Case Report

Sweet syndrome as the leading symptom in the diagnosis of gastric cancer

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Abstract: Sweet syndrome is a neutrophilic infiltration of the papillary dermis, which may be associated with the presence of unknown malignancies, either haematological or solid tumours, in 1 out of 5 cases, being considered then as a paraneoplastic syndrome. We present the case of a male with a locally advanced gastric cancer whose final diagnosis was led by the prior debut of Sweet syndrome not explained by other causes.

Keywords: Acute neutrophilic dermatosis; dermis inflammatory infiltrate; gastric cancer; paraneoplastic syndrome; Sweet syndrome

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Introduction

Paraneoplastic syndromes are defined as those which may appear when tumour cell-released substances alter the correct functions of nearby cells or tissues (1). Their incidence is not well defined, although they are more frequent in hematologic cancers and, among solid cancers, in those patients suffering from breast cancer. Moreover, they can appear either at the beginning or during the course of the disease.

We here present an old patient treated for a locally-advanced gastric cancer after the development of a paraneoplastic Sweet syndrome which lead to the final diagnosis.

Case presentation

A 72-year-old patient was admitted to the Internal Medicine Department referring syncopal episodes which were associated with dizziness, diaphoresis, nausea without vomiting, hypotension and temperature up to 39 °C. These were accompanied by macular lesions in forearms, thorax and abdomen. He also reported a weight loss of 9 kilograms in a 2-month time.

As previous medical history, underline he suffered from heart failure NYHA grade III secondary to a heart attack treated with angioplasty and vascular stent, 5 years before the present episode.

At the physical examination, he presented erythematous-oedematous fixed lesions in superior limbs, thorax and abdomen, with an average diameter of 10 mm, and with defined non-confluent oedematous borders (Figure 1). No purpuric nor infiltrative characteristics were present, as well as absence of whitening with diascopy. Other dermatologic or mucosal lesions were not present.

During the study of this episode, several tests were conducted. Blood test revealed leucocytosis (17.8×10⁹/L, normal 3.8×10⁹–10.8×10⁹/L) with neutrophilia (15,400×10⁹/L, normal 1.8×10⁹–7.5×10⁹/L), as well as lymphopenia (950×10⁹/L, normal 1.5×10⁹–4×10⁹/L) and an increased reactive C protein (23.3 mg/dL, normal 0–5 mg/dL)

A punch from de affected skin was performed, with the result of interstitial diffuse dermatitis with polymorphonuclear and histiocytic cells and intense oedema, associating CD68 positive in the interstice (Figure 2). All compatible with acute neutrophilic dermatosis.
With the suspicion of this being secondary to a systemic illness, imaging test were conducted. CT scan showed focal thickening of the gastric wall with nearby celiac and retroperitoneal lymphadenopathies as well as subcentimetric pulmonary nodules in inferior right lobe of the lung.

Diagnostic gastroscopy was performed, where an ulcerated vegetative neoplasia at the anterior gastric body with extension to the fundus was biopsied.

After all the study, the patient was diagnosed of a metastatic adenocarcinoma of the stomach poorly differentiated HER2 positive by FISH; with an associated paraneoplastic acute neutrophilic dermatosis (Sweet syndrome).

Treatment with prednisone 30 mg a day was established, with complete response of the lesions after a month of treatment and normalization of the blood parameters (leucocytes $11,000 \times 10^9/L$, neutrophils $8,000 \times 10^9/L$, reactive C protein $2 \text{ mg/dL}$).

During the surveillance of the patient, previous to the beginning of the oncologic treatment, he presented to the Emergency Department referring instability when walking. Brain MRI was performed, noticing a cystic and solid lesion on the left cerebellar hemisphere. Glucocorticoids in combination with holocraneal radiotherapy were given.

**Figure 1** Clinical presentation at Emergency Department. Pink edematous erythematous papules not confluent with poorly defined borders with undulating surface with no secondary changes in surface, located in abdomen (A) and distal extremities (B).

**Figure 2** Pathological anatomy [hematoxylin-eosin staining, ×10 (A) and ×20 (B)]. Diffuse interstitial inflammatory infiltrate composed of neutrophils affecting the papillary dermis and the superficial portion of the reticular dermis. Outbreaks of leukocytoclastia without vasculitis and marked edema of the papillary dermis are noted.
with good management of symptoms.

However, few days later, the patient presented with high digestive haemorrhage which required blood transfusions and haemostatic radiotherapy to cope with it.

Finally, after all these complications, the patient had very poor performance status and was completely dependent on his caregiver, so best supportive care was chosen as the more suitable management plan.

At the time of the writing of this report, the patient was living at home with his caregiver with a good quality of life and had had no recurrence of his Sweet syndrome nor any other paraneoplastic disorder.

**Discussion**

Acute neutrophilic dermatosis or Sweet syndrome (2) often debuts as a papular rash or as indurated and painful erythematous plaques in face, limbs and thorax. Paraneoplastic Sweet syndrome accounts for the 20% of all the cases. From these, 85% appear accompanying haematological cancers (mostly acute myeloid leukaemia), and the other 15% associated with solid tumours, such adenocarcinomas from the breast, upper genitourinary tract and upper intestinal tract (3).

Its pathogenesis is by the moment not completely understood. Nevertheless, at least three factors have been proposed to trigger it: hypersensitivity reaction to cancer neoantigens and cytokine production (4); cytokine dysregulation such as granulocyte-colony stimulating factor (G-CSF) production increasing circulating neutrophils in peripheral blood (5); and genetic susceptibility, such as alterations in chromosome 3q (6).

Clinical manifestations include fever, poor general condition and joint and muscular pain, apart from the typical dermic lesions described before. In nearly 50% of the cases extra cutaneous disease can be present, affecting the liver, the kidneys or the lungs (as a reticular or nodular infiltration with pleural effusion) (7,8).

Blood test show increased glomerular sedimentation speed and reactive C protein parameters, as well as leucocytosis with neutrophils accounting for more than the 70% of them.

Under the microscope, a dense interstitial and perivascular neutrophilic infiltration in the papillary dermis is characteristic, with oedema and nuclear fragmentation of mature neutrophils, without associated vasculitis (9).

Definitive diagnosis of paraneoplastic syndrome must be made according to the modified Su-Liu criteria (7), which include major criteria (sudden appearance of painful erythematous plaque or nodule and dense neutrophilic dermic infiltration without vasculitis in the biopsy of a lesion) and minor criteria (fever >38 °C, glomerular sedimentation speed >20 mm/h, leucocytes >8×10⁹/L, neutrophils >70%, high reactive C protein, good response to glucocorticoid therapy and history of gastrointestinal infection, neoplasia or recent immunization).

Differential diagnosis should include infectious disorders, as bacterial sepsis may induce similar symptoms and dermic affection; and other dermatologic syndromes, such as other neutrophilic dermatoses (i.e., pyoderma gangrenosum or Behçet syndrome) or medium-vessel vasculitis, among others (8).

Treatment of Sweet syndrome is based on glucocorticoids, with a starting dose of prednisone 30 mg, followed by a progressive descendent dosing being the most recommended in the literature (7). If this may not work, second line treatments may be used, such as indomethacin, cyclosporine or dapsone, among others(8).

The majority of cases will respond satisfactorily to the treatment, although up to 20–30% of recurrences have been reported. These recurrences may be indicative of disease progression (8).

In conclusion, our patient presented prior to cancer diagnosis with a paraneoplastic Sweet syndrome, with complete response to glucocorticoid treatment. Hence, a recommendation of a complete study suspecting a systemic disease, mainly cancer, can be made when this or other related syndromes related to cancer appear, so that the diagnosis is not unnoticed and the prognosis of the patient may be improved.

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None.

**Footnote**

Conflicts of Interest: The authors have no conflicts of interest to declare.

**Informed Consent**: The patient and caregivers involved gave consent to publication of this manuscript and any accompanying images.

**References**