Successful Treatment of Aortic Rupture with Endovascular Stent Grafting in a Patient with Mantle Cell Lymphoma

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Key Words
Systemic chemotherapy · Abdominal aortic rupture · Lymphoma · Endovascular stent-graft repair

Abstract
\textbf{Purpose:} To present a case of spontaneous aortic rupture in the course of mantle cell lymphoma and successful management with endovascular repair.  
\textbf{Case Report:} A 69-year-old woman presented with a cervical mass. The patient was found to have stage IIIA and Mantle Cell Lymphoma International Prognostic Index (MIPI) 4. She was placed in an intermediate-risk group. The patient received an initial cycle of systemic chemotherapy consisting of rituximab, anthracycline, vincristine and methyl prednisolone. During follow-up, she developed abdominal aortic rupture secondary to intramural hematoma which was successfully managed with endovascular exclusion.  
\textbf{Conclusion:} Hemodynamic changes can be seen during the course of lymphoma subsequent to systemic chemotherapy. These changes might be related to the spontaneous rupture of the main vessels. Endovascular repair may be a more appropriate treatment option than open surgery, especially in a patient with multiple comorbidities.

Introduction
Potential causes of aortic rupture are aneurysm, trauma and plaque rupture. Aneurysmal rupture during the course of systemic chemotherapy is an unusual and serious complication which is difficult to manage, especially in a patient whose health is compromised because of cancer and comorbid conditions. Specific therapeutic modalities have to be coordinated. One of the novel developments in interventional
radiology practice is endoluminal aortic stent grafting in the management of aortic aneurysm including aneurysmal rupture. The first encouraging experiences were published by Parodi et al. in 1991 [1]. Therein, the authors report a case with abdominal aortic rupture during the course of systemic chemotherapy and successful treatment with endovascular stent grafting in an urgent care setting.

Case Report

A 69-year-old woman presented with a cervical mass. She had been evaluated at an independent medical center and a cervical excisional biopsy had been performed. The patient was referred to our Medical Oncology Department with a pathology report. The patient had a history of mild hypertension with no need for medication. She had no history of weight loss, night fever or excessive sweating. Her physical examination was unremarkable except for cervical lymphadenopathy and obesity. Histopathological discussion of the biopsy revealed a diagnosis of mantle cell lymphoma. The patient was staged with PET-CT, endoscopic examination of the gastrointestinal system and bone marrow aspiration/biopsy. PET-CT showed multiple hypermetabolic foci at the cervical, mediastinal and retroperitoneal regions. The patient was staged as IIA. MIPI (Mantle Cell Lymphoma International Prognostic Index) of the patient was 4, so she was placed in the intermediate-risk group. The treatment plan consisted of 6 cycles of R-CHOP regimen. Two days after the first cycle of systemic chemotherapy, syncope occurred, blood pressure was 70/50 mm Hg, and her hematocrit level was 27%; the other parts of the complete blood count were normal, as were the results of other laboratory tests, including serum electrolytes and tests for renal and liver function. Cardiological and surgical evaluations were normal.

The patient was transferred to the Intensive Care Unit. Over the next 2 days, her hematocrit values progressively decreased and 6 units of packed red blood cells were transfused. Vital signs remained normal during this period. Thoracic and abdominal CT scans were performed, revealing a huge intramural hematoma and a retroperitoneal hematoma (fig. 1). The retroperitoneal hematoma was secondary to the intramural hematoma. Her aorta did not seem aneurysmatic. Cardiovascular Surgery and the Interventional Radiology Unit consulted on the patient. Due to comorbid conditions, an open surgical procedure was considered to have high preoperative and postoperative mortality; thus the patient was transferred to the Interventional Radiology Unit. A percutaneous approach was used from the right and left common femoral arteries. Diagnostic angiography revealed that the distal aorta had minimal aneurysmatic dilatation which was very mild at 30 mm in diameter (fig. 2a). A 28-mm stent graft (Medtronic, USA) was placed into the aorta, and then the contralateral leg was placed. A check angiogram showed excellent positioning of the stent graft with no leak (fig. 2b). The patient’s hemoglobin level stabilized immediately after the stent-graft placement.

Discussion

The most frequent etiology of aortic rupture is aneurysm. Risk of rupture of an aortic aneurysm increases with increased diameter of the aorta. Currently, a diameter more than 55 mm is the indication for treatment with surgical or endovascular means. There are other potential causes such as trauma and plaque rupture. Nontraumatic, noninfectious rupture of the nonaneurysmal aorta is a very rare condition and the most frequent cause is penetrating atherosclerotic ulcer [2]. Abdominal aortic aneurysms (AAA) are common and generally asymptomatic unless rupture occurs. A 3–4-cm AAA has a 1–2% risk of rupture over 5 years [3]. Aneurysms in oncologic cases are mostly seen with colorectal carcinomas and usually located in the abdominal aorta. Aneurysmal and nonaneurysmal rupture of the aorta in oncologic cases is a very rare condition and mostly seen in patients who have tumors with periaortic localization and concomitant aneurysmal dilatation [4]. For primary aortic tumors, rupture has been
described only in a single case of leiomyosarcoma of the thoracic aorta [5]. Ehata et al. [6] reported a case of ruptured AAA repair in which a pheochromocytoma was diagnosed postoperatively by CT scan. Thompson et al. [7] also reported a case in which a pheochromocytoma was found using metaiodobenzylguanidine imaging after a repaired AAA began leaking. Moreover, 4 cases of dissecting thoracic aneurysms associated with pheochromocytoma have been described [6]. Spontaneous rupture of the nonaneurysmal aorta has been reported with tumor infiltration in neurofibromatosis [8]. All reported cases of primary malignant lymphoma invading the aorta were associated with symptomatic or ruptured aortic aneurysms. Aortoiliac fistula was found in 1 case and endovascular repair was successfully performed [9].

Penetrating ulcerated plaque and hemodynamic changes such as hypertension are the most common causes of aneurysmal rupture of the aorta. Chemotherapy is a stress factor for hemodynamic parameters, but the patient presented here received her systemic chemotherapy in the oncology ward due to the risk of probable tumor lysis syndrome. During in-ward follow-up, no abnormalities were detected by means of physical examination, vital signs and laboratory results. An abdominal CT scan was performed as part of the initial staging, and radiological evaluation showed no abnormality in the abdominal aorta. A repeat CT scan at the time of diagnosis of rupture did not show an aneurysmal aorta but a big intramural hematoma. Diagnostic angiography at the beginning of endovascular intervention showed a very mild dilatation of the distal abdominal aorta.

At this point, the patient could be managed by either open surgery or endovascular stent grafting. Endovascular stent grafting has been shown to reduce procedure-related and early morbidity and mortality in a large prospective randomized trial [10]. Open surgery had the disadvantages of bleeding and infections due to cytopenia and may be hazardous because of poor conditional status and comorbid conditions of the patient. So we decided on endovascular stent grafting.

Endovascular repair provides a valuable alternative in the acute setting of aneurysmal rupture and should be an established treatment option, especially in patients with several comorbid conditions.
Fig. 1. Contrast-enhanced CT image reveals significant leak from the aorta (arrow). There is a crescent-shaped hematoma in the aortic wall causing the rupture, and a more significant hematoma in the retroperitoneum.

Fig. 2. a Aortogram before stent-graft placement reveals than the aorta is otherwise normal in size. There is a mild widening of the aorta on its left distal side. A leak is not visible on the aortogram.

b Aortogram after stent-graft placement. The stent graft seals the whole infrarenal aorta excluding the rupture point.
References


