ABSTRACT

Objectives The objective of this study was to investigate the use of, and predictors for, pharmaceutical anticancer treatment (PACT) towards the end of a patient’s life in a country with a public healthcare system.

Design Retrospective registry study.

Setting Secondary care in Norway.

Participants All Norwegian patients with cancer (International Classification of Diseases tenth revision (ICD-10) codes C00–99, D00–09, D37–48) in contact with a somatic hospital in Norway between 2009 and 2017 (N=420 655). Analyses were performed on a subsample of decedents with follow-back time of more than 1 year (2013–2017, N=52 496).

Interventions N/A.

Primary and secondary outcome measures Proportion of patients receiving PACT during the last year and month of life. We calculated CIs with block bootstrapping, while predictors of PACT were estimated with logistic regression.

Results 24.0% (95% CI 23.4% to 24.6%) of the patients received PACT during the last year of life and 3.2% (95% CI 3.0% to 3.5%) during their final month. The proportion during the last month was highest for multiple myeloma (12.7%) and breast cancer (6.5%) and lowest for urinary tract (1.1%) and prostate and kidney cancer (1.4%). Patients living in northern (OR 0.80, 95% CI 0.68 to 0.94) and western (OR 0.85, 95% CI 0.75 to 0.96) Norway had lower odds of PACT during the last month, while patients with myeloma (OR 3.0, 95% CI 2.5 to 3.7) and breast (OR 1.4, 95% CI 1.1 to 1.6) had higher odds. Kidney cancer (OR 0.25, 95% CI 0.2 to 0.4), urinary tract (OR 0.38, 95% CI 0.3 to 0.5) and prostate cancer (OR 0.4, 95% CI 0.3 to 0.5) were associated with lower probability of receiving PACT during the last month.

Conclusions The proportion of patients receiving PACT in Norway is lower than in several other industrialised countries. Age, type of cancer and area of living are significant determinants of variation in PACT.

INTRODUCTION

A fundamental problem for clinicians who consider prescribing end-of-life pharmaceutical anticancer treatment (PACT) is the assessment of the remaining lifetime of an individual patient and the expected patient benefit from the treatment. Whether oncologists should prescribe PACT near the end of life raises controversial ethical issues concerning the potential prolongation of life, quality of life for patient and their families, and the use of scarce healthcare resources. When such treatment is given near the end of life—the terminal phase, it may offer little potential to prolong life and is sometimes called futile—‘serving no useful purpose, completely ineffective’. The definition of the terminal phase has varied over time. Initially, terminal phase was defined as the time in which palliative care should be applied instead of active therapies. More recently it has become clear that palliative care should start earlier when active therapies are still in use. In this study, we explore those receiving PACT who are not in the terminal phase according to the first definition. PACT may be associated with extensive adverse effects, and may impact the patient’s ability to engage in meaningful life and prepare for death. It has been suggested that too much PACT in the last month of life may shorten life and, if side effects are not diagnosed and treated, reduce quality of life. Still, the treatment requires funds and may displace beneficial treatment for others. In
Norway, health authorities only fund new treatments that are cost-effective according to national guidelines and prioritising processes because budgets are not large enough to cover all treatments even if they are effective.8

Previous studies indicate that up to 38% of patients with cancer receive chemotherapy or other life-sustaining treatments during their last month of life.9 However, end-of-life treatment decisions may differ across countries or within jurisdictions.9 In some healthcare systems, oncologists have an incentive to prescribe PACT because their remuneration depends on it.10 Such systems typically also impose a cost on patients in terms of copayments (Norway has copayment on certain types of treatments, but a low total annual maximum of approximately US$270 in 2020). It is, therefore, unclear which system results in the highest end-of-life PACT rates. The evidence of factors influencing end-of-life care in European countries is also lacking. In a 2014 systematic review of end-of-life studies in cancer care, Langton et al9 found 15 studies that examined quality indicators (including use of chemotherapy) for end-of-life care, of which none were from Europe. More recently, however, such studies have been published for Denmark and France.11,12 The proportion receiving chemotherapy during the last 14 days of life was 4.2% in Denmark and 11.3% in France.11,12

Use of PACT towards the end of life presents a difficult medical decision and an important policy issue for patients and society. One approach to improve the quality in end-of-life care is to gain a better understanding of current patterns of end-of-life treatment at a national level in a public healthcare system where the overall goal is to offer patients equitable access to care. We used comprehensive individual patient-level nationwide data from the Norwegian Patient Registry13 to answer two research questions: First, what proportion of patients receive PACT during their last year and months of life in Norway? Second, to what extent are treatment decisions influenced by patients’ age, gender, type of cancer or geographical factors?

METHODS
We performed a retrospective cohort study using data from the Norwegian Patient Registry (delivered August 2018). Each episode of care (inpatient stay, day care or outpatient visit) in the Norwegian Patient Registry has a main International Classification of Diseases Tenth Revision (ICD-10) diagnostic code, possibly supplementary diagnostic codes, and a unique patient identifier that allows patients to be followed over time. Diagnostic codes in the Norwegian Patient Registry have proved valid when compared with the Cancer Registry of Norway.14 Each episode of care also had a diagnosis-related groups (DRG) code and an Anatomical Therapeutic Chemical (ATC) code in cases where PACT was administered. Additionally, the dataset encompassed the following variables: unique patient ID, patient age and gender, county of residence (19 counties), year/month of episode, medical and surgical procedure codes, code for infusion of PACT, days until death and level of care (hospital outpatient/day care and inpatient).

We included all episodes of care with ICD-10 codes C00–99, D00–09, D37–48. Norway has a population of approximately 5.4 million, and in 2017, a total of 33 564 new cancer cases were reported and 273 741 Norwegians were alive after having received a cancer diagnosis.15 Cancer treatment in hospice and private hospitals is negligible, thus, the Norwegian Patient Registry includes virtually all Norwegian patients with cancer. Our dataset encompassed 7 423 828 episodes for 420 655 patients. Our data included all patients with cancer who had been in contact with Norwegian hospitals (outpatient or inpatient) between 2009 and 2017 among whom 128 413 were reported dead by the end of 2017. All analyses were performed on a subset of the data (a 5 year cohort of patients who died during 2013–2017 with at least 1-year follow-back data, N=52 496) as patient administered treatment (oral and subcutaneous) were not identifiable in the registry before 2013 (see figure 1). Patient characteristics for the complete sample for the period 2013–2017 and the subsample used for analyses are presented in online supplemental table S1. The Norwegian Patient Registry has systems for data cleaning and continually check for correctness. We have further checked for correctness and consistency without detecting need for further cleaning.

Classification of cancer diagnosis
Patients with more than one cancer diagnosis were classified into mutually exclusive cancer diagnoses based on their main and supplementary diagnosis (online supplemental table S1). Patients with multiple cancer diagnoses were assigned to a single diagnosis based on their most frequently listed cancer diagnosis type. Classification of cancer diagnosis was based on data from 2009 to 2017.

Pharmaceutical anticancer treatment
PACT included cytostatic agents, cytotoxic agents, targeted therapies and immunotherapies. We classified PACT as intravenous, subcutaneous or oral by means of ATC codes and the DRG system. To calculate the proportion of patient who received PACT 1 year before death, patients with follow-back time less than 1 year from first episode of cancer until death were excluded from the analyses. We ran additional analyses to investigate the effect of limiting our study population to patients with at least 1-year follow-back.

Statistical analysis
For each patient, we registered whether he or she received PACT during the following periods prior to death: 12, 9, 6 and 3 months, and 1, 2, 3, 4, 5, 6, 7 and 8 weeks. Proportion of patients receiving PACT were defined as $\frac{PACT \times n}{N}$.
where \( PACT_t \) is the number of patients who received PACT during time period \( t \) \((t = \text{time period prior to death})\) and \( N \) is the number of patients with more than 12 months follow-back time before death. In other words, the numbers present aggregates over time periods, not treatment at a point in time.

We used block bootstrapping to estimate the SE of the proportion of patients who received PACT during time period \( t \).\(^\text{16}\) To reproduce the dependence structure of the observed data in the resampled data, we created blocks of consecutive data defined as each individual patient’s treatment course (last year before death). We estimated logistic regression models to identify predictors of receiving PACT during the last month of life. The following variables were included in the multivariable regression model: year of death, patient’s age at death, region of hospital, gender and type of cancer. The variables were chosen because they were expected to influence the use of PACT and due to availability in the obtained data.

All analyses were performed using STATA software V.14 (StataCorp).

**Patient and public involvement**

No patient involved.

**RESULTS**

**Use of end-of-life PACT**

A total of 301,611 patients with cancer received care in a hospital during the period 2013–2017. Of these patients, 52,496 patients (17.4%) were reported dead by the end of 2017 and had at least 1-year follow-back. Of this group, 12,604 (24.0%, 95% CI 23.4% to 24.6%) received PACT during the last year of life (Figure 2). The rates of PACT 1 year prior to death were highest for pancreatic cancer (60.7%, 95% CI 58.0% to 63.5%), multiple myeloma (55.0%, 95% CI 50.2% to 55.8) and lung cancer (45.7%, 95% CI 44.4% to 47.1%). Kidney cancer (11.7%, 95% CI 9.6% to 13.7%), urinary tract (12.8%, 95% CI 11.6% to 14.0%) and leukaemia (14.4%, 95% CI 3.1% to 15.8%) had low PACT rates during last year of life. In total 1691 (3.2%, 95% CI 3.0% to 3.5%) received PACT at least once during the last month before death, and patients with multiple myeloma (12.7%, 95% CI 10.9% to 14.5%), breast (6.5%, 95% CI 5.7% to 7.3%) and mouth/pharynx cancer (6.0%, 95% CI 4.3% to 7.7%) had the highest rates. Among cancers with low PACT rates during the last month before death were urinary tract (1.1%, 95% CI 0.7% to 1.5%), kidney cancer (1.4%, 95% CI 0.7% to 2.0%) and prostate cancer (1.4%, 95% CI 1.1% to 1.6%).

**Predictors of variation in treatment decisions**

The odds for receiving PACT during the last month of life were highest for patients aged 40–59 years and lowest for those aged 80+ (Table 1). Adjusted for the included covariates, patients living in Northern (OR 0.80, 95% CI 0.68 to 0.94) or Western Norway (OR 0.85, 95% CI 0.75 to 0.96) had lower odds of receiving PACT within the last month of life compared with those living in the South-East region. Also, patients diagnosed with multiple myeloma (OR 3.03, 95% CI 2.48 to 3.72) and breast cancer (OR 1.36, 95% CI 1.13 to 1.63) had higher odds of receiving PACT during the last month of life (compared with lung cancer). Kidney cancer (OR 0.25, 95% CI 0.15 to 0.43), urinary tract (OR 0.38, 95% CI 0.27 to 0.53) and prostate cancer (OR 0.59, 95% CI 0.31 to 0.49) had significantly lower odds. There was no clear trend towards
Figure 2  Proportion of patients with cancer receiving pharmaceutical anticancer treatment during last weeks or months of life, 2013–2017.
higher or lower provision of end-of-life PACT during the observation period and we found no differences between genders.

Additionally, we ran analyses for women and men separately and analyses excluding cancers that only occur for one gender (breast, uterine, prostate), but we were not able to find any differences between genders. Also, we tested for several plausible interactions between gender, age, health region and cancer type. In total, we tested for 36 interactions. Six were borderline significant, but none of the interactions were meaningful (online supplemental table S2). Finally, we investigated the impact of restricting our population to patients with at least 1-month follow-back period (compared with 1 year in our base-case analysis). The predictors were not substantially affected by the change in patient population (online supplemental table S3).

**DISCUSSION**

One in four Norwegian patients with cancer receive PACT during their last year of life, while 3% are treated within the last month. Patients who are elderly or residents of Northern or Western Norway (regions with a lower population density compared with the south-east region) are less likely to receive PACT during the last month of life. The proportion receiving PACT end-of-life in Norway are relatively low compared with other European countries with similar healthcare systems, indicating that medical culture and patient preference impact choice of treatment end of life.

Previous studies indicate high levels of PACT use near the end of life with rates up to 38% during last month in countries in North America, Asia and Europe.

Our findings of generally low PACT rates in Norway align well with results reported previously and the relative low rates found in our neighbouring country Denmark which has a similar healthcare system. Having a public healthcare system, where oncologist have no financial incentive to prescribe, may explain the lower rates in Norway. Additionally, due to strict priority setting for pharmaceuticals, not all new treatment options are available to Norwegian patients. Other factors may also explain the lower rates, such as the role of palliative care teams or culture and attitude towards end-of-life treatment in the clinical environments. Even if the negligible patient copayments in Norway should favour use of PACT, the inhibitive factors dominate the final decisions. Our findings support previous research that factors such as age, tumour site and region of residence influence the level of PACT near death.

The main strength of our study lies in the use of nationwide registry data for a period of 5 years with a range of key covariates. Almost all patients with cancer in Norway receive medical care in public hospitals, so our data cover virtually the entire Norwegian population (no selection bias). Our data also include a range of covariates allowing adjustment for patient characteristics, year of treatment, type of cancer and patient’s place of living. Another strength is that we capture both hospital-administered and patient-administered PACT. In recent years, subcutaneous and oral PACT have become important modalities, making it essential to include these treatments when studying end-of-life care.

However, the study also has several limitations. First, we do not have information on disease stage, the treating clinician, comorbidities or functional status. Complete

### Table 1  Multivariable logistic regression analysis of the odds that patients received pharmaceutical anticancer treatment last month before death

| Year of death          | OR (95% CI) | P value |
|------------------------|-------------|---------|
| Reference: 2013        | 0.99 (0.96 to 1.03) | 0.74    |
| Age                    |             |         |
| 60–69 years (reference) | 1.00        | NA      |
| 0–9 years              | 0.85 (0.95 to 1.52) | 0.82    |
| 10–19 years vs 60–69 years | 0.74 (0.54 to 1.03) | 0.62    |
| 20–29 years vs 60–69 years | 0.66 (0.29 to 1.55) | 0.36    |
| 30–39 years vs 60–69 years | 1.08 (0.73 to 1.61) | 0.70    |
| 40–49 years vs 60–69 years | 1.46 (1.19 to 1.78) | <0.001  |
| 50–59 years vs 60–69 years | 1.33 (1.16 to 1.53) | <0.001  |
| 70–79 years vs 60–69 years | 0.62 (0.55 to 0.69) | <0.001  |
| 80–89 years vs 60–69 years | 0.14 (0.12 to 0.17) | <0.001  |
| 90–99 years vs 60–69 years | 0.03 (0.14 to 0.53) | <0.001  |
| Hospital affiliation (regional health authority) | | |
| South-Eastern (reference) | 1.00        | NA      |
| Central                | 1.02 (0.90 to 1.17) | 0.74    |
| Western                | 0.85 (0.75 to 0.96) | 0.01    |
| Northern               | 0.80 (0.68 to 0.94) | 0.01    |
| Gender                 |             |         |
| Women                  | 0.97 (0.87 to 1.07) | 0.52    |
| Type of cancer          |             |         |
| Lung cancer (reference) | 1.00        | NA      |
| Mouth, pharynx          | 1.14 (0.84 to 1.55) | 0.41    |
| Colon, rectum, rectosigmoid | 0.84 (0.71 to 0.99) | 0.05    |
| Pancreatic              | 1.21 (0.94 to 1.54) | 0.14    |
| Melanoma                | 0.77 (0.58 to 1.03) | 0.08    |
| Breast                 | 1.36 (1.13 to 1.63) | <0.001  |
| Cervix uteri           | 0.69 (0.42 to 1.13) | 0.14    |
| Prostate               | 0.39 (0.31 to 0.49) | <0.001  |
| Kidney (excl. renal pelvis) | 0.25 (0.15 to 0.43) | <0.001  |
| Urinary tract          | 0.38 (0.27 to 0.53) | <0.001  |
| Non-Hodgkin’s lymphoma | 1.06 (0.81 to 1.38) | 0.69    |
| Leukaemia              | 0.72 (0.57 to 0.93) | 0.01    |
| Multiple myeloma       | 3.03 (2.48 to 3.72) | <0.001  |
| Residual group*        | 0.66 (0.57 to 0.76) | <0.001  |

**Constant: 0.10 (0.08–0.11).**

*Residual group includes all cancers not presented above. NA, not applicable; OR, odds ratio.
with similar systems have higher PACT rates during the last months of life, which indicate that not only financial incentives, but also medical culture and patient preferences may impact choice of treatment. Information on PACT rates may be useful for clinicians in order to achieve optimal end-of-life care.

Contributors CB: Data collection, study design, analysis, interpretation of results, preparation of the manuscript and approval of the manuscript. SK: Study design, interpretation of results, preparation of the manuscript and approval of the manuscript. EMS: Data collection, interpretation of results, preparation of the manuscript and approval of the manuscript. HOM: Interpretation of results, preparation of the manuscript and approval of the manuscript. ISK: Data collection, study design, interpretation of results, preparation of the manuscript and approval of the manuscript.

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Competing interests CB, EMS and ISK are affiliated with Oslo Economics and have all completed consultancy assignments for several pharmaceutical companies in recent years. None of the projects have been related to the work presented in this paper.

Patient consent for publication Not applicable.

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Data availability statement Data are available upon reasonable request. Data may be obtained from a third party and are not publicly available. The study uses registry data from the Norwegian Patient Registry. Individual patient level data can be obtained by application to the registry. Authors are willing to share access to the STATA programming code (do-file). Individual patient level data are protected by law, and reviewers have to come to Oslo to have access to data.

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ORCID iD Christoffer Bugge http://orcid.org/0000-0002-0766-2870

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