PHOTO QUIZ

A case of progressive abdominal angina

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CASE REPORT

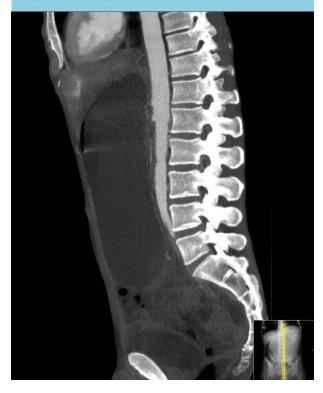
A 38-year-old patient was admitted to our hospital with signs of progressive abdominal angina, nausea, vomiting and cachexia, which had started four months earlier. His medical history reported seminoma 10 years before, for which he underwent orchiectomy followed by para-aortic radiotherapy (13 fractions of 26Gy). Within one year he presented with a local recurrence of the seminoma, which was successfully treated with chemotherapy. Further medical history included paranoid schizophrenia and drug abuse. There was no history of blood coagulation disorders or a family history of occlusive arterial disease.

The patient was referred to the psychiatric ward of our hospital by the psychiatrist from the mental health centre where he lived. The initial diagnosis was nonspecific abdominal pain combined with a delusional disorder, linked to his psychiatric and drug-related history. Thirty-eight days after presentation, a CT angiography was performed due to unsatisfactory clinical improvement, progressive abdominal pain and a high CRP (420 mg/l) (figure 1).

WHAT IS YOUR DIAGNOSIS?

See page 256 for the answer to this photo quiz.

Figure 1. CT angiography showing significant occlusion of the superior mesenteric artery (SMA) and the celiac trunk.



ANSWER TO PHOTO QUIZ (PAGE 255)

A CASE OF PROGRESSIVE ABDOMINAL ANGINA

DIAGNOSIS

Radiotherapy-induced arterial occlusion of the superior mesenteric artery (SMA) and the celiac trunk.

The absence of abnormalities of the vessel walls in other arteries, the specific location of occlusion in the radiated tissue and the age of the patient suggested that the arterial occlusion was a late complication of radiotherapy which he underwent ten years earlier. The history of nicotine and drug abuse and psychiatric comorbidity suggested that in the early stage, the abdominal pain was caused by a psychiatric disturbance, and this delayed the further diagnostic process. Smoking and intravenous drug abuse may have accelerated the process, but are unlikely to be the sole cause of localised abdominal arterial occlusion. The intervention-radiologist performed a difficult percutaneous transluminal angioplasty with stent placement (Express SD 6x19mm, Boston Scientific) in the SMA the next day (figure 2) through brachial access. During this long procedure the patient was conscious and because of agitation we decided stop after stenting the SMA.

Radiotherapy-induced arterial damage was first described in 1899. Studies show that arterial occlusion is rare and can develop many years after radiotherapy, intervals ranging from 4 to 44 years after treatment. It is notable that advanced local occlusion can be present without any signs of arterial damage in other large arteries. Arterial occlusion in the irradiated area, with sparing of the non-radiated areas, is highly suggestive of radiotherapy-induced arterial occlusion. The pathogenesis

Figure 2. Angioplasty before the intervention. (A) shows total occlusion of the SMA and celiac trunk. The SMA was successfully stented (B)



of radiotherapy-induced arterial stenosis is similar to that of accelerated atherosclerosis and can be seen in young patients many years after radiotherapy.³

Early revascularisation is important to prevent bowel ischaemia. Most patients with coronary or femoral occlusion after radiotherapy show slowly progressive nonspecific symptoms. When found early, revascularisation is successful in most cases. Intervention can be difficult, due to fibrotic changes and a slow recovery of the vessel wall in the radiated area. Surgical intervention of arterial occlusion is often complicated due to early re-occlusion. Stenting of the artery can be challenging, but seems the appropriate and preferable treatment of radiation-induced arterial occlusion.

CONCLUSION

Although arterial occlusion is rarely seen as a result of radiotherapy, it can be a serious and potentially lethal complication. It needs to be stressed that although other diseases may be distracting, it is important to be certain of the full medical history of the patient to prevent early mortality in young patients who have undergone radiotherapy.

DISCLOSURES

All authors declare no conflict of interest.

REFERENCES

- Gassmann A. Zur histologie der Roentgenulcer. Ann Intern Med. 1899:199-211.
- Scott AS, Parr LA, Johnstone PA. Risk of cerebrovascular events after neck and supraclavicular radiotherapy: a systematic review. Radiother Oncol. 2009;90:163-5.
- Jurado JA, Bashir R, Burket MW. Radiation-induced peripheral artery disease. Cathet Cardiovasc Interv. 2008;72:563-8.
- 4. Piedbois P, Becquemin JP, Blanc I, et al. Arterial occlusive disease after radiotherapy: a report of fourteen cases. Radiother Oncol. 1990;17:133-40.
- Melliere D, Desgranges P, Berrahal D, et al. [Radiation-induced aorto-iliofemoral arterial arteritis. Mediocrity of the long-term results after conventional surgery]. J Mal Vasc. 2000;25:332-5.
- Hassen-Khodja R, Kieffer E. Radiotherapy-induced supra-aortic trunk disease: early and long-term results of surgical and endovascular reconstruction. J Vasc Surg. 2004;40:254-61.