Rectosigmoid adenocarcinoma revealed by metastatic anal fistula. The visible part of the iceberg: a report of two cases with literature review

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Abstract

Colonic adenocarcinoma revealed by metastatic anorectal fistula is rare, with few cases in the literature. Such lesions can be taken for the more common manifestation of a benign perianal abscess or fistula. Once diagnosed, the management of such conditions remains controversial. We herein report two cases with perianal fistula that were subsequently found to have developed perianal adenocarcinoma on biopsy. Further colonic investigation revealed a rectosigmoid adenocarcinoma. Histology and immunohistochemical staining was identical in both primary and metastatic tumors. Preoperative chemoradiation with further rectal low anterior resection and local excision of metastatic anal fistula was performed. There is no recurrence after 3 years of follow-up.

Keywords: Rectosigmoid cancer, Anal fistula, Implantation metastasis, Abdominoperineal resection

Background

Distal implantation by a proximal adenocarcinoma into an anal fistula remains a rare condition. It has been previously reported that exfoliated colorectal cancer cells can implant in distal locations such as the staple lines [1] and hemorrhoidectomy wounds [2] resulting in tumor recurrence. The same pathological mechanism can be expected in a case of colorectal adenocarcinoma seeding into anal fistula. Cancer implantation is usually suspected when carcinoma originating in an anal fistula has a similarity of histological findings between the primary colorectal cancer and the metastatic lesion [3]. Furthermore, the fistula must be present with a synchronous colorectal cancer. The diagnosis of such conditions remains difficult mainly when the primary tumor is unknown and the chronic anal fistula seems to be benign. More challenging is the management of such perineal tumors, as the literature is divided between surgeons who prefer radical abdominoperineal resection (APR) [4-17] and others who opt for sphincter-sparing surgery [18-26].

We herein present two cases of patients diagnosed with rectosigmoid adenocarcinoma revealed by metastatic perineal fistula, and treated with neoadjuvant chemotherapy followed by low anterior resection and local excision of the perineal tumor.

Case 1

A 55-year-old man presented with a 10-year history of recurrent anal fistula. He had no history of inflammatory bowel disease. Physical examination revealed several external openings, in the right and left perianal regions. The internal opening was located posteriorly in proctoscopy. The patient underwent fistulectomy with seton placement in fistula tracts. The pathology of the fistula excision did not show any malignancy. After 3 months of follow-up, a discharging nodule appeared in the fistula tract wound. Rectal examination did not reveal any abnormalities in the anal canal or the rectum. The nodule was biopsied, and adenocarcinoma was found on histology review. Further colonic investigation revealed a rectosigmoid tumor.

Histology of both the primary tumor and the perianal nodule revealed a well-differentiated adenocarcinoma with cytokeratin 20 (CK20)+ and cytokeratin 7 (CK7)-
staining. Magnetic resonance imaging (MRI) of the pelvis (Figure 1) and endoanal ultrasound (Figure 2) showed a perianal mass with anal sphincter involvement. Computer tomography (CT) did not show any distant metastasis.

The two modalities of the treatment (including APR or neoadjuvant chemoradiation followed by low rectal resection and local excision of perineal mass with the risk of recurrence and incontinence) were discussed with the
patient and informed consent was obtained. Eight weeks after a course of chemoradiation (45 Gy + 5 Fluorouracil), the patient underwent rectal anterior resection with colorectal anastomosis and local excision of the perianal mass which responded very well to the neoadjuvant therapy. Histopathological analysis of the excised nodule and rectosigmoid tumor revealed a well-differentiated adenocarcinoma as a primary tumor without lymph node involvement (T3N0M0), while the perianal mass showed a complete response following chemoradiotherapy. There is no recurrence after 3 years follow-up.

Case 2
A 68-year-old man presented to the emergency room with a perianal abscess. He had no history of anal fistula. Examination under anesthesia was performed confirming the perianal abscess with an internal opening in the anal dentate line, proving the origin was a cryptoglandular fistula. Simple drainage was performed. Because the cavity of the abscess had an atypically indurated border, biopsy was done. Histology revealed a well-differentiated adenocarcinoma. Further investigation, including rectosigmoidoscopy, MRI of the pelvis and a CT scan, showed a large carcinoma 26 cm from the anal verge and a perianal mass extending to the anal sphincter, which was confirmed by endoanal ultrasound.

No distant metastasis was noticed. Histology of the primary tumor was the same as the perianal mass. Immunohistochemistry for CK7 and CK20 was performed on tissues to distinguish colorectal adenocarcinoma from anal gland carcinoma. Both the colorectal cancer and the perianal tumor were CK7- and CK20+ (Figure 3). The modalities of the treatment (including APR or neoadjuvant chemoradiation followed by low rectal resection and local excision of the perineal mass with the risk of recurrence and incontinence) were discussed with the patient and informed consent was obtained. Preoperative chemoradiation (45 Gy + 5 Fluorouracil), was administered. Follow-up MRI 6 weeks after chemoradiation showed dramatic shrinkage of the perineal tumor. Surgery consisted of rectal anterior resection with colorectal anastomosis and local excision of the perianal mass. Pathological results of the rectosigmoid tumor revealed well-differentiated adenocarcinoma without lymph node involvement (T2N0M0). The perianal mass was completely sterilized. After 3 years follow-up there is no recurrence.

Discussion
Cancer of the colon and rectum can be spread by direct extension, transperitoneal, lymphatic, and hematogenous spread. Spreading by implantation has been also described [2]. In 1907, Ryall reported that many solid cancers were implanted in the operative field, and described the

![Figure 3 Histology of the perineal fistula. a) Glandular proliferation (circle) surmounted by a squamous epithelium (arrow). b) Adenocarcinoma with necrosis. c) Immunohistochemical staining immunoreactivity with anti-CK20.](image-url)
Table 1 Reported cases of implantation metastasis from colorectal adenocarcinoma into a fistula in ano

| Author                        | Year | Age | Gender | Location of the primary tumor | Duration of anal fistula | Stage | Histology | ADK Different | Operation | Follow up without recurrence |
|-------------------------------|------|-----|--------|--------------------------------|--------------------------|-------|-----------|---------------|-----------|-----------------------------|
| Guiss A [4]                   | 1954 | 47  | M      | S                              | 2 m                      | Dukes A | Mod       |               | APR       | 1 y  2 m                     |
| Killingback et al. [5]         | 1965 | 63  | M      | S                              | 8 y                      | Dukes A | Well      |               | APR       | NA                          |
| Parnes [6]                    | 1976 | 47  | M      | S                              | 3 m                      | Dukes B | Well      |               | APR       | 18 m                        |
| Rollinson and Dundas [7]      | 1984 | 65  | M      | Rs                             | 20 y                     | NA     | Well      |               | APR       | 10 m                        |
| Ueta et al. [8]               | 1991 | 66  | F      | S                              | 44                       | Dukes B | Well      |               | APR       | 6 m                         |
| Thomas and Thompson [9]       | 1992 | 68  | M      | S                              | 1 y                      | Dukes B | Mod       |               | APR       | NA                          |
| Tohira et al. (Japanese) [10] | 1998 | 75  | M      | UR                             | 40 y                     | Dukes B | Well      |               | APR       | 1 y                         |
| Isbister [18]                 | 2000 | 47  | M      | Rs                             | 2 y M                    | Dukes C | Mod       | 1- LR         | NA        |                             |
| Isbister [18]                 | 2000 | 39  | M      | S                              | N 1 y M                  | NA     | NA        | Refused APR   | NA        |                             |
| Tokuhara et al. [11]          | 2001 | 69  | M      | S                              | 5 y                      | Dukes B | Mod       |               | APR       | 1 y                         |
| Yoshimura et al. [12]         | 2001 | 59  | M      | Rs                             | 29 y                     | Dukes B | Mod       |               | APR       | 3 y 7 m                     |
| Shinohara et al. [19]         | 2001 | 36  | M      | MR                             | 16 y                     | Dukes C | Mod       | AR + LR       | 6 m       |                             |
| Kouaklis et al. [13]          | 2002 | 75  | M      | S                              | 1 y                      | Dukes B | Mod       |               | APR       | NA                          |
| Yagihashi et al. [14]         | 2002 | 50  | M      | S                              | NA                       | Dukes C | Well      | TPE            | 3 y 8 m   |                             |
| Shimoyama et al. [15]         | 2003 | 61  | M      | Rs                             | 5 y                      | Dukes C | Mod       |               | APR       | 5 y                         |
| Hyman and Kida [16]           | 2003 | 66  | M      | S                              | 15 y                     | Dukes B | Mod       | CK7-/CK20     | APR       | 1 y                         |
| Zbar and Shenoy [20]          | 2004 | 72  | M      | S                              | 4 y                      | NA     | S + LR + adj CH RTH | 1 y 2 m   |                             |
| Gupta et al. [21]             | 2005 | 44  | M      | Left colon                     | 1 m                      | Dukes C | Mod       | LH=LR          | 3 y       |                             |
| Hamada et al. [23]            | 2005 | 53  | M      | S                              | 7 y                      | Dukes B | Well CK7-/CK20 | AR + LR       | 1 y       |                             |
| Ishiyama et al. [24]          | 2006 | 53  | M      | UR                             | 20 y                     | Dukes C | Mod       | AR + LR       | 10 m died from carcinomatosis |
| Sandiford et al. [25]         | 2006 | 72  | M      | Rs                             | 2 y                      | Dukes B | Mod       | S + LR + adj Ch R | 14 m     |                             |
| Gravante et al. [17]          | 2008 | 64  | M      | Left colon                     | NA                      | Dukes A | Mod CK7-/CK20 | 1-LH          | 1 y       |                             |
| Wakatsuki et al. [26]         | 2008 | 57  | M      | Rs                             | 7 y                      | Dukes B | Mod CK7-/CK20 | 1-AR          | 3y7m      |                             |
| Our report                    | 2012 | 68  | M      | Rs                             | 2 m                      | Dukes B | Well CK7-/CK20 | Neoadju chemoradia AR + LR | 3 y       |                             |
phenomena as 'cancer infection' [27]. Guiss reported the first case of cancer implantation into an anal fistula in 1954. Since then, a few other cases have been reported in the literature and the concept of cancer implantation has become widely accepted. We found 23 articles describing 24 patients with metastatic anal fistula arising from colorectal carcinoma [4-26]. Details of the cases are summarized in Table 1, along with the current patients. The diagnosis of such conditions remains difficult, mainly when the primary tumor is undiagnosed and the chronic anal fistula seems to be benign. It is, therefore, the change in behavior of the anal fistula, such as the appearance of a nodule or perianal abscess with indurated borders as is described in our patients that should raise the red flag and awareness of possible malignancy, making biopsy and further colonic investigation mandatory. Histological differentiation and immunohistochemical staining must show the same characteristic in both tumors. More specifically, immunohistochemistry of CK7 and CK20 remains the examination of choice to distinguish between colonic tumor tissues and tumors originating from the anal glands [16]. Ramalingam et al. have previously described that anal glands were strongly immunoreactive to antibodies against CK7 but not to CK20 [28]. In our case, tumor cells of both lesions were positive for CK20 and negative for CK7 antibodies. We considered the tumor of the anal fistula as metastasis of the rectosigmoid cancer in both cases.

Controversies remain in the management of metastatic anal fistula. It is difficult to define the best treatment from the literature, as some authors have performed APR, while others have chosen local resection.

After discussing the modality of treatment (either APR or neoadjuvant chemoradiation following by anterior rectal resection and local excision), emphasizing the risk and the benefit of each approach, in particular risk of recurrence and incontinence) with our patients, we opted for preoperative chemoradiation (45 Gy), followed by anterior resection of the rectum and local excision of perianal metastasis after 6 weeks. Our choice was guided by the age of patients, differentiation and immunohistochemical characteristics of the primary and metastatic tumors, evidence of sphincter involvement on MRI and endoanal ultrasound and the absence of distant metastasis on abdominal CT. Furthermore, perineal masses in both cases shrank dramatically after chemoradiation. Consistent with that, the pathology report showed a complete response to chemoradiotherapy in both cases. The primary tumors were without lymph node metastases. Chemoradiation has been indicated as an adjuvant therapy in some previous reports [17,20,25], but not as a neoadjuvant treatment. The benefit of neoadjuvant versus adjuvant chemoradiation is still unclear. Based on a literature review of colorectal cancers implanting in a fistula track, patients undergoing local resection versus APR showed similar outcomes in terms of local recurrence. Nevertheless, it is difficult to make any concrete conclusion about the best approach, as the length of follow-up of reported cases is short.

**Conclusion**

It is strongly recommended to be aware of any unusual changes in the clinical features of the fistula in ano either before or after surgery (such as increased discharge, bloody discharge, nodule formation, uncommonly slow healing or scar inductions) and not to hesitate to perform biopsy in order to exclude any malignancies that may be the visible part of the iceberg. Sphincter-saving surgery is an option that should be discussed with the patient.

**Consent**

Written informed consent was obtained from the patients for publication of this case report and any accompanying images. A copy of the written consent is available for review by the Editor-in-Chief of this journal.

**Competing interests**

The authors declare that they have no competing interests.

**Authors’ contributions**

BEB has made substantial contributions to conception, bibliography and drafting the manuscript. AS has been involved in drafting the manuscript. CL has been involved in bibliography research and conception of pathology figures. IM and KM have been involved in or revising it critically for important intellectual content. KA has given final approval of the version to be published. All authors read and approved the final manuscript.

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