Radiation induced gastroparesis—case report and literature review

Pierre Annede¹²³, Caroline Prieux-Klotz³, Thomas Dubergé², Cyrus Chargari³, Olivier Gisserot¹, Jean-pierre de Jaureguiberry¹

¹HIA Sainte Anne, Oncology Department, Toulon, France; ²Centre de Radiothérapie St. Louis, Toulon, France; ³Radiotherapy Department, Gustave Roussy, Villejuif, France

Correspondence to: Pierre Annede, MD. Radiotherapy Unit, Gustave Roussy, 114 rue Edouard Vaillant, 94800, Villejuif, France. Email: pierreannede@gmail.com.

Abstract: Radiation induced gastroparesis as well as the other autonomic nervous system radiation induced neuropathies are poorly described in the literature. A case of gastroparesis associated with phrenic and recurrent laryngeal nerves paralysis was observed in a 69-year old patient. She was already treated two times by rachis radiotherapy in a context of breast cancer with bone metastases. Anatomical and chronological correlation of lesions, concomitant nerve damage in the radiation fields and elimination of the main differential diagnoses allowed us to link this case of gastroparesis with the background of radiotherapy. It confirms the role of vagus nerve lesion in radiation induced gastroparesis. This specific diagnosis led to a successful treatment and a quality of life improvement.

Keywords: Gastroparesis; radiotherapy; neuropathy

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Introduction
Therapeutic progress in metastatic cancers improves patient survival. During disease course, patient may benefit from several successive radiotherapy treatments. Increasingly, long-term and unusual radiation induced toxicities are described. Knowing these side effects leads to provide optimum medical care.

Gastroparesis is a syndrome caused by gastric stasis unrelated to mechanical obstruction. Main symptoms are: early satiety, nausea, vomiting, abdominal pain and bloating (1). It significantly affects quality of life (2), morbidity and mortality (3). The main etiologies are metabolic (diabetes =29%), post-surgical (13%), drugs and idiopathic (36%) (3). Other etiologies are poorly described, as Parkinson, amylose, paraneoplastic syndrome and radiotherapy.

Case presentation
A 69-year-old woman was admitted in our department for nausea and vomiting in a context of breast cancer with bone metastases. She described nausea, abdominal fullness feeling and dysphonia (bitonal voice). These symptoms get progressively worse. After few weeks, every food consumed induced early vomiting. Chest film (Figure 1) and CT scan showed left diaphragmatic cupola rise, symptomatic of left phrenic nerve paralysis. Ear, nose and throat (ENT) examination revealed a left vocal cord paralysis due to left recurrent laryngeal nerve lesion. Gastroscopy revealed a gastric bezoar. An upper gastrointestinal (GI) series test showed an aspect of gastroparesis with gastroesophageal reflux. On examination, her medical treatment wasn’t described as causing gastroparesis. She wasn’t diabetic and hadn’t undergone recent surgery. On the other hand she underwent two sequential rachis radiotherapy (D3–D6 then D8–D11), both at the dose of 30 Gray in 10 fractions, 2 years and 7 months ago respectively (Figure 2). Cerebral MRI, plasmatic neuronal antibody and lumbar puncture were normal. Eventually we concluded to a radiation induced gastroparesis. Treatment with Domperidone was initiated.
without benefit after 4 weeks. Second-line treatment by intravenous erythromycin 250×3/J during 7 days and then orally over 4 weeks allowed significant reduction of GI symptoms. The patient recovered a normal nutritional status.

**Discussion**

Only three cases of radiation induced gastroparesis have been reported in the literature, *(Table 1)* (4-6). There is to our knowledge no incidence data of autonomic nervous system radiation induced neuropathy. However, many clinical presentations of late radiation-induced neuropathies have been described.

Risk factors for developing radiation induced neurotoxicity are: concomitant radiotoxic treatments,
cardiovascular risk factors, length of nerve irradiated and dose per fraction.

In this case, first dose prescribed to the spine was 30 Gy in 10 fractions of 3 Gy corresponding to an equivalent 2 Gy dose (EQD2) equal to 37.5 Gy (with an alpha/beta ratio equal to 2). A second similar dose was prescribed two years later on a nearby area. Dosimetric reconstruction showed little theoretical radiation fields overlap. However, positional uncertainties as well as morphological changes of degenerative rachis may have increased the radiation fields' superposition.

It is conventionally admitted that late radiation induced neuropathy is caused by direct nerve damage (axons, myelin and Schwann cells) or indirect lesions (ischemia, peripheral fibrosis). In this case, the dose to the stomach muscular wall was insignificant. In addition, involvement of other concomitant nerve tracts (left phrenic nerve and recurrent) suggests that a left vagus nerve lesion was the cause of digestive symptoms.

Paraneoplastic syndrome remains the principal differential diagnosis in this context. Although its exact pathogenesis remains unknown, patients' autopsy shows neural ganglia-free stomachs, and gastric nerve fibers injured by chronic inflammation. Serological detection of anti-Hu antibodies suggests an autoimmune destruction of stomach neural plexus as a key mechanism (7). This tends to make us think that paraneoplastic gastroparesis is an intrinsic disease of the stomach. Conversely, our case was probably due to a vagus nerve lesion. These assumptions seem to be confirmed by practice. Indeed, the paraneoplastic gastroparesis cases are resistant to prokinetic drugs whereas in our case erythromycin was efficient.

Conclusions

Anatomical and chronological correlation of lesions, concomitant nerve tract damage in the radiation field and elimination of the main differential diagnosis allowed us to link this case of gastroparesis with the background of radiotherapy. This specific diagnosis led to a successful treatment and a quality of life improvement.

Acknowledgements

None.

Footnote

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

Table 1 Literature review of radiation induced gastroparesis

<table>
<thead>
<tr>
<th>Author</th>
<th>Age</th>
<th>Gender</th>
<th>Primary cancer</th>
<th>Radiation fields</th>
<th>Time to onset of gastroparesis</th>
<th>Treatment</th>
<th>Treatment effectiveness</th>
</tr>
</thead>
<tbody>
<tr>
<td>Petrasch et al.</td>
<td>16</td>
<td>Male</td>
<td>Seminoma</td>
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<td>Early; then late*</td>
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<td>Total for 7 years. Relapse effectively treated with cisapride</td>
</tr>
<tr>
<td>Sturm et al.</td>
<td>75</td>
<td>Female</td>
<td>Endometrial carcinoma</td>
<td>Adjuvant radiotherapy: 30 Gy curie therapy + 30 Gy external radiotherapy</td>
<td>Early</td>
<td>IV metoclopramide (ineffective)</td>
<td>Total and sustained</td>
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<tr>
<td>Baert et al.</td>
<td>35</td>
<td>Female</td>
<td>Cerebellar medulloblastoma</td>
<td>The rachis</td>
<td>Early</td>
<td>Conventional antihistamine (ineffective)</td>
<td>Total and sustained</td>
</tr>
<tr>
<td>Annede et al.</td>
<td>69</td>
<td>Female</td>
<td>Breast</td>
<td>T3-T6 and T8-T11 rachis: 30 Gy (3 Gy fractions)</td>
<td>Late</td>
<td>IV metoclopramide (ineffective)</td>
<td>Partial and sustainable IV then oral erythromycin (1 month)</td>
</tr>
</tbody>
</table>

* within 6 months of radiotherapy; **, 6 months or more after radiotherapy. IV, intravenous.
to declare.

Informed Consent: Written informed consent was obtained from the patient for publication of this manuscript and any accompanying images.

References