

# Adult Onset Eruptive Syringoma

## Erişkin Başlangıçlı Erüptif Siringom

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### SUMMARY

Syringomas are common, benign eccrine sweat gland tumors, more frequently seen in women. Clinically, they appear as asymptomatic small firm papules skin colored to yellowish dermal papules often scattered in the periorbital area. Eruptive syringoma is a rare variant, which has been described to occur in successive crops on the anterior body surface, usually in young persons. A 38-year-old, healthy, white male presented with a 6-year history of papular eruptions. The lesions began as a few tan-colored papules on the abdomen and spread to all of the trunk and recently became mildly pruritic. Physical examination revealed multiple reddish brown-colored, flat-topped papules 1-3 mm in diameter on the anterior chest, abdomen and axillae. Also a few similar lesions were seen in the periorbital region. The biopsy specimen demonstrated a normal epidermis overlying a dermis that was filled with many small ducts of cuboidal cells with eosinophilic cytoplasm, along with cords of epithelial cells embedded in a fibrous stroma creating a 'tadpole' appearance. The lesions were diagnosed as eruptive syringoma with clinical and histopathological findings. While the localized form in the periorbital area can be clinically diagnosed, the eruptive form may not be easily considered in the differential diagnosis at the first visit. This rare entity should also be considered in the differential diagnosis of eruptive papular dermatosis.

**Key words:** Eccrine sweat gland tumor; eruptive syringoma; papular dermatosis; syringoma.

### ÖZET

Siringomlar kadınlarda daha sık görülen yaygın, selim ek rin ter bezi tümörleridir. Klinik olarak genellikle periorbi tal bölgede dağılım gösteren küçük, deri renginden sarım sı renge varan, sıkı kıvamlı asemptomatik papüller olarak belirirler. Erüptif siringom, genellikle gençlerde görülen, gövde ön yüzde çok sayıda ardışık lezyonlardan oluşan nadir bir varyantıdır. Otuz sekiz yaşında sağlıklı, beyaz er kek hasta, 6 yıldır devam eden papüler erüpsiyonlar nede niyle başvurdu. Lezyonlar karında birkaç adet bronz renkli papüller olarak başlamış ve tüm gövdeye yayılarak, son zamanlarda hafifçe kaşıntılı bir hal almıştı. Fizik muayene sinde, göğüs, karın ve aksillada çok sayıda kızıl kahveren gi, düz yüzeyle 1-3 mm çaplarında papüller vardı. Benzer birkaç adet lezyon periorbital bölgesinde de bulunmakta idi. Yapılan biyopside, fibröz stromaya gömülmüş eozinofi lik sitoplazmalı küboidal hücrelerden oluşan çok sayıda kü çük duktuslar ve 'iribaş' görüntüsü oluşturan epitelyal hü cre kordonları ile dolmuş bir dermisin üzerinde normal epi dermis saptandı. Bu klinik ve histopatolojik bulgular doğ rultusunda lezyonlar erüptif siringom olarak değerlendiril di. Periorbital bölgede yerleşim gösteren siringomlar kli nik olarak tanınabilirken, erüptif formu ilk değerlendirm e de ayırıcı tanıları arasında kolaylıkla düşünülmemektedir. Bu nadir antite de erüptif papüler dermatozların ayırıcı ta nısında düşünülmelidir.

**Anahtar sözcükler:** Ekrin ter bezi tümörü; erüptif siringom; pa püler dermatoz; siringom.

### INTRODUCTION

Syringomas are common, benign eccrine sweat gland tumors, more frequently seen in women. Clini cally, they appear as asymptomatic small firm pap-

ules skin colored to yellowish dermal papules often scattered in the periorbital area.<sup>[1,2]</sup> Eruptive syringo ma is a rare variant, which has been described to oc cur in successive crops on the anterior body surface,

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usually in young persons.<sup>[3]</sup> We present a 38-year-old man with a 6-year history of eruption, which should be considered in the differential diagnosis of eruptive papular dermatosis.

## CASE REPORT

A 38-year-old, healthy, white male presented with a 6-year history of papular eruptions. The lesions began as a few tan-colored papules on the abdomen and spread to all the trunk and recently became mildly pruritic. Since they first appeared, they had never been disappeared for 6 years. Antifungals and topical steroids had been used but the lesions did not show improvement. He had no any medical problem, was not on any medication or over-the-counter preparations, no family member had ever had a similar skin condition. Physical examination revealed multiple reddish brown-colored, flat-topped papules 1-3 mm in diameter on the anterior chest, abdomen and axillae. Also a few similar lesions were seen in the periorbital region (Fig. 1). The lesions were monomorphic, bilaterally distributed. General skin examination revealed no other significant skin lesions and systemic examination was unremarkable. Routine blood laboratory investigations were normal.

A skin biopsy was obtained with the prediagnosis of lichen planus and eruptive syringoma. The biopsy specimen demonstrated a normal epidermis overlying a dermis that was filled with many small ducts of cuboidal cells with eosinophilic cytoplasm, along with cords of epithelial cells embedded in a fibrous stroma creating a 'tadpole' appearance (Fig. 2). The ducts were lined by two rows of cuboidal epithelium and some contained amorphous debris in their lumina (Fig. 2c).

The lesions were diagnosed as eruptive syringoma with clinical and histopathological findings. He was treated with isotretinoin 40 mg/day (po) for 4 months but the lesions did not show marked improvement.

## DISCUSSION

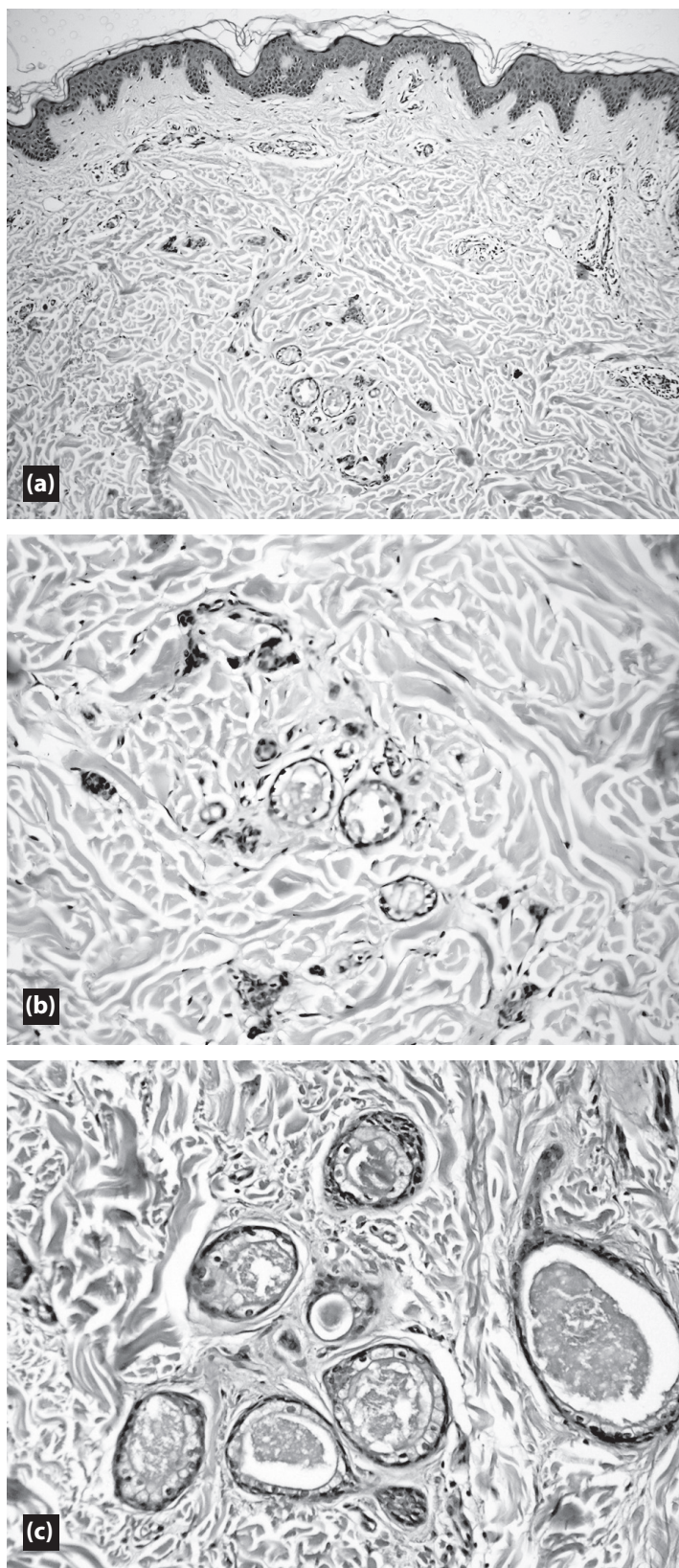
Syringoma had been classified into four principal clinical variants according to its clinical features and associations; a localized form, a familial form, a form associated with Down syndrome, and a generalized

form.<sup>[4]</sup> The most common, localized form presents with skin colored to yellow firm papules usually scattered in the periorbital region. Eruptive form is an infrequent variant, which clinically appears as multiple skin colored or slightly pigmented papules in large numbers and in successive crops on the anterior parts of the neck, chest, abdomen, axillae periumbilical region and proximal extremities. It usually starts before or during puberty.<sup>[1-3]</sup> But it has also been reported to appear after puberty as it was in our patient. The clinic, histologic and the immunohistochemical



**Fig. 1.** (a, b) Multiple reddish-brown colored flat-topped papules on the anterior chest, abdomen and axillae.





**Fig. 2.** (a-c) Cystic ducts and epithelial cords with comma like tails surrounded by collagen bundles. The lumina of the ducts contained amorphous debris.

properties of eruptive form is the same as classical syringoma; but the eruptive form may not be easily considered in the differential diagnosis at the first visit, while the localized form in the periorbital area can be clinically diagnosed.<sup>[4,5]</sup>

Eruptive syringoma treatment is done for cosmetic purposes, and the options are generally unsatisfactory. Physical techniques such as excision, cryosurgery, electrodesiccation may be frustrating with the risk of scarring. However, none of the treatment modalities eliminates the risk of recurrence; CO<sub>2</sub> laser, oral and topical retinoids and adapalene had been reported to be treatment,<sup>[3,6-8]</sup> but isotretinoin was not successful in our case.

## CONCLUSION

The clinic, histologic and the immunohistochemical properties of eruptive form is the same as classical syringoma; but the eruptive form may not be easily considered in the differential diagnosis at the first visit, while the localized form in the periorbital area can be clinically diagnosed. It should also be considered in the differential diagnosis of an adult onset eruptive papular dermatosis.

## REFERENCES

1. Friedman SJ, Butler DF. Syringoma presenting as milia. *J Am Acad Dermatol* 1987;16:310-4.
2. Tsunemi Y, Ihn H, Saeki H, et al. Generalized eruptive syringoma. *Pediatr Dermatol* 2005;22:492-3.
3. Patrizi A, Neri I, Marzaduri S, et al. Syringoma: a review of twenty-nine cases. *Acta Derm Venereol* 1998;78:460-2.
4. Pruzan DL, Esterly NB, Prose NS. Eruptive syringoma. *Arch Dermatol* 1989;125:1119-20.
5. Hashimoto K, Blum D, Fukaya T, et al. Familial syringoma. Case history and application of monoclonal anti-ecrine gland antibodies. *Arch Dermatol* 1985;121:756-60.
6. Soler-Carrillo J, Estrach T, Mascaró JM. Eruptive syringoma: 27 new cases and review of the literature. *J Eur Acad Dermatol Venereol* 2001;15:242-6.
7. Hashimoto K, DiBella RJ, Borsuk GM, et al. Eruptive hidradenoma and syringoma. Histological, histochemical, and electron microscopic studies. *Arch Dermatol* 1967;96:500-19.
8. Gómez MI, Pérez B, Azaña JM, et al. Eruptive syringoma: treatment with topical tretinoin. *Dermatology* 1994;189:105-6.
9. Frazier CC, Camacho AP, Cockerell CJ. The treatment of eruptive syringomas in an African American patient with a combination of trichloroacetic acid and CO<sub>2</sub> laser destruction. *Dermatol Surg* 2001;27:489-92.