Iron-deficiency anemia (IDA) occurs in 5% of premenopausal women and 1–2% of men and postmenopausal women. It is a common reason for referral to gastroenterologists, accounting for 4–13% of outpatient visits [1, 2]. IDA may be due to: blood loss (gastrointestinal, hemoptysis, traumatic, menorrhagia, hematuria, regular donors, post-surgery, iatrogenic repeated studies, etc.), malabsorption (celiac disease, gastrectomy or other enteropathies), or low iron intake (very low frequency in adults in our area) [1, 2]. In premenopausal women, the most common cause is gastrointestinal and/or gyneco-obstetric blood loss and in men and postmenopausal women, the most common cause is gastrointestinal blood loss. In most cases, the cause of bleeding is accessible to standard endoscopy (gastroscopy and colonoscopy) [1, 2]. However, in 2–10%, the lesions are located in the small intestine, so that these studies are negative [1, 2].

Small-bowel capsule endoscopy (CE) is a diagnostic tool that allows painless and safe assessment of the small bowel, leading to an etiologic diagnosis of small-bowel bleeding in approximately 60% of cases [3, 4]. Two meta-analyses [4, 5] have shown that CE is superior to push enteroscopy and radiological imaging for diagnosing clinically significant small-bowel pathology in patients with “suspected small-bowel bleeding (SBB)”, previously called obscure gastrointestinal bleeding.

Diagnostic yield and predictive factors of findings in small-bowel capsule endoscopy in the setting of iron-deficiency anemia

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Methods
Patients with unexplained IDA and negative upper and lower endoscopy were included. A positive diagnostic yield was considered when CE diagnosed one or more lesions that could explain the IDA. Sex, age, NSAID consumption, blood transfusion requirement, and ferritin and hemoglobin levels were recorded.

Results
In total, 120 CE were included (mean age 58.5 years; F/M 82:38). Mean hemoglobin levels were 9 g/dL and mean ferritin levels were 15.7 ng/mL. Positive findings were present in 50% of patients. The most frequent was angiodysplasia (45%). Despite several baseline variables being significantly associated with positive findings, using a logistic regression model, it was verified that male sex (OR 3.93; 95%CI 1.57–9.86), age (OR 1.03; 95%CI 1.00–1.06), and hemoglobin levels (OR 0.73; 95%CI 0.57–0.94) were the variables having an independent effect on the probability of obtaining positive findings. Age older than 50 years (OR 14.05; 95%CI 1.69–116.23) and male sex (OR 3.63; 95%CI 1.29–10.17) were the variables which increased the risk of diagnosing angiodysplasia.

Conclusions
CE is a useful technique in patients with IDA. To improve its yield, it is necessary to select patients carefully. Male sex, older age, and low hemoglobin levels were associated with a risk of positive finding in this group of patients. The risk of diagnosing angiodysplasia increased with male sex and older age.
the severity of IDA. Saurin et al. classified the lesions as highly relevant (P2), of uncertain relevance (P1), or of low relevance (P0) [8]. P2 lesions include angiodysplasia, tumors, ulcerative lesions, and varices, P1 include red spots and small or isolated erosions, and P0 include visible mucosal veins, diverticula without the presence of blood, or nodules without mucosal breaks. In a subsequent study, a P3 category was included for active bleeding [9].

Several studies have evaluated the effectiveness of CE in SBB, but few have focused on IDA alone [10 – 17]. Furthermore, in those few reports, the exact significance of the lesions identified has been poorly studied [14, 15]. Therefore, we aimed to assess the diagnostic yield of CE in patients with who had undergone previous thorough work-up. We also aimed to determine whether sex, age, NSAIDs, blood transfusions, hemoglobin (Hb), or ferritin levels could be predictors of findings.

Patients and methods

This is a retrospective cohort study of all consecutive patients undergoing CE for the evaluation of unexplained IDA conducted at the Gastroenterology Department of the “Hospital de Clínicas”, a tertiary referral center in Uruguay. All patients with unexplained IDA who had undergone at least one negative upper and lower endoscopy between January 2008 and May 2016 were included. IDA was defined, according to the laboratory distribution of these variables. The significance level in all cases was 0.05. When the variables did not follow a normal distribution, non-parametric tests were used to study their association with angiodysplasia and the presence of findings (Mann-Whitney U test). The analysis of age was made as a continuous variable and also in two groups (older and younger than 50 years). This cut-off was selected following Muhammad et al. [13]. A multivariate analysis of the variables whose P value ≤0.25 was performed. A logistic regression was performed using as dependent variables: presence of findings and the presence of angiodysplasia.

Results

In total, 120 CE were performed in 118 patients. CE completion rate was 96.7 %, defined by visualization of the cecum on the recording. Four studies (3.3 %) were defined as incomplete, in one of them CE was stopped after 3 hours due to an ulcerated stenosis that was later crossed through; in another patient CE stopped at a submucosal lesion and stayed there for 5 hours but was defecated 24 hours later. In the remaining two cases, the sensor was removed 8 hours after being swallowed (misreading of cecal image at data recorder screen). There were no complications. Of the total, 51 CE were performed with PillCamSB, 51 CE with PillCamSB2, and 18 with PillCamSB3.

The mean patient age was 58.5 years, ranging from 18 to 86 years. Eighty-seven patients (72.5 %) were older than 50 years. Eighty-two were women (68.3 %). Fifty-four patients (45 %) used NSAIDs and 54 patients (45 %) had received blood transfusions (►Table 1). Only 10 patients (8.3 %) were receiving anticoagulant therapy. The mean Hb level was 9 g/dL (range 4 – 11.9 g/dL). Twenty-one patients (17.5 %) had severe anemia (OGIB). Furthermore, CE and double-balloon enteroscopy have comparable diagnostic yields in SBB [6, 7].
(Hb ≤ 7 g/dL) and 99 patients (82.5 %) had mild to moderate anemia. The mean ferritin level was 15.7 ng/dL (range 1.5–48.0 ng/dL).

Positive findings were present in 60 patients with IDA (50 %). The distribution of positive findings is shown in ▶ Fig. 1 and images of their spectrum are shown in ▶ Fig. 4. Two patients (3.3 %) had more than one lesion. Active small-bowel bleeding was found in 5 patients (8.3 %), 3 of them without identification of a clear lesion. Three patients (5 %) had positive findings outside the small bowel (colonic active bleeding, pyloric ulcer, and cecal angiodysplasia) and were included as positive findings for statistical analysis.

In the 3 patients with bleeding lesions outside the small bowel, the following interventions were performed: a second colonoscopy was performed in the patient with colonic active bleeding and a previously missed colon cancer was diagnosed; upper gastrointestinal endoscopy was performed to take gastric and pyloric ulcer biopsies and a colonoscopy was performed to treat cecal angiodysplasias with argon plasma coagulation (APC).

Demographic, clinical, and laboratory data of patients with and without positive findings are shown in ▶ Table 1.

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Total n (%)</th>
<th>Positive findings (n = 60)</th>
<th>Without positive findings (n = 60)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sex: male</td>
<td>38 (32 %)</td>
<td>27 (45 %)</td>
<td>11 (18 %)</td>
<td>0.02</td>
</tr>
<tr>
<td>Age &gt; 50 years</td>
<td>87 (72.5 %)</td>
<td>50 (83 %)</td>
<td>37 (62 %)</td>
<td>NS</td>
</tr>
<tr>
<td>Anemia, g/dL</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>• Mild – moderate (Hb &gt; 7)</td>
<td>99 (83 %)</td>
<td>45 (75 %)</td>
<td>54 (90 %)</td>
<td>NS</td>
</tr>
<tr>
<td>• Severe (Hb ≤ 7)</td>
<td>21 (17 %)</td>
<td>15 (25 %)</td>
<td>6 (10 %)</td>
<td>NS</td>
</tr>
<tr>
<td>Ferritin, ng/dL</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>• &lt; 15</td>
<td>68 (57 %)</td>
<td>35 (58 %)</td>
<td>33 (55 %)</td>
<td>NS</td>
</tr>
<tr>
<td>• 15 – 30</td>
<td>37 (31 %)</td>
<td>18 (30 %)</td>
<td>19 (32 %)</td>
<td>NS</td>
</tr>
<tr>
<td>• 31 – 50</td>
<td>15 (12 %)</td>
<td>7 (12 %)</td>
<td>8 (13 %)</td>
<td>NS</td>
</tr>
<tr>
<td>NSAID: yes</td>
<td>54 (45 %)</td>
<td>30 (50 %)</td>
<td>24 (40 %)</td>
<td>NS</td>
</tr>
<tr>
<td>Transfusion: yes</td>
<td>54 (45 %)</td>
<td>33 (55 %)</td>
<td>21 (35 %)</td>
<td>0.03</td>
</tr>
<tr>
<td>Anticoagulant therapy: yes</td>
<td>10 (8 %)</td>
<td>4 (7 %)</td>
<td>6 (10 %)</td>
<td>NS</td>
</tr>
<tr>
<td>Capsule reached cecum</td>
<td>116 (97 %)</td>
<td>57 (95 %)</td>
<td>59 (98 %)</td>
<td>NS</td>
</tr>
</tbody>
</table>

IDA, iron deficiency anemia; NS, not significant.

Discussion

In this study, the diagnostic yield of CE for IDA was 50 %, which is in agreement with data from previous studies [17]. Our study was a phase IV clinical study, confirming the high diagnostic yield of CE found in previous phase II clinical trials [18, 19]. Most studies on CE had been performed in the USA, Europe, and the Far East. Our study is the largest on CE in IDA in South America. Therefore, our data also help define the epidemiological and etiological characteristics of IDA in additional geographic areas.

The majority of positive findings in our study were due to angiodysplasia and non-specific inflammation/ulceration, which is consistent with those found in similar studies from USA and Europe but not those from Asia. In Asia, reports of small-bowel tumors, inflammatory lesions, and multifocal stenosis enteritis are more common [20–23]. It has been recognized that CE is more sensitive than other modes of small-bowel imaging for the detection of vascular malformations [24].

All lesions included were considered to be highly relevant lesions (P2 of the Saurin classification). The “Saurin classification of small-bowel lesions on CE” allows risk stratification of patients [8, 9]. Highly relevant lesions (P2) had previously been validated by two blinded endoscopists in 100 % of cases, compared with 73 % and 27 % of cases for intermediate- (P1) and low-relevance (P0) lesions, respectively [25]. There was a necessity for therapy in a significantly higher number of P2 lesions (61 %), compared with P1 or P0 lesions (23 %) [25].

In this series, in addition to angiodysplasia, one case of multiple phlebectasias was included as vascular lesions. These are rare benign vascular anomalies that may present with occult or overt small-bowel bleeding. We have already reported two
cases of these lesions, even with active bleeding during the procedure [26].

Due to the clinical relevance, in the 50% diagnostic yield, we included 4.4% of findings outside the small bowel. Our lesion miss rate outside the small bowel is much lower than those reported in previous studies, which found that patients referred for small-bowel endoscopic imaging had a significant number of lesions within the reach of upper and lower endoscopy [27–29]. Fry et al. [30] found 24.3% of sources of gastrointestinal bleeding outside the small bowel with double balloon enteroscopy (DBE). Elijah et al. [31] also found that up to 38.8% of patients undergoing CE for OGIB had significant lesions of the upper gastrointestinal tract such as gastric antral vascular ectasias (GAVE), gastric ulcers, and Cameron’s lesions. These lesions were evidently missed during electrocardiography (EGC) and colonoscopy performed before DBE and/or CE. In our series, the lesions missed were: a pyloric ulcer whose histology revealed an adenocarcinoma, a colonic active bleeding due to a small colorectal cancer (CRC), and a cecal angiodysplasia. We believe that having performed the previous upper and lower endoscopies in our tertiary center may have resulted in a lower number of missed lesions. Indeed, despite the reported high percentages of lesions outside the small bowel and the high cost of the CE, most guidelines still do not routinely support a second-look endoscopy prior to small-bowel CE [32–34].

There is limited published data in which clinical and laboratory factors predict the ability of CE to detect pathology in IDA. In a large series of patients with OGIB of which 74% had IDA, Sisduh et al. [33] found that increasing age, use of warfarin, and liver comorbidity were significant factors which predicted a higher yield with CE. More recently, in 221 patients with IDA, Koulaouzidis et al. [17] found that age, prior blood transfusions, use of anticoagulants, and male sex were factors associated with a higher diagnostic yield for all types of small-bowel pathology. Similar to Koulaouzidis et al., we found that increas-

![Fig. 1](image1.png) Distribution of positive findings in 60 patients with IDA.

![Fig. 2](image2.png) Box plots of Hb levels in patients with and without positive findings.

![Fig. 3](image3.png) Box plots of age in patients with and without positive findings.
ing age was associated with a higher detection rate of small-bowel angiodysplasia.

All these results are partially in line with ours. Nevertheless, and to the best of our knowledge, ours is the first study to show low hemoglobin levels as a predictor of findings in patients with IDA. Furthermore, in contrast to most previous studies, our study focused exclusively on patients with IDA. We carefully select the IDA group, excluding celiac disease diagnosis in all patients by antibody detection and/or duodenal biopsies. Thus, this study demonstrated that, in IDA patients, after an intensive previous work-up, capsule endoscopy would be a very useful technique to find lesions in elderly patients with severe anemia.

Our study has several potential limitations. First, it is a retrospective study and thus prone to the potential deficits of such design. However, we carefully collected all variables including all videos of the procedures to ensure a complete database. Second, we are a tertiary center and our findings may not be reproduced by others. Third, as most of the patients were referred exclusively for CE, some data is missing. We only had data for H. pylori investigation in 29%. As H. pylori infection can be a cause of unexplained iron-deficiency anemia, this could be a limitation. Also, data on iron supplementation was not available. Fourth, we included nonspecific diagnosis as potentially explaining the bleeding leading to IDA. But until now there is no clear definition of which lesions result in bleeding and we based our definitions of bleeding lesions on existing classifications and guidelines. Nonetheless, further studies will be mandatory to clearly define which lesions result in bleeding IDA. Finally, patients were referred only for CE from different healthcare centers and the subsequent medical treatment depended on the attending physician and the availability of resources in their health service. It was not possible to perform a complete follow-up of these patients so an analysis of outcomes was not included in the study. It would be interesting to perform a study on this variable in the future.

In conclusion, CE is a useful technique in patients with IDA. However, as shown by this study, to improve its yield, it is necessary to select patients carefully. Male sex, old age, and low hemoglobin levels increased the risk of positive findings in this group of patients. The risk of diagnosing angiodysplasia increases with age. Furthermore, large multicenter, phase IV studies are needed to better define the lesions causing IDA and the outcomes of patients with IDA subjected to CE.

▶ Fig. 4 Spectrum of positive findings. a Active bleeding; b malignant tumor-like lesion; c angiodysplasia; d ulcerated stenosis; e ulcerated submucosal lesion; f multiple phlebectasia.