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Original article

^{18}F -fluorodeoxyglucose positron-emission tomography combined with computed tomography as a diagnostic tool in native valve endocarditis

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Objective The aim of the study was to investigate the value of ^{18}F -fluorodeoxyglucose positron-emission tomography combined with computed tomography (^{18}F -FDG-PET/CT) in diagnosing native valve endocarditis (NVE).

Patients and methods All patients with bacteremia and suspicion of NVE between January 2013 and June 2016 were identified from the hospitals' register and retrospectively included if echocardiography and ^{18}F -FDG-PET/CT were performed within 14 days. ^{18}F -FDG-PET/CT scans were scored independently by two nuclear medicine physicians. ^{18}F -FDG-PET/CT was compared with the modified-Duke criteria and a multidisciplinary consensus.

Results A total of 88 patients were included. In 10 patients with definite NVE according to the modified-Duke criteria, three (30.0%) patients had increased ^{18}F -FDG uptake in or around the heart valves and seven (70.0%) patients had no increased ^{18}F -FDG uptake. In patients without definite NVE according to the modified-Duke criteria, 89.7% (70/78) of the patients had no increased ^{18}F -FDG uptake in or around the heart valves. Of all 20 patients with NVE according to multidisciplinary consensus, nine (45.0%) patients had increased ^{18}F -FDG uptake in or around the heart valves and 11 (55.0%) patients had a normal ^{18}F -FDG-PET/CT result.

Conclusion A negative ^{18}F -FDG-PET/CT result should not be interpreted as an exclusion of NVE. In patients with possible or rejected NVE according to the modified-Duke criteria, ^{18}F -FDG-PET/CT could be used in case of sustained suspicion of NVE owing to its high specificity in case of abnormal FDG uptake at the valve region. ^{18}F -FDG-PET/CT is important for detecting metastatic infection which already warrants the need to perform ^{18}F -FDG-PET/CT in all patients with suspected NVE. *Nucl Med Commun* 39:747–752 Copyright © 2018 Wolters Kluwer Health, Inc. All rights reserved.

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Introduction

Infective endocarditis is a severe condition with a mortality rate up to 40% [1]. Early diagnosis of infective endocarditis is essential for successful management and improved outcome, but diagnosing endocarditis is challenging owing to a variable clinical presentation with often nonspecific symptoms. For diagnosing infective endocarditis, the modified-Duke criteria are currently used [2]. According to these criteria, diagnosis of definite endocarditis is mainly based on positive blood cultures with typical micro-organisms and/or evidence of infective endocarditis on echocardiography. However, sensitivity of echocardiography is limited, ~75% for transthoracic echocardiography (TTE) and 85–90% for transesophageal echocardiography (TEE) [3].

^{18}F -fluorodeoxyglucose (FDG) positron-emission tomography combined with computed tomography (PET/CT)

has shown effectiveness in diagnosing infectious diseases. The value of ^{18}F -FDG-PET/CT in diagnosing infective endocarditis has been reported [4] and has shown to be a valuable diagnostic technique in patients suspected of prosthetic valve endocarditis (PVE). Recently, ^{18}F -FDG-PET/CT was added to the European Society of Cardiology modified-diagnostic criteria as a major criterion for PVE [5]. In patients with native valve endocarditis (NVE), the diagnostic value of ^{18}F -FDG-PET/CT has not been studied extensively. A few studies, including often less than 10 patients with native valves, found low sensitivity for ^{18}F -FDG-PET/CT in diagnosing NVE [4]. However, these studies were often performed without prior low carbohydrate-fat-allowed diet to suppress cardiac glucose metabolism [6]. The purpose of this study was to investigate the diagnostic value of ^{18}F -FDG-PET/CT in NVE in

a large cohort of patients, prepared with a low carbohydrate-fat-allowed diet.

Patients and methods

Patients and study design

All patients in this study were retrospectively included between January 2013 and June 2016 in the Radboud University Medical Center, a Tertiary Referral Center in Nijmegen, the Netherlands. Patients with bacteremia were included when both echocardiography, because of suspicion of NVE (TTE and/or TEE), and ^{18}F -FDG-PET/CT were performed within 14 days. Exclusion criteria were an age below 18 years, prosthetic heart valve, cardiac implantable electronic devices (CIED) infections, more than 14 days between ^{18}F -FDG-PET/CT and echocardiography, when no low carbohydrate-fat-allowed diet was used before the ^{18}F -FDG-PET/CT, and when assessment of the heart was impossible owing to physiological ^{18}F -FDG uptake despite the low carbohydrate-fat-allowed diet. According to Dutch law, this study was exempt from approval by an ethics committee, because of the retrospective character of this study and the anonymous storage of data. This was confirmed by the regional ethics committee.

In all patients included in this study, the microbiologists advised for an infectious disease specialist consultation because of the positive blood culture. All patients were discussed in a multidisciplinary 'endocarditis team', including infectious disease specialists, medical microbiologists, cardiologists, and nuclear medicine physicians. In this weekly multidisciplinary meeting, the medical history, physical examination, laboratory and microbiological results, and imaging results including ^{18}F -FDG-PET/CT of all patients were presented, and based on this clinical information, a diagnosis was made. The diagnosis was not based on a scoring system but made by consensus of all physicians present at the meeting based on all clinical signs and symptoms, risk factors, blood cultures (number of positive cultures/total, micro-organism), and all imaging performed. For echocardiography, TTE was used as a first-line screening technique. TEE was advocated in all patients in whom TTE was negative for NVE, especially when imaging was hampered owing to technical or anatomical problems. NVE was diagnosed according to the modified-Duke criteria [2] as gold standard and also to the multidisciplinary consensus. NVE was treated according to international guidelines [5].

Diagnostic work-up

An integrated PET/CT scanner (Biograph 40 mCT; Siemens Healthcare, Erlangen, Germany) was used for imaging. Before ^{18}F -FDG injection, patients fasted, and any glucose or insulin-containing infusions were discontinued for at least 6 h. All patients were instructed to adhere to a low carbohydrate-fat-allowed diet 24 h before ^{18}F -FDG-PET/CT was performed. Blood glucose samples were taken from all patients before ^{18}F -FDG

administration. At the time of ^{18}F -FDG injection, glucose was below 12 mmol/l in all patients. One hour after intravenous injection of an average dosage of 3.3 MBq/kg ^{18}F -FDG (Mallinckrodt Pharmaceuticals, Petten, The Netherlands or IBA Molecular, Amsterdam, The Netherlands), a whole body low-dose CT scan was acquired for anatomic correlation and attenuation correction of the PET data. Emission images of the same area were acquired. Images of the heart were re-evaluated independently by two nuclear medicine physicians without knowledge of prior clinical evaluation. When it was not possible to reliably evaluate the valve planes, patients were excluded. Any increased ^{18}F -FDG uptake in or around the heart valves outside the area of the myocardium was considered as abnormal. Minimally increased ^{18}F -FDG uptake was defined as uptake in the region of a heart valve just above normal heart uptake. Highly increased ^{18}F -FDG uptake of a heart valve was defined as uptake in the region of the heart valve clearly distinguishable from normal heart uptake. When uptake in the region of a heart valve was not distinguishable from normal heart uptake, ^{18}F -FDG uptake was considered negative. Subsequently, discordant results were solved by consensus reading.

Echocardiography was considered positive for infective endocarditis when vegetations, defined as oscillating intracardiac structures, were visualized on the valves or their adjacent structures or in the path of a regurgitant jet in the absence of an alternative anatomic explanation.

Statistical analysis

In patients with definite NVE according to the modified-Duke criteria, any comparison was made with patients with possible and rejected NVE according to the modified-Duke criteria. The κ -statistic test was used to calculate the level of agreement between the two nuclear medicine physicians. Differences in outcomes were tested using Fisher's exact test for categorical variables. Statistical significance was defined as a *P* value less than 0.05. Statistical analysis was performed using SPSS (version 22.0; SPSS Inc., Chicago, Illinois, USA).

Results

A total of 104 patients underwent both echocardiography and ^{18}F -FDG-PET/CT within 14 days because of bacteremia. A total of 16 patients were excluded because ^{18}F -FDG-PET/CT scan quality was too limited for assessment of the heart region because of physiological ^{18}F -FDG uptake. Of all 88 included patients, baseline characteristics are shown in Table 1. Ten (11.4%) patients were diagnosed with definite NVE and 48 (54.5%) patients were diagnosed with possible NVE according to the modified-Duke criteria. In 30 patients, diagnosis of NVE was rejected (34.1%). The evaluation of ^{18}F -FDG-PET/CT for NVE showed a high level of agreement (Cohen's $\kappa = 0.97$).

Table 1 Baseline characteristics of all 88 patients with definite, possible, and rejected native valve endocarditis according to the modified-Duke criteria

	Definite endocarditis [n (%)]	Possible endocarditis [n (%)]	Rejected endocarditis [n (%)]
Number of patients	10	48	30
Male	6 (60.0)	28 (58.3)	16 (53.3)
Age [mean (range)] (years)	68.9 (35–90)	63.4 (21–88)	54.5 (17–88)
Blood culture positive	10 (100)	48 (100)	30 (100)
<i>Staphylococcus aureus</i>	3 (30.0)	40 (83.3)	13 (43.3)
<i>Coagulase-negative Staphylococcus</i> spp.	0	1 (2.1)	1 (3.3)
<i>Streptococcus</i> spp.	5 (50.0)	5 (10.4)	3 (10.0)
<i>Enterococcus</i> spp.	2 (20.0)	1 (2.1)	1 (3.3)
Gram negative	0	1 (2.1)	10 (33.3)
Other	0	0	2 (6.7)
Pacemaker	2 (20.0)	4 (8.3)	0
ICD	0	1 (2.1)	0
Echocardiography			
TTE only	2 (20.0)	19 (39.6)	23 (76.7)
TEE only	3 (30.0)	2 (4.2)	2 (6.7)
Both TTE and TEE	5 (50.0)	27 (56.3)	5 (16.7)
Metastatic infection other than endocarditis	8 (80.0)	30 (62.5)	10 (33.3)

ICD, implantable cardioverter defibrillator; TEE, transesophageal echocardiography; TTE, transthoracic echocardiography.

Definite native valve endocarditis

In 10 patients with definite NVE according to the modified-Duke criteria, three (30.0%) patients had increased ^{18}F -FDG uptake in or around the heart valves. Seven (70.0%) patients had no increased ^{18}F -FDG uptake in or around the heart valves. In patients without definite NVE according to the modified-Duke criteria, 89.7% (70/78) patients had no increased ^{18}F -FDG uptake in or around the heart valves. In all patients with definite NVE according to the modified-Duke criteria and with increased ^{18}F -FDG uptake at the heart valve region, ^{18}F -FDG uptake was highly increased.

Possible native valve endocarditis

Of all 48 patients with possible NVE according to the modified-Duke criteria, five (10.4%) patients had increased ^{18}F -FDG uptake in or around the heart valves. Although these five patients had only possible NVE according to the modified-Duke criteria, three (60.0%) of these patients were treated as having NVE based on the severity of infection, risk factors, and ^{18}F -FDG-PET/CT results (Fig. 1). Of 43 patients with possible NVE according to the modified-Duke criteria without increased ^{18}F -FDG uptake in the heart valve region, four (9.3%) patients were treated as having NVE.

Rejected native valve endocarditis

In three patients with increased ^{18}F -FDG uptake in the heart valve region, NVE was rejected according to the Duke criteria. One patient was treated for 2 years until

his death for an infected abdominal aortic graft with *Escherichia coli* and *Candida albicans*. Another patient was treated for 6 months until her death for an infected thoracic aortic aneurysm with *Salmonella dublin*. One patient was treated for 6 weeks because of *Staphylococcus aureus* bacteremia with metastatic foci in lungs and soft tissue. She died 6 months later owing to duodenum carcinoma.

Multidisciplinary consensus

NVE was diagnosed according to the multidisciplinary consensus in 20 (22.7%) patients (Fig. 2). Of all patients with NVE according to the multidisciplinary consensus, nine (45.0%) patients had increased ^{18}F -FDG uptake in or around the heart valves and 11 (55.0%) patients had normal ^{18}F -FDG-PET/CT. Of the 20 patients with definite NVE according to the multidisciplinary consensus, 10 were initially classified as definite, seven as possible, and three as rejected endocarditis by the modified-Duke criteria. In patients with increased ^{18}F -FDG uptake in or around the heart valves, metastatic infection other than endocarditis was found in 72.7% (8/11) of patients, and in patients without increased ^{18}F -FDG uptake in or around the heart valves, metastatic infection was found in 51.9% (40/77) of patients ($P=0.195$). No relapses occurred.

Classifications of all patients according to the diagnosis of NVE based on the modified-Duke criteria, modified-Duke criteria including ^{18}F -FDG-PET/CT result as a major criterion, ^{18}F -FDG-PET/CT result only, and diagnosis according to the multidisciplinary consensus are shown in Table 2.

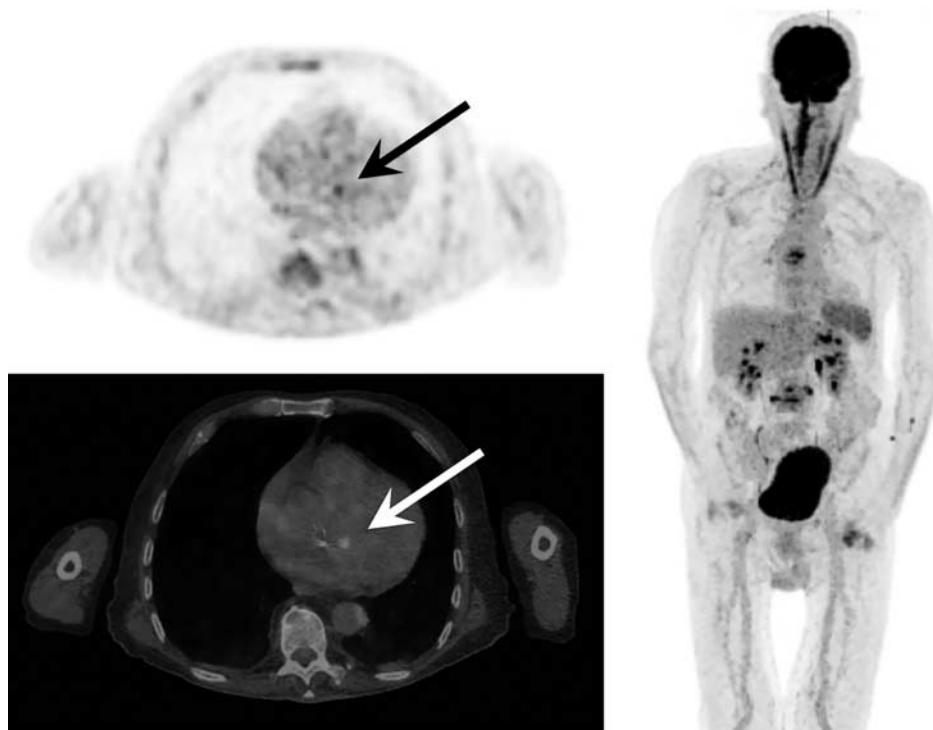
Metastatic infection

^{18}F -FDG-PET/CT detected metastatic infection other than endocarditis in 48 (54.5%) patients. ^{18}F -FDG-PET/CT was the first modality to localize infectious foci in 38 (79.2%) patients. Localizations of these metastatic infection were endovascular infection (23.7%), spondylodiscitis (23.7%), pulmonary foci (15.8%), arthritis (13.2%), soft tissue infections (13.2%), splenic abscesses (7.9%), and nonvertebral osteomyelitis (2.6%).

Discussion

Data on the value of ^{18}F -FDG-PET/CT in NVE are limited. This is the first study in a large cohort investigating the value of ^{18}F -FDG-PET/CT in patients with suspected NVE only, by excluding PVE and CIED infections. Our study shows that ^{18}F -FDG-PET/CT has a low sensitivity, with seven of 10 patients with definite NVE according to the modified-Duke criteria without increased ^{18}F -FDG uptake of the heart valve. Although ^{18}F -FDG-PET/CT is insufficient to rule out NVE, a positive finding on ^{18}F -FDG-PET/CT is sufficiently specific to imply clinical consequences, as 70 of 78 patients without definite NVE according to the modified-

Fig. 1



^{18}F -FDG-PET/CT images of a 79-year-old man who was admitted with fever and positive blood cultures with *Streptococcus gallolyticus*. Besides spondylodiscitis of Th8-9 and L3-4, ^{18}F -FDG-PET/CT showed increased ^{18}F -FDG uptake of the mitral valve (arrow). TEE was negative for endocarditis. Although this patient had only possible endocarditis according to the modified-Duke criteria, he was treated as having NVE. ^{18}F -FDG-PET/CT, ^{18}F -fluorodeoxyglucose positron-emission tomography combined with computed tomography; NVE, native valve endocarditis; TEE, transesophageal echocardiography.

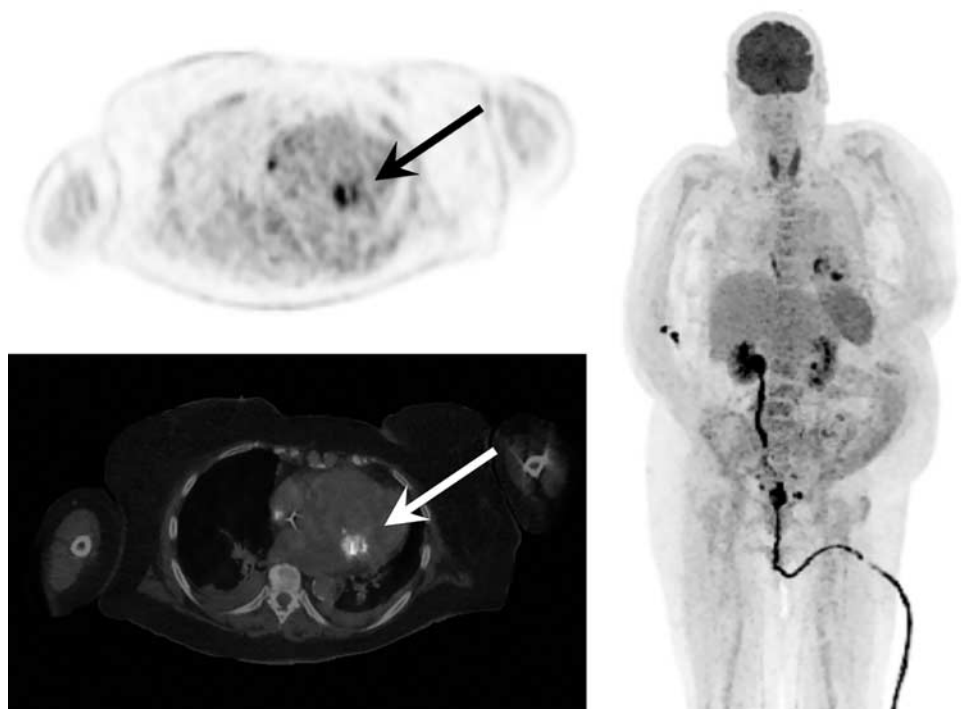
Duke criteria showed no increased ^{18}F -FDG uptake around the heart valves. The results of our study are in line with three other studies performed on the value of ^{18}F -FDG-PET/CT in NVE. These studies, mainly focusing on PVE with inclusion of only six [7] and seven [8,9] patients with NVE, found very low sensitivity for ^{18}F -FDG-PET/CT in NVE. Previously, we performed a prospectively included study on the value of ^{18}F -FDG-PET/CT in 72 patients with suspected endocarditis, of whom 66 patients had suspected NVE [10]. In this study, an older generation PET/CT scanner was used, and no low carbohydrate-fat-allowed diet was performed. In the present study, the level of agreement between two evaluating nuclear medicine physicians was much higher than in the previous study (0.97 vs. 0.36). This difference is probably explained by the increasing experience in evaluating ^{18}F -FDG-PET/CT in NVE since this previous study, and our multidisciplinary meeting performed weekly since 2013.

NVE is a clinical diagnosis that is based on risk factors, physical examination, microbiological results, and imaging. The fact that diagnosing NVE is challenging is shown by the limited sensitivity of the modified-Duke criteria [2], which remain the 'gold standard' for diagnosis. Moreover, a

large percentage of suspected NVE cases remain as possible NVE [11]. Therefore, patients with suspected NVE should be discussed in a multidisciplinary endocarditis team, to incorporate as much individualized information as possible in the final diagnosis. Our results show that by multidisciplinary consensus definite NVE was diagnosed twice as often compared with diagnosis according to the modified-Duke criteria and also the discouraging diagnosis of possible NVE was abandoned. ^{18}F -FDG-PET/CT is an imaging technique that should be assessed within the clinical context of patients. Reevaluation of ^{18}F -FDG-PET/CT without knowledge of clinical details, which is common in this type of studies, could lead to less sensitive reading.

The diagnostic value of ^{18}F -FDG-PET/CT for PVE has been extensively studied with promising results [4], and ^{18}F -FDG-PET/CT was recently added to the European Society of Cardiology modified-diagnostic criteria for PVE [5]. Because of its relatively high specificity, adding ^{18}F -FDG-PET/CT to the Duke criteria also for NVE could be valuable, as our results show an increase of NVE diagnoses when ^{18}F -FDG-PET/CT is included (Table 2). Three patients with rejected NVE according to the modified-Duke criteria and increased ^{18}F -FDG uptake of the heart valves were at high risk for NVE and could have been incorrectly

Fig. 2



¹⁸F-FDG-PET/CT images of a 75-year-old woman with a pacemaker and ovarian carcinoma who was admitted with fever and severe confusion. Blood cultures grew *Proteus mirabilis*. TEE was negative for endocarditis. ¹⁸F-FDG-PET/CT showed highly increased ¹⁸F-FDG uptake of the mitral valve (arrow). A few days later she developed splinter hemorrhages on her hand and foot. TEE was repeated 1 week after the ¹⁸F-FDG-PET/CT and confirmed the diagnosis of native mitral valve endocarditis; no vegetations were seen on her pacemaker leads. These patients died 1 week later owing to therapy-resistant mitral valve endocarditis. ¹⁸F-FDG-PET/CT, ¹⁸F-fluorodeoxyglucose positron-emission tomography combined with computed tomography; TEE, transesophageal echocardiography.

Table 2 Diagnosis of native valve endocarditis according to the modified-Duke criteria, the modified-Duke criteria including ¹⁸F-fluorodeoxyglucose positron-emission tomography combined with computed tomography, revised ¹⁸F-fluorodeoxyglucose positron-emission tomography combined with computed tomography result only, and diagnosis by multidisciplinary consensus (n = 88)

	Modified-Duke criteria [n (%)]	Modified-Duke criteria including ¹⁸ F-FDG-PET/CT [n (%)]	Revised ¹⁸ F-FDG-PET/CT result only [n (%)]	Multidisciplinary consensus ^a
Definite	10 (11.4)	18 (20.5)	11 (12.5)	20 (22.7)
Possible	48 (54.5)	43 (48.9)	0	0
Rejected	30 (34.1)	27 (30.7)	77 (87.5)	68 (77.3)

¹⁸F-FDG-PET/CT, ¹⁸F-fluorodeoxyglucose positron-emission tomography combined with computed tomography.
^aMultidisciplinary board including infectious diseases specialists, medical microbiologists, cardiologists, nuclear medicine physicians, and if needed cardiac surgeons.

diagnosed as rejected NVE. The fact that TTE only, and thus no TEE, was performed in most patients with possible or rejected NVE (Table 1) could have led to missed diagnoses of definite NVE. The diagnostic criteria of the European Society of Cardiology from 2015 recommend cardiac CT in suspected NVE [5]. In our study, we did not perform cardiac CT as part of our diagnostic protocol in NVE, partly owing to the fact that patients were included from 2013 in the present study.

Besides intracardiac lesions, the value of ¹⁸F-FDG-PET/CT has been studied on extracardiac complications of endocarditis. ¹⁸F-FDG-PET/CT has proven to be valuable and

cost effective in detecting metastatic foci in patients with bacteremia and PVE and/or NVE [12–14] and is therefore increasingly used in patients with suspected endocarditis. In our study, metastatic infection other than endocarditis was detected by ¹⁸F-FDG-PET/CT in 54.5% of patients, which is comparable to previous results [13,14].

A possible explanation for the limited sensitivity of ¹⁸F-FDG-PET/CT in diagnosing NVE is continuous movement of the cardiac valves during acquisition and the small size of vegetations, as metabolism in very small vegetations could be insufficient to be discernible above background activity, especially after being blurred by the

movement of the heart [5]. Moreover, in the subacute-chronic phase of vegetations, micro-organisms may disappear and granulomatous inflammation occurs with transformation of vegetations into calcified deposits [15]. So, a subacute course of NVE could decrease the sensitivity of ^{18}F -FDG-PET/CT as ^{18}F -FDG accumulates particularly in activated leukocytes. Gomes *et al.* [4] proposed a diagnostic algorithm in suspected infective endocarditis. The authors state that patients with highly suspected NVE should undergo both TEE and MDCTA (ECG-gated multidetector CT angiography) to image intracardiac lesions, and also ^{18}F -FDG-PET/CT for detecting extracardiac foci and for intracardiac lesions in case of sustained suspicion of NVE after inconclusive TEE and MDCTA. ^{18}F -FDG is not a specific tracer, as ^{18}F -FDG uptake could also increase in atherosclerotic plaques in the heart or physiological uptake in the myocardium and papillary muscles. Leukocyte scintigraphy with single-photon emission computed tomography/CT is known for a high specificity in detection of infectious foci, and its value has been investigated in PVE and CIED infection [16,17]. More specific PET/CT tracers, additional heparin preadministration [6], motion correction, ECG-gated scanning, respiratory gated scanning to reduce breathing motion artifacts, and/or combination with MDCTA, potentially optimize the diagnostic performance of PET/CT to detect NVE.

Limitations of this study are the retrospective study design and the small number of patients with definite NVE according to the modified-Duke criteria. In our cohort of 88 patients, 48 patients were classified as possible NVE according to the modified-Duke criteria. However, despite these small number of patients with definite NVE, this is the largest study so far on patients with NVE. Owing to our study design, the comparison between ^{18}F -FDG-PET/CT and the multidisciplinary consensus could have led to incorporation bias, because of the fact that ^{18}F -FDG-PET/CT is a part of the multidisciplinary consensus.

Conclusion

In conclusion, a negative ^{18}F -FDG-PET/CT result should not be interpreted as an exclusion of NVE. In patients with possible or rejected NVE according to the modified-Duke criteria, ^{18}F -FDG-PET/CT could be used in case of sustained suspicion of NVE owing to its high specificity in case of abnormal FDG uptake at the valve region. In patients suspected of NVE, ^{18}F -FDG-PET/CT is important for detecting metastatic infection which already warrants the need to perform ^{18}F -FDG-PET/CT in all patients with NVE.

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Conflicts of interest

There are no conflicts of interest.

References

- Murdoch DR, Corey GR, Hoen B, Miro JM, Fowler VG Jr, Bayer AS, *et al.* Clinical presentation, etiology, and outcome of infective endocarditis in the 21st century: the International Collaboration on Endocarditis-Prospective Cohort Study. *Arch Intern Med* 2009; **169**:463–473.
- Li JS, Sexton DJ, Mick N, Nettles R, Fowler VG Jr, Ryan T, *et al.* Proposed modifications to the Duke criteria for the diagnosis of infective endocarditis. *Clin Infect Dis* 2000; **30**:633–638.
- Habib G, Badano L, Tribouilloy C, Vilacosta I, Zamorano JL, Galderisi M, *et al.* Recommendations for the practice of echocardiography in infective endocarditis. *Eur J Echocardiogr* 2010; **11**:202–219.
- Gomes A, Glaudemans AW, Touw DJ, van Melle JP, Willems TP, Maass AH, *et al.* Diagnostic value of imaging in infective endocarditis: a systematic review. *Lancet Infect Dis* 2017; **17**:e1–e14.
- Habib G, Lancellotti P, Antunes MJ, Bongioanni MG, Casalta JP, Del Zotti F, *et al.* 2015ESC guidelines for the management of infective endocarditis: the Task Force for the Management of Infective Endocarditis of the European Society of Cardiology (ESC). Endorsed by: European Association for Cardiothoracic Surgery (EACTS), the European Association of Nuclear Medicine (EANM). *Eur Heart J*, 2015:3075–3128.
- Scholtens AM, Verberne HJ, Budde RP, Lam MG. Additional heparin preadministration improves cardiac glucose metabolism suppression over low-carbohydrate diet alone in (18)F-FDG PET imaging. *J Nucl Med* 2016; **57**:568–573.
- Granados U, Fuster D, Pericas JM, Llopis JL, Ninot S, Quintana E, *et al.* Diagnostic accuracy of 18F-FDG PET/CT in infective endocarditis and implantable cardiac electronic device infection: a cross-sectional study. *J Nucl Med* 2016; **57**:1726–1732.
- Salomaki SP, Saraste A, Kemppainen J, Bax JJ, Knuuti J, Nuutila P, *et al.* 18F-FDG positron emission tomography/computed tomography in infective endocarditis. *J Nucl Cardiol* 2017; **24**:195–206.
- Ricciardi A, Sordillo P, Ceccarelli L, Maffongelli G, Calisti G, Di Pietro B, *et al.* 18-Fluoro-2-deoxyglucose positron emission tomography-computed tomography: an additional tool in the diagnosis of prosthetic valve endocarditis. *Int J Infect Dis* 2014; **28**:219–224.
- Kouijzer JJ, Vos FJ, Janssen MJ, van Dijk AP, Oyen WJ, Bleeker-Rovers CP. The value of 18F-FDG PET/CT in diagnosing infectious endocarditis. *Eur J Nucl Med Mol Imaging* 2013; **40**:1102–1107.
- Habib G, Derumeaux G, Avierinos JF, Casalta JP, Jamal F, Volot F, *et al.* Value and limitations of the Duke criteria for the diagnosis of infective endocarditis. *J Am Coll Cardiol* 1999; **33**:2023–2029.
- Orvin K, Goldberg E, Bernstine H, Groshar D, Sagie A, Kornowski R, *et al.* The role of FDG-PET/CT imaging in early detection of extra-cardiac complications of infective endocarditis. *Clin Microbiol* 2015; **21**:69–76.
- Kestler M, Munoz P, Rodriguez-Creixems M, Rotger A, Jimenez-Requena F, Mari A, *et al.* Role of (18)F-FDG PET in patients with infectious endocarditis. *J Nucl Med* 2014; **55**:1093–1098.
- Vos FJ, Bleeker-Rovers CP, Sturm PD, Krabbe PF, van Dijk AP, Cuijpers ML, *et al.* 18F-FDG PET/CT for detection of metastatic infection in gram-positive bacteremia. *J Nucl Med* 2010; **51**:1234–1240.
- Thiene G, Basso C. Pathology and pathogenesis of infective endocarditis in native heart valves. *Cardiovasc Pathol* 2006; **15**:256–263.
- Rouzet F, Chequer R, Benali K, Lepage L, Ghodbane W, Duval X, *et al.* Respective performance of 18F-FDG PET and radiolabeled leukocyte scintigraphy for the diagnosis of prosthetic valve endocarditis. *J Nucl Med* 2014; **55**:1980–1985.
- Erba PA, Conti U, Lazzeri E, Sollini M, Doria R, de Tommasi SM, *et al.* Added value of 99mTc-HMPAO-labeled leukocyte SPECT/CT in the characterization and management of patients with infectious endocarditis. *J Nucl Med* 2012; **53**:1235–1243.