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Rapid Diagnosis of Azole-Resistant Aspergillosis by Direct PCR Using Tissue Specimens\textsuperscript{V}

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We report the use of PCR techniques on a formalin-fixed and paraffin-embedded tissue specimen for direct detection of one dominant azole resistance mechanism in a case of disseminated invasive aspergillosis. Rapid detection of mutations associated with azole resistance directly in tissue significantly reduces diagnostic delay.

Invasive infections due to \textit{Aspergillus fumigatus} are associated with significant morbidity and mortality, although the prognosis of patients with invasive aspergillosis has improved with the clinical use of mold-active antifungal azoles, most notably voriconazole (9, 11). However, the survival of patients may be threatened by the emergence of azole resistance of aspergilli (1, 7, 13). Resistance is commonly due to point mutations in the \textit{cyp51A} gene, which is the target for antifungal azoles (1, 4, 8, 13, 14). The isolates commonly exhibit a cross-resistant phenotype (4), and patients with azole-resistant disease may fail azole therapy (1, 7, 10, 12). One problem in the management of azole-resistant aspergillosis is the early detection of resistance as cultures are negative in up to 50\% of patients with foci pulmonary lesions (2), and in vitro susceptibility testing takes at least 5 to 7 days to complete. In this report, molecular tools were utilized to rapidly confirm the diagnosis of disseminated azole-resistant aspergillosis.

\textbf{Case report.} A 60-year-old man was diagnosed with acute myeloid leukemia and underwent an allogeneic hematopoietic stem cell transplantation. Six months later, he developed grade 2 to 3 graft-versus-host disease (GVHD) of the gastrointestinal tract and skin. He was treated with 1 mg prednisone/kg/day plus 3 mg b.i.d. Entocort. His neutrophil count was 0.77 \texttimes 10^9/liter. Diarrhea responded quickly. However, 10 days later, he developed fever, and high-resolution computed tomography (CT) of the thorax showed the presence of multiple nodular pulmonary infiltrates. \textit{Aspergillus fumigatus} was cultured from the sputum, and circulating galactomannan was detected in serum. Under the suspicion of invasive pulmonary aspergillosis, treatment with voriconazole was started immediately, and the prednisone dose was halved and, within a few weeks, further reduced to a maintenance dose of 5 mg daily. The GVHD did not relapse, but as the clinical condition of the patient did not improve and sputum cultures remained positive with \textit{A. fumigatus}, caspofungin was added. One week after the start of the antifungal therapy, paresis of the abducens nerve developed, and brain magnetic resonance imaging (MRI) showed multiple lesions. A brain biopsy was performed, and histological examination showed septate hyphae (Fig. 1); however, cultures remained negative. The sputum isolate was resistant to itraconazole and voriconazole and intermediate susceptible to posaconazole (Table 1) (13). As the cerebral lesion might be due to azole-resistant \textit{A. fumigatus}, it was decided to replace voriconazole by liposomal amphotericin B. The patient again developed fever, and posaconazole was added to the treatment regimen.

We utilized the sputum isolate and three formalin-fixed and paraffin-embedded tissue sections from the brain biopsy for identification of the resistance mechanism. Amplification of the 28S ribosomal DNA confirmed the presence of \textit{Aspergillus} in the biopsy samples. Sequence-based analysis of the \textit{cyp51A} gene (using reference sequence of strain AF338659 from GenBank) of the sputum isolate showed a substitution of the gene promoter region (TR+1-L98H). The morphological species identification of \textit{A. fumigatus} was confirmed by sequencing of the highly conserved \textit{\beta}-tubulin gene (7). Two PCR assays (Table 2) targeted at the tandem repeat and the mutation at codon 98 were performed directly on the brain biopsy specimen and were both positive (Table 1). DNA was extracted using proteinase K and the Qiagen EZ1 robot (Qiagen, The Netherlands). For the real-time detection of L98H, a 122-bp region was amplified, bridging codon 98 of the \textit{cyp51A} gene by using a forward primer and a reverse primer (Table 2). Using hybridization probes, the wild-type L98 codon could be differentiated from the mutated L98H codon. A sensor probe was designed to span the mutation and to be separated from an anchor probe by 1 nucleotide. The melting temperature (T\textsubscript{m}) of the sensor probe was approximately 6.6°C lower than that of the anchor probe; a wild-type DNA template will result in a mismatch giving a lower T\textsubscript{m} than that observed for the mutated DNA template. The LightCycler FastStart DNA Master Hybridization Probe reaction mix was used.

To detect the 34-bp repetitive insertion, a TaqMan probe (TR) (Table 2) was designed to bind to the last 13 bp and the first 9 bp of the repetitive insertion, thereby binding specifically to a mutated DNA template. To control for a successful am-
The clinical condition of the patient gradually improved, and imaging showed regression of the pulmonary and brain lesions. Liposomal amphotericin B and caspofungin therapy was discontinued after 12 weeks. Posaconazole was continued, and a plasma level of 1.7 mg/liter was achieved. Repeat CT of the brain showed further improvement of the lesions 6 months after initiation of antifungal therapy.

Azole resistance has emerged in The Netherlands, and the prevalence among clinical A. fumigatus isolates was 6% in 2007 (7). Resistance significantly complicates the treatment of inva-

FIG. 1. Histology of the brain biopsy specimen showing septate hyphae (PAS staining, ×400 magnification).

TABLE 1. Results of the diagnostic assays for the clinical specimens obtained from the patient and for three controls

<table>
<thead>
<tr>
<th>Date</th>
<th>Sample</th>
<th>Microscopy</th>
<th>Culture</th>
<th>Molecular identification</th>
<th>Detected cyp51A substitution</th>
<th>MIC (mg/liter)</th>
</tr>
</thead>
<tbody>
<tr>
<td>26 September</td>
<td>Sputum</td>
<td>Not done</td>
<td>A. fumigatus</td>
<td>A. fumigatus</td>
<td>TR + L98H</td>
<td>0.5 &gt; 16 4 0.5 0.25</td>
</tr>
<tr>
<td>2008</td>
<td>Brain biopsy sample</td>
<td>Septate hyphae</td>
<td>Negative</td>
<td>A. fumigatus</td>
<td>TR + L98H</td>
<td>0.5 &gt; 16 4 0.5 0.5</td>
</tr>
<tr>
<td></td>
<td>Lung biopsy sample</td>
<td>Septate hyphae</td>
<td>Negative</td>
<td>A. fumigatus</td>
<td>TR + L98H</td>
<td>0.25 &gt; 16 4 0.5 0.25</td>
</tr>
<tr>
<td></td>
<td>(control)</td>
<td></td>
<td>Negative</td>
<td>A. fumigatus</td>
<td></td>
<td>0.5 &gt; 16 4 0.5 0.063 0.125</td>
</tr>
<tr>
<td></td>
<td>Brain biopsy sample</td>
<td>Negative</td>
<td>Negative</td>
<td>A. fumigatus</td>
<td></td>
<td>0.5 &gt; 16 4 0.5 0.063 0.125</td>
</tr>
<tr>
<td></td>
<td>(control)</td>
<td></td>
<td>Negative</td>
<td>Negative</td>
<td></td>
<td>0.5 &gt; 16 4 0.5 0.063 0.125</td>
</tr>
</tbody>
</table>

a Molecular identification by sequencing the 28S ribosomal DNA directly on isolates if cultures were positive or directly on the tissue when cultures were negative.

b For the sputum isolate, the full sequence of the cyp51A gene was determined. For the biopsy specimens, two PCR assays were used to detect the tandem repeat and L98H substitution directly on formalin-fixed, paraffin-embedded biopsy samples. TR, 34-bp tandem repeat; L98H, substitution of leucine for histidine at codon 98.

c MIC determined using the CLSI M38-A2 protocol. AMB, amphotericin B; ITZ, itraconazole; VCZ, voriconazole; POS, posaconazole; CAS, caspofungin. For caspofungin, the minimum effective concentration was determined.
The detection of resistance in invasive aspergillosis may be problematic, as positive cultures are required to test for a resistant phenotype, while the infecting isolate is obtained in only a minority of cases. However, even if *Aspergillus* is cultured, adequate treatment may be delayed for 5 to 7 days due to the time needed for MIC results to become available. Given a median survival of central nervous system aspergillosis of only 10 days (6), this is an unacceptable delay. Although in our patient resistance was expected early due to MIC testing of the isolate, we were able to diagnose azole-resistant dissemination directly in the brain biopsy by using specific real-time PCR assays. This appears to be a feasible approach, as mutations in the *cyp51A* gene are the primary mechanism of azole resistance in *A. fumigatus*, and the TR+L98H is the dominant change in Dutch azole-resistant *A. fumigatus* isolates (7). Provided that the resistance mechanism is known, molecular techniques are suitable to detect multiple resistance mechanisms simultaneously.

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### REFERENCES


### TABLE 2. PCR assays targeted at the tandem repeat and the mutation at codon 98

<table>
<thead>
<tr>
<th>Target</th>
<th>Primer</th>
<th>Sequence</th>
<th>Binding positions (bp)</th>
</tr>
</thead>
<tbody>
<tr>
<td>TR</td>
<td>Forward primer</td>
<td>5′-CAGCACCACCTCAGAGTTGTC-3′</td>
<td>8–29</td>
</tr>
<tr>
<td></td>
<td>Reverse primer</td>
<td>5′-GTGGATGGGATGCGACACCTCG-3′</td>
<td>109–84</td>
</tr>
<tr>
<td></td>
<td>TaqMan probe (TR)</td>
<td>3′-FAM-TGCTGACCGAATGTAACGC-MTM-5′</td>
<td>26–47</td>
</tr>
<tr>
<td></td>
<td>TaqMan probe (wild-type sequence)</td>
<td>3′-FAM-ATTAGGCACCTTCATGGCTCAGC-BBQ-3′</td>
<td>78–53</td>
</tr>
<tr>
<td>L98H</td>
<td>Forward primer</td>
<td>5′-CAAAACACACAGCTTACCTG-3′</td>
<td>658–678</td>
</tr>
<tr>
<td></td>
<td>Reverse primer</td>
<td>5′-GAATTGGGACAATCATACACC-3′</td>
<td>800–780</td>
</tr>
<tr>
<td></td>
<td>Sensor probe</td>
<td>5′-ACGGCAAGCACAAGGATG-3′, fluorescein-labeled</td>
<td>707–724</td>
</tr>
<tr>
<td></td>
<td>Anchor probe</td>
<td>5′-CAATGGCGAGAGGCTTATAGTCCATTGA-3′-LC-red 640-labeled</td>
<td>723–754</td>
</tr>
</tbody>
</table>

* TR, 34-bp tandem repeat; L98H, substitution of leucine for histidine at codon 98.

* a FAM, 6-carboxyfluorescein; TMR, 6-TAMRA (6-carboxytetramethylrhodamine); BBQ, 4,4-bis-(2-hutyloctyloxy)-p-quaterphenyl; LC-red 640-aminohexyl-spacer-5′-OH.

* c Binding positions of oligonucleotides on the basis of reference strain AF 338659.