Sebaceous glands in the esophagus-new approach to metaplastic theory: a case report and review of the literature

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Abstract

Ectopic salivary glands in the esophagus have rarely been reported, representing an obscure medical condition. The endoscopic findings of the EESg indicated multiple yellowish patches or papules, which varied in size. The histopathological findings indicated several lobulated sebaceous glands in the squamous epithelium with inflammatory infiltration. Esophageal “ectopic” sebaceous glands do not require further treatment or follow-up.

Background

Metaplasia is the conversion of one differentiate cell type to another [1]. Metaplastic lesions have been recognized in many tissues of endodermal origin, especially in the stomach and in the Barrett esophagitis [2,3]. In contrast, heterotopic sebaceous glands (HSGs) in the esophagus are considered to be very rare, representing an obscure medical condition [4-8]. Some authors have tried to determine whether HSGs is result of a metaplastic process or a heterotopic congenital anomaly? However, the answer is not yet known [6-8]. Often localisation of HSGs is in the middle and lower esophagus in numbers ranging from 1 to more than 100 yellowish plaques, measuring 1-2 mm in diameter, often without any clinical symptom [8-10].

Aim

Having in the mind the pointed-out dilemma on the pathogenesis of this rare esophageal process, we have undertaken the following study: the identification of both clinical and pathohistological features in the patient with rare esophageal sebaceous process.

Case report

A 46 year-old-woman, presented with a history of discomfort during swallowing, since the last 6 months. There was no history of vomiting, hematemesis, or weight loss. The patient was not a chronic smoker. General physical examination was unremarkable. Respiratory and cardiovascular systems were normal. Symptomatic treatment, with proton pump inhibitors, has been given. No significant abnormality was found on physical examination, or laboratory tests. Clinically, carcinoma of the esophagus was suspected and the esophagogastrointestinal endoscopy has been done. Seven wart-like yellowish papillomatous projections which varied in size, in the lower esophagus, were observed.

Pinch biopsy from one, of seven projections, was taken and sent for histological examination. Paraffin biopsy blocks were cut and stained with HE and PAS methods. Microscopically, the squamous epithelial lining was fragmented, showing hiperkerathosis (Figure 1).

The presence of the esophageal mucinous glands of white color, in the subepithelial area, was revealed (two arrows).

The mostly of subepithelial space was filled with metaplastic sebaceous glands: they were lobulated. with polygonal cells, that have small nuclei and abundant clear cytoplasm, showing microvesicular vacuolisation and typical sebaceous differentiation (Figure 2). Between these cells, could be seen the remnants of the mucinous cells of small size, confirming the metaplastic theory De La Pava et al. [4]. The second microscopical proof for metaplastic theory is chronic inflammation: dense lymphocytic and macrophage infiltration, surrounded by dense fibrous, sometimes by hyalinized stroma, that provide physical support to multilobular structures [5-8]. Endly, our patient is relatively old woman, not child. On the base of pointed out criterions, histological diagnosis of the sebaceous metaplasia in the esophagus, was made.

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The patient was treated symptomatically with antacids and omeprazole for one month, and she became symptom free and remained so at one year of follow-up. Histogenesis of this rare lesion is discussed in this case report: ESGs are derived from the ectoderm. There was no history of vomiting or hematoma or, weight loss. HCGs are very rare esophageal conditions with an incidence around 0.00465% and an occurrence rate of 0.41 per year (1-6; 9-110). Despite the extreme rarity at this entity, it is considered a lesion with no malignant potential [7-14].

Discussion

Many years have passed from the tume (1962) when De La Pava et al. [4] were discovered the ectopic sebaceous glands in the middle and lower esophagus. The authors reported such occurrence in 4 of the 200 unselected cadavers. However, in that time it was first reported presence of SGs (ectodermal) in an endodermally organ. The primary endoscopic impression was papilloma, but microscopical study proved, to be sebaceous glands, but unaccompanied by hair follicles. However, in that time (1962) the same authors first reported the presence of SGs in an endodermal derived organ- esophagus, in a 44- year- old nonsmoker. Histological examination identified numerous sebaceous glands in the lamina propria, showing the characteristics of sebaceous differentiation. Histogenesis of this process is not completely clear. To explain the presence of ectodermal tissue (sebaceous gland) in the endodermal organ (esophagus), several hypotheses have been postulated [5-9].

Metaplastic theory: this is favored by the occurrence of this condition in the elderly, not in children, and by the arrangement of these glands which is like that of esophageal submucous glands. In addition, chronic inflammation with lymphocytic and macrophagic cells must be also present.

Heterotopic theory: This thesis is not supported by endoscopic studies and autopsy studies conducted on infants or children [10]. HEGs in esophagus are a very rare occurrence. It probably represents a metaplastic change in the esophageal submucosal glands. Here, we confirm this metaplastic sebaceous rare occurrence in a 45 year –old female, by the following characteristics. Histogenesis is not completely clear. Several hypotheses have been postulated: metaplasia heterotopia.

Conclusions

• Sebaceous glands in the esophagus are very rare.
• Clinical symptoms, sex and endoscopic finding are not specific.
• Tiny yellowish lesions, of submucous localization in the lamina propria of the lower esophagus, are not specific.
• The most important discover is microscopically: normal, small, mucinous submucous esophageal glands and their multilobular sebaceous metaplasia in “sebaceous” glands.

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