Informed consent was obtained from the patient. A 64-year-old man with history of smoking, alcohol abuse and peripheral artery disease presented at the Emergency department with intense pain in the lower back after a 2-week history of fever, weight loss and drenching sweats. Systemic examination was otherwise unremarkable, there were no peripheral stigmata of infective endocarditis, auscultation of the precordium identified no murmurs, chest radiography and urinalysis were negative. Blood tests revealed elevated white blood cell count (11.1×10⁹ cells/L) and C-reactive protein (103 mg/L). The patient was admitted to the ward and blood cultures were taken, which detected no bacteremia or fungemia. A magnetic resonance imaging (MRI) study of the patient’s back was performed to evaluate for possible spondylodiscitis. The study showed bone marrow edema and contrast enhancement of the lumbar vertebra L2, L3 and L4, indicative of active spondylodiscitis. As an additional finding, a saccular aneurysm protruding from the dorsal border of the abdominal aorta was observed (Figure 1).

An integrated whole body ¹⁸F-fluorodeoxyglucose (¹⁸F-FDG) positron emission tomography/MRI (PET/MRI) study was performed to evaluate the extent of the suspicious aortic lesion, its relation to the spondylodiscitis as well as for other unknown foci of infection. The ¹⁸F-FDG PET/MRI study consisted of a whole body T2 weighted study, a whole body T1 weighted study pre- and post-contrast as well as of a whole-body PET study. All MR images were performed under breathing instructions. A T2 weighted study with fat suppression and T1 weighted study pre- and post-contrast in thin slices and with cardiac triggering were performed in the area of the aortic lesion. The study was performed on a Siemens Biograph mMR 3T scanner. A total dosage of 133 MBq (¹⁸F-FDG) and 10 mL MRI-contrast (Gadovist) was administered. PET/MRI images showed a saccular aortic aneurysm of 5.6 cm, extending from the distal thoracic aorta to the level of the celiac trunk and mesenteric artery origins (type V TAAA according to modified Crawford classification). Intense ¹⁸F-FDG uptake in the vessel wall of the aortic aneurysm reaching to a maximum standard uptake value (SUVmax) of 5.7 was seen as well as contrast enhancement within the thickened aortic wall and surrounding soft tissue (Figure 2).

The patient received intravenous antibiotics for ten days...
in total. The infected aortic tissue from the distal thoracic aorta to the level of the mesenteric artery origin was resected surgically by an open approach and the aneurysm was replaced by a tube graft made of bovine pericardium. The superior mesenteric artery and celiac trunk were directly re-implanted to the graft. Genotypic analysis of the aortic wall tissue surgical specimen by PCR revealed the presence of Gram-negative bacteria (*Capnocytophaga canimorsus*), possibly transmitted through a bite from the patient’s dog. The patient made a quick recovery. Post-surgery follow-up computed tomography imaging and blood tests revealed no signs of surgical complications and no signs of residual aortic or graft infection as well as a normalization of the CRP.

Integrated *¹⁸*F-FDG PET/MRI has proven added value for the evaluation of different types of cancer (1-3). This
case shows the high potential of FDG PET/MRI in the diagnostic algorithm of aortic lesions suspicious for infected aneurysms and other infectious spread. This diagnostic tool can have direct and significant impact on the therapeutic care pathway.

Mycotic aneurysms are a rare entity, but may pose a challenging problem (4). Survival is markedly increased by prompt diagnosis and surgical treatment (5,6). In early reports, most infected aneurysms were related to valvular infection, however nowadays most mycotic aneurysms are aortitis related (7,8). *Capnocytophaga canimorsus* is found in the oral flora of most dogs and is a rare pathogen in humans. Human infections with *C. canimorsus* have been previously reported, including endocarditis, abscesses, and mycotic aneurysms. Immuno-incompetent patients are at higher risk of infections (9). Other organisms with affinity to the aortic wall include a large variety of gram-positive and gram-negative species of which *staphylococcus* species (*S. Aureus, S. pneumoniae, S. epidermidis*) and salmonella species being most commonly detected. Less common pathogens include *Coxiella burnetii* (Q fever) and mycobacteria (10). Bacterial seeding of the aortic wall can occur by haematogenous spread, lymphatic spread or direct extension from an adjacent infected focus (8). The intimal lining of the aorta is generally highly resistant to infection, however, even normal aortic intima can be subject to bacterial invasion followed by secondary degeneration of the arterial wall with aneurysm enhancement and rupture. In one study, gram negative bacteria seemed to be associated with a higher risk of aortic rupture and mortality than gram-positive (11).

The diagnosis of mycotic aneurysm is difficult, particularly in the absence of the classical signs (back or abdominal pain, fever, pulsatile mass, positive blood culture). Yet, an early diagnosis is critical as mycotic infectious aortitis is associated with a high rate of rupture and subsequent mortality as early as one week after the onset of aortitis (12).

MR imaging with gadolinium enhancement is becoming the non-invasive imaging modality of choice for aortitis (13). MR imaging demonstrates the anatomic localization, degree and extent of vascular stenosis or vascular aneurysmal dilatation. Moreover, it has an excellent resolution of the aortic wall and it can depict areas of active aortitis as crescentic or ring-like vessel wall oedema and/or thickening (14). Drawbacks of MRI are the relatively decreased sensitivity as well as a long procedure time.

More recently, the use of 18F-FDG has emerged as a potential tool for the initial diagnosis and assessment of disease activity of aortitis caused by either infectious conditions (15,16) or non-infectious giant cell arteritis or Takayasu arteritis (3,17-20). After first positive results of 18F-FDG PET imaging in infectious/mycotic aortic aneurysm (15,16), recent imaging series have reported a variable sensitivity of 60–90% and a specificity of 88–100% of 18F-FDG PET and PET CT for diagnosing active inflammation in arteritis (17,20-22). Anatomic changes, such as the presence of wall thickening, luminal thrombus and aneurysm size cannot be assessed on PET. Hence, by combining the advantages of both MRI and PET, the hybrid PET/MRI imaging is thought to be an imaging modality that could improve the sensitivity in the diagnosis of aortitis.

In conclusion, the integrated whole body 18F-FDG PET/MRI study provided all relevant diagnostic imaging information needed for the diagnosis of an infected aortic aneurysm and the successful planning of its treatment.

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**Footnote**

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

Informed Consent: Informed consent was obtained from the patient.

**References**


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