



Vasant, J. A., Maggiani, F., & Borman, A. (2017). Subcutaneous Mycotic Cyst Caused by *Roussoella percutanea* in a UK Renal Transplant Patient. *Mycopathologia*, *182*(7-8), 721-725. https://doi.org/10.1007/s11046-017-0121-0

Peer reviewed version

Link to published version (if available): 10.1007/s11046-017-0121-0

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## Mycopathologia

# Subcutaneous mycotic cyst caused by Roussoella percutanea in a UK renal transplant patient. --Manuscript Draft--

Manuscript Number:	
Full Title:	Subcutaneous mycotic cyst caused by Roussoella percutanea in a UK renal transplant patient.
Article Type:	Short Communication
Keywords:	Roussoella percutanea; emerging pathogen; rDNA sequencing; subcutaneous infection; renal transplant; Pleosporales.
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Funding Information:	
Abstract:	Fungi from more than one hundred genera have been implicated in subcutaneous fungal infections, usually following traumatic inoculation of the etiologic agent. With the advent of molecular approaches to fungal identification and taxonomy, novel agents of subcutaneous mycoses are increasingly reported. In this manner, Roussoella percutanea, a novel species in Pleosporales, was described in 2014 from a subcutaneous mass in an immunocompetent male adult. A second case was discovered after analysis of historical culture collection isolates, from a pedal mass in a renal transplant patient. Here we describe a new case of subcutaneous R. percutanea infection, causing a mycotic cyst in a renal transplant patient. Although fungal infection was confirmed histologically, viable fungal isolates could not be recovered in culture from biopsy material and identification of the causative agent relied upon PCR amplification and sequencing of fungal rDNA genes. This is only the third reported case of human infection with R. percutanea worldwide, the second in a renal transplant patient, and the first from a patient resident in the UK. The current case illustrates the importance of molecular approaches for the identification of emerging fungal pathogens in culture-negative subcutaneous fungal infections.
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7 8 9	4	Julia A. Vasant <sup>1</sup> , Francesca Maggiani <sup>2,3</sup> and Andrew M. Borman <sup>4</sup>
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26 ABSTRACT

Fungi from more than one hundred genera have been implicated in subcutaneous fungal infections, usually following traumatic inoculation of the etiologic agent. With the advent of molecular approaches to fungal identification and taxonomy, novel agents of subcutaneous mycoses are increasingly reported. In this manner, *Roussoella percutanea*, a novel species in Pleosporales, was described in 2014 from a subcutaneous mass in an immunocompetent male adult. A second case was discovered after analysis of historical culture collection isolates, from a pedal mass in a renal transplant patient. Here we describe a new case of subcutaneous *R. percutanea* infection, causing a mycotic cyst in a renal transplant patient. Although fungal infection was confirmed histologically, viable fungal isolates could not be recovered in culture from biopsy material and identification of the causative agent relied upon PCR amplification and sequencing of fungal rDNA genes. This is only the third reported case of human infection with R. percutanea worldwide, the second in a renal transplant patient, and the first from a patient resident in the UK. The current case illustrates the importance of molecular approaches for the identification of emerging fungal pathogens in culture-negative subcutaneous fungal infections.

44 191 words

### 47 Introduction

Subcutaneous fungal infections usually occur following traumatic inoculation of the etiological agent into cutaneous or subcutaneous tissue. Although infections typically remain localised in immunocompetent individuals, without treatment they frequently become chronic, with gradual expansion to involve adjacent tissues and even bone (1-3). This heterogeneous disease spectrum includes hyalohyphomycoses (infection by fungi with hyaline hyphae), phaeohyphomycoses (infection with melanised, dematiaceous fungi), chromoblastomycoses (characterised by formation of sclerotic bodies in tissue) and eumycetoma (characterised by extensive tumefaction with formation of purosanguinous sinuses extruding fungal grains) (2-5). 

Fungi from more than 100 genera have been reported from subcutaneous mycoses in humans, with novel species and genera described almost weekly (5). In some cases, the responsible fungi have only been reported from individual, isolated case reports suggesting that they are infrequent agents of subcutaneous infection (6-9). For many other species, they have been predominantly associated specifically with precise clinical presentations and have been rarely or never isolated from nature (2, 4, 5, 10-12). Although certain species are predominantly associated with a specific clinical presentation, clinical picture is also determined by host immune status, with the result that accurate definition of disease entity may be difficult (13, 14). In addition, many of the agents of subcutaneous mycoses exhibit reluctant conidiation in culture, especially when isolated from chronic cases (5, 10, 12, 14). Since accurate identification of the causative agent is paramount in directing appropriate therapeutic interventions, molecular approaches using PCR amplification and sequencing of conserved regions of fungal genomic DNA is increasingly employed (14-17). Using such approaches, Roussoella percutanea, a novel species in Pleosporales, was recently described 

from a subcutaneous mass in an immunocompetent male adult (18). Clinical presentation included hyphal masses with granulation tissue and fibrosis, but draining sinuses typical of eumycetoma were absent. A retrospective analysis of other culture collection isolates revealed one further historical isolate of R. percutanea, from a pedal mass in a renal transplant patient who also presented with necrosis and inflammation but without draining sinuses or fungal grains (13,18). In both cases, conidiation in culture was delayed or absent, and variable antifungal susceptibility profiles were reported for the two previous isolates (18).

Here we present the first case of *R. percutanea* infection from the UK, involving a subcutaneous cyst from a renal transplant patient. Attempts to culture the organism from cyst tissue were unsuccessful and diagnosis and identification were achieved by PCR amplification and sequencing of fungal genomic DNA from biopsy material.

#### 84 Case History

A 47 year old male, originally from Surat, Gujarat, but residing in the UK since 1997, underwent a renal transplant in November 2015 for end-stage renal failure of unknown cause. Post-operatively he was immune-suppressed with oral prednisolone 12.5mg once a day and tacrolimus 5mg twice daily and commenced prophylactic oral co-trimoxazole and aciclovir. Other past medical history included gout, osteoarthritis, an inguinal hernia and migraines.

Post-transplant he suffered from an episode of declining renal function secondary to
borderline T-cell mediated rejection, requiring two rounds of pulsed methylprednisolone and
addition of mycophenylate mofetil 500mg twice daily to his immunosuppressive regimen
from January 2016. His clinical course was also complicated by multiple episodes of
urosepsis, including one episode of bacteraemia, due to an extended-spectrum β-lactamase
producing *Klebsiella pneumoniae*, between January and May 2016.

During admission for an episode of *Klebsiella* urosepsis in March 2016 an incidental finding was made of a mobile, soft tissue mass over the right Achilles tendon. The lesion was non-tender and not adherent to the tendon itself with no masses elsewhere or evidence of lymphatic involvement or sinus tract formation. The patient had been unaware of the lesion and it had not caused him any pain or symptoms. Clinical suspicion of mycetoma was raised. Further travel history included trips to Dubai 2005 and Morocco in 2014. The patient could not re-call any specific injuries or possible inoculation events.

An ultra-sound scan confirmed a 2 by 1.5 centimetre heterogeneous nodule overlying but separate from the Achilles tendon. A biopsy of the lesion was performed which did not reveal any organisms on Gram stain and there was no growth by conventional bacterial culture (including Sabouraud's dextrose agar). Histology revealed a diffuse macrophage infiltrate with cytoplasmic inclusions and fungal hyphae were seen on Periodic acid-Schiff and Periodic acid-Schiff-diastase stains. Empirical antifungal treatment was not started at this point due to the patient's stable clinical condition.

Microscopic analysis of biopsy material by potassium hydroxide digestion with Calcofluor enhancer (Bactidrop, Remel) revealed the presence of moderate amounts of filamentous fungal elements. These were generally amorphous, and included fine, regularly septate and acutely branching hyphae, which were hyaline to light brown together with more common short chains of swollen cells and chlamydospores. Despite prolonged culture on a variety of mycological media, no viable fungal isolate could be recovered for further analyses. Small sections of biopsy tissue were subjected to digestion and Qiagen column purification (Qiagen Mini Blood kit) and PCR amplification of the 28S rDNA and Internal transcribed spacer 1 (ITS1) regions of the resulting extracted fungal genomic DNA using the PCR primers and conditions described previously (12, 14, 17). The sequences of the resulting PCR amplicons were 100% identical to those of the type species of Roussoella percutanea 

(isolate CBS128203; GenBank accession numbers KF366448 and KF322117) present in the public synchronised databases. Treatment with oral voriconazole 200mg twice a day was then immediately commenced and total excision of the lesion took place with 5mm margins. Histology of the excised lesions confirmed fungal hyphae on both Hematoxylin/Eosin and Grocott stains. Unfortunately, due to clinical concern regarding potential interactions between voriconazole and tacrolimus, and a falling neutrophil count (to a trough of 2.1x10<sup>9</sup>/litre) whilst being treated with voriconazole, anti-fungal treatment was discontinued after only 2 weeks. Despite this, the patient's wound is healing and dry at 6 weeks post-excision with no evidence of recurrence so far.

We report here the first known UK case of subcutaneous infection by *R. percutanea*. To our knowledge, this is only the third case reported worldwide, and the second to have affected a renal transplant patient (13,18). In all three cases, presentation differed from that seen with eumycetoma due to the presence only of inflammatory infiltrates and discrete fungal elements and absence of fungal grains or draining sinuses, Subcutaneous mycoses in immunocompetent patients frequently present diagnostic challenges due to the heterogeneous nature of clinical presentations and the fact that clinically detectable infections can take years to develop. This situation is further compounded in transplant patients, where relatively dormant fungi that have been harboured subcutaneously for many years or decades become clinically active after immunosuppression. Although most subcutaneous mycoses are thought to result from traumatic inoculation, it is hardly surprising given this protracted and variable incubation period that most patients are unable to recall any history of injury at the affected sites. This was the case with the current patient and with the two previous reports of R. *percutanea* subcutaneous mycoses published to date (13, 18).

Previous reports have suggested strain-specific variations in antifungal susceptibility in R. percutanea (18), indicating that specific testing of patient isolates should be used to guide antifungal therapeutic decisions. Since the current case was culture-recalcitrant and no isolate was available for testing, the patients was treated with total excision of the lesion and commenced on voriconazole therapy. Unfortunately, antifungal therapy was stopped after only 2 weeks due to drug interactions. Although the patient is currently well, close follow up will be required, especially in the light of the reports of recurrences in both previous cases of R. percutanea (13,18). Finally, the current case further highlights the utility of molecular diagnostic and identification approaches in medical mycology, especially in cases of subcutaneous fungal infections where conventional identification approaches are hindered by 

difficulties in recovering viable fungi from biopsy samples and organisms that fail to producedistinctive features in culture.

References 1. Romano C. Subcutaneous mycoses. Part 1: subcutaneous mycoses due to non-dermatophytes. G Ital Dermatol Venereol. 2013; 148: 621-631. 2. Garnica M, Nucci M, Queiroz-Telles F. Difficult mycoses of the skin: advances in the epidemiology and management of eumycetoma, phaeohyphomycosis and chromoblastomycosis. Curr Opin Infect Dis. 2009; 22: 559-563. 3. Queiroz-Telles F, McGinnis MR, Salkin I, Graybill JR. Subcutaneous mycoses. Infect Dis Clin N Am 2003; 17:59-85. 4. Isa-Isa R, García C, Isa M, Arenas R. Subcutaneous phaeohyphomycosis (mycotic cyst). Clin Dermatol. 2012; 30: 425-31. 5. de Hoog GS, Guarro J, Gené J, Figueras MJ. Atlas of Clinical Fungi. 2nd edn.Utrecht, The Netherlands: Centraalbureau voor Schimmelcultures; 2004. 6. Mhmoud NA, Ahmed SA, Fahal AH, de Hoog GS, Gerrits van den Ende AH, van de Sande WW. Pleurostomophora ochracea, a novel agent of human eumycetoma with vellow grains. J Clin Microbiol. 2012; 50: 2987-2994. 7. McGrogan D, David MD, Roberts C, Borman AM, Nath J, Inston NG, Mellor S. Pseudotumoral presentation of fungating mycetoma caused by Phaeoacremonium fuscum in a renal transplant patient. Transpl Infect Dis. 2015; 17: 897-903.

8. Ponnuswamy K, Muthureddy Y, Sigamani K. Two cases of multiple subcutaneous cystic phaeohyphomycosis in immunocompromised patients with a rare causative organism. Indian J Dermatol. 2014; 59: 421.

- 9. Ge YP, Lv GX, Shen YN, Li M, Deng SW, De Hoog S, Samerpitak K, Liu WD. First report of subcutaneous phaeohyphomycosis caused by Ochroconis tshawytschae in an immunocompetent patient. Med Mycol. 2012; 50: 637-640. 8 10. Ahmed SA, van de Sande WW, Stevens DA, Fahal A, van Diepeningen AD, Menken SB, de Hoog GS. Revision of agents of black-grain eumycetoma in the order Pleosporales. Persoonia. 2014; 33: 141-154. 11. van de Sande WW. Global burden of human mycetoma: a systematic review and meta-analysis. PLoS Negl Trop Dis. 2013; 7 :e2550. 12. Borman AM, Desnos-Ollivier M, Campbell CK, Bridge PD, Dannaoui E, Johnson EM. Novel taxa associated with human fungal black-grain mycetomas -Emarellia grisea gen nov. et sp. nov. and Emarellia paragrisea sp. nov. J Clin Microbiol. 2016; 54: 1738-1745. 13. Meis JF, Schouten RA, Verweij PE, Dolmans W, Wetzels JF. Atypial presentation of Madurella mycetomatis mycetoma in a renal transplant patient. Transpl Infect Dis 2000; 2: 96-98. 14. Abdolrasouli A, Gonzalo X, Jatan A, McArthur GJ, Francis N, Azadian BS, Borman AM, Johnson EM. Subcutaneous phaeohyphomycosis cyst associated with Medicopsis romeroi in an immunocompromised host. Mycopathologia. 2016; 181: 717-721. 15. Santos DW, Padovan AC, Melo AS, Gonçalves SS, Azevedo VR, Ogawa MM, Freitas TV, Colombo AL. Molecular identification of melanised non-sporulating moulds: useful studying epidemiology of a tool for the phaeohyphomycosis. Mycopathologia 2013; 175: 445-454.

1	204	16. Ahmed AO, Desplaces N, Leonard P, Goldstein F, de Hoog S, Verbrugh H, van
2 3	205	Belkum A. Molecular detection and identification of agents of eumycetoma: detailed
4 5 6	206	report of two cases. J Clin Microbiol 2003; 41: 5813-5816.
7 8	207	17. Borman AM, Linton CJ, Miles SJ, Johnson EM. Molecular identification of
9 10	208	pathogenic fungi. J Antimicrob Chemother. 2008; 61 Suppl 1:i7-12,
11 12 13	209	18. Ahmed SA, Stevens DA, van de Sande WW, Meis JF, de Hoog, GS. Roussoella
14 15	210	percutanea, a novel opportunistic pathogen causing subcutaneous mycoses. Med
16 17 19	211	Mycol. 2014; 52: 689-698.
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28 29 30	216	Figure Legend.
31 32	217	Figure 1. Histopathological appearance of biopsy material excised from the subcutaneous
33 34 25	218	cyst. H&E stain. Panel B is a magnification of a section of panel A. Scale Bars = 5 $\mu$ m.
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