

# Esophageal variceal ligation in patients on anticoagulation and factors that affect post-procedure bleeding risk

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### Introduction

- Anticoagulation medication use in patients with cirrhosis has increased in recent years.
- The presence of esophageal varices and necessity of band ligation (EVL) complicates their use.
- Optimal agents and recommendations for periprocedural management are lacking.

# Aim

 Better understand specific risk factors associated with post-EVL bleeding events in patients on anticoagulation.

# Methods

- Retrospective chart review identified all patients with cirrhosis undergoing EVL and requiring concurrent periprocedural anticoagulant management from 1/1/2015-2/16/2022.
- Patients with non-cirrhotic portal hypertension were excluded.
- Information about bleeding events were collected four weeks after EVL.
- Major and minor bleeding events were defined by the International Society on Thrombosis and Haemostasis criteria.
- Comparative statistics were performed using the Fisher exact test and Wilcoxon test.

| Characteristics                  | Bleeding group | Non-bleeding group | P-value |
|----------------------------------|----------------|--------------------|---------|
| MELD-Na                          | 24             | 12                 | <0.01   |
| Child Pugh                       | 11             | 7                  | <0.01   |
| Avg # days AC discontinued prior | 2.8            | 2.7                | 0.3     |
| Avg # days until<br>AC restarted | 3.0            | 3.7                | 0.6     |
| Avg Na                           | 135            | 138                | 0.03    |
| Avg Bilirubin                    | 8.0            | 1.2                | < 0.01  |
| Avg Creatinine                   | 1.0            | 0.9                | 0.6     |
| Avg INR                          | 2.3            | 1.4                | < 0.01  |
| Avg Platelets                    | 117            | 119                | 0.8     |
| Avg Hgb                          | 11.7           | 11.8               | 1       |

| Table 1 | l. Charac | teristics | of b | leeding | and | non-bl | leeding | graiin |
|---------|-----------|-----------|------|---------|-----|--------|---------|--------|
|         | i. Charac |           | OI U | iccumg  | and | HOH U  | ccumg   | group. |

| Gender                          | M 100%<br>W 0%  | M 70%<br>W 30%  | 0.3 |  |
|---------------------------------|---|---|-----|--|
| Indication for Endoscopy        | Primary Prophylaxis - 4 Secondary Prophylaxis - 1 Active Bleeding - 1 | Primary Prophylaxis - 24 Secondary Prophylaxis - 14 Active Bleeding - 2 |     |  |
| Subtherapeutic dosing           | 20%   | 27%   | 1   |  |
| Choice of AC                    | DOAC 75%<br>Other 25%   | DOAC 81%<br>Other 19%   | 0.6 |  |
| Indication for AC               | PVT - 3<br>DVT/PE - 2<br>Arrhythmia - 0<br>Other- 1                   | PVT - 27<br>DVT/PE - 9<br>Arrhythmia - 2<br>Other- 2                    |     |  |
| High Risk<br>Esophageal Varices | 100%  | 85%   | 0.6 |  |

Non-bleeding group

Table 2. Characteristics of bleeding and non-bleeding group.

**Characteristics** Bleeding group

| Bleeding<br>Event | MELD-<br>Na | Child<br>Pugh<br>Class | AC         | Therapeutic AC | Days<br>Discontinu<br>ed Prior | Days<br>Started<br>After | Major<br>bleed | Days Bleeding Started After Procedure | Cause of Bleed     | 6 Week<br>Mortality |
|-------------------|-------------|------------------------|------------|----------------|--------------------------------|--------------------------|----------------|---------------------------------------|--------------------|---------------------|
| 1                 | 12          | A (6)                  | Apixaban   | Yes            | 2                              | Unknow<br>n              | Yes            | 15                                    | Post banding ulcer | Alive               |
| 2                 | 30          | C (13)                 | Enoxaparin | Yes            | 3                              | 1                        | Yes            | 9                                     | Post banding ulcer | Alive               |
| 3                 | 30          | C (11)                 | Enoxaparin | Yes            | N/A                            | 1                        | Yes            | 5                                     | EV Rebleed         | Alive               |
| 4                 | 17          | C (12)                 | Coumadin   | Yes            | 4                              | 1                        | Yes            | 20                                    | PHG/GAVE           | Alive               |
| 5*                | 25          | B (9)                  | Apixaban   | Yes            | 2                              | 7                        | No             | 7                                     | PHG/GAVE           | Deceased            |
| 6*                | 27          | B (9)                  | Apixaban   | Yes            | Unknown                        | 5                        | Yes            | 12                                    | Post banding ulcer | Deceased            |

Table 3. Characteristics of patients with bleeding events. Asterisk indicates bleeding event associated with the same patient.

#### Results

- Forty six procedures were identified that met inclusion criteria.
- There were six bleeding events (13%)
- Bleeding events occurred between 5-20 days post procedure.
- Bleeding events were associated patients with more severe liver disease, as represented by higher mean MELD-Na (24 vs. 12, p<0.01) and Child-Pugh score (11 vs. 7, p<0.01).
- Bleeding events were not associated with timing of pre-procedural anticoagulation discontinuation (p=0.30), timing of anticoagulation initiation or resumption (p=0.60), type of anticoagulant (DOAC vs. other, p=0.60), subtherapeutic dosing (p=1.00), or the presence of high-risk esophageal varices (p=0.60).

## Conclusions

- Bleeding events were not associated with the timing of starting or resuming anticoagulation, type of anticoagulation, subtherapeutic dosing, or high-risk esophageal varices.
- Bleeding was associated only with severity of liver disease.
- Our results may help inform recommendations for EVL periprocedural anticoagulation management.
- Further research involving a larger sample size is needed to more fully characterize the risk factors for bleeding post-EVL.