Rectal Carcinoma Metastasis to Oral Cavity: A Rare Presentation

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ABSTRACT

Colorectal cancer is 3rd most common cancer all over the world. Metastasis in colorectal cancer is common and seen in approximately 30% cases. Metastatic pattern in colorectal cancer depends upon the primary tumour site and histological subtypes. Liver is the most common site for metastasis. Oral cavity is very rare site for metastasis. Here, we report a case of 52 year old male diagnosed with rectal adenocarcinoma with liver and lung metastases and presenting with oral ulcer. Histopathological examination of oral ulcer revealed metastatic adenocarcinoma. We are presenting it as a rare case report and reviewing similar published cases.

Keywords: Oral Ulcer, Colorectal Cancer, Metastasis

Introduction

Worldwide, colorectal cancers (CRC) are 3rd most common cancer. Metastasis is common in CRC and indicates grim prognosis with low survival rates. CRC most commonly metastasize to the liver, lung and peritoneum but may also spread to lymph nodes, bone and other sites. CRC metastasis to soft tissue is very rare presentation. Metastasis to oral cavity from any primary tumour in the body is rare, comprising less than 1% of oral cavity tumours. Most common source are tumours of lung, breast and kidney. The jawbones are twice as common for metastasis than the oral mucosa.

Thus, metastasis to oral cavity soft tissue is rare presentation and even rarer is the CRC metastasis to oral cavity. Here we present one such case with rectal adenocarcinoma metastasis to gingiva.

Case

A 52 year old male patient presented with altered bowel symptoms and mucus in stool, since 1 month, in May 2017. He also complained of decreased appetite and weight loss. His general condition was good and per rectal examination showed a proliferative growth in rectum, 2 cm from anal verge. Computed Tomography (CT) abdomen and pelvis reported rectal carcinoma with liver metastases and metastatic lymphadenopathy involving perirectal, retroperitoneal, pre-aortic and para-aortic lymph nodes. Rectal biopsy was reported as adenocarcinoma. He was immediately started on chemotherapy FOLFOX-4 (Oxaliplatin + Folinic Acid + 5-Fluorouracil). In August 2017, after completing six cycles of FOLFOX-4, he complained of cough. CT scan was advised for reassessment and revealed progression of disease with new lung metastases. Patient clinical condition and prognosis was explained to the relatives and chemotherapy regimen was changed to FOLFIRI (Irinotecan + Folinic Acid + 5-Fluorouracil). CT scan after six cycles of FOLFIRI revealed partial response to chemotherapy, so six more cycles of FOLFIRI were advised. His carcino-embryonic antigen (CEA) level was 10.96ng/ml. But, patient defaulted and came after 4 months in March 2018 with complaints of constipation and difficulty in passing urine. CT scan findings were suggestive of progression of disease and his CEA level rose to 362.75ng/ml. Chemotherapy was resumed with FOLFIRI. Six cycles of FOLFIRI were completed. He visited in June 2018 with lower abdomen pain and distension. Ascitic tapping was done. He also showed 1x1 cm oral ulcer at right upper alveolus near last molar. Oral ulcer biopsy was taken and was reported as metastatic adenocarcinoma (Fig 1). Immunohistochemistry was positive for CK20, CDX2 and negative for CK7 (Fig 2). CT scan revealed progression of diseases. In July 2018, patient succumbed to the disease and died at home.

Discussion

Metastases are common in CRC. Riihimaki et al studied 49096 CRC cases, and found metastases in 30.48% cases. Similarly, Hugen et al reported metastases in 28.9% cases. Metastasis in CRC can be synchronous or metachronous. Metastatic pattern in CRC varies with the primary tumour site and histological subtypes. The frequency of liver metastases does not differ between colon and rectal cancer patients. However, in comparison, colon cancer metastasizes more frequently to intra-abdominal sites and rectal cancer metastasizes more frequently to extra-
### Table 1: Reported cases of colorectal carcinoma metastasis to oral cavity.

| Author            | Age in yrs | Sex | Primary tumour | Interval between primary and oral lesion | Sy/ Me | Lesion | Site | Other metastasis                                                                 | Death after oral lesion |
|-------------------|------------|-----|----------------|------------------------------------------|--------|--------|------|---------------------------------------------------------------------------------|-------------------------|
| Lagha A et al\(^5\) | 46         | M   | ACR            | 12 mths                                   | Me     | Mass   | Gingiva | Liver, lung                                                                     | 4 mths                  |
| Usman N et al\(^6\) | 60         | M   | ACR            | 24 mths                                   | Me     | Mass   | Buccal mucosa, GB sulcus, gingiva | LN, bone, adrenals, spinal cord | 4-6 mths             |
| Almangush A et al\(^7\) | 61        | F   | ACR            | 15 mths                                   | Me     | Mass   | GB sulcus | Liver, spine, skin                                                               | 3 mths                  |
| Gill KS et al\(^8\) | 57         | M   | ACR            | 24 mths                                   | Me     | Mass   | Tongue | LN, lung                                                              | NK                      |
| Singh T et al\(^9\)  | 42         | F   | ACR            | 38 mths                                   | Me     | Mass   | Floor   | Recurrence                            | 20 days                 |
| Pelissari C et al\(^10\) | 64      | F   | ACC            | 36 mths                                   | Me     | Mass   | Gingiva | No mets                                | 6 mths                  |
| Bhaskaran A et al\(^11\) | 92     | M   | ACRS           | NA                                        | Sy     | Ulcer  | Tongue | LN, liver, lung, brain, bone, retroperitoneum | NK                      |
| Yamauchi M et al\(^12\) | 50       | M   | ACR            | NA                                        | Sy     | Ulcer  | Gingiva | Lung, liver, spleen                                                               | NK                      |
| Baranović M et al\(^13\) | 78      | M   | ACR            | 18 mths                                   | Me     | Mass   | Gingiva | Liver                                  | 4 mths                  |
| Watanabe M et al\(^14\) | 64       | M   | ACR            | 84 mths                                   | Me     | Mass   | Gingiva | Lung, vertebral body                                                               | 2 mths                  |
| Ren QG et al\(^15\)  | 60         | M   | ACC            | 24 mths                                   | Me     | Mass   | Gingiva | Liver                                  | NK                      |
| Present case       | 52         | M   | ACR            | 14 mths                                   | Me     | Ulcer  | Gingiva | LN, liver, lung,                                                                  | 1 mth                   |

M: Male; F: Female; ACR: Adenocarcinoma rectum; ACC: Adenocarcinoma colon; ACRS: Adenocarcinoma rectosigmoid; Sy: Synchronous; Me: Metachronous; NA: Not applicable; NK: Not known; GB: Gingivo-buccal; LN: Lymph node

**Fig. 1:** Metastatic deposits of adenocarcinoma rectum in alveolar mucosa (H&E 40X).
abdominal sites. The higher rate of distant metastases in rectal cancer can be explained by the venous drainage of the rectum bypassing the liver and draining into the inferior vena cava. Adenocarcinoma more frequently metastasizes to the liver, while mucinous and signet ring cell carcinoma metastases more frequently to peritoneal surface.

The oral cavity is uncommon site for metastatic deposit for any primary tumour in the body. Metastasis to oral cavity is usually a late complication and indicates widely disseminated disease with poor prognosis. Lung is the most common primary in males and breast is most common primary in females for oral cavity metastasis. CRC metastasis to oral cavity is rare and when present, it is usually associated with multiple site metastases and thus carry unfavourable prognosis. Oral cavity metastasis may involve jaw bone or soft tissue. Metastasis is more common in jaw bone than oral cavity soft tissue. In the oral soft tissue, gingiva is the most common site because of its rich blood supply and chronically inflamed nature. It presents as ulcerated lesion or exophytic growth. Tongue is the second common site. Patients usually present with pain, bleeding, dysphagia and disfigurement. Possible mechanism for oral metastasis can be attributed to Batson’s venous plexus extending from the skull to the sacrum. This valve less system theoretically offers less resistance to the spread of tumour emboli.

With known history of primary tumour elsewhere, diagnosis becomes easy with histopathology for metastasis in oral cavity. But, in cases of occult primary, oral lesion could be the only complaint. In such cases immunohistochemistry is useful in making the diagnosis. CRC are positive for CK20, CDX2 and negative for CK7 on immunohistochemistry.

Chemotherapy is the treatment of choice for metastatic CRC. Surgical treatment is planned for patients with limited involvement of distant organs particularly, liver and lung. Mean survival time after diagnosis of metastasis in oral cavity is 6-8 months.

As the CRC metastasis to oral cavity soft tissue is rare presentation, there are very few published case reports. We found 11 such published cases with Google search engine. A brief review of 11 published cases and the present case is given in Table 1.

It is observed that, sixth and seventh decade is the most common age group affected. Men are more commonly affected than women. Rectum is the most common primary site. The minimum time interval between primary tumour and development of oral lesion is 1 year and maximum time interval is 7 years. Gingiva is the most common site for metastasis. Oral lesion mostly present as a growth and sometimes with ulcerated lesion. Metastases to oral cavity usually appear after the primary tumour. But in two cases, oral ulcer was the initial clinical presentation, which led to the diagnosis of primary tumour. Most of the cases show at least one other metastatic site along with oral metastases. Singh et al reported a case with no other metastases but with history of recurrence at presacral region. Pelissari et al reported a rare case with no history of recurrence or other site metastases. Ten patients died within six months of development of oral metastasis.

**Conclusion**

Colorectal carcinoma metastases to oral cavity soft tissue are rare presentation. Rectal adenocarcinoma show more tendencies to spread to oral cavity soft tissue than colon cancer. These lesions usually present as gingival growth,
but can present with an ulcerative lesion. These are usually accompanied with multiple site metastases, thus carrying poor prognosis with less than six months survival. So, metastasis should be considered in differential diagnosis for oral ulcer, particularly in patients with known primary elsewhere.

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