ID Design Press, Skopje, Republic of Macedonia Open Access Macedonian Journal of Medical Sciences. 2018 Jan 25; 6(1):108-109. Special Issue: Global Dermatology-2 https://doi.org/10.3889/oamjms.2018.032 elSSN: 1857-9655 Case Report



# **Idiopathic Scrotal Calcinosis – A Case Report**

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#### Abstract

Citation: Wollina U, Schönlebe J, França K, Tchernev G, Lotti T. Idiopathic Scrotal Calcinosis – A Case Report. Open Access Maced J Med Sci. 2018 Jan 25; 6(1):108-109. https://doi.org/10.3889/oamjms.2018.032

**Keywords:** Scrotum; Idiopathic calcinosis; Scrotal cysts. Surgery; Histopathology

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Received: 22-Aug-2017; Revised: 23-Sep-2017; Accepted: 24-Sep-2017; Online first: 10-Jan-2018

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Funding: This research did not receive any financial support

Competing Interests: The authors have declared that no

Idiopathic scrotal calcinosis is a rare disorder presenting with firm and painless nodules on the scrotal skin. The most common site is the frontal aspect of the scrotum whereas the dorsal aspect with the transition to the perineum is rarely involved. Surgery is the gold standard of treatment.

## Introduction

Inguinoscrotal disorders can be chronic disorders or emergencies like testicular trauma or torsions [1].

Table 1: Scrotal cysts and tumours

Entity	Remarks
Epidermal Cyst	Stratified lining epithelium, filled with keratin and debris
	May occur in Gardner syndrome
	Can lead to secondary calcinosis
	Cancerization is very rare
Cutaneous ciliated cyst	Rare benign lesion, very rare in males
	Female predominance (here on the legs)
Steatocystoma	Uncommon benign tumours of the pilosebaceous unit
multiplex	Stratified squamous epithelium without granular layer
	Filled with sebum
	Mutations in KRT17 gene
Eruptive vellus hair cyst	Stratified squamous epithelium with granular layer
	Multiple vellus hair shafts inside
Pilomatricoma	Rarely on scrotal skin
	Firm nodules, mostly single tumours
	Islands of epithelial cells composed of ghost cells in the
	centre surrounded by basaloid cells
Idiopathic scrotal	No epithelial lining
calcinosis	· · · · · · · · · · · · · · · · · · ·
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Scrotal nodules and cysts are rare findings. If they are asymptomatic, the diagnostic delay may be for several years or even decades. Table 1 provides an overview of scrotal cysts and tumours [2].

We report on a rare case of extensive idiopathic scrotal calcinosis treated surgically.

### Case report

A 46-year-old male patient presented with asymptomatic nodules of the scrotal skin for diagnosis and treatment. He reported the slow development of multiple lesions within the last ten years. He was otherwise healthy and did not have any medications or allergies.

On examination, we observed more than 30

firm subcutaneous cysts of variable size attached to the scrotal skin. On palpation, they were firm but painless. Their size varied form 3 mm to 4 cm (Fig. 1). Inguinal lymph nodes were impalpable. We performed surgical excision in general anaesthesia.



Figure 1: Multiple scrotal tumours – idiopathic scrotal calcinosis of the anterior aspect of the scrotum

The tumours were subjected to histopathological examination. On examination, pseudocystic formations with a fibrotic tissue around calcium deposits of variable size could be seen. There was no epithelial lining (Fig 2).

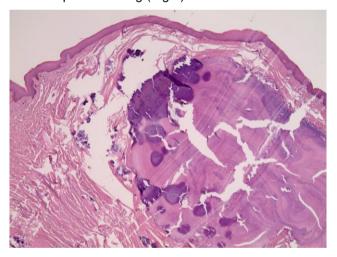


Figure 2: Histopathology of idiopathic scrotal calcinosis with coarse calcifications (hematoxylin-eosin x 10)

Healing was unremarkable. The patient was discharged on the second day after surgery.

#### **Discussion**

Firm nodules of the scrotal skin are rare. They can arise from pre-existing cysts like sebaceous cysts or steatocystoma multiplex or develop de novo. The latter is designated idiopathic scrotal calcinosis. The

major difference to calcified cysts is the complete absence of a lining epithelium [3]. Surgery is the treatment of choice.

The senior author of this paper (UW) noted during his decades of experience in clinical dermatology that idiopathic calcinosis and scrotal cysts are mainly localised an the anterior aspect of scrotal skin. Scrotal skin is a product of cloacal membrane ectoderm forming the labioscrotal folds [4].

There are some differential diagnoses to idiopathic scrotal calcinosis (Table 1). Multiple epidermal cysts of the scrotum [5][6][7], sebaceous cysts [8], steatocystoma multiplex [9]. Larger cysts need surgery; smaller ones can be subjected to laser therapy with either carbon dioxide or diode laser [10][11][12]. A linear nick with a radiofrequency electrode works well in enucleating the cysts intact as long as they are not melded together with the surrounding tissue [12].

#### References

- 1. Guerra L, Leonard M. Inguinoscrotal pathology. Can Urol Assoc J. 2017; 11(1-2Suppl1): S41-S46.
- 2. Redondo Martínez E, Rey López A, Sánchez Lobo V. Surgical pathology of the scrotum. An analysis of a series of 56 cases. Arch Esp Urol. 1999; 52(1):11-6. PMid:10101882
- 3. Killedar MM, Shivani AA, Shinde U. Idiopathic scrotal calcinosis. Indian J Surg. 2016; 78(4):329-30. <a href="https://doi.org/10.1007/s12262-016-1463-4">https://doi.org/10.1007/s12262-016-1463-4</a> PMid:27574356 PMCid:PMC4987564
- 4. Yiee JH, Baskin LS. Penile embryology and anatomy. Scientific World Journal. 2010; 10:1174-9. https://doi.org/10.1100/tsw.2010.112 PMid:20602076
- 5. Prasad KK, Manjunath RD. Multiple epidermal cysts in the scrotum. Indian J Med Res. 2014; 140(2):318. PMid:25297369 PMCid:PMC4216510
- 6. Ząbkowski T, Wajszczuk M. Epidermoid cyst of the scrotum: a clinical case. Urol J. 2014; 11(3):1706-9. PMid:25015622
- 7. Solanki A, Narang S, Kathpalia R, Goel A. Scrotal calcinosis: pathogenetic link with epidermal cyst. BMJ Case Rep. 2015; 2015. pii: bcr2015211163.
- 8. Angus W, Mistry R, Floyd MS Jr, Machin DG. Multiple large infected scrotal sebaceous cysts masking Fournier's gangrene in a 32-year-old man. BMJ Case Rep. 2012; 2012. pii: bcr1120115253.
- 9. Rahman MH, Islam MS, Ansari NP. Atypical steatocystoma multiplex with calcification. SRN Dermatol. 2011; 2011:381901. https://doi.org/10.5402/2011/381901
- 10. Bakkour W, Madan V. Carbon dioxide laser perforation and extirpation of steatocystoma multiplex. Dermatol Surg. 2014; 40(6):658-62. PMid:24852470
- 11. Wollina U. Three hundred patients treated with ultrapulsed 980 nm diode laser for skin disorders. Indian J Dermatol. 2016; 61(5):540-4. https://doi.org/10.4103/0019-5154.190111 PMid:27688445 PMCid:PMC5029241
- 12. Kamra HT, Gadgil PA, Ovhal AG, Narkhede RR. Steatocystoma multiplex-a rare genetic disorder: a case report and review of the literature. J Clin Diagn Res. 2013; 7(1):166-8. PMid:23449619 PMCid:PMC3576779