ABSTRACT – Background – The diagnosis of corrosion cancer should be suspected in patients with corrosive ingestion if after a latent period of negligible symptoms there is development of dysphagia, or poor response to dilatation, or if respiratory symptoms develop in an otherwise stable patient of esophageal stenosis. Narrow Band Imaging detects superficial squamous cell carcinoma more frequently than white-light imaging, and has significantly higher sensitivity and accuracy compared with white-light. Objective – To determinate the clinical applicability of Narrow Band Imaging versus Lugol’s solution chromendoscopy for detection of early esophageal cancer in patients with caustic/corrosive agent stenosis. Methods – Thirty-eight patients, aged between 28-84 were enrolled and examined by both Narrow Band Imaging and Lugol’s solution chromendoscopy. A 4.9mm diameter endoscope was used facilitating examination of a stenotic area without dilation. Narrow Band Imaging was performed and any lesion detected was marked for later biopsy. Then, Lugol’s solution chromendoscopy was performed and biopsies were taken at suspicious areas. Patients who had abnormal findings at the routine, Narrow Band Imaging or Lugol’s solution chromoscopy exam had their stenotic ring biopsied. Results – We detected nine suspicious lesions with Narrow Band Imaging and 14 with Lugol’s solution chromendoscopy. The sensitivity and specificity of the Narrow Band Imaging was 100% and 80.6%, and with Lugol’s chromoscopy 100% and 66.67%, respectively. Five (13%) suspicious lesions were detected both with Narrow Band Imaging and Lugol’s chromoscopy, two (40%) of these lesions were confirmed carcinoma on histopathological examination. Conclusion – Narrow Band Imaging is an applicable option to detect and evaluate cancer in patients with caustic/corrosive stenosis compared to the Lugol’s solution chromoscopy. HEADINGs – Esophageal neoplasms. Squamous cell carcinoma. Caustics, adverse effects. Iodides, therapeutic use.

INTRODUCTION

In 1904, Teleky described the first case of esophageal cancer (EC) after corrosive ingestion. It is known in this patient population the incidence of EC is 1%-4% and the duration between caustic ingestion and development of EC is 15-30 years(11,18,21). Alkalis corrosive agents combine with tissue proteins and cause liquefactive necrosis and saponification, and penetrate deeper into tissues, helped by a higher viscosity and a longer contact time through the esophagus. Additionally, alkali absorption leads to thrombosis in blood vessels, impeding blood flow to already damaged tissue(35).

Esophageal neoplasms (both adenocarcinoma and squamous cell carcinoma) may develop as a late complication of caustic injury at a rate 1000-3000 times higher than expected in patients of a similar age(22). When squamous cell carcinoma (SCC) occurs in these patients it appears in the stenotic ring and may be due to repeated trauma of dilation, slow passage of food (stasis), reflux of the food due to the shortening of the esophagus and of the scarring process itself(1,9,18,19).

Different to non caustic cancer, the long-term survival for scar carcinoma patient was excellent: 45.6% of the patients after resection were alive after 5 years and 14.4% after 10 years(17). Three reason can explain this: first, caustic-injured patients are followed quite closely and this may contribute to a diagnosis of cancer at an earlier stage, as in all patients at risk for developing neoplasms, who are in a close follow-up programme. Second, oesophageal carcinoma developing after caustic ingestion may be diagnosed earlier because the tumour grows only intramurally due to the surrounding scarring. A smaller lesion, developing in a lye stricture, is responsible for earlier dysphagia, although the duration of symptoms is longer in these patients, who are already suffering from dysphagia. Third, the intra- and peri-tumoral fibrosis might prevent the early cancer spread, with lymph node and visceral metastases developing only at a later phase(6,21).
One-stage resection and reconstruction is the best way to treat the radically operable patients. In patients with esophageal corrosive stricture in need of operation, both a bypass procedure and resection can be performed, but it should be pointed out that malignancy may develop even years after the operation in the remaining part of the gullet. Total esophagectomy is therefore suggested instead of by-pass(17).

The diagnosis of corrosive cancer should be suspected in patients with corrosive ingestion if after a latent period of negligible symptoms there is development of dysphagia, or poor response to dilatation, or if respiratory symptoms develop in an otherwise stable patient of esophageal stenosis(20).

Currently, there are many diagnostic methods available to facilitate the early detection of EC, among which upper GI endoscopy, is the most important. Upper endoscopy when combined with imaging magnification and chromoscopy may be the key to improving diagnosis and may subsequently improve survival. The need to develop techniques to improve the early diagnosis of esophageal SSC without side effects has spawned research into new endoscopic optical techniques. For example, NBI which was demonstrated first in 1994, which uses spectroscopy(2,14).

The technique known as “Narrow Band Imaging” (NBI), uses narrow light spectrum, which penetrates in the superficial tissues, allowing the identification of small blood vessels and details of the mucosal pattern(12). With the NBI technique the normal mucosa appears light blue/green and the vessels brown, lighting up suspicious lesions, similarly to Lugol’s chromoscopy(5,16). This allows for demarcation of suspicious areas and hence targeted biopsy(2,18). NBI detects superficial SCC more frequently than white-light imaging, and has significantly higher sensitivity and accuracy compared with white-light. In addition, narrowband imaging magnified endoscopy (NBI-ME) is useful for differentiating cancers from non-cancerous lesions and assessing tumor depth and invasion by analysis of the microvascular patterns. Generally, the presence of irregular loop-shaped microvessels suggests noninvasive cancer in the mucosal layer, and thick non-looped vessels suggest invasive cancer in typical SCC(13).

The aim of this study is to determine the clinical applicability of Narrow Band Imaging versus Lugol’s solution chromoscopy for detection of early esophageal cancer in patients with caustic/corrosive agent stenosis.

METHODS

This is a single center, observational transversal trial, cross-over study, that enrolled 38 patients with esophageal stenosis secondary to corrosive ingestion who underwent upper GI endoscopy with NBI and Lugol’s iodine chroomendoscopy at the Gastrointestinal Endoscopy Unit of Hospital das Clínicas da Universidade de São Paulo. This project was granted approval by the Ethics Committee, and was registered in the Sao Paulo University Ethics Committee number 1087/06.

Patients were included if they had their ingestion >20 years ago and had esophageal stenosis secondary to caustic lesion caused by ingestion. Informed consent was obtained in all patients. Patients were excluded if they had their caustic injury in the last 20 years, refused to consent, had an iodine allergy, had under 21 years age or refused to undergo upper endoscopy.

The gold standard were histopathological, surgery and follow-up.

Endoscopic procedure

Upper gastrointestinal endoscopy was performed with the videoendoscope (Olympus 180 – GIF Type N180 “slimlight”) with 4.9mm diameter by only one endoscopist (E.G.H.M., with more than 25 years’ experience in experience of conventional endoscopy). He had experienced more than 70,000 esophagogastroduodenoscopies and had more than 6 years of experience with NBI.

The exam sequence was:

- Monitorization with pulse oximetry and oxygen catheter, intravenous access, sedation with midazolam (0.025 – 0.1 mg/kg), associated to fentanyl (0.7 - 2µg/kg) and propofol (0.5 – 1mg/kg).
- Conventional endoscopic exam and removal of fluids and secretion of the esophagus with saline and N-Acetylcysteine.
- Exam with image enhanced endoscopy (NBI) and when the eventually lesions were found, were taken notes about their localization in relation to incisors and esophageal walls.
- Then, Lugol’s iodine solution 1.25% chromoscopy was performed and eventual found lesion were taken notes similarly to NBI exam.
- Previously marked suspicious lesions were biopsied with disposable forceps. If no lesion was identified, then the stenotic ring was biopsied and sent for histopathological examination.
- The lesion was considered suspicious (positive findings) when presents Lugol voiding area >5 mm in Lugol’s chroomoscopy and brownish and dark brown spots in NBI chroomoscopy (Figures 1 and 2).

No suspicious areas were considered negative findings. The chroomendoscopy exam with Lugol’s iodine and NBI have had the same execution time.

Statistical analysis

The results were submitted to statistical analysis to evaluate sensitivity, specificity, positive and negative predictive values, likelihood ratio of a positive test, likelihood ratio of a negative test and accuracy of esophageal chromoscopy with NBI and Lugol.
RESULTS

Thirty-eight patients (22 female – 57.8%) with a median age of 56 years (28-84) were enrolled. In this study, abnormal (pre-malignant or malignant) lesions no were detected during regular endoscopy. The median length of follow-up in the entire cohort was 4 months. Of these, 29 (76.3%) patients had negative findings and 9 (23.7%) positive findings with NBI. Lugol identified 24 (63.2%) negative findings and 14 (36.8%) positive findings identified. All the suspected lesions were located adjacent to stenosis. Histopathologic analysis showed 36 (94.7%) no cancer biopsies versus 02 (5.3%) SCC anatomiopathological diagnosis. All the patients are followed annually, and additional controls exams were performed on patients with negative findings or dysplasia, that not confirmed positive cancer samples.

Lugol iodine chromoendoscopy identified 14 suspicious areas (Lugol voiding area) and two (14.3%) were confirmed as SCC. There was no case of neoplastic disease not identified by Lugol iodine chromoendoscopy resulting in a sensitivity was 100% (exact Fisher’s Test 0.1294). The specificity was 66.67%. Therefore, there were 12 (85.7%) false positives. The positive predicted value was of 14.3% and the negative predictive value was 100%. The likelihood ratio of a positive test was of 2.9, and its negative correspondent doesn’t exist. The accuracy of this diagnostic test was of 73% (Table 1).

TABLE 1. Evaluation of conformity between chromoendoscopy with Lugol’s solution and the result of the histopathological exam in the detection of early esophageal cancer in patients with esophagus stenosis secondary to caustic ingestion. According to this, there is a week conformity between the tests (Kappa = 0.174).

<table>
<thead>
<tr>
<th>Kappa</th>
<th>0.174</th>
<th>[-0.213 – 0.561]</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sensitivity</td>
<td>100.00%</td>
<td>[34.2 –100.0]</td>
</tr>
<tr>
<td>Specificity</td>
<td>66.67%</td>
<td>[50.3 – 79.8]</td>
</tr>
<tr>
<td>Positive predictive value</td>
<td>14.30%</td>
<td>[4.0 – 39.9]</td>
</tr>
<tr>
<td>Negative predictive value</td>
<td>100.00%</td>
<td>[86.2 – 100.0]</td>
</tr>
<tr>
<td>Likelihood ratio of a positive test</td>
<td>2.9</td>
<td></td>
</tr>
<tr>
<td>Likelihood ratio of a negative test</td>
<td>Inexistent</td>
<td></td>
</tr>
<tr>
<td>Accuracy</td>
<td>0.73</td>
<td></td>
</tr>
</tbody>
</table>

NBI identified nine suspicious areas (brownish and dark brown spots) considered positive findings which 2 (22.2%) were confirmed as SCC in the histopathological examination. Seven of nine were false positives, since brownish areas were negative for SCC in the histopathological examination. According to the table there was no case of neoplasia that wasn’t identified by the NBI technique, 29 were true negatives (exact Fisher’s Test 0.0512) (Table 2).

TABLE 2. Evaluation of conformity between the NBI technique and the histopathological exam. According to this, there is a week conformity between the tests (Kappa = 0.239).

<table>
<thead>
<tr>
<th>Kappa</th>
<th>0.204</th>
<th>[-0.162 – 0.77]</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sensitivity</td>
<td>100%</td>
<td>[34.2 –100.0]</td>
</tr>
<tr>
<td>Specificity</td>
<td>80.56%</td>
<td>[65.0 – 790.2]</td>
</tr>
<tr>
<td>Positive predictive value</td>
<td>22.20%</td>
<td>[6.3 – 54.7]</td>
</tr>
<tr>
<td>Negative predictive value</td>
<td>100%</td>
<td>[74.1 – 100.0]</td>
</tr>
<tr>
<td>Likelihood ratio of a positive test</td>
<td>5</td>
<td></td>
</tr>
<tr>
<td>Likelihood ratio of a negative test</td>
<td>0.0</td>
<td></td>
</tr>
<tr>
<td>Accuracy</td>
<td>0.81</td>
<td></td>
</tr>
</tbody>
</table>

When assessing the conformity between the NBI and the Lugol’s chromoendoscopy in diagnosis of EC, both methods revealed five (55.6%) positive findings, two of those were cases of neoplasia. Nine (31%) negative findings with the NBI technique and positive with Lugol’s solution; four (44.4%) negative findings with Lugol’s solution and positive with NBI technique. Twenty (69%) were negative findings in both methods (Table 3A and 3B).

TABLE 3A. Evaluation of conformity between the NBI technique and the Lugol’s chromoendoscopy for detection of early esophageal cancer in patients with esophageal stenosis secondary to caustic lesion.

<table>
<thead>
<tr>
<th>Esophagus neoplasy</th>
<th>NBI</th>
<th>Lugol</th>
</tr>
</thead>
<tbody>
<tr>
<td>Positive test</td>
<td>5 (55.6%)</td>
<td>4 (44.4%)</td>
</tr>
<tr>
<td>Negative test</td>
<td>9 (51.0%)</td>
<td>20 (69.0%)</td>
</tr>
</tbody>
</table>

TABLE 3B. Evaluation of conformity between the NBI technique and the Lugol’s chromoendoscopy for detection of early esophageal cancer in patients with esophageal stenosis secondary to caustic lesion. According to this, there is a weak conformity between the techniques (Kappa = 0.206).

<table>
<thead>
<tr>
<th>Kappa</th>
<th>0.206</th>
<th>[-0.144 – 0.556]</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sensitivity</td>
<td>35.70%</td>
<td>[16.3 – 61.2]</td>
</tr>
<tr>
<td>Specificity</td>
<td>83.30%</td>
<td>[64.1 – 93.3]</td>
</tr>
<tr>
<td>Positive predictive test</td>
<td>55.60%</td>
<td>[26.7 – 81.1]</td>
</tr>
<tr>
<td>Negative predictive test</td>
<td>69.00%</td>
<td>[50.8 – 82.7]</td>
</tr>
</tbody>
</table>

To improve the data analysis the Fagan’s Nomogram was used with the likelihood ratio of a positive test for both methods. Therefore, the likelihood ratio of a positive test was 2.9 for the Lugol and 5 for the NBI (Figure 3).

FIGURE 3. Fagan’s Nomogram. “A” (Black line) is NBI and “B” (red line) is Lugol.
Figure 3 shows that the pre test probability of EC in these patients was 5%. Using the likelihood ratio of a positive test for the NBI (A, Black line) of 5 and for the Lugol (B, red line) of 2.9, the post test probability obtained is 22% for the NBI and for the Lugol. The clinical use of these values indicates that a patient with esophageal stenosis secondary of caustic ingestion has 18% and 22% higher chance of having SCC diagnosed by Lugol and NBI, respectively, than a healthy patient.

According to Figure 4, we notice that the pre test probability for the SCC in patients with stenosis secondary to caustic lesion was 5%. Using the likelihood ratio of a positive test for the NBI (A, Black line) and for Lugol (B, red line), we have achieved a post test probability of 22% for the NBI and 18% for the Lugol. Trying to improve the current data analysis, considering the pre test probability of 5% and applying a likelihood ratio of a positive test of 5 for the NBI, there is a post test probability of 22% Following the same process, considering the pre test probability now of 22% and the likelihood ratio of a positive test for the Lugol, there is a post test probability of 52%, highly improving the odds of diagnosis using both methods.

**FIGURE 4.** Fagan’s Nomogram. “A” (Black line) is NBI and “B” (red line) is Lugol.

**DISCUSSION**

This is an observational study, assessing 38 patients with esophageal stenosis secondary to caustic lesion. We compared the use of NBI and Lugol chromoendoscopy for its ability to diagnose EC. The study was performed by one proceduralist and both methods were performed during the same endoscopic exam. We identified two cases of SCC among the 38 selected patients, which gave the prevalence of EA of 5%. This is consistent with a study Goldman presented a prevalence of 0.8%-4%.

In this study we used an ultra slim gastroscope, with 4.9 mm diameter, which allowed the evaluation of the stenotic ring’s mucosa, without needing dilatation, excluding some distortion factors caused by the dilation. A majority suspected lesions and esophageal neoplasms are diagnosed at oral side of stenosis or adjacent to stenosis. So, about early cancer diagnosis, no difference existing between standard adult gastroscope or ultrathin gastroscope. A large review of lye corrosion carcinoma of the esophagus with 63 cases showed that majority of corrosion carcinomas occur at the site of tracheal bifurcation, possibly due to stasis and more severe injury at this site because of anatomical narrowing. Fifty-three of the 63 patients were not after the occurrence of SCC is seen in conventional endoscopy. Reported a case of a 65-year-old man who had a flat and small squamous cell carcinoma in the esophagus and the endoscopic observation with the NBI system was useful for detecting the lesion.

In the current study the sensitivity and specificity of the Lugol’s chromoendoscopy was 100% and 66.67%, respectively. The sensitivity was high since all the cases of carcinoma were identified. The probable explanation for the lower specificity, at least with regards to Lugol’s iodine is that due to the healing process caused by the caustic injury, areas of vascular neof ormation and fibrosis, presenting low glycogen and therefore do not color by iodine dye. The positive predictive value was low, 14.29%, since it relates indirectly to the specificity; and the negative predictive value was high, 100%, since all the cases of cancer were diagnosed. The findings were consistent to the literature, what makes NBI suitable for screening, once the chance of false negatives is minimal. The test’s accuracy was of 73% (12,30,31,28).

Another comparative study between NBI technique and Lugol’s chromoendoscopy in the diagnosis of SCC in patients with history of head and neck neoplasia showed that the NBI and Lugol’s chromoendoscopy were equally sensitive and easy to perform (23). We also found these results in our study, which confirm the useful of both methods.

The accuracy of NBI endoscopy in screening for esophageal cancer has been reported to be comparable to that of Lugol chromoendoscopy, with low rates of adverse events. Especially, the specificity of NBI endoscopy with or without magnifying imaging was higher than that of Lugol chromoendoscopy. Although Lugol chromoendoscopy is the current gold standard for screening for esophageal cancer, NBI endoscopy might be the first-choice endoscopy for screening in the future (15).

In the current study the sensitivity and specificity of the Lugol’s chromoendoscopy was 100% and 66.67%, respectively. The sensitivity was high since all the cases of carcinoma were identified. The probable explanation for the lower specificity, at least with regards to Lugol’s iodine is that due to the healing process caused by the caustic injury, areas of vascular neof ormation and fibrosis, presenting low glycogen and therefore do not color by iodine dye. The positive predictive value was low, 14.29%, since it relates indirectly to the specificity; and the negative predictive value was high, 100%, since all the cases of cancer were diagnosed. The findings were consistent to the literature, what makes NBI suitable for screening, once the chance of false negatives is minimal. The test’s accuracy was of 73% (12,30,31,28).

**CONCLUSION**

NBI is an applicable option to detect and evaluate esophageal carcinoma patients with caustic lesion and corrosive stenosis compared to the Lugol’s solution chromoendoscopy.

**Authors’ contributions**

Guedes HG, Moura DTH, Pennacchi CMPS, Kumbhari V wrote the article, did review of the literature and Moura EGH did the exams; Pennacchi CMPS and Amorim RBP collected cases’ outcomes; Moura EGH critically revised the manuscript; final version was approved by all authors.
Pennachi CMPS, Moura DTH, Amorim RBP, Guedes HG, Kumbhari V, Moura EGH. Lugol’s iodine chromoendoscopy versus Narrow Band Image enhanced endoscopy for the detection of esophageal cancer in patients with stenosis secondary to caustic/corrosive agent ingestion

RESUMO – Contexto – A suspeita do câncer de esôfago na lesão cáustica ocorre quando os pacientes com estenoses previamente estáveis, após um período latente sem sintomas, apresentam disfagia, baixa resposta às dilatações ou sintomas respiratórios. A cromoscopia com luz de banda estreita detecta o câncer superficial de esôfago mais frequentemente que a luz branca, com alta sensibilidade e acurácia. Objetivo – Determinar a aplicabilidade clínica da luz de banda estreita versus a cromoscopia vital com Lugol na detecção do câncer precoce de esôfago em pacientes com lesões cáusticas.

Métodos – Um total de 38 pacientes, entre 28 e 84 anos, foram alocados seguidamente e submetidos à cromoscopia com luz de banda estreita e com Lugol. Um gasto-cópico de 4,9 mm de diâmetro foi usado para facilitar o exame da área estenosada, sem necessidade de dilatação. A cromoscopia com luz de banda estreita era realizada primeiro e as áreas suspeitas anotadas. Depois, a cromoscopia com Lugol era realizada e as áreas suspeitas biopsiadas. Resultados – Detectamos nove lesões suspeitas com a luz de banda estreita e 14 com o Lugol. A sensibilidade e especificidade da cromoscopia com luz de banda estreita foi de 100% e 80,6%, e a do Lugol foi de 100% e 66,67% respectivamente. Cinco (13%) lesões suspeitas foram detectadas coincidentemente pelos dois métodos, sendo duas (40%) com diagnóstico anatomo-patológico de câncer de esôfago. Conclusão – A cromoscopia com luz de banda estreita é opção concreta para o diagnóstico de câncer em pacientes com estenoses esofágicas por corrosões cáusticas, comparado a cromoscopia com Lugol.


REFERENCES