A 38-year-old woman with a 25-year history of numerous cystic nodules and papules localized on the face, neck, chest, axillae, inguinal regions and vulva was referred to our department. Her mother, brother, sister and daughter had similar, but fewer lesions. In the last few years new cysts had appeared in the inguinal region and on the vulva impairing her sexual activity (Fig. 1) (Dermatology Life Quality Index (DLQI) score = 16 points). She was concerned about being seen in public, feeling physically unattractive and sometimes depressed.

Histopathology of the lesion excised from the axillary fold demonstrated a folded cystic lesion located in the mid-dermis (Fig. 2). In some parts of the lumen a small amount of eosinophilic horny material was present. No connection between the cyst wall and overlying epidermis was found. Immunohistochemical study showed the presence of keratin 17 in upper layer cells in the cyst wall.

What is your diagnosis?
Cystic Nodules Affecting Sexual Activity: Comment
Acta Derm Venereol 2010; 90: XX–XX (contd)

**Diagnosis: Steatocystoma multiplex**

Steatocystoma multiplex (SM) is an uncommon disorder characterized by development of numerous dermal cysts, which are benign nevoid pilosebaceous gland tumours. Although the condition is inherited in an autosomal dominant fashion, many sporadic cases have also been reported. The onset of SM tends to occur during adolescence or early adulthood, and there is no evidence of any predilection as to gender or ethnicity. Clinically, the lesions appear as smooth, round, skin-coloured to yellowish papules or nodules, especially in the axillae, as in this patient (Fig. 3). The cyst may exude an oily or creamy fluid when punctured. Cysts range in diameter from a few millimetres to several centimetres, and are typically located in areas where the pilosebaceous apparatus is well developed (1, 2). Cases of SM with predominantly acral and vulvar distribution have also been reported (3–5). The lesions are usually asymptomatic, but may become inflamed, rupture and heal with scarring, as was reported by our patient.

The histopathology of SM is characterized by the presence of empty cyst formed by a thin wall of stratified squamous epithelium without a granular layer. Sebaceous glands or atrophic sebaceous elements are presented adjacent to or within the cyst wall (1, 2, 6).

The origin of SM is uncertain, but it is probably derived from either transformed sebaceous ducts or cells differentiating toward the sebaceous duct because of comparable keratin expression (7). Immunohistochemical study reveals the presence of keratin 17 in upper layer cells in the cyst wall, whereas basal and parabasal layers are positive for keratin 14 (7). Treatment is not usually required, but in many cases the disease affects the psychological well-being of the patients. Surgical care, e.g. aspiration, excision, carbon dioxide laser therapy or liquid nitrogen cryotherapy, performed for cosmetic reasons, is sufficient in less severe cases. Treatment is not usually required, but in many cases the disease affects the psychological well-being of the patients. Surgical care, e.g. aspiration, excision, carbon dioxide laser therapy or liquid nitrogen cryotherapy, performed for cosmetic reasons, is sufficient in less severe cases. In the present case surgical excision of the eyelid cysts was performed successfully. At follow-up after 4 weeks, the patient reported an improvement in the quality of her sex life. The DLQI score decreased from 16 to 7 points. In inflammatory type of SM oral antibiotics and isotretinoin could be used, but this treatment is not fully effective (1, 2, 8).

**REFERENCES**


---

**Fig. 3.** Numerous smooth-surfaced, skin-coloured nodules in the axillary fold.