ANALYSIS OF SURVIVAL IN PATIENTS DIAGNOSED WITH METASTATIC

B. Isla-Tejera, J. Lopez Santamaria Donoso, P. Montejano Hervas, I. Reyes, A. Gago, S. De la Fuente

Department of Pharmacy, Reina Sofia University Hospital, Córdoba, Spain. University of Córdoba, Córdoba. Spain

BACKGROUND & OBJECTIVES

Following an observational retrospective design, data of patients who received at least one dose of eribulin from February 2014 until July 2016 were obtained from the computerised physician order entry system. resistance to capecitabine and vinorelbine who have previously received treatment lines A data collection form was designed to record patient's demographics, time since including taxanes and anthracyclines. diagnosis, sites of metastases, previous lines of treatment, number of cycles of eribulin, progression free survival (PFS), and overall survival (OS) ajusted by age, previous treatment lines (anthracyclines, taxanes, capecitabine and vinorelbine), administration of subsequent lines, and types and number of metastases. Scraphs were produced and statistics were performed using several packages of the R patients treated. language (R Development Core Team, <u>http://www.R-project.org</u>) RESULTS first year HR (95%CI) HR (95%CI) results p (longrak test) p (longrak test) 2.275 (0.823, 6.287) 2.275 (0.823, 6.287) 0,103 0,103 Age <55 years 1.45 (0.553, 3.919) 1.544 (0.586, 4.065) Comply with previous lines 0,461 0,375 Comply >=2 y <5previous lines 0,079 2.417 (0.757, 7.716) 0,126 3.021 (0.830, 10.99) 0.605 (0.134, 2.737) 0.605 (0.134, 2.737) 0,51 0,51 Previous anthracyclines 1.461 (0.331, 6.445) 0,614 1.679 (0.381, 7.396) 0,488 Previous taxol 1.273 (0.363, 4.459) 0,705 0,851 revious anthracyclines and taxol 1.128 (0.320, 3.975) 1.117 (0.405, 3.077) Previous vinorelbine 1.117 (0.405, 3.077) 0,831 0,831 0,356 1.982 (0.449, 8.737) 0,356 Previous capecitabine 1.982 (0.449, 8.737) 0.326 (0.104, 1.022) 0,0546 0.424 (0.147, 1.22) 0,101 Posterior lines Metastases 1.803 (0.686, 4.733) lung 2.039 (0.748, 5.55) 0,155 0,225 0.758 (0.242, 2.373) bone 0.961 (0.269, 3.423) 0,633 0,951 0.969 (0.126, 7.424) 0,976 0,976 Carcinomatosis peritoneal 0.9696 (0.126, 7.424) 0,571 0,571 0.7424 (0.263, 2.091) Lymph node 0.7424 (0.263, 2.091) 4.495 (1.011, 19.99) 0,062 3.72 (0.844, 16.38) 0,031 2.762 (0.767, 9.945) brain 2.762 (0.767, 9.945) 0,104 0,104 0,573 0,756 0.558 (0.071, 4.354) 0.726 (0.095, 5.523) 0,877 0,877 skin/Soft tissues 1.124 (0.253, 4.993) 1.124 (0.253, 4.993) 1.88 (0.355, 9.936) 0,457 Number of metastases >3 3.271 (0.381, 28.07) 0,28 Number of metastases 2-3 2.634 (0.336, 20.65) 0,665 1.401 (0.304, 6.446) 0,357 ard ratio: 95%CI: 95% coenficient Interval CONCLUSIONS

breast cancer who have progressed after at least one chemotherapeutic regimen for advanced disease. However, in our hospital its use is limited to a subgroup of patients with cancer in a clinical setting. In addition, we explored factors that might influence survival of results | reviewed.

BACKGROUND Eribulin has been indicated for the treatment of patients with locally advanced or metastatic PURPOSE The aim was to analyze the effectiveness of eribulin for the treatment of metastatic breast Clinical data of 40 patients [97,5% women, 54 years old (range 33-85)] were finally U With a median of time since breast cancer diagnosis of 8.1 years, they had been received a median of 4.6 (range 2-7) treatment lines. U We detected that most patients did not fulfill local criteria for eribulin use (67.5%). However, they received 3.5 (range 1–16) cycles for metastasic diseas (location were 75% bone, 50% lung, 65% liver, and 10% brain). Median PFS was 2.4 months (0.5-16.5) and OS with 45% of events was 4.2 months (0.5-20.5). 17,5% of the patients died before 3 months. \Box Only liver metastases predicted OS [hazard ratio 4.495; 95% CI 1.011-19.99; p = 0.031).

In our case, the effectiveness of eribulin in the clinical setting was modest. V PFS and OS values were lower than published in literature. W Survival analysis did not identify a subgroup of patients that could benefit of this treatment in our population.

METHODS



