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Background:

Fosaprepitant contains the synthetic surfactant polysorbate 80, which has been associated with infusion-site adverse events (ISAEs) and hypersensitivity systemic reactions (HSRs) during or after administration.¹ Previously, ISAEs have been reported in 15%-67% of patients receiving fosaprepitant.²⁻⁶

Study design

- A retrospective chart review of patients (n=127) who had received doxorubicin plus anthracycline-based (AC) chemotherapy via a peripheral line from 14 US oncology community practices¹
- Patients were treated with fosaprepitant as part of a three-drug antiemetic combination regimen including a 5-HT₃ RA and dexamethasone¹

Primary objective

- To investigate and describe the incidence of ISAEs and HSRs during and following the administration of fosaprepitant with anthracycline-containing chemotherapy¹



Similar to other publications, rates of HSRs and ISAEs with fosaprepitant have been found to be higher than originally reported¹⁻⁶

Results/Conclusions

- The results of this chart review found that 35 of 127 patients (28%) developed ISAEs/HSRs during or after the infusion of antiemetics and chemotherapy¹
 - The rate of ISAEs appears to be higher than shown in the prescribing information, which reports ISAEs in 3% of HEC-treated patients and 2.2% of MEC-treated patients¹
- Several institutions have changed their antiemetic clinical practices due to the rate of ISAEs reported with fosaprepitant¹

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