

MATION CENTRE ON HPV AND **Human Papillomavirus** and **Related Diseases Report**



JIVARC IN

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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 Poland 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			
Women at risk for cervical cancer (Female population aged >=15 y	rs)		16.9 millior
Burden of cervical cancer and other HPV-related cancers	15)		10.0 1111101
Annual number of cervical cancer cases			3862
Annual number of cervical cancer deaths			2137
Crude incidence rates per 100,000 population:		Male	Female
	al cancer	-	19.8
	al cancer	1.18	1.44
	va cancer		3.90
	al cancer		0.55
•	le cancer	2.40	
Oropharynge		6.87	2.05
Oral cavi		16.3	6.68
	al cancer	14.8	2.17
Burden of cervical HPV infection			
Prevalence (%) of HPV 16 and/or HPV 18 among women with:			
		Normal cytology	3.4
Low-gra	ade cervical le	sions (LSIL/CIN-1)	27.1
High-grade cervical	lesions (HSII	/CIN-2/CIN-3/CIS)	54.5
		Cervical cancer	88.1
Other factors contributing to cervical cancer			
Smoking prevalence (%) [95% UI], women			21.7 [17-26.3]
Total fertility rate (live births per women)			1.5
Oral contraceptive use (%)			17
HIV prevalence (%) [95% UI], women (15-49 years)			- [—
Sexual behaviour			
Percentage of 15-year-old who have had sexual intercourse (men/v	vomen)		16.0/18.0
Range of median age at first sexual intercourse (men/women)			17.0/18.0-18.7
Cervical screening practices and recommendations			
Existence of official national recommendations			Yes
Starting year of current recommendations			2016
Active invitation to screening	· · · 1 · C		No
Screening ages (years), primary screening test used, and screening	g interval or h	requency of screen-	25-59 (cytology, 3
ings HPV vaccine in females			years
HPV vaccine in temates HPV vaccination programme			
Year of introduction			
Year of estimation of HPV vaccination coverage			
HPV coverage – first dose (%)			
HPV coverage – last dose (%)			
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Contents

E	xecu	tive su	Immary	iii
1	Inti	roduct	ion	2
2	Der	nograj	phic and socioeconomic factors	4
3	Bur	den o	f HPV related cancers	5
	3.1	HPV	related cancers incidence	5
	3.2	HPV	related cancers mortality	7
	3.3	Cervie		9
		3.3.1	Cervical cancer incidence in Poland	9
		3.3.2	Cervical cancer incidence by histology in Poland	12
		3.3.3	Cervical cancer mortality in Poland	14
		3.3.4	Cervical cancer incidence and mortality comparison in Poland	16
	3.4	Anoge		18
		3.4.1		18
			3.4.1.1 Anal cancer incidence in Poland	
			3.4.1.2 Anal cancer mortality in Poland	
			3.4.1.3 Anal cancer incidence and mortality comparison in Poland	
		3.4.2	Vulva cancer	
			3.4.2.1 Vulva cancer incidence in Poland	
			3.4.2.2 Vulva cancer mortality in Poland	
			3.4.2.3 Vulva cancer incidence and mortality comparison in Poland	
		3.4.3	Vaginal cancer	
			3.4.3.1 Vaginal cancer incidence in Poland	
			3.4.3.2 Vaginal cancer mortality in Poland	
		044	3.4.3.3 Vaginal cancer incidence and mortality comparison in Poland	
		3.4.4	Penile cancer	
			3.4.4.1 Penile cancer incidence in Poland	
			3.4.4.2Penile cancer mortality in Poland	
	3.5	Head	and neck cancers	
	3.0	пеац 3.5.1		
		5.5.1	Oropharyngeal cancer	38
			3.5.1.1Oropharyngeal cancer incidence in Poland	
			3.5.1.2 Oropharyngeal cancer incidence and mortality comparison in Poland	
		3.5.2	Oral cavity cancer	
		0.0.2	3.5.2.1 Oral cavity cancer incidence in Poland	
			3.5.2.2 Oral cavity cancer incidence and mortality comparison in Poland	
			3.5.2.3 Oral cavity cancer incidence and mortality comparison in Poland	
		3.5.3	Laryngeal cancer	
		0.0.0	3.5.3.1 Laryngeal cancer incidence in Poland	
			3.5.3.2 Laryngeal cancer incidence and mortality comparison in Poland	
			3.5.3.3 Laryngeal cancer incidence and mortality comparison in Poland	
4				53
	4.1		burden in women with normal cervical cytology, cervical precancerous lesions or	
				53
				54
		4.1.2	HPV type distribution among women with normal cervical cytology, precancerous	
				55
		4.1.3	HPV type distribution among HIV+ women with normal cervical cytology	65

	4.2 4.3 4.4	4.1.4 Terminology6HPV burden in anogenital cancers other than cervix64.2.1 Anal cancer and precancerous anal lesions64.2.2 Vulvar cancer and precancerous vulvar lesions74.2.3 Vaginal cancer and precancerous vaginal lesions74.2.4 Penile cancer and precancerous penile lesions7HPV burden in men7HPV burden in the head and neck74.4.1 Burden of oral HPV infection in healthy population74.4.2 HPV burden in head and neck cancers7	67 68 70 72 74 76 77 77
5	Fac	tors contributing to cervical cancer 8	80
6	Sex	ual and reproductive health behaviour indicators	81
7	HP	V preventive strategies	84
	7.1	Cervical cancer screening practices	34
	7.2	HPV vaccination	36
8	Pro	tective factors for cervical cancer	88
9	Anr	nex 8	89
	9.1	Incidence	39
		9.1.1 Cervical cancer incidence in Poland across Eastern Europe	
		9.1.2 Anal cancer incidence in Poland across Eastern Europe	
		9.1.3 Vulva cancer incidence in Poland across Eastern Europe	
		9.1.4 Vaginal cancer incidence in Poland across Eastern Europe	
		9.1.5 Penile cancer incidence in Poland across Eastern Europe	
		9.1.6 Oropharyngeal cancer incidence in Poland across Eastern Europe	
		 9.1.7 Oral cavity cancer incidence in Poland across Eastern Europe 9.1.8 Laryngeal cancer incidence in Poland across Eastern Europe 1.1 	
	9.2	Mortality	
	9.4	9.2.1 Cervical cancer mortality in Poland across Eastern Europe	
		9.2.2 Anal cancer mortality in Poland across Eastern Europe	
		9.2.3 Vulva cancer mortality in Poland across Eastern Europe	
		9.2.4 Vaginal cancer mortality in Poland across Eastern Europe	
		9.2.5 Penile cancer mortality in Poland across Eastern Europe	
		9.2.6 Oropharyngeal cancer mortality in Poland across Eastern Europe	
		9.2.7 Oral cavity cancer mortality in Poland across Eastern Europe	
		9.2.8 Laryngeal cancer mortality in Poland across Eastern Europe	48

10 Glossary

153

List of Figures

1	Poland and Eastern Europe	2
2	Population pyramid of Poland for 2022	4
3	Population trends in four selected age groups in Poland	4
4	Comparison of HPV related cancers incidence to other cancers in men and women of all ages in Poland (estimates	_
-	for 2020)	5
5	Comparison of HPV related cancers incidence to other cancers among men and women 15-44 years of age in	c
G	Poland (estimates for 2020)	6
6	for 2020)	7
7	Comparison of HPV related cancers mortality to other cancers among men and women 15-44 years of age in	'
•	Poland (estimates for 2020)	8
8	Age-specific incidence rates of cervical cancer in Poland (estimates for 2020)	11
9	Annual number of new cases of cervical cancer in Poland (estimates for 2020)	11
10	Time trends in cervical cancer incidence in Poland (cancer registry data)	13
11	Age-specific mortality rates of cervical cancer in Poland (estimates for 2020)	15
12	Annual number of deaths of cervical cancer in Poland (estimates for 2020)	15
13	Comparison of age-specific cervical cancer incidence and mortality rates in Poland (estimates for 2020)	16
14	Comparison of annual premature deaths and disability from cervical cancer in Poland to other cancers among	
	women (estimates for 2019)	17
15	Age-specific incidence rates of anal cancer in Poland (estimates for 2020)	19
16	Annual number of new cases of anal cancer in Poland (estimates for 2020)	19
17	Age-specific mortality rates of anal cancer in Poland (estimates for 2020)	21
18	Annual number of deaths of of anal cancer in Poland (estimates for 2020)	21
19	Comparison of age-specific anal cancer incidence and mortality rates among men in Poland (estimates for 2020)	22
20	Comparison of age-specific anal cancer incidence and mortality rates among women in Poland (estimates for 2020 Age-specific incidence rates of vulva cancer in Poland (estimates for 2020)) 22 24
21 22	Age-specific incidence rates of vulva cancer in Poland (estimates for 2020)	$\frac{24}{24}$
22	Age-specific mortality rates of vulva cancer in Poland (estimates for 2020)	24 26
23 24	Annual number of deaths of vulva cancer in Poland (estimates for 2020)	20 26
2 4 25	Comparison of age-specific vulva cancer incidence and mortality rates in Poland (estimates for 2020)	20 27
26	Age-specific incidence rates of vaginal cancer in Poland (estimates for 2020)	29
27	Annual number of new cases of vaginal cancer in Poland (estimates for 2020)	29
28	Age-specific mortality rates of vaginal cancer in Poland (estimates for 2020)	31
29	Annual number of deaths of vaginal cancer in Poland (estimates for 2020)	31
30	Comparison of age-specific vaginal cancer incidence and mortality rates in Poland (estimates for 2020)	32
31	Age-specific incidence rates of penile cancer in Poland (estimates for 2020)	34
32	Annual number of new cases of penile cancer in Poland (estimates for 2020)	34
33	Age-specific mortality rates of penile cancer in Poland (estimates for 2020)	36
34	Annual number of deaths of penile cancer in Poland (estimates for 2020)	36
35	Comparison of age-specific penile cancer incidence and mortality rates in Poland (estimates for 2020)	
36	Age-specific incidence rates of oropharyngeal cancer in Poland (estimates for 2020)	39
37	Annual number of new cases of oropharyngeal cancer in Poland (estimates for 2020)	39
38	Age-specific mortality rates of oropharyngeal cancer in Poland (estimates for 2020)	41
39	Annual number of deaths of oropharyngeal cancer in Poland (estimates for 2020)	41
40	Comparison of age-specific oropharyngeal cancer incidence and mortality rates among men in Poland (estimates for 2020)	42
41	Comparison of age-specific oropharyngeal cancer incidence and mortality rates among women in Poland (esti-	42
41	mates for 2020)	42
42	Age-specific incidence rates of oral cavity cancer in Poland (estimates for 2020)	44
43	Annual number of new cases of oral cavity cancer in Poland (estimates for 2020)	44
44	Age-specific mortality rates of oral cavity cancer in Poland (estimates for 2020)	46
45	Annual number of deaths of oral cavity cancer in Poland (estimates for 2020)	46
46	Comparison of age-specific oral cavity cancer incidence and mortality rates among men in Poland (estimates for	
	2020)	47
47	Comparison of age-specific oral cavity cancer incidence and mortality rates among women in Poland (estimates	
	for 2020)	47
4 8	Age-specific incidence rates of laryngeal cancer in Poland (estimates for 2020)	49
49	Annual number of new cases of laryngeal cancer in Poland (estimates for 2020)	49
50	Age-specific mortality rates of laryngeal cancer in Poland (estimates for 2020)	51
51	Annual number of deaths of of laryngeal cancer in Poland (estimates for 2020)	51
52	Comparison of age-specific laryngeal cancer incidence and mortality rates among men in Poland (estimates for	- -
	2020)	52

53	Comparison of age-specific laryngeal cancer incidence and mortality rates among women in Poland (estimates for 2020)	52
54	Crude age-specific HPV prevalence (%) and 95% confidence interval in women with normal cervical cytology in	
	Poland	54
55	HPV prevalence among women with normal cervical cytology in Poland, by study	54
56	HPV 16 prevalence among women with normal cervical cytology in Poland, by study	55
57	HPV 16 prevalence among women with low-grade cervical lesions in Poland, by study	56
58	HPV 16 prevalence among women with high-grade cervical lesions in Poland, by study	56
59	HPV 16 prevalence among women with invasive cervical cancer in Poland, by study	57
60	Comparison of the ten most frequent HPV oncogenic types in Poland among women with and without cervical	
	lesions	58
61	Comparison of the ten most frequent HPV oncogenic types in Poland among women with invasive cervical cancer	
	by histology	60
62	Comparison of the ten most frequent HPV types in anal cancer cases in Europe and the World	69
63	Comparison of the ten most frequent HPV types in AIN 2/3 cases in Europe and the World	69
64	Comparison of the ten most frequent HPV types in cases of vulvar cancer in Europe and the World	71
65	Comparison of the ten most frequent HPV types in VIN 2/3 cases in Europe and the World	71
66	Comparison of the ten most frequent HPV types in cases of vaginal cancer in Europe and the World	73
67	Comparison of the ten most frequent HPV types in VaIN 2/3 cases in Europe and the World	73
68	Comparison of the ten most frequent HPV types in cases of penile cancer in Europe and the World	75
69	Comparison of the ten most frequent HPV types in PeIN 2/3 cases in Europe and the World	75 75
	Estimated coverage* of cervical cancer screening in Poland	85
70		
71	HPV vaccination coverage in females by year in Poland	86
72	HPV vaccination coverage in males by year in Poland	87
73	Age-standardised incidence rates of cervical cancer of Poland (estimates for 2020)	89
74	Annual number of new cases of cervical cancer by age group in Poland (estimates for 2020)	90
75	Comparison of age-specific cervical cancer incidence rates in Poland, within the region, and the rest of world	91
76	Age-standardised incidence rates of anal cancer of Poland (estimates for 2020)	92
77	Annual number of new cases of anal cancer among men by age group in Poland (estimates for 2020)	93
78	Annual number of new cases of anal cancer among women by age group in Poland (estimates for 2020)	94
79	Comparison of age-specific anal cancer incidence rates among men by age in Poland, within the region, and the	
	rest of world	95
80	Comparison of age-specific anal cancer incidence rates among women by age in Poland, within the region, and	
	the rest of world	96
81	Age-standardised incidence rates of vulva cancer of Poland (estimates for 2020)	97
82	Annual number of new cases of vulva cancer by age group in Poland (estimates for 2020)	98
83	Comparison of age-specific vulva cancer incidence rates in Poland, within the region, and the rest of world	99
84	Age-standardised incidence rates of vaginal cancer of Poland (estimates for 2020)	100
85	Annual number of new cases of cervical cancer by age group in Poland (estimates for 2020)	101
86	Comparison of age-specific vaginal cancer incidence rates in Poland, within the region, and the rest of world	102
87	Age-standardised incidence rates of penile cancer of Poland (estimates for 2020)	103
88	Annual number of new cases of penile cancer by age group in Poland (estimates for 2020)	104
89	Comparison of age-specific penile cancer incidence rates in Poland, within the region, and the rest of world	105
90	Age-standardised incidence rates of oropharyngeal cancer of Poland (estimates for 2020)	106
91	Annual number of new cases of oropharyngeal cancer among men by age group in Poland (estimates for 2020) .	
92	Annual number of new cases of oropharyngeal cancer among women by age group in Poland (estimates for 2020)	
93	Comparison of age-specific oropharyngeal cancer incidence rates among men by age in Poland, within the region,	100
	and the rest of world	109
94	Comparison of age-specific oropharyngeal cancer incidence rates among women by age in Poland, within the	100
01	region, and the rest of world	110
95	Age-standardised incidence rates of oral cavity cancer of Poland (estimates for 2020)	111
95 96	Annual number of new cases of oral cavity cancer among men by age group in Poland (estimates for 2020)	112
97	Annual number of new cases of oral cavity cancer among men by age group in Poland (estimates for 2020)	112
98	Comparison of age-specific oral cavity cancer incidence rates among men by age in Poland, within the region,	110
30	and the rest of world	114
00	Comparison of age-specific oral cavity cancer incidence rates among women by age in Poland, within the region,	114
99		115
100	and the rest of world	115
	Age-standardised incidence rates of laryngeal cancer of Poland (estimates for 2020)	116
	Annual number of new cases of laryngeal cancer among men by age group in Poland (estimates for 2020)	117
	Annual number of new cases of laryngeal cancer among women by age group in Poland (estimates for 2020)	118
103	Comparison of age-specific laryngeal cancer incidence rates among men by age in Poland, within the region, and	.
	the rest of world	119
104	Comparison of age-specific laryngeal cancer incidence rates among women by age in Poland, within the region,	
	and the rest of world	
105	Age-standardised mortality rates of cervical cancer of Poland (estimates for 2020)	121

	Annual number of deaths of cervical cancer by age group in Poland (estimates for 2020)	122
		123
108	Age-standardised mortality rates of anal cancer of Poland (estimates for 2020)	124
109		125
		126
111	Comparison of age-specific anal cancer mortality rates among men by age in Poland, within the region, and the	
	rest of world	127
112	Comparison of age-specific anal cancer mortality rates among women by age in Poland, within the region, and	
	the rest of world	128
	Age-standardised mortality rates of vulva cancer of Poland (estimates for 2020)	129
	Annual number of deaths of vulva cancer by age group in Poland (estimates for 2020)	130
	$Comparison \ of \ age-specific \ vulva \ cancer \ mortality \ rates \ in \ Poland, \ within \ the \ region, \ and \ the \ rest \ of \ world \ \ . \ .$	131
116	Age-standardised mortality rates of vaginal cancer of Poland (estimates for 2020)	132
		133
118	$Comparison \ of \ age-specific \ vaginal \ cancer \ mortality \ rates \ in \ Poland, \ within \ the \ region, \ and \ the \ rest \ of \ world \ \ . \ .$	134
119	Age-standardised mortality rates of penile cancer of Poland (estimates for 2020)	135
120	Annual number of new deaths of penile cancer by age group in Poland (estimates for 2020)	136
121	Comparison of age-specific penile cancer mortality rates in Poland, within the region, and the rest of world	137
	Age-standardised mortality rates of oropharyngeal cancer of Poland (estimates for 2020)	138
	Annual number of deaths of oropharyngeal cancer among men by age group in Poland (estimates for 2020)	139
	Annual number of deaths of oropharyngeal cancer among women by age group in Poland (estimates for 2020) $$.	140
125	Comparison of age-specific oropharyngeal cancer mortality rates among men by age in Poland, within the region,	
	and the rest of world	141
126	Comparison of age-specific oropharyngeal cancer mortality rates among women by age in Poland, within the	
	region, and the rest of world	142
127	Age-standardised mortality rates of oral cavity cancer of Poland (estimates for 2020)	143
	Annual number of deaths of oral cavity cancer among men by age group in Poland (estimates for 2020)	144
129	Annual number of deaths of oral cavity cancer among women by age group in Poland (estimates for 2020)	145
130	Comparison of age-specific oral cavity cancer mortality rates among men by age in Poland, within the region,	
	and the rest of world	146
131	Comparison of age-specific oral cavity cancer mortality rates among women by age in Poland, within the region,	
	and the rest of world	147
132	Age-standardised mortality rates of laryngeal cancer of Poland (estimates for 2020)	148
133	Annual number of deaths of laryngeal cancer among men by age group in Poland (estimates for 2020)	149
134	Annual number of deaths of laryngeal cancer among women by age group in Poland (estimates for 2020)	150
135	Comparison of age-specific laryngeal cancer mortality rates among men by age in Poland, within the region, and	
	the rest of world	151
136	Comparison of age-specific laryngeal cancer mortality rates among women by age in Poland, within the region,	
	and the rest of world	152

List of Tables

1	Key Statistics
2	Cervical cancer incidence in Poland (estimates for 2020)
3	Cervical cancer incidence in Poland by cancer registry 10
4	Age-standardised incidence rates of cervical cancer in Poland by histological type and cancer registry 12
5	Cervical cancer mortality in Poland (estimates for 2020) 14
6	Premature deaths and disability from cervical cancer in Poland, Europe and the rest of the world (estimates for
	2019)
7	Anal cancer incidence in Poland (estimates for 2020) 18
8	Anal cancer mortality in Poland (estimates for 2020)
9	Vulva cancer incidence in Poland (estimates for 2020)
10	Vulva cancer mortality in Poland (estimates for 2020)
11	Vaginal cancer incidence in Poland (estimates for 2020) 28
12	Vaginal cancer mortality in Poland (estimates for 2020) 30
13	Penile cancer incidence in Poland (estimates for 2020)
14	Penile cancer mortality in Poland (estimates for 2020) 35
15	Oropharyngeal cancer incidence in Poland (estimates for 2020)
16	Oropharyngeal cancer mortality in Poland (estimates for 2020)
17	Oral cavity cancer incidence in Poland (estimates for 2020)
18	Oral cavity cancer mortality in Poland (estimates for 2020)
19	Laryngeal cancer incidence in Poland (estimates for 2020)
20	Laryngeal cancer mortality in Poland (estimates for 2020)
21	Prevalence of HPV16 and HPV18 by cytology in Poland
22	Type-specific HPV prevalence in women with normal cervical cytology, precancerous cervical lesions and invasive
	cervical cancer in Poland
23	Type-specific HPV prevalence among invasive cervical cancer cases in Poland by histology
24 24	Studies on HPV prevalence among HIV+ women with normal cytology in Poland
25	Studies on HPV prevalence among anal cancer cases in Poland (male and female)
26	Studies on HPV prevalence among cases of AIN2/3 in Poland
27	Studies on HPV prevalence among vulvar cancer cases in Poland
28	Studies on HPV prevalence among VIN 2/3 cases in Poland
20 29	Studies on HPV prevalence among value cases in Poland
30	Studies on HPV prevalence among Valin 2/3 cases in Poland
31	Studies on HPV prevalence among penile cancer cases in Poland
32	Studies on HPV prevalence among PeIN 2/3 cases in Poland
33	Studies on HPV prevalence among men in Poland
34	Studies on HPV prevalence among men from special subgroups in Poland
35	Studies on oral HPV prevalence among healthy in Poland
36	Studies on HPV prevalence among cases of oral cavity cancer in Poland
30 37	Studies on HPV prevalence among cases of oropharyngeal cancer in Poland
38	Studies on HPV prevalence among cases of oropharyngeal or laryngeal cancer in Poland
	Factors contributing to cervical carcinogenesis (cofactors) in Poland
39 40	
40	Percentage of 15-year-olds who have had sexual intercourse in Poland 81 Median age at first sex in Poland 81
41	
42	Marriage patterns in Poland
43	Average number of sexual partners in Poland 82 Lifetime number of sexual intervence on a second partners in Poland 85
44	Lifetime prevalence of anal intercourse among women in Poland
45	Main characteristics of cervical cancer screening in Poland
46	National HPV Immunization programme in Poland
47	Prevalence of male circumcision in Poland
48	Prevalence of condom use in Poland
49	Glossary

1 Introduction

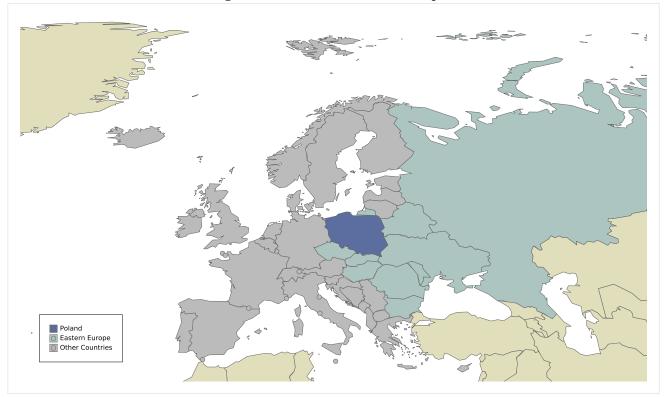


Figure 1: Poland and Eastern Europe

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 Poland 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 Poland. For analytical purposes, Poland is classified in the geographical region of Eastern Europe (Figure 1, lighter blue), which is composed of the following countries: Belarus, Czechia, Hungary, Republic of Moldova, Romania, Russian Federation, Slovakia, and Ukraine. Throughout the report, Poland 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 Poland 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 Poland, 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) 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**

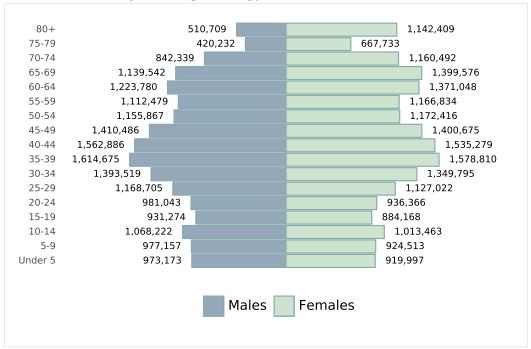


Figure 2: Population pyramid of Poland for 2022

Data accessed on 30 Jul 2022

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].

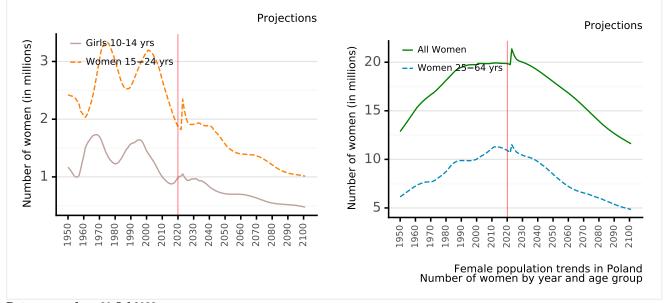


Figure 3: Population trends in four selected age groups in Poland

Data accessed on 30 Jul 2022

Please refer to original source for methods of estimation. Year of estimate: 2022

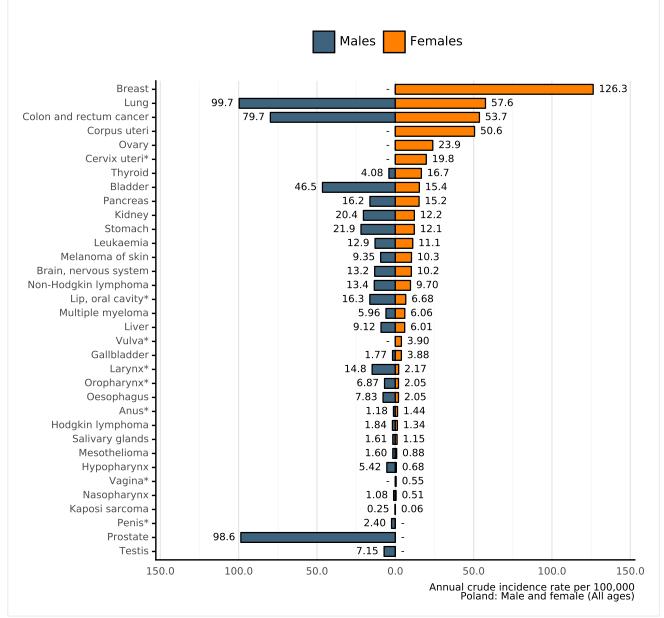
Data Sources: United Nations, Department of Economic and Social Affairs, Population Division (2022). World Population Prospects 2022, Online Edition. [Accessed on July 30, 2022].

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 Poland (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

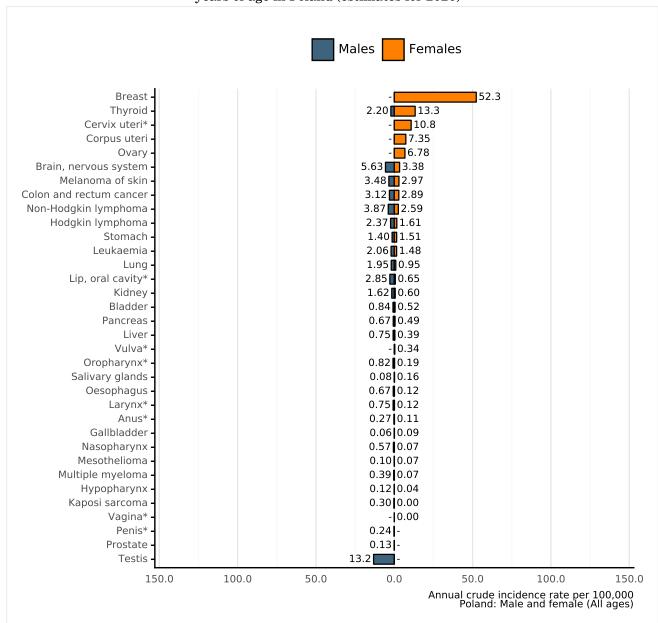


Figure 5: Comparison of HPV related cancers incidence to other cancers among men and women 15-44 years of age in Poland (estimates for 2020)

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

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

Rates per 100,000 women per year.

Data Sources:

3.2 HPV related cancers mortality

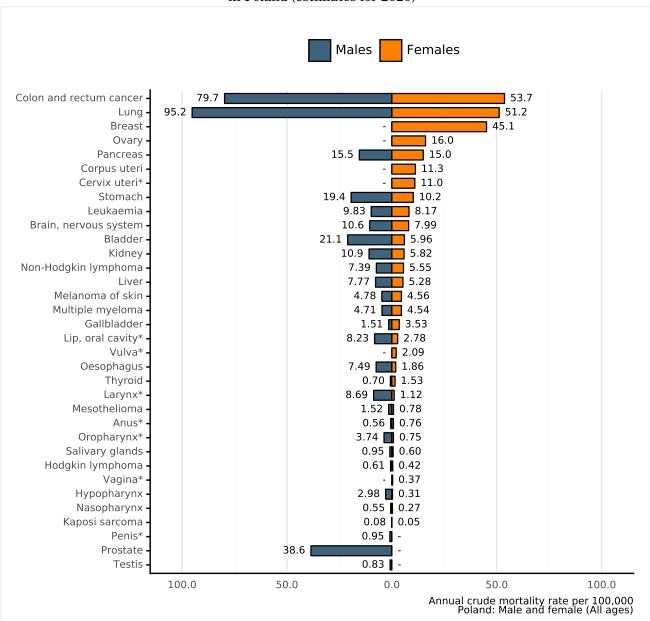
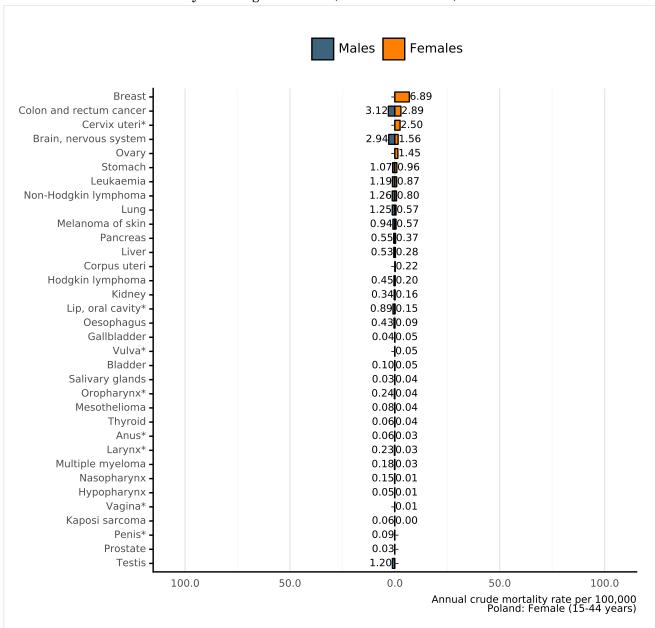


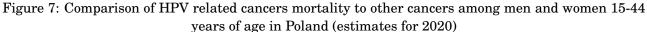
Figure 6: Comparison of HPV related cancers mortality to other cancers in men and women of all ages in Poland (estimates for 2020)

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.





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 Poland 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 Poland

Key Stats.	
About 3,862 new cervical cancer cases are diagnosed annually in Poland (estimations for 2020).	
Cervical cancer ranks* as the 6th leading cause of female cancer in Poland .	
Cervical cancer is the 3rd most common female cancer in women aged 15 to 44 years in Poland .	

* 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

Table 2. Cervical cancer incluence in Forand (estimates for 2020)					
Poland	Eastern Europe	World			
3,862	32,348	604,127			
[3,583-4,163]	[31,584-33,131]	[582,031-627,062]			
19.8	20.8	15.6			
12.3	14.5	13.3			
1.30	1.42	1.39			
	Poland 3,862 [3,583-4,163] 19.8 12.3	PolandEastern Europe3,86232,348[3,583-4,163][31,584-33,131]19.820.812.314.5			

Table 2: Cervical cancer incidence in Poland (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 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 Poland by cancer registry					
Cancer registry	Period	N cases ^a	Crude rate ^b	ASR ^b	
Cieszyn ¹	1973 - 1977	77	21	17.6	
Cieszyn and Nowy Sacz ²	1968-1972	171	19.9	18.6	
Cracow ³	2003-2006	333	20.7	13.9	
Cracow City ⁴	1993-1997	550	27.9	19.6	
Cracow City and District ²	1968 - 1972	1466	22	19.5	
Four Rural Areas ⁵	1965 - 1966	116	20.9	19.4	
Katowice District ¹	1973-1974	939	25	20.9	
Kielce ³	2003-2007	578	17.4	12.1	
Lower Silesia ³	2003-2007	1418	18.9	12.6	
Nowy Sacz Rural Areas ⁶	1983-1986	123	14.3	12.3	
Opole ⁶	1985-1987	304	19.4	16.3	
Warsaw City ⁷	1998-2002	871	19.7	12.5	
Warsaw Rural Areas ⁶	1983-1987	258	16.9	14.6	
Podkarpackie ³	2003-2007	799	14.8	11.1	
Kielce ⁸	2008-2012	467	14.2	9.5	
Lower Silesia ⁸	2008-2012	1273	16.9	10.8	
Lublin ⁸	2008-2012	850	15.2	10.3	
Podkarpackie ⁸	2008-2012	645	11.9	8.1	
Poznan ⁸	2008-2012	1205	13.7	9.4	

Table 3: Carvical cancer incidence in Poland by cancer registry

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. ^a Accumulated number of cases during the period in the population covered by the corresponding registry.

b Rates per 100,000 women per year.

Data Sources:

T Waterhouse, J., Muir, C.S., Shanmugaratnam, K., Powell, J., eds (1982). Cancer Incidence in Five Continents, Vol. IV. IARC Scientific Publications No. 42, Lyon, IARC.

² 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.

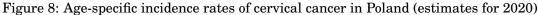
³ Forman D, Bray F, Brewster DH, Gombe Mbalawa C, Kohler B, Piñeros M, Steliarova-Foucher E, Swaminathan R and Ferlay J eds (2013). Cancer Incidence in Five Continents, Vol. X (electronic version) Lyon, IARC. http://ci5.iarc.fr

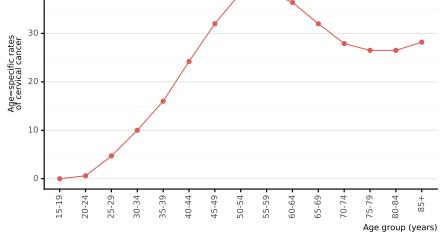
⁴ Parkin, D.M., Whelan, S.L., Ferlay, J., Teppo, L., and Thomas, D.B., eds (2002). Cancer Incidence in Five Continents, Vol. VIII. IARC Scientific Publications No. 155, Lyon, IARC.
 ⁵ Doll, R., Muir, C.S., Waterhouse, J.A.H., eds (1970). Cancer Incidence in Five Continents, Vol. II. Union Internationale Contre le Cancer, Geneva.

⁶ 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. 7 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,

¹ Lyon, IARC.
 ⁸ 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].



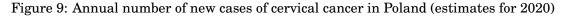


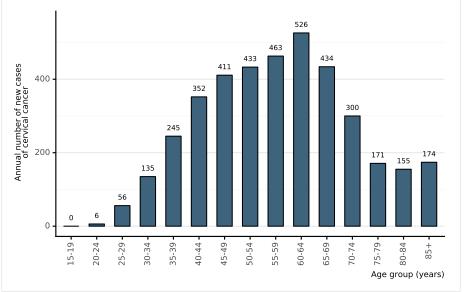


For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods^a Rates per 100,000 women per year.

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].





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

3.3.2 Cervical cancer incidence by histology in Poland

Table 4: Age-standardised incidence rates of cervical cancer in Poland by histological type and cancer registry

	registry				
Cancer registry ¹	Period	Squamo	Adeno	Other	Unspec.
Greater Poland	2008-2012	7.8	0.9	0.1	0.2
Kielce	2008-2012	7.5	1.1	0.1	0.5
Lower Silesia	2008-2012	9.2	0.8	0.1	0.3
Lublin	2008-2012	8	1.2	0.1	0.2
Podkarpackie	2008-2012	6	1.3	0.2	0.2

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: Carcinoma; Unspecified carcinoma;

Adeno: autocationia, outor outor enclosed, galaxies, gal

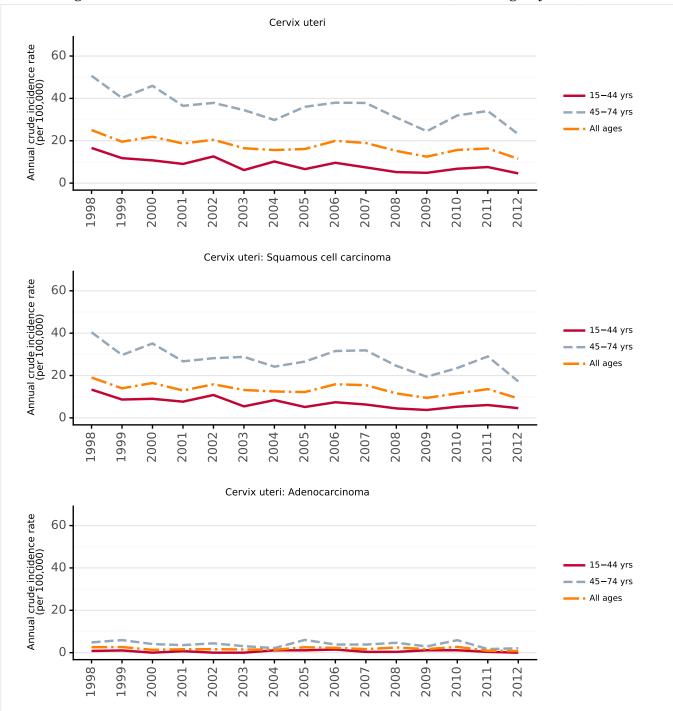


Figure 10: Time trends in cervical cancer incidence in Poland (cancer registry data)

Data accessed on 28 Aug 2018 The following regional cancer registries provided data and contributed to their national estimate: Kielce ^a Estimated annual percentage change based on the trend variable from the net drift for 15 years, from 1988-2002.

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.

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 Poland

Key Stats. About 2,137 cervical cancer deaths occur annually in Poland are diagnosed annually (estimations for 2020). Cervical cancer ranks* as the 7th leading cause of cancer deaths of female cancer deaths in Poland. Cervical cancer is the 3rd leading cause of cancer deaths in women aged 15 to 44 years in Poland.

* 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.

Indicator	Poland	Eastern Europe	World
Annual number of deaths	2,137	15,854	341,831
Uncertainty intervals of mortal- ity cancer cases [95% UI]	[2,012-2,270]	[15, 373 - 16, 350]	[324,231-360,386]
Crude mortality rate ^b	11.0	10.2	8.84
Age-standardized mortality rate ^b	5.86	6.06	7.25
Cumulative risk (%) at 75 years old ^a	0.65	0.65	0.82

Table 5. Commissi com con montality in Dal 1 / f.

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.

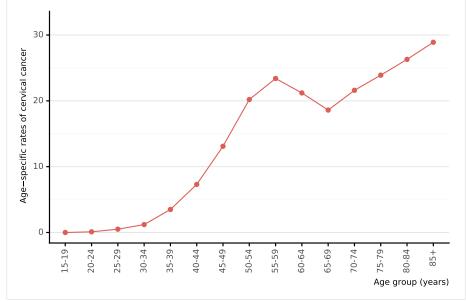


Figure 11: Age-specific mortality rates of cervical cancer in Poland (estimates for 2020)

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: https://gco.iarc.fr/today, accessed [27 January 2021].

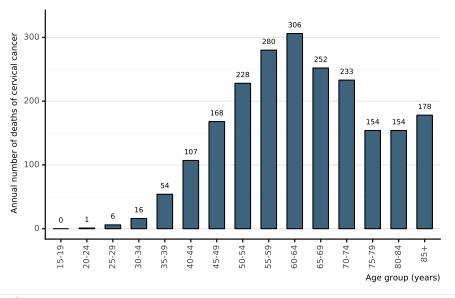


Figure 12: Annual number of deaths of cervical cancer in Poland (estimates for 2020)

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 mortality rates of cervical cancer of Poland (estimates for 2020) please refer to Figure 105
- For annual number of deaths of cervical cancer by age group in Poland (estimates for 2020) please refer to Figure 106
- For comparison of age-specific cervical cancer mortality rates in Poland, within the region, and the rest of world please refer to Figure 107

3.3.4 Cervical cancer incidence and mortality comparison in Poland

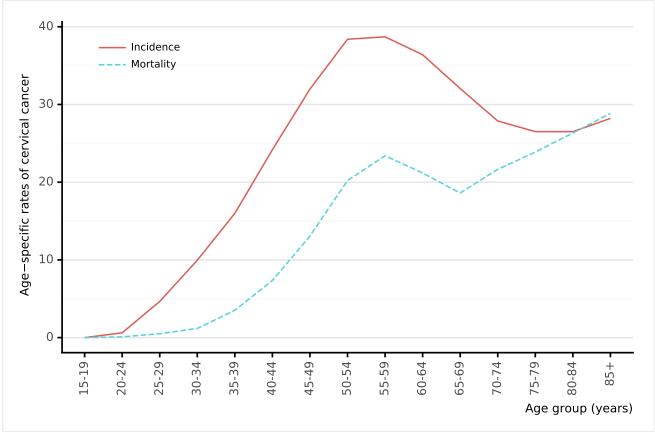


Figure 13: Comparison of age-specific cervical cancer incidence and mortality rates in Poland (estimates for 2020)

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 Sourc

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 Poland, Europe and the rest of the world (estimates for 2019)

Polan	l Europe		World		
Number	Rate	Number	Rate	Number	Rate
55,300	279	824,336	189	8,955,013	232
(43,001-70,366)	(217-355)	(726, 198-913, 992)	(166-209)	(7,547,733-9,978,462)	(196-259)
54,062	273	793,756	182	8,712,962	226
(41,708-69,049)	(210-348)	(703,004-877,841)	(161-201)	(7, 365, 279 - 9, 728, 886)	(191-252)
1,238 (795-1,778)	6 (4-9)	30,580 (21,266-42,064)	7 (5-10)	242,051 (171,644-326,024)	6 (4-8)
	Number 55,300 (43,001-70,366) 54,062 (41,708-69,049) 1,238	$\begin{array}{cccc} 55,300 & 279 \\ (43,001\text{-}70,366) & (217\text{-}355) \\ \hline 54,062 & 273 \\ (41,708\text{-}69,049) & (210\text{-}348) \\ \hline 1,238 & 6 (4\text{-}9) \end{array}$	Number Rate Number 55,300 279 824,336 (43,001-70,366) (217-355) (726,198-913,992) 54,062 273 793,756 (41,708-69,049) (210-348) (703,004-877,841) 1,238 6 (4-9) 30,580	Number Rate Number Rate 55,300 279 824,336 189 (43,001-70,366) (217-355) (726,198-913,992) (166-209) 54,062 273 793,756 182 (41,708-69,049) (210-348) (703,004-877,841) (161-201) 1,238 6 (4-9) 30,580 7 (5-10)	Number Rate Number Rate Number 55,300 279 824,336 189 8,955,013 (43,001-70,366) (217-355) (726,198-913,992) (166-209) (7,547,733-9,978,462) 54,062 273 793,756 182 8,712,962 (41,708-69,049) (210-348) (703,004-877,841) (161-201) (7,365,279-9,728,886) 1,238 6 (4-9) 30,580 7 (5-10) 242,051

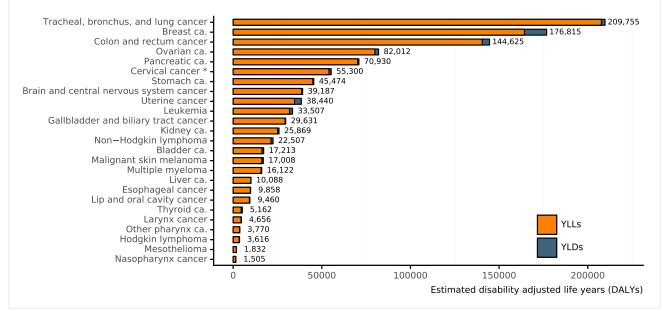
Data accessed on 29 Apr 2021

^a 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) Data Sources:

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



Data accessed on 29 Apr 2021

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

Data Sources: 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

3.4 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 Poland

Table 7: Anal cancer incidence in Poland (estimates for 2020)				
Indicator	Poland	Eastern Europe	World	
MEN				
Annual number of new cancer cases	216	900	21,706	
Uncertainty intervals of new cancer cases [95% UI]	[154-303]	[648-1,250]	[18,432-25,561]	
Crude incidence rate ^b	1.18	0.65	0.55	
Age-standardized incidence rate ^b	0.65	0.41	0.49	
Cumulative risk (%) at 75 years old ^a	0.08	0.05	0.06	
WOMEN				
Annual number of new cancer cases	280	1,558	29,159	
Uncertainty intervals of new cancer cases [95% UI]	[192-408]	[1,228-1,977]	[25,656-33,140]	
Crude incidence rate ^c	1.44	1.00	0.75	
Age-standardized incidence rate ^c	0.59	0.50	0.58	
Cumulative risk (%) at 75 years old^a	0.07	0.06	0.07	

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.

Data Sources:

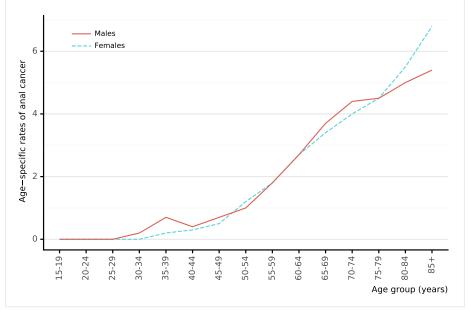
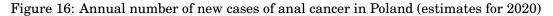


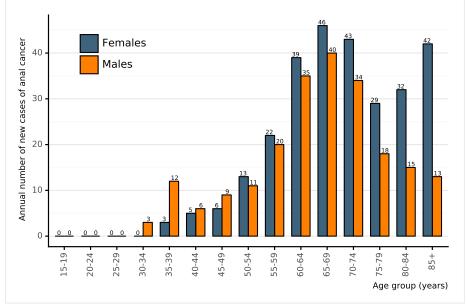
Figure 15: Age-specific incidence rates of anal cancer in Poland (estimates for 2020)

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.

^b 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 $Research \ on \ Cancer. \ Available \ from: \ \texttt{https://gco.iarc.fr/today} \ , \ accessed \ [27 \ January \ 2021].$





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

3.4.1.2 Anal cancer mortality in Poland

Table 8. Anal	concor mortalit	v in P	bland ((estimates for 2020)
Table 0. Allal	cancer mortant	y 111 1	ofallu v	(countaico 101 2020)

Indicator	Poland	Eastern Europe	World
MEN			
Annual number of new cancer cases	103	523	9,416
Uncertainty intervals of new cancer cases [95% UI]	[79-134]	[356-769]	[7,282-12,175]
Crude incidence rate ^b	0.56	0.38	0.24
Age-standardized incidence rate ^b	0.29	0.23	0.21
Cumulative risk (%) at 75 years old ^a	0.03	0.03	0.02
WOMEN			
Annual number of new cancer cases	148	715	9,877
Uncertainty intervals of new cancer cases [95% UI]	[110-199]	[515-993]	[7,795-12,516]
Crude incidence rate ^c	0.76	0.46	0.26
Age-standardized incidence rate ^c	0.26	0.20	0.19
Cumulative risk (%) at 75 years old ^a	0.03	0.02	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.

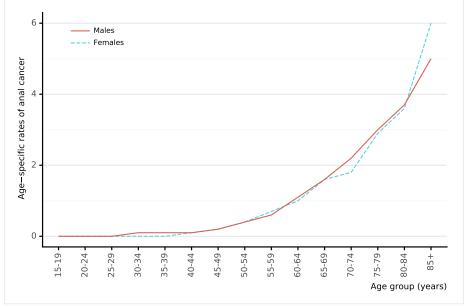
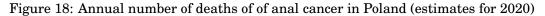


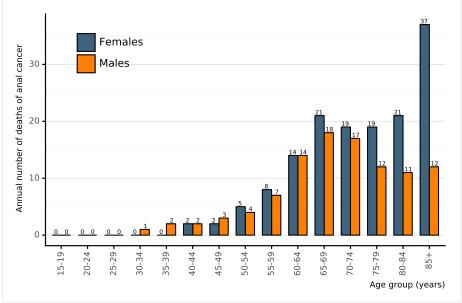
Figure 17: Age-specific mortality rates of anal cancer in Poland (estimates for 2020)

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.

^b 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 $Research \ on \ Cancer. \ Available \ from: \ \texttt{https://gco.iarc.fr/today} \ , \ accessed \ [27 \ January \ 2021].$





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

3.4.1.3 Anal cancer incidence and mortality comparison in Poland

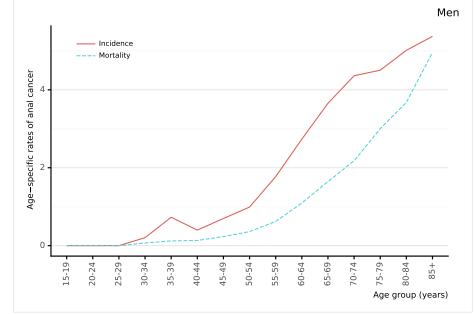
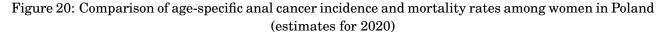


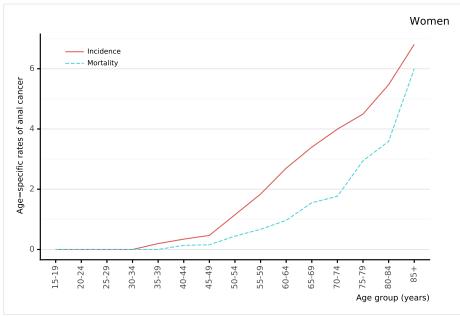
Figure 19: Comparison of age-specific anal cancer incidence and mortality rates among men in Poland (estimates for 2020)

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].





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:

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 Poland

Table 9: Vulva cancer incidence in Poland (estimates for 2020)			
Indicator	Poland	Eastern Europe	World
Annual number of new cancer cases	761	5,012	45,240
Uncertainty intervals [95% UI]	[627-924]	[4,689-5,357]	[40,656-50,342]
Crude incidence rate ^b	3.90	3.23	1.17
Age-standardized incidence rate ^b	1.41	1.30	0.85
Cumulative risk (%) at 75 years old ^a	0.16	0.15	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 causes.

^b Rates per 100,000 women per year. <u>Data Sources</u>:

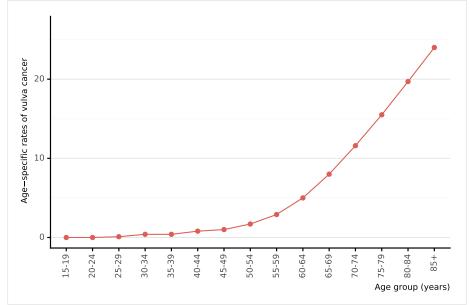


Figure 21: Age-specific incidence rates of vulva cancer in Poland (estimates for 2020)

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: https://gco.iarc.fr/today, accessed [27 January 2021].

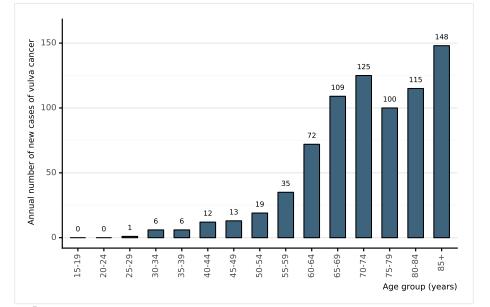


Figure 22: Annual number of new cases of vulva cancer in Poland (estimates for 2020)

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

3.4.2.2 Vulva cancer mortality in Poland

Indicator	Poland	Eastern Europe	World
Annual number of deaths	407	$2,\!561$	$17,\!427$
Uncertainty intervals [95% UI]	[350-474]	[2,144-3,059]	[14,497-20,950]
Crude mortality rate ^b	2.09	1.65	0.45
Age-standardized mortality rate ^b	0.65	0.59	0.30
Cumulative risk (%) at 75 years old ^a	0.07	0.06	0.03

tality in Poland (estimates for 2020) Table 10. Wals

Data accessed on 27 Jan 2021

Data accessed on 2/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.

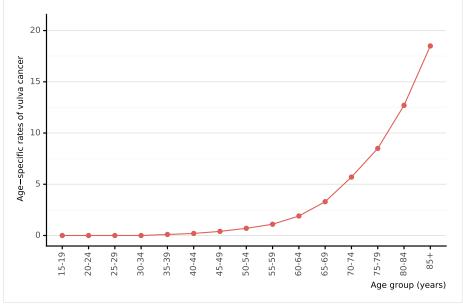
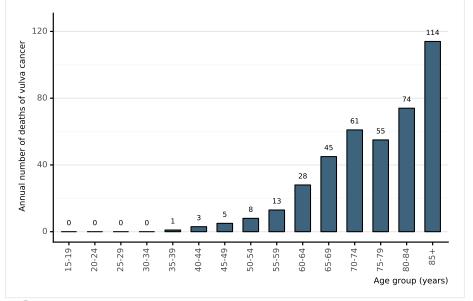
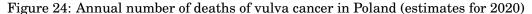


Figure 23: Age-specific mortality rates of vulva cancer in Poland (estimates for 2020)

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: 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 Rates per 100,000 women per year.

Data Sources:

3.4.2.3 Vulva cancer incidence and mortality comparison in Poland

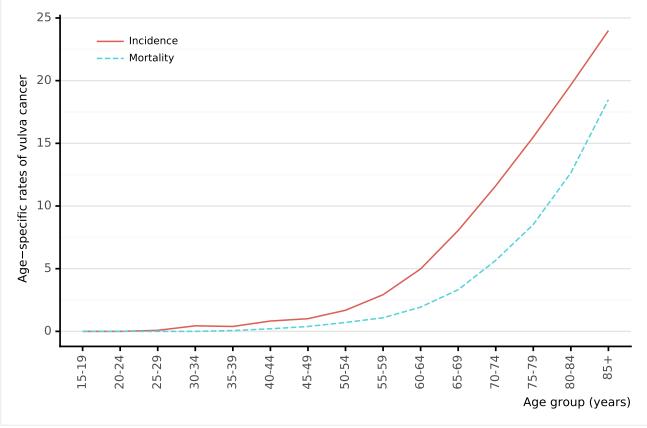


Figure 25: Comparison of age-specific vulva cancer incidence and mortality rates in Poland (estimates for 2020)

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.

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 Poland

Indicator	Poland	Eastern Europe	World
Annual number of new cancer cases	108	1,005	17,908
Uncertainty intervals [95% UI]	[78-149]	[877-1,152]	[14,678-21,848]
Crude incidence rate ^b	0.55	0.65	0.46
Age-standardized incidence rate ^b	0.20	0.31	0.36
Cumulative risk (%) at 75 years old ^a	0.02	0.04	0.04

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 women per year. Data Sources

erlay 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].

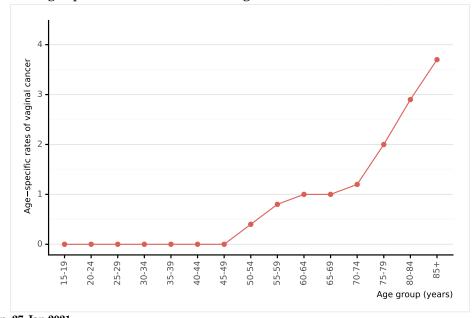
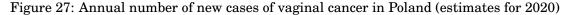
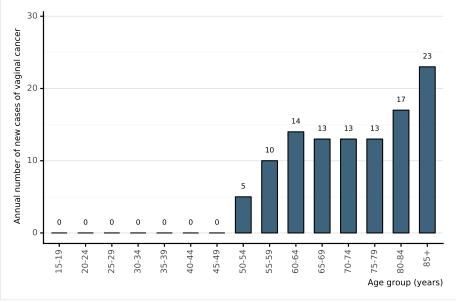


Figure 26: Age-specific incidence rates of vaginal cancer in Poland (estimates for 2020)

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 $Research \ on \ Cancer. \ Available \ from: \ \texttt{https://gco.iarc.fr/today} \ , \ accessed \ [27 \ January \ 2021].$





Data accessed on 27 Jan 2021

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

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 $Research \ on \ Cancer. \ Available \ from: \ \texttt{https://gco.iarc.fr/today} \ , \ accessed \ [27 \ January \ 2021].$

3.4.3.2 Vaginal cancer mortality in Poland

Indicator	Poland	Eastern Europe	World
Annual number of deaths	73	485	7,995
Uncertainty intervals [95% UI]	[56-96]	[322-730]	[5,983-10,684]
Crude mortality rate ^b	0.37	0.31	0.21
Age-standardized mortality rate ^b	0.12	0.13	0.16
Cumulative risk (%) at 75 years old ^a	0.01	0.01	0.02

Table 19	Vaginal cancer	mortality in Pol	and (actimator)	for 2020)
1able 12.	vaginai cancei	moreancy mint of	and (estimates)	

Data accessed on 27 Jan 2021

Data accessed on 2/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.

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].

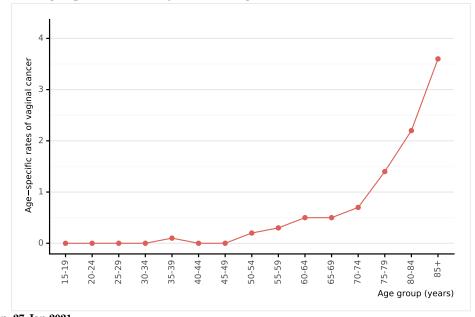
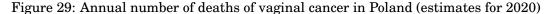
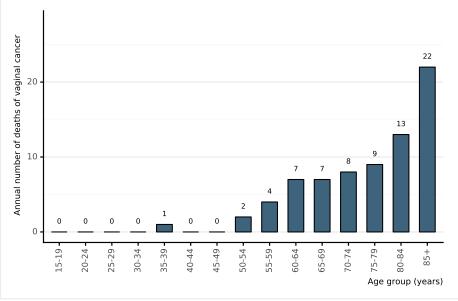


Figure 28: Age-specific mortality rates of vaginal cancer in Poland (estimates for 2020)

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 $Research \ on \ Cancer. \ Available \ from: \ \texttt{https://gco.iarc.fr/today} \ , \ accessed \ [27 \ January \ 2021].$





Data accessed on 27 Jan 2021

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

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 $Research \ on \ Cancer. \ Available \ from: \ \texttt{https://gco.iarc.fr/today} \ , \ accessed \ [27 \ January \ 2021].$

3.4.3.3 Vaginal cancer incidence and mortality comparison in Poland

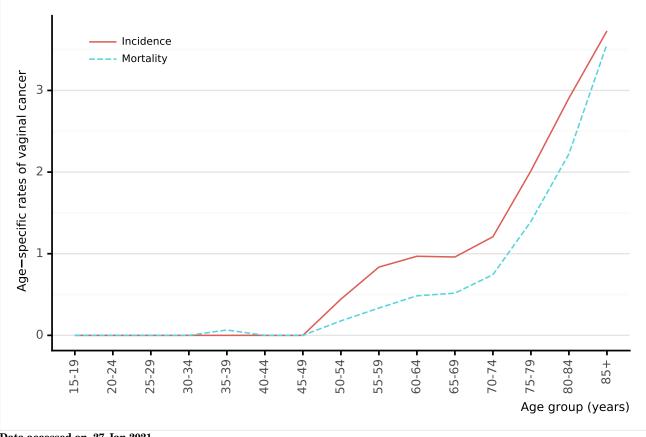


Figure 30: Comparison of age-specific vaginal cancer incidence and mortality rates in Poland (estimates for 2020)

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.

Pata 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 Poland

Table 13: Penile cancer incidence in Poland (estimates for 2020)				
Indicator	Poland	Eastern Europe	World	
Annual number of new cancer cases	440	2,038	36,068	
Uncertainty intervals [95% UI]	[314-617]	[1,828-2,272]	[30,963-42,015]	
Crude incidence rate ^b	2.40	1.48	0.92	
Age-standardized incidence rate ^b	1.25	0.91	0.80	
Cumulative risk (%) at 75 years old ^a	0.15	0.11	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 causes.

^b Rates per 100,000 men per year. Data Sources

Perlay 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].

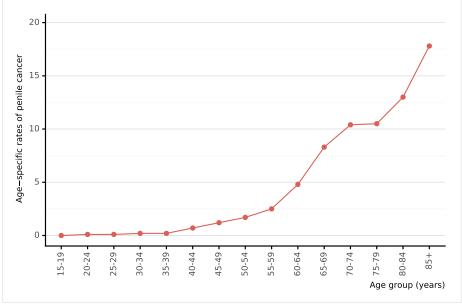
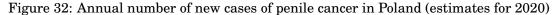


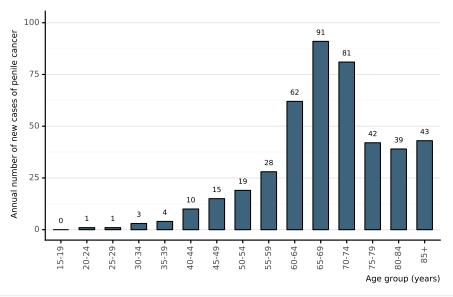
Figure 31: Age-specific incidence rates of penile cancer in Poland (estimates for 2020)

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].





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 $Research \ on \ Cancer. \ Available \ from: \ \texttt{https://gco.iarc.fr/today} \ , \ accessed \ [27 \ January \ 2021].$

3.4.4.2 Penile cancer mortality in Poland

Indicator	Poland	Eastern Europe	World
Annual number of deaths	174	762	13,211
Uncertainty intervals [95% UI]	[134-227]	[551-1,055]	[10,687-16,332]
Crude mortality rate ^b	0.95	0.55	0.34
Age-standardized mortality rate ^b	0.49	0.33	0.29
Cumulative risk (%) at 75 years old ^a	0.05	0.04	0.03

Table 14: Penile cancer mortality in Poland (estimates for 2020)

Data accessed on 27 Jan 2021

Data accessed on 2/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.

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].

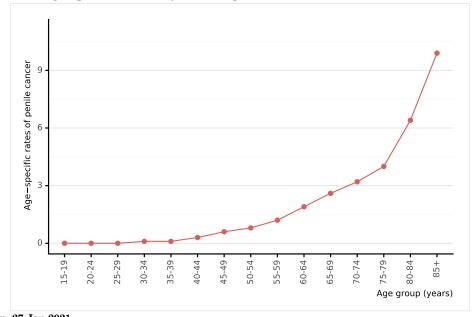


Figure 33: Age-specific mortality rates of penile cancer in Poland (estimates for 2020)

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 $Research \ on \ Cancer. \ Available \ from: \ \texttt{https://gco.iarc.fr/today} \ , \ accessed \ [27 \ January \ 2021].$

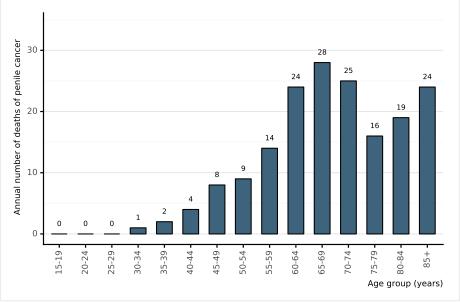


Figure 34: Annual number of deaths of penile cancer in Poland (estimates for 2020)

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 $Research \ on \ Cancer. \ Available \ from: \ \texttt{https://gco.iarc.fr/today} \ , \ accessed \ [27 \ January \ 2021].$

3.4.4.3 Penile cancer incidence and mortality comparison in Poland

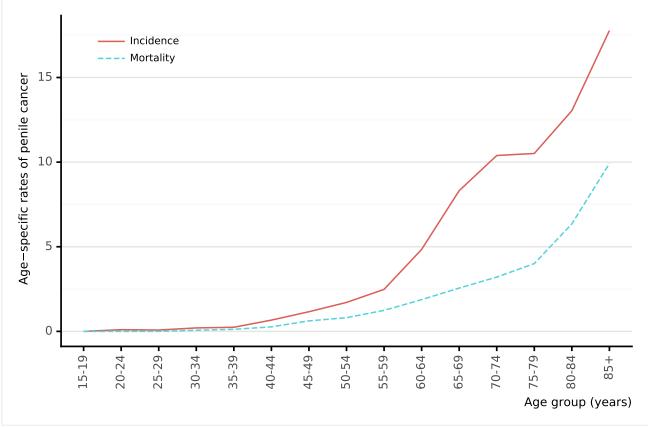


Figure 35: Comparison of age-specific penile cancer incidence and mortality rates in Poland (estimates for 2020)

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].

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 Poland

Table 15: Oropharyngeal cancer incidence in Poland (estimates for 2020)				
Indicator	Poland	Eastern Europe	World	
MEN				
Annual number of new cancer cases	1,260	9,200	79,045	
Uncertainty intervals of new cancer cases [95% UI]	[1,076-1,476]	[8,757-9,665]	[72,769-85,862]	
Crude incidence rate sa ^b	6.87	6.67	2.01	
Age-standardized incidence rate sa ^b	4.13	4.36	1.79	
Cumulative risk (%) at 75 years old ^a	0.52	0.54	0.22	
WOMEN				
Annual number of new cancer cases	399	1,891	19,367	
Uncertainty intervals of new cancer cases [95% UI]	[273-584]	[1,696-2,108]	[16,279-23,041]	
Crude incidence rate sa ^c	2.05	1.22	0.50	
Age-standardized incidence rate sa ^c	1.09	0.68	0.40	
Cumulative risk (%) at 75 years old^a	0.14	0.08	0.05	

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.

Data Sources

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].

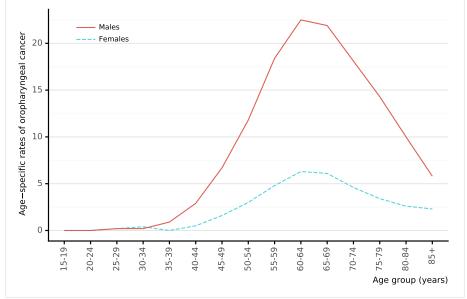


Figure 36: Age-specific incidence rates of oropharyngeal cancer in Poland (estimates for 2020)

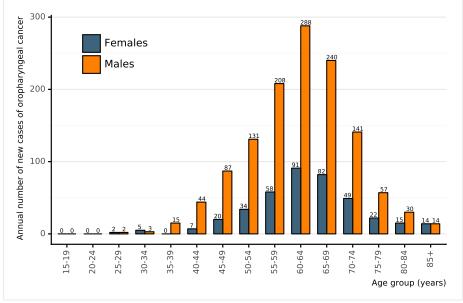
Data accessed on 27 Jan 2021

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: 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 37: Annual number of new cases of oropharyngeal cancer in Poland (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].

3.5.1.2 Oropharyngeal cancer mortality in Poland

Indicator	Poland	Eastern Europe	World
MEN			
Annual number of deaths	686	4,925	39,590
Uncertainty intervals of mortality cancer cases [95% UI]	[606-777]	[4,331-5,601]	[35,255-44,458]
Crude mortality rate sa ^b	3.74	3.57	1.01
Age-standardized mortality rate sa ^b	2.16	2.31	0.89
Cumulative risk (%) at 75 years old ^a	0.27	0.29	0.11
WOMEN			
Annual number of deaths	147	710	8,553
Uncertainty intervals of mortality cancer cases [95% UI]	[109-198]	[506-996]	[6,684-10,945]
Crude mortality rate sa ^c	0.75	0.46	0.22
Age-standardized mortality rate sa ^c	0.36	0.23	0.17
Cumulative risk (%) at 75 years old ^a	0.05	0.03	0.02

Table 16: Oropharyngeal cancer mortality in Poland (estimates for 2020
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Data accessed on 27 Jan 2021

Data accessed OI 21 0 all 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:
Data Source

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 $Research \ on \ Cancer. \ Available \ from: \ \texttt{https://gco.iarc.fr/today} \ , \ accessed \ [27 \ January \ 2021].$

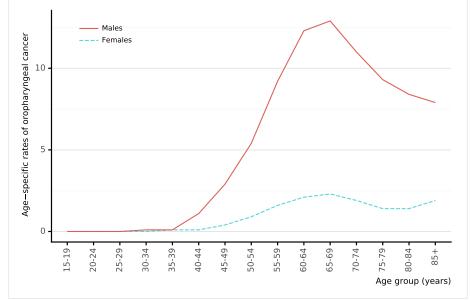


Figure 38: Age-specific mortality rates of oropharyngeal cancer in Poland (estimates for 2020)

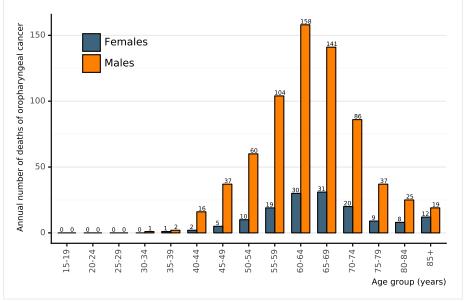
Data accessed on 27 Jan 2021

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: 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 39: Annual number of deaths of oropharyngeal cancer in Poland (estimates for 2020)

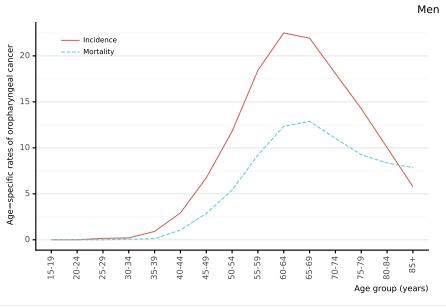


Data accessed on 27 Jan 2021 For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods Data Sources:

Farlay 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].

3.5.1.3 Oropharyngeal cancer incidence and mortality comparison in Poland

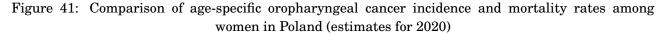
Figure 40: Comparison of age-specific oropharyngeal cancer incidence and mortality rates among men in Poland (estimates for 2020)

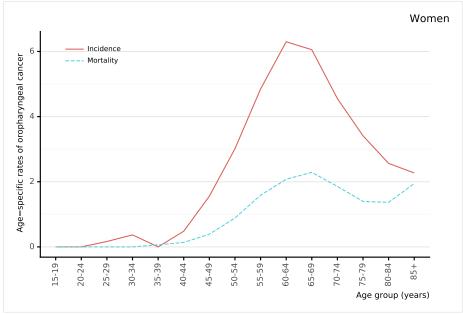


Data accessed on 27 Jan 2021

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

Farlay 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 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].

3.5.2 Oral cavity cancer

3.5.2.1 Oral cavity cancer incidence in Poland

Indicator	Poland	Eastern Europe	World
MEN			
Annual number of new cancer cases	2,985	19,884	264,211
Uncertainty intervals of new cancer cases [95% UI]	[2,606-3,419]	[19,288-20,498]	[251,153- 277,948]
Crude incidence rate sa ^b	16.3	14.4	6.72
$\begin{array}{llllllllllllllllllllllllllllllllllll$	9.57	9.22	5.96
Cumulative risk (%) at 75 years old^a	1.14	1.12	0.68
WOMEN			
Annual number of new cancer cases	1,304	6,199	113,502
Uncertainty intervals of new cancer cases [95% UI]	[1,063-1,599]	[5,864-6,553]	[105,599- 121,997]
Crude incidence rate sa ^c	6.68	3.99	2.94
Age-standardized incidence rate sa ^c	2.83	1.87	2.28
Cumulative risk (%) at 75 years old^a	0.33	0.21	0.26

incidence in Poland (actimates for 2020) Table 17 Oral ovitv a

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.

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].

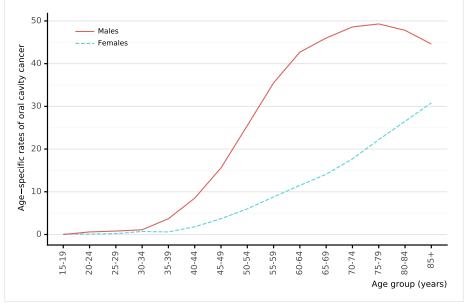


Figure 42: Age-specific incidence rates of oral cavity cancer in Poland (estimates for 2020)

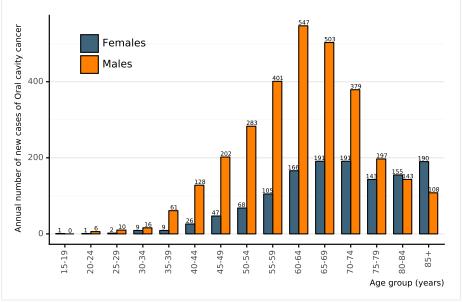
Data accessed on 27 Jan 2021

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: 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 43: Annual number of new cases of oral cavity cancer in Poland (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:

Parlay JErvik 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].

3.5.2.2 Oral cavity cancer incidence and mortality comparison in Poland

Table 18: Oral cavity cancer mortality in Poland (estimates for 2020)				
Indicator	Poland	Eastern Europe	World	
MEN				
Annual number of deaths	1,509	9,761	125,022	
Uncertainty intervals of mortality cancer cases [95% UI]	[1,357-1,678]	[8,901-10,704]	[116,573- 134,084]	
Crude mortality rate sa ^b	8.23	7.08	3.18	
Age-standardized mortality rate sa ^b	4.80	4.54	2.82	
Cumulative risk (%) at 75 years old ^a	0.57	0.56	0.32	
WOMEN				
Annual number of deaths	542	2,729	52,735	
Uncertainty intervals of mortality cancer cases [95% UI]	[462-636]	[2,286-3,258]	[47,690-58,313]	
Crude mortality rate sa ^c	2.78	1.76	1.36	
Age-standardized mortality rate sa ^c	1.13	0.82	1.04	
Cumulative risk (%) at 75 years old^a	0.13	0.09	0.12	

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, 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].

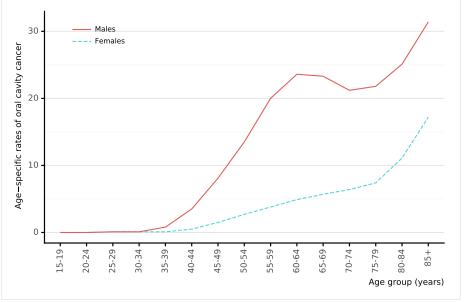


Figure 44: Age-specific mortality rates of oral cavity cancer in Poland (estimates for 2020)

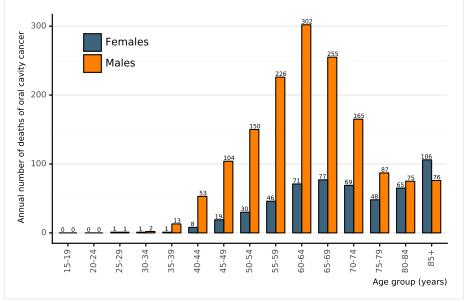
Data accessed on 27 Jan 2021

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: 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 45: Annual number of deaths of oral cavity cancer in Poland (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:

Parlay JErvik 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].

3.5.2.3 Oral cavity cancer incidence and mortality comparison in Poland

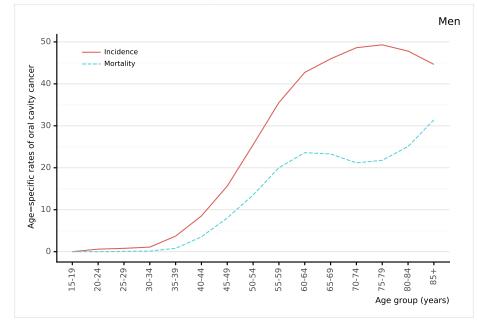
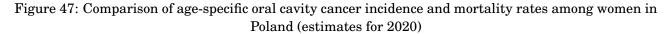


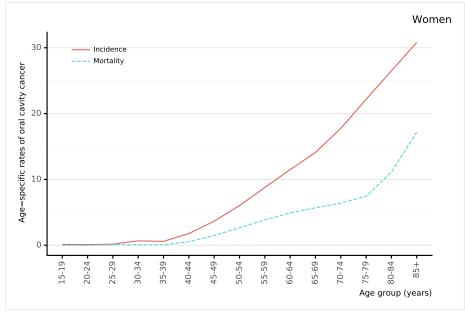
Figure 46: Comparison of age-specific oral cavity cancer incidence and mortality rates among men in Poland (estimates for 2020)

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].





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].

3.5.3 Laryngeal cancer

3.5.3.1 Laryngeal cancer incidence in Poland

Indicator	Poland	Eastern Europe	World
MEN			
Annual number of new cancer cases	2,712	16,580	160,265
Uncertainty intervals of new cancer cases [95% UI]	[2,442-3,012]	[16,000-17,181]	[150,633- 170,513]
Crude incidence rate sa ^b	14.8	12.0	4.08
Age-standardized incidence rate sa ^b	8.39	7.58	3.59
Cumulative risk (%) at 75 years old^a	1.05	0.97	0.45
WOMEN			
Annual number of new cancer cases	423	1,550	24,350
Uncertainty intervals of new cancer cases [95% UI]	[326-549]	[1,367-1,757]	[20,845-28,444]
Crude incidence rate sa ^c	2.17	1.00	0.63
Age-standardized incidence rate sa ^c	1.11	0.53	0.49
Cumulative risk (%) at 75 years old ^a	0.14	0.07	0.06

ar incidence in Poland (estimates for 2020) Table 10. Lawrongeal cane

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.

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].

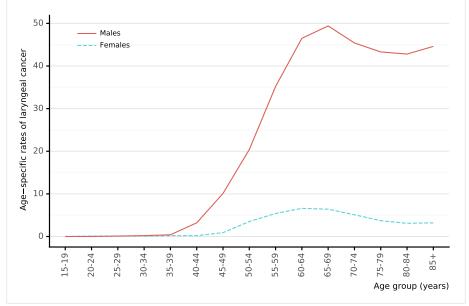


Figure 48: Age-specific incidence rates of laryngeal cancer in Poland (estimates for 2020)

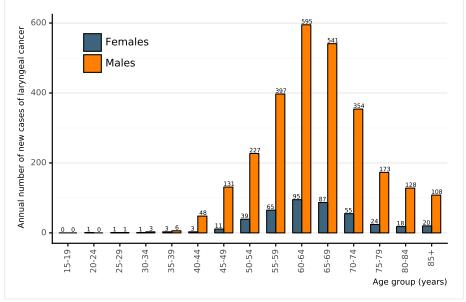
Data accessed on 27 Jan 2021

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: 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 49: Annual number of new cases of laryngeal cancer in Poland (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:

Farlay 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].

3.5.3.2 Laryngeal cancer incidence and mortality comparison in Poland

Table 20: Laryngeal cancer mortality in Poland (estimates for 2020)				
Poland	Eastern Europe	World		
1,593	9,518	85,351		
[1,467-1,730]	[9,130-9,923]	[78,895-92,335]		
8.69	6.91	2.17		
4.65	4.26	1.89		
0.59	0.55	0.23		
218	741	14,489		
[178-268]	[633-868]	[11,902-17,639]		
1.12	0.48	0.37		
0.52	0.24	0.28		
0.07	0.03	0.03		
	Poland	Poland Eastern Europe 1,593 9,518 [1,467-1,730] [9,130-9,923] 8.69 6.91 4.65 4.26 0.59 0.55 218 741 [178-268] [633-868] 1.12 0.48 0.52 0.24		

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, 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].

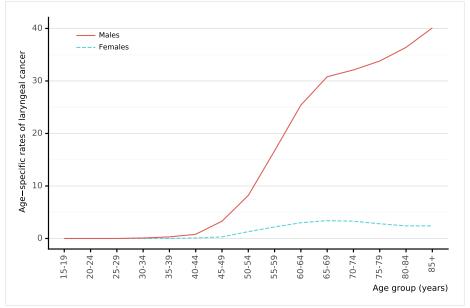


Figure 50: Age-specific mortality rates of laryngeal cancer in Poland (estimates for 2020)

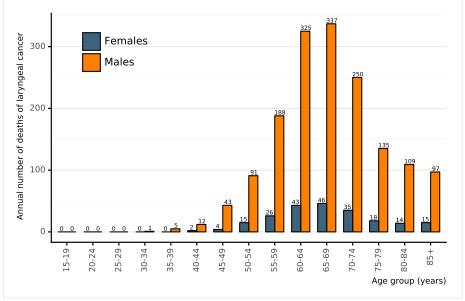
Data accessed on 27 Jan 2021

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: 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].

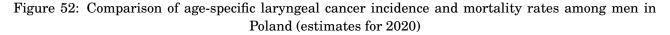
Figure 51: Annual number of deaths of of laryngeal cancer in Poland (estimates for 2020)

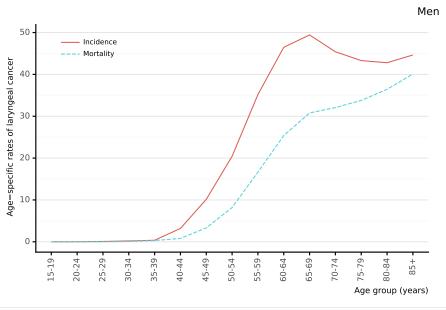


Data accessed on 27 Jan 2021 For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods Data Sources:

Farlay 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].

3.5.3.3 Laryngeal cancer incidence and mortality comparison in Poland

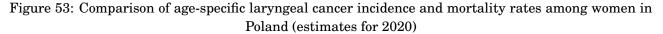


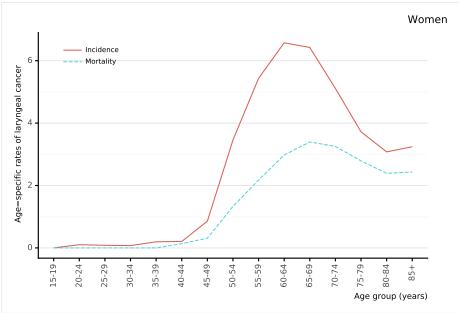


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For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods a Rates per 100,000 men per year. Data Sources

Farlay 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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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].

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

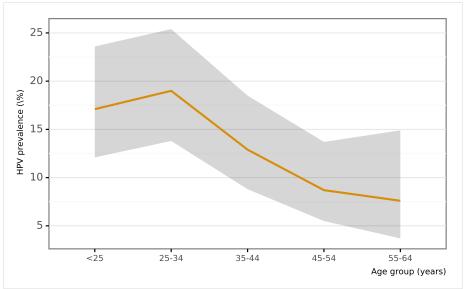
The statistics shown in this section focus on HPV infection in the cervix uteri. HPV cervical infection results in cervical morphological lesions ranging from normalcy (cytologically normal women) to different stages of precancerous lesions (CIN-1, CIN-2, CIN-3/CIS) and invasive cervical cancer. HPV 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 Poland



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

Data Sources: Bardin A, Eur J Cancer 2008; 44: 557

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 Poland, 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: Bardin A, Eur J Cancer 2008; 44: 557

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

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

	No. tested	HPV 16/18 Prevalence % (95% CI)
Normal cytology ^{1,2}	799	3.4 (2.3-4.9)
Low-grade lesions ³	-	
High-grade lesions ⁴	-	
Cervical cancer ^{5,6}	1010	88.1 (86.0-90.0)

Table 21: Prevalence of HPV16 and HPV18 by cytology in Poland

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)

Number of women tested ^b 95% Confidence Interval

Data Sources

Bardin A, Eur J Cancer 2008; 44: 557

² 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

³ 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

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. ⁵ Contributing studies: Baay MF, Eur J Gynaecol Oncol 2009; 30: 162 | Bardin A, Eur J Cancer 2008; 44: 557 | Biesaga B, Folia Histochem Cytobiol 2012; 50: 239 | Bosch FX, J Natl

Cancer Inst 1995; 87: 796 | Dybikowska A, Oncol Rep 2002; 9: 871 | Kwasniewska A, Eur J Gynaecol Oncol 2009; 30: 65 | Pirog EC, Am J Pathol 2000; 157: 1055 ⁶ 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 56: HPV 16 prevalence among women with normal cervical cytology in Poland, by study

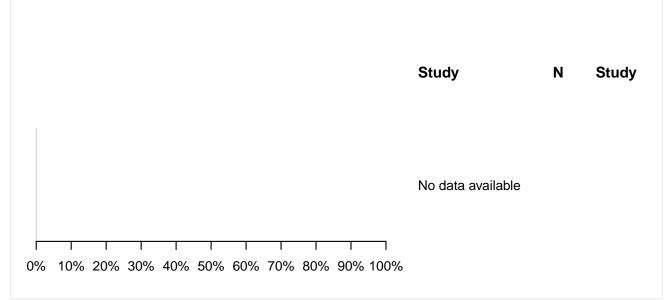
	Baudry 2008 799 26(958-4.4)
0%	10%

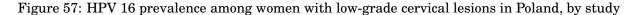
Data updated on 30 Jun 2015 (data as of 30 Jun 2014)

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

Data Sources: Bardin A, Eur J Cancer 2008; 44: 557

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





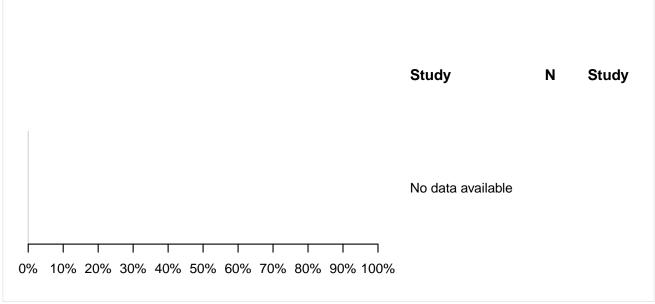
Data updated on 27 Jan 2017 (data as of 30 Jun 2015)

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

Data Sources:

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 Poland, by study



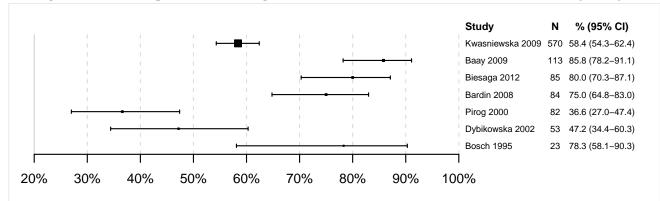
Data updated on 27 Jan 2017 (data as of 30 Jun 2015)

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

Data Sources:

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.

Figure 59: HPV 16 prevalence among women with invasive cervical cancer in Poland, 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)

^a Number of women tested Data Sources:

Data Sources: Baay MF, Eur J Gynaecol Oncol 2009; 30: 162 | Bardin A, Eur J Cancer 2008; 44: 557 | Biesaga B, Folia Histochem Cytobiol 2012; 50: 239 | Bosch FX, J Natl Cancer Inst 1995; 87: 796 | Dyblikowska A, Oncol Rep 2002; 9: 871 | Kwasniewska A, Eur J Gynaecol Oncol 2009; 30: 65 | Pirog EC, Am J Pathol 2000; 157: 1055 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) 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.

16 56 45 52 31 51 33 58 18 59	2.8 1.8 1.5 1.3 1.3 0.9 0.9 0.6 0.6 0.6 0.4							Normal cytology (5,6)
1st 2nd 3th 4th 5th 6th 7th 8th 9th 10th	* * * * * *							Low-grade lesions (4)
AH 1st 2nd 3th 4th 5th 6th 7th 8th 9th 10th	* * * * * * * * * * * * *							High-grade lesions (3)
16 18 45 52 31 33 56 58 39 35	2.3 2.3 1.7 1.3 1.1 1 1	7		25.3			62.	Cervical cancer (1,2)
	0%	10%	20%	^{30%} Preva	40% alence	50%	60%	70%

Figure 60: Comparison of the ten most frequent HPV oncogenic types in Poland among women with and without cervical lesions

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

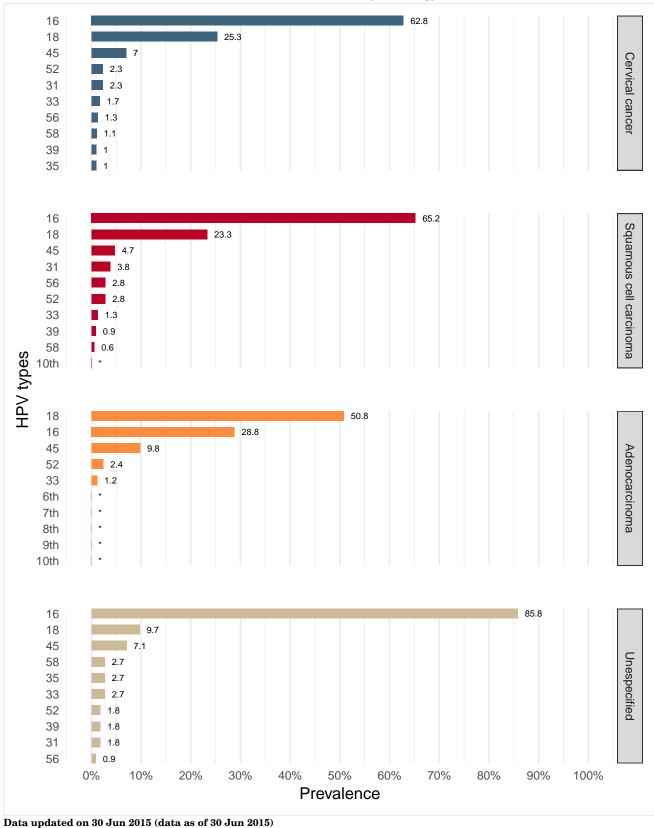
Data Sources:

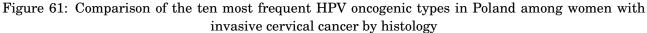
Data Sources:
 ¹ Contributing studies: Baay MF, Eur J Gynaecol Oncol 2009; 30: 162 | Bardin A, Eur J Cancer 2008; 44: 557 | Biesaga B, Folia Histochem Cytobiol 2012; 50: 239 | Bosch FX, J Natl Cancer Inst 1995; 87: 796 | Dybikowska A, Oncol Rep 2002; 9: 871 | Kwasniewska A, Eur J Gynaecol Oncol 2009; 30: 65 | Pirog EC, Am J Pathol 2000; 157: 1055
 ² 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

³ 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.

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) Clifford GM, Cancer Epidemiol Biomarkers Prev 2005;14:1157 ⁵ Bardin A, Eur J Cancer 2008; 44: 557 ⁶ 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





 st No data available. No more types than shown were tested or were positive

¹Contributing studies: Baay MF, Eur J Gynaecol Oncol 2009; 30: 162 | Bardin A, Eur J Cancer 2008; 44: 557 | Biesaga B, Folia Histochem Cytobiol 2012; 50: 239 | Bosch FX, J Natl

² 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;88:63 5) Clifford

GM, Br J Cancer 2003;89:101.
³ 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

Data Sources:

GM, Br J Cancer 2003;89:101.

lesions and invasive cervical cancer in Poland									
	Normal cytology ^{1,2}		Low-grade lesions ³		High-grade lesions ⁴		Cervical cancer ^{5,6}		
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)	
ONCOG	ENIC HPV	TYPES							
High-r	risk HPV ty	pes							
16	799	2.8 (1.8-4.1)	-	-		-	1010	62.8 (59.7-65.7)	
18	799	0.6 (0.3-1.5)	-	-	-	-	1010	25.3 (22.8-28.1)	
31	799	1.3 (0.7-2.3)	-	-	-	-	355	2.3 (1.1-4.4)	
33	799	0.9 (0.4-1.8)	-	-	-	-	355	1.7 (0.8-3.6)	
35	799	0.3 (0.1-0.9)	-	-	-	-	302	1.0 (0.3-2.9)	
39	799	0.1 (0.0-0.7)	-	-	-	-	302	1.0 (0.3-2.9)	
45	799	1.5 (0.9-2.6)	-	-	-	-	302	7.0 (4.6-10.4)	
51	799	0.9 (0.4-1.8)	-	-	-	-	302	0.0 (0.0-1.3)	
52	799	1.3 (0.7-2.3)	-	-	-	-	302	2.3 (1.1-4.7)	
56	799	1.8 (1.0-2.9)	-	-	-	-	302	1.3 (0.5-3.4)	
58	799	0.6 (0.3-1.5)	-	-	-	-	355	1.1 (0.4-2.9)	
59	799	0.4 (0.1-1.1)	-	-	-	-	302	0.3 (0.1-1.9)	
	ble/possible	e carcinogen							
26	799	0.3 (0.1-0.9)	-	-	-		220	0.0 (0.0-1.7)	
30	799	0.0 (0.0-0.5)	-	-		-	84	0.0 (0.0-4.4)	
34	799	0.0 (0.0-0.5)	-	-			166	0.0 (0.0-2.3)	
53	799	0.5 (0.2-1.3)	-	-		-	302	0.0 (0.0-1.3)	
66	799	1.0 (0.5-2.0)	-	-		_	302	0.3 (0.1-1.9)	
67	799	0.3 (0.1-0.9)	- -	-		-	84	0.0 (0.0-4.4)	
68	799	0.4 (0.1-1.1)		-		-	302	0.3 (0.1-1.9)	
69	799	0.0 (0.0-0.5)	_	-				-	
70	799	0.6 (0.3-1.5)		-			189	0.0 (0.0-2.0)	
73	799	0.8 (0.3-1.6)		-	-	-	105	0.0 (0.0-3.5)	
82	799	0.1 (0.0-0.7)		-	-	-	107	0.0 (0.0-3.5)	
85	799	0.0 (0.0-0.5)		-	-	-	-		
97	-	-		-		-		-	
	- SK HPV TY		-	-	-	-	-	-	
6	799	0.4 (0.1-1.1)	· .				355	0.0 (0.0-1.1)	
11	799	0.3 (0.1-0.9)					355	0.0 (0.0-1.1)	
32	799		-		-	-	-	-	
40	799	0.0 (0.0-0.5)	-	-	-	-		- 0.0 (0.0-3.3)	
			-	-	-	-	113		
42	799	2.3 (1.4-3.5)	-	-	-	-	113	0.0 (0.0-3.3)	
43	799	0.3 (0.1-0.9)	-	-	-	-	113	0.0 (0.0-3.3)	
44	799	0.4 (0.1-1.1)	-	-	-	-	113	0.0 (0.0-3.3)	
54	799	0.1 (0.0-0.7)	-	-	-	-	-	-	
55	-	-	-	-	-	-	-	-	
57	799	0.0 (0.0-0.5)	-	-	-	-	-	-	
61	799	0.1 (0.0-0.7)	-	-	-	-	-	-	
62	-	-	-	-	-	-	-	-	
64	-	-	-	-	-	-	-	-	
71	799	0.0 (0.0-0.5)	-	-	-	-	-	-	
72	799	0.0 (0.0-0.5)	-	-	-	-	-	-	
74	-	-	-	-	-	-	-	-	
81	799	0.1 (0.0-0.7)	-	-	-	-	-	-	
83	799	0.9 (0.4-1.8)	-	-	-	-	-	-	
84	799	0.1 (0.0-0.7)	-	-	-	-	-	-	
86	799	0.1 (0.0-0.7)	-	-	-	-	-	-	
87	-	-	-	-	-	-	-	-	
89	799	0.1 (0.0-0.7)	-	-	-	-	-	-	
90	799	0.8 (0.3-1.6)	-	-	-	-	-	-	
91	-	-	-	-	-	-	-	-	

Table 22: Type-specific HPV prevalence in women with normal cervical cytology, precancerous cervical lesions and invasive cervical cancer in Poland

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)

Data Sources:

¹ Bardin A, Eur J Cancer 2008; 44: 557

² 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

³ 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

 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.
 ⁵ Contributing studies: Baay MF, Eur J Gynaecol Oncol 2009; 30: 162 | Bardin A, Eur J Cancer 2008; 44: 557 | Biesaga B, Folia Histochem Cytobiol 2012; 50: 239 | Bosch FX, J Natl

Cancer Inst 1995; 87: 796 | Dybikowska A, Oncol Rep 2002; 9: 871 | Kwasniewska A, Eur J Gynaecol Oncol 2009; 30: 65 | Pirog EC, Am J Pathol 2000; 157: 1055 ⁶ 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 Poland by histolog Any Histology Squamous cell carcinoma Adenocarcinoma Unespecified								
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)
ONCOG	ENIC HPV	TYPES						
High-r	risk HPV ty	pes						
16	1010	62.8 (59.7-65.7)	765	65.2 (61.8-68.5)	132	28.8 (21.8-37.0)	113	85.8 (78.2-91.1)
18	1010	25.3 (22.8-28.1)	765	23.3 (20.4-26.4)	132	50.8 (42.3-59.1)	113	9.7 (5.5-16.6)
31	355	2.3 (1.1-4.4)	160	3.8 (1.7-7.9)	82	0.0 (0.0-4.5)	113	1.8 (0.5-6.2)
33	355	1.7 (0.8-3.6)	160	1.3 (0.3-4.4)	82	1.2 (0.2-6.6)	113	2.7 (0.9-7.5)
35	302	1.0 (0.3-2.9)	107	0.0 (0.0-3.5)	82	0.0 (0.0-4.5)	113	2.7 (0.9-7.5)
39	302	1.0 (0.3-2.9)	107	0.9 (0.2-5.1)	82	0.0 (0.0-4.5)	113	1.8 (0.5-6.2)
45	302	7.0 (4.6-10.4)	107	4.7 (2.0-10.5)	82	9.8 (5.0-18.1)	113	7.1 (3.6-13.4)
51	302	0.0 (0.0-1.3)	107	0.0 (0.0-3.5)	82	0.0 (0.0-4.5)	113	0.0 (0.0-3.3)
52	302	2.3 (1.1-4.7)	107	2.8 (1.0-7.9)	82	2.4 (0.7-8.5)	113	1.8 (0.5-6.2)
56	302	1.3 (0.5-3.4)	107	2.8 (1.0-7.9)	82	0.0 (0.0-4.5)	113	0.9 (0.2-4.8)
58	355	1.1 (0.4-2.9)	160	0.6 (0.1-3.5)	82	0.0 (0.0-4.5)	113	2.7 (0.9-7.5)
59	302	0.3 (0.1-1.9)	107	0.0 (0.0-3.5)	82	0.0 (0.0-4.5)	113	0.9 (0.2-4.8)
Proba	ble/possibl	e carcinogen						
26	220	0.0 (0.0-1.7)	-	-	-	-	-	-
30	84	0.0 (0.0-4.4)	84	0.0 (0.0-4.4)	-	-	-	-
34	166	0.0 (0.0-2.3)	84	0.0 (0.0-4.4)	82	0.0 (0.0-4.5)	-	-
53	302	0.0 (0.0-1.3)	-	-	-	-	-	-
66	302	0.3 (0.1-1.9)	107	0.0 (0.0-3.5)	82	0.0 (0.0-4.5)	113	0.9 (0.2-4.8)
67	84	0.0 (0.0-4.4)	84	0.0 (0.0-4.4)	-	-	-	-
68	302	0.3 (0.1-1.9)	107	0.9 (0.2-5.1)	82	0.0 (0.0-4.5)	113	0.0 (0.0-3.3)
69	-	-	-	-	-	-	-	-
70	189	0.0 (0.0-2.0)	-	-	-	-	-	-
73	107	0.0 (0.0-3.5)	-	-	-	-	-	-
82	107	0.0 (0.0-3.5)	107	0.0 (0.0-3.5)	-	-	-	-
85	-	-	-	-	-	-	-	-
97	-	-	-	-	-	-	-	-
LOW RIS	SK HPV TY	PES						
6	355	0.0 (0.0-1.1)	-	-	-	-	-	-
11	355	0.0 (0.0-1.1)	-	-	-	-	-	-
32	-	-	-	-	-	-	-	-
40	113	0.0 (0.0-3.3)	-	-	-	-	-	-
42	113	0.0 (0.0-3.3)	-	-	-	-	113	0.0 (0.0-3.3)
43	113	0.0 (0.0-3.3)	-	-	-	-	-	-
44	113	0.0 (0.0-3.3)	-	-	-	-	113	0.0 (0.0-3.3)
54	-	-	-	-	-	-	-	-
55	-	-	-	-	-	-	-	-
57	-	-	-	-	-	-	-	-
61	-	-	-	-	-	-	-	-
62	-	-	-	-	-	-	-	-
64	-	-	-	-	-	-	-	-
71	-	-	-	-	-	-	-	-
72	-	-	-	-	-	-	-	-
74	-	-	-	-	-	-	-	-
81	-	-	-	-	-	-	-	-
83	-	-	-	-	-	-	-	-
84	-	-	-	-	-	-	-	-
86	-	-	-	-	-	-	-	-
87	-	-	-	-	-	-	-	-
89	-	-	-	-	-	-	-	-
90	-	-	-	-	-	-	-	-
91	-	-	-	-	-	-	-	-

Table 23: Type-specific HPV prevalence among invasive cervical cancer cases in Poland by histology

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) a Number of women tested b 95% Confidence Interval

Data Sources:

Data Sources: Contributing studies: Baay MF, Eur J Gynaecol Oncol 2009; 30: 162 | Bardin A, Eur J Cancer 2008; 44: 557 | Biesaga B, Folia Histochem Cytobiol 2012; 50: 239 | Bosch FX, J Natl Cancer Inst 1995; 87: 796 | Dybikowska A, Oncol Rep 2002; 9: 871 | Kwasniewska A, Eur J Gynaecol Oncol 2009; 30: 65 | Pirog EC, Am J Pathol 2000; 157: 1055 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) 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.

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 Poland

			HPV	Prevalence	
Study	HPV detection method and targeted HPV types	No. Tested ^a	%	(95% CI) ^b	Prevalence of 5 most frequent HPVs, HPV type (%)
-	-	-	-	-	

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

DBH: Dot Blot Hybridization; EIA: Enzyme ImmunoAssay; HC2: Hybrid Capture 2; PCR: Polymerase Chain Reaction; TS: Type Specific ^a Number of women tested ^b 95% Confidence Interval

Data Sources:

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).

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.

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 Poland are presented.

Table 25: Studies on HPV prevalence among anal cancer cases in Poland (male and female)

Study ^b	HPV detection method and targeted HPV types	No. Tested	%	(95% CI) ^a	Prevalence of 5 most frequent HPVs, HPV type (%)
Alemany 2015	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)

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; 95% Confidence Interval

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

Data Sources:

Alemany L. Int J Cancer 2015: 136: 98

Hermany J, into Source 2010, 100. 50 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 amon	g cases of AIN2/3 in Poland	

HPV Prevalence						
Study ^b	HPV detection method and targeted HPV types	No. Tested	%	(95% CI) ^a	Prevalence of 5 most frequent HPVs, HPV type (%)	
Alemany 2015	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)	

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

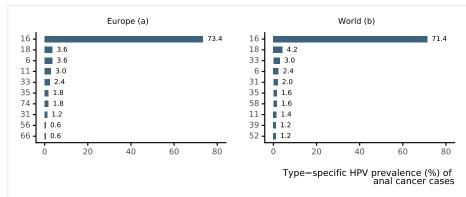
^a 95% Confidence Interval
 ^b Includes cases from Bosnia-Herzegovina, Czech Republic, France, Germany, Poland, Portugal, Slovenia, Spain and United Kingdom

Data Sources

Alemany L. Int J Cancer 2015: 136: 98

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



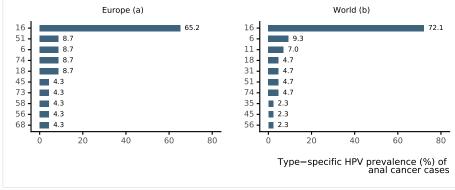
Data updated on 9 Feb 2017 (data as of 30 Jun 2014)

^a 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 and United States); Africa (Mali, Nigeria and Senegal); Asia (Bangladesh, India and South Korea) 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.

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

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)

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

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 Poland are presented.

Table 27: Studies on HPV prevalence among vulvar cancer cases in Poland

			HPV Prevalence				
Study	HPV detection method and targeted HPV types	No. Tested	%	(95% CI) ^a	Prevalence of 5 most frequent HPVs, HPV type (%)		
Bujko 2012	PCR, LBA (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)	44	34.1	(21.9-48.9)	HPV 16 (20.5), HPV 11 (11.4), HPV 44 (4.5), HPV 52 (4.5), HPV 58 (4.5)		
de Sanjosé 2013 ^b	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)		
Liss 1998	PCR-MY09/11, PCR L1-Consensus primer, PCR-E6, PCR-E7, RFLP (HPV 6, 11, 16, 18, 31, 33, 35, 45, 52, 58)	18	16.7	(5.8-39.2)	HPV 16 (16.7)		

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; 95% Confidence Interval

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

Data Sources: Buiko M, Acta Obstet Gynecol Scand 2012; 91: 391 | de Saniosé S, Eur J Cancer 2013; 49: 3450 | Liss J, Ginekol Pol 1998; 69: 330

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 28: Studies on HPV prevalence among VIN 2/3 cases in Poland

Study ^b	HPV detection method and targeted HPV types	No. Tested	%	(95% CI) ^a	Prevalence of 5 most frequent HPVs, HPV type (%)
de Sanjosé 2013	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)

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

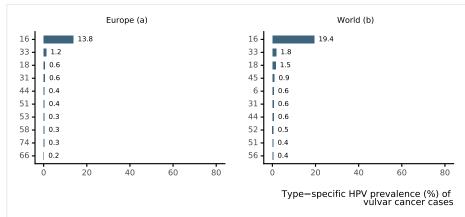
VIN 2/3: Vulvar intraepithelial neoplasia of grade 2/3

^a 95% Confidence Interval
^b Includes cases from Austria, Belarus, Bosnia-Herzegovina, Czech Republic, France, Germany, Greece, Italy, Poland, Portugal, Spain and United Kingdom

Data Sources de Sanjosé 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 2009:124:1626

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

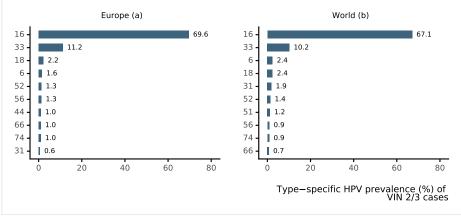


Data updated on 30 Jun 2015 (data as of 30 Jun 2015)

^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, 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, 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.

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 4.2.3

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 Poland are presented.

Table 29: Studies on HPV prevalence among vaginal cancer cases in Poland

		HPV	HPV Prevalence			
Study ^b	HPV detection method and targeted HPV types	No. Tested	%	(95% CI) ^a	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; 95% Confidence Interval

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

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 Poland

HPV Prevalence						
Study	HPV detection method and targeted HPV types	No. Tested	%	(95% CI) ^a	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

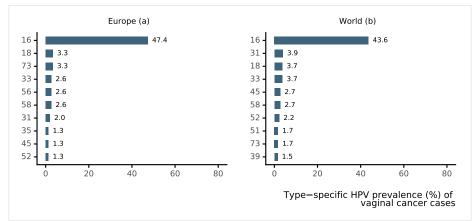
a 95% Confidence Interval

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 2009;124:1626

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



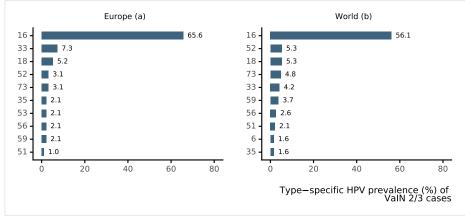
Data updated on 30 Jun 2015 (data as of 30 Jun 2015)

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

^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)

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

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

b Includes cases from Flaveria, Belarus, Status, Statu

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.

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 Poland are presented.

Table 31: Studies on HPV prevalence among penile cancer cases in Poland

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

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:

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 2009;62:870

Table 32: Studies on HPV prevalence among PeIN 2/3 cases in Poland

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

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

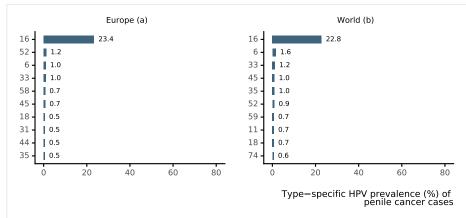
PeIN 2/3: Penile intraepithelial neoplasia of grade 2/3

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

<u>Data Sources:</u> 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



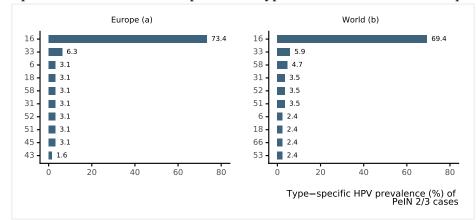
Data updated on 9 Feb 2017 (data as of 30 Jun 2015)

^a 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

Data Sources 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)

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

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. Data Sources

Herrero R, J Natl Cancer Inst 2003; 95: 1772 | Ribeiro KB, Int J Epidemiol 2011; 40: 489 | Snietura M, Pol J Pathol 2010; 61: 133 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

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 Poland 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.

		HPV Prevaler						
Study	Anatomic sites samples	HPV detection method	Population	Age (years)	No. Tested	%	(95% CI) ^a	
-	-	-	-	-	-	-	-	

Table 33: Studies on HPV prevalence among men in Poland

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 diseases ^a 95% Confidence Interval

Data Sources:

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 Poland

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

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:

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 Poland is presented.

4.4.1 Burden of oral HPV infection in healthy population

				· F-			, J J		
Study	Specimen collection method / anatomic site	HPV detec- tion method ^a	Population	% males	Age (years) ^b	No. tested ^c	HPV prevalence % (95% CI)	High-Risk HPV prevalence % (95% CI)	5 most frequent HPVs, HPV type (n) ^d
-	-	-	-	-	-	-	-	-	-

Table 35: Studies on oral HPV prevalence among healthy in Poland

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

20,510 0 0	5: Studies on HPV prevalence an	out		Prevalence	
Study	HPV detection method and targeted HPV types	No. Tested	%	(95% CI) ^a	Prevalence of 5 most frequent HPVs, HPV type (%)
MEN					
Herrero 2003	GP5+/GP6+ (L1) Hybridization with EIA oligonucleotide probes (2. 6. 11. 16. 18. 31. 33. 35. 39. 40. 42. 43. 44. 45. 51. 52. 56. 58. 59. 66. 68)	53	0	-	-
WOMEN					
Herrero 2003	 GP5+/GP6+ (L1) Hybridization with EIA oligonucleotide probes (2. 6. 11. 16. 18. 31. 33. 35. 39. 40. 42. 43. 44. 45. 51. 52. 56. 58. 59. 66. 68) 	30	0	-	-
BOTH OR UNSPECIF	IED				
Herrero 2003	 GP5+/GP6+ (L1) Hybridization with EIA oligonucleotide probes (2. 6. 11. 16. 18. 31. 33. 35. 39. 40. 42. 43. 44. 45. 51. 52. 56. 58. 59. 66. 68) 	83	0	-	-
Ribeiro 2011 ^b	PGMY09/11 (L1) Amplification with TS primers (16)	132	0	-	-
Snietura 2010	Real-time High Risk HPV test (Abbott Molecular) using L1 consensus primers Amplification with TS primers (16. 18. 31. 33. 35. 39. 45. 51. 52. 56. 58. 59. 66 and 68 - the technique only differentiates 16-18-other)	45	4.4	(1.2-14.8)	HPV 16 (4.4)

Table 36: Studies on HPV provalance among eases of oral cavity concer in Poland

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
^b Includes cases from Argentina, Brazil, Cuba, Russia, Slovakia, Czech Republic, Romania and Poland

Data Sources Herrero R, J Natl Cancer Inst 2003; 95: 1772 | Ribeiro KB, Int J Epidemiol 2011; 40: 489 | Snietura M, Pol J Pathol 2010; 61: 133

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 Poland

			HPV	Prevalence	
Study	HPV detection method and targeted HPV types	No. Tested	%	(95% CI) ^a	Prevalence of 5 most frequent HPVs, HPV type (%)
MEN					
No data available	-	-	-	-	-
WOMEN					
No data available	-	-	-	-	-
BOTH OR UNSPECIFI	ED				
Ribeiro 2011	PGMY09/11 (L1) Amplification with TS primers (16)	136	0.7	(0.1-4.0)	HPV 16 (0.7)
Snietura 2010	Real-time High Risk HPV test (Abbott Molecular) using L1 consensus primers Amplification with TS primers (16. 18. 31. 33. 35. 39. 45. 51. 52. 56. 58. 59. 66 and 68 - the technique only differentiates 16-18-other)	14	50.0	(26.8-73.2)	HPV 16 (50.0)
Szkaradkiewicz 2002	MY09/MY11 (L1) Amplification with TS primers (16. 18)	28	10.7	(3.7-27.2)	-

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: Ribeiro KB, Int J Epidemiol 2011; 40: 489 | Snietura M, Pol J Pathol 2010; 61: 133 | Szkaradkiewicz A, Clin Exp Med 2002; 2: 137

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

			HPV	Prevalence	
Study	targeted HPV types		%	(95% CI) ^a	Prevalence of 5 most frequent HPVs, HPV type (%)
MEN					
Morshed 2008	SPF10 (L1) LiPA 25	78	34.6	(25.0-45.7)	-
WOMEN					
Morshed 2008	SPF10 (L1) LiPA 25	15	40.0	(19.8-64.3)	-
BOTH OR UNSPECIFI	ED				
Morshed 2008	SPF10 (L1) LiPA 25	93	35.5	(26.5-45.6)	HPV 16 (30.1) HPV 18 (6.5) HPV 33 (5.4)
Ribeiro 2011 ^b	PGMY09/11 (L1) Amplification with TS primers (16)	239	0.8	(0.2-3.0)	HPV 16 (0.8)
Snietura 2011	Real-time High Risk HPV test (Abbott Molecular) using L1 consensus primers RT-PCR (16. 18. 31. 33. 35. 39. 45. 51. 52. 56. 58. 59. 66. 68)	65	0.0	-	-

Table 38: Studies on HPV prevalence among cases of hypopharyngeal or laryngeal cancer in Poland

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 ^b Includes cases from Argentina, Brazil, Cuba, Russia, Slovakia, Czech Republic, Romania and Poland

Data Sources:

Marshed K, Eur Arch Otorhinolaryngol 2008; 265 Suppl 1: S89 | Ribeiro KB, Int J Epidemiol 2011; 40: 489 | Snietura M, Eur Arch Otorhinolaryngol 2011; 268: 721 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 Poland are presented.

INDICATOR		MALE	FEMALE	TOTAL
Smoking				
Smoking of any tobacco adjusted	Current ^a	33.6 [27.2-42.6]	21.7 [17-26.3]	27.4 [21.9-34.1]
prevalence (%) [95% UI]	Daily ^b	28.7 [22.7-35.2]	17.7 [14.2-21.6]	23 [18.3-28.1]
Cigarette smoking adjusted	Current ^c	33.6 [27.2-42.6]	21.7 [17-26.3]	27.4 [21.9-34.1]
prevalence (%) [95% UI]	Daily ^d	28.7 [22.7-35.2]	17.7 [14.2-21.6]	23 [18.3-28.1]
Parity			10	
Total fertility rate per woman	15.10	-	1.3	-
	15-19 yrs	-	11.0	-
	20-24 yrs	-	50.9	-
Age-specific fertility rate	25-29 yrs	-	100.2	-
(per 1000 women)	30-34 yrs	-	88.8	-
	35-39 yrs	-	38.4	-
	40-44 yrs	-	7.6	-
	45-49 yrs	-	0.3	-
Hormonal contraception Oral contraceptive use (%) among w married or in union		-	17	-
Injectable contraception use (%) a who are married or in union	among women	-	-	-
Implant contraceptive use (%) amon are married or in union	ng women who	-	-	-
HIV				
Estimated percent of adults aged living with HIV [95% UI]	15-49 who are	- [—]	- [—]	- [—]
Estimated percent of young adults a are living with HIV [95% UI]	iged 15-24 who	- [—]	- [—]	- [—]
HIV prevalence (%) among sex worl		-	-	-
HIV prevalence (%) among men who men		7.1999998	-	7.1999998
Estimated number of people living v UI]	with HIV [95%	-	-	- [—]
Estimated number of adults (15+ y HIV [95% UI]	rs) living with	- [—]	- [—]	- [—]
Estimated number of AIDS-related	d deaths [95%	-	-	- [—]

Table 39: Factors contributing to cervical carcinogenesis (cofactors) in Poland

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

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. c "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.

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/ WHO global report on terms in prevalence of tobacco use 2000-2020, and central organization, rest internation, rest inte

frate. [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/ufd2017.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]. UNAIDS database [internet]. Available at: http://aidsinfo.unaids.org/ [Accessed on November 21, 2019]

- 80 -

6 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 Poland are presented.

Table 40: Percentage of 15-year-olds who have had sexual intercourse in Poland

Indicator	Male	Female
Percentage of 15-year-old subjects who report sexual intercourse	16.0	18.0
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: Growing up 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 Poland								
MALE FEMALE TOTAL								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
Crochard 2009 ¹	2006-2007	1982 - 1988	489	17.0	409	18.0	-	-
Olszewski 2010 ^{2,a,b}	2008	1981-1990	-	-	993	18.7	-	-

Data accessed on 16 Mar 2017

Please refer to original source for methods of estimation ^a Data pertain to high school and secondary schools students.

^b Data pertain to high school and secondary schools studen
 ^b Mean age at first sex.

Data Sources:

¹ Crochard A, Luyts D, di Nicola S, Gonçalves MAG. Self-reported sexual debut and behavior in young adults aged 18-24 years in seven European countries: implications for HPV vaccination programs. Gynecol. Oncol. 2009 Dic;115(3 Suppl):S7-S14.

² Olszewski J, Olszewska H, Abacjew-Chmylko A, Chmylko L, Gaworska-Krzeminska A, Wydra D. Sexual behavior and contraception among young Polish women. Acta Obstet Gynecol Scand. 2010 Nov;89(11):1447-52.

Indicator		Male	Female
Average age at first marriage ¹		28.7	26.6
Age-specific % of ever married ²	15-19 years	0.15	1.18
	20-24 years	8.23	21.75
	25-29 years	41.32	61.53
	30-34 years	69.37	81
	35-39 years	80.16	87.73
	40-44 years	84.18	91.14
	45-49 years	85.71	92.76
	50-54 years	87.69	93.67
	55-59 years	90.75	94.13
	60-64 years	93.68	94.53
	65-69 years	95.23	95.37
	70-74 years	95.92	96.07
	+75	97.28	95.8

Table 42: Marriage patterns in Poland

Data accessed on 20 Feb 2020 Please refer to original source for methods of estimation.

^a 2011 Census ^b UNSD

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 ² 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.

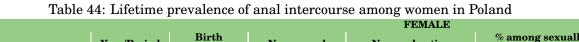
Table 43: Average number of sexual partners in Poland

$\mathbf{Study}^{1,a}$	Period of estimate	Year/Period	Birth cohort	Male Mean(N)	Female Mean(N)	Total Mean(N)
Crochard 2009	Lifetime	2006-2007	(1982-1989)	3.0(481)	2.0(404)	-(-)

Data accessed on 8 Aug 2013 Please refer to original source for methods of estimation ^a Median number of sexual partners.

Data Sources:

1 Crochard A, Luyts D, di Nicola S, Gonçalves MAG. Self-reported sexual debut and behavior in young adults aged 18-24 years in seven European countries: implications for HPV vaccination programs. Gynecol. Oncol. 2009 Dic;115(3 Suppl):S7-S14.



Study	Year/Period	Birth cohort	N surveyed	N sexual active	% among sexually active
-	-	-	-	-	-
Data accessed on 8 Aug 20	013				

Please refer to original source for methods of estimation

HPV preventive strategies 7

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 Poland.

Cervical cancer screening practices 7.1

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 Foland								
gion	Existence of official national	Starting year of current	Active invitation	Screening ages (yes primary screening tes and screening interv				

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
Poland	Yes	2016	No	25-59 (cytology, 3 years)

Table 45. Main characteristics of conviced concernation in Deland

Data accessed on 31 Aug 2022

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.

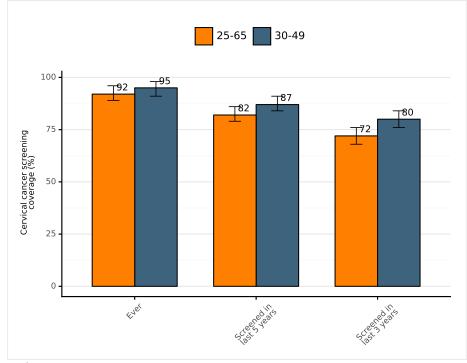


Figure 70: Estimated coverage* of cervical cancer screening in Poland

Data accessed on 31 Aug 2022 * Estimated coverage and 95% confidence interval in 2019 <u>Data Sources:</u> 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 Poland

Female	Male
Not Available/Not Introduced	Not Available/Not Introduced
-	-
-	-
-	-
-	-
	Not Available/Not Introduced - -

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 71: HPV vaccination coverage in females by year in Poland

No data available

Data accessed on 24 Oct 2022

Data accessed of 24 Oct 2022 <u>Junan 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.</u>



Figure 72: HPV vaccination coverage in males by year in Poland

Data accessed on 24 Oct 2022

Data accessed on D4 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.

Protective factors for cervical cancer 8

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

Table 47: Prevalence of male circumcision in Poland					
Reference	Prevalence % (95% CI)	Methods			
WHO 2007	<20	Data from Demographic and Health Surveys (DHS) and other publications to categorize the country-wide prevalence of male circumcision as <20%, 20-80%, or >80%.			

Data accessed on 31 Aug 2015

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

Data Sources: 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.

Indicator	Age range	Year of estimate	Prevalence % ^a
Condom use	15-49	2014	27.1

Data accessed on 18 Nov 2019

Please refer to original source for methods of estimation. ^a 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: 2014 HIS

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].

9 Annex

9.1 Incidence

9.1.1 Cervical cancer incidence in Poland across Eastern Europe

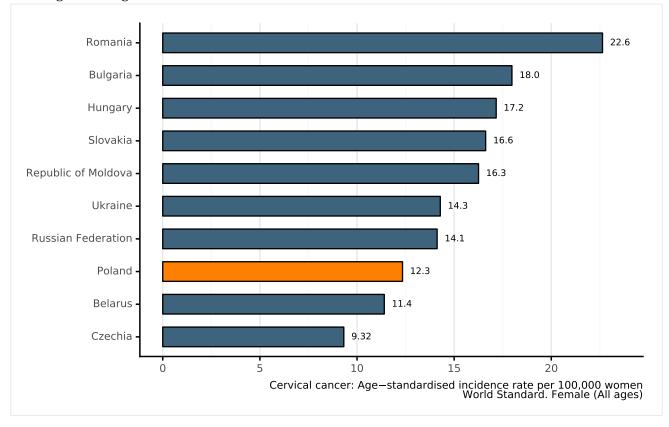


Figure 73: Age-standardised incidence rates of cervical cancer of Poland (estimates for 2020)

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.

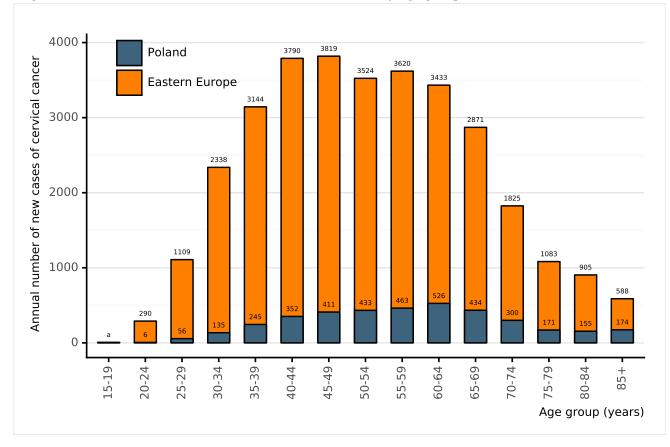
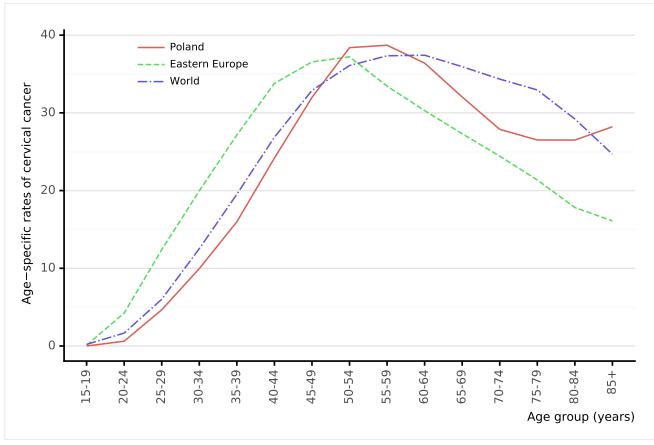
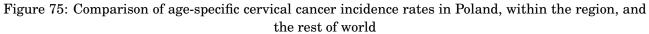


Figure 74: Annual number of new cases of cervical cancer by age group in Poland (estimates for 2020)

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 Poland and 7 cases for Eastern Europe in the 15-19 age group.





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.

9.1.2 Anal cancer incidence in Poland across Eastern Europe

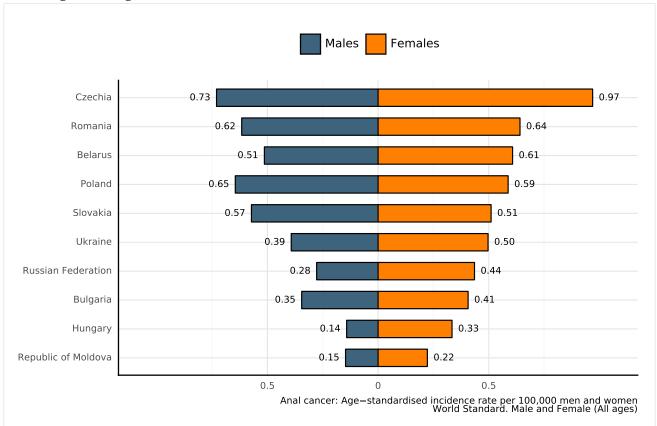


Figure 76: Age-standardised incidence rates of anal cancer of Poland (estimates for 2020)

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.

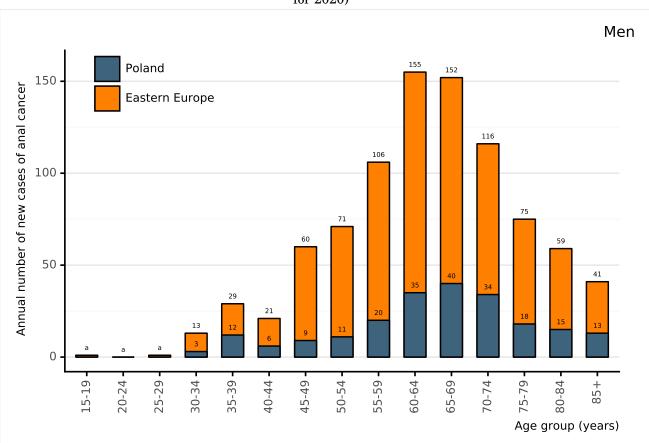


Figure 77: Annual number of new cases of anal cancer among men by age group in Poland (estimates for 2020)

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 Poland and 1 cases for Eastern Europe in the 15-19 age group. 0 cases for Poland and 0 cases for Eastern Europe in the 20-24 age group. 0 cases for Poland and 1 cases for

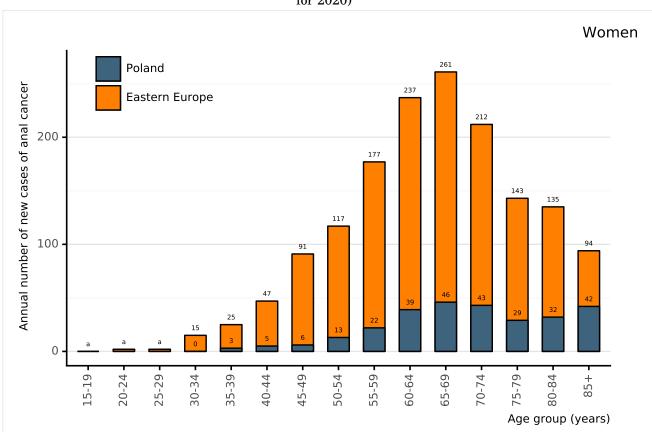


Figure 78: Annual number of new cases of anal cancer among women by age group in Poland (estimates for 2020)

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods ^a 0 cases for Poland and 0 cases for Eastern Europe in the 15-19 age group. 0 cases for Poland and 2 cases for Eastern Europe in the 20-24 age group. 0 cases for Poland and 2 cases for Eastern Europe in the 25-29 age group. Data Sources:

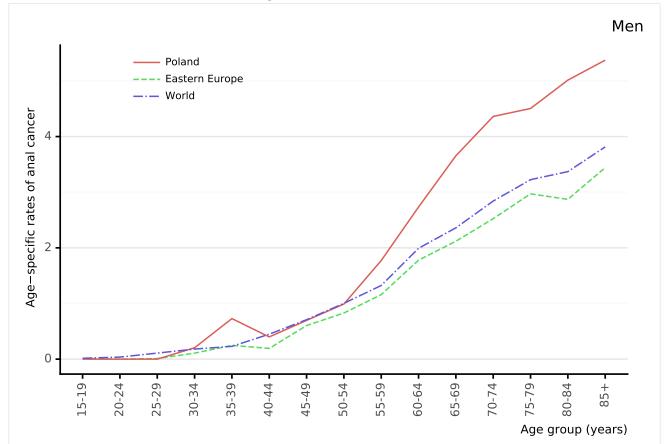


Figure 79: Comparison of age-specific anal cancer incidence rates among men by age in Poland, within the region, and the rest of world

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. <u>Data Sources:</u> 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].

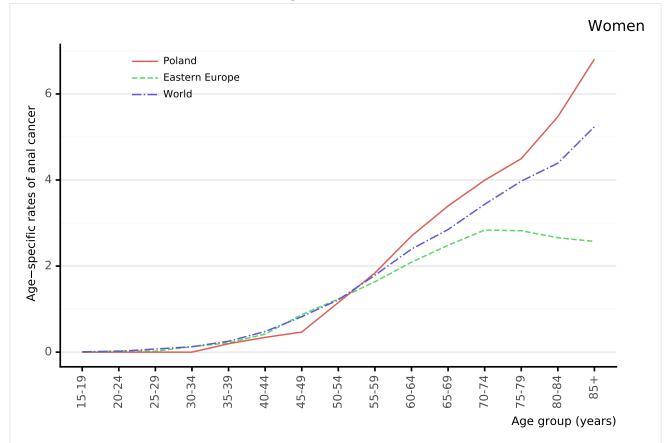


Figure 80: Comparison of age-specific anal cancer incidence rates among women by age in Poland, within the region, and the rest of world

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.

9.1.3 Vulva cancer incidence in Poland across Eastern Europe

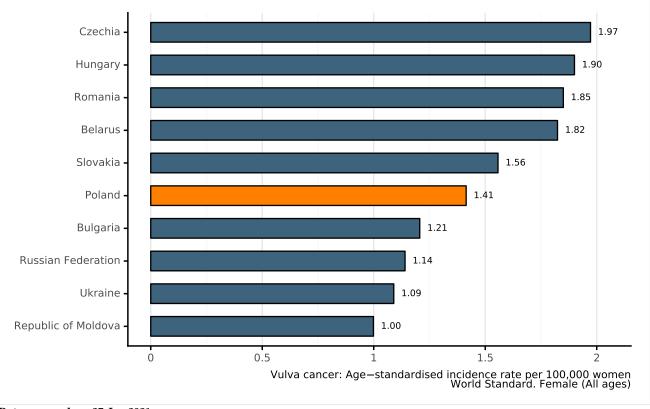


Figure 81: Age-standardised incidence rates of vulva cancer of Poland (estimates for 2020)

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:

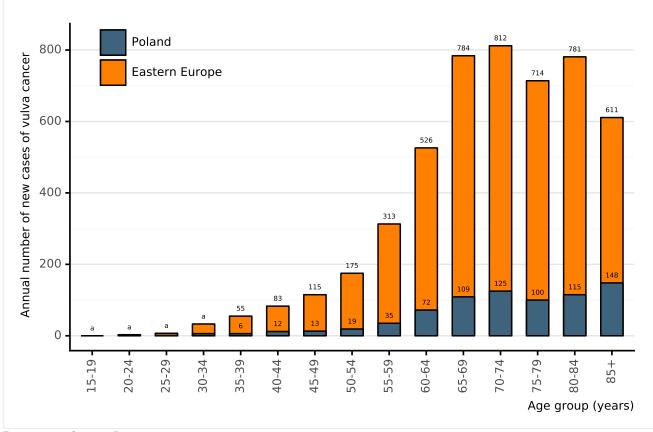


Figure 82: Annual number of new cases of vulva cancer by age group in Poland (estimates for 2020)

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 Poland and 0 cases for Eastern Europe in the 15-19 age group. 0 cases for Poland and 3 cases for Eastern Europe in the 20-24 age group. 1 cases for Poland and 7 cases for Eastern Europe in the 25-29 age group. 6 cases for Poland and 33 cases for Eastern Europe in the 30-34 age group.

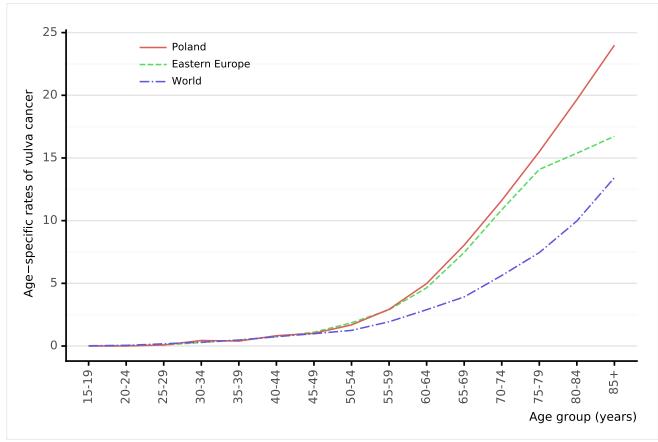


Figure 83: Comparison of age-specific vulva cancer incidence rates in Poland, within the region, and the rest of world

Data accessed on 27 Jan 2021

Data accessed on 2/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].

9.1.4 Vaginal cancer incidence in Poland across Eastern Europe

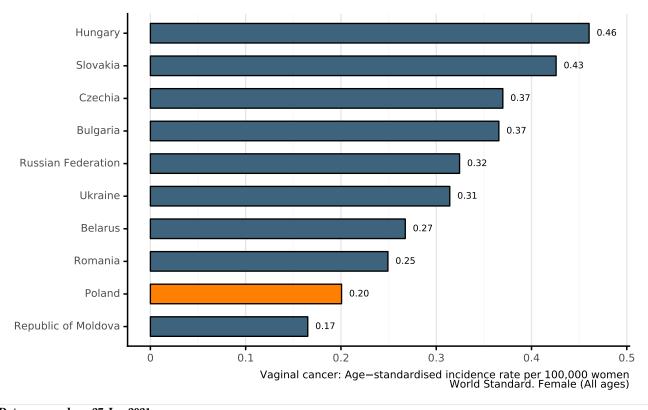
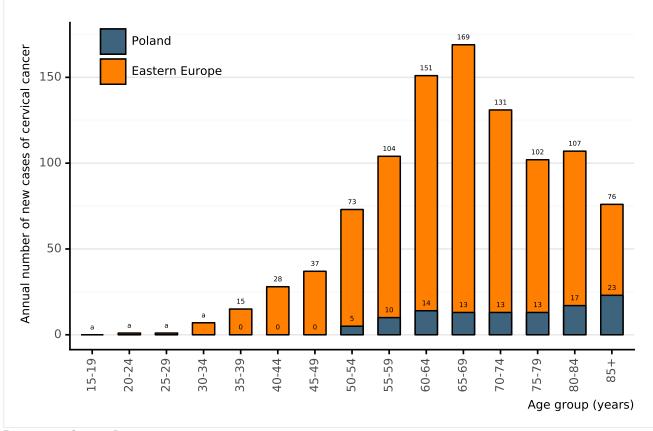


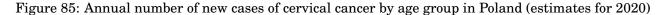
Figure 84: Age-standardised incidence rates of vaginal cancer of Poland (estimates for 2020)

Data accessed on 27 Jan 2021

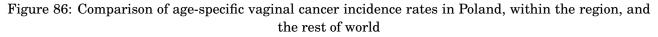
For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods ^a Rates per 100,000 women per year.

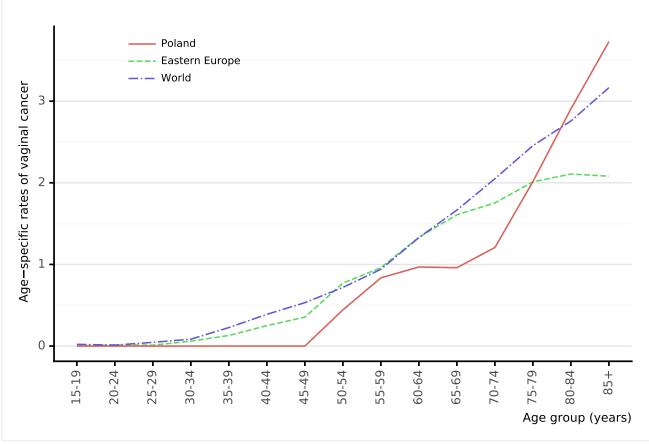
Data Sources:





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 Poland and 0 cases for Eastern Europe in the 15-19 age group. 0 cases for Poland and 1 cases for Eastern Europe in the 20-24 age group. 0 cases for Poland and 1 cases for Eastern Europe in the 25-29 age group. 0 cases for Poland and 7 cases for Eastern Europe in the 30-34 age group.





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. <u>Data Sources</u>: 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].

9.1.5 Penile cancer incidence in Poland across Eastern Europe

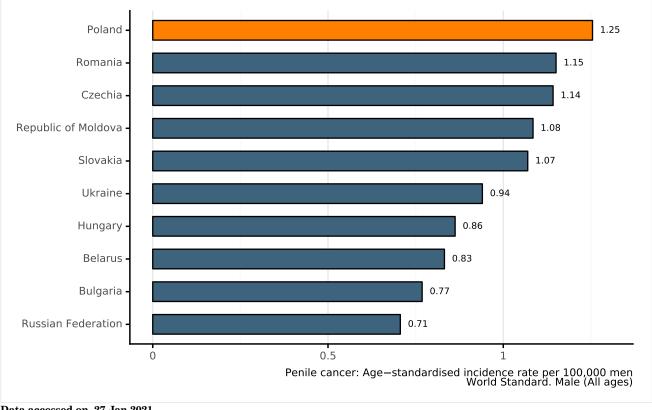


Figure 87: Age-standardised incidence rates of penile cancer of Poland (estimates for 2020)

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 Rates per 100,000 men per year.

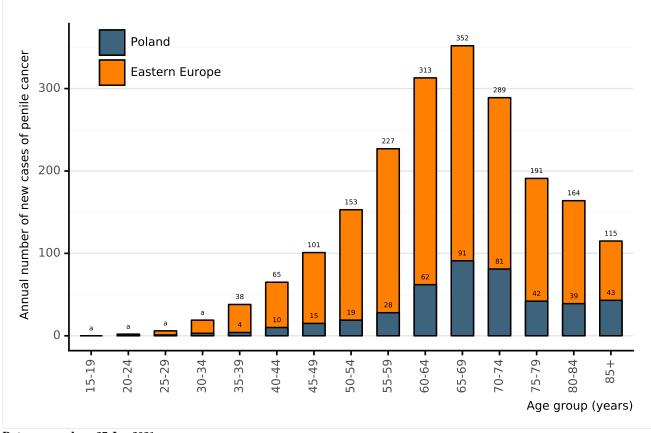


Figure 88: Annual number of new cases of penile cancer by age group in Poland (estimates for 2020)

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 Poland and 0 cases for Eastern Europe in the 15-19 age group. 1 cases for Poland and 2 cases for Eastern Europe in the 20-24 age group. 1 cases for Poland and 6 cases for Eastern Europe in the 25-29 age group. 3 cases for Poland and 19 cases for Eastern Europe in the 30-34 age group.

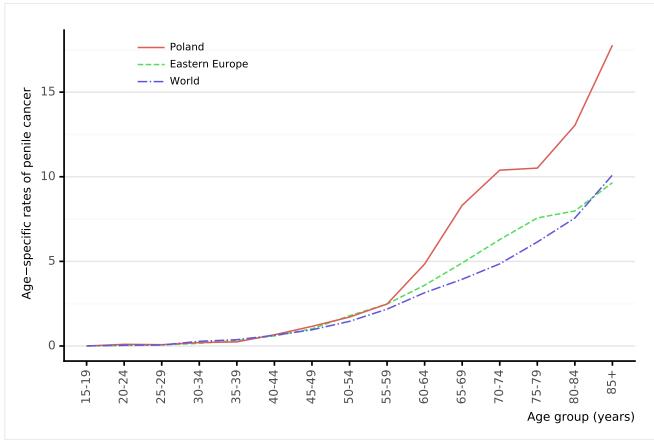


Figure 89: Comparison of age-specific penile cancer incidence rates in Poland, within the region, and the rest of world

Data accessed on 27 Jan 2021

Data accessed on 2/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].

9.1.6 Oropharyngeal cancer incidence in Poland across Eastern Europe

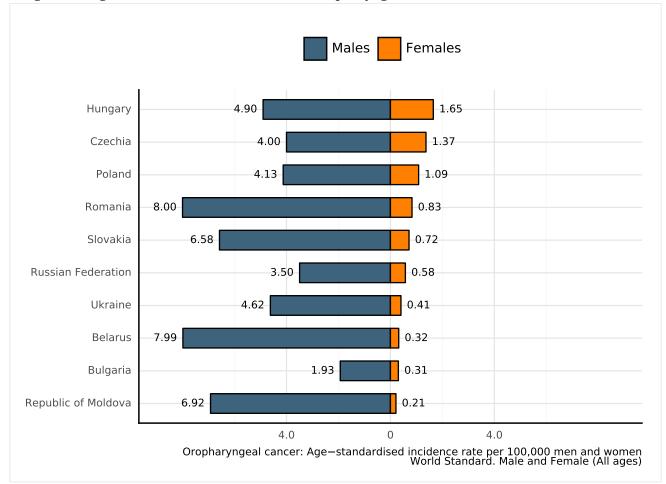


Figure 90: Age-standardised incidence rates of oropharyngeal cancer of Poland (estimates for 2020)

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.

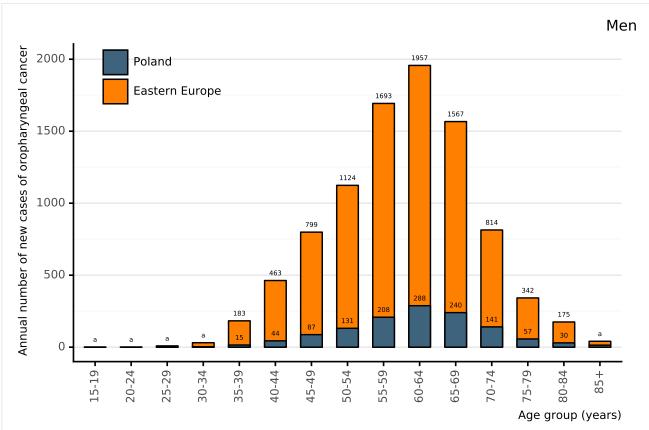


Figure 91: Annual number of new cases of oropharyngeal cancer among men by age group in Poland (estimates for 2020)

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 Poland and 1 cases for Eastern Europe in the 15-19 age group. 0 cases for Poland and 1 cases for Eastern Europe in the 20-24 age group. 2 cases for Poland and 9 cases for Eastern Europe in the 25-29 age group. 3 cases for Poland and 31 cases for Eastern Europe in the 30-34 age group. 14 cases for Poland and 41 cases for Eastern Europe in the 85+ age group.

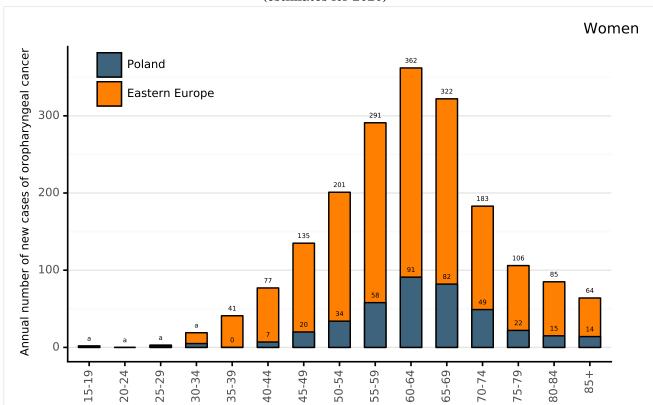


Figure 92: Annual number of new cases of oropharyngeal cancer among women by age group in Poland (estimates for 2020)

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 Poland and 2 cases for Eastern Europe in the 15-19 age group. 0 cases for Poland and 0 cases for Eastern Europe in the 20-24 age group. 2 cases for Poland and 3 cases for Eastern Europe in the 25-29 age group. 5 cases for Poland and 19 cases for Eastern 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 Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

Age group (years)

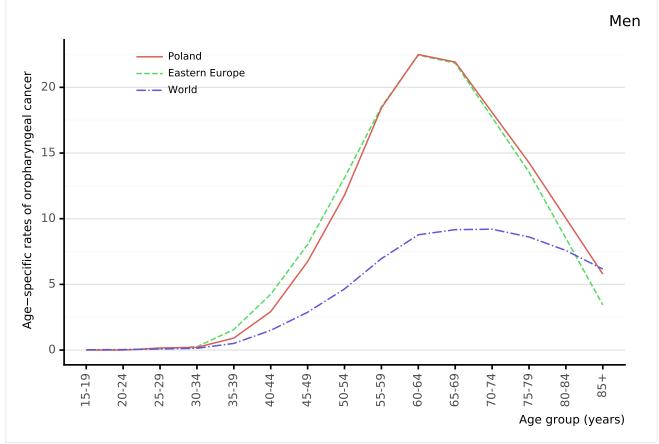
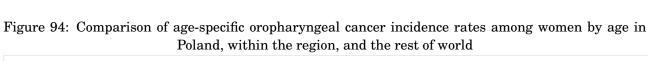
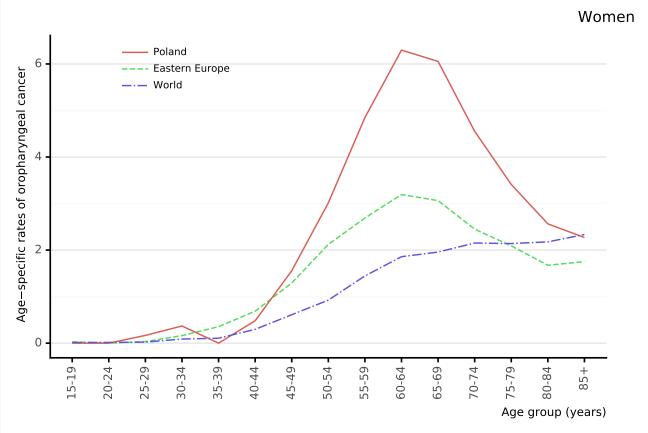


Figure 93: Comparison of age-specific oropharyngeal cancer incidence rates among men by age in Poland, within the region, and the rest of world

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.

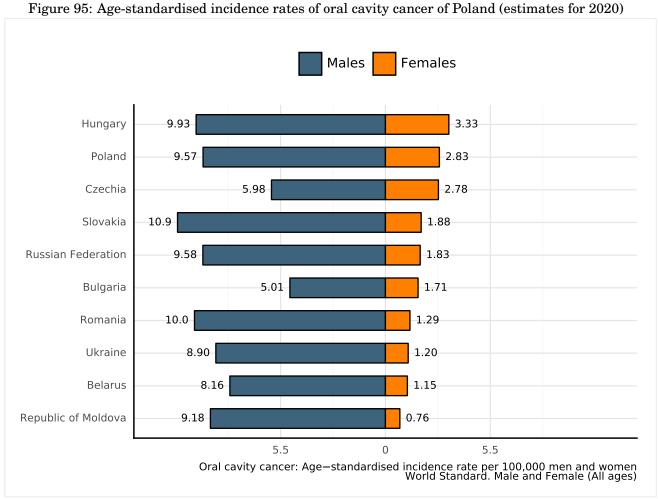




Data accessed on 27 Jan 2021

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.

9.1.7 Oral cavity cancer incidence in Poland across Eastern Europe



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.

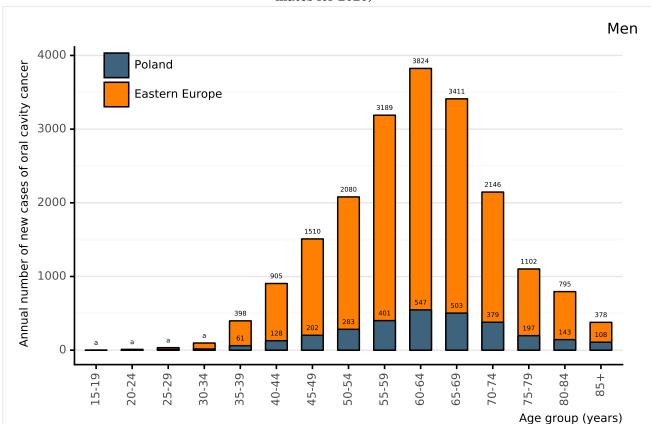


Figure 96: Annual number of new cases of oral cavity cancer among men by age group in Poland (estimates for 2020)

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 Poland and 1 cases for Eastern Europe in the 15-19 age group. 6 cases for Poland and 11 cases for Eastern Europe in the 20-24 age group. 10 cases for Poland and 34 cases for Eastern Europe in the 25-29 age group. 16 cases for Poland and 97 cases for Eastern Europe in the 30-34 age group.

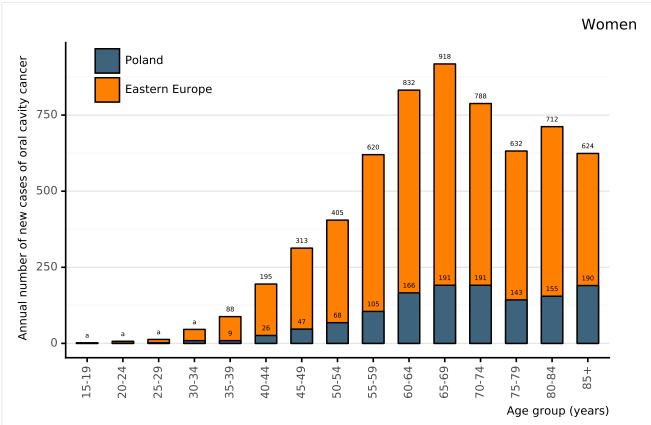


Figure 97: Annual number of new cases of oral cavity cancer among women by age group in Poland (estimates for 2020)

Data accessed on 27 Jan 2021

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods ^a 1 cases for Poland and 2 cases for Eastern Europe in the 15-19 age group. 1 cases for Poland and 7 cases for Eastern Europe in the 20-24 age group. 2 cases for Poland and 13 cases for Eastern Europe in the 25-29 age group. 9 cases for Poland and 46 cases for Eastern Europe in the 30-34 age group.

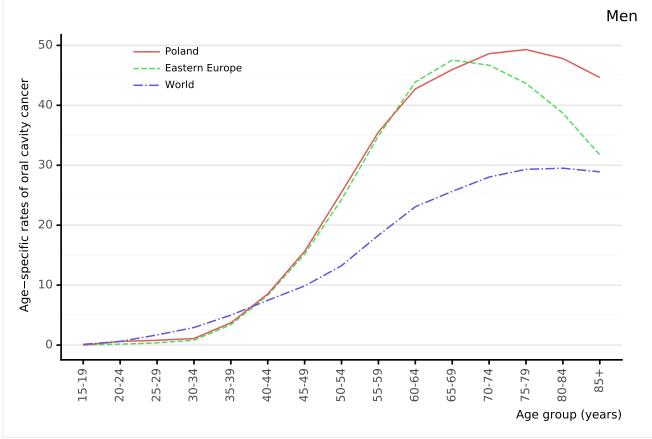


Figure 98: Comparison of age-specific oral cavity cancer incidence rates among men by age in Poland, within the region, and the rest of world

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.

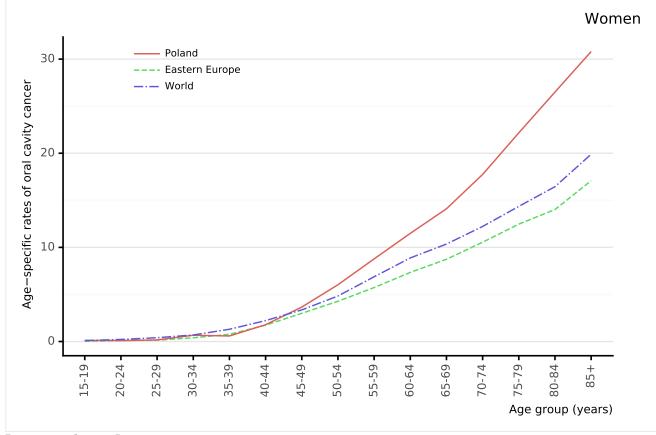


Figure 99: Comparison of age-specific oral cavity cancer incidence rates among women by age in Poland, within the region, and the rest of world

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.

Males Females Hungary 8.23 1.78 Poland 8.39 1.11 Republic of Moldova 11.9 0.72 Romania 11.4 0.61 7.67 Bulgaria 0.61 Czechia 4.27 0.60 Slovakia 8.69 0.43 **Russian Federation** 6.61 0.38 Ukraine 7.74 0.30 0.19 Belarus 9.22 6.0 0 6.0 Laryngeal cancer: Age-standardised incidence laryngeal per 100,000 men and women World Standard. Male and Female (All ages)

9.1.8 Laryngeal cancer incidence in Poland across Eastern Europe

Figure 100: Age-standardised incidence rates of laryngeal cancer of Poland (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.

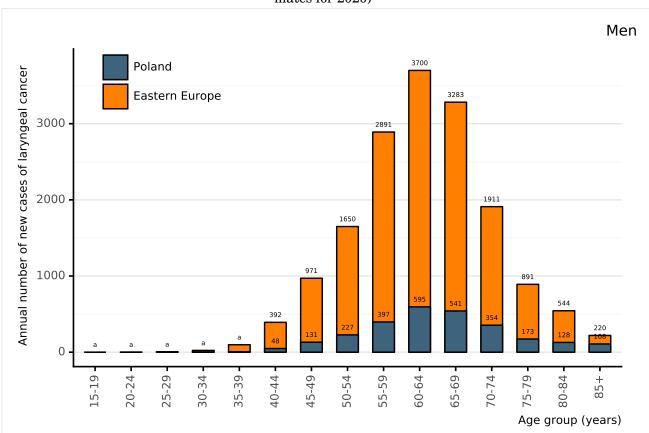


Figure 101: Annual number of new cases of laryngeal cancer among men by age group in Poland (estimates for 2020)

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 Poland and 0 cases for Eastern Europe in the 15-19 age group. 0 cases for Poland and 1 cases for Eastern Europe in the 20-24 age group. 1 cases for Poland and 5 cases for Eastern Europe in the 25-29 age group. 3 cases for Poland and 23 cases for Eastern Europe in the 30-34 age group. 6 cases for Poland and 98 cases for Eastern Europe in the 35-39 age

Eastern Europe in the 20-20 age group. 5 cases for Foland and 20 cases for Eastern Europe in the Correst Eastern Eastern

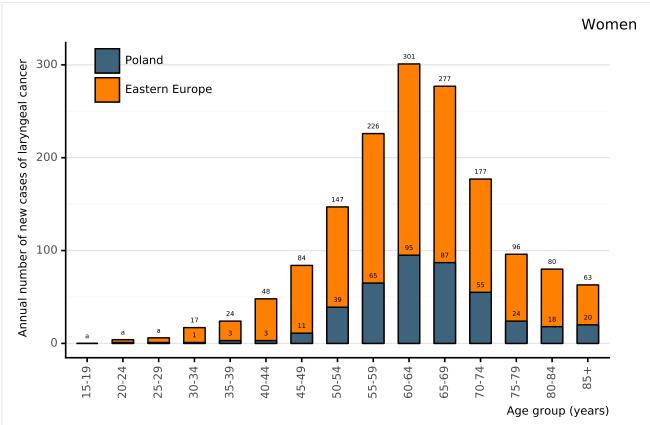


Figure 102: Annual number of new cases of laryngeal cancer among women by age group in Poland (estimates for 2020)

For more detailed methods of estimation please refer to http://gco.iarc.fr/today/data-sources-methods ^a 0 cases for Poland and 0 cases for Eastern Europe in the 15-19 age group. 1 cases for Poland and 4 cases for Eastern Europe in the 20-24 age group. 1 cases for Poland and 6 cases for Eastern Europe in the 25-29 age group. Poto Source:

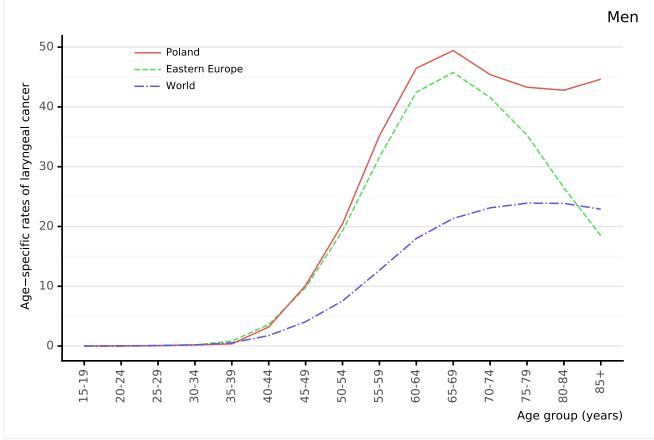


Figure 103: Comparison of age-specific laryngeal cancer incidence rates among men by age in Poland, within the region, and the rest of world

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.

- 120 -

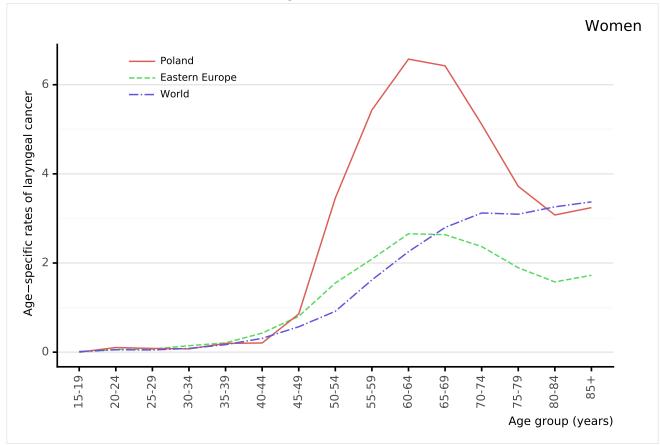


Figure 104: Comparison of age-specific laryngeal cancer incidence rates among women by age in Poland, within the region, and the rest of world

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.

9.2 Mortality

9.2.1 Cervical cancer mortality in Poland across Eastern Europe

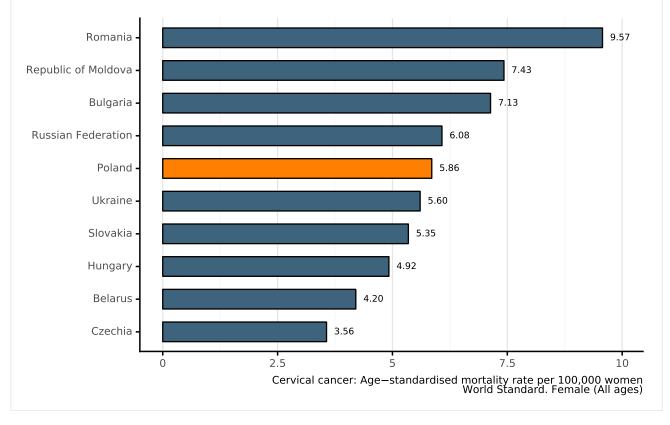


Figure 105: Age-standardised mortality rates of cervical cancer of Poland (estimates for 2020)

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:

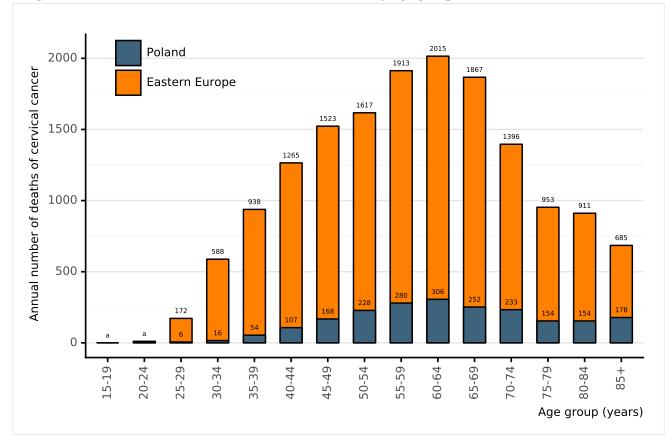


Figure 106: Annual number of deaths of cervical cancer by age group in Poland (estimates for 2020)

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 Poland and 0 cases for Eastern Europe in the 15-19 age group. 1 cases for Poland and 11 cases for Eastern Europe in the 20-24 age group.

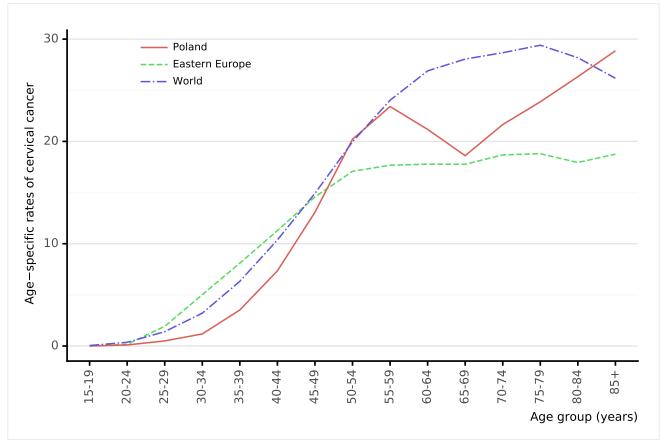


Figure 107: Comparison of age-specific cervical cancer mortality rates in Poland, within the region, and the rest of world

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.

9.2.2 Anal cancer mortality in Poland across Eastern Europe

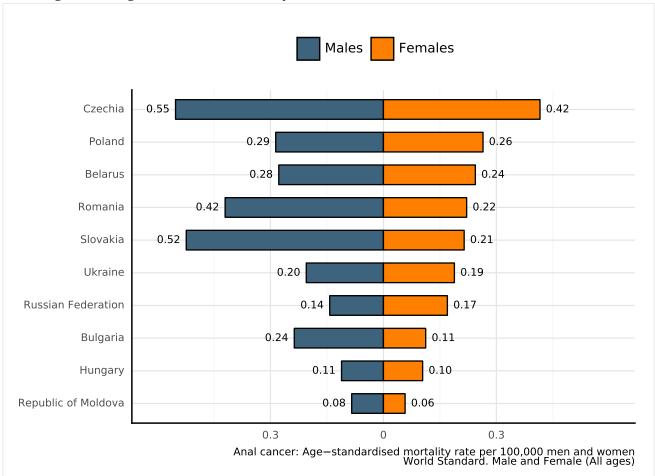


Figure 108: Age-standardised mortality rates of anal cancer of Poland (estimates for 2020)

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.

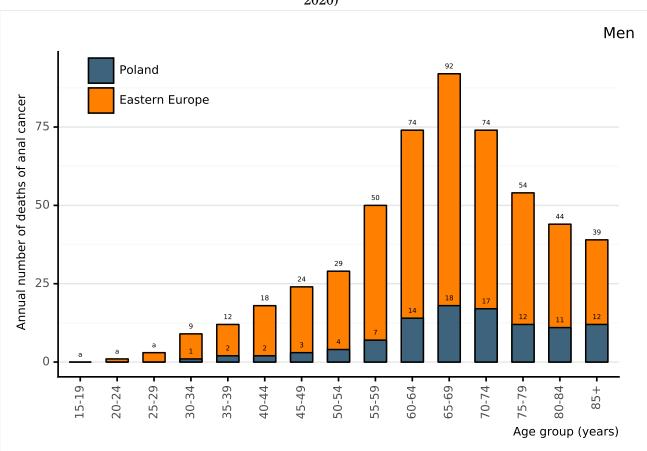


Figure 109: Annual number of deaths of anal cancer among men by age group in Poland (estimates for 2020)

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 Poland and 0 cases for Eastern Europe in the 15-19 age group. 0 cases for Poland and 1 cases for Eastern Europe in the 20-24 age group. 0 cases for Poland and 3 cases for Eastern Europe in the 25-29 age group.

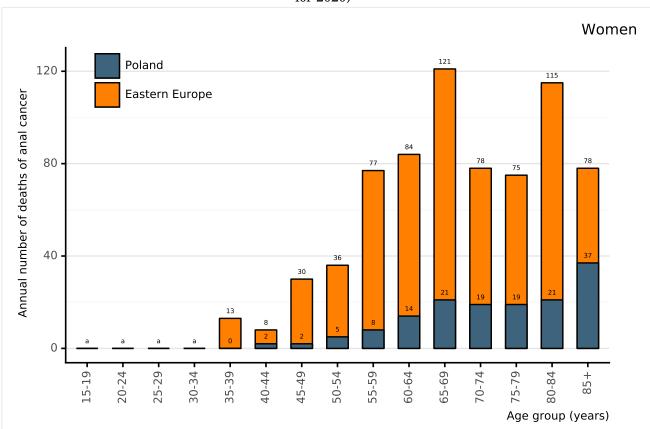


Figure 110: Annual number of deaths of anal cancer among women by age group in Poland (estimates for 2020)

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 Poland and 0 cases for Eastern Europe in the 15-19 age group. 0 cases for Poland and 0 cases for Eastern Europe in the 20-24 age group. 0 cases for Poland and 0 cases for Eastern Europe in the 25-29 age group. 0 cases for Poland and 0 cases for Eastern Europe in the 30-34 age group.

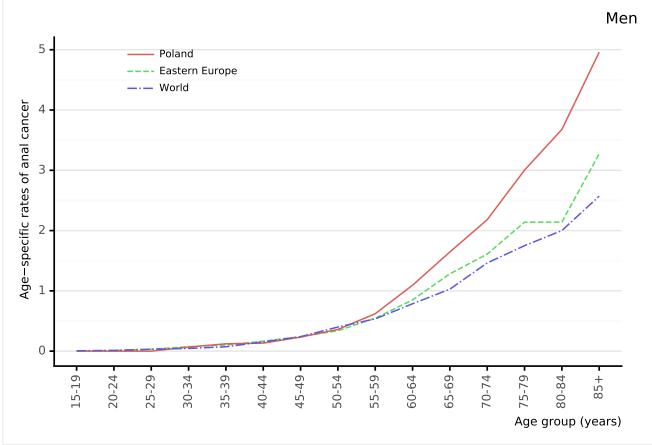


Figure 111: Comparison of age-specific anal cancer mortality rates among men by age in Poland, within the region, and the rest of world

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. <u>Data Sources:</u> 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].

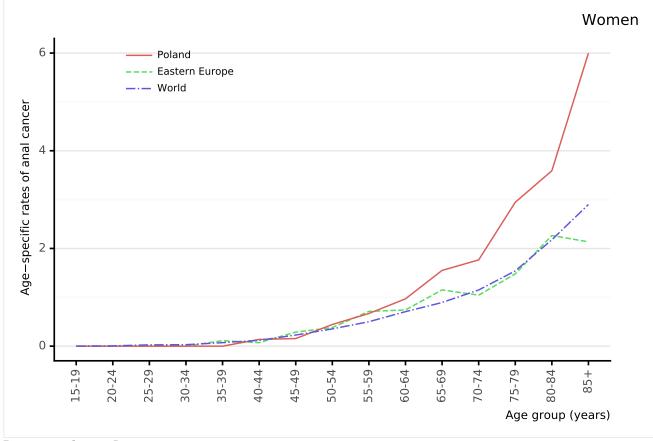


Figure 112: Comparison of age-specific anal cancer mortality rates among women by age in Poland, within the region, and the rest of world

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.

9.2.3 Vulva cancer mortality in Poland across Eastern Europe

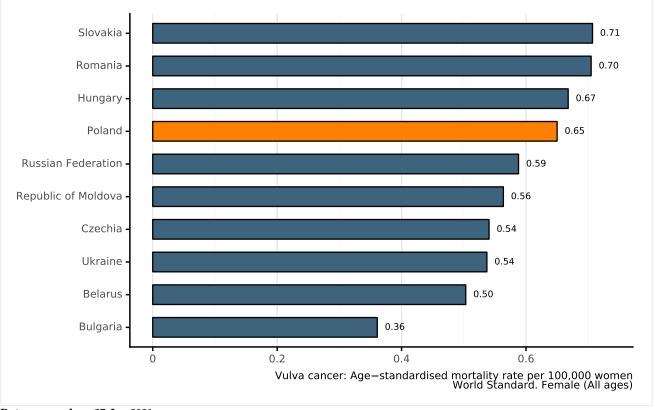


Figure 113: Age-standardised mortality rates of vulva cancer of Poland (estimates for 2020)

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:

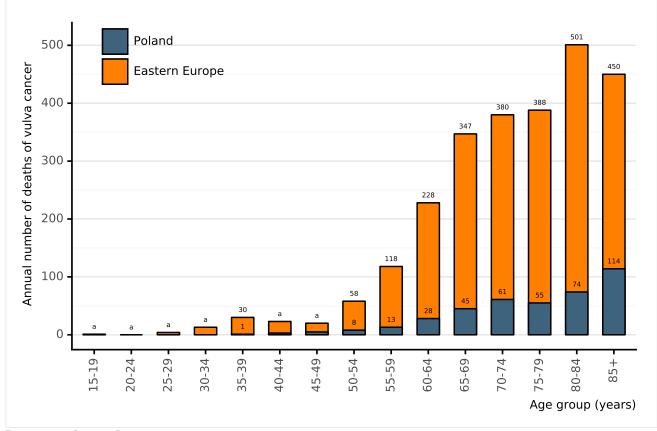


Figure 114: Annual number of deaths of vulva cancer by age group in Poland (estimates for 2020)

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 Poland and 1 cases for Eastern Europe in the 15-19 age group. 0 cases for Poland and 0 cases for Eastern Europe in the 20-24 age group. 0 cases for Poland and 4 cases for Eastern Europe in the 25-29 age group. 0 cases for Poland and 13 cases for Eastern Europe in the 30-34 age group. 3 cases for Poland and 23 cases for Eastern Europe in the 40-44 age group. 5 cases for Poland and 20 cases for Eastern Europe in the 45-49 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 Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

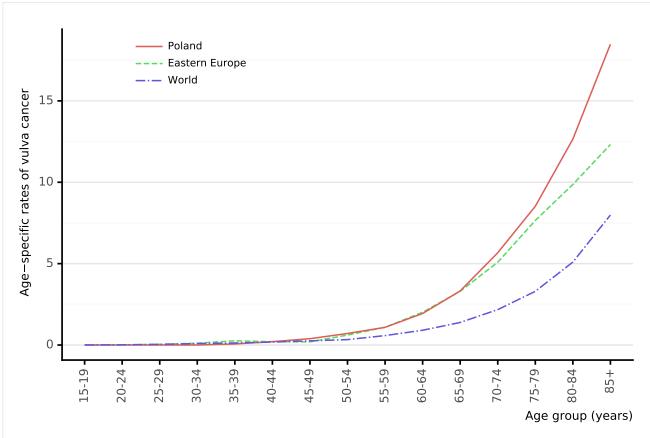


Figure 115: Comparison of age-specific vulva cancer mortality rates in Poland, within the region, and the rest of world

Data accessed on 27 Jan 2021

Data accessed on 2/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].

9.2.4 Vaginal cancer mortality in Poland across Eastern Europe

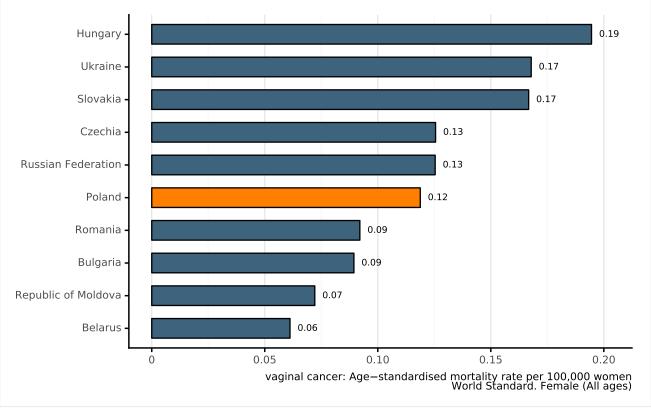


Figure 116: Age-standardised mortality rates of vaginal cancer of Poland (estimates for 2020)

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:

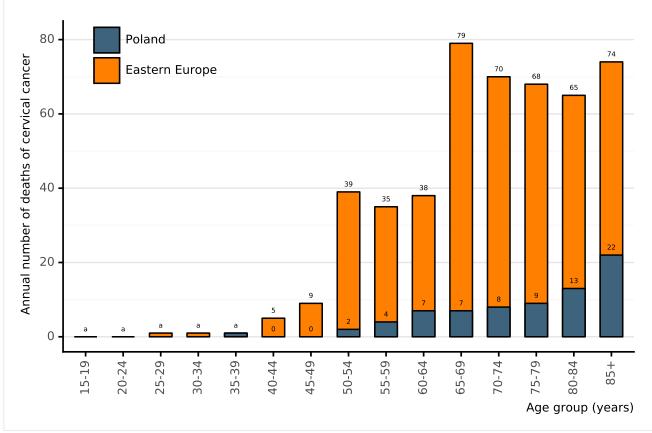


Figure 117: Annual number of deaths of cervical cancer by age group in Poland (estimates for 2020)

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 Poland and 0 cases for Eastern Europe in the 15-19 age group. 0 cases for Poland and 0 cases for Eastern Europe in the 20-24 age group. 0 cases for Poland and 1 cases for Eastern Europe in the 25-29 age group. 0 cases for Poland and 1 cases for Eastern Europe in the 30-34 age group. 1 cases for Poland and 1 cases for Eastern Europe in the 35-39 age group.

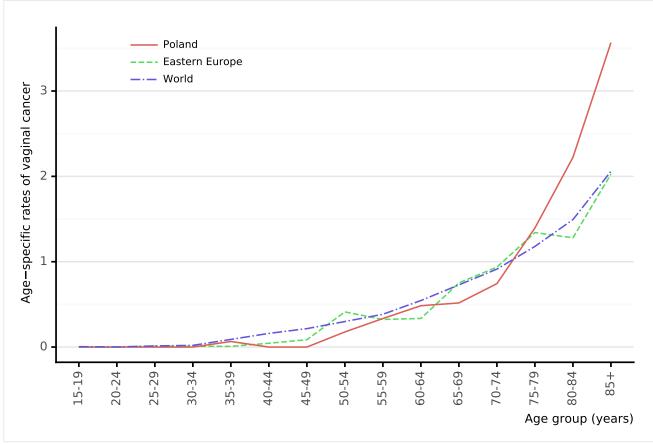


Figure 118: Comparison of age-specific vaginal cancer mortality rates in Poland, within the region, and the rest of world

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. <u>Data Sources</u>: 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].

9.2.5 Penile cancer mortality in Poland across Eastern Europe

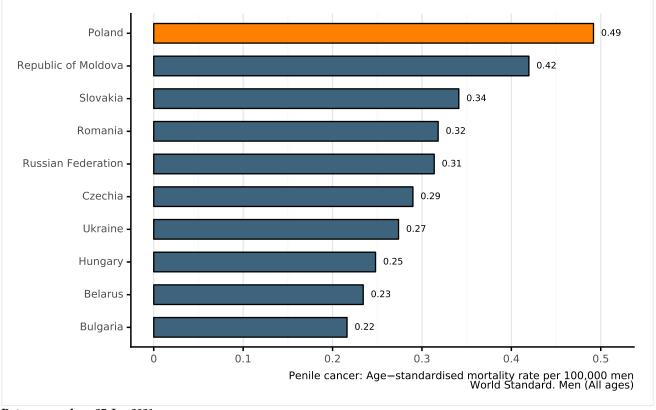
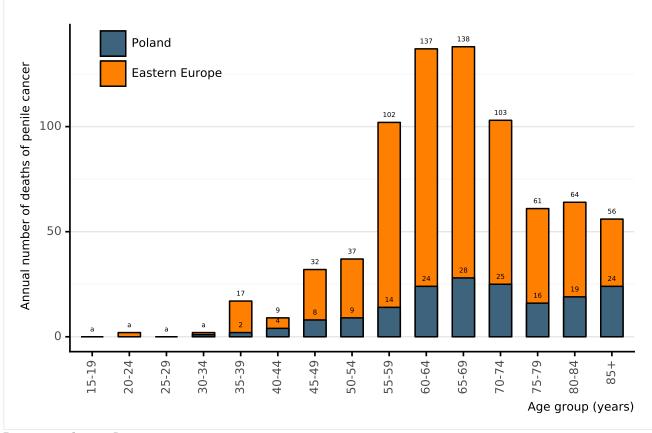


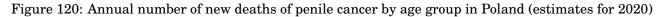
Figure 119: Age-standardised mortality rates of penile cancer of Poland (estimates for 2020)

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:





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 Poland and 0 cases for Eastern Europe in the 15-19 age group. 0 cases for Poland and 2 cases for Eastern Europe in the 20-24 age group. 0 cases for Poland and 0 cases for Eastern Europe in the 25-29 age group. 1 cases for Poland and 2 cases for Eastern Europe in the 30-34 age group.

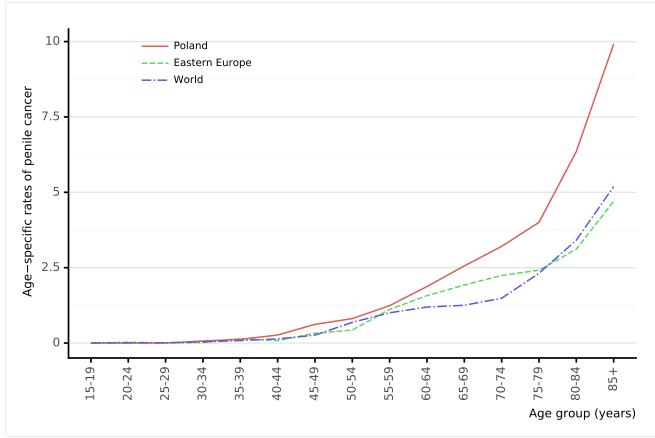


Figure 121: Comparison of age-specific penile cancer mortality rates in Poland, within the region, and the rest of world

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. Deta Sources

9.2.6 Oropharyngeal cancer mortality in Poland across Eastern Europe

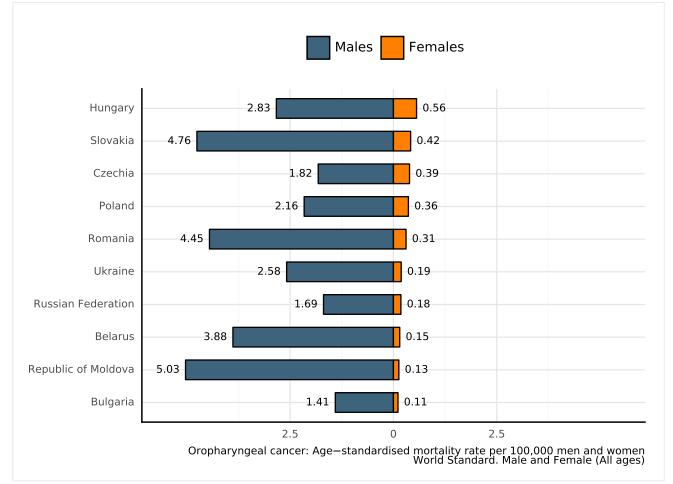


Figure 122: Age-standardised mortality rates of oropharyngeal cancer of Poland (estimates for 2020)

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.

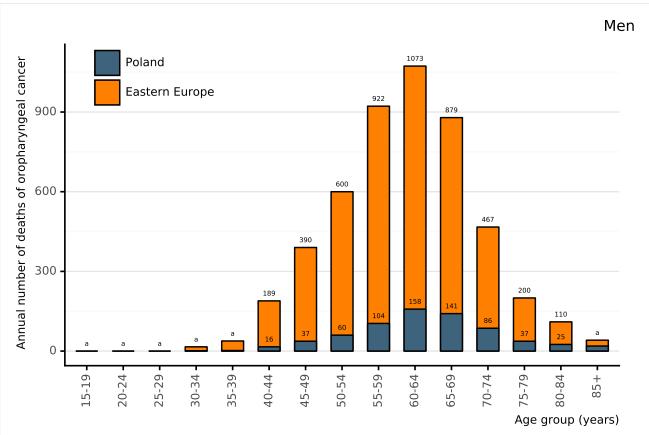
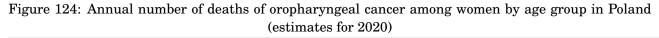
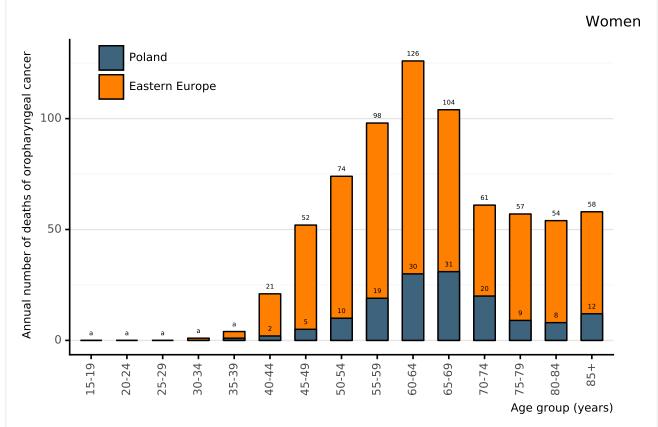


Figure 123: Annual number of deaths of oropharyngeal cancer among men by age group in Poland (estimates for 2020)

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 Poland and 0 cases for Eastern Europe in the 15-19 age group. 0 cases for Poland and 0 cases for Eastern Europe in the 20-24 age group. 0 cases for Poland and 0 cases for Eastern Europe in the 25-29 age group. 1 cases for Poland and 16 cases for Eastern Europe in the 30-34 age group. 2 cases for Poland and 38 cases for Eastern Europe in the 35-39 age group. 19 cases for Poland and 41 cases for Eastern Europe in the 85+ age group.





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 Poland and 0 cases for Eastern Europe in the 15-19 age group. 0 cases for Poland and 0 cases for Eastern Europe in the 25-29 age group. 0 cases for Poland and 1 cases for Eastern Europe in the 35-39 age group.

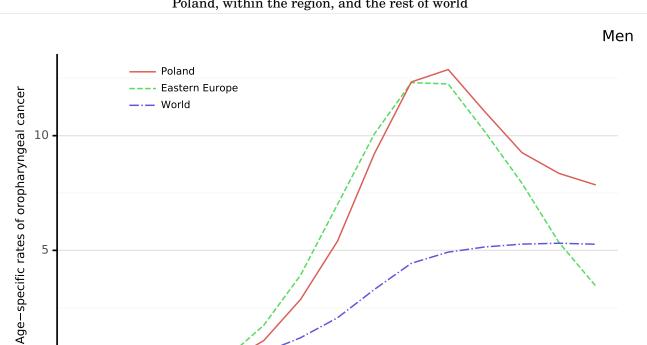


Figure 125: Comparison of age-specific oropharyngeal cancer mortality rates among men by age in Poland, within the region, and the rest of world

Data accessed on 27 Jan 2021

15-19.

20-24

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.

30-34

35-39 -

40-44

45-49.

25-29.

5

0

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].

50-54

55-59

60-64

62-69

70-74

75-79

80-84

Age group (years)

85+

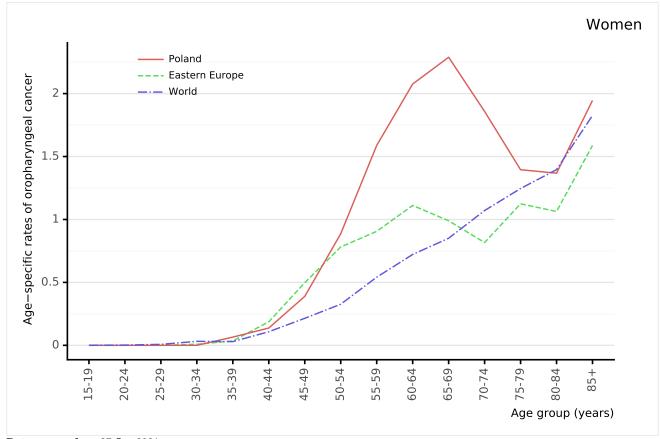


Figure 126: Comparison of age-specific oropharyngeal cancer mortality rates among women by age in Poland, within the region, and the rest of world

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. <u>Data Sources:</u> 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].

9.2.7 **Oral cavity cancer mortality in Poland across Eastern Europe**

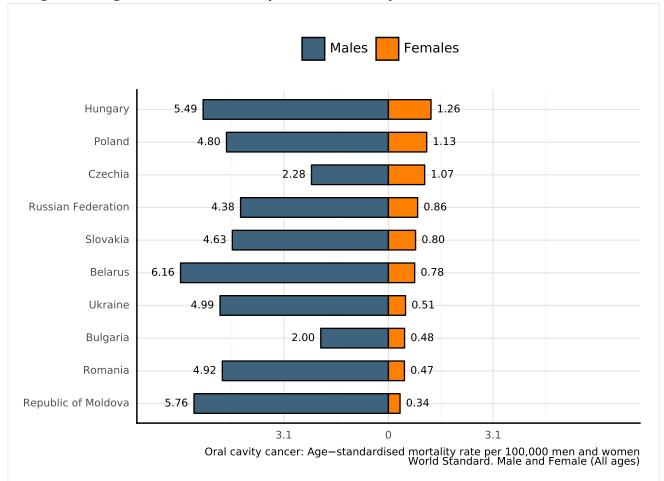


Figure 127: Age-standardised mortality rates of oral cavity cancer of Poland (estimates for 2020)

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.

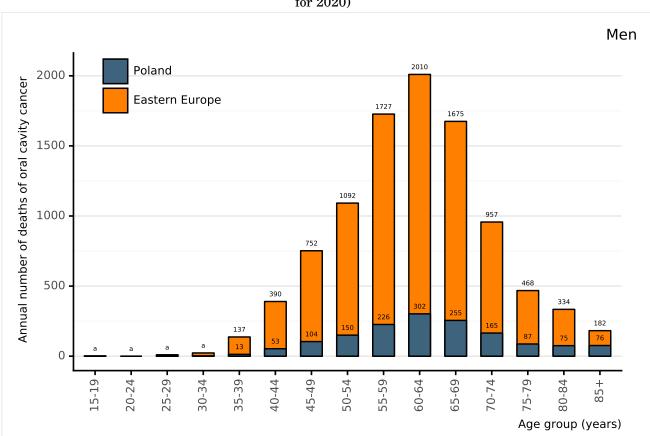


Figure 128: Annual number of deaths of oral cavity cancer among men by age group in Poland (estimates for 2020)

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 Poland and 2 cases for Eastern Europe in the 15-19 age group. 0 cases for Poland and 0 cases for Eastern Europe in the 20-24 age group. 1 cases for Poland and 10 cases for Eastern Europe in the 25-29 age group. 2 cases for Poland and 23 cases for Eastern Europe in the 30-34 age group.

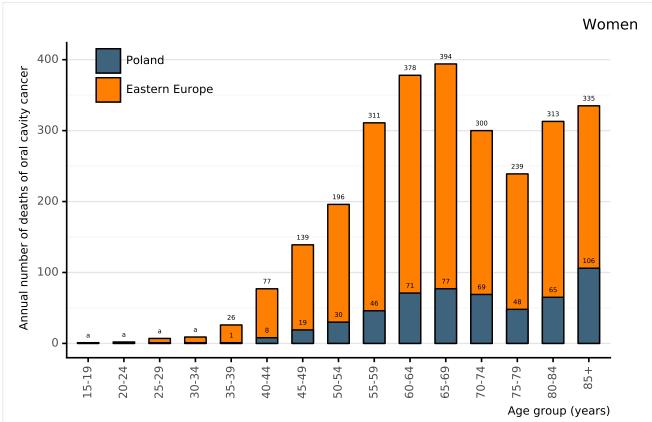


Figure 129: Annual number of deaths of oral cavity cancer among women by age group in Poland (estimates for 2020)

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 Poland and 1 cases for Eastern Europe in the 15-19 age group. 0 cases for Poland and 2 cases for Eastern Europe in the 20-24 age group. 1 cases for Poland and 7 cases for Eastern Europe in the 25-29 age group. 1 cases for Poland and 9 cases for Eastern Europe in the 30-34 age group.

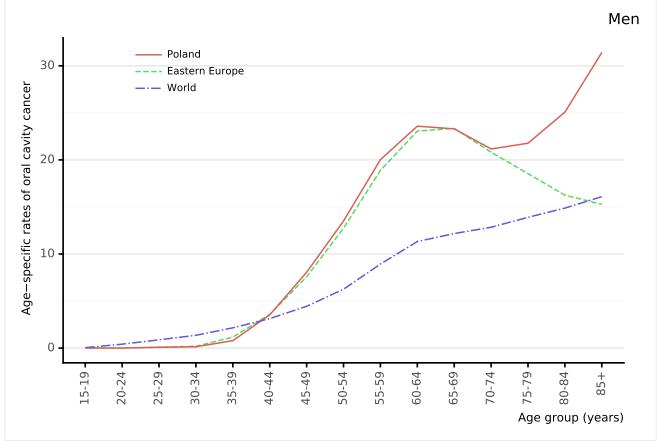


Figure 130: Comparison of age-specific oral cavity cancer mortality rates among men by age in Poland, within the region, and the rest of world

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. <u>Data Sources</u>: 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].

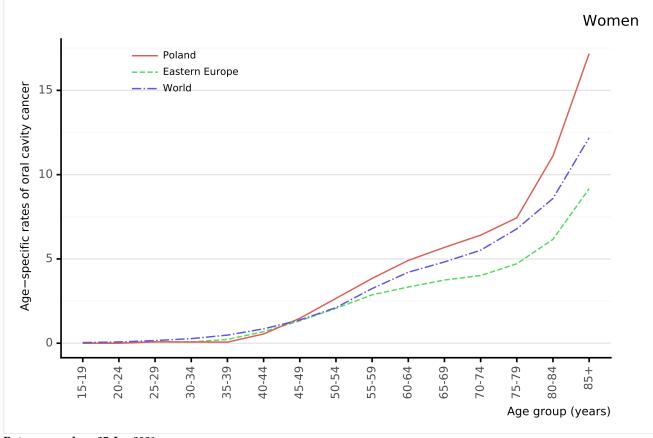


Figure 131: Comparison of age-specific oral cavity cancer mortality rates among women by age in Poland, within the region, and the rest of world

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.

9.2.8 Laryngeal cancer mortality in Poland across Eastern Europe

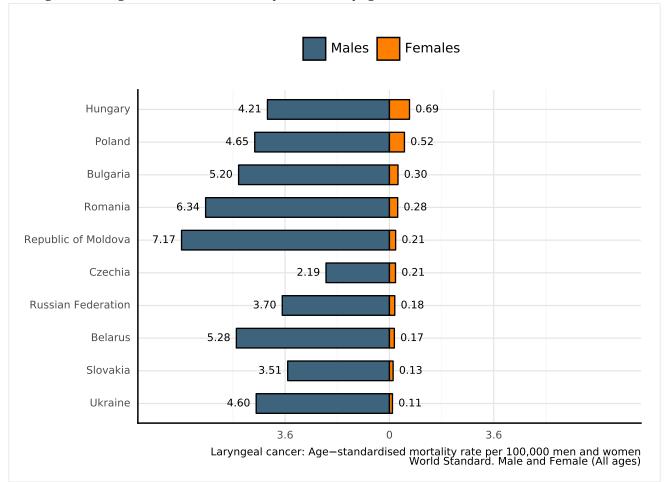


Figure 132: Age-standardised mortality rates of laryngeal cancer of Poland (estimates for 2020)

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.

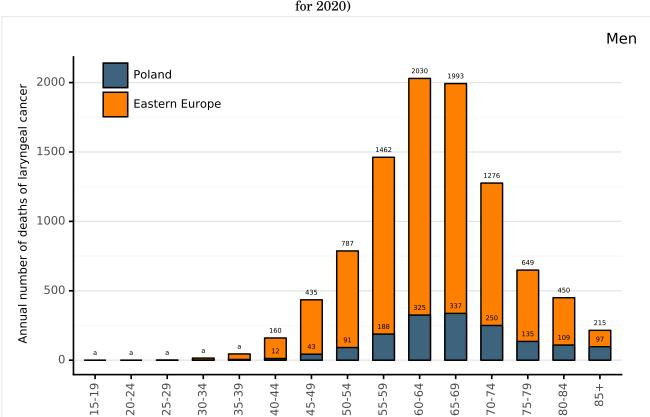


Figure 133: Annual number of deaths of laryngeal cancer among men by age group in Poland (estimates for 2020)

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 Poland and 0 cases for Eastern Europe in the 15-19 age group. 0 cases for Poland and 0 cases for Eastern Europe in the 20-24 age group. 0 cases for Poland and 1 cases for Eastern Europe in the 25-29 age group. 1 cases for Poland and 15 cases for Eastern Europe in the 30-34 age group. 5 cases for Poland and 45 cases for Eastern Europe in the 35-39 age

Eastern Europe in the 20-20 age group. I cases for Foland and To cases for Eastern Europe in the Correst Eastern Eastern

Age group (years)

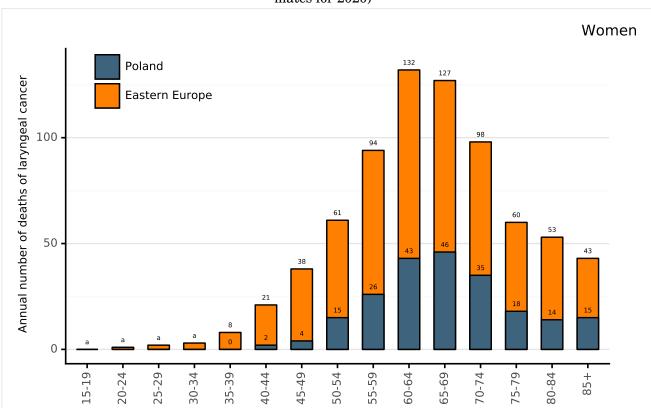


Figure 134: Annual number of deaths of laryngeal cancer among women by age group in Poland (estimates for 2020)

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 Poland and 0 cases for Eastern Europe in the 15-19 age group. 0 cases for Poland and 1 cases for Eastern Europe in the 20-24 age group. 0 cases for Poland and 2 cases for Eastern Europe in the 25-29 age group. 0 cases for Poland and 3 cases for Eastern 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 Research on Cancer. Available from: https://gco.iarc.fr/today, accessed [27 January 2021].

Age group (years)

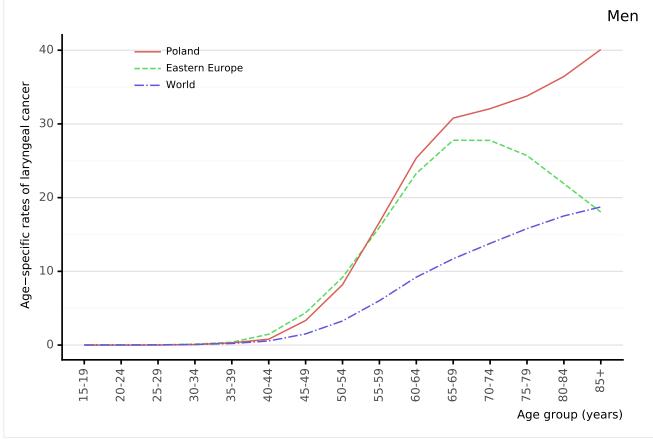


Figure 135: Comparison of age-specific laryngeal cancer mortality rates among men by age in Poland, within the region, and the rest of world

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.

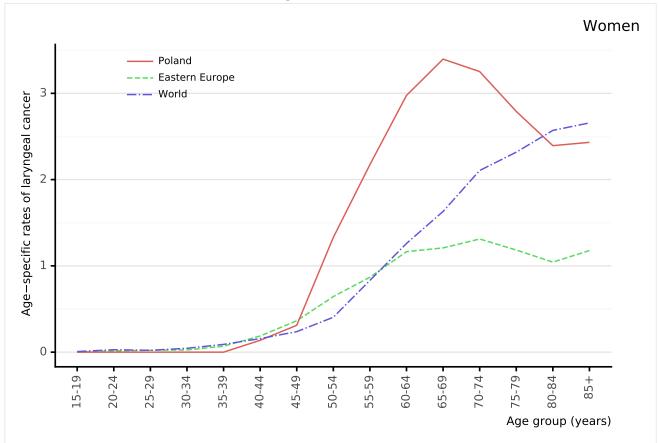


Figure 136: Comparison of age-specific laryngeal cancer mortality rates among women by age in Poland, within the region, and the rest of world

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.

10 Glossary

Table 49: Glossary		
Term	Definition	
Incidence	Incidence is the number of new cases arising in a given period in a speci- fied 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 per- sons 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. Com- plete 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 con- sidered 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 rele- vance 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

Table 49 – continued from previous page		
Term	Definition	
Cumulative risk	Cumulative incidence/mortality is the probability or risk of individuals get- ting/dying from the disease during a specified period. For cancer, it is ex- pressed 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- peting causes.	
Cytologically normal women	No abnormal cells are observed on the surface of their cervix upon cytology.	
Cervical Intraepithe- lial Neoplasia (CIN) / Squamous Intraepithe- lial 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 tis- sue 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 le- sions (LSIL/CIN-1)	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 le- sions (HSIL / CIN-2 / CIN-3 / CIS)	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 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 can- cer (ICC) / Cervical	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	
cancer Adenocarcinoma	only) to stage IV (the cancer has spread to distant organs, such as the liver). Invasive tumour with glandular and squamous elements intermingled	

Acknowledgments

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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)

Note to the reader

Anyone who is aware of relevant published data that may not have been included in the present report is encouraged to contact the HPV Information Centre for potential contributions.

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