Primary anorectal melanoma is a rare condition accounting for 0.2-3% of all melanomas and approximately 1% of cancers afflicting this particular area.

The mean survival time after diagnosis is 24 months and only 10% of all patients will be alive at 5 years. Nearly 60% of patients present with metastatic disease.

The most frequent presentation symptom is rectal bleeding followed by change in defecatory habits.

Albeit the large majority of anorectal melanomas have evident gross or microscopic pigmentation there are 30% of which are amelanotic.

Currently abdomino-perineal resection with bilateral inguinal lymphadenectomy remains the first option procedure especially in large or obstructing lesions.

**CASE DESCRIPTION**

A 77-year old female living in a nursing home with Alzheimer’s, diabetes and hypertension was referred to the Gastroenterology Department for a large prolapsed anorectal mass in December of 2007.

Three months previously the patient had undergone a left-sided colonoscopy performed outside the hospital after persistent rectal bleeding had ensued. A sessile...
polyp with 1 cm was then registered in the distal rectum.

Our physical examination identified a polypoid, prolapsed, pedunculated and mobile mass with white coloured areas alternating with necrotic ones, measuring 4-5 cm in diameter.

Laboratory data was: hematocrit 29.2%; platelet count 367.6 x10^3/µl; INR 1.09; aPTT 24.2 seconds.

Total colonoscopy did not detect further lesions along the remaining colon.

The lesion was almost completely excised by fragmentation using a large polypectomy snare (Fig. 1a, 1b).

Histological findings were consistent with malignant melanoma (Fig. 2) and immunohistochemical stain was positive for S100 and HMB45 confirming the diagnosis.

Accordingly the patient was referred to a specialized oncology centre where abdomino-perineal resection was performed in January of 2008 showing no involvement of regional lymph nodes. Six months after surgery liver metastasis were diagnosed and the patient eventually died one month later.

The authors emphasize the rarity of this condition and the rapid tumour growth in a very short period of time.

BIBLIOGRAFIA