Abstract. Paediatric invasive fungal infections have significantly increased over the past few decades, in particular among the immunocompromised population. Candida and Aspergillus spp. are still the most commonly isolated organisms. Image-based assessment of fungal infections can indeed be challenging especially in oncological patients where the differential diagnosis relative to other infections and neoplastic lesions cannot be often obvious. Therefore, the knowledge of the main radiological features associated with fungal infections is crucial to achieve an early correct diagnosis and address the most appropriate therapeutic approach. Thus, our aim was to review the main radiological features of paediatric fungal infections with particular focus on positron emission tomography/magnetic resonance imaging (PET/MRI), referring to the experience of our tertiary level hospital.

Invasive fungal disease (IFD) can be a life-threatening condition, in particular in paediatric patients who undergo chemotherapeutic or immunosuppressive treatments, in children with congenital or acquired immunodeficiencies, in premature neonates and in patients with fixed prostheses or catheters. IFD incidence has increased over the past few decades among the paediatric population, due to the improvement of oncological therapies which have significantly increased the survival but lead to an inevitable weakening of immune response. The most frequent pathogens involved are Candida albicans/non-albicans spp, Aspergillus spp and Cryptococcus neoformans, Histoplasma capsulatum, Zygomycetes, Fusarium, and other pathogens (1, 2). Fungal infections have a significant impact on patients’ outcome, with a mortality rate of about 15.8% for children affected by candidemia and about 18% for children with invasive aspergillosis (3). Therefore, an early diagnosis is essential to assure a prompt treatment. Since clinical manifestations are often non-specific, the radiological examination plays a key role for the detection of IFD and for the assessment of the disease extent. In addition to conventional imaging [ultrasound (US), plain X ray films, computed tomography (CT) and magnetic resonance imaging (MRI)], in the last years an increasing interest has been developed in hybrid whole-body imaging techniques [i.e. positron emission tomography/CT (PET/CT), PET/MRI], in particular in PET/MRI which combines the benefits of high soft tissue contrast resolution of MR with the metabolic information of PET, providing also the intrinsic advantage of a low dose radiation exposure. Therefore, integrated, simultaneously acquired PET/MRI can be an innovative and highly promising diagnostic tool for the evaluation of systemic fungal infections in paediatric patients. Nevertheless, the differential diagnosis with other infections and with malignant lesions can be challenging and histopathological confirmation is still strongly required when feasible to provide a definite diagnosis.

PET/CT and the Potential Role of PET/MRI

Whole-body hybrid imaging techniques such as 18F-labeled fluoro-2-deoxyglucose (18F-FDG)-PET/CT and PET/MRI can allow an early diagnosis of fungal infection foci mainly because of the high glucose consumption of these lesions, also providing a morphological definition of the disease extent. Moreover, the combination of the metabolic and the morphological information can be crucial in guiding the
surgeons to choose the best target for a biopsy and to assess
the response to antifungal treatment, especially if no
morphological modifications are seen on conventional imaging.
Nevertheless, the characterization of hypermetabolic areas can
be difficult when using PET/CT because of the low contrast
resolution especially in case of soft tissues involvement and
besides PET/CT exposes children to a higher dose of radiation
compared to CT alone MRI. Therefore, in the last few years,
an increasing interest has been developed towards hybrid
PET/MRI because of its potential to overcome these limits (4).

Lung

The lungs are the most commonly affected site in IFD and
fungal pulmonary involvement is associated with high
mortality. Aspergillus is the most common pathogen found
in fungal pulmonary infections, in particular in
haematological malignancies. The clinical manifestations
include fever, cough, dyspnea and haemoptysis (5). Chest X-
ray examination is the first-choice imaging modality but it
often shows non-specific signs, such as ill-defined
parenchymal opacities usually in both lungs and pleural
effusion (Figure 1); sometimes well-defined and cavitated
nodules can be evident (Figure 2).

The radiological standard of reference in pulmonary
fungal involvement is the chest CT scan; common findings
in invasive aspergillosis are the presence of multiple
bilateral nodules usually surrounded by ground glass opacity
(halo sign) (Figure 3) and wedge-shaped areas of
consolidation corresponding to haemorrhagic infarcts. The
air-crescent sign, due to the central necrotic area inside a
consolidation, can also be seen in some cases of aspergillosis
during recovery (Figure 4). In candidiasis CT scan, bilateral
pulmonary consolidations with possible cavitation and
nodules can be found as well (Figure 5) (2).

The reverse halo sign instead, defined as the presence of
ground glass attenuation surrounded by consolidation, can be
more frequently seen in haematological patients with Mucor
mycosis (2).

Lung visualization with MRI is definitely still a challenge,
even though in recent years the introduction of lung-
dedicated gradient echo (GE) sequences with short echo time
(TE) (such as VIBE, UTE) in PET/MRI protocols has
improved the chances for the detection of pulmonary fungal
involvement. Parenchymal consolidations and nodules can be
seen as relatively high intensity signal lesions depending
on their composition (proper solid lesions appear more
hypointense than ground glass lesions in low TE gradient
echo sequences because of their higher density); excavations
are usually seen as hypointense areas within (Figures 6 and
7). Furthermore, the combination with PET, adding the
metabolic information, improves the detection of pulmonary
fungal involvement in case of avid lesions and allows the
assessment of treatment response. Indeed, the evidence of a
decrease in the hypermetabolic activity of fungal foci after
antifungal therapy helps to confirm the diagnosis especially
when no morphological changes within the lesion are seen
and demonstrates the efficacy of the treatment protocol (6).

Figure 1. Chest X-ray showing bilateral lung parenchymal
consolidations and left pleural effusion in a 12-year-old girl, with
pulmonary aspergillosis.

Figure 2. Chest X-ray demonstrating two cavitated lesions (arrows) in
the right lung in a 16-year-old boy with pulmonary aspergillosis.
Nasal Cavities and Sinuses

Nasal and paranasal fungal infections are quite common in immunocompromised paediatric patients with a mortality rate of approximately 50%. The clinical signs and symptoms include fever, headache, facial pain and oedema, nasal congestion and epistaxis.

Middle turbinate, ethmoid cells, maxillary and sphenoid sinuses are more frequently involved. The assessment of the disease extent is crucial in these sites especially because of the risk of CNS involvement. CT scan usually shows thickening of the nasal or paranasal mucosa, with infiltration of the periantral fat planes. In late disease, bone erosion with orbital invasion and intracranial extension can be seen. On MRI, in addition to the mucosal thickening and fat planes infiltration, focal loss of contrast enhancement can be appreciated after gadolinium injection in the infected sino-nasal mucosa (Figure 8). The functional information given by PET can improve the diagnostic accuracy of MRI and also provides a valuable tool for treatment response evaluation detecting a potential FDG uptake reduction or disappearance (7-9).

Central Nervous System

Fungal infections affecting the central nervous system have a very poor prognosis with high mortality, especially in immunocompromised patients (almost 100%). The brain is the third most common site of fungal infection in patients with haematological malignancies after lungs and paranasal sinuses. Clinical symptoms are non-specific (confusion, hemiparesis, dysarthria, lethargy, seizures, possible fever) but their presence in immunocompromised children should suggest the diagnosis. Two main patterns can be present: focal lesions or diffuse meningoencephalitis. Fungal brain focal lesions are more common in Candida and Aspergillus infections; they are frequently multiple and usually involve the basal ganglia. CT findings are non-specific for fungal infection, such as focal hypodensities with a peripheral rim of contrast enhancement and often surrounded by oedema. On MRI, fungal abscesses show a central core, hypointense on T1-weighted and hyperintense on T2-weighted sequences, while the peripheral rim is iso-hyperintense on T1-weighted and hypointense on T2-weighted images. Peripheral enhancement after gadolinium injection and signal restriction on diffusion-weighted images (DWI) usually help to characterize the lesions (Figure 9). Fungal meningoencephalitis is most frequently caused by Cryptococcus, followed by Aspergillus and Candida. In encephalitis, CT scan of the brain may show ill-defined hypodense lesions with areas of increased attenuation, due to haemorrhage or increased metal ions. On MRI examinations, the infected areas are iso-hypointense on T1-weighted, hyperintense on T2-weighted/FLAIR images and with signal...
restiction on DWI. Micro-haemorrhages and focal deposits of metals are usually found within the lesions as decreased signal intensity on T2-weighted sequences. In fungal meningitis, radiological examinations show meningeal thickening and enhancement, both on CT and MRI, but the demonstration of fungi in CSF is necessary for the diagnosis. On PET, the visualization of lesions showing FDG uptake may be difficult due to the high intrinsic hypermetabolism of the cerebral cortex. Nevertheless, in case of fungal abscesses, especially when localized in the white matter, the combination of PET and MRI might help their detection, and again giving information about their response to treatment (Figures 10 and 11) (10, 11).

Liver and Spleen

Liver and spleen can be involved in paediatric IFD, in particular in children with haematological malignancies. Candida species are the most frequent pathogens affecting these organs with a mortality rate up to 30% (12). US can show four different patterns of fungal infection: an hypoechoic necrotic lesion surrounded by a hyperechoic rim of inflammatory cells and an external fibrotic hypoechoic ring, an inner hyperechoic nidus surrounded by a hypoechoic fibrotic ring; hypoechoic lesion that is expression of fibrotic changes after inflammation; hyperechoic scar or calcification (13). CT scan usually shows rounded hypodense lesions, sometimes with a hyperdense core. These nodules can demonstrate contrast enhancement of the whole lesion or an enhanced peripheral rim (14). On MRI, these lesions appear slightly hypointense on T1w images and strongly
Figure 8. A 11-year-old boy affected by Mucor mycosis: a large mycotic lesion can be seen in the right maxillary sinus (A), causing diffuse bone erosion of the sinus medial wall (B) and intrathecal spread of the fungal infection (arrow in C) surrounded by oedema. Orbital cellulitis can be also appreciated (D), involving the right medial rectus muscle.
hyperintense on T2w sequences with an intense and early peripheral enhancement after gadolinium injection. Some studies have demonstrated that $^{18}$F-FDG-PET may be a feasible and sensitive tool for the detection of liver and splenic fungal lesions and the assessment of residual disease if compared with conventional imaging. Therefore, the integrated PET/MRI, especially in paediatric patients, can play an important role (15).

Figure 9. Fungal brain abscess (arrows) in a 16-year-old boy characterized by central restricted diffusion on ADC map (A) and peripheral rim of enhancement after Gadolinium (B).

Figure 10. Candida cerebral abscess (arrows) characterized by surrounding oedema on T2-weighted image (A) and significant hypermetabolism on fused PET/MRI (B) in a 17-year-old boy.
Other Organs

Rarely, other organs can be affected by IFD. The hematogenous spread of pathogens makes possible the localization of infection also in soft tissues (skin, subcutaneous tissue, muscles), bones, kidney, etc. In these cases, a whole-body scan with PET/MR can reasonably help the detection of uncommon sites of infection (Figures 12-14).
Figure 13. A hypermetabolic Aspergillus lesion in the subcutaneous tissue (arrow) on fused PET/MR image, in a 12-year-old girl.

Figure 14. Left kidney Aspergillus lesion (arrows) in a 16-year-old girl on MRI T2-weighted and fused PET/MR images at diagnosis (A and C) and after four weeks of antifungal treatment (B and D): the lesion shows a slight reduction in size and a significant decrease in metabolic activity.
Conclusion

The knowledge of the most common radiological features of IFD is crucial to obtain a correct and early diagnosis. Nevertheless, the differential diagnosis with other infections and with neoplastic lesions can be challenging since fungal infections often do not show specific radiological characteristics; the histological examination is still frequently required when feasible for a definite diagnosis. Whole-body \(^{18}\)F-FDG-PET/MRI allows the simultaneous acquisition of both morphological and metabolic information with a low-dose radiation exposure, which represents a significant advantage in comparison with PET-CT especially among the paediatric population. PET/MRI can indeed be considered a useful diagnostic tool in the assessment of disease extent and the evaluation of treatment response in paediatric patients affected by IFD.

Conflicts of Interest

The Authors declare that no honorarium, grant nor other form of payment was given to anyone to produce the manuscript. The Authors declare that they have no potential conflicts of interest to disclose.

Authors’ Contributions

Varotto A, Olsatti G, Crimi F: review design, data collection, and manuscript drafting; Cecchin D, Toffolutti T: data collection, and manuscript revision; Stramare R, Zucchetta P: review design, and manuscript drafting; Varotto A, Orsatti G, Crimi F: review design, data collection, and manuscript revision; Cecchin D, Toffolutti T: data collection, and manuscript revision.

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