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Retrospective Study

Eribulin for heavily pre-treated metastatic breast cancer patients

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Abstract

AIM: To discuss the experience with eribulin in clinical practice outside a clinical trial.

METHODS: Archives of patients treated for metastatic breast cancer were reviewed and 21 patients treated with the new chemotherapeutic eribulin mesylate, a synthetic analog of a natural marine product, were identified. Information on patients characteristics and treatment outcomes was extracted. Treatment with eribulin mesylate was initiated at the recommended dose of 1.4 mg/m² on day 1 and 8 of a 21 days cycle in 17 patients and at a decreased dose of 1.1 mg/m² day 1 and 8 of a 21 cycle in 4 patients due to comorbidities and frailty. Efficacy of the drug was evaluated using to the revised RECIST (Response

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