The Off-Label Use of Antineoplastics in Oncology Is Limited But Has Notable Scientific Support in a University Hospital Setting

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Purpose: The off-label (OL) use of antineoplastic drugs for the treatment of various types of tumors in patients of different disease stages is becoming a common occurrence. The objective of this study was to analyze these patterns by quantification and characterization of the OL use of antineoplastic drugs and their level of scientific evidence in a medium/high-complexity Spanish general university hospital.

Method: All oncology patients who underwent OL treatment with one or several antineoplastics during the 10 years from 2002 to 2012 were retrospectively selected. The use of these drugs was considered OL if they were used for indications, stages, lines of treatment, or chemotherapy schemes not reflected in the summary of product characteristics published by the European Medicines Agency at the time of prescription. To calculate the prevalence of patients who received one or more OL treatments during the study period, all patients whose primary or secondary diagnosis had been coded with the diagnoses included in the study were selected through the minimum basic data set (MBDS). This database was cross-referenced with that of the Farmatools® program (Dominion®), which collects information on all patients receiving chemotherapy to obtain the total number of patients who received chemotherapy in the hospital during this period.

Results: In total, 684 patients and 866 OL treatments were included. The prevalence of patients undergoing OL treatment with one or several antineoplastics during the 10 years from 2002 to 2012 were retrospectively selected. The use of these drugs was considered OL if they were used for indications, stages, lines of treatment, or chemotherapy schemes not reflected in the summary of product characteristics published by the European Medicines Agency at the time of prescription. To calculate the prevalence of patients who received one or more OL treatments during the study period, all patients whose primary or secondary diagnosis had been coded with the diagnoses included in the study were selected through the minimum basic data set (MBDS). This database was cross-referenced with that of the Farmatools® program (Dominion®), which collects information on all patients receiving chemotherapy to obtain the total number of patients who received chemotherapy in the hospital during this period.

Conclusion: The OL use of antineoplastics in oncology patients is limited; their use is mainly focused in a small group of tumors and at advanced stages of disease. OL use of antineoplastics occurs under palliative therapeutic strategies with a limited number of
drugs, preferably off-patent drugs. In addition, these OL treatments have high levels of clinical evidence.

**Keywords:** off-label use, antineoplastic agents, evidence, prevalence, oncology

**INTRODUCTION**

Cancer is one of the major health problems in countries with developed healthcare systems and is currently the leading cause of death worldwide. Advances and improvements in diagnoses and therapies are contributing to the control and reduction of the death rate from this disease in the United States (USA) and Europe (de Angelis et al., 2014; Ferlay et al., 2015; Organización Mundial de la Salud, 2018).

Increasing patient survival and quality of life in turn increases the likelihood that patients will receive additional lines of treatment (Hillner and Smith, 2009; Schickedanz, 2010). However, the guidelines and lines of chemotherapy approved by the regulatory agencies are not sufficient to treat the different stages and clinical forms of disease among affected patients. This limitation of the approved therapeutic offerings causes physicians to resort to the use of antineoplastic drugs for conditions that are different from those specified in the product’s technical sheet, which is known as off-label (OL) use.

In clinical practice, OL use in the treatment of various types of tumors and progressing stages of disease is a frequent and relevant reality for patients, prescribing physicians, and the economic cost of the healthcare system. Despite its importance, to date, analyses of the efficacy and efficiency of the OL use of drugs in oncology have been limited. Most of the data available are estimates based on a survey conducted in 1991 by the General Accounting Office (GAO) (Laetz and Silberman, 1991; United States General Accounting Office, 1991) of the USA among oncologist members of the American Society of Clinical Oncology (ASCO). According to the results, more than half of the patients (56%) underwent OL treatment with at least one drug, and 33% of all drugs administered were under conditions other than those specified on the data sheet. In 2007, the ASCO and the European Society for Medical Oncology (ESMO) reported that approximately 50% of the use of antineoplastics was for indications that were not reflected on the data sheet (American Society of Clinical Oncology, 2006; Casali and Executive Committee of ESMO, 2007). According to the National Comprehensive Cancer Network (NCCN®) estimate, in the USA, 50%–75% of all uses of antineoplastics in oncology are OL (Benson and Brown, 2008; Cohen et al., 2009).

Several descriptive studies of OL use in oncology have been performed, such as a study conducted by Levêque et al. (2005) in France, which estimated an annual prevalence of 6.7% by analyzing OL prescriptions of 10 antineoplastic drugs for 10 tumor types. In Switzerland, Joerger et al. (2014) determined that 27.2% of antineoplastic administrations were OL, and a study by Mellor et al. (2009) in Australia reported that 35% of the prescriptions were OL and that the prevalence increased from 22% in 2001 to 35% in 2008.

The present study analyzes the OL use of any antineoplastic agent for patients treated over 10 years. We attempted to minimize the transitory impact of periods related to the upcoming approval of a specific drug by analyzing a long period of time. This approach also allowed us to evaluate OL use in all tumors without limiting the analysis to neoplasms. The study was conducted in an oncology service in the setting of a medium/high-complexity general university hospital. The pattern of OL use was analyzed by quantification and characterization according to the tumor type and stage and the progressive clinical phase of the patient. We also investigated the level of scientific evidence that supports OL use in oncology.

**MATERIALS AND METHODS**

This observational and retrospective study was conducted at the Prince of Asturias University Hospital located in Alcalá de Henares (Madrid, Spain) with 500 beds, specialized diagnosis and treatment units, and more than 10 highly differentiated clinical specialties, which offers healthcare coverage to a population of approximately 400,000 inhabitants. This study analyzed the period between January 1, 2002, and December 31, 2012.

All patients with OL use of one or several antineoplastic drugs, either as a single agent or in combination, during the study period were included. The use of these drugs was considered OL if they were used in indications, stages, lines of treatment, or chemotherapy schemes that were not reflected in the summary of product characteristics published by the European Medicines Agency at the time of prescription. We excluded patients who received treatments with experimental drugs or drugs unauthorized in Spain and those uses that were considered OL due to their use in posologies or routes of administration other than those authorized. Each patient was counted only once; in the cases of patients with more than one OL use of a drug, each treatment was counted separately.

The clinical histories and the dispensing data of the selected patients were reviewed. Information on each patient’s age, sex, Eastern Cooperative Oncology Group (ECOG) status, diagnosis, tumor stage, metastasis, line of treatment, OL drug, type of OL use (monotherapy or combination), patent status and route of administration of the drug, and whether the treatment had curative or palliative intent was collected. To evaluate the status of the patent, we defined a drug “not protected by the patent” if it had expired before or during the study period. We examined the level of evidence of the chemotherapy schemes with OL use (CSOLs) according to the NCCN v1.2015 and Micromedex® 2015 compendia.

**Abbreviations:** OL, off-label; SmPC, summary of product characteristics; CSOLs, chemotherapy schemes with off-label use; MBDS, minimum basic data set; AEMPS, Spanish Agency for Medicines and Health Products.
The diagnoses were classified according to the ICD-9.10.MC (Ministerio de Sanidad and Servicios Sociales e Igualdad, 2014) classification and were grouped into 14 primary tumor locations or types. A miscellaneous group was created for OL use in rare tumors and those with an absolute frequency lower than 5. A “patent-protected drug” was defined as a drug that, at the end of the study (December 31, 2012), was still under a valid patent.

The types of OL use that were classified as monotherapy were additionally classified as “not indicated for the tumor” if the drug did not have any indication for that tumor type or in any of the stages of the tumor or as “indicated in combination” if the drug had an indication for a certain tumor type in combination with other agents. Likewise, for OL uses classified as combination therapy, those “with other indications” were distinguished if they were indicated for that tumor type as monotherapy in another line of treatment or in combination with other agents and as “without indication” when at least one drug in the combination had no indication for that type of tumor.

To calculate the prevalence of patients who received one or more OL treatments during the study period, all patients whose primary or secondary diagnosis had been coded with the diagnoses included in the study were selected through the minimum basic data set. This database was cross-referenced with that of the Farmatools® program (Dominion®), which collects information on all patients receiving chemotherapy to obtain the total number of patients who received chemotherapy in the hospital during this period.

**Statistical Analysis**

A descriptive statistical analysis of the data was conducted. Conformity tests of the categorical variables related to the characteristics of the patients and the OL use of antineoplastic agents (tumor location, drug used, type of OL, intention of treatment, and patent status of the drug) were performed by comparing the proportion of the sample using a chi-square test. A value of p ≤ 0.05 was considered significant. All statistical analyses were performed with SPSS version 18.0 (SPSS Inc., Chicago, IL, USA).

![Inclusion criteria: 794 patients and 980 OL treatments. Exclusion criteria: 110 patients and 114 OL treatments eliminated.](image)

**RESULTS**

**Demographic and Clinical Characteristics of the Sample With Oncological Disease**

This study included 794 patients with oncological disease and 980 OL treatments during the 10 years of the study. After the exclusion criteria were applied to this sample, 684 patients and 866 OL treatments were included in the analysis (Figure 1).

The study showed a significantly higher amount of OL use in women with an average age of less than 65 years and good general condition (p < 0.001). OL use was predominant in patients with stage IV disease and the presence of metastasis (p < 0.001) (Table 1).

**OL treatment Is Preferably Used in Oncology for a Small Group of Tumor Types and With Limited Use of Agents**

Among all cancer patients treated with chemotherapy (n = 11,385), the prevalence of those receiving OL use of antineoplastic drugs was 6% (684/11,385).

OL treatments were used mainly in four locations or tumor types: breast cancer (25.2%), gynecological tumors (16.1%), lung cancer (11.8%), and gastric cancer (10.2%). The OL use in these tumor types was significantly greater than that in all other tumor types (63.3% vs 36.7%, respectively; p < 0.001) (Table 2).

We found an OL use of 27 antineoplastic agents, and the most commonly used was paclitaxel (19.2%), followed by gemcitabine (10.9%), carboplatin (9.6%), vinorelbine (8.8%), and capecitabine (8.7%). The OL use of these five drugs was significantly more frequent than that of the other 22 drugs (57.2% vs 42.8%, respectively; p < 0.001) and was essentially focused on the four...
most frequent tumor types. The OL use of drugs as palliative treatment predominated over their use as curative treatment (79% vs 21%, p < 0.001) (Table 3).

The wide OL use of paclitaxel was due to its use in monotherapy in nonsmall cell lung cancer; cancers of the cervix, bladder, and esophagus; and head and neck tumors. The OL use of gemcitabine was notable as a single agent in breast and ovarian tumors, for which it has an indication when combined with other agents. OL use of carboplatin occurred for breast cancer, both as monotherapy and combined with other agents. The OL use of vinorelbine was due to its use as a single agent for the treatment of gynecological and prostate tumors; the OL use of capecitabine mainly included its use in a neoadjuvant form in rectal cancer and as an adjuvant in high-risk stage II colon cancer and with palliative intent in pancreatic cancer.

Most of the drugs used were not protected by patents (73% vs 27%; p < 0.001). The most commonly used route of administration was intravenous. Regarding the line of treatment for OL use of antineoplastics, no significant differences were observed between the first four lines used. However, 50.8% of OL uses were as second- and third-line treatments.

The OL treatments were mainly used in monotherapy (70.3% vs 29.7%; p < 0.001), with a notable use of drugs without any type of indication approved for the tumor on which they were used.

### Table 1 | General characteristics of patients who received off-label treatments.

| Patient characteristics | N = 866 | % |
|-------------------------|---------|---|
| Gender* | | |
| Male | 370 | 42.7 |
| Female | 496 | 57.3 |
| Age* | | |
| Mean age of patients (years) | 58.21 ± 11.57 |
| Age <65 | 601 | 69.4 |
| Age ≥65 | 265 | 30.6 |
| Performance status (ECOG)* | | |
| 0 | 402 | 46.4 |
| 1 | 308 | 35.6 |
| 2 | 156 | 18 |
| Presence of metastatic disease* | | |
| Yes | 684 | 79 |
| No | 182 | 21 |
| Disease stage* | | |
| I | 39 | 4.5 |
| II | 59 | 6.8 |
| III | 84 | 9.7 |
| IV | 684 | 79 |
| Previous antineoplastic treatment line | | |
| 0-3 | 704 | 81.3 |
| 3-6 | 142 | 16.4 |
| 6-9 | 19 | 2.2 |
| >9 | 1 | 0.1 |

*p < 0.001.

### Table 2 | Primary location of tumors involved in off-label use of antineoplastic drugs.

| Tumor location | Treatment (n = 866) | % |
|----------------|---------------------|---|
| Breast | 218 | 25.2 |
| Gynecological | 139 | 16.1 |
| Lung | 102 | 11.8 |
| Gastric | 88 | 10.2 |
| Head and neck | 69 | 8 |
| Biliopancreatic | 64 | 7.4 |
| Colorectal | 53 | 6.1 |
| Soft tissue sarcoma | 29 | 3.3 |
| Bladder-urothelial | 25 | 2.9 |
| Prostate and testicle | 24 | 2.8 |
| Melanoma | 22 | 2.5 |
| Esophagus | 15 | 1.7 |
| Miscellaneous | 12 | 1.4 |
| Liver | 6 | 0.7 |

### Table 3 | Characteristics of treatments with off-label use of antineoplastics.

| Drug | Treatment (n = 866) | % |
|------|---------------------|---|
| Paclitaxel | 195 | 22.9 |
| Gemcitabine | 111 | 12.9 |
| Carboplatin | 98 | 11.6 |
| Vinorelbine | 89 | 10.3 |
| Capecitabine | 88 | 10.3 |
| Irinotecan | 77 | 8.8 |
| Trastuzumab | 74 | 8.5 |
| Oxaliplatin | 71 | 8.2 |
| Docetaxel | 54 | 6.3 |
| Bevacizumab | 36 | 4.2 |
| Liposomal doxorubicin | 9 | 1.1 |
| Cetuximab | 16 | 1.9 |
| Etoposide | 13 | 1.5 |
| Lapatinib | 11 | 1.3 |
| Cisplatin | 8 | 0.9 |
| Topotecan | 8 | 0.9 |
| IL-2 | 5 | 0.6 |
| Pemetrexed | 5 | 0.6 |
| Adriamycin | 4 | 0.5 |
| Albumin-paclitaxel | 4 | 0.5 |
| Pegylated liposomal doxorubicin | 3 | 0.4 |
| Nilotinib | 2 | 0.2 |
| Cyclophosphamide | 1 | 0.1 |
| Imatinib | 1 | 0.1 |
| Sorafenib | 1 | 0.1 |
| Sunitinib | 1 | 0.1 |

*Type of off-label use
- Monotherapy
- Not indicated for the tumor
  - Indicated in combination
  - Combinations
  - With other indications
  - Without indication

*Intent of off-label treatment
- Curative
- Palliative

*Patent status
- Patent protected
- Off-patent

*Line of off-label treatment
- 1st line
- 2nd line
- 3rd line
- 4th line
- Successive lines
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used (52.9%). Table 4 indicates the type of OL use for each tumor site.

**OL Treatments Are Mostly Supported by Clinical Evidence**

The analysis of the degree of evidence of the 114 CSOLs included in the study showed the following: the NCCN® indicated a 2A level of evidence for 56.1% of the schemes, while the Micromedex® indicated that practically the same percentage (55.3%) of schemes had a 2B level of evidence. The same analysis was performed on the 866 OL treatments included in the study; in this case, 64% of the treatments used had a 2A level of evidence according to the NCCN® and a 2B level of evidence according to Micromedex®.

The percentages of CSOLs that were not reflected in either of the two compendia were 22.8% in NCCN® and 34.2% in Micromedex® however, these CSOLs accounted for only 11.6% and 19.4% of 866 OL treatments analyzed, respectively (Table 5).

**DISCUSSION**

The results of this study showed that in the setting of a general university hospital, the OL use of antineoplastic drugs in patients with tumor pathology is limited and is focused on a small group of tumors (breast, gynecological, lung, and gastric) in patients under 65 years old with good general condition and advanced-disease stages. The OL use of antineoplastic drugs mainly occurs in a palliative therapeutic strategy with a limited number of drugs, preferably off-patent drugs. In addition, these OL treatments attain high levels of clinical evidence.

**TABLE 4 | Type of off-label use for each location/tumor type.**

| Location/Tumor type | Type of Label Treatment | Monotherapy | Combination |
|---------------------|-------------------------|-------------|-------------|
|                     |                         | Not indicated for the tumor | Patients (n) | (%) | Indicated in combination | Patients (n) | (%) | Without indication | Patients (n) | (%) | With other indications | Patients (n) | (%) |
| Biliopancreatic      |                         | Pancreas     | 46 | 86.8 | 0 | 0 | 7 | 13.2 | 0 | 0 |
|                     |                         | Biliary      | 10 | 90.9 | 0 | 0 | 1 | 9.1 | 0 | 0 |
|                     |                         | Bladder-urothelial | 24 | 96 | 0 | 0 | 1 | 4 | 0 | 0 |
|                     |                         | Breast       | 27 | 12.39 | 41 | 18.81 | 46 | 21.1 | 104 | 47.7 |
| Colorectal           |                         | Colon        | 17 | 77.3 | 0 | 0 | 5 | 22.7 | 0 | 0 |
|                     |                         | Rectal       | 30 | 96.8 | 0 | 0 | 1 | 3.2 | 0 | 0 |
|                     |                         | Esophagus    | 15 | 100 | 0 | 0 | 0 | 0 | 0 | 0 |
|                     |                         | Gastric      | 70 | 79.6 | 17 | 19.3 | 1 | 1.1 | 0 | 0 |
| Gynecological        |                         | Cervix       | 22 | 66.7 | 0 | 0 | 11 | 33.3 | 0 | 0 |
|                     |                         | Endometrium  | 5 | 23.8 | 0 | 0 | 16 | 76.2 | 0 | 0 |
|                     |                         | Ovary        | 43 | 50.57 | 32 | 37.63 | 10 | 11.8 | 0 | 0 |
|                     |                         | Head and Neck| 37 | 73.9 | 14 | 20.3 | 18 | 26.1 | 0 | 0 |
|                     |                         | Liver        | 6 | 100 | 0 | 0 | 0 | 0 | 0 | 0 |
| Lung                |                         | Microcytic   | 28 | 87.5 | 4 | 12.5 | 0 | 0 | 0 | 0 |
|                     |                         | Not microcytic| 5 | 7.15 | 43 | 61.45 | 0 | 0 | 22 | 31.4 |
|                     |                         | Melanoma     | 17 | 77.3 | 0 | 0 | 5 | 22.7 | 0 | 0 |
| Miscellaneous       |                         | GIST (gastrointestinal stromal tumor) | 2 | 100 | 0 | 0 | 0 | 0 | 0 | 0 |
|                     |                         | Neuroendocrine | 0 | 0 | 0 | 0 | 2 | 100 | 0 | 0 |
|                     |                         | Osteosarcoma  | 0 | 0 | 0 | 0 | 1 | 100 | 0 | 0 |
|                     |                         | Kidney       | 1 | 100 | 0 | 0 | 0 | 0 | 0 | 0 |
|                     |                         | Adrenal      | 1 | 100 | 0 | 0 | 0 | 0 | 0 | 0 |
|                     |                         | Thymoma      | 2 | 66.7 | 0 | 0 | 1 | 33.3 | 0 | 0 |
|                     |                         | TOD (tumor of unknown origin) | 2 | 100 | 0 | 0 | 0 | 0 | 0 | 0 |
| Prostate and Testicle|                         | Prostate     | 11 | 100 | 0 | 0 | 0 | 0 | 0 | 0 |
|                     |                         | Testicle     | 12 | 100 | 0 | 0 | 0 | 0 | 0 | 0 |
|                     |                         | Soft tissue sarcoma | 24 | 82.8 | 0 | 0 | 5 | 17.2 | 0 | 0 |
The OL use of antineoplastic drugs in oncology is a therapeutic practice that is used in healthcare in a highly variable manner according to the published studies. Such studies have specifically analyzed the prescriptions for patients with specific tumors without specifying whether they are different patients or successive lines used in the same patient. This fact can explain the variability observed in the reflected prevalence of OL use—between 6% and 35% (Leveque et al., 2005; Mellor et al., 2009; Joerger et al., 2014). Our results of the analysis of all patients who received OL antineoplastic treatment show that the frequency of use of this therapeutic strategy is limited and is focused on 6% of patients. These data contrast with those obtained in the GAO survey conducted on a sample of North American medical oncologists (Laetz and Silberman, 1991; United States General Accounting Office, 1991). These differences can be explained, in addition to the methodological reasons previously described, by other reasons, including accessibility and control of the OL use of antineoplastic drugs in the setting in which this study was conducted: a hospital of a national universal healthcare system for the population served by the center with free access by the ill. However, until 2009, the management of requests and approval for the OL use of antineoplastic drugs in our national healthcare system was centralized in a review committee external to the hospital; thus, this management strategy for OL drug use could have influenced the low comparative prevalence found in our study.

To assess the OL treatments used in routine clinical practice, the epidemiology of the tumor and advances in the clinical and therapeutic management of the disease must be taken into account (Hanahan and Weinberg, 2000; Kocs and Fendrick, 2003; Newell, 2005; Hillner and Smith, 2009; Schickedanz, 2010). Likewise, improvements in supportive care have also increased the number of candidates for additional chemotherapy lines (Kocs and Fendrick, 2003). Our study shows that the OL use of antineoplastic drugs is significantly greater in palliative oncological treatment strategies, in metastatic stages of the disease, and especially as second- and third-line treatments.

These findings are consistent with those obtained in the USA by the GAO (Laetz and Silberman, 1991; United States General Accounting Office, 1991; Eastman, 2005) on the OL use of antineoplastic agents and those of other studies with smaller sample sizes (Kocs and Fendrick, 2003; Peppercorn et al., 2008; Roila et al., 2009; Cioffi et al., 2012; Cohen et al., 2013; Joerger et al., 2014).

This work indicates that the OL use of antineoplastic drugs is mainly focused on breast cancer, gynecological tumors, lung cancer, and gastric cancer. These four neoplasms share the characteristics of high prevalence and/or mortality (Ferlay et al., 2013; de Angelis et al., 2014; Ferlay et al., 2015; Organización Mundial de la Salud_OMS, 2018; SEOM, 2019). In previous studies, such as that of Joerger et al. (2014), which included 10 tumor types, the OL use of antineoplastic was more prevalent for gastrointestinal, breast, lung, and gynecological tumors. In the study conducted by Leveque et al. (2005), of 10 tumor types, prostate, breast, bladder, and ovarian cancers had the most OL prescriptions.

Our findings indicate that the results obtained by other authors who performed selective analysis of some tumor types included those more prevalent in a general analysis of all tumors (Leveque et al., 2005; Mellor et al., 2009; Joerger et al., 2014).

In our study, OL use was mainly focused on the use of 5 of 27 drugs, including paclitaxel, gemcitabine, carboplatin, vinorelbine, and capecitabine. All five drugs are cytotoxic agents with recognized activity in different tumors, and their use is supported by more and less extensive studies and for advanced pathologies. We found that paclitaxel had the most OL use; in contrast, OL use of docetaxel was limited. These data could be explained by the introduction of generic drugs and the changes observed in toxicity patterns, which could have changed the attitudes toward medical prescription. This possibility could affect the discrepancy between the OL use of docetaxel observed in our study and that found by other authors (Leveque et al., 2005; Joerger et al., 2014). Joerger et al. (2014) examined the OL use of 10 antineoplastics and 437 prescriptions in their analysis and found that OL use was more frequent with paclitaxel, followed by carboplatin and docetaxel. In the study by Leveque et al. (2005), the drug most commonly prescribed for OL use was docetaxel, followed by oxaliplatin, fluorouracil, carboplatin, gemcitabine, paclitaxel, and irinotecan. The authors explained that the wide use of docetaxel was mainly due to its use in prostate cancer; in 2002, docetaxel still did not have an indication for this type of tumor in France.

The evaluation of our results on the OL use of antineoplastic in relation to other studies requires consideration of some relevant aspects of oncological treatment. Among them, the analyzed period is notable due to the advances made in the management of neoplasms and the scope of the study, which includes the healthcare system in which the study is conducted, the country, and the hospital level. From a general perspective, the distribution of the OL use of antineoplastic according to the type or tumor location reflected in our study shows patterns of indication and use similar to those described in two previous studies.

### TABLE 5 | Evidence for off-label treatment.

| NCCN® | Off-label chemotherapy regimens | % | Treatments n = 866 | % |
|-------|---------------------------------|---|--------------------|---|
| 1     | 6                               | 5.3 | 113                | 13 |
| 2A    | 64                              | 56.1| 554                | 64 |
| 2B    | 16                              | 14  | 104                | 12 |
| 3     | 2                               | 1.8 | 3                  | 0.3|
| N/A   | 26                              | 22.8| 92                 | 10.6|

| MICROMEDEX® | Off-label chemotherapy regimens | % | Treatments n = 866 | % |
|-------------|---------------------------------|---|--------------------|---|
| 1           | 1                               | 0.9 | 30                 | 3.5|
| 2A          | 8                               | 7   | 101                | 11.7|
| 2B          | 63                              | 55.3| 555                | 64.1|
| 3           | 3                               | 2.6 | 6                  | 0.7|
| N/A         | 39                              | 34.2| 174                | 20.1|

N/A (Not applicable): chemotherapy regimens for which evidence is not included in the compendium that was evaluated.
studies. It should be noted that these three studies were conducted in European healthcare settings.

Notably, the OL use of antineoplastic drugs is more common in monotherapy than in combination therapies, with a predominant use in groups of drugs without any authorized indication for the tumor for which they were used. This finding can be related to the palliative intent of treatment and the use of these drugs as second- and third-line treatments. In this type of clinical situation, an increase in patient survival is sought via good control of the disease, with quality of life prevailing over other factors. In addition, the previous lines of treatment will limit the available options according to the health results obtained with the use of these drugs and any toxicities developed during the treatment.

In our analysis, a majority of the drugs used for OL treatments were not protected by patents. This finding is consistent with that of other publications that establish that an increase in OL use may occur among antineoplastic drugs after the patent has expired. This loss of patent could lead to a lack of interest on the part of the pharmaceutical industry to obtain new indications (Radley et al., 2006; Casali and Executive Committee of ESMO, 2007; Rola et al., 2009; Gota and Patial, 2011; Lerose et al., 2012; Wittich et al., 2012; Conti et al., 2013). Likewise, other drugs are not authorized for all indications for which they could be effective, mainly due to the high economic cost and time required to obtain a new indication (Boos, 2003; Gupta and Nayak, 2014).

Currently, several authors have established that in clinical practice, a chemotherapy scheme is supported by evidence and therefore is “medically accepted” if it is considered category 1 or 2A by the NCCN compendium or class 1, 2A, or 2B by Micromedex® (American Society of Clinical Oncology, 2006; Benson and Brown, 2008; Mehr, 2012). Independent clinical investigations, despite being considered of lower scientific quality than clinical trials, generate experience and provide data on the efficacy and safety of OL use of antineoplastics (Gazarian et al., 2006; Casali and Executive Committee ESMO, 2007; Mellor et al., 2009; Feliu and Espinosa, 2013; Gota and Nayak, 2014).

Despite the debate that exists regarding the evidence supporting the OL use of antineoplastic drugs (Gazarian et al., 2006; Abernethy et al., 2009; Irwin et al., 2012; Pfister, 2012), our results indicate that most of these OL uses are performed with significant scientific support, such as that collected in the North American therapeutic compendia. This coincides with studies previously published by Gota and Patial (2011) and Mellor et al. (2012) that used the NCCN to assess the degree of evidence of the OL schemes analyzed. Regarding the OL use of antineoplastic drugs that were not included in these compendia, drugs or regimens used in marginal cases were included, in which the justification for their use is based on results published in scientific journals without having reached the levels of so-called recognized therapeutic clinical evidence. It should also be noted that the compendia used originated from the USA, while the usual clinical practice was performed in Europe, and the approved indications for each drug are different between the two continents.

Our analysis presents some important limitations such as a one-center study, and therefore, the results focus on the type of population that our hospital serves, and epidemiologically infrequent or “rare” tumors do not reach a relevant qualitative significance. Multicentric studies should be carried out to confirm our results and to analyze the use of OL antineoplastics in less frequent tumors.

**DATA AVAILABILITY STATEMENT**

All datasets generated for this study are included in the article/Supplementary Files.

**ETHICS STATEMENT**

The studies involving human participants were reviewed and approved by Comité Ético de Investigacion Clinica del Hospital Universitario Príncipe de Asturias. The ethics committee waived the requirement of written informed consent for participation.

**AUTHOR CONTRIBUTIONS**

MF designed the study, performed the experiments, analyzed the data, and wrote the manuscript. RV designed the study, performed the experiments, and wrote the manuscript. MY performed the experiments. FE performed the experiments. MA designed the study and wrote the manuscript.

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Conflict of Interest: The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

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