Azole-Resistant Aspergillus fumigatus, Iran

To the Editor: Aspergillus fumigatus causes a variety of diseases in humans. The drugs recommended for treatment of Aspergillus diseases are the mold-active azole antifungal drugs (1). However, a wide range of mutations in A. fumigatus confer azole resistance, which commonly involves modifications in the cyp51A gene (2), the target for azole antifungal drugs.

Azole resistance is thought to be selected for as a result of patient therapy or exposure to azole compounds in the environment; resistance in clinical A. fumigatus isolates has been increasingly reported in several European countries, Asia, and the United States (3–7). The most frequently reported resistance mechanism is a 34-bp tandem repeat (TR34) in combination with a substitution at codon 98 (TR34/L98H) (4); this mechanism is believed to have been selected for through environmental exposure to azole fungicides.

Because routine in vitro susceptibility testing of clinical Aspergillus isolates is not common in many centers worldwide, the prevalence of azole resistance might be underestimated. We investigated the prevalence of azole resistance in clinical A. fumigatus isolates stored for 6 years (2003–2009) at Tehran University Mycology Reference Centre and Islamic Azad University, Ardabil Branch, Iran.

We investigated 124 clinical A. fumigatus isolates obtained from patients with Aspergillus diseases (online Technical Appendix Table 1, wwwnc.cdc.gov/EID/article/19/5/13-0075-Techapp1.pdf). We conducted strain identification, in vitro antifungal susceptibility testing, and sequence-based analysis of the Cyp51A gene, as described (4). We performed microsatellite genotyping of all A. fumigatus isolates for which the MIC of itraconazole was $\geq$16 mg/L (8) by using a short tandem repeat A. fumigatus assay, and we compared the results with those reported for the Netherlands (20 isolates) and other European countries (24 isolates) (online Technical Appendix Figure).

The distribution of azole-resistant and wild-type A. fumigatus isolates examined in this study, according to year of isolation, is shown in online Technical Appendix Table 1. Of 124 A. fumigatus isolates, 4 grew on the wells containing itraconazole and voriconazole, indicating a multidrug-resistant phenotype. Of these resistant isolates, 3 were from patients with chronic pulmonary aspergillosis and 1 was from a patient with allergic bronchopulmonary aspergillosis (Table).

Sequence analysis of the CYP51A gene indicated the presence of TR34/L98H in 3 isolates and no mutations in the other isolate (Table). The first TR34/L98H isolate had been recovered in 2005, which is relatively early compared with reported isolations in other countries (online Technical Appendix Table 2). Microsatellite typing of 6 short tandem repeat loci demonstrated identical patterns for 2 of the 3 azole-resistant isolates from Iran, but the TR34/L98H isolates from Iran did not cluster with those from the Netherlands and other European countries, indicating no close genetic relatedness (online Technical Appendix Figure).

The TR34/L98H azole resistance mechanism was first described in 1998 in the Netherlands; since then, its presence in clinical and environmental A. fumigatus isolates in multiple European countries and recently in Asia has been increasingly reported (online Technical Appendix Table 2) (3–7). In the study reported here, prevalence of azole resistance in clinical A. fumigatus isolates obtained from patients in Iran was 3.2%; most isolates exhibited identical patterns for 2 of the 3 azole-resistant isolates from Iran, but the TR34/L98H isolates from Iran did not cluster with those from the Netherlands and other European countries, indicating no close genetic relatedness (online Technical Appendix Figure).

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References


of azole resistance might be underestimated in many countries because in vitro susceptibility testing of *A. fumigatus* is not routinely performed.

Microsatellite genotypic analysis of *A. fumigatus* isolates from the Netherlands and various European countries showed that the genetic diversity of TR<sub>L98H</sub> isolates is lower than that of wild-type controls (8). It has been suggested that TR<sub>L98H</sub> isolates might have a common ancestor that developed locally and subsequently migrated across Europe. In contrast, genotyping of TR<sub>L98H</sub> originating from India suggested a different dynamic; all environmental and clinical TR<sub>L98H</sub> isolates from India shared the same multilocus microsatellite genotype not found in any other analyzed samples, from within India or from the Netherlands, France, Germany, or the People’s Republic of China (9). The molecular epidemiology of the TR<sub>L98H</sub> isolates from Iran suggests that they cluster apart from the European isolates, indicating that migration from Europe to Iran, or vice versa, is unlikely. Genotyping of more TR<sub>L98H</sub> isolates from the Middle East and comparison with those from India would enhance understanding of the origin and geographic spread of TR<sub>L98H</sub>.

Our study indicates that TR<sub>L98H</sub> was in Iran in 2005; this finding adds to the growing list of regions where acquired resistance in *A. fumigatus* of environmental origin is documented. From a global perspective, fungicide use is second highest in the Asia–Pacific regions (24%), preceded only by western Europe (37%) (10).

For a bettering understand of the scale of this emerging public health problem and for insight into the dynamics of geographic migration, surveys of fungal culture collections for TR<sub>L98H</sub> and molecular typing studies are warranted. These data would be useful not only for clinical management of *Aspergillus* diseases but also for enabling policy makers to develop strategies that prevent resistance selection by the environmental route.

**Acknowledgments**

We thank Jacques F. Meis and Ferry Hagen for performing additional real-time PCRs to screen mutations in 4 azole-resistant isolates.

This work was supported in part by a research grant from the Faculty of Medicine and Medical Sciences, Islamic Azad University, Ardabil Branch, Iran, and the School of Hygiene & Institute of Public Health Research, Tehran University of Medical Sciences, Tehran, Iran.

S.J.M., E.Z., M.T.H., J.Z., and W.J.G.M. have no conflicts of interest. S.S. received a research grant from the Faculty of Medicine and Medical Sciences, Islamic Azad University, Ardabil Branch, Iran. J.W.M. and P.E.V. have served as consultants to and have received research grants from Astellas, Basilea, Gilead Sciences, and Amphotericin B (AmBisome) from Astellas, Basilea, Gilead Sciences, and Janssen Pharmaceutica N.V. and have received research grants from Astellas, Basilea, Gilead Sciences, and Janssen Pharmaceutica N.V. J.W.M. and P.E.V. have served as consultants to and have received research grants from Astellas, Basilea, Gilead Sciences, and Janssen Pharmaceutica N.V. J.W.M. and P.E.V. have served as consultants to and have received research grants from Astellas, Basilea, Gilead Sciences, and Janssen Pharmaceutica N.V.

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DOI: [http://dx.doi.org/10.3201/eid1905.130075](http://dx.doi.org/10.3201/eid1905.130075)

**References**

DNA–positive. The Q fever outbreak in cats, DNA–positive bovine outbreak, 1 genotype of in the Netherlands affected ≈4,000 
ella burnetii 
ucleus caused by the bacterium Coxi- 
C. burnetii 
\[C \text{sufficient DNA load (cycle threshold }
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\]samples revealed a (partial) genotype, which is related to the outbreak genotype (sheep 192, Figure). Seven of the 
\]DNA was extracted from the al- lantochorion of the placenta and ana 
lized as described (2). Samples with sufficient DNA load (cycle threshold \[C \text{ value }<32\text{ were typed by using 2 multilocus variable-number tandem-repeat analyses (MLVA) genotyping methods (MLVA-12 and MLVA-6), and the multispacer sequence typing method (3–5). Two } C. burnetii \text{ strains from the Netherlands representing the outbreak genotype (X09003262, 3345937) and the Nine Mile RSA 493 were included as reference. For prevalence calculations, the Netherlands was divided in a southern part, comprising the Q fever hot spot area of notified cases in humans and small ruminants during the 2007–2010 epidemic (1,6), and a northern part, comprising the rest of the country. }
\[C. burnetii \text{ DNA was not detected in placentas from cats, goats, or pigs. } C. burnetii \text{ DNA was detected in 4 (7} \% [95\% \text{ CI 0.4–14.4}] \text{ of 54 canine pla}
\]from the north of the country, without known abortion history. } C. burnetii \text{ DNA was detected in 7 (26}\% [95\% \text{ CI 9.4–42.5}] \text{ of 27 ovine and in 33 (24}\% [95\% \text{ CI 16.7–30.8}] \text{ of 139 bovine placenta. The prevalence of } C. burnetii \text{ DNA–positive ovine and bovine placenta from the north and the south did not differ significantly.}

The } C. burnetii \text{ DNA load in the placentas from dogs (C, value 37.4–38.0) and horses (C, value 35.4–37.4) was too low to be suitable for genotyping. Typing of 1 positive sheep sample resulted in an incomplete genotype, which is related to the outbreak genotype (sheep 192, Figure). Seven of the 33 } C. burnetii \text{ DNA–positive bovine placenta were suitable for typing. One sample had a genotype similar to the outbreak genotype (2,3). Six other samples revealed a (partial) genotype related to bovine genotypes from the Netherlands (2,5,7), including a novel one. MLVA-6 and multispacer sequence typing results were consistent with the MLVA-12 results (Figure). Results give no indication for major reservoirs of } C. burnetii \text{ in cats, goats, and pigs in the Netherlands in 2011. However, the low numbers of placenta may have biased the results. Dogs and horses should be considered as reservoirs for } C. burnetii \text{. The detection of } C. burnetii \text{ DNA–positive placentas in dogs and horses in the northern part of the country indicates the presence of a true reservoir rather than a spillover effect from the contaminated environment in the south. This observation is consistent with a reported seroprevalence of 13\% in dogs in the Netherlands in 1992 (1). Until now, horses had been discussed as a risk factor in the Q fever outbreak in the Netherlands (8).}

Prevalence data from sheep and cattle suggest that } C. burnetii \text{ is present in placentas in 25\% of the abortion cases in these species. Presence of the outbreak genotype of } C. burnetii \text{ in sheep has been observed (2,5), indicating sheep are a reservoir for Q fe-