

EVALUATION OF PREMEDICATION USE IN ADVERSE DRUG REACTIONS OCCURRENCE IN PATIENTS WHO RECEIVED INFlixIMAB TO TREAT INFLAMMATORY BOWEL DISEASE

X. Larrea Urtaran¹, Q López Noguera¹, G. Espín Martín², L. Torrealba Medina², C. Díez Vallejo¹, À. Castelló Nòria¹, M. Bruguera Teixidor¹, E. Nogué Pujadas¹, D. Busquets Casal², R. Sacrest Güell¹.

1. University Hospital Dr. Josep Trueta, Pharmacy Department, Girona, Spain.
2. University Hospital Dr. Josep Trueta, Digestive Department, GIRONA, SPAIN.



Abstract number: 4CPS-142
ATC code: L04- IMMUNOSUPPRESSANTS

AIM AND OBJECTIVES

Infliximab can cause infusion-related reactions like delayed hypersensitivity or anaphylactic shock. Using corticosteroids or antihistamines as premedication can reduce adverse drug reactions (ADRs) frequency.

OBJECTIVE: evaluate premedication impact on ADRs occurrence in patients with inflammatory bowel disease (IBD) who received infliximab.

MATERIAL AND METHODOS



Retrospective, observational study.
• Patients with IBD who received intravenous infliximab from January 2016 to December 2020.



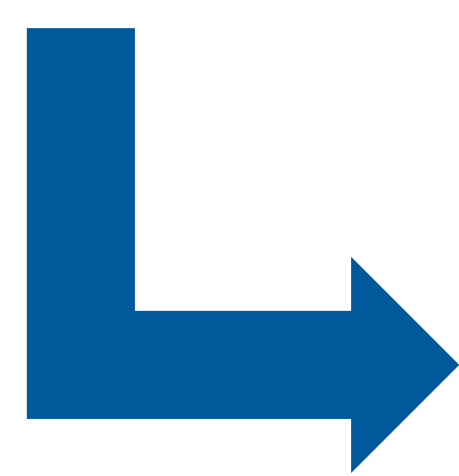
- Demographic variables (age and sex).
- Type of IBD.
- Harvey index.
- Mayo index.
- Premedication used (type of drug and number of administrations).
- Number of infliximab administrations.
- ADRs characteristics.



Statistical analysis:
• Mean.
• Standard deviation (SD).
• Absolute risk (AR).

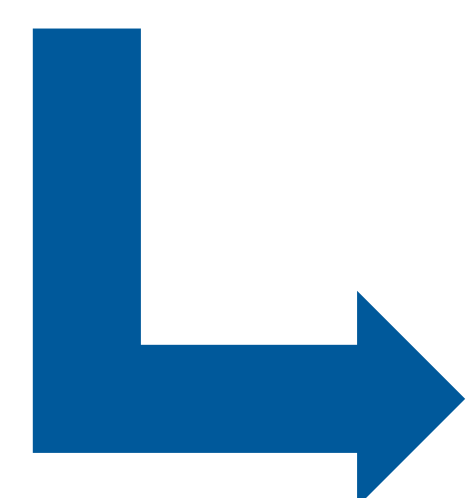
RESULTS

119 patients were included with an average age of 46 ± 17 years and 42% were women.

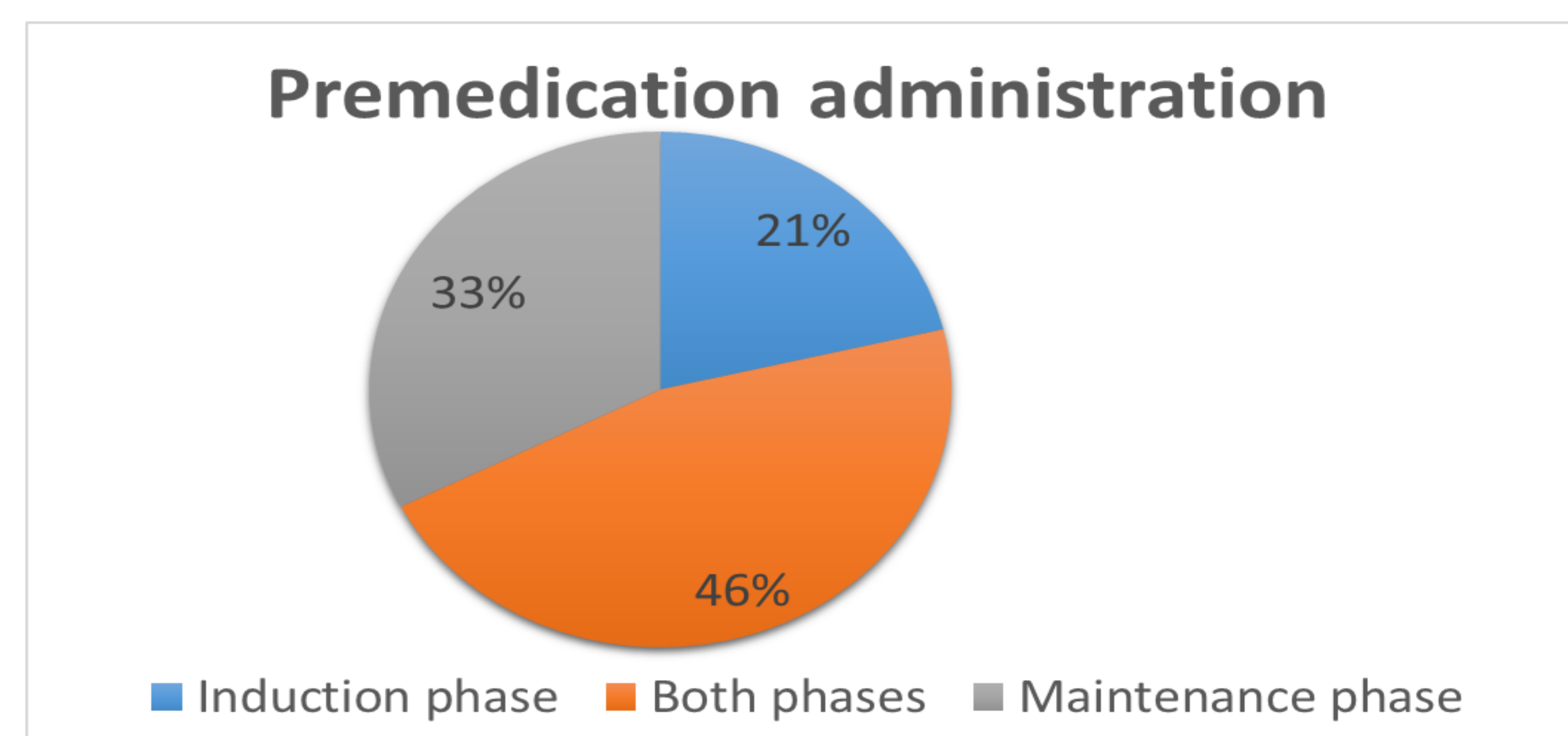


42 patients had **ulcerative colitis**: partial Mayo score mean of 3.7 ± 2.3 .
74 patients had **Crohn's disease**: Harvey score mean of 7.1 ± 3.7 .
3 patients had **indeterminate colitis**.

A total of **1909 infliximab infusions** were administrated: **premedication was used in 1185 administration in 80 patients**.



Glucocorticoids were used as a premedication in **97.5% of cases**.



25 ADRs were recorded in **21 patients**.

- The patients (n=17) who received **premedication** had 21 ADRs and an AR of 10.3% (CI95, 0.7%-19.8%).
- The patients who **didn't receive premedication** had 4 ADRs (n=4) and an AR of 21.3% (CI95, 12.3%- 30.2%).

44% of ADRs occurred in induction phase and 56% in maintenance phase.

The **main symptoms** of ADRS registered:

skin manifestations (n=16)
cardiovascular (n=6)
respiratory symptoms (n=3).

CONCLUSIONS

No lower absolute ADR risk were observed in patients who **received premedication** compared to patients who did not receive premedication. More studies should needed in order to evaluate the impact of premedication on ADRs occurrence.