

REFERENCES

1. Shalhoub J, Naughton P, Lau N, Tsang JS, Kelly CJ, Leahy AL, et al. Concurrent colorectal malignancy and abdominal aortic aneurysm: a multicentre experience and review of the literature. *Eur J Vasc Endovasc Surg.* 2009;37:544-56.
2. Eliescu A, Bratucu E. The management of colon cancer in case of coexistence with an abdominal aortic aneurysm. *Chirurgia.* 2012;107:785-90.
3. Minicozzi A, Veraldi GF, Sboarina A, Lombardo F, Osmani H, Scudo G, et al. One stage or two stage treatment of colorectal cancer associated to abdominal aortic aneurysm: morbidity and mortality. *Minerva Chir.* 2012;67:453-7.
4. Lin PH, Barshes NR, Albo D, Koungias P, Berger DH, Huynh TT, et al. Concomitant colorectal cancer and abdominal aortic aneurysm: evolution of treatment paradigm in the endovascular era. *J Am Coll Surg.* 2008;206:1065-73.
5. Bali C, Matsagas M, Harissis H, Lagos N, Kappas AM. Management of synchronous abdominal aortic aneurysm and complicating colorectal cancer. *Vascular.* 2006;14:119-22.
6. Szilagyi DE, Elliott JP, Berguer R. Coincidental malignancy and abdominal aortic aneurysm: problems of management. *Arch Surg.* 1967;95:402-12.
7. Moll FL, Powell JT, Fredrich G, Verzini F, Haulon S, Waltham M, et al. Management of abdominal aortic aneurysms. Clinical practice guidelines of the European Society for Vascular Surgery. *Eur J Vasc Endovasc Surg.* 2011;41:S13.
8. Baxter NN, Noel AA, Cherry K, Wolff BG. Management of patients with colorectal cancer and concomitant abdominal aortic aneurysm. *Dis Colon Rectum.* 2002;45:165-70.
9. Porcellini M, Nastro P, Bracale U, Brearley S, Giordano P. Endovascular versus open surgical repair of abdominal aortic aneurysm with concomitant malignancy. *J Vasc Surg.* 2007;46:16-23.
10. Amato B, Esposito G, Serra R, Compagna R, Vigliotti G, Bianco T, et al. One-step mini-invasive treatment of abdominal aortic-iliac aneurysm associated with colo-rectal cancer. *Int J Surg.* 2014;12 Suppl. 2:S193-6.

Irene M. López Arquillo*, Jorge Vidal Rey,
Jose Manuel Encisa de Sá

Servicio de Angiología y Cirugía Vascular, Hospital Álvaro
Cunqueiro, Vigo, Pontevedra, Spain

*Corresponding author.

E-mail address: irenearquillo@hotmail.com

(I.M. López Arquillo).

2173-5077/

© 2016 AEC. Published by Elsevier España, S.L.U. All rights reserved.

A Case Report of Penile Metastases From Rectal Carcinoma[☆]



Presentación de un caso de metástasis en pene de carcinoma rectal

Metastatic disease of the penis is a rare occurrence. In a large series of autopsies performed in the Royal London Hospital on 623 patients who died due to secondary genitourinary tract malignancies, only 5 were located in the penis.¹ This clinical condition represents an advanced stage of the primary neoplasm and the overall outcome is generally poor. The most frequent sites of the primary tumor are bladder (34.7%), prostate (29.8%), recto-sigmoid colon (15.7%), and kidney (6.5%).² Few cases of penile metastasis from colorectal cancer have been reported.³⁻⁶ We present a case of penile metastasis from rectal carcinoma. Written informed consent for scientific use of the images was obtained from the patient.

A 70-year-old man was admitted to the hospital with 4 months of penis pain and voiding dysfunction. He had a past history, eight years before, of prostate carcinoma treated with radiotherapy, chemotherapy and hormonotherapy, with no evidence of disease recurrence in routine postoperative follow-up. Thirty months before he had been diagnosed of rectal cancer and underwent neoadjuvant chemotherapy,

robot-assisted laparoscopic abdominoperineal resection (Miles' procedure), colostomy and adjuvant chemotherapy. Radiotherapy was not administered due to the previous high-dose pelvic radiotherapy treatment that had been given for prostate carcinoma. Pathological examination of the rectal cancer specimen revealed a moderately differentiated adenocarcinoma infiltrating the perirectal fat (pT3), with a radial margin of 0.3 cm and perineural and vascular. All the 20 lymph nodes removed were negative for metastasis (N0).

Physical examination revealed a painful induration along the shaft of the penis and ulcers on the glans. Biochemical blood analysis evidenced a progressively rising serum CEA levels from 2.3 ng/mL to 10.9 ng/mL over the last 8 months. Penile ultrasound showed heterogeneous masses in both corpora cavernosa with interruption of the tunica albuginea. Additionally, three hypochoic urethral masses were detected, which produced dilation of the proximal urethra. Ultrasound scan findings suggested penile metastases. A suprapubic catheter was inserted into the urinary bladder.

[☆] Please cite this article as: Gómez-de la Fuente F-J, Martínez-Rodríguez I, Alonso-Martín J, Jiménez-Bonilla J, Banzo I. Presentación de un caso de metástasis en pene de carcinoma rectal. *Cir Esp.* 2017;95:116-118.



Fig. 1 – ^{18}F -FDG PET/CT scan. Selected images show increased FDG metabolism in the penile and glans metastases (arrows) and in multiple bone metastases.

A computerized tomography showed postsurgical changes in the pelvis and no other findings. A whole body ^{18}F -FDG PET/CT scan was performed for restaging. Images showed hypermetabolic lesions in the proximal shaft of the penis, distal penis and glans. Furthermore, multiple hypermetabolic bone metastases were detected in the pelvis. Local tumor recurrence was not observed (Fig. 1). Pelvic magnetic resonance imaging showed multiple low-signal-intensity lesions on T1-weighted sequences, involving the corpora cavernosa and the glans. After gadolinium administration, the lesions showed peripheral enhancement (Fig. 2). Pelvic bone metastases were also confirmed. Biopsies of the urethra, glans and corpora cavernosa revealed infiltration of adenocarcinoma. Immunohistochemical staining of the tumor cells was positive for cytokeratine 20, CDX2 and racemase, and negative for cytokeratin 7 and PSA. These findings supported the diagnosis of metastatic lesions from rectal cancer. Surgery, radiotherapy and chemotherapy would have been valid therapeutic options in such a case. Considering the previous pelvic radiotherapy administered for the prostatic carcinoma, the patient underwent chemotherapy with FOLFIRI® (folinic acid, fluorouracil and irinotecan) plus Bevacizumab. Twelve months after the diagnosis,

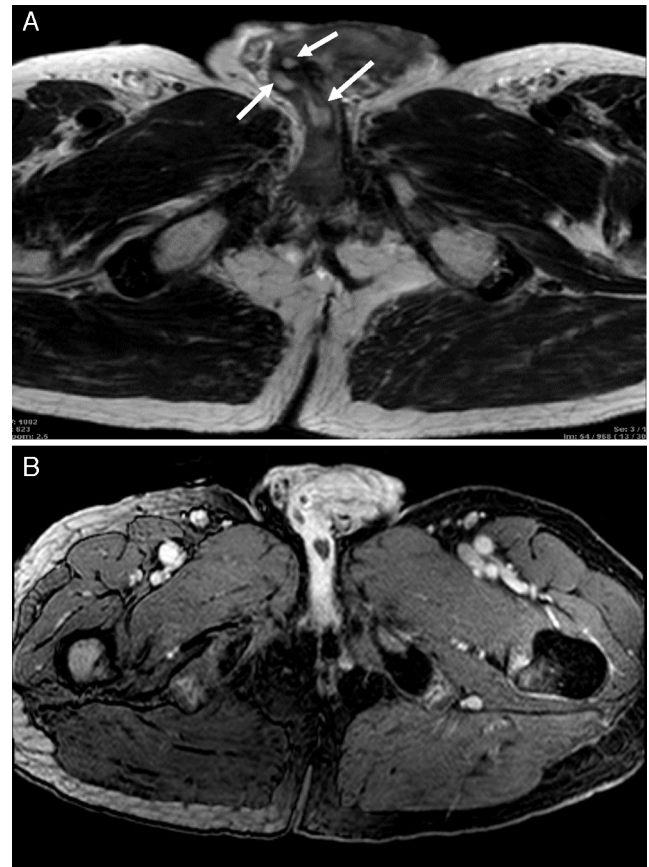


Fig. 2 – Penile magnetic resonance image. T1-weighted imaging shows lesions in the corpora cavernosa and glans (arrows) (A). After gadolinium administration, penile lesions shows peripheral enhancement (B).

the patient is still undergoing treatment with minor side effects.

Oncologic male patients can seek medical attention for penile complaints. Most of them come from minor benign problems, but may mask a serious underlying health condition. The most common clinical manifestations of penile metastasis are priapism, penile or perineal pain, and voiding disturbances.⁴ Despite its rich vascularization, the penis is rarely involved in metastatic spread of tumors located in other organs. In 1956, Paquin and Roland⁷ described the possible mechanisms of tumor spread to the penis as follows: direct extension, retrograde venous route, retrograde lymphatic route, direct tumor extension into branches of the hypogastric arterial pathway, metastatic tumor emboli from secondary tumor deposits in the lungs, tertiary embolism from liver metastasis producing metastasis to the lung and subsequent penis tumor emboli, and spread by instrumentation (iatrogenic).

The management of patients with serial CEA elevation after resection of rectal cancer includes physical examination, colonoscopy, and chest, abdominal and pelvic computerized tomography. Recently, the National Comprehensive Cancer Network guideline for rectal cancer, version 1.2016, took into consideration the use of ^{18}F -FDG PET/CT scan in patients with rising serum CEA levels after curative resection to detect tumor recurrence.⁸ Giacomobono et al.⁹ reported that ^{18}F -FDG

PET/CT allowed the identification of distant metastasis in patients with unexplained CEA rise after curative surgery of colorectal cancer. In a meta-analysis on the diagnostic performance of ^{18}F -FDG PET or PET/CT in patients with elevated CEA serum levels, Lu et al.¹⁰ found a sensitivity and specificity of 94.1% and 77.2%, respectively of ^{18}F -FDG PET/CT in the detection of tumor recurrence.

Diagnosis of penile metastasis is usually performed by biopsy or fine-needle aspiration. Penile metastasis represents a spread from the primary tumor. Non-invasive imaging methods are performed to determine the extent of the disease.^{4,6} In the case presented here, penile ultrasound was the first procedure to identify tumor involvement of the penis and the urethra. In addition to penile metastases, ^{18}F -FDG PET/CT scan revealed unknown bone metastatic spread to pelvis, not detected by computerized tomography, but confirmed by magnetic resonance imaging.

Conflict of Interest

They have not received support in the form of scholarships for study.

The information of the manuscript has not been previously presented at a conference.

REFERENCES

- Bates AW, Baithun SI. Secondary tumors of the penis. *J R Soc Med.* 2002;95:162-3.
- Hizli F, Berkmen F. Penile metastasis from other malignancies. A study of ten cases and review of the literature. *Urol Int.* 2006;76:118-21.
- Chaux A, Amin M, Cubilla AL, Young RH. Metastatic tumors to the penis: a report of 17 cases and review of the literature. *Int J Surg Pathol.* 2011;19:597-606.
- Mearini L, Colella R, Zucchi A, Nunzi E, Porrozzzi C, Porena M. A review of penile metastasis. *Oncol Rev.* 2012;6:80-7.
- Zhang K, Da J, Yao HJ, Zheng DC, Cai ZK, Jiang YK, et al. Metastatic tumors of the penis. A report of 8 cases and review of the literature. *Medicine (Baltimore).* 2015;94:e132.
- Seo HS, Kim ES, Kim S, Im SJ, Park YH, Lee JH, et al. A case of urethral metastasis from sigmoid colon cancer diagnostically and prognostically indicated by ^{18}F -FDG PET/CT. *Nucl Med Mol Imaging.* 2011;45:319-23.
- Paquin AJ, Roland SI. Secondary carcinoma of the penis. A review of the literature and a report of nine new cases. *Cancer.* 1956;9:626-32.
- NCCN.org [database on the Internet]. New York: National Comprehensive Cancer Network, Inc.; 2015. Available from: <http://www.NCCN.org/> [actualized 11.04.15, cited 22.02.16]
- Giacomobono S, Gallicchio R, Capacchione D, Nardelli A, Gattozzi D, Lettini G, et al. ^{18}F -FDG PET/CT in the assessment of patients with unexplained CEA rise after surgical curative resection for colorectal cancer. *Int J Colorectal Dis.* 2013;28:1699-705.
- Lu YY, Chen JH, Chien CR, Chen WT, Tsai SC, Lin WY, et al. Use of FDG-PET or PET/CT to detect recurrent colorectal cancer in patients with elevated CEA: a systematic review and meta-analysis. *Int J Colorectal Dis.* 2013;28:1039-47.

Francisco-Javier Gómez-de la Fuente,^a
Isabel Martínez-Rodríguez,^a Joaquín Alonso-Martín,^b
Julio Jiménez-Bonilla,^a Ignacio Banzo^{a,*}

^aServicio de Medicina Nuclear, Grupo de Investigación Imagen Molecular IDIVAL, H. U. Marqués de Valdecilla, Universidad de Cantabria, Santander, Spain

^bServicio de Cirugía General y Digestiva, Unidad de Cirugía Colorrectal, H. U. Marqués de Valdecilla, Universidad de Cantabria, Santander, Spain

*Corresponding author.

E-mail address: mnumbj@humv.es (I. Banzo).

2173-5077/

© 2016 AEC. Published by Elsevier España, S.L.U. All rights reserved.

Surgical Management of a Complete Section of the Oesophagus During Total Thyroidectomy[☆]



Tratamiento de una sección completa esofágica ocasionada en el curso de una tiroidectomía total

Oesophageal perforation or division in the course of thyroidectomy is an extremely uncommon but potentially serious complication that requires complex treatment. Only 7 cases have been published to date,¹⁻⁶ and none of them

reports a circumferential oesophageal lesion, as the case we present.

A 62-year-old woman with no relevant medical history had undergone total thyroidectomy due to multinodular goitre

[☆] Please cite this article as: Maupoey Ibáñez J, Ballester Pla N, García-Domínguez R, Vaqué Urbaneja J, Mingol Navarro F. Tratamiento de una sección completa esofágica ocasionada en el curso de una tiroidectomía total. *Cir Esp.* 2017;95:118-120.