

H2-antagonists for prevention of taxane infusion reactions: are they required?

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Background

- Taxane-based chemotherapy is used in the treatment of multiple malignancies
- Due to the high risk of hypersensitivity reactions, product information for paclitaxel and cabazitaxel recommends pre-medication with antihistamine, corticosteroid and H2-antagonist
- In September 2019, ranitidine became unavailable due to possible contamination. Prior to routine availability of alternate H2-antagonists our unit administered taxane chemotherapy without H2-antagonists. This practice has continued for weekly paclitaxel and 3-weekly cabazitaxel protocols

Aims

- To determine if the rate or severity of taxane infusion reactions changed following removal of ranitidine as a standard pre-medication within our institution

Method

- A retrospective review of case notes was conducted of all patients who received weekly paclitaxel or 3-weekly cabazitaxel for one year preceding unavailability of ranitidine (August 2018 to August 2019) and one year post unavailability (October 2019 to October 2020)
- Case notes were reviewed for the first three cycles of all patients who received weekly paclitaxel and all cycles for patients who received cabazitaxel

Results

- 191 patients received 593 infusions of weekly paclitaxel or 3-weekly cabazitaxel
- No patient receiving cabazitaxel experienced an infusion reaction
- No patient experienced more than one reaction

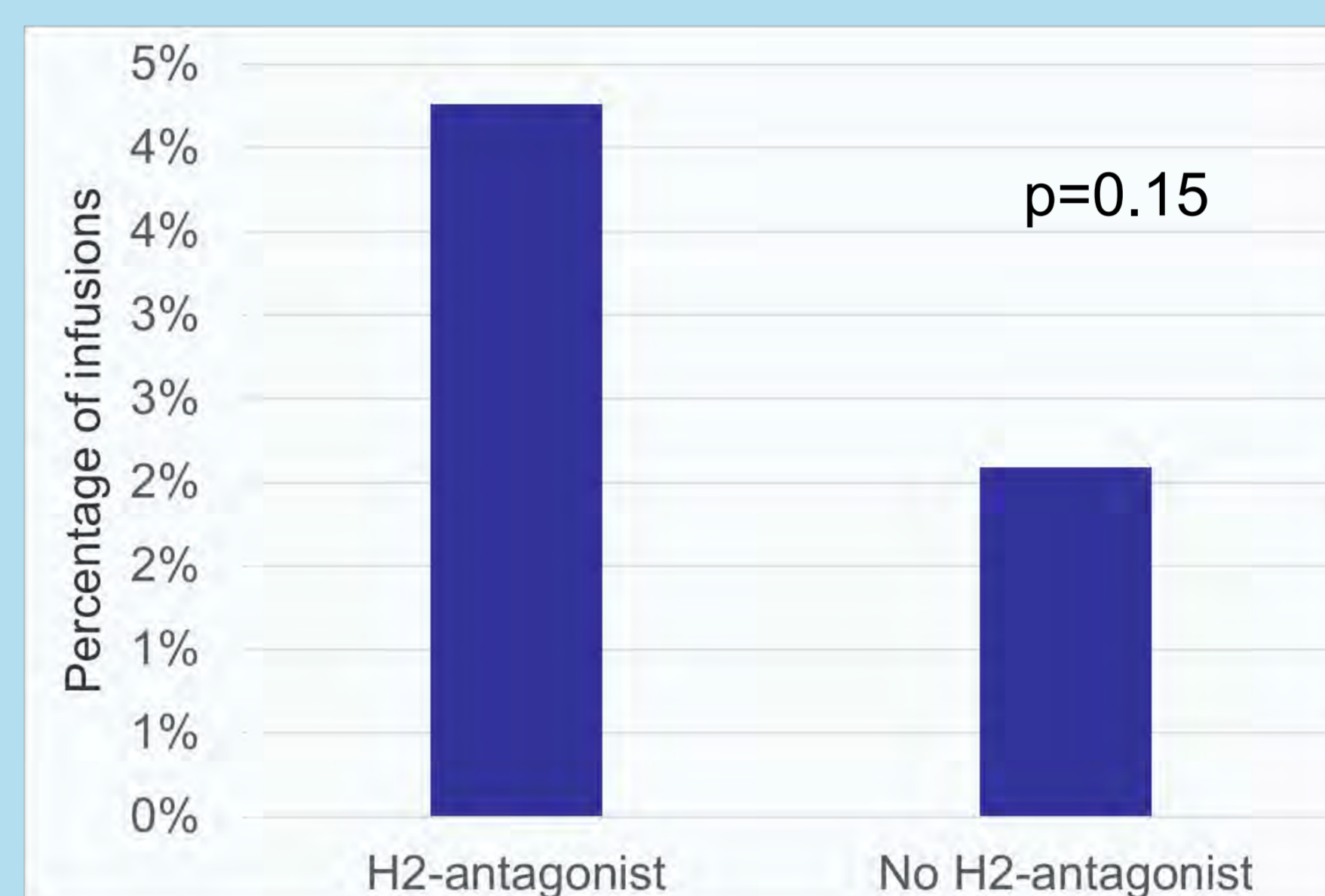


Figure 1: Infusion reactions

Characteristic	H2-antagonist (n=258)	No H2-antagonist (n=335)
Paclitaxel	243 (94%)	290 (87%)
Cabazitaxel	15 (6%)	45 (13%)
Monotherapy	114 (44%)	170 (51%)
Combination therapy	144 (56%)	165 (49%)

Table 1: Patient characteristics

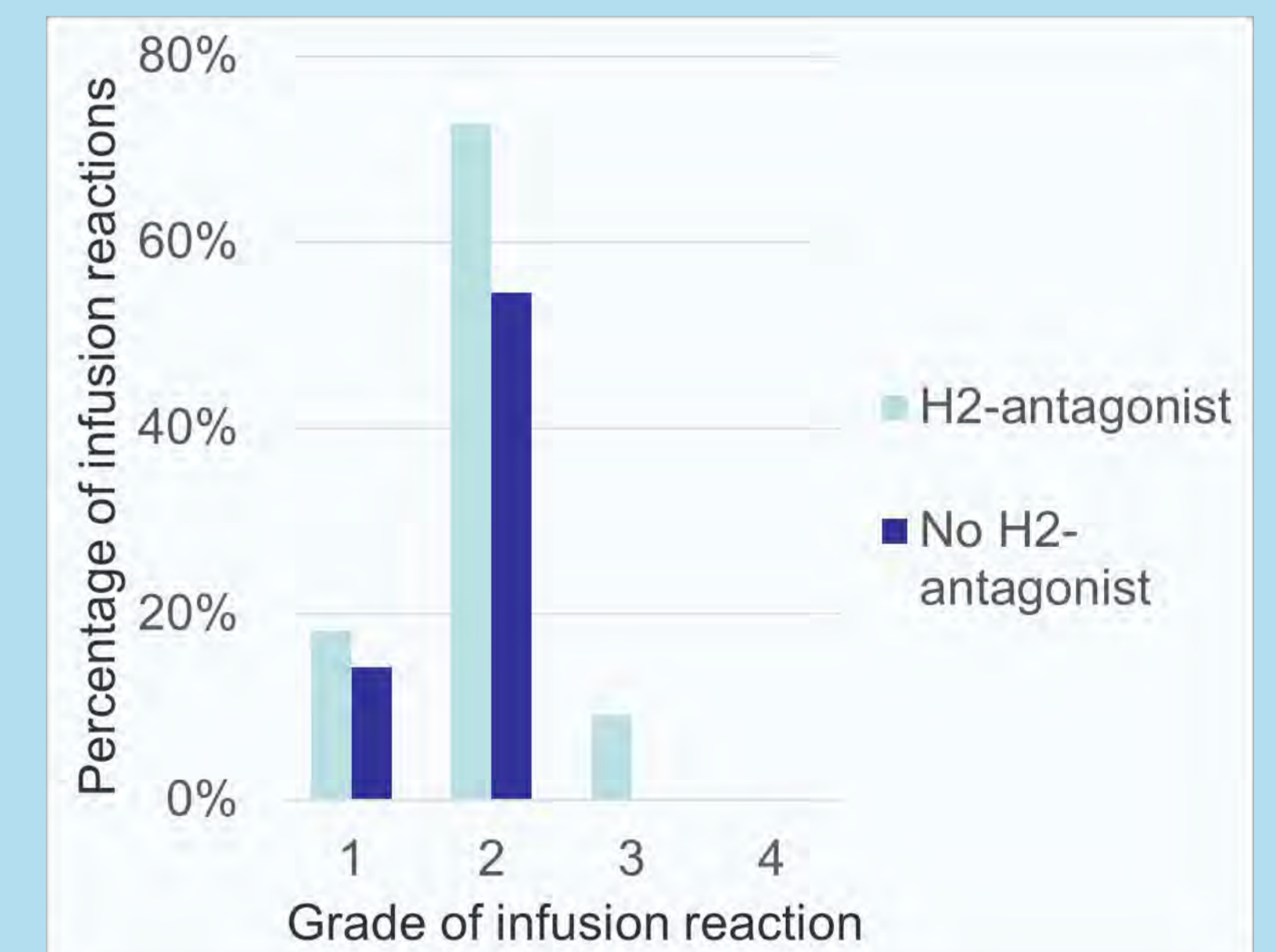


Figure 2: Grade of Infusion Reactions

Conclusion

- We observed no increase in the rate of infusion reactions with weekly paclitaxel and 3-weekly cabazitaxel despite cessation of routine H2-antagonists administration
- **Our study suggests patients may be able to safely receive these treatments without routine H2-antagonist**

For more information

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