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## 1.1. Cancer patients who received chemotherapy in the last 14 days of life (EOL-3)

### 1.1.1. Documentation sheet

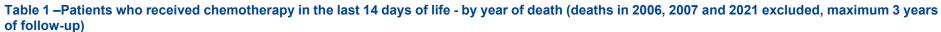
Description	Patients with cancer receiving chemotherapy in the last 14 days of life (% of cancer patients with poor prognosis who died)						
Calculation	Numerator: number of patients receiving chemotherapy in the last 14 days of life						
	Denominator: number of patients diagnosed with cancer with poor prognosis that died within the studied time period						
Rationale	The main goal of palliative care is to improve - or at least maintain - quality of life in patients near their death. In this way burdensome curative treatments, such as active cancer treatment in case of oncological patients, are stopped and the main focus of therapy becomes alleviation of pain and other symptoms. This indicator measures the aggressiveness of care in the last days of life of cancer patients.						
Data source	Belgian Cancer Registry (BCR), linked with IMA data						
Technical definitions	It is currently not possible to identify all palliative patients in administrative databases or in registries. Therefore, the indicate been restricted to patients diagnosed with cancer who have a poor prognosis (based on relative survival probability) and deceduring the study period.						
	Inclusion criteria						
	Incidence years: 2006-2020						
	Tumour selection based on the Pallcare project <sup>1</sup> : combination of topography and morphology according to Eurocare-4 <sup>2-3</sup> , a Eurocare-5 <sup>4</sup> : see EOL-1						
	Patients deceased before July 1 <sup>st</sup> , 2021						
	Age at diagnosis ≥18 years						
	Exclusion criteria						
	Patients with more than one invasive tumour (until 2020)						
	Patients without a Belgian residence at time of diagnosis						
	Patients without national social security number						
	Patients for whom no IMA data of the year of death were available (=4.1%)						
	Maximum 3 years of follow-up						
	Analyses were limited to patients who died before January 1st of the third year following the year of incidence. For example patients diagnosed in 2006 were included in case they died in 2006, 2007 or 2008; patients diagnosed in 2007 were included in case they died in 2007, 2008 or 2009, etc. These cohorts were defined to assure that for every diagnosis a similar follow-up period could be taken into account and to avoid bias by varying length of the follow-up period.						
	<u>Chemotherapy</u>						

·	Drug (ATC) selection: ATC category L01, minus some products that are (predominantly) used in non-oncological settings (e.g.						
	methotrexate (Ledertrexate), celecoxib (Celebrex), everolimus (Certican, Afinitor), Alemtuzumab (MabCampath)).						
Limitation	The true proportion is probably higher than the presented results, as patients treated within clinical trials can receive chemotherap provided by the sponsoring company and therefore it will not be detected within reimbursement (IMA-AIM) data.						
	No information on aggressiveness of care in terminally ill patients not suffering from cancer.						
	Very diverse agents (i.e. ATC L01 all together) are included in the analyses, with varying toxicity-profile and way of administration (orally versus IV).						
International comparability	This is not an international indicator. Some results are available in national reports or in specific scientific articles.						
Dimension	End-of-life care; Quality (appropriateness)						
Related indicators							
Reviewers	Cindy De Gendt (BCR), Lien Van Walle (BCR)						

#### 1.1.2. Results

An average of 10.5% of the cancer patients who died in the period between 2008 and 2020, received chemotherapy in the last 14 days of their life (see Table 1). The proportion was slightly higher in patients with chronic tumour types compared to patients with acute tumour types (13.5% versus 10.1%). More variation is noted between the individual tumour types: the proportion of patients who received chemotherapy in the last 14 days of life ranged from 3.8% for brain cancer to 35.5% for acute lymphatic leukaemia and chronic myeloid leukaemia and 33.2% for acute myeloid leukaemia (see Table 2). Generally higher percentages are observed in patients with haematological tumours.

A slightly higher proportion of patients received chemotherapy just before death in Wallonia compared to Brussels and Flanders (12.3% vs 11.1% and 9.5%, respectively).



	All Tumours			Acut	e Tumours	Chron	Chronic Tumours			
	Total	N with chemotherapy in the last 14 days of life		Total	N with chemotherapy in the last 14 days of life		Total	N with chemotherapy in the last 14 days of life		
	N	n	%	N	n	%	N	n	%	
2008	9597	1048	10.9	8385	902	10.8	1212	146	12.0	
2009	9455	1033	10.9	8243	876	10.6	1212	157	13.0	
2010	9873	1167	11.8	8614	995	11.6	1259	172	13.7	
2011	9950	1116	11.2	8770	961	11.0	1180	155	13.1	
2012	10059	1118	11.1	8813	925	10.5	1246	193	15.5	
2013	10120	1061	10.5	8906	872	9.8	1214	189	15.6	
2014	10048	1005	10.0	8882	860	9.7	1166	145	12.4	
2015	9876	913	9.2	8677	775	8.9	1199	138	11.5	
2016	9773	834	8.5	8614	693	8.0	1159	141	12.2	
2017	9472	956	10.1	8326	788	9.5	1146	168	14.7	
2018	9409	1092	11.6	8299	915	11.0	1110	177	15.9	
2019	9467	995	10.5	8335	849	10.2	1132	146	12.9	
2020	9643	1021	10.6	8449	858	10.2	1194	163	13.7	
Total	126742	13359	10.5	111313	11269	10.1	15429	2090	13.5	

Source: BCR linked to IMA-AIM data

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Table 2 –Patients receiving chemotherapy in the last 14 days of life by tumour type (all patients 2006-2020, maximum 3 years of follow-up)

	Total	n with chemotherapy
	N	n %
Acute	124900	12768 10.2
Oesophagus	7435	589 7.9
• Stomach	9875	577 5.8
Liver, primary	6657	374 5.6
Gallbladder and biliary Tract	3567	175 4.9
• Pancreas	16271	1512 9.3
Lung, bronchus and trachea	68069	7941 11.7
Pleura	2523	116 4.6
Brain	6826	262 3.8
Acute myeloid leukaemia	3677	1222 33.2
Chronic	17141	2324 13.6
Head and Neck	7238	835 11.5
Small Intestine	1011	50 5.0
Nasal cavities and sinuses	535	34 6.4
Ovary and uterine adnexa	4269	495 11.6
Multiple Myeloma	3056	544 17.8
Acute lymphatic leukaemia	138	49 35.5
Chronic myeloid leukaemia	894	317 35.5
Total	142041	15092 10.6

Source: BCR linked to IMA-AIM data

Table 3 -Patients who received chemotherapy in the last 14 days of life - by region (year of death=2020, incidence years 2018-2020 included)

	All Tumours			Acute T	Acute Tumours			Chronic Tumours		
	Total	N with chemotherapy in the last 14 days of life		Total	N with chemotherapy in the last 14 days of life		Total	N with chemotherapy in the last 14 days of life		
	N	n	%	N	n	%	N	n	%	
Brussels	721	80	11.1	623	74	11.9	98	6	6.1	
Flanders	5537	525	9.5	4896	443	9	641	82	12.8	
Wallonia	3385	416	12.3	2930	341	11.6	455	75	16.5	
Total	9643	1021	10.6	8449	858	10.2	1194	163	13.7	

Source: BCR linked to IMA-AIM data

#### **Key points**

- The administration of chemotherapy during the last days of life of patients dying from cancer is an indicator of the aggressiveness of care.
- Belgian data demonstrate that at least one out of ten cancer patients received chemotherapy in the last 14 days of life (average of 10.5% of the cancer patients who died in the period between 2008 and 2020).
- Substantial variation in the administration of chemotherapy in the last 14 days of life is observed between different tumour types, with generally higher percentages in patients with haematological tumour types in comparison to other tumour types.

#### References

- 1. Gielen B, De Gendt C, De Schutter H, Henin E, Ceuppens A, Peltier A, et al. Hospitalisaties bij het levenseinde van kankerpatiënten. 2013.
- 2. De Angelis R, Francisci S, Baili P, Marchesi F, Roazzi P, Belot A, et al. The EUROCARE-4 database on cancer survival in Europe: data standardisation, quality control and methods of statistical analysis. Eur J Cancer. 2009;45(6):909-30.
- 3. Sant M, Allemani C, Santaquilani M, Knijn A, Marchesi F, Capocaccia R. EUROCARE-4. Survival of cancer patients diagnosed in 1995-1999. Results and commentary. Eur J Cancer. 2009;45(6):931-91.
- Rossi S, Baili P, Capocaccia R, Caldora M, Carrani E, Minicozzi P, et al. The EUROCARE-5 study on cancer survival in Europe 1999–2007: Database, quality checks and statistical analysis methods. Eur J Cancer. 2015;Oct 51(15):2104-19.