Endoscopic Submucosal Dissection of Seborrheic Keratosis-Like Lesion of the Esophagus: A New Entity?

Nese EKİNCİ¹ 💿, Eylül GÜN¹ 💿, Fatih ASLAN² 💿

¹Department of Pathology, Izmir Katip Celebi University Ataturk Training and Research Hospital, IZMIR, TURKEY ²Department of Gastroenterology, Koc University Hospital, ISTANBUL, TURKEY

This study has been presented as a poster at the 27th Turkish National Congress of Pathology which took place in Antalya on 15-18th November, 2017.

ABSTRACT

Seborrheic keratosis, one of the most common lesions of the epidermis, is rarely seen on mucosal surfaces. We report a case of a distinctive epithelial neoplasm of the esophagus showing close resemblance to seborrheic keratosis that was resected with endoscopic submucosal dissection. A 65-year-old patient's previous esophageal biopsy showed suspicious low grade dysplasia and the patient was referred for endoscopic submucosal dissection of a flat lesion in the mid-esophagus. Macroscopic examination revealed a well circumscribed, pigmented and elevated lesion with a diameter of 20 mm. Microscopically, the lesion was well circumscribed, with plaque-like elevation, and showed hyperkeratosis, acanthosis, and papillomatosis. Broad coalescing solid sheets and interconnecting trabeculae of basaloid cells were the consistent feature throughout the lesion. Squamous eddies and occasional central keratinization were present. Mitotic activity and koilocytes were not identified. Immunohistochemically, the lesion showed diffuse nuclear positivity with p63 and negativity with p16. Ki-67 index was confined to the basal cell layer. With the help of histopathologic and immunohistochemical findings, we diagnosed this morphologically benign case as "seborrheic keratosis-like lesion of the esophagus". It should be kept in mind that seborrheic keratosis-like lesions might be rarely seen on mucosal surfaces such as the esophagus. Endoscopic submucosal dissection is a new, curative, and safe endoscopic resection technique in en-bloc resection of superficial esophageal lesions. To our knowledge, this is the first case of the aforementioned lesion in the esophagus being resected with endoscopic submucosal dissection.

Key Words: Endoscopic submucosal dissection, Esophagus, Seborrheic keratosis

INTRODUCTION

Seborrheic keratosis is one of the most common lesions in dermatopathology but it is rarely seen on mucosal surfaces (1). Curative endoscopic resection methods such as endoscopic mucosal resection (EMR) and endoscopic submucosal dissection (ESD) are new resection techniques initially developed in Japan and allow en bloc resection and treatment of superficial gastrointestinal lesions. Although application of ESD to the esophagus is limited in early stage esophageal neoplasia because of its greater technical difficulty, it avoids the high morbidity and mortality rates of surgical treatment (2) and offers a highly effective, safe and less expensive way for the detection and treatment of esophageal neoplastic lesions. It is considered the cornerstone of endoscopic treatment of Barrett's esophagus and early squamous cell carcinoma of the esophagus (3). It is therefore increasingly and successfully in use all around the world. We report a case of distinctive epithelial neoplasm of the esophagus, which shows close resemblance to seborrheic keratosis, one of the most common benign epidermal tumors, resected with ESD.

CASE REPORT

A 65-year-old man with slight dyspnea was referred to the gastroenterology clinic of a tertiary hospital in 2013. Gastro-esophagoscopy was done and no abnormality was found. The follow-up endoscopic biopsy in November 2016 showed low grade dysplasia and the patient was referred to our hospital for further examination and treatment. After pre-procedural assessment with narrow band imaging and chromoendoscopy with Lugol's solution, a flat lesion of the esophagus with a diameter of 20 mm at approximately 28 cm from the incisor teeth was seen (Figure 1). The resection borders were marked with dual knife and after submucosal injection of indigo-carmine and sodium hyaluronate solution, and en bloc resection of the lesion 2-3 mm away from the margins was successfully performed. Intra- and post-procedural prophylactic coagulation with hemostatic forceps followed. Complications such as delayed bleeding or perforation did not occur after the ESD and the patient was discharged 2 days after the treatment. The specimen was pinned against a plate peripherally by stainless-steel pins and entirely immersed in formaldehyde overnight to preserve the tissue shape and configuration (Figure 2).

Correspondence: Eylül GÜN Izmir Katip Celebi University, Ataturk Training and Research Hospital, Department of Pathology, Izmir, TURKEY E-mail: dreylulgun@gmail.com Phone: +90 506 828 40 48

(Turk Patoloji Derg 2020, 36:73-76)

Received: 05.07.2018 Accepted: 08.09.2018

Copyright © 2020 The Author(s). This is an open-access article published by Federation of Turkish Pathology Societies under the terms of the Creative Commons Attribution License which permits unrestricted use, distribution, and reproduction in any medium or format, provided the original work is properly cited. No use, distribution or reproduction is permitted which does not comply with these terms.

Macroscopic examination revealed a well circumscribed, slightly pigmented and elevated plague-like lesion with a diameter of 20 mm on a velvety mucosal surface. Alcian blue staining was used for macroscopic delineation of mucosal margins and the specimen was then serially sectioned perpendicularly at 2 mm intervals. All sections were subjected to histopathologic review. The lesion was well circumscribed, with plaque-like elevation on low power magnification and the base of the lesion was rough on an imaginary axis drawn between two mucosalsubmucosal junctions at both ends of normal esophageal

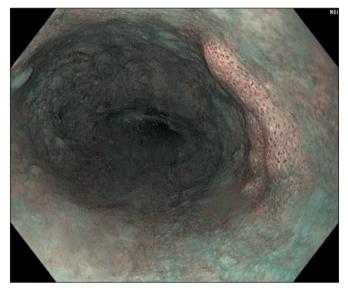


Figure 1: Endoscopic view of the flat lesion with a diameter of 20 mm at approximately 28 cm from the incisor teeth of the esophagus.

tissue. On 10x and 20x magnification, hematoxylin and eosin-stained sections revealed hyperkeratosis, acanthosis, and papillomatosis. Broad coalescing solid sheets and interconnecting trabeculae of basaloid cells were the consistent feature throughout the lesion (Figure 3). Squamous eddies and occasional central keratinization were present (Figure 4A). Mitotic activity and koilocytes were not identified. Immunohistochemically, the lesion showed negativity with p16, diffuse positivity with cytokeratin 5/6, and diffuse nuclear positivity with p63 (Figure 4B-C). The Ki-67 labeling index was confined only to the basal cell layer of the lesion and normal esophageal squamous epithelium (Figure 4D). No dysplasia was identified.

DISCUSSION

Seborrheic keratosis is one of the most common lesions seen by dermatologists and it is considered as a benign epidermal tumor but it may be a sign of concomitant skin cancer and internal malignancies. They are sharply demarcated, slightly elevated, hyperpigmented patch or plaque-like lesions and are seen commonly in areas such as the trunk, neck, face and upper extremities. They are considered as hyperkeratotic lesions of the epidermis and reported not to be seen on the mucosal surfaces (1). However, there are several case reports in the literature presenting seborrheic keratosis on the conjunctiva (4) and the nasal vestibule (5). All of the dermatological lesions with hyperkeratosis, acanthosis and papillomatosis might be considered in the clinical and pathological differential diagnosis. Our patient did not show any similar lesions on the skin and there was no internal malignancies detected with the imaging

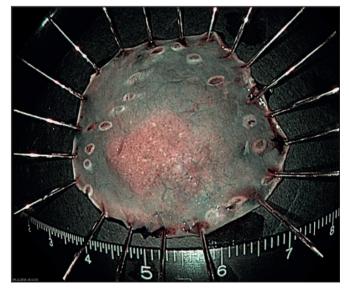


Figure 2: Pinned endoscopic submucosal dissection specimen.

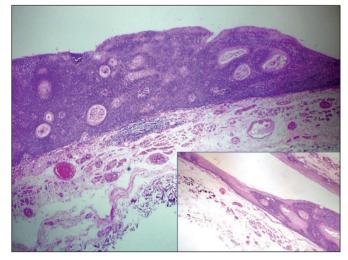


Figure 3: Circumscribed plaque-like lesion with acanthosis and the transition zone between the lesion and normal esophageal mucosa (inset) (H&E; x10).

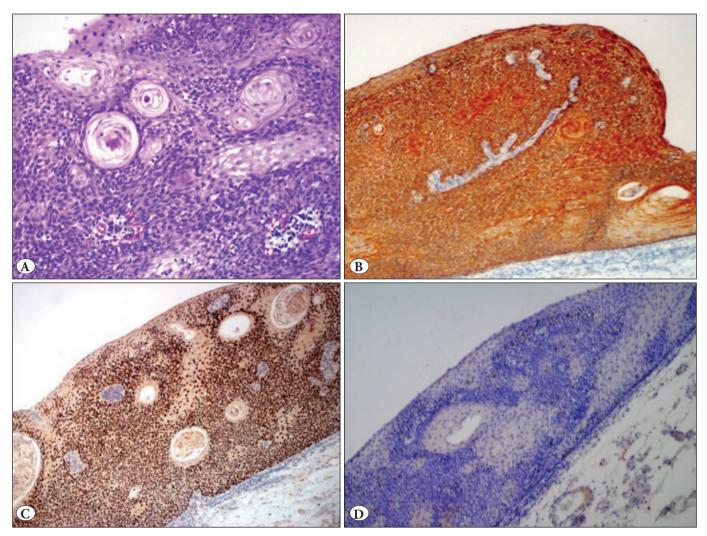


Figure 4: A) Squamous eddies, central keratinization and basaloid cells (H&E; x40). **B)** Cytokeratin 5/6 positivity (IHC; x20). **C)** Diffuse p63 positivity (IHC; x20). **D)** Ki-67 confined to basal layer (IHC; x20).

procedures performed before the endoscopic procedure. The Leser-Trélat syndrome, which is characterized by the eruptive appearance of multiple seborrheic keratoses in association with an underlying malignant disease, was therefore not considered during the differential diagnosis.

The term "seborrheic keratosis-like lesion" as a new entity was previously used in a case series by Talia and McCluggage that included a total of 7 cases of the cervix and vagina and a relationship with the human papilloma virus (HPV) was shown in two of these cases (6). However we did not find any similar lesions of the esophagus reported in the literature.

ESD is an effective method for neoplastic lesions of the esophagus and it is a safe treatment modality in the

management of early esophageal squamous cell neoplasms. However, great skill in this technique is definitely required (2). ESD is suggested to be performed rather than EMR while dissecting lesions that are larger than 15 mm because it ensures *en bloc* resections and the recurrence rates are lower (7). A study by Chen et al. on 296 patients with early esophageal squamous cell neoplasms and high-grade intraepithelial neoplasms showed no cancer-related deaths and it was concluded that ESD is a well-accomplished and secure procedure (8).

Rare lesions of the esophagus resected with ESD or EMR reported in the literature include Barrett's esophageal cancer (3), granular cell tumor (9) and leiomyomas in the category of stromal tumors (10).

Herein, we report a case of a superficial esophageal lesion resected with ESD. We diagnosed this morphologically benign case as a "seborrheic keratosis-like lesion of the esophagus" with the help of histopathologic and immunohistochemical findings. To our knowledge, this is the first case of the aforementioned lesion in the esophagus.

In conclusion, it should be kept in mind that seborrheic keratosis-like lesions might be rarely seen on mucosal surfaces such as the esophagus and that ESD is a safe procedure in *en bloc* resection of superficial esophageal lesions.

CONFLICT of INTEREST

The authors declare no conflicts of interest.

REFERENCES

- Phulari RG, Buddhdev K, Rathore R, Patel S. Seborrheic keratosis. Journal of Oral and Maxillofacial Pathology: JOMFP. 2014;18:327-30.
- 2. Chaves DM, Maluf Filho F, de Moura EGH, Santos MEL, Arrais LRG, Kawaguti F, Sakai P. Endoscopic submucosal dissection for the treatment of early esophageal and gastric cancer initial experience of a western center. Clinics. 2010;65:377-82.
- Ishihara R, Yamamoto S, Hanaoka N, Takeuchi Y, Higashino K, Uedo N, Iishi H. Endoscopic submucosal dissection for superficial Barrett's esophageal cancer in the Japanese state and perspective. Ann Transl Med. 2014;2:24.

- Kim JH, Bae HW, Lee KK, Kim TI, Kim EK. Seborrheic keratosis of the conjunctiva: a case report. Korean J Ophthalmol. 2009;23:306-8.
- Buyuklu F, Aydin H, Tarhan E, Ada S, Cakmak O.Seborrheic keratosis of the nasal vestibule. Kulak Burun Bogaz Ihtis Derg. 2007;17:298-300.
- 6. Talia KL, McCluggage WG. Seborrheic Keratosis-like lesions of the cervix and vagina: Report of a new entity possibly related to low-risk human papillomavirus infection. Am J Surg Pathol. 2017;41:517-24.
- Kesavan K, Chedgy FQ, Sharmila S, Sreedhari T, Arun K, Pradeep B. Early squamous neoplasia of the esophagus: The endoscopic approach to diagnosis and management. Saudi J Gastroenterol. 2017;23:75-81.
- Chen Y, Zhao Y, Zhao X, Shi R. Clinical outcomes of endoscopic submucosal dissection for early esophageal squamous cell neoplasms: A retrospective single-center study in China. Gastroenterology Research and Practice. 2016;2016:3741456.
- 9. Hulagu S, Senturk O, Aygun C, Gurbuz Y, Kocaman O, Celebi A, Konduk T. Granular cell tumor of esophagus removed with endoscopic submucosal dissection. Turk J Gastroenterol. 2007;18:188-91.
- Huang ZG, Zhang XS, Huang SL, Yuan XG. Endoscopy dissection of small stromal tumors emerged from the muscularis propria in the upper gastrointestinal tract: Preliminary study. World J Gastrointest Endosc. 2012 16;4:565-70.