccine 2006, Vol. 24, Suppl 3; Vaccine 2008, Vol. 26, Suppl 10; Vaccine 2012, Vol. 30, Suppl 5; IARC Monographs 2007, Vol. 90).

#### 3.4.1 Anal cancer

Anal cancer is rare in the general population with an average worldwide incidence of 1 per 100,000, but is reported to be increasing in more developed regions. Globally, there are an estimated 29,000 new cases in 2018 every year (de Martel C et al. Lancet Glob Health 2020;8(2):e180-e190). Women have higher incidences of anal cancer than men. Incidence is particularly high among populations of men who have sex with men (MSM), women with history of cervical or vulvar cancer, and immunosuppressed populations, including those who are HIV-infected and patients with a history of organ transplantation. These cancers are predominantly squamous cell carcinoma, adenocarcinomas, or basaloid and cloacogenic carcinomas.

### 3.4.1.1 Anal cancer incidence in United Kingdom of Great Britain and Northern Ireland

Table 7: Anal cancer incidence in United Kingdom of Great Britain and Northern Ireland (estimates for

Indicator	United Kingdom of Great Britain and Northern Ireland	Northern Europe	World
MEN			
Annual number of new cancer cases	544	750	21,706
Uncertainty intervals of new cancer cases [95% UI]	[434-682]	[676-832]	[18,432-25,561]
Crude incidence rate <sup>b</sup>	1.62	1.43	0.55
Age-standardized incidence rate <sup>b</sup>	0.93	0.79	0.49
Cumulative risk (%) at 75 years old <sup>a</sup>	0.11	0.09	0.06
WOMEN			
Annual number of new cancer cases	1,141	1,557	29,159
Uncertainty intervals of new cancer cases [95% UI]	[1,048-1,242]	[1,450-1,672]	[25,656-33,140]
Crude incidence rate <sup>c</sup>	3.32	2.90	0.75
Age-standardized incidence rate <sup>c</sup>	1.85	1.57	0.58
Cumulative risk (%) at 75 years old <sup>a</sup>	0.22	0.19	0.07

#### Data accessed on 27 Jan 2021

Data Sources:

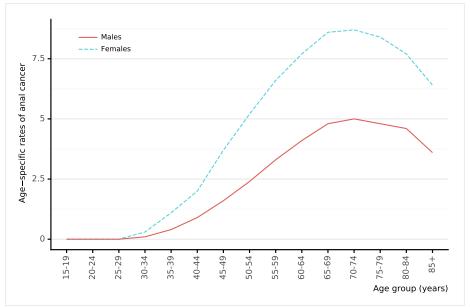
For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a Cumulative risk (incidence) is the probability or risk of individuals getting from the disease during ages 0-74 years. For cancer, it is expressed as the % of new born children who would be expected to develop from a particular cancer before the age of 75 if they had the rates of cancer observed in the period in the absence of competing causes

Rates per 100,000 men per year.

<sup>&</sup>lt;sup>c</sup> Rates per 100,000 women per year.

Figure 15: Age-specific incidence rates of anal cancer in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



#### Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

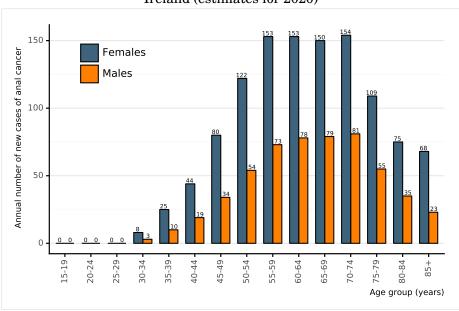
a Rates per 100,000 men per year.

 $b\,$  Rates per 100,000 women per year.

Data Sources:

Fernay J. Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

Figure 16: Annual number of new cases of anal cancer in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

## 3.4.1.2 Anal cancer mortality in United Kingdom of Great Britain and Northern Ireland

Table 8: Anal cancer mortality in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

Indicator	United Kingdom of Great Britain and Northern Ireland	Northern Europe	World
MEN			
Annual number of new cancer cases	231	301	9,416
Uncertainty intervals of new cancer cases [95% UI]	[174-308]	[255-356]	[7,282-12,175]
Crude incidence rate <sup>b</sup>	0.69	0.57	0.24
Age-standardized incidence rate <sup>b</sup>	0.34	0.27	0.21
Cumulative risk (%) at 75 years old <sup>a</sup>	0.04	0.03	0.02
WOMEN			
Annual number of new cancer cases	344	452	9,877
Uncertainty intervals of new cancer cases [95% UI]	[289-409]	[395-518]	[7,795-12,516]
Crude incidence rate <sup>c</sup>	1.00	0.84	0.26
Age-standardized incidence rate <sup>c</sup>	0.41	0.33	0.19
Cumulative risk (%) at 75 years old <sup>a</sup>	0.04	0.04	0.02

Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a Cumulative risk (mortality) is the probability or risk of individuals dying from the disease during ages 0-74 years. For cancer, it is expressed as the % of new born children who would be expected to die from a particular cancer before the age of 75 if they had the rates of cancer observed in the period in the absence of competing causes.

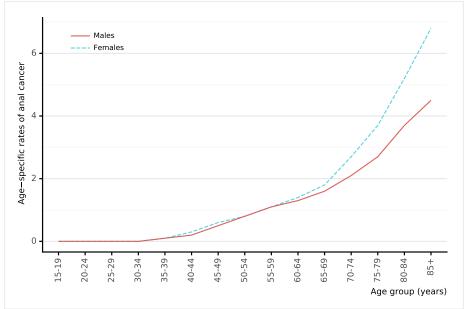
b Rates per 100,000 men per year.

c Rates per 100,000 women per year.

Data Sources:

Ferlay J. Farrik M. Lam F. Colombet M. Mery L. Piñerce M. Znear A. Socrience and L. Parrik M. Lam F. Colombet M. Mery L. Piñerce M. Znear A. Socrience and L. Parrik M. Lam F. Colombet M. Mery L. Piñerce M. Znear A. Socrience and L. Parrik M. Lam F. Colombet M. Mery L. Piñerce M. Znear A. Socrience and L. Parrik M. Lam F. Colombet M. Mery L. Piñerce M. Znear A. Socrience and L. Parrik M. Lam F. Colombet M. Mery L. Piñerce M. Znear A. Socrience and L. Parrik M. Lam F. Colombet M. Mery L. Piñerce M. Znear A. Socrience and L. Parrik M. Lam F. Colombet M. Mery L. Piñerce M. Znear A. Socrience and L. Parrik M. Lam F. Colombet M. Mery L. Piñerce M. Znear A. Socrience and L. Parrik M. Lam F. Colombet M. Mery L. Piñerce M. Znear A. Socrience and L. Parrik M. Lam F. Colombet M. Mery L. Piñerce M. Znear A. Socrience and L. Parrik M. Lam F. Colombet M. Mery L. Piñerce M. Znear A. Socrience and L. Parrik M. Lam F. Colombet M. Mery L. Piñerce M. Znear A. Socrience and L. Parrik M. Lam F. Colombet M. Mery L. Piñerce M. Znear A. Socrience and L. Parrik M. Lam F. Colombet M. Mery L. Piñerce M. Znear A. Socrience and L. Parrik M. Lam F. Colombet M. Mery L. Piñerce M. Znear A. Socrience and L. Parrik M. Lam F. Colombet M. Mery L. Piñerce M. Znear A. Socrience and L. Parrik M. Lam F. Colombet M. Mery L. Piñerce M. Znear A. Socrience and L. Parrik M. Lam F. Colombet M. Mery L. Piñerce M. Znear A. Socrience and L. Parrik M. Lam F. Colombet M. Lam F. Colombet M. Lam F. Colombet M.

Figure 17: Age-specific mortality rates of anal cancer in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

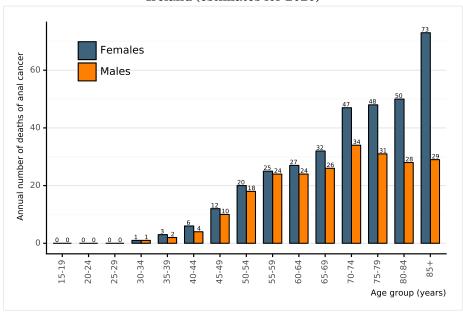


Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods a Rates per 100,000 men per year.

Data Sources:
Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

Figure 18: Annual number of deaths of of anal cancer in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



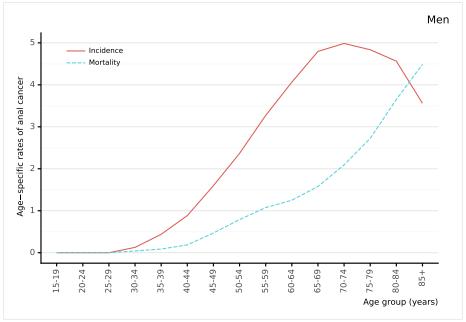
Data accessed on 27 Jan 2021

 $For more \ detailed \ methods \ of \ estimation \ please \ refer \ to \ http://gco.iarc.fr/today/data-sources-methods$ 

b Rates per 100,000 women per year

## 3.4.1.3 Anal cancer incidence and mortality comparison in United Kingdom of Great Britain and Northern Ireland

Figure 19: Comparison of age-specific anal cancer incidence and mortality rates among men in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

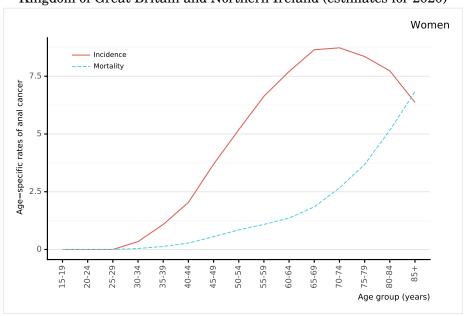


Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods a Rates per 100,000 men per year.

Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

Figure 20: Comparison of age-specific anal cancer incidence and mortality rates among women in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods <sup>a</sup> Rates per 100,000 women per year.

#### 3.4.2 Vulva cancer

Cancer of the vulva is rare among women worldwide, with an estimated 44,000 new cases in 2018, representing 6% of all gynaecologic cancers (de Martel C et al. Lancet Glob Health 2020;8(2):e180e190). Worldwide, about 60% of all vulvar cancer cases occur in more developed countries. Vulvar cancer has two distinct histological patterns with two different risk factor profiles: (1) basaloid/warty types (2) keratinising types. Basaloid/warty lesions are more common in young women, are very often associated with HPV DNA detection (75-100%), and have a similar risk factor profile as cervical cancer. Keratinising vulvar carcinomas represent the majority of the vulvar lesions (>60%), they occur more often in older women and are more rarely associated with HPV (IARC Monograph Vol 100B).

#### 3.4.2.1 Vulva cancer incidence in United Kingdom of Great Britain and Northern Ireland

Table 9: Vulva cancer incidence in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

Indicator	United Kingdom of Great Britain and Northern Ireland	Northern Europe	World
Annual number of new cancer cases	1,521	2,227	45,240
Uncertainty intervals [95% UI]	[1,399-1,653]	[2,092-2,371]	[40,656-50,342]
Crude incidence rate <sup>b</sup>	4.43	4.14	1.17
Age-standardized incidence rate <sup>b</sup>	2.09	1.85	0.85
Cumulative risk (%) at 75 years old <sup>a</sup>	0.23	0.20	0.09

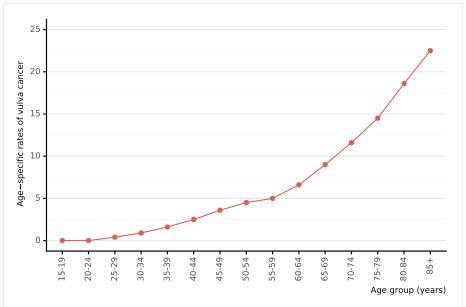
#### Data accessed on 27 Jan 2021

 $\stackrel{.}{b}$  Rates per 100,000 women per year.

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a Cumulative risk (incidence) is the probability or risk of individuals getting from the disease during ages 0-74 years. For cancer, it is expressed as the % of new born children who would be expected to develop from a particular cancer before the age of 75 if they had the rates of cancer observed in the period in the absence of competing causes.

Figure 21: Age-specific incidence rates of vulva cancer in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

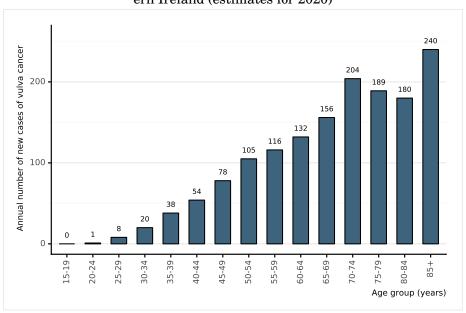


 $For more \ detailed \ methods \ of \ estimation \ please \ refer \ to \ http://gco.iarc.fr/today/data-sources-methods \ detailed \ methods \ of \ estimation \ please \ refer \ to \ http://gco.iarc.fr/today/data-sources-methods \ detailed \ methods \ of \ estimation \ please \ refer \ to \ http://gco.iarc.fr/today/data-sources-methods \ detailed \ methods \ of \ estimation \ please \ refer \ to \ http://gco.iarc.fr/today/data-sources-methods \ detailed \ methods \ of \ estimation \ please \ refer \ to \ http://gco.iarc.fr/today/data-sources-methods \ detailed \ methods \ of \ estimation \ please \ refer \ to \ http://gco.iarc.fr/today/data-sources-methods \ detailed \ detailled \ detailed \ detailed \ detailed \ detailed \ detailed \ det$ 

a Rates per 100,000 women per year.

Data Sources:
Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

Figure 22: Annual number of new cases of vulva cancer in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

Bata Doutes.
Ferlay J. Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

## 3.4.2.2 Vulva cancer mortality in United Kingdom of Great Britain and Northern Ireland

Table 10: Vulva cancer mortality in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

Indicator	United Kingdom of Great Britain and Northern Ireland	Northern Europe	World
Annual number of deaths	526	795	17,427
Uncertainty intervals [95% UI]	[456-607]	[713-886]	[14,497-20,950]
Crude mortality rate <sup>b</sup>	1.53	1.48	0.45
Age-standardized mortality rate <sup>b</sup>	0.49	0.45	0.30
Cumulative risk (%) at 75 years old <sup>a</sup>	0.05	0.04	0.03

Data accessed on 27 Jan 2021

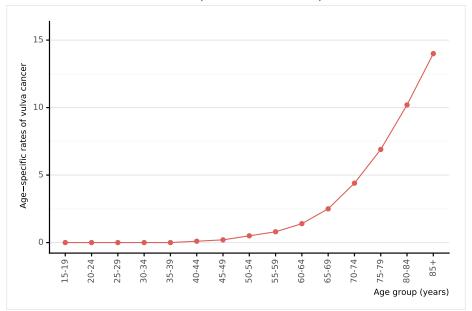
For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

Cumulative risk (mortality) is the probability or risk of individuals dying from the disease during ages 0-74 years. For cancer, it is expressed as the % of new born children who would be expected to die from a particular cancer before the age of 75 if they had the rates of cancer observed in the period in the absence of competing causes.

Bates per 100,000 women per year.

Data Sources:
Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for  $Research \ on \ Cancer. \ Available \ from: \ \verb|https://gco.iarc.fr/today|, \ accessed \ [27 \ January \ 2021].$ 

Figure 23: Age-specific mortality rates of vulva cancer in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

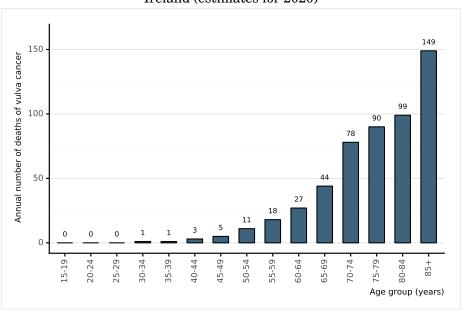


 $For more \ detailed \ methods \ of \ estimation \ please \ refer \ to \ http://gco.iarc.fr/today/data-sources-methods \ detailed \ methods \ of \ estimation \ please \ refer \ to \ http://gco.iarc.fr/today/data-sources-methods \ detailed \ methods \ of \ estimation \ please \ refer \ to \ http://gco.iarc.fr/today/data-sources-methods \ detailed \ methods \ of \ estimation \ please \ refer \ to \ http://gco.iarc.fr/today/data-sources-methods \ detailed \ methods \ of \ estimation \ please \ refer \ to \ http://gco.iarc.fr/today/data-sources-methods \ detailed \ methods \ of \ estimation \ please \ refer \ to \ http://gco.iarc.fr/today/data-sources-methods \ detailed \ detailled \ detailed \ detailed \ detailed \ detailed \ detailed \ det$ 

 $^a$  Rates per 100,000 women per year.

Data Sources:
Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for  $Research \ on \ Cancer. \ Available \ from: \ \verb|https://gco.iarc.fr/today|, \ accessed \ [27 \ January \ 2021].$ 

Figure 24: Annual number of deaths of vulva cancer in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



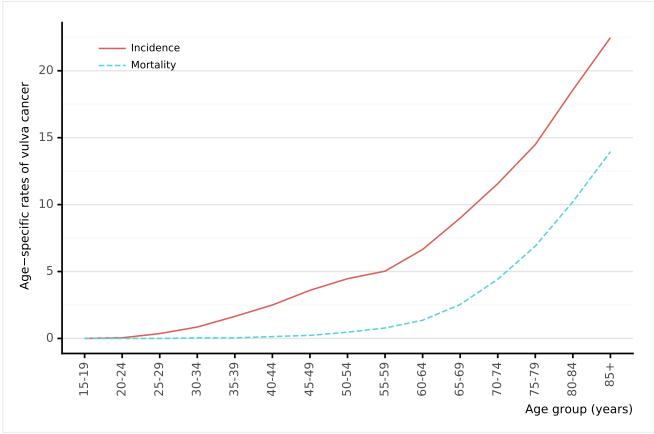
Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods a Rates per 100,000 women per year.

Data Sources:
Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

## 3.4.2.3 Vulva cancer incidence and mortality comparison in United Kingdom of Great Britain and Northern Ireland

Figure 25: Comparison of age-specific vulva cancer incidence and mortality rates in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods a Rates per 100,000 women per year.

Data Sources:

Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

#### 3.4.3 Vaginal cancer

Cancer of the vagina is a rare cancer, with an estimated 18,000 new cases in 2018, representing 3% of all gynaecologic cancers (de Martel C et al. Lancet Glob Health 2020;8(2):e180-e190). Similar to cervical cancer, the majority of vaginal cancer cases (68%) occur in less developed countries. Most vaginal cancers are squamous cell carcinoma (90%) generally attributable to HPV, followed by clear cell adenocarcinomas and melanoma. Vaginal cancers are primarily reported in developed countries. Metastatic cervical cancer can be misclassified as cancer of the vagina. Invasive vaginal cancer is diagnosed primarily in old women (>= 65 years) and the diagnosis is rare in women under 45 years whereas the peak incidence of carcinoma in situ is observed between ages 55 and 70 (Vaccine 2008, Vol. 26, Suppl 10).

#### 3.4.3.1 Vaginal cancer incidence in United Kingdom of Great Britain and Northern Ireland

Table 11: Vaginal cancer incidence in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

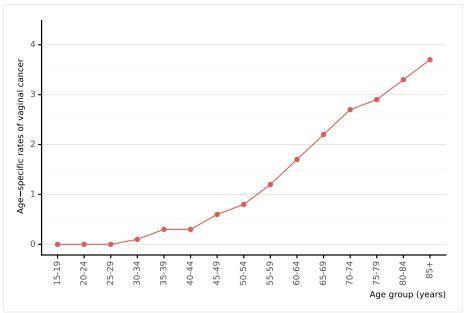
Indicator	United Kingdom of Great Britain and Northern Ireland	Northern Europe	World
Annual number of new cancer cases	306	450	17,908
Uncertainty intervals [95% UI]	[241-388]	[390-519]	[14,678-21,848]
Crude incidence rate <sup>b</sup>	0.89	0.84	0.46
Age-standardized incidence rate <sup>b</sup>	0.43	0.38	0.36
Cumulative risk (%) at 75 years old <sup>a</sup>	0.05	0.04	0.04

#### Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to <a href="http://gco.iarc.fr/today/data-sources-methods">http://gco.iarc.fr/today/data-sources-methods</a>
<sup>a</sup> Cumulative risk (incidence) is the probability or risk of individuals getting from the disease during ages 0.74 years. For cancer, it is expressed as the % of new born children who would be expected to develop from a particular cancer before the age of 75 if they had the rates of cancer observed in the period in the absence of competing causes.

Data Sources:
Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

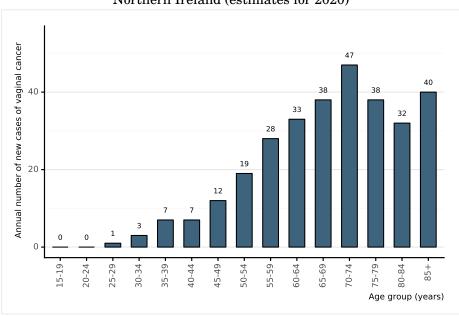
Figure 26: Age-specific incidence rates of vaginal cancer in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods a Rates per 100,000 women per year.

Data Sources:
Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for  $Research \ on \ Cancer. \ Available \ from: \ \texttt{https://gco.iarc.fr/today}\ ,\ accessed\ [27\ January\ 2021].$ 

Figure 27: Annual number of new cases of vaginal cancer in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

 $For more \ detailed \ methods \ of \ estimation \ please \ refer \ to \ http://gco.iarc.fr/today/data-sources-methods$ 

Data Sources:

Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

## 3.4.3.2 Vaginal cancer mortality in United Kingdom of Great Britain and Northern Ireland

Table 12: Vaginal cancer mortality in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

Indicator	United Kingdom of Great Britain and Northern Ireland	Northern Europe	World
Annual number of deaths	125	185	7,995
Uncertainty intervals [95% UI]	[93-168]	[148-232]	[5,983-10,684]
Crude mortality rate <sup>b</sup>	0.36	0.34	0.21
Age-standardized mortality rate <sup>b</sup>	0.15	0.13	0.16
Cumulative risk (%) at 75 years old <sup>a</sup>	0.02	0.01	0.02

Data Sources:
Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

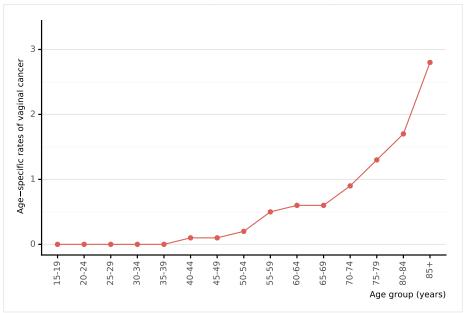
Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a Cumulative risk (mortality) is the probability or risk of individuals dying from the disease during ages 0-74 years. For cancer, it is expressed as the % of new born children who would be expected to die from a particular cancer before the age of 75 if they had the rates of cancer observed in the period in the absence of competing causes.

b Rates per 100,000 women per year.

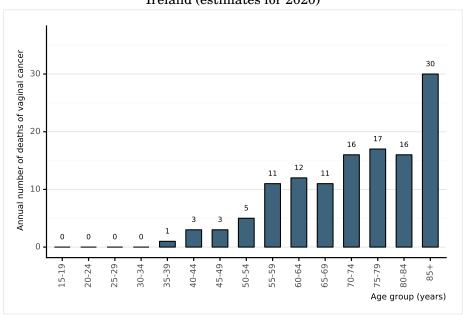
Figure 28: Age-specific mortality rates of vaginal cancer in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods a Rates per 100,000 women per year.

Data Sources:
Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for  $Research \ on \ Cancer. \ Available \ from: \ \texttt{https://gco.iarc.fr/today}\ ,\ accessed\ [27\ January\ 2021].$ 

Figure 29: Annual number of deaths of vaginal cancer in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

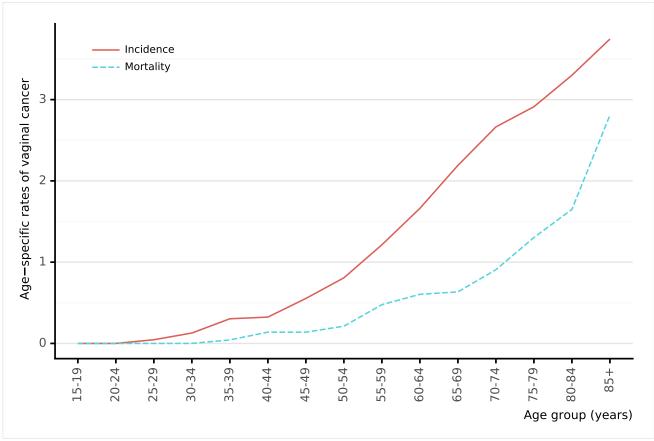
 $For more \ detailed \ methods \ of \ estimation \ please \ refer \ to \ http://gco.iarc.fr/today/data-sources-methods$ 

Data Sources:

Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

#### 3.4.3.3 Vaginal cancer incidence and mortality comparison in United Kingdom of Great **Britain and Northern Ireland**

Figure 30: Comparison of age-specific vaginal cancer incidence and mortality rates in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods a Rates per 100,000 women per year.

Bata Sources. Ferlay J. Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

#### 3.4.4 Penile cancer

The annual burden of penile cancer has been estimated to be 34,000 cases in 2018 worldwide with incidence rates strongly correlating with those of cervical cancer (de Martel C et al. Lancet Glob Health 2020;8(2):e180-e190). Penile cancer is rare and most commonly affects men aged 50-70 years. Incidence rates are higher in less developed countries than in more developed countries, accounting for up to 10% of male cancers in some parts of Africa, South America and Asia. Precursor cancerous penile lesions (PeIN) are rare.

Cancers of the penis are primarily of squamous cell carcinomas (SCC) (95%) and the most common penile SCC histologic sub-types are keratinising (49%), mixed warty-basaloid (17%), verrucous (8%) warty (6%), and basaloid (4%). HPV is most commonly detected in basaloid and warty tumours but is less common in keratinising and verrucous tumours. Approximately 60-100% of PeIN lesions are HPV DNA positive.

## 3.4.4.1 Penile cancer incidence in United Kingdom of Great Britain and Northern Ireland

Table 13: Penile cancer incidence in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

Indicator	United Kingdom of Great Britain and Northern Ireland	Northern Europe	World
Annual number of new cancer cases	763	1,155	36,068
Uncertainty intervals [95% UI]	[600-970]	[1,056-1,264]	[30,963-42,015]
Crude incidence rate <sup>b</sup>	2.27	2.20	0.92
Age-standardized incidence rate <sup>b</sup>	1.18	1.11	0.80
Cumulative risk (%) at 75 years old <sup>a</sup>	0.14	0.13	0.09

#### Data accessed on 27 Jan 2021

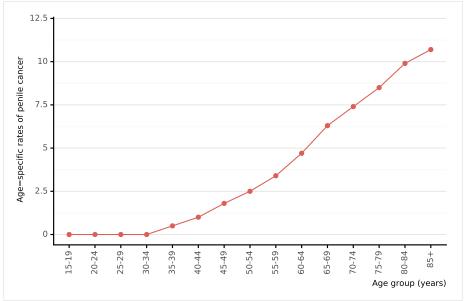
For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a Cumulative risk (incidence) is the probability or risk of individuals getting from the disease during ages 0-74 years. For cancer, it is expressed as the % of new born children who would be expected to develop from a particular cancer before the age of 75 if they had the rates of cancer observed in the period in the absence of competing cause

 $<sup>^{\</sup>it b}$ Rates per 100,000 men per year.

Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

Figure 31: Age-specific incidence rates of penile cancer in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



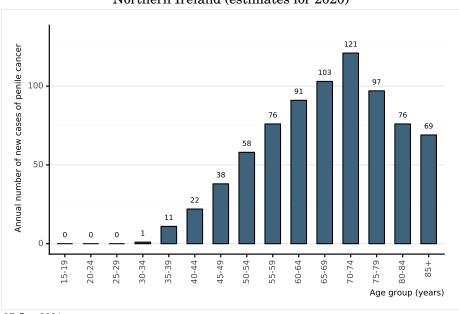
Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a Rates per 100,000 men per year.

Data Sources:
Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

Figure 32: Annual number of new cases of penile cancer in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

 $For more \ detailed \ methods \ of \ estimation \ please \ refer \ to \ http://gco.iarc.fr/today/data-sources-methods$ 

Feriay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

#### 3.4.4.2 Penile cancer mortality in United Kingdom of Great Britain and Northern Ireland

Table 14: Penile cancer mortality in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

Indicator	United Kingdom of Great Britain and Northern Ireland	Northern Europe	World
Annual number of deaths	150	252	13,211
Uncertainty intervals [95% UI]	[118-192]	[204-311]	[10,687-16,332]
Crude mortality rate <sup>b</sup>	0.45	0.48	0.34
Age-standardized mortality rate <sup>b</sup>	0.19	0.21	0.29
Cumulative risk (%) at 75 years old <sup>a</sup>	0.02	0.02	0.03

Data accessed on 27 Jan 2021

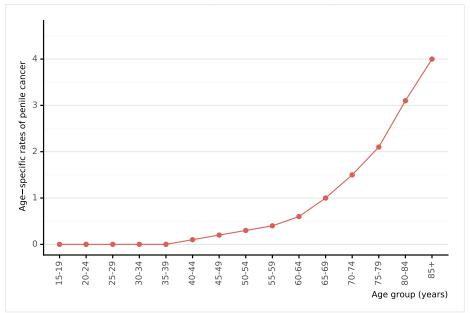
For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

<sup>a</sup> Cumulative risk (mortality) is the probability or risk of individuals dying from the disease during ages 0-74 years. For cancer, it is expressed as the % of new born children who would be expected to die from a particular cancer before the age of 75 if they had the rates of cancer observed in the period in the absence of competing causes.

<sup>b</sup> Rates per 100,000 men per year.

Data Sources:
Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for  $Research \ on \ Cancer. \ Available \ from: \ \verb|https://gco.iarc.fr/today|, \ accessed \ [27 \ January \ 2021].$ 

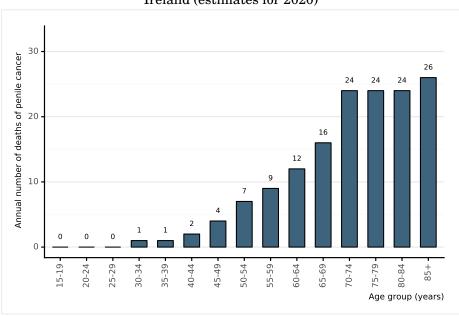
Figure 33: Age-specific mortality rates of penile cancer in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods a Rates per 100,000 men per year.

Data Sources:
Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for  $Research \ on \ Cancer. \ Available \ from: \ \texttt{https://gco.iarc.fr/today}\ ,\ accessed\ [27\ January\ 2021].$ 

Figure 34: Annual number of deaths of penile cancer in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



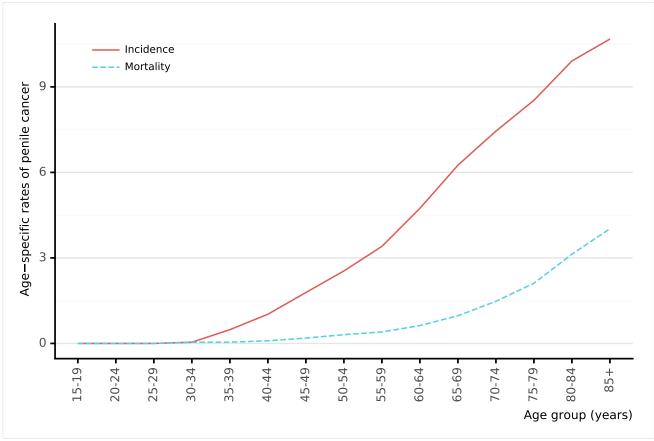
Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods Data Sources:

Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

## 3.4.4.3 Penile cancer incidence and mortality comparison in United Kingdom of Great Britain and Northern Ireland

Figure 35: Comparison of age-specific penile cancer incidence and mortality rates in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods Rates per 100,000 men per year.

Data Sources:
Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

#### 3.5 Head and neck cancers

The majority of head and neck cancers are associated with high tobacco and alcohol consumption. However, increasing trends in the incidence at specific sites suggest that other aetiological factors are involved, and infection by certain high-risk types of HPV (i.e. HPV16) have been reported to be associated with head and neck cancers, in particular with oropharyngeal cancer. Current evidence suggests that HPV16 is associated with tonsil cancer (including Waldeyer ring cancer), base of tongue cancer and other oropharyngeal cancer sites. Associations with other head and neck cancer sites such as oral cancer are neither strong nor consistent when compared to molecular-epidemiological data on HPV and oropharyngeal cancer. Association with laryngeal cancer is still unclear (IARC Monograph Vol 100B)

#### 3.5.1 Oropharyngeal cancer

#### 3.5.1.1 Oropharyngeal cancer incidence in United Kingdom of Great Britain and Northern **Ireland**

Table 15: Oropharyngeal cancer incidence in United Kingdom of Great Britain and Northern Ireland

Indicator	United Kingdom of Great Britain and Northern Ireland	Northern Europe	World
MEN			
Annual number of new cancer cases	2,110	3,342	79,045
Uncertainty intervals of new cancer cases [95% UI]	[1,880-2,369]	[3,163-3,531]	[72,769-85,862]
Crude incidence rate sa <sup>b</sup>	6.29	6.37	2.01
Age-standardized incidence rate sa <sup>b</sup>	4.08	4.00	1.79
Cumulative risk (%) at 75 years old <sup>a</sup>	0.48	0.48	0.22
WOMEN			
Annual number of new cancer cases	700	1,090	19,367
Uncertainty intervals of new cancer cases [95% UI]	[545-899]	[990-1,200]	[16,279-23,041]
Crude incidence rate sa <sup>c</sup>	2.04	2.03	0.50
Age-standardized incidence rate sa <sup>c</sup>	1.25	1.21	0.40
Cumulative risk (%) at 75 years old <sup>a</sup>	0.15	0.15	0.05

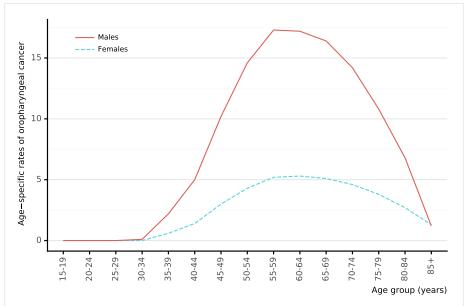
#### Data accessed on 27 Jan 2021

Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods  $^a$  Cumulative risk (incidence) is the probability or risk of individuals getting from the disease during ages 0-74 years. For cancer, it is expressed as the % of new born children who would be expected to develop from a particular cancer before the age of 75 if they had the rates of cancer observed in the period in the absence of competing causes.

Rates per 100,000 men per year. <sup>c</sup> Rates per 100,000 women per year.

Figure 36: Age-specific incidence rates of oropharyngeal cancer in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

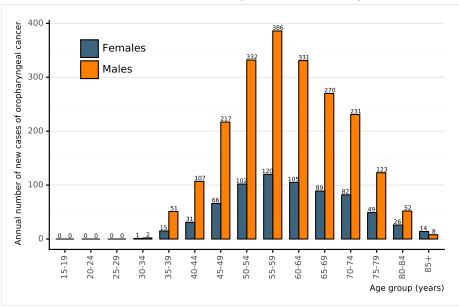


For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

 $^a$  Rates per 100,000 men per year.

Data Sources:
Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

Figure 37: Annual number of new cases of oropharyngeal cancer in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

 $For more \ detailed \ methods \ of \ estimation \ please \ refer \ to \ http://gco.iarc.fr/today/data-sources-methods$ 

Data Sources

Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for  $Research \ on \ Cancer. \ Available \ from: \ \verb|https://gco.iarc.fr/today|, \ accessed \ [27 \ January \ 2021].$ 

b Rates per 100,000 women per year

## 3.5.1.2 Oropharyngeal cancer mortality in United Kingdom of Great Britain and Northern **Ireland**

Table 16: Oropharyngeal cancer mortality in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

Indicator	United Kingdom of Great Britain and Northern Ireland	Northern Europe	World
MEN			
Annual number of deaths	768	1,154	39,590
Uncertainty intervals of mortality cancer cases [95% UI]	[686-860]	[1,056-1,261]	[35,255-44,458]
Crude mortality rate sa <sup>b</sup>	2.29	2.20	1.01
Age-standardized mortality rate sa <sup>b</sup>	1.21	1.16	0.89
Cumulative risk (%) at 75 years old <sup>a</sup>	0.15	0.15	0.11
WOMEN			
Annual number of deaths	236	328	8,553
Uncertainty intervals of mortality cancer cases [95% UI]	[192-289]	[279-386]	[6,684-10,945]
Crude mortality rate sa <sup>c</sup>	0.69	0.61	0.22
Age-standardized mortality rate sa <sup>c</sup>	0.35	0.30	0.17
Cumulative risk (%) at 75 years old <sup>a</sup>	0.04	0.04	0.02

#### Data accessed on 27 Jan 2021

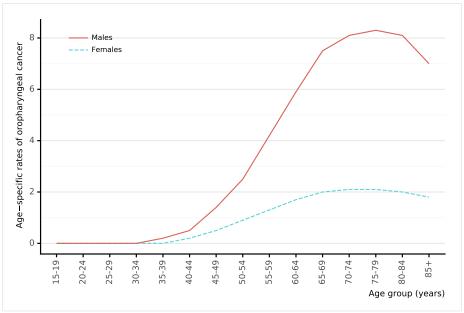
Data Sources:
Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for  $Research \ on \ Cancer. \ Available \ from: \ \texttt{https://gco.iarc.fr/today} \ , \ accessed \ [27 \ January \ 2021].$ 

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a Cumulative risk (mortality) is the probability or risk of individuals dying from the disease during ages 0-74 years. For cancer, it is expressed as the % of new born children who would be expected to die from a particular cancer before the age of 75 if they had the rates of cancer observed in the period in the absence of competing causes b Rates per 100,000 men per year.

c Rates per 100,000 women per year.

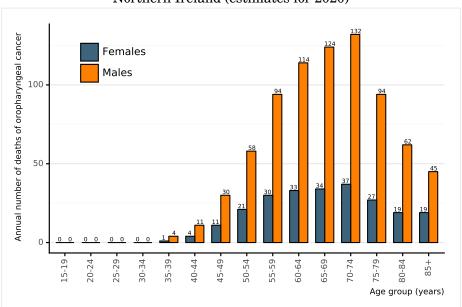
Figure 38: Age-specific mortality rates of oropharyngeal cancer in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



For more detailed methods of estimation please refer to  $\texttt{http://gco.iarc.fr/today/data-sources-methods}^a$  Rates per 100,000 men per year.

Data Sources:
Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

Figure 39: Annual number of deaths of oropharyngeal cancer in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



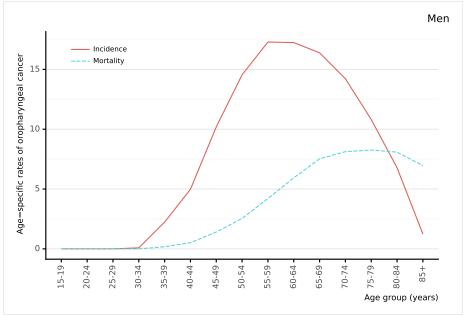
#### Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for  $Research \ on \ Cancer. \ Available \ from: \ \texttt{https://gco.iarc.fr/today}, \ accessed \ [27 \ January \ 2021].$ 

#### 3.5.1.3 Oropharyngeal cancer incidence and mortality comparison in United Kingdom of **Great Britain and Northern Ireland**

Figure 40: Comparison of age-specific oropharyngeal cancer incidence and mortality rates among men in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

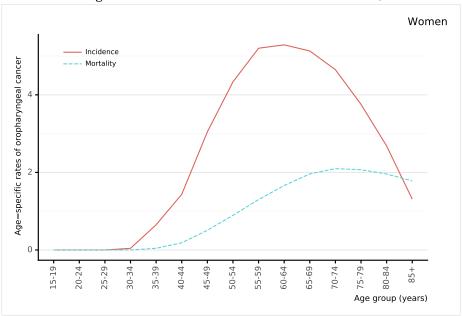


Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to  $\frac{1}{2} \frac{1}{2} \frac{1}$ 

Data Sources:
Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

Figure 41: Comparison of age-specific oropharyngeal cancer incidence and mortality rates among women in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

Rates per 100,000 women per year.

Ferlay J. Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

#### 3.5.2 Oral cavity cancer

#### 3.5.2.1 Oral cavity cancer incidence in United Kingdom of Great Britain and Northern Ireland

Table 17: Oral cavity cancer incidence in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

Indicator	United Kingdom of Great Britain and Northern Ireland	Northern Europe	World
MEN			
Annual number of new cancer cases	3,931	5,582	264,211
Uncertainty intervals of new cancer cases [95% UI]	[3,725-4,148]	[5,372-5,801]	[251,153- 277,948]
Crude incidence rate sa <sup>b</sup>	11.7	10.6	6.72
Age-standardized incidence rate sa <sup>b</sup>	6.77	6.01	5.96
Cumulative risk (%) at 75 years old <sup>a</sup>	0.81	0.72	0.68
WOMEN			
Annual number of new cancer cases	2,386	3,457	113,502
Uncertainty intervals of new cancer cases [95% UI]	[2,111-2,696]	[3,284-3,640]	[105,599- 121,997]
Crude incidence rate sa <sup>c</sup>	6.95	6.43	2.94
Age-standardized incidence rate sa <sup>c</sup>	3.50	3.10	2.28
Cumulative risk (%) at 75 years old <sup>a</sup>	0.41	0.36	0.26

#### Data accessed on 27 Jan 2021

Data Sources:
Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

Data accessed on 27 Jan 2021

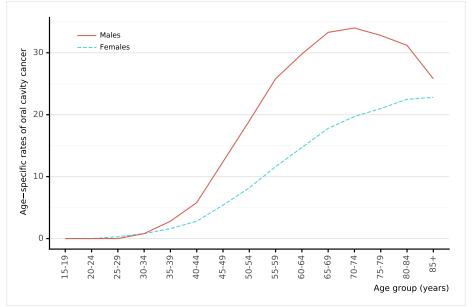
For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a Cumulative risk (incidence) is the probability or risk of individuals getting from the disease during ages 0-74 years. For cancer, it is expressed as the % of new born children who would be expected to develop from a particular cancer before the age of 75 if they had the rates of cancer observed in the period in the absence of competing causes.

b Rates per 100,000 men per year.

c Rates per 100,000 women per year.

Figure 42: Age-specific incidence rates of oral cavity cancer in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

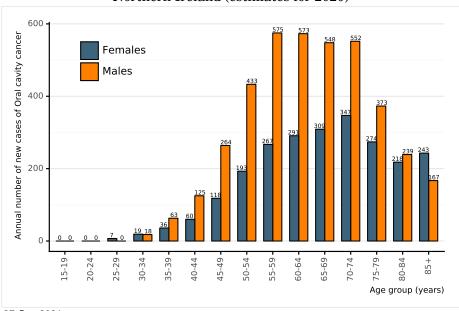


For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

 $^a$  Rates per 100,000 men per year.

Data Sources:
Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for  $Research \ on \ Cancer. \ Available \ from: \ \verb|https://gco.iarc.fr/today|, \ accessed \ [27 \ January \ 2021].$ 

Figure 43: Annual number of new cases of oral cavity cancer in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

 $For more \ detailed \ methods \ of \ estimation \ please \ refer \ to \ http://gco.iarc.fr/today/data-sources-methods$ 

Data Sources

Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for  $Research \ on \ Cancer. \ Available \ from: \ \verb|https://gco.iarc.fr/today|, \ accessed \ [27 \ January \ 2021].$ 

b Rates per 100,000 women per year

## 3.5.2.2 Oral cavity cancer incidence and mortality comparison in United Kingdom of Great **Britain and Northern Ireland**

Table 18: Oral cavity cancer mortality in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

Indicator	United Kingdom of Great Britain and Northern Ireland	Northern Europe	World
MEN			
Annual number of deaths	1,074	1,714	125,022
Uncertainty intervals of mortality cancer cases [95% UI]	[968-1,192]	[1,591-1,847]	[116,573- 134,084]
Crude mortality rate sa <sup>b</sup>	3.20	3.27	3.18
Age-standardized mortality rate sa <sup>b</sup>	1.63	1.67	2.82
Cumulative risk (%) at 75 years old <sup>a</sup>	0.19	0.20	0.32
WOMEN			
Annual number of deaths	656	1,004	52,735
Uncertainty intervals of mortality cancer cases [95% UI]	[584-737]	[911-1,107]	[47,690-58,313]
Crude mortality rate sa <sup>c</sup>	1.91	1.87	1.36
Age-standardized mortality rate sa <sup>c</sup>	0.76	0.72	1.04
Cumulative risk (%) at 75 years old <sup>a</sup>	0.08	0.08	0.12

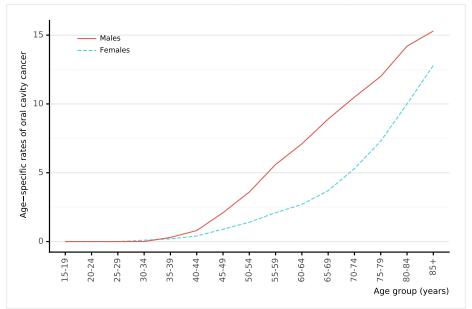
#### Data accessed on 27 Jan 2021

Data Sources:
Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for  $Research \ on \ Cancer. \ Available \ from: \ \texttt{https://gco.iarc.fr/today}, \ accessed \ [27 \ January \ 2021].$ 

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a Cumulative risk (mortality) is the probability or risk of individuals dying from the disease during ages 0-74 years. For cancer, it is expressed as the % of new born children who would be expected to die from a particular cancer before the age of 75 if they had the rates of cancer observed in the period in the absence of competing causes b Rates per 100,000 men per year. c Rates per 100,000 women per year.

Figure 44: Age-specific mortality rates of oral cavity cancer in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

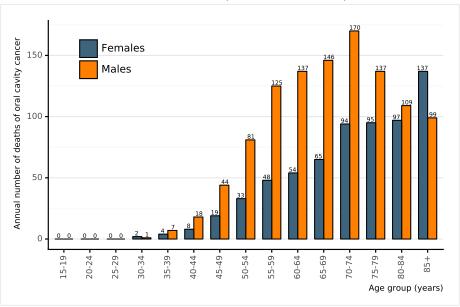


For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

 $^a$  Rates per 100,000 men per year.

Data Sources:
Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for  $Research \ on \ Cancer. \ Available \ from: \ \verb|https://gco.iarc.fr/today|, \ accessed \ [27 \ January \ 2021].$ 

Figure 45: Annual number of deaths of oral cavity cancer in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

 $For more \ detailed \ methods \ of \ estimation \ please \ refer \ to \ http://gco.iarc.fr/today/data-sources-methods$ 

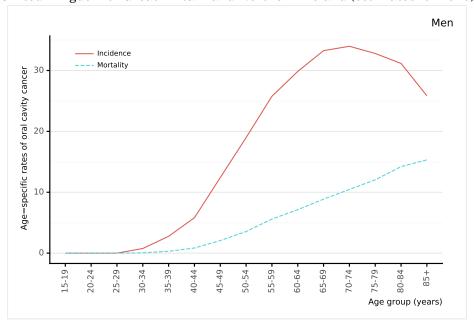
Data Sources

Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for  $Research \ on \ Cancer. \ Available \ from: \ \verb|https://gco.iarc.fr/today|, \ accessed \ [27 \ January \ 2021].$ 

b Rates per 100,000 women per year

#### 3.5.2.3 Oral cavity cancer incidence and mortality comparison in United Kingdom of Great **Britain and Northern Ireland**

Figure 46: Comparison of age-specific oral cavity cancer incidence and mortality rates among men in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

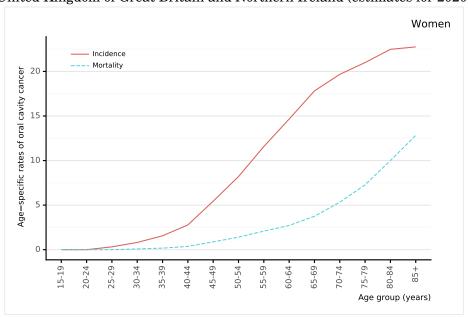


#### Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to  $\frac{1}{2} \frac{1}{2} \frac{1}$ 

Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

Figure 47: Comparison of age-specific oral cavity cancer incidence and mortality rates among women in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



#### Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods a Rates per 100,000 women per year.

Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

#### 3.5.3 Laryngeal cancer

#### 3.5.3.1 Laryngeal cancer incidence in United Kingdom of Great Britain and Northern Ireland

Table 19: Laryngeal cancer incidence in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

Indicator	United Kingdom of Great Britain and Northern Ireland	Northern Europe	World	
MEN				
Annual number of new cancer cases	2,115	3,160	160,265	
Uncertainty intervals of new cancer cases [95% UI]	[1,899-2,355]	[2,996-3,334]	[150,633- 170,513]	
Crude incidence rate sa <sup>b</sup>	6.31	6.02	4.08	
Age-standardized incidence rate sa <sup>b</sup>	3.30	3.16	3.59	
Cumulative risk (%) at 75 years old <sup>a</sup>	0.41	0.39	0.45	
WOMEN				
Annual number of new cancer cases	503	666	24,350	
Uncertainty intervals of new cancer cases [95% UI]	[416-608]	[594-747]	[20,845-28,444]	
Crude incidence rate sa <sup>c</sup>	1.46	1.24	0.63	
Age-standardized incidence rate sa <sup>c</sup>	0.74	0.62	0.49	
Cumulative risk (%) at 75 years old <sup>a</sup>	0.09	0.08	0.06	

#### Data accessed on 27 Jan 2021

Data Sources:
Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

Data accessed on 27 Jan 2021

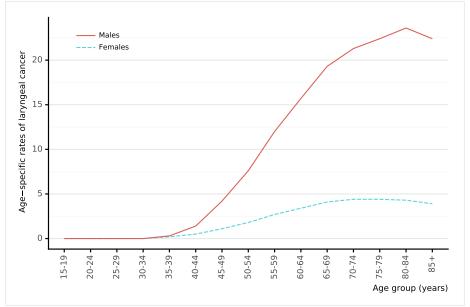
For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a Cumulative risk (incidence) is the probability or risk of individuals getting from the disease during ages 0-74 years. For cancer, it is expressed as the % of new born children who would be expected to develop from a particular cancer before the age of 75 if they had the rates of cancer observed in the period in the absence of competing causes.

b Rates per 100,000 men per year.

c Rates per 100,000 women per year.

Figure 48: Age-specific incidence rates of laryngeal cancer in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

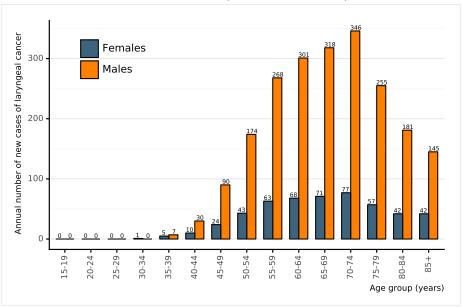


For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

 $^a$  Rates per 100,000 men per year.

Data Sources:
Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for  $Research \ on \ Cancer. \ Available \ from: \ \verb|https://gco.iarc.fr/today|, \ accessed \ [27 \ January \ 2021].$ 

Figure 49: Annual number of new cases of laryngeal cancer in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

 $For more \ detailed \ methods \ of \ estimation \ please \ refer \ to \ http://gco.iarc.fr/today/data-sources-methods$ 

Data Sources

Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for  $Research \ on \ Cancer. \ Available \ from: \ \verb|https://gco.iarc.fr/today|, \ accessed \ [27 \ January \ 2021].$ 

b Rates per 100,000 women per year

## 3.5.3.2 Laryngeal cancer incidence and mortality comparison in United Kingdom of Great **Britain and Northern Ireland**

Table 20: Laryngeal cancer mortality in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

Indicator	United Kingdom of Great Britain and Northern Ireland	Northern Europe	World
MEN			
Annual number of deaths	723	1,223	85,351
Uncertainty intervals of mortality cancer cases [95% UI]	[650-804]	[1,116-1,340]	[78,895-92,335]
Crude mortality rate sa <sup>b</sup>	2.16	2.33	2.17
Age-standardized mortality rate sa <sup>b</sup>	0.96	1.07	1.89
Cumulative risk (%) at 75 years old <sup>a</sup>	0.11	0.13	0.23
WOMEN			
Annual number of deaths	192	251	14,489
Uncertainty intervals of mortality cancer cases [95% UI]	[149-248]	[208-303]	[11,902-17,639]
Crude mortality rate sa <sup>c</sup>	0.56	0.47	0.37
Age-standardized mortality rate sa <sup>c</sup>	0.23	0.19	0.28
Cumulative risk (%) at 75 years old <sup>a</sup>	0.03	0.02	0.03

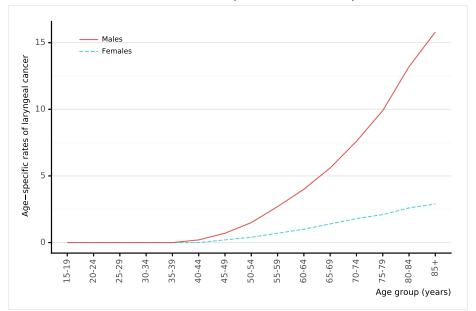
#### Data accessed on 27 Jan 2021

Data Sources:
Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for  $Research \ on \ Cancer. \ Available \ from: \ \texttt{https://gco.iarc.fr/today}, \ accessed \ [27 \ January \ 2021].$ 

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a Cumulative risk (mortality) is the probability or risk of individuals dying from the disease during ages 0-74 years. For cancer, it is expressed as the % of new born children who would be expected to die from a particular cancer before the age of 75 if they had the rates of cancer observed in the period in the absence of competing causes <sup>b</sup> Rates per 100,000 men per year.
<sup>c</sup> Rates per 100,000 women per year.

Figure 50: Age-specific mortality rates of laryngeal cancer in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

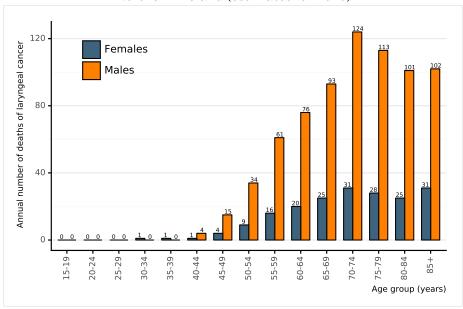


For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

 $^a$  Rates per 100,000 men per year.

Data Sources:
Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for  $Research \ on \ Cancer. \ Available \ from: \ \verb|https://gco.iarc.fr/today|, \ accessed \ [27 \ January \ 2021].$ 

Figure 51: Annual number of deaths of of laryngeal cancer in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

 $For more \ detailed \ methods \ of \ estimation \ please \ refer \ to \ http://gco.iarc.fr/today/data-sources-methods$ 

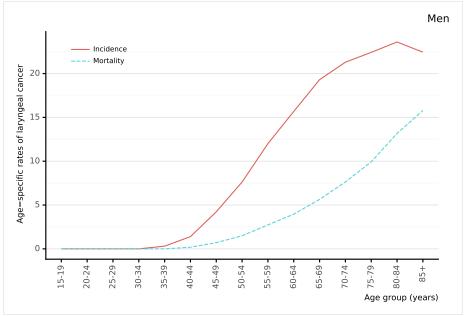
Data Sources

Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for  $Research \ on \ Cancer. \ Available \ from: \ \verb|https://gco.iarc.fr/today|, \ accessed \ [27 \ January \ 2021].$ 

b Rates per 100,000 women per year

#### 3.5.3.3 Laryngeal cancer incidence and mortality comparison in United Kingdom of Great **Britain and Northern Ireland**

Figure 52: Comparison of age-specific laryngeal cancer incidence and mortality rates among men in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

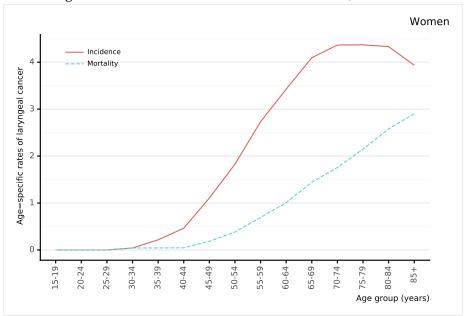


Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to  $\frac{1}{2} \frac{1}{2} \frac{1}$ 

Data Sources:
Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

Figure 53: Comparison of age-specific laryngeal cancer incidence and mortality rates among women in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

Rates per 100,000 women per year.

Ferlay J. Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

#### 4 HPV related statistics

HPV infection is commonly found in the anogenital tract of men and women with and without clinical lesions. The aetiological role of HPV infection among women with cervical cancer is well-established, and there is growing evidence of its central role in other anogenital sites. HPV is also responsible for other diseases such as recurrent juvenile respiratory papillomatosis and genital warts, both mainly caused by HPV types 6 and 11 (Lacey CJ, Vaccine 2006; 24(S3):35). For this section, the methodologies used to compile the information on HPV burden are derived from systematic reviews and meta-analyses of the literature. Due to the limitations of HPV DNA detection methods and study designs used, these data should be interpreted with caution and used only as a guide to assess the burden of HPV infection within the population. (Vaccine 2006, Vol. 24, Suppl 3; Vaccine 2008, Vol. 26, Suppl 10; Vaccine 2012, Vol. 30, Suppl 5; IARC Monographs 2007, Vol. 90).

## 4.1 HPV burden in women with normal cervical cytology, cervical precancerous lesions or invasive cervical cancer

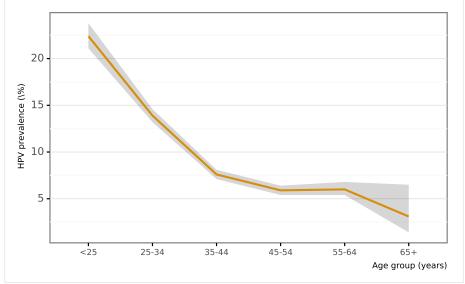
The statistics shown in this section focus on HPV infection in the cervix uteri. HPV cervical infection results in cervical morphological lesions ranging from normalcy (cytologically normal women) to different stages of precancerous lesions (CIN-1, CIN-2, CIN-3/CIS) and invasive cervical cancer. HPV infection is measured by HPV DNA detection in cervical cells (fresh tissue, paraffin embedded or exfoliated cells). The prevalence of HPV increases with lesion severity. HPV causes virtually 100% of cervical cancer cases, and an underestimation of HPV prevalence in cervical cancer is most likely due to the limitations of study methodologies. Worldwide, HPV16 and 18 (the two vaccine-preventable types) contribute to over 70% of all cervical cancer cases, between 41% and 67% of high-grade cervical lesions and 16-32% of low-grade cervical lesions. After HPV16/18, the six most common HPV types are the same in all world regions, namely 31, 33, 35, 45, 52 and 58; these account for an additional 20% of cervical cancers worldwide (Clifford G, Vaccine 2006;24(S3):26).

# Methods: Prevalence and type distribution of human papillomavirus in cervical carcinoma, low-grade cervical lesions, high-grade cervical lesions and normal cytology: systematic review and meta-analysis

A systematic review of the literature was conducted regarding the worldwide HPV-prevalence and type distribution for cervical carcinoma, low-grade cervical lesions, high-grade cervical lesions and normal cytology from 1990 to 'data as of' indicated in each section. The search terms for the review were 'HPV' AND cerv\* using Pubmed. There were no limits in publication language. References cited in selected articles were also investigated. Inclusion criteria were: HPV DNA detection by means of PCR or HC2, a minimum of 20 cases for cervical carcinoma, 20 cases for low-grade cervical lesions, 20 cases for highgrade cervical lesions and 100 cases for normal cytology and a detailed description of HPV DNA detection and genotyping techniques used. The number of cases tested and HPV positive extracted for each study were pooled to estimate the prevalence of HPV DNA and the HPV type distribution globally and by geographical region. Binomial 95% confidence intervals were calculated for each HPV prevalence. For more details refer to the methods document.

#### 4.1.1 HPV prevalence in women with normal cervical cytology

Figure 54: Crude age-specific HPV prevalence (%) and 95% confidence interval in women with normal cervical cytology in United Kingdom of Great Britain and Northern Ireland



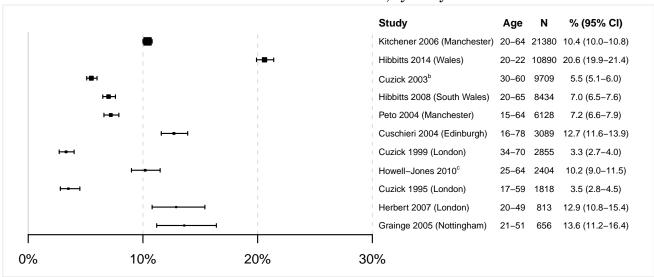
Data updated on 30 Jun 2015 (data as of 30 Jun 2014)

Data Source

Cuschieri KS, J Clin Pathol 2004; 57: 68 | Grainge MJ, Emerging Infect Dis 2005; 11: 1680 | Herbert A, J Fam Plann Reprod Health Care 2007; 33: 171 | Howell-Jones R, Br J Cancer 2010; 103: 209 | Kitchener HC, Br J Cancer 2006; 95: 56 | Peto J, Br J Cancer 2004; 91: 942
Based on systematic reviews and meta-analysis performed by ICO. The ICO HPV Information Centre has updated data until June 2014. Reference publications: 1) Bruni L, J Infect Dis

2010; 202: 1789. 2) De Sanjosé S, Lancet Infect Dis 2007; 7: 453

Figure 55: HPV prevalence among women with normal cervical cytology in United Kingdom of Great Britain and Northern Ireland, by study



Data updated on 30 Jun 2015 (data as of 30 Jun 2014)

The samples for HPV testing come from cervical specimens (fresh/fixed biopsies or exfoliated cells)

Data Sources

 $\overline{\text{Cuschieri KS}}, J \text{ Clin Pathol 2004; } 57: 68 \mid \text{Cuzick J}, \text{Br J Cancer 1999; } 81: 554 \mid \text{Cuzick J}, \text{Lancet 1995; } 345: 1533 \mid \text{Cuzick J}, \text{Lancet 2003; } 362: 1871 \mid \text{Grainge MJ}, \text{Emerging Infect Dis 2005; } 11: 1680 \mid \text{Herbert A}, \text{J Fam Plann Reprod Health Care 2007; } 33: 171 \mid \text{Hibbitts S}, \text{Br J Cancer 2008; } 99: 1929 \mid \text{Hibbitts S}, \text{J Clin Virol 2014; } 59: 109 \mid \text{Howell-Jones R}, \text{Br J Cancer 2008; } 99: 1929 \mid \text{Hibbitts S}, \text{J Clin Virol 2014; } 59: 109 \mid \text{Howell-Jones R}, \text{Br J Cancer 2008; } 99: 1929 \mid \text{Hibbitts S}, \text{J Clin Virol 2014; } 59: 109 \mid \text{Howell-Jones R}, \text{Br J Cancer 2008; } 99: 1929 \mid \text{Hibbitts S}, \text{J Clin Virol 2014; } 59: 109 \mid \text{Howell-Jones R}, \text{Br J Cancer 2008; } 99: 1929 \mid \text{Hibbitts S}, \text{J Clin Virol 2014; } 59: 109 \mid \text{Howell-Jones R}, \text{J Clin Virol 2014; } 59: 10$ 2010; 103: 209 | Kitchener HC, Br J Cancer 2006; 95: 56 | Peto J, Br J Cancer 2004; 91: 942

Based on systematic reviews and meta-analysis performed by ICO. The ICO HPV Information Centre has updated data until June 2014. Reference publications: 1) Bruni L, J Infect Dis 2010; 202: 1789. 2) De Sanjosé S, Lancet Infect Dis 2007; 7: 453

a Number of women tested b Birmingham, Edinburg, London, Manchester and Mansfield  $^{\it c}$  Gateshead, Birmingham, London, Gloucestershire and Norfolk

#### 4.1.2 HPV type distribution among women with normal cervical cytology, precancerous cervical lesions and cervical cancer

Table 21: Prevalence of HPV16 and HPV18 by cytology in United Kingdom of Great Britain and Northern Ireland

	No. tested	HPV 16/18 Prevalence % (95% CI)
Normal cytology <sup>1,2</sup>	42449	3.2 (3.0-3.4)
Low-grade lesions <sup>3,4</sup>	2768	29.6 (27.9-31.3)
High-grade lesions <sup>5,6</sup>	2927	58.6 (56.8-60.4)
Cervical cancer <sup>7,8</sup>	3140	79.0 (77.6-80.4)

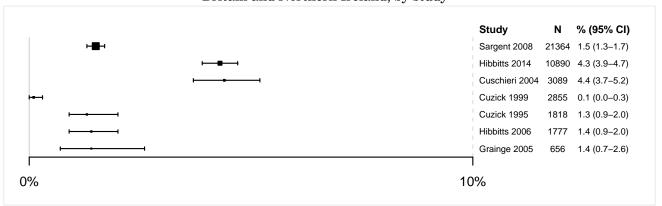
#### Data updated on 19 May 2017 (data as of 30 Jun 2015 / 30 Nov 2014)

The samples for HPV testing come from cervical specimens (fresh/fixed biopsies or exfoliated cells)

a Number of women tested b 95% Confidence Interval

Data Sources:

Figure 56: HPV 16 prevalence among women with normal cervical cytology in United Kingdom of Great Britain and Northern Ireland, by study



#### Data updated on 30 Jun 2015 (data as of 30 Jun 2014)

The samples for HPV testing come from cervical specimens (fresh/fixed biopsies or exfoliated cells)

a Number of women tested

Data Sources

Cuschieri KS, J Clin Pathol 2004; 57: 68 | Cuzick J, Br J Cancer 1999; 81: 554 | Cuzick J, Lancet 1995; 345: 1533 | Grainge MJ, Emerging Infect Dis 2005; 11: 1680 | Hibbitts S, Br J Cancer 2006; 95: 226 | Hibbitts S, J Clin Virol 2014; 59: 109 | Sargent A, Br J Cancer 2008; 98: 1704

Based on systematic reviews and meta-analysis performed by ICO. The ICO HPV Information Centre has updated data until June 2014. Reference publications: 1) Bruni L, J Infect Dis 2010; 202: 1789. 2) De Sanjosé S, Lancet Infect Dis 2007; 7: 453

<sup>1</sup> Cuschieri KS, J Clin Pathol 2004; 57: 68 | Cuzick J, Br J Cancer 1999; 81: 554 | Cuzick J, Lancet 1995; 345: 1533 | Grainge MJ, Emerging Infect Dis 2005; 11: 1680 | Hibbitts S, Br J Cancer 2006; 95: 226 | Hibbitts S, J Clin Virol 2014; 59: 109 | Sargent A, Br J Cancer 2008; 98: 1704

<sup>2</sup> Based on systematic reviews and meta-analysis performed by ICO. The ICO HPV Information Centre has updated data until November 2014. Reference publications: 1) Bruni L, J Infect Dis 2010; 202: 1789. 2) De Sanjosé S, Lancet Infect Dis 2007; 7: 453

<sup>3</sup> Contributing studies: Anderson L, J Med Virol 2013; 85: 295 | Arends MJ, Hum Pathol 1993; 24: 432 | Cuschieri KS, J Clin Pathol 2004; 57: 68 | Cuzick J, Br J Cancer 1994; 69: 167 | Cuzick J, Br J Cancer 1999; 81: 554 | Giannoudis A, Int J Cancer 1999; 83: 66 | Hibbitts S, Br J Cancer 2008; 99: 1929 | Howell-Jones R, Br J Cancer 2010; 103: 209 | Jamison J, Cytopathology 2009; 20: 242 | Sargent A, Br J Cancer 2008; 98: 1704 | Southern SA, Hum Pathol 2001; 32: 1351 | Woo YL, Int J Cancer 2010; 126: 133

<sup>4</sup> Based on meta-analysis performed by IARC's Infections and Cancer Epidemiology Group up to November 2011, the ICO HPV Information Centre has updated data until June 2015. Reference publications: 1) Guan P, Int J Cancer 2012;131:2349 2) Clifford GM, Cancer Epidemiol Biomarkers Prev 2005;14:1157

<sup>&</sup>lt;sup>5</sup> Contributing studies: Anderson L, J Med Virol 2013; 85: 295 | Arends MJ, Hum Pathol 1993; 24: 432 | Cuschieri KS, J Clin Pathol 2004; 57: 68 | Cuzick J, Br J Cancer 1994; 69: 167 | Cuzick J, J Clin Virol 2014; 60: 44 | Geraets DT, J Clin Microbiol 2014; 52: 3996 | Herrington CS, Br J Cancer 1995; 71: 206 | Hibbitts S, Br J Cancer 2008; 99: 1929 | Howell-Jones R, Br J Cancer 2010; 103: 209 | Jamison J, Cytopathology 2009; 20: 242 | Sargent A, Br J Cancer 2008; 98: 1704 | Southern SA, Diagn Mol Pathol 1998; 7: 114

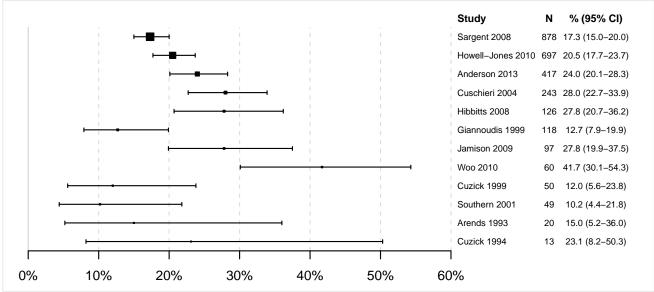
Based on meta-analysis performed by IARC's Infections and Cancer Epidemiology Group up to November 2011, the ICO HPV Information Centre has updated data until June 2015.

Reference publications: 1) Guan P, Int J Cancer 2012;131:2349 2) Smith JS, Int J Cancer 2007;121:621 3) Clifford GM, Br J Cancer 2003;89:101.

<sup>7</sup> Contributing studies: Arends MJ, Hum Pathol 1993; 24: 432 | Crook T, Lancet 1992; 339: 1070 | Cuschieri K, Br J Cancer 2010; 102: 930 | Cuschieri K, Int J Cancer 2014; 135: 2721 | Cuzick J, Br J Cancer 2000; 82: 1348 | Giannoudis A, Int J Cancer 1999; 83: 66 | Howell-Jones R, Br J Cancer 2010; 103: 209 | Mesher D, J Clin Pathol 2015; 68: 135 | Powell N, Int J Cancer 2009; 125: 2425 | Tawfik El-Mansi M, Int J Gynecol Cancer 2006; 16: 1025

<sup>8</sup> Based on meta-analysis performed by IARC's Infections and Cancer Epidemiology Group up to November 2011, the ICO HPV Information Centre has updated data until June 2015. Reference publications: 1) Guan P, Int J Cancer 2012;131:2349 2) Li N, Int J Cancer 2011;128:927 3) Smith JS, Int J Cancer 2007;121:621 4) Clifford GM, Br J Cancer 2003;88:63 5) Clifford GM, Br J Cancer 2003;89:101.

Figure 57: HPV 16 prevalence among women with low-grade cervical lesions in United Kingdom of Great Britain and Northern Ireland, by study



Data updated on 27 Jan 2017 (data as of 30 Jun 2015)

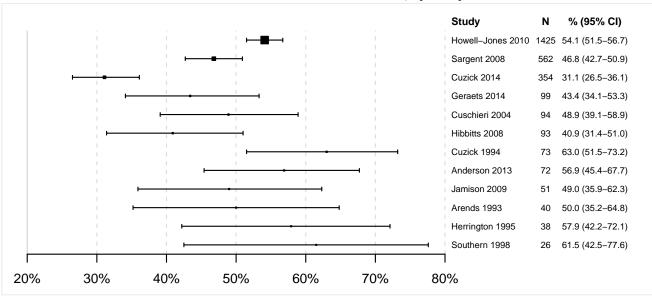
The samples for HPV testing come from cervical specimens (fresh/fixed biopsies or exfoliated cells)  $^{a}\,$  Number of women tested

Data Sources

Anderson L, J Med Virol 2013; 85: 295 | Arends MJ, Hum Pathol 1993; 24: 432 | Cuschieri KS, J Clin Pathol 2004; 57: 68 | Cuzick J, Br J Cancer 1994; 69: 167 | Cuzick J, Br J Cancer 1999; 81: 554 | Giannoudis A, Int J Cancer 1999; 83: 66 | Hibbitts S, Br J Cancer 2008; 99: 1929 | Howell-Jones R, Br J Cancer 2010; 103: 209 | Jamison J, Cytopathology 2009; 20: 242 | Sargent A, Br J Cancer 2008; 98: 1704 | Southern SA, Hum Pathol 2001; 32: 1351 | Woo YL, Int J Cancer 2010; 126: 133

Based on meta-analysis performed by IARC's Infections and Cancer Epidemiology Group up to November 2011, the ICO HPV Information Centre has updated data until June 2015. Reference publications: 1) Guan P, Int J Cancer 2012;131:2349 2) Clifford GM, Cancer Epidemiol Biomarkers Prev 2005;14:1157

Figure 58: HPV 16 prevalence among women with high-grade cervical lesions in United Kingdom of Great Britain and Northern Ireland, by study



Data updated on 27 Jan 2017 (data as of 30 Jun 2015)

The samples for HPV testing come from cervical specimens (fresh/fixed biopsies or exfoliated cells) a Number of women tested

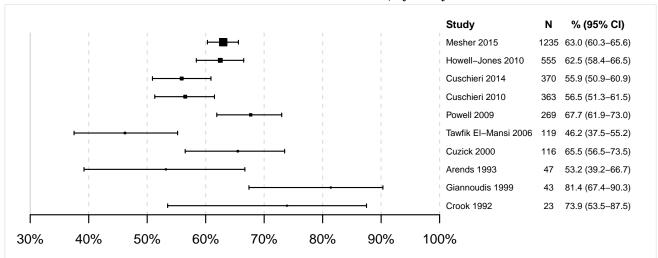
Data Sources

Anderson L, J Med Virol 2013; 85: 295 | Arends MJ, Hum Pathol 1993; 24: 432 | Cuschieri KS, J Clin Pathol 2004; 57: 68 | Cuzick J, Br J Cancer 1994; 69: 167 | Cuzick J, J Clin Virol 2014; 60: 44 | Geraets DT, J Clin Microbiol 2014; 52: 3996 | Herrington CS, Br J Cancer 1995; 71: 206 | Hibbitts S, Br J Cancer 2008; 99: 1929 | Howell-Jones R, Br J Cancer 2010; 103: 209 | Jamison J, Cytopathology 2009; 20: 242 | Sargent A, Br J Cancer 2008; 98: 1704 | Southern SA, Diagn Mol Pathol 1998; 7: 114
Based on meta-analysis performed by IARC's Infections and Cancer Epidemiology Group up to November 2011, the ICO HPV Information Centre has updated data until June 2015. Refer-

ence publications: 1) Guan P, Int J Cancer 2012;131:2349 2) Smith JS, Int J Cancer 2007;121:621 3) Clifford GM, Br J Cancer 2003;89:101

- 57 -

Figure 59: HPV 16 prevalence among women with invasive cervical cancer in United Kingdom of Great Britain and Northern Ireland, by study



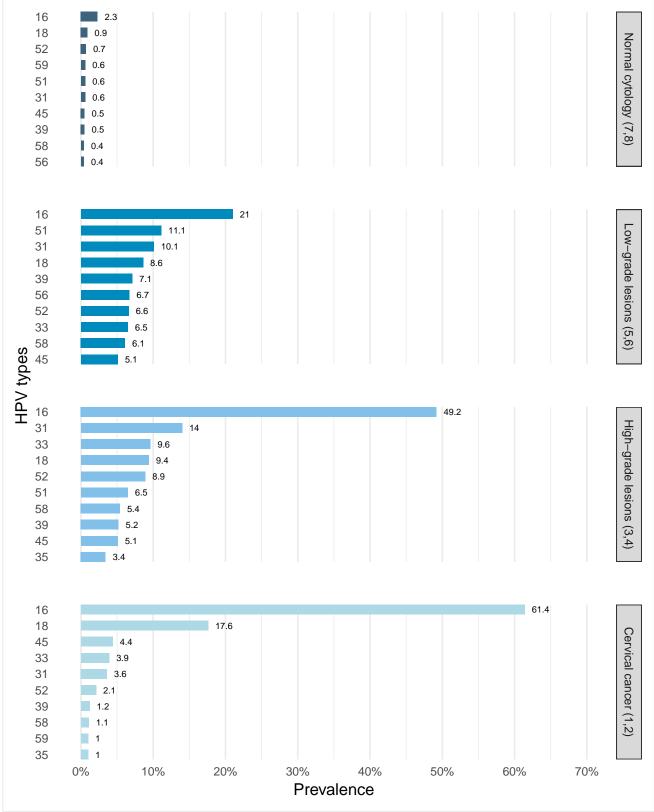
Data updated on 19 May 2017 (data as of 30 Jun 2015)

The samples for HPV testing come from cervical specimens (fresh/fixed biopsies or exfoliated cells)  $^{\it a}$  Number of women tested

<u>Data Sources:</u>
Arends MJ, Hum Pathol 1993; 24: 432 | Crook T, Lancet 1992; 339: 1070 | Cuschieri K, Br J Cancer 2010; 102: 930 | Cuschieri K, Int J Cancer 2014; 135: 2721 | Cuzick J, Br J Cancer 2000; 82: 1348 | Giannoudis A, Int J Cancer 1999; 83: 66 | Howell-Jones R, Br J Cancer 2010; 103: 209 | Mesher D, J Clin Pathol 2015; 68: 135 | Powell N, Int J Cancer 2009; 125: 2425 | Tawfik El-Mansi M, Int J Gynecol Cancer 2006; 16: 1025

Based on meta-analysis performed by IARC's Infections and Cancer Epidemiology Group up to November 2011, the ICO HPV Information Centre has updated data until June 2015. Reference publications: 1) Guan P, Int J Cancer 2012;131:2349 2) Li N, Int J Cancer 2011;128:927 3) Smith JS, Int J Cancer 2007;121:621 4) Clifford GM, Br J Cancer 2003;88:63 5) Clifford GM, Br J Cancer 2003:89:101.

Figure 60: Comparison of the ten most frequent HPV oncogenic types in United Kingdom of Great Britain and Northern Ireland among women with and without cervical lesions



Data updated on 30 Jun 2015 (data as of 30 Jun 2015)

Data Sources

<sup>1</sup> Contributing studies: Arends MJ, Hum Pathol 1993; 24: 432 | Crook T, Lancet 1992; 339: 1070 | Cuschieri K, Br J Cancer 2010; 102: 930 | Cuschieri K, Int J Cancer 2014; 135: 2721 | Cuzick J, Br J Cancer 2000; 82: 1348 | Giannoudis A, Int J Cancer 1999; 83: 66 | Howell-Jones R, Br J Cancer 2010; 103: 209 | Mesher D, J Clin Pathol 2015; 68: 135 | Powell N, Int J Cancer 2009; 125: 2425 | Tawfik El-Mansi M, Int J Gynecol Cancer 2006; 16: 1025

Cancer 2009; 125: 2425 | Tawthk El-Mansı M, Int J Gynecol Cancer 2006; 16: 1025

Based on meta-analysis performed by IARC's Infections and Cancer Epidemiology Group up to November 2011, the ICO HPV Information Centre has updated data until June 2015.

Reference publications: 1) Guan P, Int J Cancer 2012;131:2349 2) Li N, Int J Cancer 2011;128:927 3) Smith JS, Int J Cancer 2007;121:621 4) Clifford GM, Br J Cancer 2003;88:63 5) Clifford GM. Br J Cancer 2003;88:101.

GM, Br J Cancer 2003;89:101.

3 Contributing studies: Anderson L, J Med Virol 2013; 85: 295 | Arends MJ, Hum Pathol 1993; 24: 432 | Cuschieri KS, J Clin Pathol 2004; 57: 68 | Cuzick J, Br J Cancer 1994; 69: 167 | Cuzick J, J Clin Virol 2014; 60: 44 | Geraets DT, J Clin Microbiol 2014; 52: 3996 | Herrington CS, Br J Cancer 1995; 71: 206 | Hibbitts S, Br J Cancer 2008; 99: 1929 | Howell-Jones R, Br

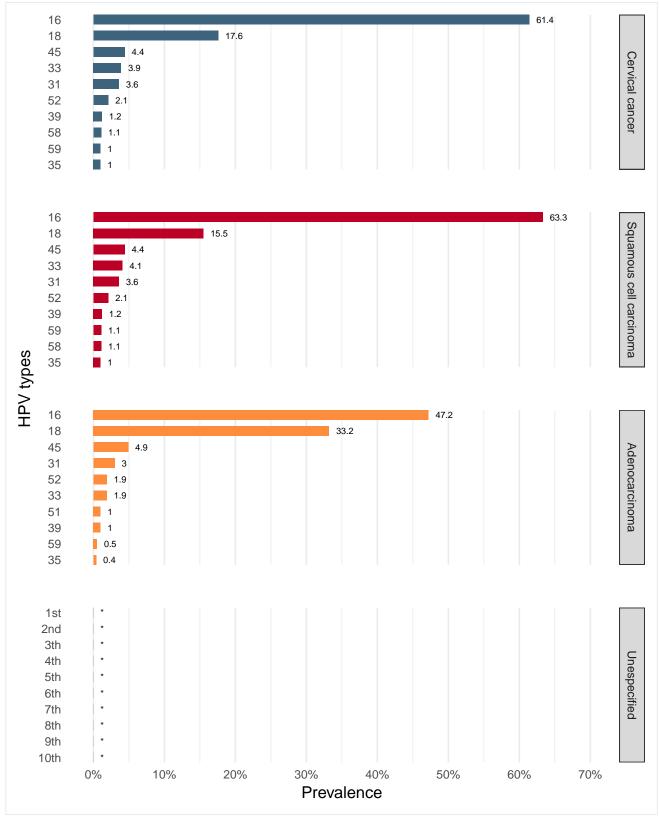
- J Cancer 2010; 103: 209 | Jamison J, Cytopathology 2009; 20: 242 | Sargent A, Br J Cancer 2008; 98: 1704 | Southern SA, Diagn Mol Pathol 1998; 7: 114
- 4 Based on meta-analysis performed by IARC's Infections and Cancer Epidemiology Group up to November 2011, the ICO HPV Information Centre has updated data until June 2015.  $Reference\ publications:\ 1)\ Guan\ P,\ Int\ J\ Cancer\ 2012; 131:2349\ 2)\ Smith\ JS,\ Int\ J\ Cancer\ 2007; 121:621\ 3)\ Clifford\ GM,\ Br\ J\ Cancer\ 2003; 89:101.$
- Reference publications: 1) Guan P, Int J Cancer 2012;131:2349 2) Smith JS, Int J Cancer 2007;121:621 3) Clifford GM, Br J Cancer 2003;89:101.

  Contributing studies: Anderson L, J Med Virol 2013; 85: 295 | Arends MJ, Hum Pathol 1993; 24: 432 | Cuschieri KS, J Clin Pathol 2004; 57: 68 | Cuzick J, Br J Cancer 1994; 69: 167 | Cuzick J, Br J Cancer 1999; 81: 554 | Giannoudis A, Int J Cancer 1999; 83: 66 | Hibbitts S, Br J Cancer 2008; 99: 1929 | Howell-Jones R, Br J Cancer 2010; 103: 209 | Jamison J, Cytopathology 2009; 20: 242 | Sargent A, Br J Cancer 2008; 98: 1704 | Southern SA, Hum Pathol 2001; 32: 1351 | Woo YL, Int J Cancer 2010; 126: 133

  Based on meta-analysis performed by IARC's Infections and Cancer Epidemiology Group up to November 2011, the ICO HPV Information Centre has updated data until June 2015. Reference publications: 1) Guan P, Int J Cancer 2012;131:2349 2) Clifford GM, Cancer Epidemiol Biomarkers Prev 2005;14:1157
- 7 Cuschieri KS, J Clin Pathol 2004; 57: 68 | Cuzick J, Br J Cancer 1999; 81: 554 | Cuzick J, Lancet 1995; 345: 1533 | Grainge MJ, Emerging Infect Dis 2005; 11: 1680 | Hibbitts S, Br J Cancer 2006; 95: 226 | Hibbitts S, J Clin Virol 2014; 59: 109 | Sargent A, Br J Cancer 2008; 98: 1704

  8 Based on systematic reviews and meta-analysis performed by ICO. The ICO HPV Information Centre has updated data until November 2014. Reference publications: 1) Bruni L, J Infect
- Dis 2010; 202: 1789. 2) De Sanjosé S, Lancet Infect Dis 2007; 7: 453

Figure 61: Comparison of the ten most frequent HPV oncogenic types in United Kingdom of Great Britain and Northern Ireland among women with invasive cervical cancer by histology



Data updated on  $30~\mathrm{Jun}~2015$  (data as of  $30~\mathrm{Jun}~2015$ )

 $<sup>^{\</sup>ast}$  No data available. No more types than shown were tested or were positive <code>Data Sources</code>:

<sup>1</sup> Contributing studies: Arends MJ, Hum Pathol 1993; 24: 432 | Crook T, Lancet 1992; 339: 1070 | Cuschieri K, Br J Cancer 2010; 102: 930 | Cuschieri K, Int J Cancer 2014; 135: 2721 | Cuzick J, Br J Cancer 2000; 82: 1348 | Giannoudis A, Int J Cancer 1999; 83: 66 | Howell-Jones R, Br J Cancer 2010; 103: 209 | Mesher D, J Clin Pathol 2015; 68: 135 | Powell N, Int J Cancer 2009; 125: 2425 | Tawfik El-Mansi M, Int J Gynecol Cancer 2006; 16: 1025

<sup>&</sup>lt;sup>2</sup> Based on meta-analysis performed by IARC's Infections and Cancer Epidemiology Group up to November 2011, the ICO HPV Information Centre has updated data until June 2014. Reference publications: 1) Guan P, Int J Cancer 2012;131:2349 2) Li N, Int J Cancer 2011;128:927 3) Smith JS, Int J Cancer 2007;121:621 4) Clifford GM, Br J Cancer 2003;89:101.

<sup>3</sup> Based on meta-analysis performed by IARC's Infections and Cancer Epidemiology Group up to November 2011, the ICO HPV Information Centre has updated data until June 2015.

Reference publications: 1) Guan P, Int J Cancer 2012;131:2349 2) Li N, Int J Cancer 2011;128:927 3) Smith JS, Int J Cancer 2007;121:621 4) Clifford GM, Br J Cancer 2003;88:63 5) Clifford GM, Br J Cancer 2003;89:101.

Table 22: Type-specific HPV prevalence in women with normal cervical cytology, precancerous cervical lesions and invasive cervical cancer in United Kingdom of Great Britain and Northern Ireland

ies	lesions and invasive cervical cancer in United Kingdom of Great Britain and Northern Ireland							
TIPY		al cytology <sup>1,2</sup>	_	grade lesions <sup>3,4</sup>		grade lesions <sup>5,6</sup>		ical cancer <sup>7,8</sup>
HPV Type	No. tested	HPV Prev % (95% CI)	No. tested	HPV Prev % (95% CI)	No. tested	HPV Prev % (95% CI)	No. tested	HPV Prev % (95% CI)
	ENIC HPV		testeu	(99% C1)	testeu	(95% C1)	testeu	(95% C1)
	risk HPV ty							
16	42449	2.3 (2.2-2.5)	2768	21.0 (19.5-22.6)	2927	49.2 (47.4-51.0)	3140	61.4 (59.7-63.1)
18	42449	0.9 (0.8-1.0)	2708	8.6 (7.6-9.8)	2927	9.4 (8.4-10.5)	3140	17.6 (16.3-19.0)
31	30903	0.6 (0.6-0.7)	2688	10.1 (9.0-11.3)	2887	14.0 (12.8-15.3)	2974	3.6 (3.0-4.3)
33	30903	0.3 (0.3-0.4)	2708	6.5 (5.7-7.5)	2927	9.6 (8.6-10.7)	3021	3.9 (3.3-4.7)
35	29085	0.2 (0.2-0.3)	2688	3.7 (3.0-4.5)	2849	3.4 (2.8-4.2)	2951	1.0 (0.7-1.4)
39	26230	0.5 (0.5-0.6)	$\frac{2605}{2625}$	7.1 (6.2-8.1)	2776	5.2 (4.4-6.0)	2835	1.2 (0.9-1.7)
45	26230	0.5 (0.5-0.6)	2675	5.1 (4.3-6.0)	2776	5.1 (4.3-6.0)	2835	4.4 (3.7-5.2)
51	29085	0.6 (0.5-0.7)	2675	11.1 (9.9-12.3)	2776	6.5 (5.6-7.5)	2835	0.8 (0.5-1.2)
52	29085	0.7 (0.6-0.8)	2578	6.6 (5.7-7.6)	2725	8.9 (7.9-10.0)	2835	2.1 (1.6-2.7)
56	29085	0.4 (0.3-0.5)	2675	6.7 (5.8-7.7)	2776	2.9 (2.3-3.6)	2951	0.5 (0.3-0.8)
58	29085	0.4 (0.3-0.4)	2675	6.1 (5.3-7.1)	2776	5.4 (4.6-6.3)	2951	1.1 (0.7-1.5)
59	26230	0.6 (0.5-0.7)	2625	4.9 (4.1-5.8)	2750	3.2 (2.6-3.9)	2835	1.0 (0.7-1.5)
		e carcinogen	2020	4.0 (4.1-0.0)	2100	0.2 (2.0-0.0)	2000	1.0 (0.1-1.0)
26	3089	0.0 (0.0-0.1)	1915	0.3 (0.1-0.7)	1596	0.4 (0.2-0.9)	918	0.0 (0.0-0.4)
30	-	-	-	-	- 1000	0.4 (0.2-0.5)		-
34			-					
53	3089	0.5 (0.3-0.8)	1672	9.4 (8.1-10.9)	1502	2.7 (2.0-3.6)	1288	0.8 (0.4-1.4)
66	4866	0.6 (0.4-0.9)	2382	8.2 (7.2-9.4)	2120	2.9 (2.2-3.7)	1600	0.3 (0.1-0.7)
67	-	-	1672	2.4 (1.8-3.2)	1476	0.5 (0.3-1.1)	555	0.0 (0.0-0.7)
68	26230	0.2 (0.2-0.3)	2576	2.1 (1.6-2.8)	2750	1.7 (1.3-2.3)	2472	0.9 (0.6-1.4)
69		-	1672	0.3 (0.1-0.7)	1476	0.3 (0.1-0.7)	918	0.0 (0.0-0.4)
$\frac{-60}{70}$			1672	4.3 (3.4-5.4)	1476	1.3 (0.8-2.0)	918	0.2 (0.1-0.8)
73	3089	0.8 (0.5-1.2)	1915	5.0 (4.1-6.0)	2132	2.2 (1.6-2.9)	925	0.5 (0.2-1.3)
82	3089	0.1 (0.0-0.3)	1915	1.6 (1.1-2.2)	2132	1.2 (0.8-1.7)	918	0.0 (0.0-0.4)
85	-	-	-	-	-	-	-	-
97		-		-		-	370	0.3 (0.0-1.5)
	SK HPV TY	PES						***************************************
6	3089	0.6 (0.3-0.9)	1872	4.6 (3.8-5.7)	1615	1.6 (1.1-2.3)	1401	1.3 (0.8-2.0)
11	3089	0.2 (0.1-0.4)	1872	1.4 (0.9-2.0)	1615	0.3 (0.1-0.7)	1401	0.3 (0.1-0.7)
32	-	-	-	-	-	-	-	-
40	3089	0.1 (0.1-0.3)	-	-	-	-	363	0.0 (0.0-1.0)
42	3089	0.7 (0.5-1.1)	-	-	-	-	370	0.0 (0.0-1.0)
43	-	-	-	-	-	-	363	0.0 (0.0-1.0)
44	3089	0.4 (0.2-0.7)		-	-	-	733	0.3 (0.1-1.0)
54	3089	0.6 (0.4-1.0)	-	-	-	-	733	1.9 (1.1-3.2)
55	-	-	-	-	-	-	-	-
57	3089	0.1 (0.0-0.3)	-	-	-	-	-	-
61	-	-	-	-	-	-	-	-
62	-	-	-	-	-	-	-	-
64	-	-	-	-	-	-	-	-
71	-	-	-	-	-	-	363	0.0 (0.0-1.0)
72	-	-	-	-	-	-	-	-
74	-	-	-	-	-	-	363	0.0 (0.0-1.0)
81	-	-	-	-	-	-	-	-
83	3089	0.3 (0.1-0.5)	-	-	-	-	-	-
84	3089	0.4 (0.2-0.6)	-	-	-	-	-	-
86	-	-	-	-	-	-	-	-
87	-	-	-	-	-	-	-	-
89	-	-	-	-	-	-	-	-
90	-	-	-	-	-	-	-	-
91	-	-	-	-	-	-	-	-

### Data updated on 30 Jun 2015 (data as of 30 Jun 2015 / 30 Nov 2014)

The samples for HPV testing come from cervical specimens (fresh/fixed biopsies or exfoliated cells)

<sup>&</sup>lt;u>Data Sources</u>:

1 Cuschieri KS, J Clin Pathol 2004; 57: 68 | Cuzick J, Br J Cancer 1999; 81: 554 | Cuzick J, Lancet 1995; 345: 1533 | Grainge MJ, Emerging Infect Dis 2005; 11: 1680 | Hibbitts S, Br J

Cancer 2006; 95: 226 | Hibbitts S, J Clin Virol 2014; 59: 109 | Sargent A, Br J Cancer 2008; 98: 1704

Based on systematic reviews and meta-analysis performed by ICO. The ICO HPV Information Centre has updated data until November 2014. Reference publications: 1) Bruni L, J Infect

Dis 2010; 202: 1789. 2) De Sanjosé S, Lancet Infect Dis 2007; 7: 453 Contributing studies: Anderson L, J Med Virol 2013; 85: 295 | Arends MJ, Hum Pathol 1993; 24: 432 | Cuschieri KS, J Clin Pathol 2004; 57: 68 | Cuzick J, Br J Cancer 1994; 69: 167 Contributing studies: Anderson L, J Med Virol 2013; 88: 299 | Arenos MJ, Hum Pathol 1993; 24: 432 | Cuschier RS, J Clin Pathol 2004; 97: 68 | Cuzick J, Br J Cancer 1999; 81: 554 | Giannoudis A, Int J Cancer 1999; 83: 66 | Hibbitts S, Br J Cancer 2008; 99: 1929 | Howell-Jones R, Br J Cancer 2010; 103: 209 | Jamison J, Cytopathology 2009; 20: 242 | Sargent A, Br J Cancer 2008; 98: 1704 | Southern SA, Hum Pathol 2001; 32: 1351 | Woo YL, Int J Cancer 2010; 126: 133

Based on meta-analysis performed by IARC's Infections and Cancer Epidemiology Group up to November 2011, the ICO HPV Information Centre has updated data until June 2015. Reference publications: 1) Guan P, Int J Cancer 2012;131:2349 2) Clifford GM, Cancer Epidemiol Biomarkers Prev 2005;14:1157

<sup>&</sup>lt;sup>5</sup> Contributing studies: Anderson L, J Med Virol 2013; 85: 295 | Arends MJ, Hum Pathol 1993; 24: 432 | Cuschieri KS, J Clin Pathol 2004; 57: 68 | Cuzick J, Br J Cancer 1994; 69: 167 |

Cuzick J, J Clin Virol 2014; 60: 44 | Geraets DT, J Clin Microbiol 2014; 52: 3996 | Herrington CS, Br J Cancer 1995; 71: 206 | Hibbitts S, Br J Cancer 2008; 99: 1929 | Howell-Jones R, Br

Cuzick J, J Cin Viol 2014; 60: 44 | Geraets DI, J Cin Microbiol 2014; 52: 3996 | Herrington CS, Br J Cancer 1995; 17: 206 | Hibbits S, Br J Cancer 2008; 98: 1929 | Howell-Jones K, Br J Cancer 2010; 103: 209 | Jamison J, Cytopathology 2009; 20: 242 | Sargent A, Br J Cancer 2008; 98: 1704 | Southern SA, Diagn Mol Pathol 1998; 7: 114

Based on meta-analysis performed by IARC's Infections and Cancer Epidemiology Group up to November 2011, the ICO HPV Information Centre has updated data until June 2015.

Reference publications: 1) Guan P, Int J Cancer 2012;131:2349 2) Smith JS, Int J Cancer 2007;121:621 3) Clifford GM, Br J Cancer 2003;89:101.

Contributing studies: Arends MJ, Hum Pathol 1993; 24: 432 | Crook T, Lancet 1992; 339: 1070 | Cuschieri K, Br J Cancer 2010; 102: 930 | Cuschieri K, Int J Cancer 2014; 135: 2721 |

Couzick J, Br J Cancer 2000; 82: 1348 | Giannoudis A, Int J Cancer 1999; 83: 66 | Howell-Jones R, Br J Cancer 2010; 103: 209 | Mesher D, J Clin Pathol 2015; 68: 135 | Powell N, Int J Cancer 2009; 125: 2425 | Tawfik El-Mansi M, Int J Gynecol Cancer 2006; 16: 1025

8 Based on meta-analysis performed by IARC's Infections and Cancer Epidemiology Group up to November 2011, the ICO HPV Information Centre has updated data until June 2015. Reference publications: 1) Guan P, Int J Cancer 2012;131:2349 2) Li N, Int J Cancer 2011;128:927 3) Smith JS, Int J Cancer 2007;121:621 4) Clifford GM, Br J Cancer 2003;88:63 5) Clifford GM, Br J Cancer 2003;89:101.

Table 23: Type-specific HPV prevalence among invasive cervical cancer cases in United Kingdom of Great Britain and Northern Ireland by histology

	An	y Histology		us cell carcinoma		nocarcinoma	Ur	nespecified
HPV	No.	HPV Prev %	No.	HPV Prev %	No.	HPV Prev %	No.	HPV Prev %
Туре	tested	(95% CI)	tested	(95% CI)	tested	(95% CI)	tested	(95% CI)
	ENIC HPV		Jesteu	(00% 02)	testea	(00% 01)	testea	(0070 01)
	risk HPV ty							
16	3140	61.4 (59.7-63.1)	2763	63.3 (61.5-65.1)	377	47.2 (42.2-52.3)		
18	3140	17.6 (16.3-19.0)	2763	15.5 (14.2-16.9)	377	33.2 (28.6-38.1)		
31	2974	3.6 (3.0-4.3)	2737	3.6 (3.0-4.4)	237	3.0 (1.4-6.0)		<u>-</u>
33	3021	3.9 (3.3-4.7)	2763	4.1 (3.4-4.9)	258	1.9 (0.8-4.5)		<u>-</u>
35	2951	1.0 (0.7-1.4)	2714	1.0 (0.7-1.5)	$\frac{238}{237}$	0.4 (0.1-2.4)	<del>-</del>	<u> </u>
		1.2 (0.9-1.7)						-
39	2835	, ,	2629	1.2 (0.9-1.7)	206	1.0 (0.3-3.5)		-
45	2835	4.4 (3.7-5.2)	2629	4.4 (3.7-5.2)	206	4.9 (2.7-8.7)		-
51	2835	0.8 (0.5-1.2)	2629	0.8 (0.5-1.2)	206	1.0 (0.3-3.5)		-
52	2835	2.1 (1.6-2.7)	2629	2.1 (1.6-2.8)	206	1.9 (0.8-4.9)		-
56	2951	0.5 (0.3-0.8)	2714	0.6 (0.3-0.9)	237	0.0 (0.0-1.6)		-
58	2951	1.1 (0.7-1.5)	2714	1.1 (0.8-1.6)	237	0.0 (0.0-1.6)		-
59	2835	1.0 (0.7-1.5)	2629	1.1 (0.7-1.5)	206	0.5 (0.1-2.7)		-
		e carcinogen						
26	918	0.0 (0.0-0.4)	-	-	-	-		-
30	-	-	-	-	-	-	-	-
34	-	-	-	-	-	-	-	-
53	1288	0.8 (0.4-1.4)	-	-	-	-	-	-
66	1600	0.3 (0.1-0.7)	1394	0.4 (0.2-0.8)	206	0.0 (0.0-1.8)	-	-
67	555	0.0 (0.0-0.7)	450	0.0 (0.0-0.8)	105	0.0 (0.0-3.5)	-	-
68	2472	0.9 (0.6-1.4)	2320	0.9 (0.6-1.4)	152	0.7 (0.1-3.6)	-	-
69	918	0.0 (0.0-0.4)	-	-	-	-	-	-
70	918	0.2 (0.1-0.8)	-	-	-	-	-	-
73	925	0.5 (0.2-1.3)	-	-	-	-	-	-
82	918	0.0 (0.0-0.4)	759	0.0 (0.0-0.5)	159	0.0 (0.0-2.4)		-
85		-	-	-	-	-		-
97	370	0.3 (0.0-1.5)	370	0.3 (0.0-1.5)	-	-		-
LOW RIS	SK HPV TY							
6	1401	1.3 (0.8-2.0)	-	-				-
11	1401	0.3 (0.1-0.7)						-
32		-						-
40	363	0.0 (0.0-1.0)						_
42	370	0.0 (0.0-1.0)	370	0.0 (0.0-1.0)		-		
43	363	0.0 (0.0-1.0)		-				
44	733	0.3 (0.1-1.0)	679	0.3 (0.1-1.1)	54	0.0 (0.0-6.6)		-
54	733	1.9 (1.1-3.2)	-	-		- 0.0 (0.0-0.0)		-
55		1.0 (1.1-0.2)						
57								
61		-		-		-		-
62		-		-		-		-
64		-		-		-		-
	- 262	- 0.0 (0.0 1.0)	-	-	-	-		-
$\frac{71}{72}$	363	0.0 (0.0-1.0)	-	-		-		-
	- 000	- 0.0 (0.0 1.0)		-	-	-		-
74	363	0.0 (0.0-1.0)		-	-	-		-
81		-		-	-	-		-
83		-	-	-	-	-		-
84	-	-	-		-	-	-	-
86	-	-	-		-		-	-
87	-	-	-	-	-		-	-
89	-	-	-	-	-	-	-	-
90	-	-	-	-	-	-	-	-
91	-	-	-	-	-	-	-	-

Data updated on 19 May 2017 (data as of 30 Jun 2015)

The samples for HPV testing come from cervical specimens (fresh/fixed biopsies or exfoliated cells)

Data Sources:

Contributing studies: Arends MJ, Hum Pathol 1993; 24: 432 | Crook T, Lancet 1992; 339: 1070 | Cuschieri K, Br J Cancer 2010; 102: 930 | Cuschieri K, Int J Cancer 2014; 135: 2721 |

Contributing studies: Arenas MJ, rium Fathol 1993; 24: 452 f - Crook 1, Lancer 1992; 339: 1107 l Cuscineri K, Br J Cancer 2010; 102: 930 l Cuscineri K, int J Cancer 1998, 83: 66 l 1907 l Cuscineri K, Br J Cancer 2010; 103: 209 l Mesher D, J Clin Pathol 2015; 68: 135 l Powell N, Int J Cancer 2009; 125: 2425 l Tawfik El-Mansi M, Int J Gynecol Cancer 2006; 16: 1025

Based on meta-analysis performed by IARC's Infections and Cancer Epidemiology Group up to November 2011, the ICO HPV Information Centre has updated data until June 2015. Reference publications: 1) Guan P, Int J Cancer 2012;131:2349 2) Li N, Int J Cancer 2011;128:927 3) Smith JS, Int J Cancer 2007;121:621 4) Clifford GM, Br J Cancer 2003;88:63 5) Clifford GM, Br J Cancer 2003;89:101.

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a Number of women tested b 95% Confidence Interval

# 4.1.3 HPV type distribution among HIV+ women with normal cervical cytology

Table 24: Studies on HPV prevalence among HIV+ women with normal cytology in United Kingdom of Great Britain and Northern Ireland

HPV Prevalence							
Study <sup>1</sup>	HPV detection method and targeted HPV types	No. Tested <sup>a</sup>	%	(95% CI) <sup>b</sup>	Prevalence of 5 most frequent HPVs, HPV type (%)		
Cubie 2000	HC2 (HPV 6, 11, 16, 18, 31, 33, 35, 39, 42, 43, 44, 45, 51, 52, 56, 58, 59, 68), No genotyping	44	25.0	(13.2-40.3)			

#### Data updated on 31 Dec 2011 (data as of 31 Dec 2011)

 $DBH: Dot\ Blot\ Hybridization;\ EIA:\ Enzyme\ Immuno Assay;\ HC2:\ Hybrid\ Capture\ 2;\ PCR:\ Polymerase\ Chain\ Reaction;\ TS:\ Type\ Specific Capture\ 2;\ PCR:\ Polymerase\ Chain\ Reaction;\ TS:\ Type\ Specific Capture\ 2;\ PCR:\ Polymerase\ Chain\ Reaction;\ TS:\ Type\ Specific\ Capture\ 2;\ PCR:\ Polymerase\ Chain\ Reaction;\ TS:\ Type\ Specific\ Capture\ 2;\ PCR:\ Polymerase\ Chain\ Reaction;\ TS:\ Type\ Specific\ Capture\ 2;\ PCR:\ Polymerase\ Chain\ Reaction;\ TS:\ Type\ Specific\ Capture\ 2;\ PCR:\ Polymerase\ Chain\ Reaction;\ TS:\ Type\ Specific\ Capture\ 2;\ PCR:\ Polymerase\ Chain\ Reaction;\ TS:\ Type\ Specific\ Capture\ 2;\ PCR:\ Polymerase\ Chain\ Reaction;\ TS:\ Type\ Specific\ Capture\ 2;\ PCR:\ Polymerase\ Chain\ Reaction;\ TS:\ Type\ Specific\ Capture\ 2;\ PCR:\ Polymerase\ Chain\ Reaction;\ TS:\ Type\ Specific\ Capture\ 2;\ PCR:\ Polymerase\ Chain\ Reaction;\ TS:\ Type\ Specific\ Capture\ 2;\ PCR:\ Polymerase\ Chain\ Reaction;\ TS:\ Type\ Specific\ Capture\ 2;\ PCR:\ Polymerase\ Chain\ Reaction;\ TS:\ Type\ Specific\ Capture\ 2;\ PCR:\ Polymerase\ Chain\ Reaction;\ TS:\ Type\ Specific\ Capture\ 2;\ PCR:\ Polymerase\ 2;\ PCR:\ PCR:\$ 

Data Sources:

Systematic review and meta-analysis were performed by the ICO HPV Information Centre up to December 2011. Selected studies had to include at least 20 HIV positive women who had both normal cervical cytology and HPV test results (PCR or HC2).

<sup>&</sup>lt;sup>1</sup> Cubie HA, Sex Transm Infect 2000;76:257

# 4.1.4 Terminology

# Cytologically normal women

No abnormal cells are observed on the surface of their cervix upon cytology.

# Cervical Intraepithelial Neoplasia (CIN) / Squamous Intraepithelial Lesions (SIL)

SIL and CIN are two commonly used terms to describe precancerous lesions or the abnormal growth of squamous cells observed in the cervix. SIL is an abnormal result derived from cervical cytological screening or Pap smear testing. CIN is a histological diagnosis made upon analysis of cervical tissue obtained by biopsy or surgical excision. The condition is graded as CIN 1, 2 or 3, according to the thickness of the abnormal epithelium (1/3, 2/3 or the entire thickness).

# Low-grade cervical lesions (LSIL/CIN-1)

Low-grade cervical lesions are defined by early changes in size, shape, and number of abnormal cells formed on the surface of the cervix and may be referred to as mild dysplasia, LSIL, or CIN-1.

# High-grade cervical lesions (HSIL/CIN-2/CIN-3/CIS)

High-grade cervical lesions are defined by a large number of precancerous cells on the surface of the cervix that are distinctly different from normal cells. They have the potential to become cancerous cells and invade deeper tissues of the cervix. These lesions may be referred to as moderate or severe dysplasia, HSIL, CIN-2, CIN-3 or cervical carcinoma in situ (CIS).

# Carcinoma in situ (CIS)

Preinvasive malignancy limited to the epithelium without invasion of the basement membrane. CIN 3 encompasses the squamous carcinoma in situ.

# Invasive cervical cancer (ICC) / Cervical cancer

If the high-grade precancerous cells invade the basement membrane is called ICC. ICC stages range from stage I (cancer is in the cervix or uterus only) to stage IV (the cancer has spread to distant organs, such as the liver).

# Invasive squamous cell carcinoma

Invasive carcinoma composed of cells resembling those of squamous epithelium.

# Adenocarcinoma

Invasive tumour with glandular and squamous elements intermingled.

# 4.2 HPV burden in anogenital cancers other than cervix

Methods: Prevalence and type distribution of human papillomavirus in carcinoma of the vulva, vagina, anus and penis: systematic review and meta-analysis

A systematic review of the literature was conducted on the worldwide HPV-prevalence and type distribution for anogenital carcinomas other than cervix from January 1986 to 'data as of' indicated in each section. The search terms for the review were 'HPV' AND (anus OR anal) OR (penile) OR vagin\* OR vulv\* using Pubmed. There were no limits in publication language. References cited in selected articles were also investigated. Inclusion criteria were: HPV DNA detection by means of PCR, a minimum of 10 cases by lesion and a detailed description of HPV DNA detection and genotyping techniques used. The number of cases tested and HPV positive cases were extracted for each study to estimate the prevalence of HPV DNA and the HPV type distribution. Binomial 95% confidence intervals were calculated for each HPV prevalence.

# 4.2.1 Anal cancer and precancerous anal lesions

Anal cancer is similar to cervical cancer with respect to overall HPV DNA positivity, with approximately 100% of anal squamous cell carcinoma cases associated with HPV infection worldwide (de Martel C et al. Lancet Glob Health 2020;8(2):e180-e190). HPV16 is the most common type detected, representing 73% of all HPV-positive tumours. HPV18 is the second most common type detected and is found in approximately 5% of cases. HPV DNA is also detected in the majority of precancerous anal lesions (AIN) (91.5% in AIN1 and 93.9% in AIN2/3) (De Vuyst H et al. Int J Cancer 2009; 124: 1626-36). In this section, the burden of HPV among cases of anal cancers and precancerous anal lesions in United Kingdom of Great Britain and Northern Ireland are presented.

Table 25: Studies on HPV prevalence among anal cancer cases in United Kingdom of Great Britain and Northern Ireland (male and female)

		HPV Prevalence					
Study	HPV detection method and targeted HPV types	No. Tested	%	(95% CI) <sup>a</sup>	Prevalence of 5 most frequent HPVs, HPV type (%)		
Alemany 2015 <sup>b</sup>	PCR-SPF10, EIA, (HPV 6, 11, 16, 18, 26, 30, 31, 33, 34, 35, 39, 40, 42, 43, 44, 45, 51, 52, 53, 54, 56, 58, 59, 61, 66, 67, 68, 69, 70, 73, 74, 82, 83, 87, 89, 91)	169	87.6	(81.8-91.7)	HPV 16 (73.4), HPV 18 (3.6), HPV 6 (3.6), HPV 11 (3.0), HPV 33 (2.4)		
Baricevic 2015	PCR-L1C1/C2, PCR L1-Consensus primer, PCR-E6, PCR-E7, PCR- MULTIPLEX (HPV 6, 11, 16, 18, 31, 33, 45, 52, 58)	151	95.4	(90.7-97.7)	HPV 16 (88.7), HPV 6 (11.9), HPV 33 (6.6), HPV 18 (4.6), HPV 58 (4.6)		

#### Data updated on 30 Jun 2015 (data as of 30 Jun 2015)

DBH: Dot Blot Hybridization; EIA: Enzyme ImmunoAssay; HC2: Hybrid Capture 2; ISH: In Situ Hybridization; LBA: Line-Blot Assay; LiPA: Line Probe Assay; PCR: Polymerase Chain Reaction; RFLP: Restriction Fragment Length Polymorphism; RLBH: Reverse Line Blot Hybridization; RF-PCR: Real Time Polymerase Chain Reaction; SBH: Southern Blot Hybridization; SFF: Short Primer Fragment; TS: Type Specific;

a 95% Confidence Interval

Alemany L, Int J Cancer 2015; 136: 98 | Baricevic I, Eur J Cancer 2015; 51: 776

Based on systematic reviews (up to 2008) performed by ICO for the IARC Monograph on the Evaluation of Carcinogenic Risks to Humans volume 100B and IARC's Infections and Cancer Epidemiology Group. The ICO HPV Information Centre has updated data until June 2015. Reference publications: 1) Bouvard V, Lancet Oncol 2009;10:321 2) De Vuyst H, Int J Cancer 2009;124:1626

Table 26: Studies on HPV prevalence among cases of AIN2/3 in United Kingdom of Great Britain and Northern Ireland

			HPV		
Study	HPV detection method and targeted HPV types	No. Tested	%	(95% CI) <sup>a</sup>	Prevalence of 5 most frequent HPVs, HPV type (%)
Alemany 2015 <sup>b</sup>	PCR-SPF10, EIA, (HPV 6, 11, 16, 18, 26, 30, 31, 33, 34, 35, 39, 40, 42, 43, 44, 45, 51, 52, 53, 54, 56, 58, 59, 61, 66, 67, 68, 69, 70, 73, 74, 82, 83, 87, 89, 91)	23	95.7	(79.0-99.2)	HPV 16 (65.2), HPV 18 (8.7), HPV 51 (8.7), HPV 6 (8.7), HPV 74 (8.7)
Fox 2005 <sup>c</sup>	, PCR-MY09/11, (HPV 16, 18, 31, 33, 35, 39, 45, 51, 52, 56, 58, 59)	74	97.3	(90.7-99.3)	HPV 16 (64.9), HPV 18 (25.7), HPV 33 (24.3), HPV 58 (21.6), HPV 31 (18.9)

# Data updated on 30 Jun 2015 (data as of 30 Jun 2015)

DBH: Dot Blot Hybridization; EIA: Enzyme ImmunoAssay; HC2: Hybrid Capture 2; ISH: In Situ Hybridization; LBA: Line-Blot Assay; LiPA: Line Probe Assay; PCR: Polymerase Chain Reaction; RFLP: Restriction Fragment Length Polymorphism; RLBH: Reverse Line Blot Hybridization; RT-PCR: Real Time Polymerase Chain Reaction; SBH: Southern Blot Hybridization; SPF: Short Primer Fragment; TS: Type Specific;

AIN 2/3: Anal intraepithelial neoplasia of grade 2/3  $^a$  95% Confidence Interval

c HIV positive cases

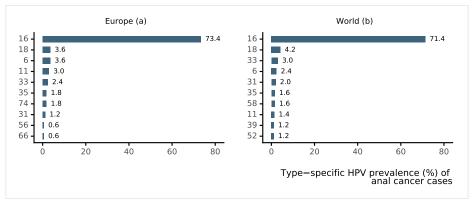
Alemany L, Int J Cancer 2015; 136: 98 | Fox PA, Sex Transm Infect 2005; 81: 142

Based on systematic reviews (up to 2008) performed by ICO for the IARC Monograph on the Evaluation of Carcinogenic Risks to Humans volume 100B and IARC's Infections and Cancer Epidemiology Group. The ICO HPV Information Centre has updated data until June 2015. Reference publications: 1) Bouvard V, Lancet Oncol 2009;10:321 2) De Vuyst H, Int J Cancer 2009;124:1626

b Includes cases from Bosnia-Herzegovina, Czech Republic, France, Germany, Poland, Portugal, Slovenia, Spain and United Kingdom

b Includes cases from Bosnia-Herzegovina, Czech Republic, France, Germany, Poland, Portugal, Slovenia, Spain and United Kingdom

Figure 62: Comparison of the ten most frequent HPV types in anal cancer cases in Europe and the World

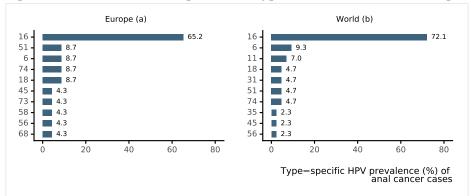


## Data updated on 9 Feb 2017 (data as of 30 Jun 2014)

<sup>a</sup> Includes cases from Bosnia-Herzegovina, Czech Republic, France, Germany, Poland, Portugal, Slovenia, Spain and United Kingdom

Data from Alemany L, Int J Cancer 2015; 136: 98. This study has gathered the largest international series of anal cancer cases and precancerous lesions worldwide using a standard protocol with a highly sensitive HPV DNA detection assay.

Figure 63: Comparison of the ten most frequent HPV types in AIN 2/3 cases in Europe and the World



# Data updated on 7 Feb 2017 (data as of 30 Jun 2014)

AIN 2/3: Anal intraepithelial neoplasia of grade 2/3

Data Sources:
Data from Alemany L, Int J Cancer 2015; 136: 98. This study has gathered the largest international series of anal cancer cases and precancerous lesions worldwide using a standard protocol with a highly sensitive HPV DNA detection assay

b Includes cases from Europe (Bosnia-Herzegovina, Czech Republic, France, Germany, Poland, Portugal, Slovenia, Spain and United Kingdom); America (Chile, Colombia, Ecuador, Guatemala, Honduras, Mexico, Paraguay and United States); Africa (Mali, Nigeria and Senegal); Asia (Bangladesh, India and South Korea) Data Sources:

Includes cases from Bosnia-Herzegovina, Czech Republic, France, Germany, Poland, Portugal, Slovenia, Spain and United Kingdom

b Includes cases from Europe (Bosnia-Herzegovina, Czech Republic, France, Germany, Poland, Portugal, Slovenia, Spain and United Kingdom); America (Chile, Colombia, Ecuador, Guatemala, Honduras, Mexico, Paraguay)

# 4.2.2 Vulvar cancer and precancerous vulvar lesions

HPV attribution for vulvar cancer is 48% among age 15-54 years, 28% among age 55-64 years, and 15% among age 65+ worldwide (de Martel C et al. Lancet Glob Health 2020;8(2):e180-e190). Vulvar cancer has two distinct histological patterns with two different risk factor profiles: (1) basaloid/warty types (2) keratinising types. Basaloid/warty lesions are more common in young women, are frequently found adjacent to VIN, are very often associated with HPV DNA detection (86%), and have a similar risk factor profile as cervical cancer. Keratinising vulvar carcinomas represent the majority of the vulvar lesions (>60%). These lesions develop from non HPV-related chronic vulvar dermatoses, especially lichen sclerosus and/or squamous hyperplasia, their immediate cancer precursor lesion is differentiated VIN, they occur more often in older women, and are rarely associated with HPV (6%) or with any of the other risk factors typical of cervical cancer. HPV prevalence is frequently detected among cases of high-grade VIN (VIN2/3) (85.3%). HPV 16 is the most common type detected followed by HPV 33 (De Vuyst H et al. Int J Cancer 2009; 124: 1626-36). In this section, the HPV burden among cases of vulvar cancer cases and precancerous vulvar lesions in United Kingdom of Great Britain and Northern Ireland are presented.

Table 27: Studies on HPV prevalence among vulvar cancer cases in United Kingdom of Great Britain and Northern Ireland

		HPV Prevalence				
Study	HPV detection method and targeted HPV types	No. Tested	%	(95% CI) <sup>a</sup>	Prevalence of 5 most frequent HPVs, HPV type (%)	
Abdel-Hady 2001	TS (HPV 6, 11, 16, 18, 31, 33)	11	27.3	(9.7-56.6)	HPV 16 (27.3), HPV 33 (18.2), HPV 18 (9.1)	
de Sanjosé 2013 <sup>b</sup>	PCR-SPF10, EIA, (HPV 6, 11, 16, 18, 26, 30, 31, 33, 34, 35, 39, 40, 42, 43, 44, 45, 51, 52, 53, 54, 56, 58, 59, 61, 66, 67, 68, 69, 70, 73, 74, 82, 83, 87, 89, 91)	903	19.3	(16.8-22.0)	HPV 16 (13.8), HPV 33 (1.2), HPV 18 (0.6), HPV 31 (0.6), HPV 44 (0.4)	

Data updated on 30 Jun 2015 (data as of 30 Jun 2015)

DBH: Dot Blot Hybridization; EIA: Enzyme ImmunoAssay; HC2: Hybrid Capture 2; ISH: In Situ Hybridization; LBA: Line-Blot Assay; LiPA: Line Probe Assay; PCR: Polymerase Chain Reaction; RFLP: Restriction Fragment Length Polymorphism; RLBH: Reverse Line Blot Hybridization; RF-PCR: Real Time Polymerase Chain Reaction; SBH: Southern Blot Hybridization; SFF: Short Primer Fragment; TS: Type Specific;

a 95% Confidence Interval

<u>Data Sources</u>:

Abdel-Hady ES, Cancer Res 2001: 61: 192 | de Saniosé S, Eur J Cancer 2013: 49: 3450

Based on systematic reviews (up to 2008) performed by ICO for the IARC Monograph on the Evaluation of Carcinogenic Risks to Humans volume 100B and IARC's Infections and Cancer Epidemiology Group. The ICO HPV Information Centre has updated data until June 2015. Reference publications: 1) Bouvard V, Lancet Oncol 2009;10:321 2) De Vuyst H, Int J Cancer 2009;124:1636

Table 28: Studies on HPV prevalence among VIN 2/3 cases in United Kingdom of Great Britain and Northern Ireland

			HPV	Prevalence	
Study	HPV detection method and targeted HPV types	No. Tested	%	(95% CI) <sup>a</sup>	Prevalence of 5 most frequent HPVs, HPV type (%)
Abdel-Hady 2001	TS (HPV 06/11, 16, 18, 31, 33)	32	71.9	(54.6-84.4)	HPV 16 (62.5), HPV 6/11 (18.8), HPV 31 (3.1), HPV 33 (3.1)
Baldwin 2003	PCR L1-Consensus primer, Sequencing (HPV 6, 11, 16, 18, 31, 33)	11	100	(74.1-100.0)	HPV 16 (90.9), HPV 33 (9.1)
Bryant 2011	PCR- MULTIPLEX (HPV 6, 11, 16, 18, 31, 33, 35, 40, 42, 43, 44, 45, 51, 52, 53, 56, 58, 59, 66, 73)	49	81.6	(68.6-90.0)	HPV 16 (67.3), HPV 33 (16.3), HPV 6 (10.2), HPV 18 (2.0), HPV 31 (2.0)
Daayana 2010	EIA, (HPV 6, 11, 16, 26, 31, 33, 34, 35, 39, 40, 42, 43, 44, 45, 51, 52, 53, 54, 56, 57, 58, 59, 61, 66, 68, 70, 71, 72, 73, 81, 82, 83, 84)	19	78.9	(56.7-91.5)	HPV 16 (73.7), HPV 33 (5.3), HPV 42 (5.3), HPV 84 (5.3)
de Sanjosé 2013 <sup>b</sup>	PCR-SPF10, EIA, (HPV 6, 11, 16, 18, 26, 30, 31, 33, 34, 35, 39, 40, 42, 43, 44, 45, 51, 52, 53, 54, 56, 58, 59, 61, 66, 67, 68, 69, 70, 73, 74, 82, 83, 87, 89, 91)	312	86.9	(82.7-90.2)	HPV 16 (69.6), HPV 33 (11.2), HPV 18 (2.2), HPV 6 (1.6), HPV 52 (1.3)

Continued on next page

<sup>50 %</sup> Collinearie: Interval

Includes cases from Austria, Belarus, Bosnia-Herzegovina, Czech Republic, France, Germany, Greece, Italy, Poland, Portugal, Spain and United Kingdom

Table 28 - continued from previous page

Prevalence						
Study	HPV detection method and targeted HPV types	No. Tested	%	(95% CI) <sup>a</sup>	Prevalence of 5 most frequent HPVs, HPV type (%)	
Winters 2008	EIA, (HPV 6, 11, 16, 18, 26, 31, 33, 34, 35, 39, 40, 42, 43, 44, 45, 51, 52, 53, 54, 56, 57, 58, 59, 61, 66, 68, 70, 71, 72, 73, 81, 82, 83, 84)	20	85	(64.0-94.8)	HPV 16 (75.0), HPV 18 (5.0), HPV 33 (5.0)	

# Data updated on 30 Jun 2015 (data as of 30 Jun 2015)

DBH: Dot Blot Hybridization; EIA: Enzyme ImmunoAssay; HC2: Hybrid Capture 2; ISH: In Situ Hybridization; LBA: Line-Blot Assay; LiPA: Line Probe Assay; PCR: Polymerase Chain Reaction; RFLP: Restriction Fragment Length Polymorphism; RLBH: Reverse Line Blot Hybridization; RT-PCR: Real Time Polymerase Chain Reaction; SBH: Southern Blot Hybridization; SPF: Short Primer Fragment; TS: Type Specific;
VIN 2/3: Vulvar intraepithelial neoplasia of grade 2/3

a 95% Confidence Interval

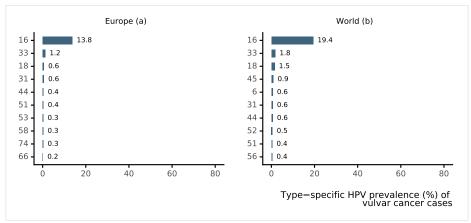
b Includes cases from Austria, Belarus, Bosnia-Herzegovina, Czech Republic, France, Germany, Greece, Italy, Poland, Portugal, Spain and United Kingdom

<u>Data Sources:</u>
Abdel-Hady ES, Cancer Res 2001; 61: 192 | Baldwin PJ, Clin Cancer Res 2003; 9: 5205 | Bryant D, J Med Virol 2011; 83: 1358 | Daayana S, Br J Cancer 2010; 102: 1129 | de Sanjosé S,

Eur J Cancer 2013; 49: 3450 | Winters U, Clin Cancer Res 2008; 14: 5292

Based on systematic reviews (up to 2008) performed by ICO for the IARC Monograph on the Evaluation of Carcinogenic Risks to Humans volume 100B and IARC's Infections and Cancer Epidemiology Group. The ICO HPV Information Centre has updated data until June 2015. Reference publications: 1) Bouvard V, Lancet Oncol 2009;10:321 2) De Vuyst H, Int J Cancer 2009;124:1626

Figure 64: Comparison of the ten most frequent HPV types in cases of vulvar cancer in Europe and the World



## Data updated on 30 Jun 2015 (data as of 30 Jun 2015)

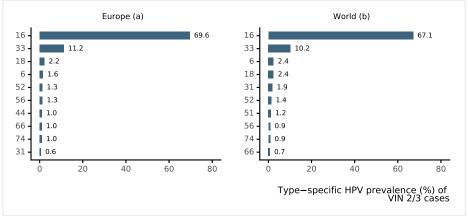
<sup>a</sup> Includes cases from Austria, Belarus, Bosnia-Herzegovina, Czech Republic, France, Germany, Greece, Italy, Poland, Portugal, Spain and United Kingdom.

b Includes cases from America (Argentina, Brazil, Chile, Colombia, Ecuador, Guatemala, Honduras, Mexico, Paraguay, Uruguay, United States of America and Venezuela); Africa (Mali, Mozambique, Nigeria, and Senegal); Oceania (Australia and New Zealand); Europe (Austria, Belarus, Bosnia-Herzegovina, Czech Republic, France, Germany, Greece, Italy, Poland, Portugal, Spain and United Kingdom); and in Asia (Bangladesh, India, Israel, South Korea, Kuwait, Lebanon, Philippines, Taiwan and Turkey)
Data Sources:

Data Sources:

Data from de Sanjosé S, Eur J Cancer 2013; 49: 3450. This study has gathered the largest international series of vulva cancer cases and precancerous lesions worldwide using a standard protocol with a highly sensitive HPV DNA detection assay.

Figure 65: Comparison of the ten most frequent HPV types in VIN 2/3 cases in Europe and the World



# Data updated on 30 Jun 2014 (data as of 30 Jun 2014)

VIN 2/3: Vulvar intraepithelial neoplasia of grade 2/3

a Includes cases from Austria, Belarus, Bosnia-Herzegovina, Czech Republic, France, Germany, Greece, Italy, Poland, Portugal, Spain and United Kingdom.

b Includes cases from America (Argentina, Brazil, Chile, Colombia, Ecuador, Guatemala, Honduras, Mexico, Paraguay, Uruguay and Venezuela); Oceania (Australia and New Zealand); Europe (Austria, Belarus, Bosnia-Herzegovina, Czech Republic, France, Germany, Greece, Italy, Poland, Portugal, Spain and United Kingdom); and in Asia (Bangladesh, India, Israel, South Korea, Kuwait, Lebanon, Philippines, Taiwan and Turkey)
Data Sources:

Data from de Sanjosé S, Eur J Cancer 2013; 49: 3450. This study has gathered the largest international series of vulva cancer cases and precancerous lesions worldwide using a standard protocol with a highly sensitive HPV DNA detection assay.

# Vaginal cancer and precancerous vaginal lesions

Vaginal and cervical cancers share similar risk factors and it is generally accepted that both carcinomas share the same aetiology of HPV infection although there is limited evidence available. Women with vaginal cancer are more likely to have a history of other ano-genital cancers, particularly of the cervix, and these two carcinomas are frequently diagnosed simultaneously. HPV DNA is detected among 78% of invasive vaginal carcinomas and 91% of high-grade vaginal neoplasias (VaIN2/3). HPV16 is the most common type in high-grade vaginal neoplasias and it is detected in at least 78% of HPV-positive carcinomas (de Martel C et al. Lancet Glob Health 2020;8(2):e180-e190; De Vuyst H et al. Int J Cancer 2009; 124:1626-36). In this section, the HPV burden among cases of vaginal cancer cases and precancerous vaginal lesions in United Kingdom of Great Britain and Northern Ireland are presented.

Table 29: Studies on HPV prevalence among vaginal cancer cases in United Kingdom of Great Britain and Northern Ireland

HPV Prevalence					
Study <sup>b</sup>	HPV detection method and targeted HPV types	No. Tested	%	(95% CI) <sup>a</sup>	Prevalence of 5 most frequent HPVs, HPV type (%)
Alemany 2014	PCR-SPF10, EIA, (HPV 6, 11, 16, 18, 26, 30, 31, 33, 35, 39, 42, 45, 51, 52, 53, 56, 58, 59, 66, 67, 68, 69, 73, 82)	152	71.1	(63.4-77.7)	HPV 16 (47.4), HPV 18 (3.3), HPV 73 (3.3), HPV 33 (2.6), HPV 56 (2.6)

### Data updated on 30 Jun 2015 (data as of 30 Jun 2015)

DBH: Dot Blot Hybridization; EIA: Enzyme ImmunoAssay; HC2: Hybrid Capture 2; ISH: In Situ Hybridization; LBA: Line-Blot Assay; LiPA: Line Probe Assay; PCR: Polymerase Chain Reaction; RFLP: Restriction Fragment Length Polymorphism; RLBH: Reverse Line Blot Hybridization; RT-PCR: Real Time Polymerase Chain Reaction; SBH: Southern Blot Hybridization; SPF: Short Primer Fragment; TS: Type Specific;

Data Sources

Alemany L, Eur J Cancer 2014; 50: 2846

Based on systematic reviews (up to 2008) performed by ICO for the IARC Monograph on the Evaluation of Carcinogenic Risks to Humans volume 100B and IARC's Infections and Cancer Epidemiology Group. The ICO HPV Information Centre has updated data until June 2015. Reference publications: 1) Bouvard V, Lancet Oncol 2009;10:321 2) De Vuyst H, Int J Cancer 2009;124:1626

Table 30: Studies on HPV prevalence among VaIN 2/3 cases in United Kingdom of Great Britain and Northern Ireland

HPV Prevalence					
Study	HPV detection method and targeted HPV types	No. Tested	%	(95% CI) <sup>a</sup>	Prevalence of 5 most frequent HPVs, HPV type (%)
Alemany 2014	PCR-SPF10, EIA, (HPV 6, 11, 16, 18, 26, 30, 31, 33, 35, 39, 42, 45, 51, 52, 53, 56, 58, 59, 66, 67, 68, 69, 73, 82)	96	97.9	(92.7-99.4)	HPV 16 (65.6), HPV 33 (7.3), HPV 18 (5.2), HPV 52 (3.1), HPV 73 (3.1)

# Data updated on 30 Jun 2015 (data as of 30 Jun 2015)

DBH: Dot Blot Hybridization; EIA: Enzyme ImmunoAssay; HC2: Hybrid Capture 2; ISH: In Situ Hybridization; LBA: Line-Blot Assay; LiPA: Line Probe Assay; PCR: Polymerase Chain Reaction; RFLP: Restriction Fragment Length Polymorphism; RLBH: Reverse Line Blot Hybridization; RT-PCR: Real Time Polymerase Chain Reaction; SBH: Southern Blot Hybridization; SPF: Short Primer Fragment; TS: Type Specific;

VAIN 2/3: Vaginal intraepithelial neoplasia of grade 2/3

 $^a$  95% Confidence Interval

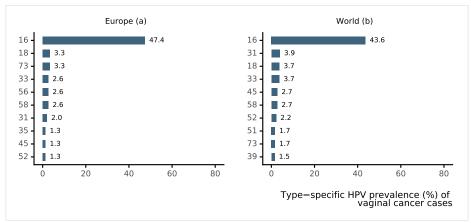
Data Sources

Alemany L. Eur J Cancer 2014; 50: 2846

Based on systematic reviews (up to 2008) performed by ICO for the IARC Monograph on the Evaluation of Carcinogenic Risks to Humans volume 100B and IARC's Infections and Cancer Epidemiology Group. The ICO HPV Information Centre has updated data until June 2015. Reference publications: 1) Bouvard V, Lancet Oncol 2009;10:321 2) De Vuyst H, Int J Cancer

<sup>95%</sup> Confidence Interval b Includes cases from Austria, Belarus, Czech Republic, France, Germany, Greece, Poland, Spain and United Kingdom

Figure 66: Comparison of the ten most frequent HPV types in cases of vaginal cancer in Europe and the World

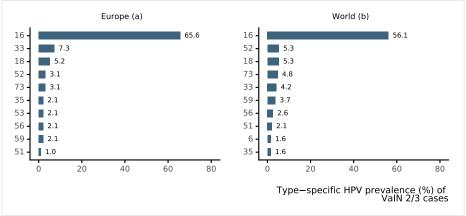


## Data updated on 30 Jun 2015 (data as of 30 Jun 2015)

a Includes cases from Austria, Belarus, Czech Republic, France, Germany, Greece, Poland, Spain and United Kingdom.

Data Sources:
Data from Alemany L, Eur J Cancer 2014; 50: 2846. This study has gathered the largest international series of vaginal cancer cases and precancerous lesions worldwide using a standard protocol with a highly sensitive HPV DNA detection assay.

Figure 67: Comparison of the ten most frequent HPV types in VaIN 2/3 cases in Europe and the World



# Data updated on 30 Jun 2014 (data as of 30 Jun 2014)

VAIN 2/3: Vaginal intraepithelial neoplasia of grade 2/3

Data Sources

Data from Alemany L, Eur J Cancer 2014; 50: 2846. This study has gathered the largest international series of vaginal cancer cases and precancerous lesions worldwide using a standard protocol with a highly sensitive HPV DNA detection assay.

b Includes cases from Europe (Austria, Belarus, Czech Republic, France, Germany, Greece, Poland, Spain and United Kingdom); America (Argentina, Brazil, Chile, Colombia, Ecuador, Guatemala, Mexico, Paraguay, Uruguay, United states of America and Venezuela); Africa (Mozambique, Nigeria); Asia (Bangladesh, India, Israel, South Korea, Kuwait, Philippines, Taiwan and Turkey); and Oceania (Australia)

a Includes cases from Austria, Belarus, Czech Republic, France, Germany, Greece, Poland, Spain and United Kingdom.

b Includes cases from Europe (Austria, Belarus, Czech Republic, France, Germany, Greece, Poland, Spain and United Kingdom); America (Argentina, Brazil, Chile, Colombia, Ecuador, Guatemala, Mexico, Paraguay, Uruguay, United states of America and Venezuela); Asia (Bangladesh, India, Israel, South Korea, Kuwait, Philippines, Taiwan and Turkey); and Oceania

# 4.2.4 Penile cancer and precancerous penile lesions

HPV DNA is detectable in approximately 51% of all penile cancers (de Martel C et al. Lancet Glob Health 2020;8(2):e180-e190). Among HPV-related penile tumours, HPV16 is the most common type detected, followed by HPV18 and HPV types 6/11 (Miralles C et al. J Clin Pathol 2009;62:870-8). Over 95% of invasive penile cancers are SCC and the most common penile SCC histologic sub-types are keratinising (49%), mixed warty-basaloid (17%), verrucous (8%), warty (6%), and basaloid (4%). HPV is commonly detected in basaloid and warty tumours but is less common in keratinising and verrucous tumours. In this section, the HPV burden among cases of penile cancer cases and precancerous penile lesions in United Kingdom of Great Britain and Northern Ireland are presented.

Table 31: Studies on HPV prevalence among penile cancer cases in United Kingdom of Great Britain and Northern Ireland

Study	HPV detection method and targeted HPV types	No. Tested	%	(95% CI) <sup>a</sup>	Prevalence of 5 most frequent HPVs, HPV type (%)
Prowse 2008	PCR-SPF10, LIPA, (HPV 6, 11, 16, 18, 26, 31, 33-35, 39, 40, 42-45, 51-54, 56, 58, 59, 66, 68-71, 73, 74)	26	53.8	(35.5-71.2)	
Stankiewicz 2011	PCR-SPF10, LIPA, (HPV 6, 11, 16, 18, 26, 31, 33-35, 39, 40, 42-45, 51-54, 56, 58, 59, 66, 68-71, 73, 74)	102	55.9	(46.2-65.1)	HPV 16 (45.1), HPV 11 (9.8), HPV 45 (5.9), HPV 6 (5.9), HPV 31 (4.9)

#### Data updated on 5 Mar 2015 (data as of 30 Jun 2014)

DBH: Dot Blot Hybridization; EIA: Enzyme ImmunoAssay; HC2: Hybrid Capture 2; ISH: In Situ Hybridization; LBA: Line-Blot Assay; LiPA: Line Probe Assay; PCR: Polymerase Chain Reaction; RFLP: Restriction Fragment Length Polymorphism; RLBH: Reverse Line Blot Hybridization; RT-PCR: Real Time Polymerase Chain Reaction; SBH: Southern Blot Hybridization; SPF: Short Primer Fragment; TS: Type Specific;

a 95% Confidence Interval

Data Sources

Prowse DM, Br J Dermatol 2008; 158: 261 | Stankiewicz E, Histopathology 2011; 58: 433
The ICO HPV Information Centre has updated data until June 2014. Reference publications (up to 2008): 1) Bouvard V, Lancet Oncol 2009;10:321 2) Miralles-Guri C,J Clin Pathol

Table 32: Studies on HPV prevalence among PeIN 2/3 cases in United Kingdom of Great Britain and Northern Ireland

	HPV Prevalence					
Study	HPV detection method and targeted HPV types	No. Tested	%	(95% CI) <sup>a</sup>	Prevalence of 5 most frequent HPVs, HPV type (%)	
No data available	-	-	-	-		

# Data updated on 10 Feb 2015 (data as of 30 Jun 2014)

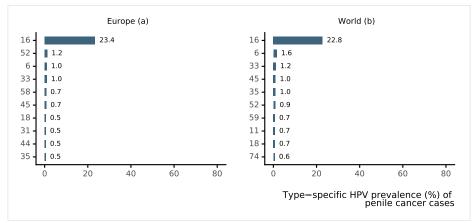
PeIN 2/3: Penile intraepithelial neoplasia of grade 2/3

DBH: Dot Blot Hybridization: EIA: Enzyme ImmunoAssay: HC2: Hybrid Capture 2: ISH: In Situ Hybridization: LBA: Line-Blot Assay: LiPA: Line Probe Assay: PCR: Polymerase Chain Reaction; RFLP: Restriction Fragment Length Polymorphism; RLBH: Reverse Line Blot Hybridization; RT-PCR: Real Time Polymerase Chain Reaction; SBH: Southern Blot Hybridization; SPF: Short Primer Fragment: TS: Type Specific:

a 95% Confidence Interval

The ICO HPV Information Centre has updated data until June 2014. Reference publication (up to 2008): Bouvard V, Lancet Oncol 2009;10:321

Figure 68: Comparison of the ten most frequent HPV types in cases of penile cancer in Europe and the World



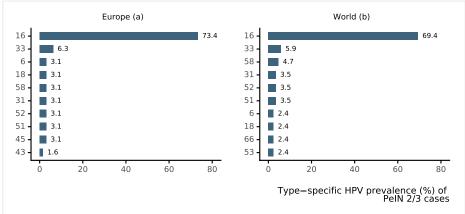
## Data updated on 9 Feb 2017 (data as of 30 Jun 2015)

<sup>a</sup> Includes cases from Czech Republic, France, Greece, Poland, Portugal, Spain and United Kingdom

b Includes cases from Australia, Bangladesh, India, South Korea, Lebanon, Philippines, Chile, Colombia, Ecuador, Guatemala, Honduras, Mexico, Paraguay, Venezuela and United States, Mozambique, Nigeria, Senegal, Czech Republic, France, Greece, Poland, Portugal, Spain and United Kingdo

Alemany L, Eur Urol 2016; 69: 953

Figure 69: Comparison of the ten most frequent HPV types in PeIN 2/3 cases in Europe and the World



# Data updated on 9 Feb 2017 (data as of 30 Jun 2015)

Pe<br/>IN 2/3: Penile intraepithelial neoplasia of grade 2/3<br/>  $\,$ 

Lopes V, Oral Oncol 2011; 47: 698 | Snijders PJ, Int J Cancer 1996; 66: 464 | Yeudall WA, J Gen Virol 1991; 72 ( Pt 1): 173

Based on systematic reviews and meta-analysis performed by ICO. Reference publications: 1) Ndiaye C, Lancet Oncol 2014; 15: 1319 2) Kreimer AR, Cancer Epidemiol Biomarkers Prev 2005: 14: 467

Includes cases from Czech Republic, France, Greece, Poland, Portugal, Spain and United Kingdom

b Includes cases from Australia, Bangladesh, India, South Korea, Lebanon, Philippines, Chile, Colombia, Ecuador, Guatemala, Honduras, Mexico, Paraguay, Venezuela, Mozambique, Nigeria, Senegal, Czech Republic, France, Greece, Poland, Portugal, Spain and United Kingdom.

# 4.3 HPV burden in men

The information to date regarding anogenital HPV infection is primarily derived from cross-sectional studies of selected populations such as general population, university students, military recruits, and studies that examined husbands of control women, as well as from prospective studies. Special subgroups include mainly studies that examined STD (sexually transmitted diseases) clinic attendees, MSM (men who have sex with men), HIV positive men, and partners of women with HPV lesions, CIN (cervical intraepithelial neoplasia), cervical cancer or cervical carcinoma in situ. Globally, prevalence of external genital HPV infection in men is higher than cervical HPV infection in women, but persistence is less likely. As with genital HPV prevalence, high numbers of sexual partners increase the acquisition of oncogenic HPV infections (Vaccine 2012, Vol. 30, Suppl 5). In this section, the HPV burden among men in United Kingdom of Great Britain and Northern Ireland is presented.

# **Methods**

HPV burden in men was based on published systematic reviews and meta-analyses (Dunne EF, J Infect Dis 2006; 194: 1044, Smith JS, J Adolesc Health 2011; 48: 540, Olesen TB, Sex Transm Infect 2014; 90: 455, and Hebnes JB, J Sex Med 2014; 11: 2630) up to October 31, 2015. The search terms for the review were human papillomavirus, men, polymerase chain reaction (PCR), hybrid capture (HC), and viral DNA. References cited in selected articles were also investigated. Inclusion criteria were: HPV DNA detection by means of PCR or HC (ISH if data are not available for the country), and a detailed description of HPV DNA detection and genotyping techniques used. The number of cases tested and HPV positive cases were extracted for each study to estimate the anogenital prevalence of HPV DNA. Binomial 95% confidence intervals were calculated for each anogenital HPV prevalence.

Table 33: Studies on HPV prevalence among men in United Kingdom of Great Britain and Northern Ireland

						HPV	Prevalence	
Study	Anatomic sites samples	HPV detection method	Population	Age (years)	No. Tested	%	(95% CI) <sup>a</sup>	
-	-	-	-	-	-	-	-	

Data updated on 31 Oct 2015 (data as of 31 Oct 2015)

HC2: Hybrid Capture 2; ISH: In Situ Hybridization; PCR: Polymerase Chain Reaction; RT-PCR: Real Time Polymerase Chain Reaction; SPF: Short Primer Fragment; TS: Type Specific; MSM: Men who have sex with men; MSW:Men who have sex with women; STD: sexually transmitted disease 95% Confidence Interval

Based on published systematic reviews, the ICO HPV Information Centre has updated data until October 2015. Reference publications: 1) Dunne EF, J Infect Dis 2006; 194: 1044 2) Smith JS, J Adolesc Health 2011; 48: 540 3) Olesen TB, Sex Transm Infect 2014; 90: 455 4) Hebnes JB, J Sex Med 2014; 11: 2630

Table 34: Studies on HPV prevalence among men from special subgroups in United Kingdom of Great Britain and Northern Ireland

						HPV	Prevalence
Study	Anatomic sites samples	HPV detection method	Population	Age (years)	No. Tested	%	(95% CI) <sup>a</sup>
Bissett 2011	Glans, prepuce, shaft, scrotum	PCR-General primers (GP5 + /6+), Bio-Plex array technology for typing	Genitourinary clinic attendees with multiple sexual partners or diagnosis of genital warts within 6 months	-	87	49.4	(38.5-60.4)
Cuschieri 2011	Shaft	PCR-INNO-LiPA	Drop-in sexual health service attendees	16-25	117	29.1	(21.0-38.2)
Hillman 1993	Urethra	PCR-GP5+/6+	Men infected with gonorrhea	17-55.6	100	18.0	(11.0-26.9)
Jalal 2007	Urethra	PCR-General primers for L1 (MY09/11, GP5 + /6+) and RLH	Genitourinary clinic attendees	15-77	437	20.8	(17.1-24.9)
King 2015	Anus	PCR-Multiplex and Bio-Plex Any nonavalent vaccine HPV types	MSM	Median 30 (IQR=25- 35)	454	40.1	(35.5-44.8)
						Continu	ed on next nage

Continued on next page

Table 34 - continued from previous page

						HPV I	Prevalence
Study	Anatomic sites samples	HPV detection method	Population	Age (years)	No. Tested	%	(95% CI) <sup>a</sup>
King 2015	Coronal sulcus, glans, penis shaft, scrotum and perianal area	PCR-Multiplex and Bio-Plex Any nonavalent vaccine HPV types	MSM	Median 30 (IQR=25- 35)	446	36.1	(31.6-40.7)
Lacey 1999	Anal canal	PCR-GP5+/6+	HIV+ MSM	19-62	57	84.2	(72.1-92.5)

# Data updated on 31 Oct 2015 (data as of 31 Oct 2015)

DBH: Dot Blot Hybridization; EIA: Enzyme ImmunoAssay; HC2: Hybrid Capture 2; LiPA: Line Probe Assay; PCR: Polymerase Chain Reaction; RFLP: Restriction Fragment Length Polymorphism; RLH: Reverse Line Hybridisation; RT-PCR: Real Time Polymerase Chain Reaction; SPF: Short Primer Fragment; TS: Type Specific; MSM: Men who have sex with men; MSW:Men who have sex with women; STD: sexually transmitted diseases

a 95% Confidence Interval

Data Sources:
Bissett SL, J Med Virol 2011; 83: 1744 | Cuschieri K, J Med Virol 2011; 83: 1983 | Hillman RJ, Genitourin Med 1993; 69: 187 | Jalal H, Int J STD AIDS 2007; 18: 617 | King EM, Br J

Cancer 2015; 112: 1585 | Lacey HB, Sex Transm Infect 1999; 75: 172
Based on published systematic reviews, the ICO HPV Information Centre has updated data until October 2015. Reference publications: 1) Dunne EF, J Infect Dis 2006; 194: 1044 2) Smith JS, J Adolesc Health 2011; 48: 540 3) Olesen TB, Sex Transm Infect 2014; 90: 455 4) Hebnes JB, J Sex Med 2014; 11: 2630.

# 4.4 HPV burden in the head and neck

The last evaluation of the International Agency for Research in Cancer (IARC) on the carcinogenicity of HPV in humans concluded that (a) there is enough evidence for the carcinogenicity of HPV type 16 in the oral cavity, oropharynx (including tonsil cancer, base of tongue cancer and other oropharyngeal cancer sites), and (b) limited evidence for laryngeal cancer (IARC Monograph Vol 100B). There is increasing evidence that HPV-related oropharyngeal cancers constitute an epidemiological, molecular and clinical distinct form as compared to non HPV-related ones. Some studies indicate that the most likely explanation for the origin of this distinct form of head and neck cancers associated with HPV is a sexually acquired oral HPV infection that is not cleared, persists and evolves into a neoplastic lesion. Around 30% of oropharyngeal cancers (which mainly comprises the tonsils and base of tongue sites) are caused by HPV with HPV16 being the most frequent type (de Martel C et al. Int J Cancer 2017;141(4):664-670). Attributable fraction varies greatly worldwide, being highest in more developed countries (60% in Republic of Korea, 51% in North America, 50% in Eastern Europe, 46% in Japan, 42% in North-Western Europe, 41% in Australia/New Zealand, 24% in South Europe, 23% in China, 22% in India, and 13% in elsewhere) (de Martel C et al. Lancet Glob Health 2020;8(2):e180-e190). In this section, the HPV burden in the head and neck in United Kingdom of Great Britain and Northern Ireland is presented.

# 4.4.1 Burden of oral HPV infection in healthy population

Table 35: Studies on oral HPV prevalence among healthy in United Kingdom of Great Britain and Northern Ireland

Study	Specimen collection method / anatomic site	HPV detection methoda	Population	% males	$\begin{array}{c} \textbf{Age} \\ (\textbf{years})^b \end{array}$	No. $\mathbf{tested}^c$	HPV prevalence % (95% CI)	High-Risk HPV prevalence % (95% CI)	$5 \text{ most}$ frequent HPVs, HPV type $(\mathbf{n})^d$
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# Data updated on 19 Oct 2021 (data as of 19 May 2015)

(95% CI): 95% Confidence Interval

 $^{a}$  TS: type-specific; RT-PCR: real-time PCR; qPCR: quantitative PCR

b NS: not specified

c number of cases tested for HPV DNA

d number of cases positive for the specific HPV-type

Data Sources:

Systematic review and meta-analysis was performed by ICO HPV Information Centre until May 19, 2015. Reference publication: Mena M et al. J Infect Dis 2019;219(10):1574-1585

# 4.4.2 HPV burden in head and neck cancers

Table 36: Studies on HPV prevalence among cases of oral cavity cancer in United Kingdom of Great Britain and Northern Ireland

	HPV Prevalence						
Study	HPV detection method and targeted HPV types	No. Tested	%	(95% CI) <sup>a</sup>	Prevalence of 5 most frequent HPVs, HPV type (%)		
MEN							
No data available	-	-	-	-	-		
WOMEN							
No data available	-	-	-	-	-		
BOTH OR UNSPECIFIE	CD CO						
Lopes 2011	GP5+/GP6+ (L1) and qPCR for 16/18 Hybridization with TS probes (16. 18)	142	3.5	(1.5-8.0)	HPV 16 (2.1) HPV 18 (2.1)		
Snijders 1996	GP5+/GP6+ (L1) Amplification with TS primers and SBH with TS probes (6. 11. 16. 18. 31. 33)	25	20	(8.9-39.1)	HPV 16 (20.0)		
Yeudall 1991	TS-PCR E6/E7 for 16. E6 for 18 and specific for 4 Hybridization with TS probes (4. 16. 18)	39	46.2	(31.6-61.4)	HPV 16 (25.6) HPV 18 (20.5)		

#### Data updated on 9 May 2016 (data as of 31 Dec 2015)

DBH: Dot Blot Hybridization; EIA: Enzyme ImmunoAssay; HC2: Hybrid Capture 2; ISH: In Situ Hybridization; LBA: Line-Blot Assay; LiPA: Line Probe Assay; PCR: Polymerase Chain Reaction; RFLP: Restriction Fragment Length Polymorphism; RLBH: Reverse Line Blot Hybridization; RT-PCR: Real Time Polymerase Chain Reaction; SBH: Southern Blot Hybridization; SPF: Short Primer Fragment; TS: Type Specific;

Only for European countries a 95% Confidence Interval

Data Sources:
Lopes V, Oral Oncol 2011; 47: 698 | Snijders PJ, Int J Cancer 1996; 66: 464 | Yeudall WA, J Gen Virol 1991; 72 ( Pt 1): 173

Based on systematic reviews and meta-analysis performed by ICO. Reference publications: 1) Ndiaye C, Lancet Oncol 2014; 15: 1319 2) Kreimer AR, Cancer Epidemiol Biomarkers Prev 2005; 14: 467

Table 37: Studies on HPV prevalence among cases of oropharyngeal cancer in United Kingdom of Great Britain and Northern Ireland

			HPV	Prevalence	
Study	HPV detection method and targeted HPV types	No. Tested	%	(95% CI) <sup>a</sup>	Prevalence of 5 most frequent HPVs, HPV type (%)
MEN					
No data available	-	-	-	-	-
WOMEN					
No data available	-	-	-	-	-
BOTH OR UNSPECIFIE	E <b>D</b>				
Anderson 2007	GP5+/GP6+ (L1) Hybridization with Roche LBA (6. 11. 16. 18. 26. 31. 33. 35. 39. 40. 42. 45. 51. 52. 53. 54. 55. 56. 58. 59. 61. 62. 64. 66. 67. 68. 69. 70. 71. 72. 73. 81. 82. 83. 84. 89)	36	22.2	(11.7-38.1)	HPV 16 (19.4) HPV 11 (2.8)
Schache 2011	TS-PCR E6 for 16 Amplification with TS primers (16)	98	40.8	(31.6-50.7)	HPV 16 (40.8)
Thavaraj 2011	GP5+/GP6+ (L1) Luminex 200 IS (16. 18. 26. 31. 33. 35. 39. 45. 51. 52. 53. 56. 58. 59. 66. 68. 73. 82)	142	70.4	(62.5-77.3)	HPV 16 (64.1) HPV 33 (2.1) HPV 18 (0.7) HPV 35 (0.7)

# Data updated on 9 May 2016 (data as of 31 Dec 2015)

DBH: Dot Blot Hybridization; EIA: Enzyme ImmunoAssay; HC2: Hybrid Capture 2; ISH: In Situ Hybridization; LBA: Line-Blot Assay; LiPA: Line Probe Assay; PCR: Polymerase Chain Reaction; RFLP: Restriction Fragment Length Polymorphism; RLBH: Reverse Line Blot Hybridization; RT-PCR: Real Time Polymerase Chain Reaction; SBH: Southern Blot Hybridization; SPF: Short Primer Fragment; TS: Type Specific

Only for European countries <sup>a</sup> 95% Confidence Interval

Data Sources:
Anderson CE, J Clin Pathol 2007; 60: 439 | Schache AG, Clin Cancer Res 2011; 17: 6262 | Thavaraj S, J Clin Pathol 2011; 64: 308

Based on systematic reviews and meta-analysis performed by ICO. Reference publications: 1) Ndiaye C, Lancet Oncol 2014; 15: 1319 2) Kreimer AR, Cancer Epidemiol Biomarkers Prev 2005; 14: 467

Table 38: Studies on HPV prevalence among cases of hypopharyngeal or laryngeal cancer in United Kingdom of Great Britain and Northern Ireland

			HPV	Prevalence	
Study	HPV detection method and targeted HPV types	No. Tested	%	(95% CI) <sup>a</sup>	Prevalence of 5 most frequent HPVs, HPV type (%)
MEN					
No data available	<u>-</u>	-	-	-	-

Continued on next page

Table 38 - continued from previous page

	Tubic do continue.		P		
Study	HPV detection method and targeted HPV types	No. Tested	%	(95% CI) <sup>a</sup>	Prevalence of 5 most frequent HPVs, HPV type (%)
WOMEN					
No data available	-	-	-	-	-
BOTH OR UNSPECIFII	ED				
Anderson 2007	GP5+/GP6+ (L1) Hybridization with Roche LBA (6. 11. 16. 18. 26. 31. 33. 35. 39. 40. 42. 45. 51. 52. 53. 54. 55. 56. 58. 59. 61. 62. 64. 66. 67. 68. 69. 70. 71. 72. 73. 81. 82. 83. 84. 89)	64	0.0	-	-
Conway 2012	PCR-GP5+/6+, TS, Sequencing (HPV 16, 18, 20, 21, 22, 23, 26, 30, 31, 32, 33, 34, 35, 38, 39, 40, 42, 43, 44, 45, 51, 52, 53, 54, 56, 57, 58, 59, 60, 61, 62, 66, 67, 68, 69, 70, 71, 72, 73, 74, 80, 81, 82, 83, 84, 85, 86, 87, 89, 90, 91)	12	0.0	-	-
Salam 1995	MY09/MY11 (L1) RFLP (6. 11. 16. 18. 33)	36	22.2	(11.7-38.1)	HPV 6 (8.3) HPV 16 (5.6) HPV 11 (2.8)
Snijders 1996	GP5+/GP6+ (L1) Amplification with TS primers and SBH with TS probes (6. 11. 16. 18. 31. 33)	31	19.4	(9.2-36.3)	HPV 16 (19.4)

# Data updated on 9 May 2016 (data as of 31 Dec 2015)

DBH: Dot Blot Hybridization; EIA: Enzyme ImmunoAssay; HC2: Hybrid Capture 2; ISH: In Situ Hybridization; LBA: Line-Blot Assay; LiPA: Line Probe Assay; PCR: Polymerase Chain Reaction; RFLP: Restriction Fragment Length Polymorphism; RLBH: Reverse Line Blot Hybridization; RT-PCR: Real Time Polymerase Chain Reaction; SBH: Southern Blot Hybridization; SPF: Short Primer Fragment, TS: Type Specific Only for European countries

a 95% Confidence Interval

Data Sources:
Anderson CE, J Clin Pathol 2007; 60: 439 | Conway C, J Mol Diagn 2012; 14: 104 | Salam M, Eur J Surg Oncol 1995; 21: 290 | Snijders PJ, Int J Cancer 1996; 66: 464
Based on systematic reviews and meta-analysis performed by ICO. Reference publications: 1) Ndiaye C, Lancet Oncol 2014; 15: 1319 2) Kreimer AR, Cancer Epidemiol Biomarkers Prev 2005; 14: 467

#### 5 Factors contributing to cervical cancer

HPV is a necessary cause of cervical cancer, but it is not a sufficient cause. Other cofactors are necessary for progression from cervical HPV infection to cancer. Tobacco smoking, high parity, long-term hormonal contraceptive use, and co-infection with HIV have been identified as established cofactors. Co-infection with Chlamydia trachomatis and herpes simplex virus type-2, immunosuppression, and certain dietary deficiencies are other probable cofactors. Genetic and immunological host factors and viral factors other than type, such as variants of type, viral load and viral integration, are likely to be important but have not been clearly identified. (Muñoz N, Vaccine 2006; 24(S3): 1-10). In this section, the prevalence of smoking, parity (fertility), oral contraceptive use, and HIV in United Kingdom of Great Britain and Northern Ireland are presented.

Table 39: Factors contributing to cervical carcinogenesis (cofactors) in United Kingdom of Great Britain and Northern Ireland

		and Northern freta		
INDICATOR		MALE	FEMALE	TOTAL
Smoking				
Smoking of any tobacco adjusted	Current <sup>a</sup>	23.3 [18.6-28.4]	18.5 [14.6-22.4]	20.8 [16.6-25.3]
prevalence (%) [95% UI]	Daily <sup>b</sup>	18.5 [12.2-24.6]	14.3 [9.5-19.5]	16.3 [10.8-22]
Cigarette smoking adjusted	Current <sup>c</sup>	23.3 [18.6-28.4]	18.5 [14.6-22.4]	20.8 [16.6-25.3]
prevalence (%) [95% UI]	Daily <sup>d</sup>	18.5 [12.2-24.6]	14.3 [9.5-19.5]	16.3 [10.8-22]
Parity				
Total fertility rate per woman		-	1.9	-
	15-19 yrs	-	12.6	_
	20-24 yrs	-	52.7	_
	25-29 yrs	-	94.0	_
Age-specific fertility rate	30-34 yrs	-	108.8	-
per 1000 women)	35-39 yrs	-	64.7	_
	40-44 yrs	-	14.5	-
	45-49 yrs	-	1	_
Hormonal contraception Oral contraceptive use (%) among w	romen who are	-	28	-
married or in union Injectable contraception use (%) a who are married or in union	imong women	-	2	-
Implant contraceptive use (%) amor are married or in union	ng women who	-	1	-
HIV				
Estimated percent of adults aged living with HIV [95% UI]		0.3 [0.2-0.3]	0.1 [0.1-0.1]	0.2 [0.2-0.2]
Estimated percent of young adults a are living with HIV [95% UI]	_	- [—]	- [—]	- [—]
HIV prevalence (%) among sex worl	cers	-	-	-
HIV prevalence (%) among men who men	have sex with	7.6999998	-	7.6999998
Estimated number of people living v UI]	with HIV [95%	-	-	100000 [85000-120000]
Estimated number of adults (15+ y HIV [95% UI]	rs) living with	72000 [58000-84000]	30000 [26000-34000]	100000 [85000-120000]
Estimated number of AIDS-related UI]	l deaths [95%	-	-	<200 [<200-<500]

# Data accessed on 12 Nov 2019

Crude adjusted prevalence (%) estimates of tabacco use among people aged >= 15 years by country, for the year 2016.

Excluding Northern Ireland.

Year of estimate: 2016

Data Sources:

WHO global report on trends in prevalence of tobacco use 2000-2025, third edition. Geneva: World Health Organization; 2019. Available at https://www.who.int/publications/i/ ends-in-prevalence-

Eurostat - Statistical office of the European Comission [web site]. Luxembourg: European Commission; 2017. Available at: https://ec.europa.eu/eurostat/web/products-datasets/-/demofrate. [Accessed on November 13, 2019].

United Nations, Department of Economic and Social Affairs, Population Division (2017). World Population Prospects: The 2017 Revision, DVD Edition. Available at: https://www.un.org/

en/development/desa/population/publications/dataset/fertility/wfd2017.asp. [Accessed on November 13, 2019].
United Nations, Department of Economic and Social Affairs, Population Division (2019). World Contraceptive Use 2019 (POP/DB/CP/Rev2019). https://www.un.org/en/development/desa/population/publications/dataset/contraception/wcu2019.asp. Available at: [Accessed on November 18, 2019].

a "Current" means smoking at the time of the survey, including both daily and non-daily or occasional smoking. "Tobacco smoking" means smoking any form of tobacco, including cigarettes, cigars, pipes, or any other smoked tobacco products and excluding smokeless products.

b "Daily" means smoking every day at the time of the survey. "Tobacco smoking" means smoking any form of tobacco, including cigarettes, cigars, pipes, or any other smoked tobacco products and excluding smokeless products

<sup>&</sup>quot;Current" means smoking at the time of the survey, including both daily and non-daily or occasional smoking

d "Daily" means smoking every day at the time of the survey.

 $UNAIDS\ database\ [internet].\ Available\ at: \verb|http://aidsinfo.unaids.org/|\ [Accessed\ on\ November\ 21,2019]$ 

# Sexual and reproductive health behaviour indicators

Sexual intercourse is the primary route of transmission of genital HPV infection. Information about sexual and reproductive health behaviours is essential to the design of effective preventive strategies against anogenital cancers. In this section, we describe sexual and reproductive health indicators that may be used as proxy measures of risk for HPV infection and anogenital cancers. Several studies have reported that earlier sexual debut is a risk factor for HPV infection, although the reason for this relationship is still unclear. In this section, information on sexual and reproductive health behaviour in United Kingdom of Great Britain and Northern Ireland are presented.

Table 40: Percentage of 15-year-olds who have had sexual intercourse in United Kingdom of Great Britain and Northern Ireland

Indicator <sup>a</sup>	Male	Female
Percentage of 15-year-old subjects who report sexual intercourse	21.0	32.0
Percentage of 15-year-old subjects who report sexual intercourse	24.0	27.0
Percentage of 15-year-old subjects who report sexual intercourse	18.0	23.0

#### Data accessed on 16 Mar 2017

Please refer to original source for methods of estimation

Fifteen-year-olds teenagers only were asked whether they had ever had sexual intercourse

Year of estimation: 2013-2014

Data Sources:

unequal: gender and socioeconomic differences in young people's health and well-being. Health Behaviour in School-aged Children (HBSC) study: international report from the 2013/2014 survey. Inchley J, Currie D, Young T, et al. Copenhagen, WHO Regional Office for Europe, 2016 (Health Policy for Children and Adolescents, No. 7). Available at: http://www.euro.who.int/\_data/assets/pdf\_file/0003/303438/HSBC-No.7-Growing-up-unequal-Full-Report.pdf?ua=1

Table 41: Median age at first sex in United Kingdom of Great Britain and Northern Ireland

				MALE		FEMALE		TOTAL
Study	Year/period	Birth cohort N	N	Median age at first sex	N	Median age at first sex	N	Median age at first sex
Hubert 1998 <sup>1,a</sup>	1991	1962-1966	1202	17.2	1629	17.9	-	-
Hubert 1998 <sup>1,a</sup>	1991	1952-1961	2268	17.5	3031	18.2	-	-
Hubert 1998 <sup>1,a</sup>	1991	1942-1951	1924	18.3	2306	19.5	-	-
Hubert 1998 <sup>1,a</sup>	1991	1967-1971	864	17.1	1125	17.4	-	-
Hubert 1998 <sup>1,a</sup>	1991	1932-1941	1318	19.1	1980	20.9	-	-
Hubert 1998 <sup>1,a</sup>	1991	1972-1973	288	17.0	350	17.3	-	-
Wellings 2001 <sup>2,b</sup>	2001	1972-1976	981	17.0	1357	17.0	-	-
Wellings 2001 <sup>2,b</sup>	2001	1977-1981	903	17.0	1279	16.0	-	-
Wellings 2001 <sup>2,b</sup>	2001	1967-1971	808	17.0	1160	17.0	-	-
Wellings 2001 <sup>2,b</sup>	2001	1982-1985	827	16.0	1131	16.0	-	-
Wellings 2001 <sup>2,b</sup>	2001	1957-1961	578	17.0	613	17.0	-	-
Wellings 2001 <sup>2,b</sup>	2001	1957-1985	4743	17.0	6364	17.0	-	-
Wellings 2001 <sup>2,b</sup>	2001	1962-1966	646	17.0	824	17.0	-	-
Wellings 2013 <sup>3,b</sup>	2010-2012	1956-1965	780	17.0	1106	17.0	-	-
Wellings 2013 <sup>3,b</sup>	2010-2012	1936-1945	657	18.0	850	19.0	-	-
Wellings 2013 <sup>3,b</sup>	2010-2012	1936-1994	6207	17.0	8746	17.0	-	-
Wellings 2013 <sup>3,b</sup>	2010-2012	1976-1985	1508	17.0	2469	16.0	-	-
Wellings 2013 <sup>3,b</sup>	2010-2012	1966-1975	792	17.0	1196	17.0	-	-
Wellings 2013 <sup>3,b</sup>	2010-2012	1946-1955	758	18.0	1015	18.0	-	-
Wellings 2013 <sup>3,b</sup>	2010-2012	1986-1994	1712	16.0	2110	16.0	-	-

### Data accessed on 16 Mar 2017

Please refer to original source for methods of estimation

 $<sup>^{</sup>a}$  Indicates a significant gender difference (at p<0.05).

Not especified if estimations are among sexually active or surveyed.

b Number of subjects refers to the number of surveyed men/women (not all sexually active). Data Sources

 $<sup>^{1}</sup>$  Hubert M, Bajos N, Sandfort T. Sexual behaviour and HIV/AIDS in Europe: comparisons of national surveys. London: UCL Press; 1998

<sup>2</sup> Wellings K, Nanchahal K, Macdowall W, McManus S, Erens B, Mercer CH, Johnson AM, Copas AJ, Korovessis C, Fenton KA, Field J. Sexual behaviour in Britain: early heterosexual experience. Lancet. 2001 Dec 1;358(9296):1843-50.

<sup>&</sup>lt;sup>3</sup> Wellings K, Jones KG, Mercer CH, Tanton C, Clifton S, Datta J, et al. The prevalence of unplanned pregnancy and associated factors in Britain: fi ndings from the third National Survey of Sexual Attitudes and Lifestyles (Natsal-3). Lancet 2013; 382: 1807-16.

Table 42: Marriage patterns in United Kingdom of Great Britain and Northern Ireland

Indicator		Male	Female
Average age at first marriage <sup>1</sup>		28.7	27
Age-specific % of ever married <sup>2</sup>	15-19 years	1.24	2.97
	20-24 years	15.05	26.21
	25-29 years	44.12	57.53
	30-34 years	65.69	73.74
	35-39 years	75.97	80.9
	40-44 years	80.98	85.11
	45-49 years	84.47	88.5
	50-54 years	87.72	91.58
	55-59 years	90.04	93.71
	60-64 years	92.02	95.27
	65-69 years	93.25	95.88
	70-74 years	94.04	95.81
	+75	94.4	94.37

# Data accessed on 20 Feb 2020

Please refer to original source for methods of estimation.

Table 43: Average number of sexual partners in United Kingdom of Great Britain and Northern Ireland

Study	Period of estimate	Year/Period	Birth cohort	Male Mean(N)	Female Mean(N)	Total Mean(N)
Almonte 2011 <sup>1,a,b</sup>	Lifetime	2003-2007	(1959-1972)	-(-)	5.6(195)	-(-)
Almonte 2011 <sup>1,a,b</sup>	Lifetime	2003-2007	(1959-1990)	-(-)	7.5(436)	-(-)
Almonte 2011 <sup>1,a,b</sup>	Lifetime	2003-2007	(1969-1982)	-(-)	7.7(168)	-(-)
Almonte 2011 <sup>1,a,b</sup>	Lifetime	2003-2007	(1979-1990)	-(-)	12.0(73)	-(-)
Hubert 1998 <sup>2,b</sup>	Lifetime	1990-1991	(1941-1973)	10.4(6414)	3.7(8049)	-(-)
Hubert 1998 <sup>2,c</sup>	Last year	1990-1991	(1941-1973)	1.3(6134)	1.1(7790)	-(-)
Johnson 2001 <sup>3,d</sup>	Lifetime	1999-2001	(1955-1966)	16.0(1691)	6.8(2356)	-(-)
Johnson 2001 <sup>3,d</sup>	Last 5 years	1999-2001	(1955-1966)	2.2(1669)	1.5(2332)	-(-)
Johnson 2001 <sup>3,d</sup>	Lifetime	1999-2001	(1955-1985)	12.7(4762)	6.5(6399)	-(-)
Johnson 2001 <sup>3,d</sup>	Last 5 years	1999-2001	(1955-1985)	3.8(4762)	2.4(6399)	-(-)
Johnson 2001 <sup>3,d</sup>	Lifetime	1999-2001	(1965-1976)	13.6(1759)	7.3(2486)	-(-)
Johnson 2001 <sup>3,d</sup>	Last 5 years	1999-2001	(1965-1976)	4.2(1751)	2.2(2474)	-(-)
Johnson 2001 <sup>3,d</sup>	Lifetime	1999-2001	(1975-1985)	6.9(1211)	5.0(1433)	-(-)
Johnson 2001 <sup>3,d</sup>	Last 5 years	1999-2001	(1975-1985)	5.3(1200)	3.8(1422)	-(-)
Mercer 2009 <sup>4,d</sup>	Last year	1999-2001	(1955-1985)	1.8(4762)	1.3(6399)	-(-)

## Data accessed on 8 Aug 2013

a 2011 Census

b Eurostat

Data Sources:

The world bank: health nutrition and population statistics. Updated 20-Dec-2019. Accessed on February 20 2020. Available at http://data.worldbank.org/data-catalog/ health-nutrition- and-population- statistics

United Nations, Department of Economic and Social Affairs, Population Division (2019). World Marriage Data 2019 (POP/DB/Marr/Rev2019). Available at: https://population.un.

org/MarriageData/Index.html#/home Accessed on February 24, 2020.

Please refer to original source for methods of estimation

The authors used the following formula to estimate the Number of lifetime partners in year 2000 in women aged 17-45 years= Number of lifetime partners - Number of new partners in

b Number of surveyed people (not all sexually active).

<sup>&</sup>lt;sup>c</sup> Data among responders who ever had a heterosexual partner.

 $d\,$  Number of heterosexual partners among surveyed (not all sexually active).

<sup>&</sup>lt;sup>1</sup> Almonte M, Silva Idos S, Asare A, Gilham C, Sargent A, Bailey A, Turner A, Desai M, Kitchener HC, Peto J. Sexual behavior and HPV infection in British women, by postal questionnaires and telephone interviews. J Med Virol. 2011 Jul;83(7):1238-46.  $^2$  Hubert M, Bajos N, Sandfort T. Sexual behaviour and HIV/AIDS in Europe: comparisons of national surveys. London: UCL Press; 1998.

<sup>&</sup>lt;sup>3</sup> Johnson AM, Mercer CH, Erens B, Copas AJ, McManus S, Wellings K, et al. Sexual behaviour in Britain: partnerships, practices, and HIV risk behaviours. Lancet. 2001 Dec

<sup>1;358(9296):1835-42.</sup>Mercer CH, Copas AJ, Sonnenberg P, Johnson AM, McManus S, Erens B, Cassell JA. Who has sex with whom? Characteristics of heterosexual partnerships reported in a national probability survey and implications for STI risk.Int J Epidemiol. 2009 Feb;38(1):206-14.

Table 44: Lifetime prevalence of anal intercourse among women in United Kingdom of Great Britain and Northern Ireland

FEMALE				FEMALE	
$\mathbf{Study}^a$	Year/Period	Birth cohort	N surveyed	N sexual active	% among sexually active
Johnson $2001^{1,b}$	1990-1991	(1946-1975)	7765	-	6.5
Johnson $2001^{1,b}$	1999-2001	(1955-1985)	6399	-	11.3
Stone 2006 <sup>2,c</sup>	2003-2005	(1984-1990)	765	-	9.3

# Data accessed on 8 Aug 2013

Please refer to original source for methods of estimation <sup>a</sup> Instead of number of women sexually active, this time was number of women who answered the question on anal intercourse.

a Instead of number of women sexually active, this time was number of women who answered the question on anal intercourse.
 b Data pertain to heterosexual women.
 c Data pertain to full and part time students.
 Data Sources:
 1 Johnson AM, Mercer CH, Erens B, Copas AJ, McManus S, Wellings K, et al. Sexual behaviour in Britain: partnerships, practices, and HIV risk behaviours. Lancet. 2001 Dec 1;358(9296):1835-42.
 2 Stone N, Hatherall B, Ingham R, McEachran. J Oral sex and condom use among young people in the United Kingdom. Perspect Sex Reprod Health. 2006 Mar;38(1):6-12.

# 7 HPV preventive strategies

It is established that well-organised cervical screening programmes or widespread good quality cytology can reduce cervical cancer incidence and mortality. The introduction of HPV vaccination could also effectively reduce the burden of cervical cancer in the coming decades. This section presents indicators on basic characteristics and performance of cervical cancer screening, status of HPV vaccine licensure and introduction in United Kingdom of Great Britain and Northern Ireland.

# 7.1 Cervical cancer screening practices

Screening strategies differ between countries. Some countries have population-based programmes, where in each round of screening women in the target population are individually identified and invited to attend screening. This type of programme can be implemented nationwide or only in specific regions of the country. In opportunistic screening, invitations depend on the individual's decision or on encounters with health-care providers. The most frequent method for cervical cancer screening is cytology, and there are alternative methods such as HPV DNA tests and visual inspection with acetic acid (VIA). VIA is an alternative to cytology-based screening in low-resource settings (the 'see and treat' approach). HPV DNA testing is being introduced into some countries as an adjunct to cytology screening ('co-testing') or as the primary screening test to be followed by a secondary, more specific test, such as cytology.

Table 45: Main characteristics of cervical cancer screening in United Kingdom of Great Britain and Northern Ireland

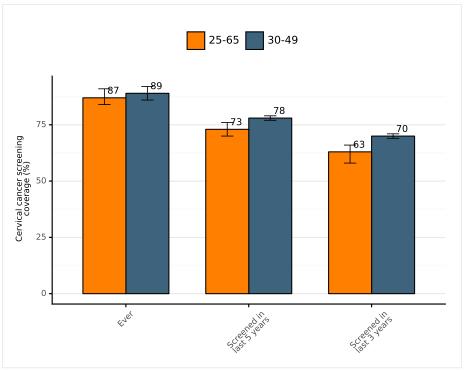
Region	Existence of official national recommendations	Starting year of current recommendations	Active invitation to screening	Screening ages (years), primary screening test used, and screening interval or frequency of screenings
England	Yes	2020	Yes	25-49 (HPV test, 3 years); 50-64 (HPV test, 5 years)
Northern Ireland	Yes	2011	Yes	25-49 (cytology, 3 years); 50-64 (cytology, 5 years)
Scotland	Yes	2020	Yes	25-64 (HPV test, 5 years)
Wales	Yes	2019	Yes	25-49 (HPV test, 3 years); 50-64 (HPV test, 5 years)

Data accessed on 31 Aug 2022

Data Source

Bruni L, Serrano B, Roura E, Alemany L, Cowan M, Herrero R, et al. Cervical cancer screening programmes and age-specific coverage estimates for 202 countries and territories worldwide: a review and synthetic analysis. Lancet Glob Health. 2022;10(8):e1115.

Figure 70: Estimated coverage\* of cervical cancer screening in United Kingdom of Great Britain and Northern Ireland



Data accessed on 31 Aug 2022

\* Estimated coverage and 95% confidence interval in 2019

Data Sources:
Bruni L, Serrano B, Roura E, Alemany L, Cowan M, Herrero R, et al. Cervical cancer screening programmes and age-specific coverage estimates for 202 countries and territories worldwide: a review and synthetic analysis. Lancet Glob Health. 2022;10(8):e1115.

# 7.2 HPV vaccination

Table 46: National HPV Immunization programme in United Kingdom of Great Britain and Northern Ireland

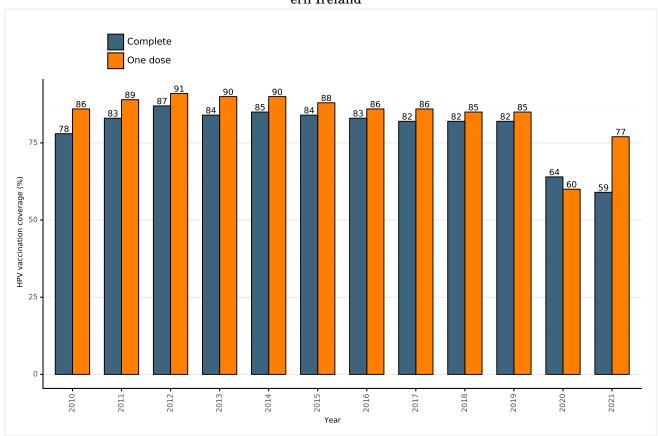
	Female	Male
HPV vaccination programme	Introduced	Introduced
Year of introduction	2008	2019
Year of estimation of HPV vaccination coverage	2021	2021
HPV coverage – first dose (%)	77	71
HPV coverage – last dose (%)	59	48

#### Data accessed on 24 Oct 2022

<u>Data Sources</u>: Human papillomavirus (HPV) vaccination coverage. World Health Organization. 2022. Available from: https://immunizationdata.who.int/pages/coverage/hpv.html, accessed [24 Oct 2022]

Bruni L, Saura-Lázaro A, Montoliu A, Brotons M, Alemany L, Diallo MS, et al. HPV vaccination introduction worldwide and WHO and UNICEF estimates of national HPV immunization coverage 2010-2019. Prev Med. 2021;144(106399):106399.

Figure 71: HPV vaccination coverage in females by year in United Kingdom of Great Britain and Northern Ireland

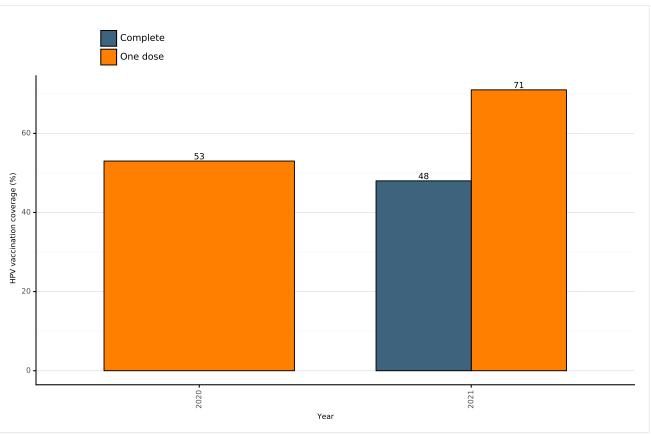


Data accessed on 24 Oct 2022

Data Sources:
Human papillomavirus (HPV) vaccination coverage. World Health Organization. 2022. Available from: https://immunizationdata.who.int/pages/coverage/hpv.html, accessed [24 Oct 2022]

Bruni L, Saura-Lázaro A, Montoliu A, Brotons M, Alemany L, Diallo MS, et al. HPV vaccination introduction worldwide and WHO and UNICEF estimates of national HPV immunization coverage 2010-2019. Prev Med. 2021;144(106399):106399.

Figure 72: HPV vaccination coverage in males by year in United Kingdom of Great Britain and Northern Ireland



# Data accessed on 24 Oct 2022

Data Sources:
Human papillomavirus (HPV) vaccination coverage. World Health Organization. 2022. Available from: https://immunizationdata.who.int/pages/coverage/hpv.html, accessed [24 Oct 2022]

Bruni L, Saura-Lázaro A, Montoliu A, Brotons M, Alemany L, Diallo MS, et al. HPV vaccination introduction worldwide and WHO and UNICEF estimates of national HPV immunization coverage 2010-2019. Prev Med. 2021;144(106399):106399.

#### 8 Protective factors for cervical cancer

Male circumcision and the use of condoms have shown a significant protective effect against HPV transmission.

Table 47: Prevalence of male circumcision in United Kingdom of Great Britain and Northern Ireland

Reference	Prevalence % (95% CI)	Methods	
Oriel 1971	24.0 (18.9-29.6)	N=263: STD Clinics patients	
		Data fron National Health System (NHS),	
		medically indicated operative	
Rickwood 2000	3.8	circumcisions, circumcision rate about	
		12200 procedures annually in boys by 15	
		years old	
		N=5,746: British National Survey of	
Dave 2003	15.8 (14.7-17.1)	Sexual Attitudes and Lifestyles (Natsa	
		2000), british men aged 16-44 years old	
Thornton 2011	28.9 (25.5-32.6)	N=653: Men who have sex with men	
Doerner 2013	16.7 (14.9-18.7)	N=1,521: White men who have sex with	
Doerner 2019	10.7 (14.3-10.7)	men	
		Data from Demographic and Health	
		Surveys (DHS) and other publications to	
WHO 2007	<20	categorize the country-wide prevalence of	
		male circumcision as <20%, 20-80%, or	
		>80%.	
Cathcart 2006	(2.1/1,000 boys)	N=75,868: General population, boys below	
Califai i 2000	(2.1/1,000 boys)	15 years of age (2003)	

#### Data accessed on 31 Aug 2015

Please refer to country-specific reference(s) for full methodologies.

Data Sources:

Tathcart P, Br J Surg 2006; 93: 885 | Dave SS, Sex Transm Infect 2003; 79: 499 | Doerner R, Arch Sex Behav 2013; 42: 1319 | Oriel JD, Br J Vener Dis 1971; 47: 1 | Rickwood AM, BMJ 2000; 321: 792 | Thornton AC, Sex Transm Dis 2011; 38: 928 | WHO 2007: Male circumcision: Global trends and determinants of prevalence, safety and acceptability
Based on systematic reviews and meta-analysis performed by ICO. The ICO HPV Information Centre has updated data until August 2015. Reference publication: Albero G, Sex Transm
Dis. 2012 Feb;39(2):104-13.

Table 48: Prevalence of condom use in United Kingdom of Great Britain and Northern Ireland

Indicator	Age range	Year of estimate	Prevalence $\%^a$
Condom use	16-49	2008	27

# Data accessed on 18 Nov 2019

Please refer to original source for methods of estimation

Excluding Northern Ireland.

<sup>a</sup> Condom use: Proportion of male partners who are using condoms with their female partners of reproductive age to whom they are married or in union by country. Data Sources: Annual OS

United Nations, Department of Economic and Social Affairs, Population Division (2019). World Contraceptive Use 2019 (POP/DB/CP/Rev2019). https://www.un.org/en/development/ desa/population/publications/dataset/contraception/wcu2019.asp. Available at: [Accessed on November 18, 2019].

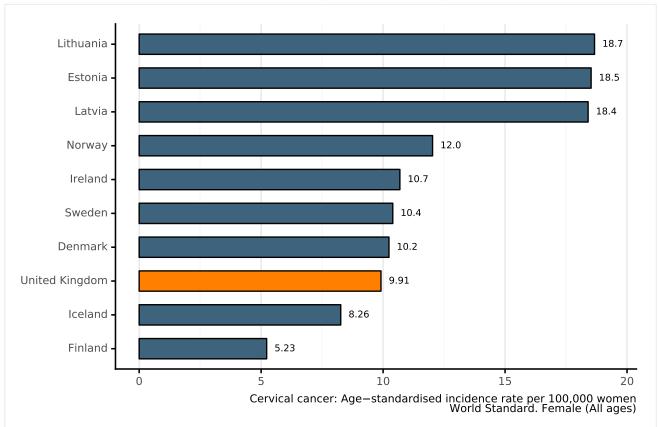
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#### 9 Annex

# 9.1 Incidence

# 9.1.1 Cervical cancer incidence in United Kingdom of Great Britain and Northern Ireland across Northern Europe

Figure 73: Age-standardised incidence rates of cervical cancer of United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



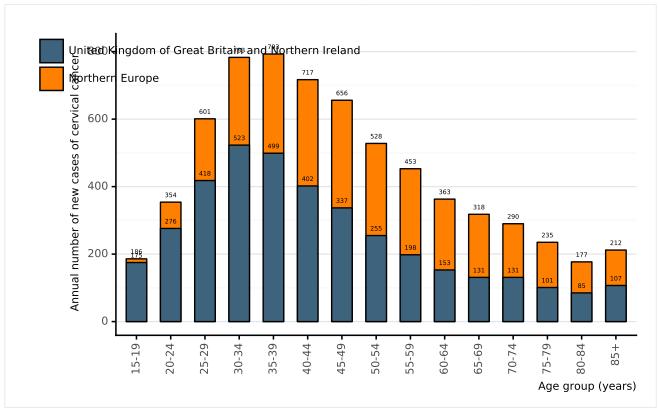
# Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods a Rates per 100,000 women per year.

Data Sources:
Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

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Figure 74: Annual number of new cases of cervical cancer by age group in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

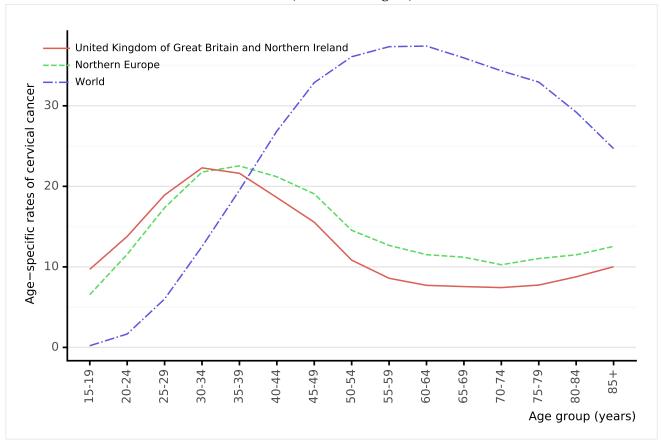


Data accessed on 27 Jan 2021
For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods
Data Sources:

Bata Doutes.
Ferlay J. Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

9 ANNEX - 95 -

Figure 75: Comparison of age-specific cervical cancer incidence rates in United Kingdom of Great Britain and Northern Ireland, within the region, and the rest of world



For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods a Rates per 100,000 women per year.

Data Sources:

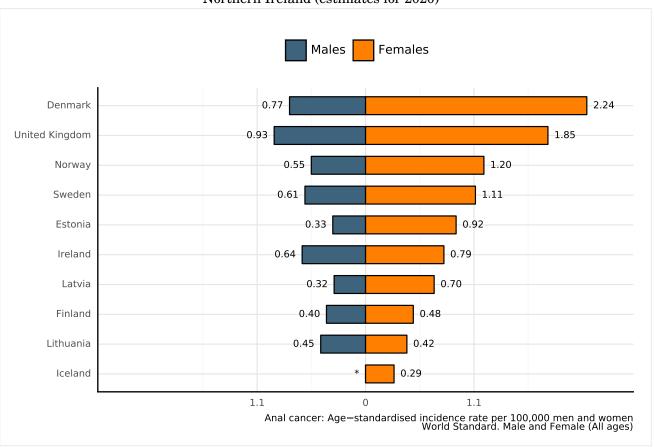
Early, I. P. 3. 2. 5.

Ferlay J. Evrik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

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# 9.1.2 Anal cancer incidence in United Kingdom of Great Britain and Northern Ireland across **Northern Europe**

Figure 76: Age-standardised incidence rates of anal cancer of United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

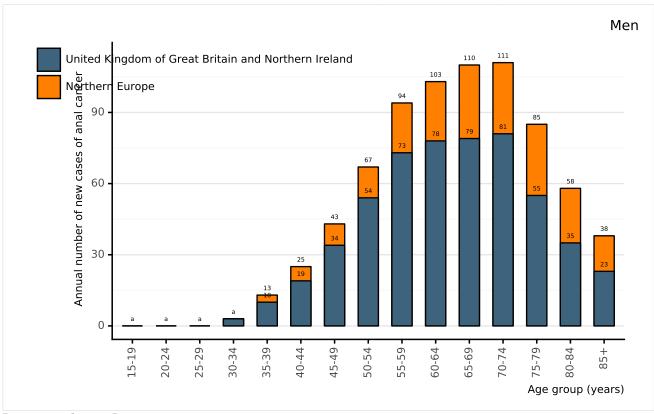
For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods a Rates per 100,000 men per year.

Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

b Rates per 100,000 women per year.
Rates are not available

9 ANNEX - 97 -

Figure 77: Annual number of new cases of anal cancer among men by age group in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

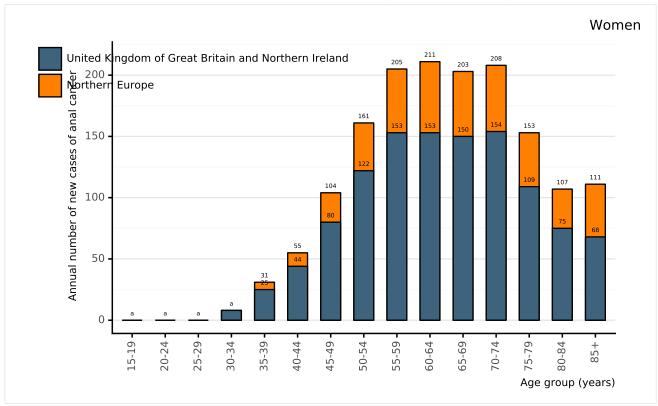
Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 15-19 age group. 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 20-24 age group. 0 cases for United Kingdom of Great Britain and Northern Europe in the 25-29 age group.
3 cases for United Kingdom of Great Britain and Northern Ireland and 3 cases for Northern Europe in the 30-34 age group.

9 ANNEX - 98 -

Figure 78: Annual number of new cases of anal cancer among women by age group in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

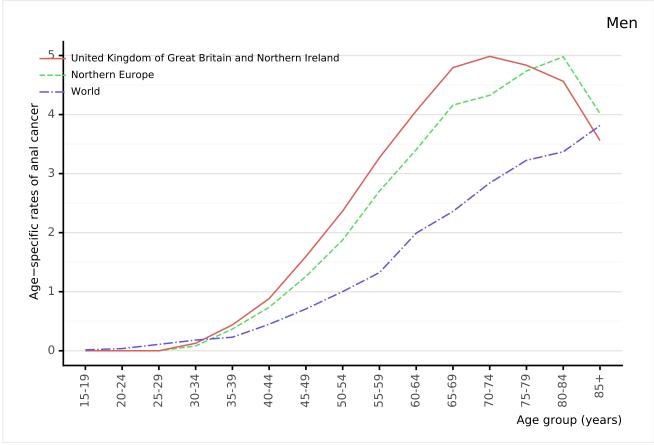
For more detailed methods of estimation please refer to <a href="http://gco.iarc.fr/today/data-sources-methods">http://gco.iarc.fr/today/data-sources-methods</a>

a 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 15-19 age group. 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 20-24 age group. 0 cases for United Kingdom of Great Britain and Northern Europe in the 25-29 age group. 8 cases for United Kingdom of Great Britain and Northern Ireland and 8 cases for Northern Europe in the 30-34 age group.

Data Sources:
Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for

9 ANNEX - 99 -

Figure 79: Comparison of age-specific anal cancer incidence rates among men by age in United Kingdom of Great Britain and Northern Ireland, within the region, and the rest of world

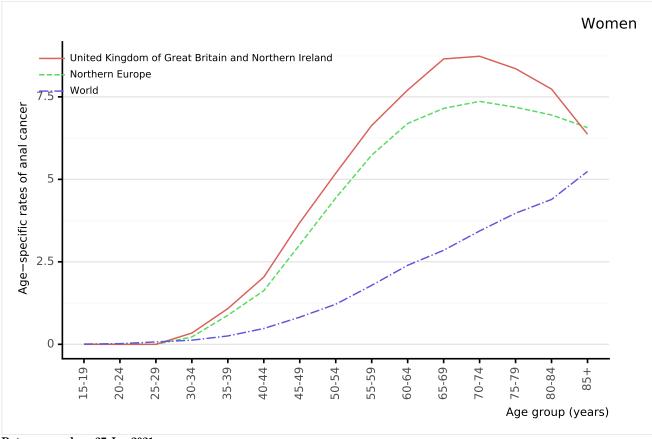


Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to  $\frac{\text{http://gco.iarc.fr/today/data-sources-methods}}{a} \text{ Rates per } 100,000 \text{ men per year.}$ 

9 ANNEX - 100 -

Figure 80: Comparison of age-specific anal cancer incidence rates among women by age in United Kingdom of Great Britain and Northern Ireland, within the region, and the rest of world



Data accessed on 27 Jan 2021

Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

<sup>a</sup> Rates per 100,000 women per year.

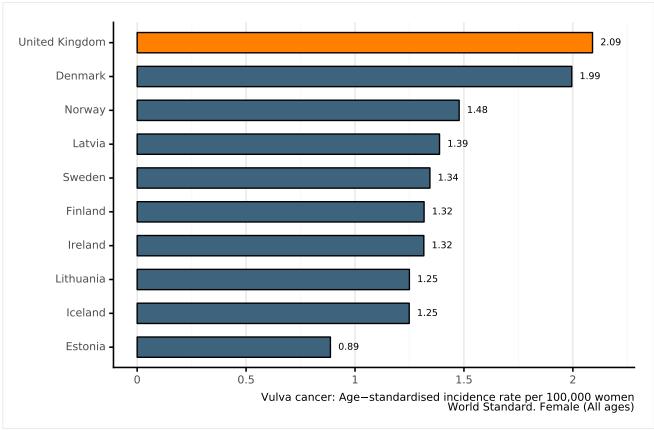
Data Sources:

Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

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## 9.1.3 Vulva cancer incidence in United Kingdom of Great Britain and Northern Ireland across Northern Europe

Figure 81: Age-standardised incidence rates of vulva cancer of United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

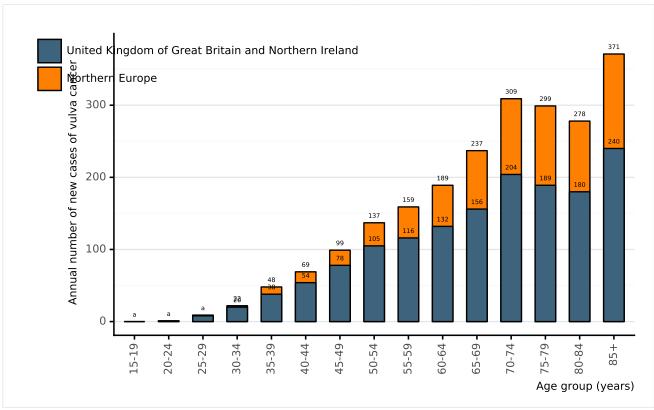


Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods a Rates per 100,000 women per year.

9 ANNEX - 102 -

Figure 82: Annual number of new cases of vulva cancer by age group in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



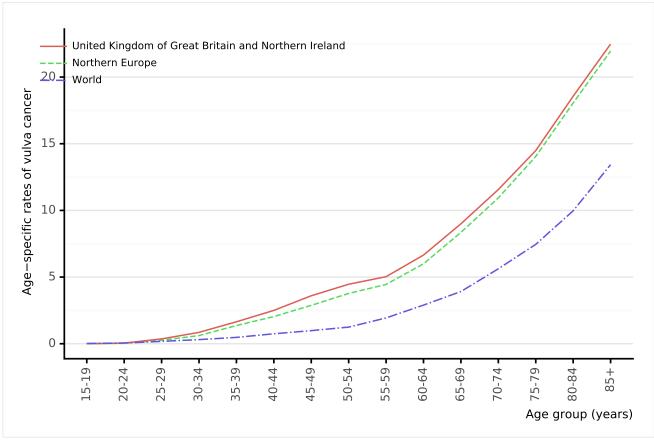
Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 15-19 age group. 1 cases for United Kingdom of Great Britain and Northern Ireland and 1 cases for Northern Europe in the 20-24 age group. 8 cases for United Kingdom of Great Britain and Northern Ireland and 9 cases for Northern Europe in the 25-29 age group.

9 ANNEX - 103 -

Figure 83: Comparison of age-specific vulva cancer incidence rates in United Kingdom of Great Britain and Northern Ireland, within the region, and the rest of world



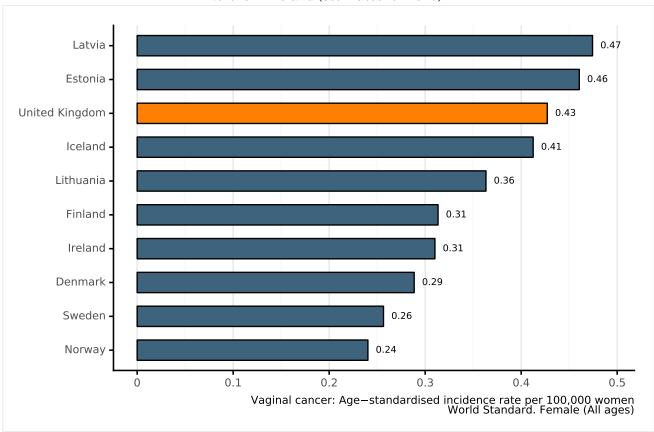
Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to  $\frac{1}{2} \frac{1}{2} \frac{1}$ 

9 ANNEX - 104 -

# 9.1.4 Vaginal cancer incidence in United Kingdom of Great Britain and Northern Ireland across Northern Europe

Figure 84: Age-standardised incidence rates of vaginal cancer of United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

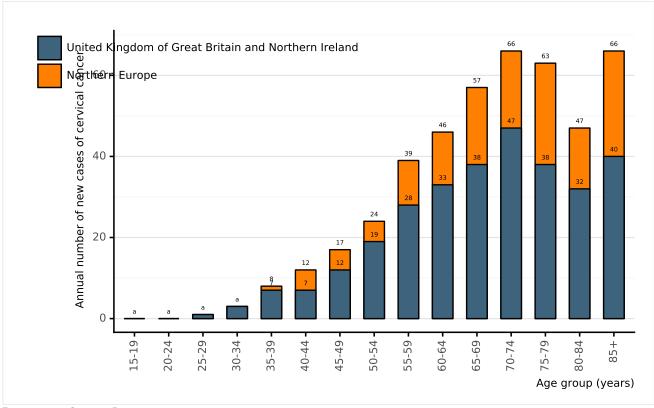


Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods a Rates per 100,000 women per year.

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Figure 85: Annual number of new cases of cervical cancer by age group in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



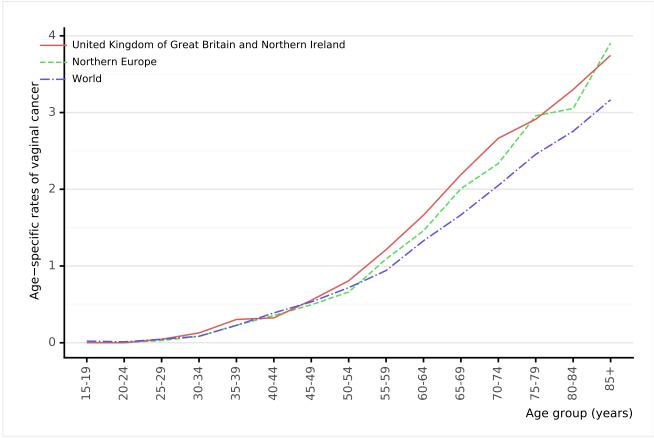
Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 15-19 age group. 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 20-24 age group. 1 cases for United Kingdom of Great Britain and Northern Ireland and 1 cases for Northern Europe in the 25-29 age group.  $3\ cases\ for\ United\ Kingdom\ of\ Great\ Britain\ and\ Northern\ Ireland\ and\ 3\ cases\ for\ Northern\ Europe\ in\ the\ 30-34\ age\ group.$ 

9 ANNEX - 106 -

Figure 86: Comparison of age-specific vaginal cancer incidence rates in United Kingdom of Great Britain and Northern Ireland, within the region, and the rest of world



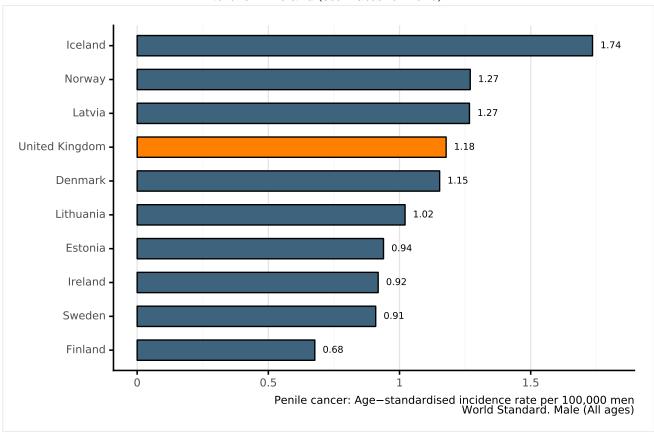
Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to  $\frac{1}{2} \frac{1}{2} \frac{1}$ 

9 ANNEX - 107 -

## 9.1.5 Penile cancer incidence in United Kingdom of Great Britain and Northern Ireland across Northern Europe

Figure 87: Age-standardised incidence rates of penile cancer of United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

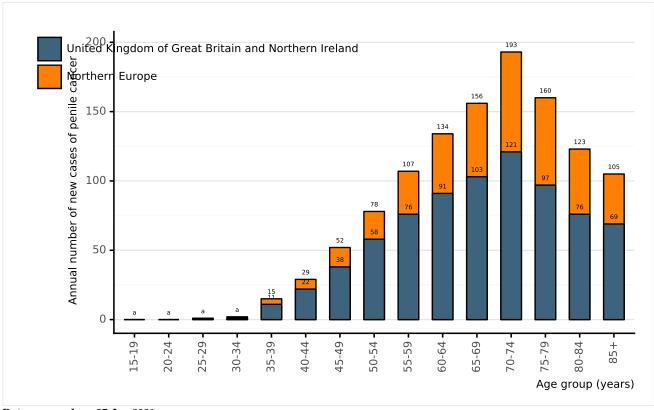


Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods a Rates per 100,000 men per year.

9 ANNEX - 108 -

Figure 88: Annual number of new cases of penile cancer by age group in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



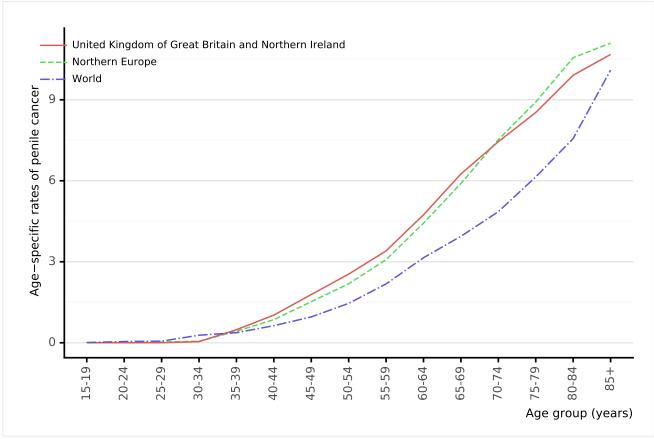
Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 15-19 age group. 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 20-24 age group. 0 cases for United Kingdom of Great Britain and Northern Europe in the 25-29 age group.  $1\ cases\ for\ United\ Kingdom\ of\ Great\ Britain\ and\ Northern\ Ireland\ and\ 2\ cases\ for\ Northern\ Europe\ in\ the\ 30-34\ age\ group.$ 

9 ANNEX - 109 -

Figure 89: Comparison of age-specific penile cancer incidence rates in United Kingdom of Great Britain and Northern Ireland, within the region, and the rest of world



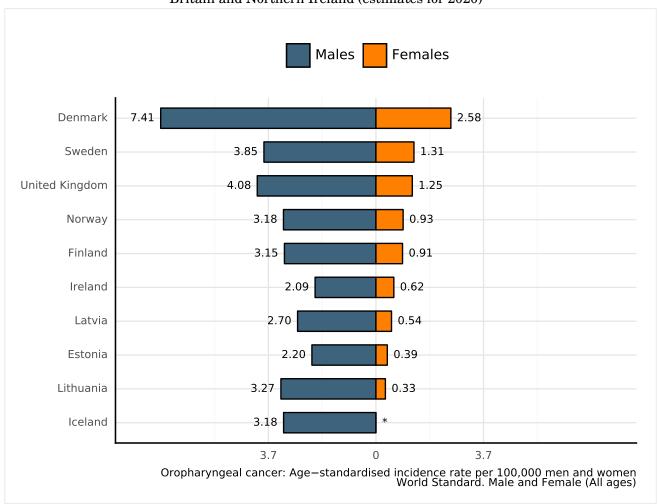
Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to  $\frac{1}{2} \frac{1}{2} \frac{1}$ 

9 ANNEX - 110 -

# 9.1.6 Oropharyngeal cancer incidence in United Kingdom of Great Britain and Northern **Ireland across Northern Europe**

Figure 90: Age-standardised incidence rates of oropharyngeal cancer of United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



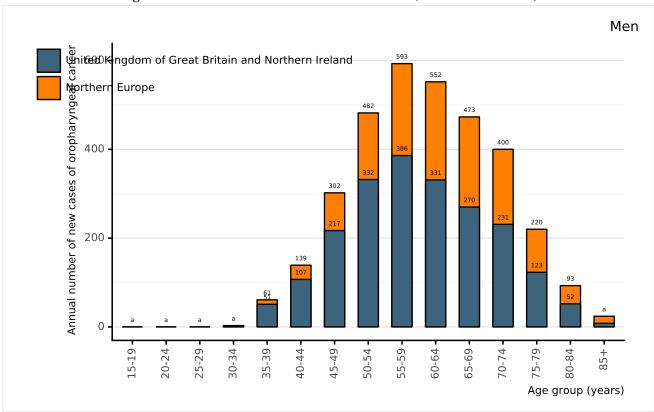
Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods Rates per 100,000 men per year.

b Rates per 100,000 women per year
Rates are not available

9 ANNEX -111-

Figure 91: Annual number of new cases of oropharyngeal cancer among men by age group in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

Data accessed on 27 Jan 2021

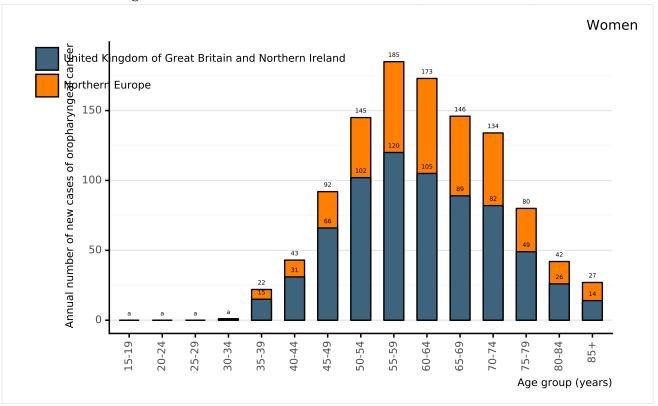
For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 15-19 age group. 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 20-24 age group. 0 cases for United Kingdom of Great Britain and Northern Europe in the 25-29 age group.

2 cases for United Kingdom of Great Britain and Northern Ireland and 3 cases for Northern Europe in the 30-34 age group. 8 cases for United Kingdom of Great Britain and Northern

9 ANNEX - 112 -

Figure 92: Annual number of new cases of oropharyngeal cancer among women by age group in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

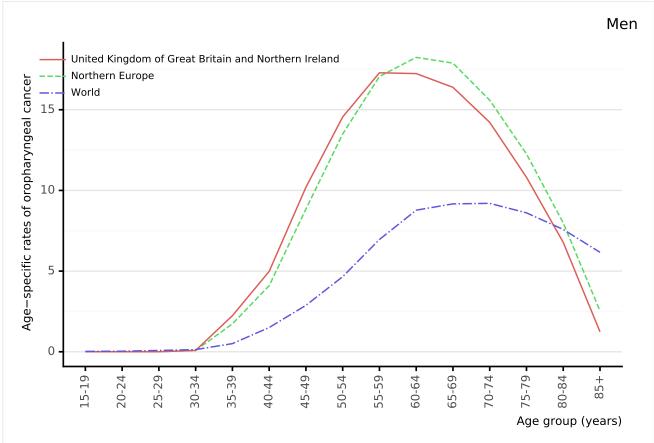
a 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 15-19 age group. 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 20-24 age group. 0 cases for United Kingdom of Great Britain and Northern Europe in the 25-29 age group.

1 cases for United Kingdom of Great Britain and Northern Ireland and 1 cases for Northern Europe in the 30-34 age group.

Data Sources:
Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for

9 ANNEX - 113 -

Figure 93: Comparison of age-specific oropharyngeal cancer incidence rates among men by age in United Kingdom of Great Britain and Northern Ireland, within the region, and the rest of world

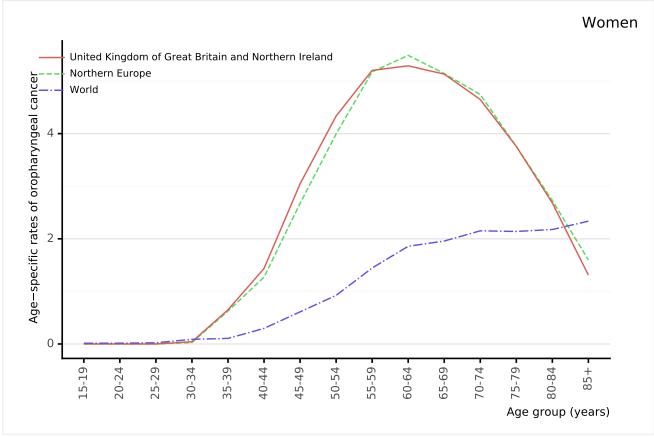


Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to  $\frac{\text{http://gco.iarc.fr/today/data-sources-methods}}{a} \text{ Rates per } 100,000 \text{ men per year.}$ 

9 ANNEX - 114 -

Figure 94: Comparison of age-specific oropharyngeal cancer incidence rates among women by age in United Kingdom of Great Britain and Northern Ireland, within the region, and the rest of world



Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a Rates per 100,000 women per year.

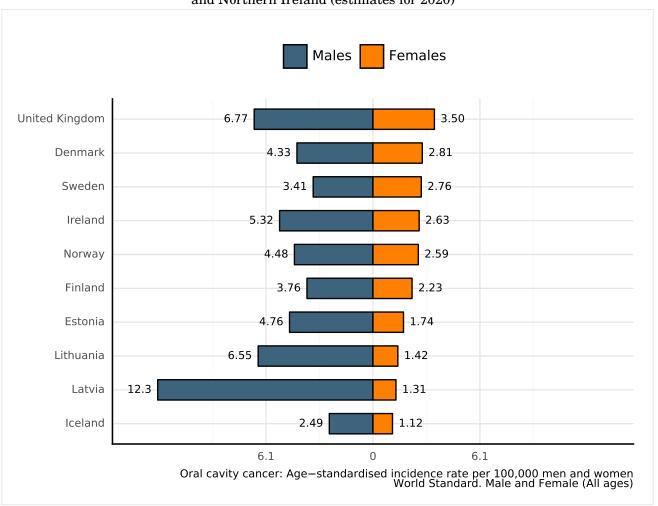
Data Sources:

Ferlay J. Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

9 ANNEX - 115 -

## 9.1.7 Oral cavity cancer incidence in United Kingdom of Great Britain and Northern Ireland across Northern Europe

Figure 95: Age-standardised incidence rates of oral cavity cancer of United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

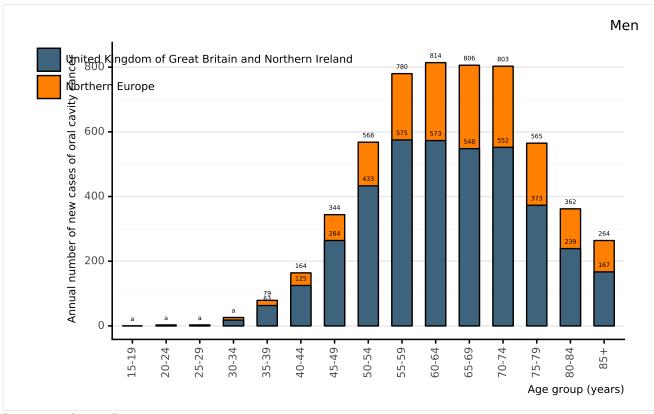
For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a Rates per 100,000 men per year.

b Rates per 100,000 women per year.

9 ANNEX - 116 -

Figure 96: Annual number of new cases of oral cavity cancer among men by age group in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

Data accessed on 27 Jan 2021

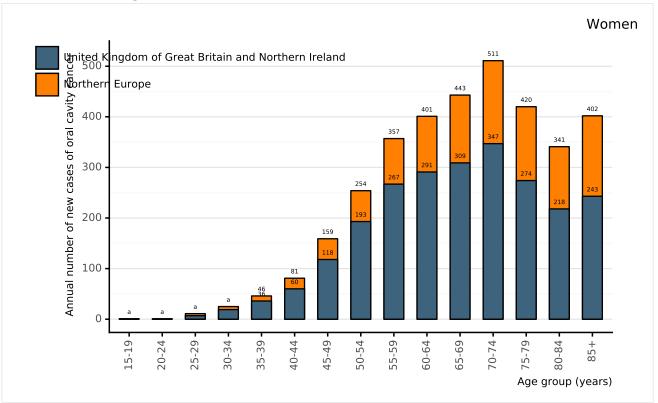
For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 15-19 age group. 0 cases for United Kingdom of Great Britain and Northern Ireland and 3 cases for Northern Europe in the 20-24 age group. 0 cases for United Kingdom of Great Britain and Northern Europe in the 25-29 age group.

18 cases for United Kingdom of Great Britain and Northern Ireland and 26 cases for Northern Europe in the 30-34 age group.

9 ANNEX - 117 -

Figure 97: Annual number of new cases of oral cavity cancer among women by age group in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

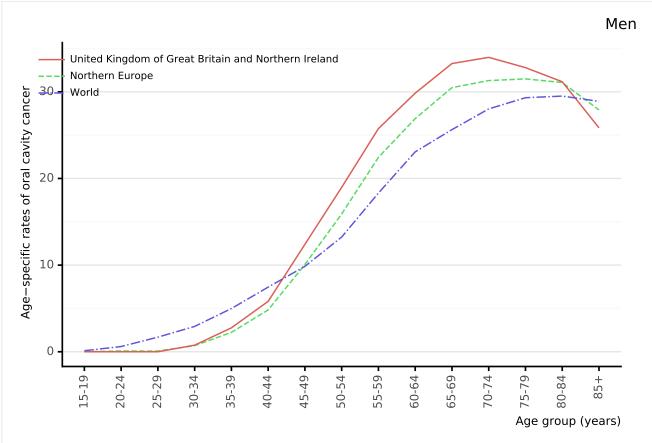
a 0 cases for United Kingdom of Great Britain and Northern Ireland and 1 cases for Northern Europe in the 15-19 age group. 0 cases for United Kingdom of Great Britain and Northern Ireland and 1 cases for Northern Europe in the 20-24 age group. 7 cases for United Kingdom of Great Britain and Northern Ireland and 11 cases for Northern Europe in the 25-29 age group.

19 cases for United Kingdom of Great Britain and Northern Ireland and 25 cases for Northern Europe in the 30-34 age group.

Data Sources:
Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for

9 ANNEX -118-

Figure 98: Comparison of age-specific oral cavity cancer incidence rates among men by age in United Kingdom of Great Britain and Northern Ireland, within the region, and the rest of world

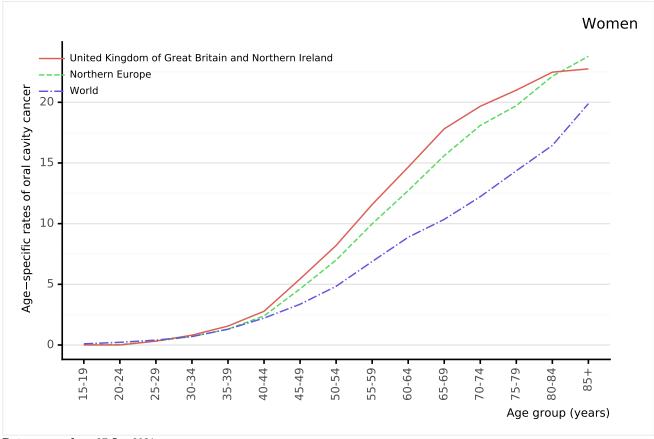


Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to  $\frac{\text{http://gco.iarc.fr/today/data-sources-methods}}{a} \text{ Rates per } 100,000 \text{ men per year.}$ 

9 ANNEX - 119 -

Figure 99: Comparison of age-specific oral cavity cancer incidence rates among women by age in United Kingdom of Great Britain and Northern Ireland, within the region, and the rest of world



Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a Rates per 100,000 women per year.

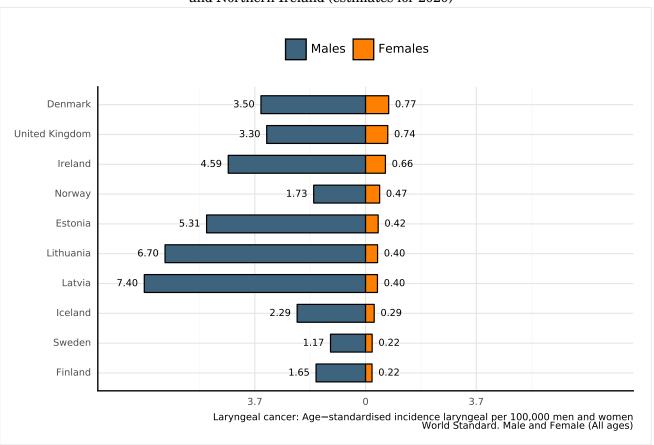
Data Sources:

Ferlay J. Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

9 ANNEX - 120 -

# 9.1.8 Laryngeal cancer incidence in United Kingdom of Great Britain and Northern Ireland across Northern Europe

Figure 100: Age-standardised incidence rates of laryngeal cancer of United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



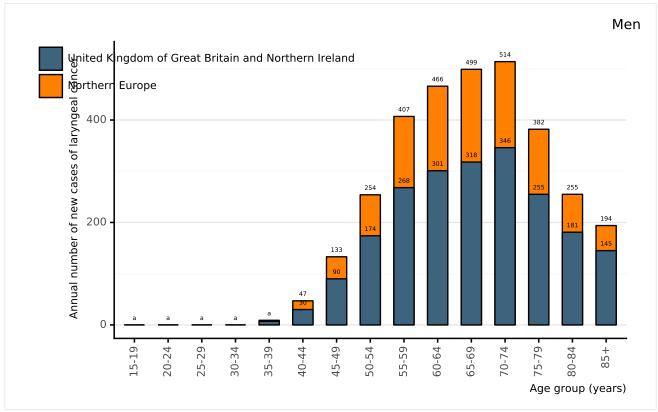
Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods a Rates per 100,000 men per year.

b Rates per 100,000 women per year.

9 ANNEX - 121 -

Figure 101: Annual number of new cases of laryngeal cancer among men by age group in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

Data accessed on 27 Jan 2021

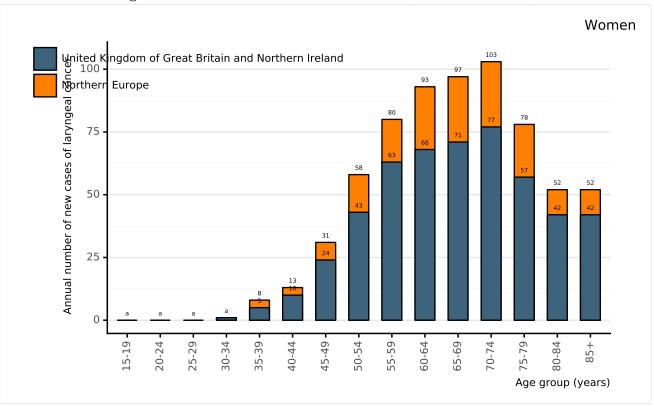
For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 15-19 age group. 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 20-24 age group. 0 cases for United Kingdom of Great Britain and Northern Europe in the 25-29 age group.

0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 30-34 age group. 7 cases for United Kingdom of Great Britain and Northern Ireland and Northern Europe in the 30-34 age group. Ireland and 9 cases for Northern Europe in the 35-39 age group.

9 ANNEX - 122 -

Figure 102: Annual number of new cases of laryngeal cancer among women by age group in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

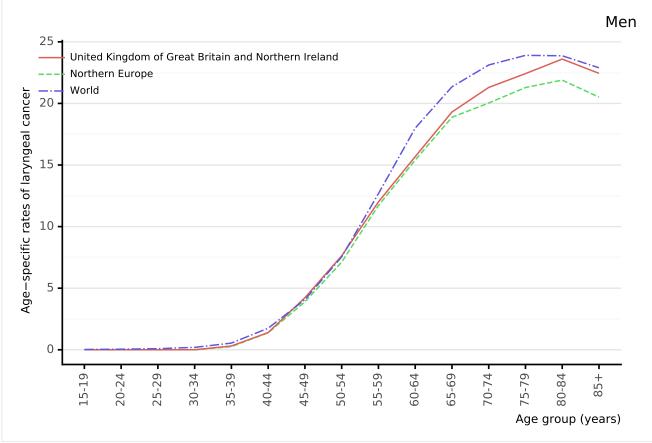
a 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 15-19 age group. 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 20-24 age group. 0 cases for United Kingdom of Great Britain and Northern Europe in the 25-29 age group.

1 cases for United Kingdom of Great Britain and Northern Ireland and 1 cases for Northern Europe in the 30-34 age group.

Data Sources:
Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for

9 ANNEX - 123 -

Figure 103: Comparison of age-specific laryngeal cancer incidence rates among men by age in United Kingdom of Great Britain and Northern Ireland, within the region, and the rest of world

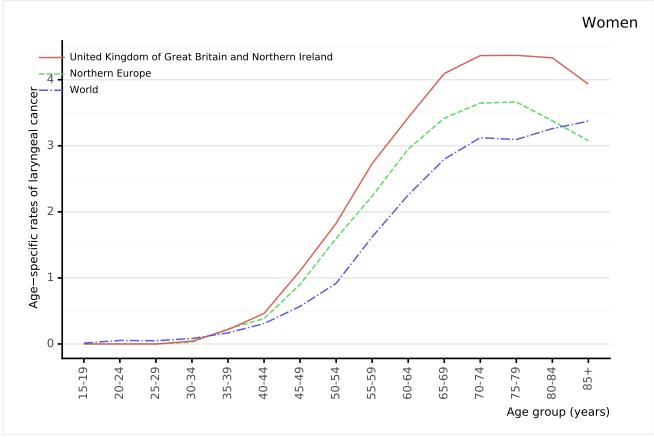


Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to  $\frac{\text{http://gco.iarc.fr/today/data-sources-methods}}{a} \text{ Rates per } 100,000 \text{ men per year.}$ 

9 ANNEX - 124 -

Figure 104: Comparison of age-specific laryngeal cancer incidence rates among women by age in United Kingdom of Great Britain and Northern Ireland, within the region, and the rest of world



Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a Rates per 100,000 women per year.

Data Sources:

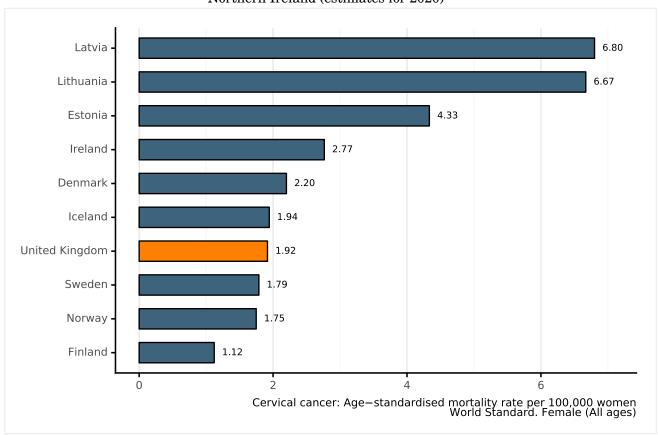
Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

9 ANNEX - 125 -

### 9.2 Mortality

## 9.2.1 Cervical cancer mortality in United Kingdom of Great Britain and Northern Ireland across Northern Europe

Figure 105: Age-standardised mortality rates of cervical cancer of United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

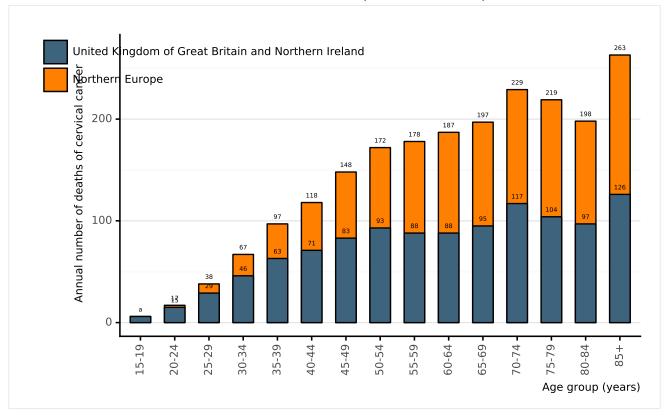


#### Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods Rates per 100,000 women per year.

9 ANNEX - 126 -

Figure 106: Annual number of deaths of cervical cancer by age group in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



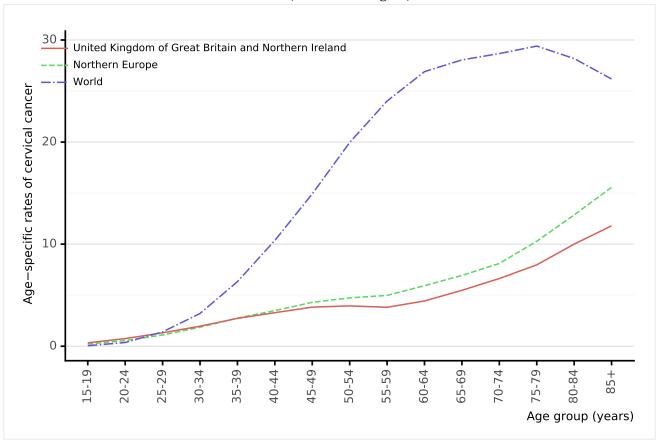
#### Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a 6 cases for United Kingdom of Great Britain and Northern Ireland and 6 cases for Northern Europe in the 15-19 age group.

9 ANNEX - 127 -

Figure 107: Comparison of age-specific cervical cancer mortality rates in United Kingdom of Great Britain and Northern Ireland, within the region, and the rest of world



### Data accessed on 27 Jan 2021

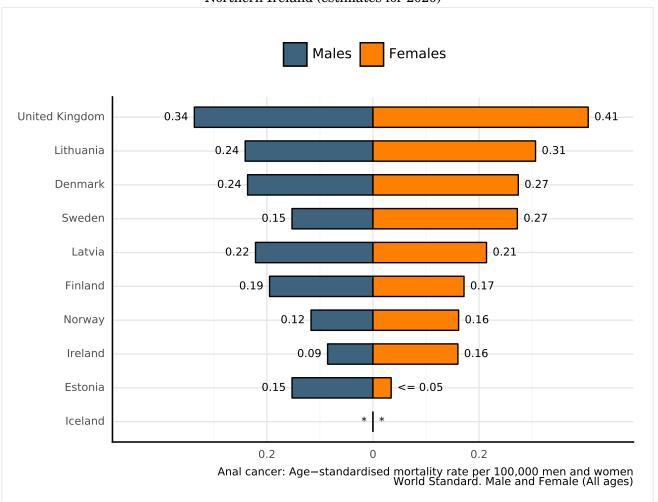
For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods a Rates per 100,000 women per year.

Data Sources:

9 ANNEX - 128 -

# 9.2.2 Anal cancer mortality in United Kingdom of Great Britain and Northern Ireland across **Northern Europe**

Figure 108: Age-standardised mortality rates of anal cancer of United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

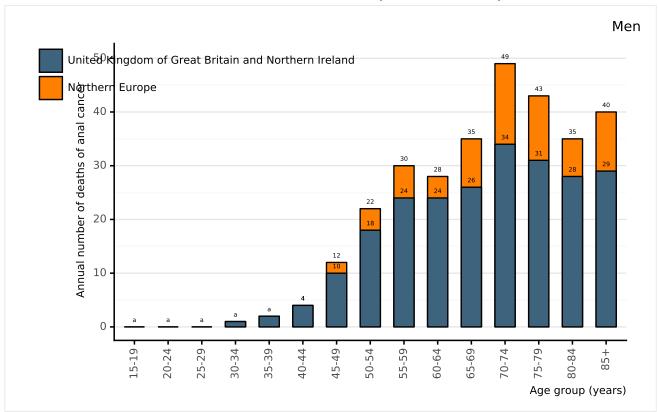
For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a Rates per 100,000 men per year.

b Rates per 100,000 women per year.
\* Rates are not available

9 ANNEX - 129 -

Figure 109: Annual number of deaths of anal cancer among men by age group in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

Data accessed on 27 Jan 2021

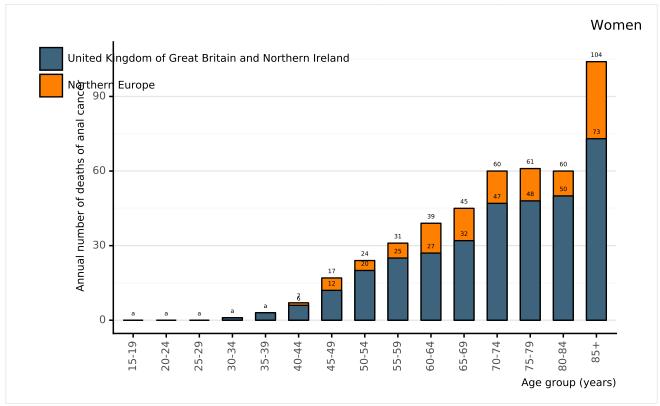
For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 15-19 age group. 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 20-24 age group. 0 cases for United Kingdom of Great Britain and Northern Europe in the 25-29 age group.

1 cases for United Kingdom of Great Britain and Northern Ireland and 1 cases for Northern Europe in the 30-34 age group. 2 cases for United Kingdom of Great Britain and Northern Ireland and 2 cases for Northern Europe in the 35-39 age group.

9 ANNEX - 130 -

Figure 110: Annual number of deaths of anal cancer among women by age group in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

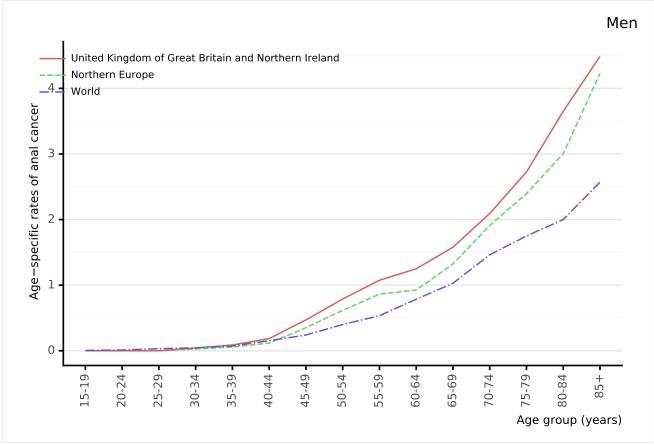
For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 15-19 age group. 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 20-24 age group. 0 cases for United Kingdom of Great Britain and Northern Europe in the 25-29 age group.

1 cases for United Kingdom of Great Britain and Northern Ireland and 1 cases for Northern Europe in the 30-34 age group. 3 cases for United Kingdom of Great Britain and Northern Ireland and 1 cases for Northern Europe in the 30-34 age group. Ireland and 3 cases for Northern Europe in the 35-39 age group.

9 ANNEX - 131 -

Figure 111: Comparison of age-specific anal cancer mortality rates among men by age in United Kingdom of Great Britain and Northern Ireland, within the region, and the rest of world

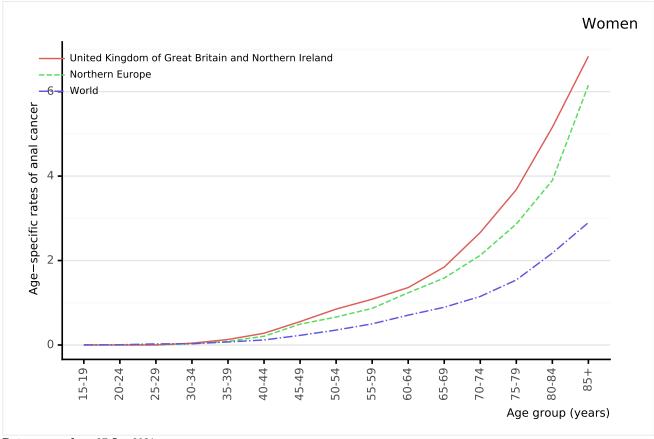


Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to  $\frac{\text{http://gco.iarc.fr/today/data-sources-methods}}{a} \text{ Rates per } 100,000 \text{ men per year.}$ 

9 ANNEX - 132 -

Figure 112: Comparison of age-specific anal cancer mortality rates among women by age in United Kingdom of Great Britain and Northern Ireland, within the region, and the rest of world



Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a Rates per 100,000 women per year.

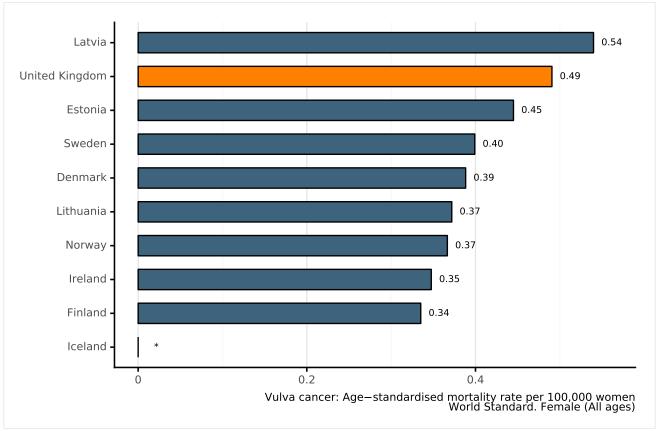
Data Sources:

Ferlay J. Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

9 ANNEX - 133 -

### 9.2.3 Vulva cancer mortality in United Kingdom of Great Britain and Northern Ireland across Northern Europe

Figure 113: Age-standardised mortality rates of vulva cancer of United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

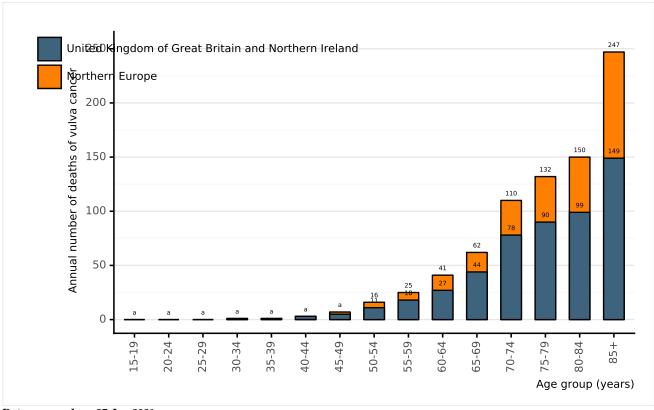


Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods a Rates per 100,000 women per year. \* Rates are not available

9 ANNEX - 134 -

Figure 114: Annual number of deaths of vulva cancer by age group in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 15-19 age group. 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 20-24 age group. 0 cases for United Kingdom of Great Britain and Northern Europe in the 25-29 age group.

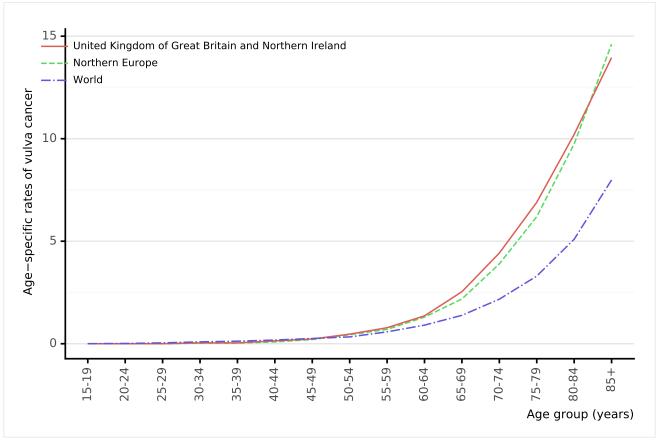
1 cases for United Kingdom of Great Britain and Northern Ireland and 1 cases for Northern Europe in the 35-39 age group. 3 cases for United Kingdom of Great Britain and Northern Ireland and 1 cases for Northern Europe in the 45-49 age group.

5 cases for United Kingdom of Great Britain and Northern Ireland and 7 cases for Northern Europe in the 45-49 age group.

That Sources.

9 ANNEX - 135 -

Figure 115: Comparison of age-specific vulva cancer mortality rates in United Kingdom of Great Britain and Northern Ireland, within the region, and the rest of world



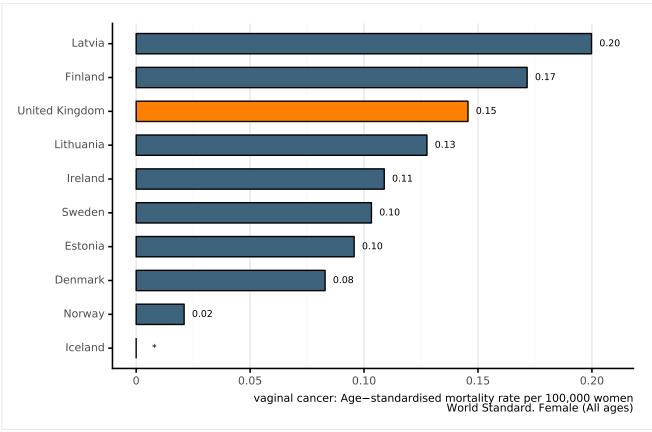
Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to  $\frac{1}{2} \frac{1}{2} \frac{1}$ 

9 ANNEX - 136 -

## 9.2.4 Vaginal cancer mortality in United Kingdom of Great Britain and Northern Ireland across Northern Europe

Figure 116: Age-standardised mortality rates of vaginal cancer of United Kingdom of Great Britain and Northern Ireland (estimates for 2020)

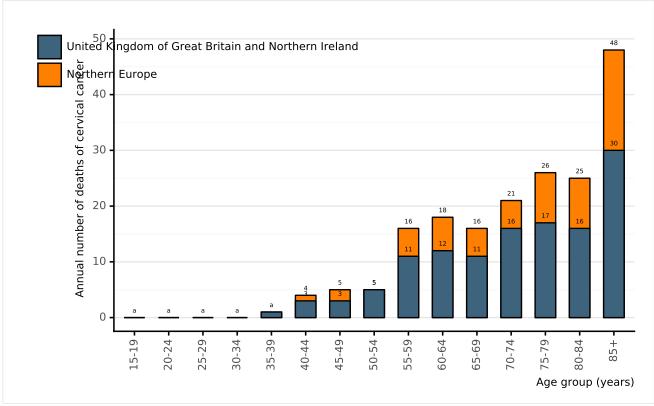


Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods a Rates per 100,000 women per year. \* Rates are not available

9 ANNEX - 137 -

Figure 117: Annual number of deaths of cervical cancer by age group in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

Data accessed on 27 Jan 2021

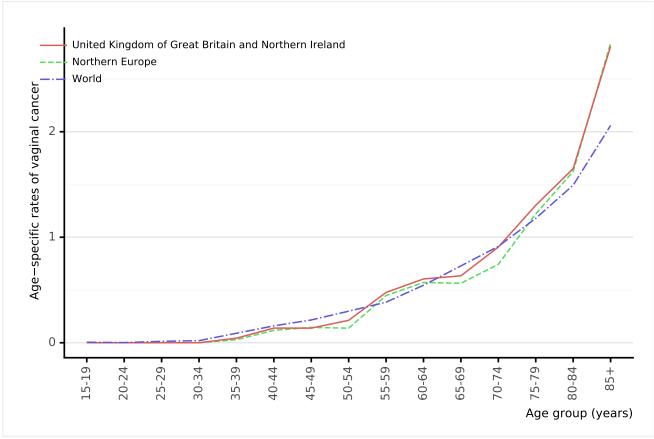
For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 15-19 age group. 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 20-24 age group. 0 cases for United Kingdom of Great Britain and Northern Europe in the 25-29 age group.

0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 30-34 age group. 1 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 30-34 age group. Ireland and 1 cases for Northern Europe in the 35-39 age group.

9 ANNEX - 138 -

Figure 118: Comparison of age-specific vaginal cancer mortality rates in United Kingdom of Great Britain and Northern Ireland, within the region, and the rest of world



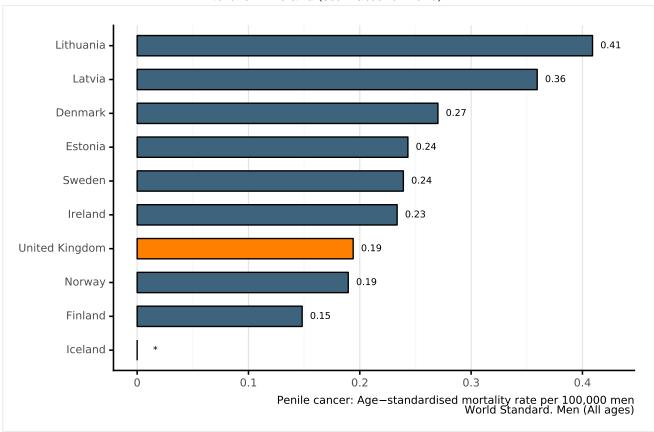
Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods <sup>a</sup> Rates per 100,000 women per year.

9 ANNEX - 139 -

### 9.2.5 Penile cancer mortality in United Kingdom of Great Britain and Northern Ireland across Northern Europe

Figure 119: Age-standardised mortality rates of penile cancer of United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



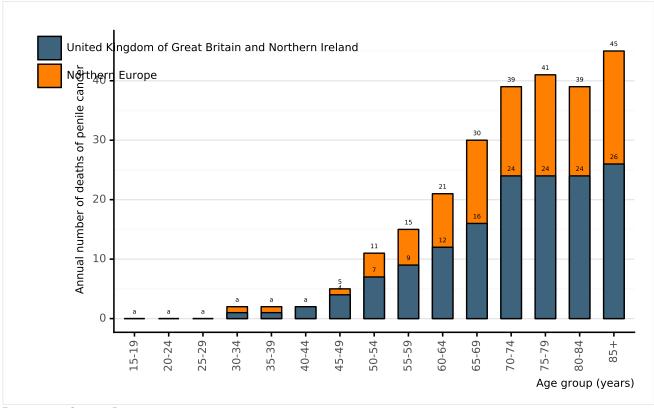
Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods a Rates per 100,000 men per year.

\* Rates are not available

9 ANNEX - 140 -

Figure 120: Annual number of new deaths of penile cancer by age group in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

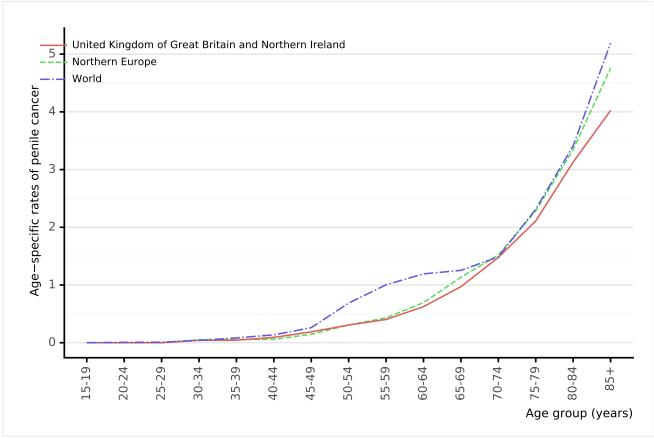
a 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 15-19 age group. 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 20-24 age group. 0 cases for United Kingdom of Great Britain and Northern Europe in the 25-29 age group.

1 cases for United Kingdom of Great Britain and Northern Ireland and 2 cases for Northern Europe in the 35-39 age group. 2 cases for Vnited Kingdom of Great Britain and Northern Europe in the 40-44 age group.

1 Cases for Northern Europe in the 35-39 age group. 2 cases for United Kingdom of Great Britain and Northern Ireland and 2 cases for Northern Europe in the 40-44 age group.

9 ANNEX - 141 -

Figure 121: Comparison of age-specific penile cancer mortality rates in United Kingdom of Great Britain and Northern Ireland, within the region, and the rest of world



Data accessed on 27 Jan 2021

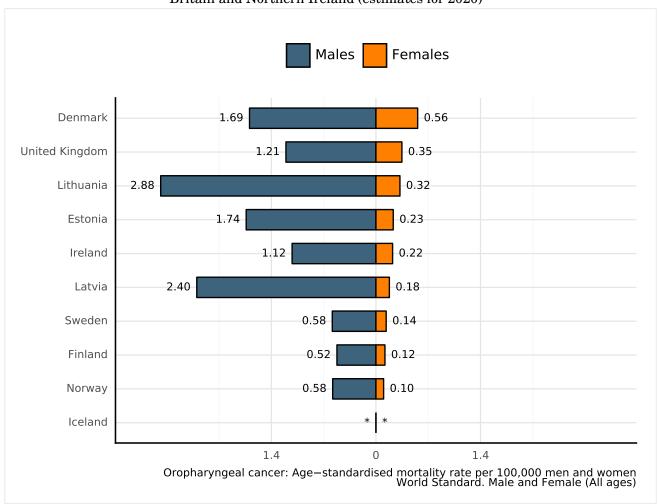
For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a Rates per 100,000 men per year.

9 ANNEX - 142 -

### 9.2.6 Oropharyngeal cancer mortality in United Kingdom of Great Britain and Northern **Ireland across Northern Europe**

Figure 122: Age-standardised mortality rates of oropharyngeal cancer of United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



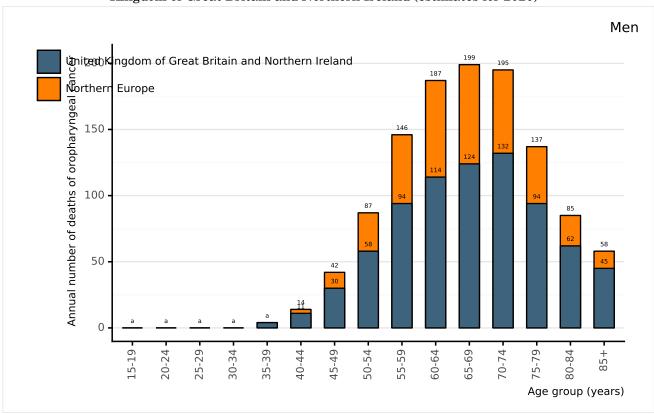
Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods Rates per 100,000 men per year.

b Rates per 100,000 women per year
Rates are not available

9 ANNEX - 143 -

Figure 123: Annual number of deaths of oropharyngeal cancer among men by age group in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

Data accessed on 27 Jan 2021

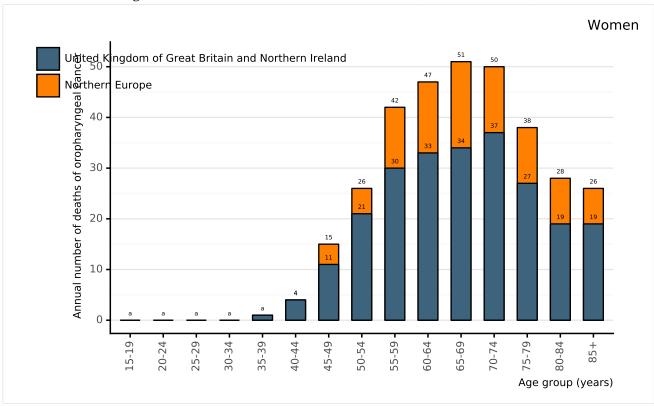
For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 15-19 age group. 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 20-24 age group. 0 cases for United Kingdom of Great Britain and Northern Europe in the 25-29 age group.

0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 25-29 age group. Ireland and 4 cases for Northern Europe in the 35-39 age group.

9 ANNEX - 144 -

Figure 124: Annual number of deaths of oropharyngeal cancer among women by age group in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



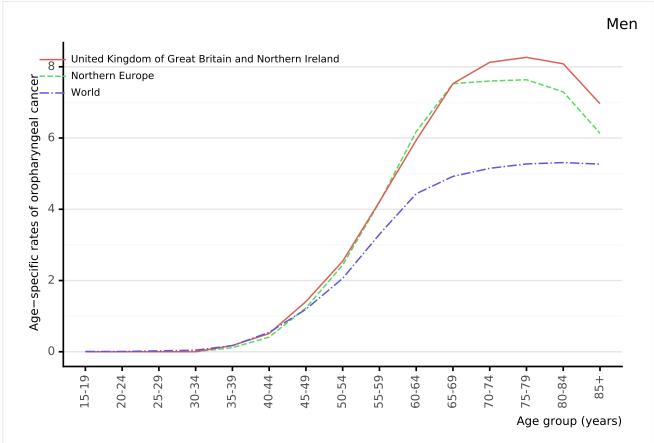
#### Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 15-19 age group. 0 cases for United Kingdom of Great Britain and Northern Europe in the 20-24 age group. 0 cases for United Kingdom of Great Britain and Northern Europe in the 25-29 age group.
0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 25-29 age group.
1 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 30-34 age group. 1 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 30-34 age group. Ireland and 1 cases for Northern Europe in the 35-39 age group.

9 ANNEX - 145 -

Figure 125: Comparison of age-specific oropharyngeal cancer mortality rates among men by age in United Kingdom of Great Britain and Northern Ireland, within the region, and the rest of world

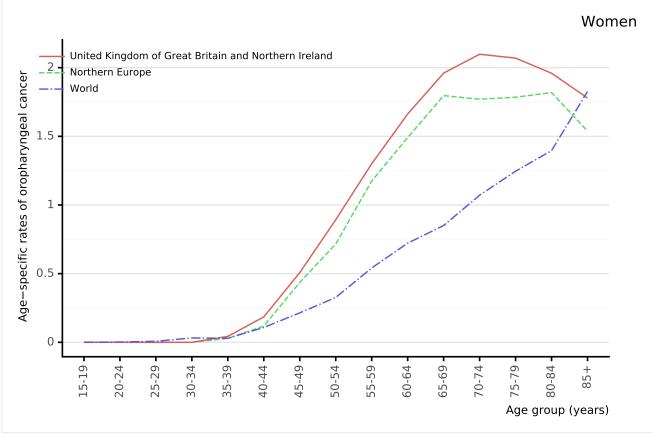


Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to  $\frac{\text{http://gco.iarc.fr/today/data-sources-methods}}{a} \text{ Rates per } 100,000 \text{ men per year.}$ 

9 ANNEX - 146 -

Figure 126: Comparison of age-specific oropharyngeal cancer mortality rates among women by age in United Kingdom of Great Britain and Northern Ireland, within the region, and the rest of world



Data accessed on 27 Jan 2021

Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

<sup>a</sup> Rates per 100,000 women per year.

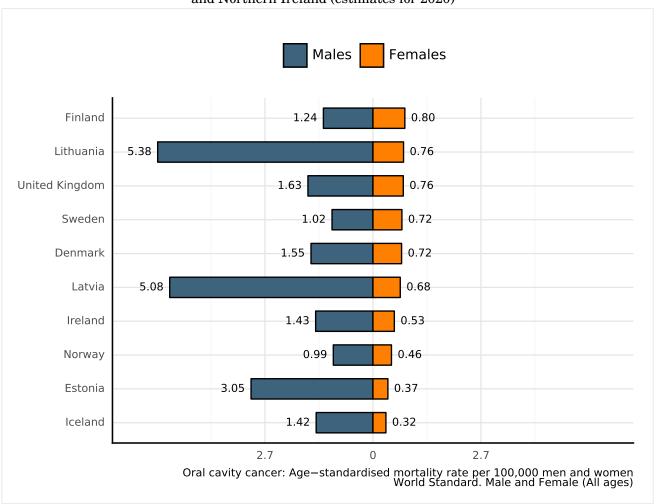
Data Sources:

Ferlay J, Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

9 ANNEX - 147 -

## 9.2.7 Oral cavity cancer mortality in United Kingdom of Great Britain and Northern Ireland across Northern Europe

Figure 127: Age-standardised mortality rates of oral cavity cancer of United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

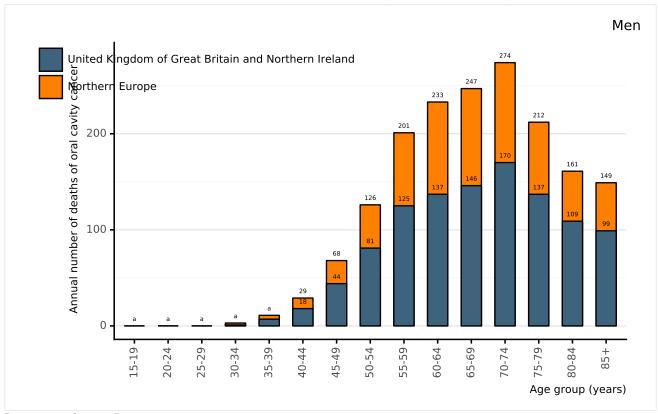
For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a Rates per 100,000 men per year.

b Rates per 100,000 women per year.

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Figure 128: Annual number of deaths of oral cavity cancer among men by age group in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

Data accessed on 27 Jan 2021

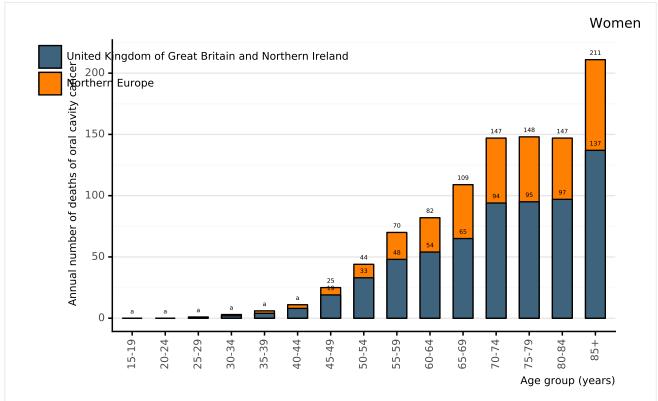
For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 15-19 age group. 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 20-24 age group. 0 cases for United Kingdom of Great Britain and Northern Europe in the 25-29 age group.

1 cases for United Kingdom of Great Britain and Northern Ireland and 3 cases for Northern Europe in the 30-34 age group. 7 cases for United Kingdom of Great Britain and Northern

9 ANNEX - 149 -

Figure 129: Annual number of deaths of oral cavity cancer among women by age group in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

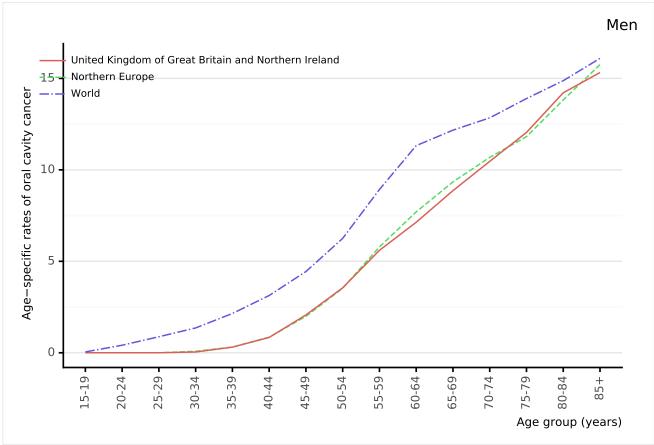
For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 15-19 age group. 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 20-24 age group. 0 cases for United Kingdom of Great Britain and Northern Europe in the 25-29 age group.

2 cases for United Kingdom of Great Britain and Northern Ireland and 3 cases for Northern Europe in the 30-34 age group. 4 cases for United Kingdom of Great Britain and Northern Ireland and 6 cases for Northern Europe in the 35-39 age group. 8 cases for United Kingdom of Great Britain and Northern Ireland and 11 cases for Northern Europe in the 40-44 age group.

9 ANNEX - 150 -

Figure 130: Comparison of age-specific oral cavity cancer mortality rates among men by age in United Kingdom of Great Britain and Northern Ireland, within the region, and the rest of world

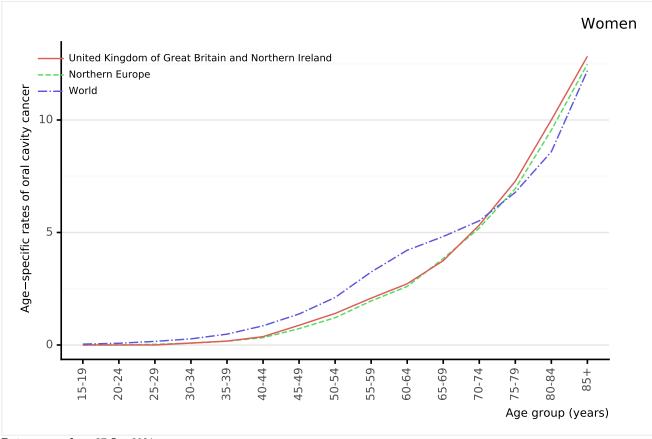


Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to  $\frac{\text{http://gco.iarc.fr/today/data-sources-methods}}{a} \text{ Rates per } 100,000 \text{ men per year.}$ 

9 ANNEX - 151 -

Figure 131: Comparison of age-specific oral cavity cancer mortality rates among women by age in United Kingdom of Great Britain and Northern Ireland, within the region, and the rest of world



Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a Rates per 100,000 women per year.

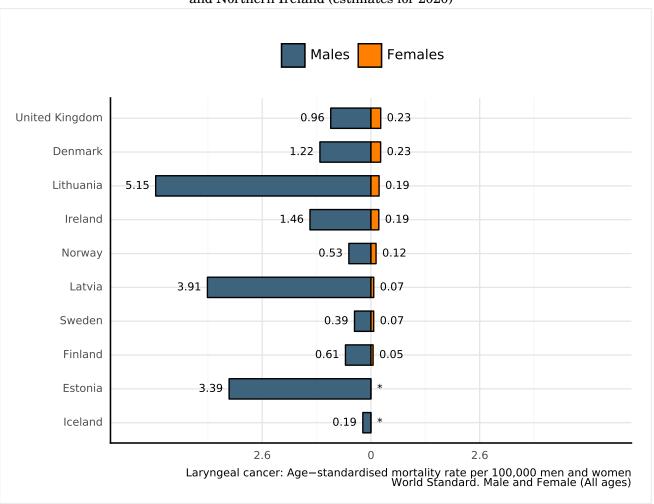
Data Sources:

Ferlay J. Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

9 ANNEX - 152 -

## 9.2.8 Laryngeal cancer mortality in United Kingdom of Great Britain and Northern Ireland across Northern Europe

Figure 132: Age-standardised mortality rates of laryngeal cancer of United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



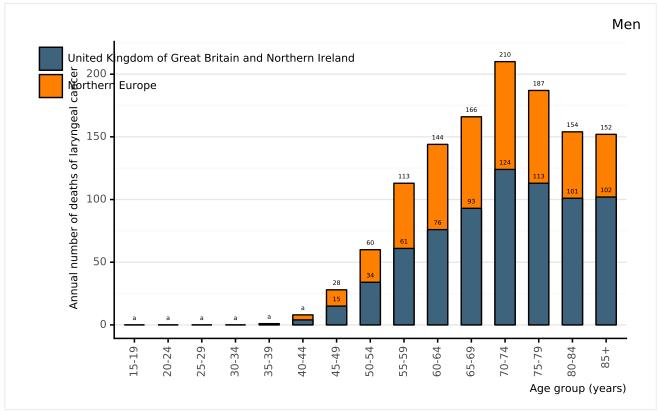
Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to  $\texttt{http://gco.iarc.fr/today/data-sources-methods}^a$  Rates per 100,000 men per year.

b Rates per 100,000 women per year
Rates are not available

9 ANNEX - 153 -

Figure 133: Annual number of deaths of laryngeal cancer among men by age group in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

Data accessed on 27 Jan 2021

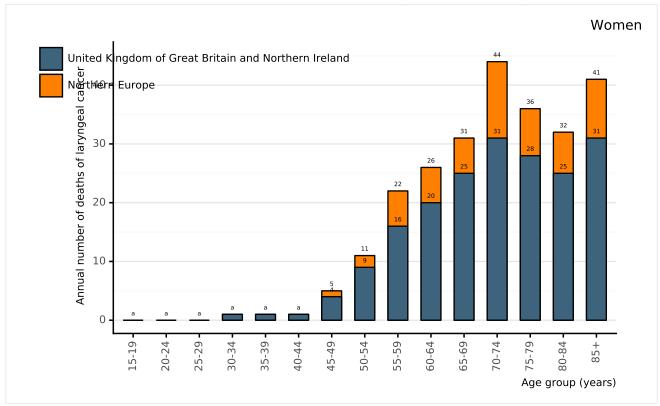
For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 15-19 age group. 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 20-24 age group. 0 cases for United Kingdom of Great Britain and Northern Europe in the 25-29 age group.

0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 30-34 age group. 0 cases for United Kingdom of Great Britain and Northern Ireland and 1 cases for Northern Europe in the 35-39 age group. 4 cases for United Kingdom of Great Britain and Northern Ireland and 8 cases for Northern Europe in the 40-44 age group.

9 ANNEX - 154 -

Figure 134: Annual number of deaths of laryngeal cancer among women by age group in United Kingdom of Great Britain and Northern Ireland (estimates for 2020)



Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

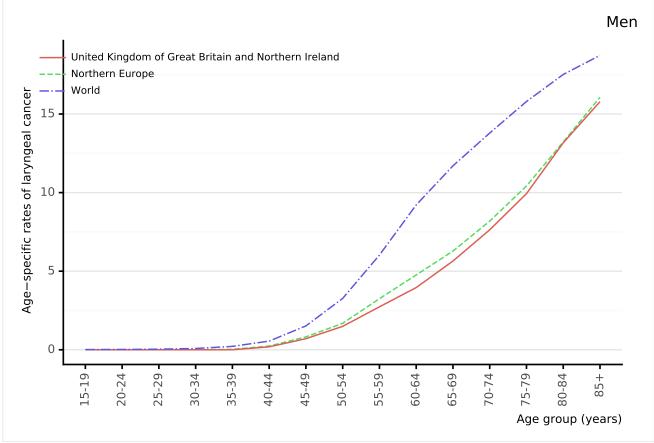
a 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 15-19 age group. 0 cases for United Kingdom of Great Britain and Northern Ireland and 0 cases for Northern Europe in the 20-24 age group. 0 cases for United Kingdom of Great Britain and Northern Europe in the 25-29 age group.

1 cases for United Kingdom of Great Britain and Northern Ireland and 1 cases for Northern Europe in the 35-39 age group. 1 cases for United Kingdom of Great Britain and Northern Europe in the 40-44 age group.

1 cases for Northern Europe in the 35-39 age group. 1 cases for United Kingdom of Great Britain and Northern Europe in the 40-44 age group.

9 ANNEX - 155 -

Figure 135: Comparison of age-specific laryngeal cancer mortality rates among men by age in United Kingdom of Great Britain and Northern Ireland, within the region, and the rest of world

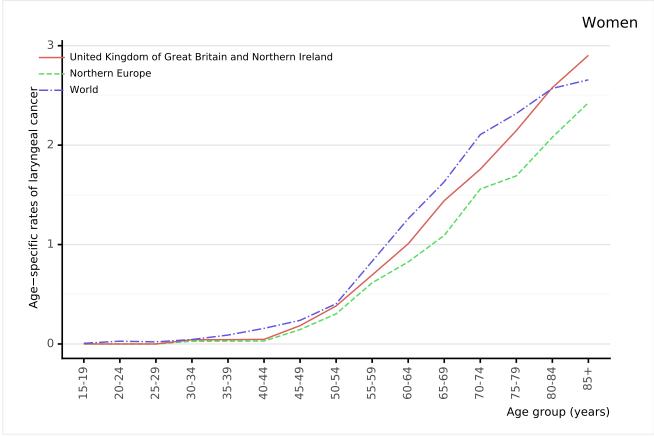


Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to  $\frac{\text{http://gco.iarc.fr/today/data-sources-methods}}{a} \text{ Rates per } 100,000 \text{ men per year.}$ 

9 ANNEX - 156 -

Figure 136: Comparison of age-specific laryngeal cancer mortality rates among women by age in United Kingdom of Great Britain and Northern Ireland, within the region, and the rest of world



Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods

a Rates per 100,000 women per year.

Data Sources:

Ferlay J. Ervik M, Lam F, Colombet M, Mery L, Piñeros M, Znaor A, Soerjomataram I, Bray F (2020). Global Cancer Observatory: Cancer Today. Lyon, France: International Agency for Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

10 GLOSSARY -157-

## 10 Glossary

Table 49: Glossary

Term	Definition
Incidence	Incidence is the number of new cases arising in a given period in a specified population. This information is collected routinely by cancer registries. It can be expressed as an absolute number of cases per year or as a rate per 100,000 persons per year (see Crude rate and ASR below). The rate provides an approximation of the average risk of developing a cancer.
Mortality	Mortality is the number of deaths occurring in a given period in a specified population. It can be expressed as an absolute number of deaths per year or as a rate per 100,000 persons per year.
Prevalence	The prevalence of a particular cancer can be defined as the number of persons in a defined population who have been diagnosed with that type of cancer, and who are still alive at the end of a given year, the survivors. Complete prevalence represents the number of persons alive at certain point in time who previously had a diagnosis of the disease, regardless of how long ago the diagnosis was, or if the patient is still under treatment or is considered cured. Partial prevalence, which limits the number of patients to those diagnosed during a fixed time in the past, is a particularly useful measure of cancer burden. Prevalence of cancers based on cases diagnosed within one, three and five are presented as they are likely to be of relevance to the different stages of cancer therapy, namely, initial treatment (one year), clinical follow-up (three years) and cure (five years). Patients who are still alive five years after diagnosis are usually considered cured since the death rates of such patients are similar to those in the general population. There are exceptions, particularly breast cancer. Prevalence is presented for the adult population only (ages 15 and over), and is available both as numbers and as proportions per 100,000 persons.
Crude rate	Data on incidence or mortality are often presented as rates. For a specific tumour and population, a crude rate is calculated simply by dividing the number of new cancers or cancer deaths observed during a given time period by the corresponding number of person years in the population at risk. For cancer, the result is usually expressed as an annual rate per 100,000 persons at risk.
ASR (age-standardised rate)	An age-standardised rate (ASR) is a summary measure of the rate that a population would have if it had a standard age structure. Standardization is necessary when comparing several populations that differ with respect to age because age has a powerful influence on the risk of cancer. The ASR is a weighted mean of the age-specific rates; the weights are taken from population distribution of the standard population. The most frequently used standard population is the World Standard Population. The calculated incidence or mortality rate is then called age-standardised incidence or mortality rate (world). It is also expressed per 100,000. The world standard population used in GLOBOCAN is as proposed by Segi [1] and modified by Doll and al. [2]. The age-standardised rate is calculated using 10 age-groups. The result may be slightly different from that computed using the same data categorised using the traditional 5 year age bands.

Continued on next page

10 GLOSSARY -158-

Table 49 - continued from previous page

Cumulative risk  Cumulative risk of individuals getting/dying from the disease during a specified period. For cancer, it is expressed as the number of new born children (out of 100, or 1000) who would be expected to develop/die from a particular cancer before the age of 75 if they had the rates of cancer observed in the period in the absence of competing causes.  Cytologically normal women  SIL and CIN are two commonly used terms to describe precancerous lesions or the abnormal growth of squamous cells observed in the cervix. SIL is an abnormal result derived from cervical cytological screening or Pap smear testing. CIN is a histological diagnosis made upon analysis of cervical tissue obtained by biopsy or surgical excision. The condition is graded as CIN 1, 2 or 3, according to the thickness of the abnormal epithelium (1/3, 2/3 or the entire thickness).  Low-grade cervical lesions are defined by early changes in size, shape, and number of ab-normal cells formed on the surface of the cervix and may be referred to as mild dysplasia, LSIL, or CIN-1.  High-grade cervical lesions are defined by a large number of precancerous cells on the sur-face of the cervix that are distinctly different from normal cells. They have the potential to become cancerous cells and invade deeper tissues of the cervix. These lesions may be referred to as moderate or severe tissues of the cervix. These lesions may be referred to as moderate or severe	Table 45 - continued from previous page	
Cumulative risk biting/dying from the disease during a specified period. For cancer, it is expressed as the number of new born children (out of 100, or 1000) who would be expected to develop/die from a particular cancer before the age of 75 if they had the rates of cancer observed in the period in the absence of competing causes.  Cytologically normal women  SIL and CIN are two commonly used terms to describe precancerous lesions or the abnormal growth of squamous cells observed in the cervix. SIL is an abnormal result derived from cervical cytological screening or Pap smear testing. CIN is a histological diagnosis made upon analysis of cervical tissue obtained by biopsy or surgical excision. The condition is graded as CIN 1, 2 or 3, according to the thickness of the abnormal epithelium (1/3, 2/3 or the entire thickness).  Low-grade cervical lesions are defined by early changes in size, shape, and number of ab-normal cells formed on the surface of the cervix and may be referred to as mild dysplasia, LSIL, or CIN-1.  High-grade cervical lesions are defined by a large number of precancerous cells on the sur-face of the cervix that are distinctly different from normal cells. They have the potential to become cancerous cells and invade deeper tissues of the cervix. These lesions may be referred to as moderate or severe	Term	Definition
SIL and CIN are two commonly used terms to describe precancerous lesions or the abnormal growth of squamous cells observed in the cervix. SIL is an abnormal result derived from cervical cytological screening or Pap smear testing. CIN is a histological diagnosis made upon analysis of cervical tissue obtained by biopsy or surgical excision. The condition is graded as CIN 1, 2 or 3, according to the thickness of the abnormal epithelium (1/3, 2/3 or the entire thickness).  Low-grade cervical lesions are defined by early changes in size, shape, and number of ab-normal cells formed on the surface of the cervix and may be referred to as mild dysplasia, LSIL, or CIN-1.  High-grade cervical lesions are defined by a large number of precancerous cells on the sur-face of the cervix that are distinctly different from normal cells. They have the potential to become cancerous cells and invade deeper tissues of the cervix. These lesions may be referred to as moderate or severe	Cumulative risk	ting/dying from the disease during a specified period. For cancer, it is expressed as the number of new born children (out of 100, or 1000) who would be expected to develop/die from a particular cancer before the age of 75 if they had the rates of cancer observed in the period in the absence of com-
Cervical Intraepithe-lial Neoplasia (CIN) / Squamous Intraepithe-lial Lesions (SIL)  Squamous Intraepithe-lial Lesions (SIL)  Low-grade cervical lesions (LSIL/CIN-1)  Low-grade cervical lesions (LSIL/CIN-1)  High-grade cervical lesions are defined by a large number of precancerous cells on the sur-face of the cervix that are distinctly different from normal cells. They have the potential to become cancerous cells and invade deeper tissues of the cervix. SIL is an abnormal growth of squamous cells observed in the cervix. SIL is an abnormal growth of squamous cells observed in the cervix. SIL is an abnormal cells formed upon analysis of cervical tissue obtained by biopsy or surgical excision. The condition is graded as CIN 1, 2 or 3, according to the thickness of the abnormal epithelium (1/3, 2/3 or the entire thickness).  Low-grade cervical lesions are defined by early changes in size, shape, and number of ab-normal cells formed on the surface of the cervix and may be referred to as mild dysplasia, LSIL, or CIN-1.  High-grade cervical lesions are defined by a large number of precancerous cells on the sur-face of the cervix that are distinctly different from normal cells. They have the potential to become cancerous cells and invade deeper tissues of the cervix. These lesions may be referred to as moderate or severe		No abnormal cells are observed on the surface of their cervix upon cytology.
number of ab-normal cells formed on the surface of the cervix and may be referred to as mild dysplasia, LSIL, or CIN-1.  High-grade cervical lesions are defined by a large number of precancerous cells on the surface of the cervix that are distinctly different from normal cells. They have the potential to become cancerous cells and invade deeper tissues of the cervix. These lesions may be referred to as moderate or severe	lial Neoplasia (CIN) / Squamous Intraepithe-	or the abnormal growth of squamous cells observed in the cervix. SIL is an abnormal result derived from cervical cytological screening or Pap smear testing. CIN is a histological diagnosis made upon analysis of cervical tissue obtained by biopsy or surgical excision. The condition is graded as CIN 1, 2 or 3, according to the thickness of the abnormal epithelium (1/3, 2/3 or
High-grade cervical lesions (HSIL / CIN-2 / CIN-3 / CIS) cells on the sur-face of the cervix that are distinctly different from normal cells. They have the potential to become cancerous cells and invade deeper tissues of the cervix. These lesions may be referred to as moderate or severe	O .	number of ab-normal cells formed on the surface of the cervix and may be
	sions (HSIL / CIN-2 / CIN-3 / CIS)	cells on the sur-face of the cervix that are distinctly different from normal cells. They have the potential to become cancerous cells and invade deeper tissues of the cervix. These lesions may be referred to as moderate or severe dysplasia, HSIL, CIN-2, CIN-3 or cervical carcinoma in situ (CIS).
Carcinoma in situ Preinvasive malignancy limited to the epithelium without invasion of the (CIS) basement membrane. CIN 3 encompasses the squamous carcinoma in situ.		
Invasive cervical cancer (ICC) / Cervical cancer (ICC)	cer (ICC) / Cervical	called ICC. ICC stages range from stage I (cancer is in the cervix or uterus only) to stage IV (the cancer has spread to distant organs, such as the liver).
Adenocarcinoma Invasive tumour with glandular and squamous elements intermingled	Adenocarcinoma	Invasive tumour with glandular and squamous elements intermingled

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# Cancer Epidemiology Research Program, Catalan Institute of Oncology (ICO), Institut d'Investigació Biomèdica de Bellvitge (IDIBELL), in alphabetic order

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**International Agency for Research on Cancer (IARC)** 

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