TRIAZOLE RESISTANCE IN ASPERGILLUS FUMIGATUS - A COMPREHENSIVE REVIEW

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ABSTRACT

Aspergillus fumigatus is a common ubiquitous fungal pathogen present in the environment including soil, decaying leaves, compost, grain crops and household dust. It can turn opportunistic in immunocompromised individuals causing allergic diseases, respiratory problems and invasive blood stream infections. Being saprophytic, it propagates by producing minute grey-green conidia that can become airborne and reach humans. In majority of cases, aspergillosis due to this organism can heal with treatment by antifungals. The azole compounds, one of the most important antifungal drugs, generally target the 14α-sterol demethylase enzyme, encoded by the cyp51A gene of the fungus, causing impaired membrane ergosterol synthesis. However in recent years the situation has turned grave due to development of azole resistant fungal species both in the environment as well as in the clinics. The main mechanism of resistance in A. fumigatus involves incorporation of tandem repeats in the promoter region of cyp51A, along with point mutations in the gene sequence. Resistant isolates tend to arise either due to excessive use of azole fungicides in the agricultural fields, from where they reach human through sporulation; or by prolonged treatment of patients withazole drugs. The early diagnosis of aspergillosis is very crucial for successful treatment of a patient. The article brings forth a comprehensive study of this organism, the mechanism of development and spread of azole resistant fungal isolates, genetic basis of development of azole resistance, detection of resistant strains and finally drugs used to combat the fungal infection.

Keywords: Antifungal, Aspergillosis, Azoles, CYP51A

INTRODUCTION

A. fumigatus is a saprophytic fungus that grows on dead and decaying parts of the organic matter. It is not a primary pathogen; however the use of immunosuppressive therapies or prolonged use of antifungals can turn it into an opportunistic fungal pathogen (Chowdhary et al., 2013). The genus Aspergillus is divided into five sections- fumigati, nidulante, flavi, niger, and terreii, but only few species such as A. fumigatus and A. flavus are known to be the major pathogenic fungi in humans and animals. A. fumigatus is the most common airborne opportunistic fungal pathogen. Individuals suffering from asthma or cystic fibrosis are at a major risk to develop allergic bronchopulmonary aspergillosis. Invasive aspergillosis (IA) represents the severe form of the disease and targets mainly immunocompromised patients (Van de veerdonk et al., 2017; Garcia rubio et al., 2017). Treatment options are limited to only three antifungal classes such as polyenes (amphotericin B), azole drugs and echinocandins. Triazole compound which are the most commonly used azoles, Itraconazol along with point mutations in the azole resistant mutation in cyp51A (Mellado et al., 2004). Presence of tandem repeats in the promoter region of cyp51A along with point mutations in the azole binding site of the enzyme, causes reduced sensitivity of the fungi to azoles. The agricultural use of azole fungicides is thought to be a major route responsible for the emergence and spread of TR34/L98H and TR46/Y121F/T289V mutant isolates (Chowdhary et al., 2012a). Apart from alterations in cyp51A gene, non-cyp51A mutations causing drug resistance have also been reported in clinical isolates (Frazek et al. 2013; Hagiwara et al., 2018; Gsaller et al., 2016; Camps et al., 2012a,b). The development of azole resistance among clinical isolates imparts a big challenge for the treatment of Aspergillus-related disease and demands a strong need for the development of alternative antifungal compounds.

BIOLOGY OF A. FUMIGATUS

A. fumigatus has enormous amount of ability to adapt to diverse environmental conditions. This property makes it a common pathogenic organism in human host. It produces millions of spores that ensure its survival and dispersal. The development of infection in human hosts starts through the inhalation of conidia (asexual spores) and its deposition in bronchioles and alveoli of the lungs. The conidia of A. fumigatus appear to be bluish-green or grey-green in color and have a globose to sub-globose like structure. These asexual spores are produced on a specialized hyphal structure called as conidiophores. They are hydrophobic in nature and their dispersal by air currents occurs very easily. The size of conidia plays an important role in the development of infection. The average size of Aspergillus conidia is about 2-3 μm that is best suited for its deposition in the lungs, though large-sized conidia of A. flavus and A. niger is removed by the movement of mucociliary membrane in the lungs. In immunocompromised patients, the mucosal membrane of lung is not able to clear out inhaled conidia which can further lead to fungal growth and colonization (Dagenais and Keller, 2009). Apart from this, Aspergillus shows thermotolerance. It can grow well at 37°C and has the ability to withstand temperature upto 50°C. The cell wall of conidia is made up of hydrophobic protein (expressed by well characterized gene RodA). This hydrophobin layer masks the fungal virulence factors due to which conidia are not recognized by immune cells of the host. The conidia also contain negatively charged carbohydrates such as sialic acid which helps them to adhere to the pulmonary wall containing fibroconnectin and laminin proteins. Besides this, the presence of melanin on the surface of conidium appears to protect it from ultra violet light and reactive oxygen species (ROS). The ROS are used by alveolar macrophages to phagocytose the conidia. The absence of melanin in the cell wall of conidium makes it less virulent and more susceptible towards phagocytosis. The melanin component of a cell wall is synthesized from the 1,3-dihydroxynaphthalene (called as DHN melanin) (Boral et al., 2018). The inner cell wall of A. fumigatus is composed of β-1,3-D-glucan to which chitin and galactosaminoglycan (GAG) are covalently attached. During germination, the conidia shed out its hydrophobin and melanin layer and swells up to form hyphae in the lungs (Van de veerdonk et
al., 2017). During this growth period, *A. fumigatus* is exposed to various stress conditions like iron acquisition, nutrient limitation, glucose deprivation and alkaline stress in host body. *A. fumigatus* adapt to these stress conditions by activating the expression of ‘Giltoxin’. Giltoxin is a secondary metabolite produced by the fungus that shows immunosuppressive and cytotoxic activity in host body. The fungus also gets benefits from the nutrients released by Giltoxin-mediated killing of host cells (Kwon Chung and Suguil, 2013).

**RISK FACTORS**

The spectrum of diseases associated with *A. fumigatus* is very broad. It causes chronic pulmonary aspergillosis (CPA), acute invasive aspergillosis and allergic syndromes. The immunodeficiency of the host, which includes neutropenia, corticosteroid induced immunosuppression, hematological diseases, long term use of cytotoxic drugs like cyclophosphamide (prevents graft rejection in patients), is the main predisposing risk factor responsible for the development of invasive aspergillosis (IA). The patients undergoing hematopoietic stem cell transplantation (HSCT) and solid organ transplantations (SOTs) may acquire some kind of immune defects which also contributes to the development of Aspergillus-related disease (Van de veerder et al., 2017). Patients suffering from severe lung disease such as chronic obstructive pulmonary disease (COPD) are at major risk to develop CPA and chronic necrotizing pulmonary aspergillosis (CPNA) that can cause fungal growth in damaged tissues. Patients having long stay in intensive care units (ICU) and those infected with tuberculosis or Human immunodeficiency virus (HIV) are also at great risk to develop aspergillosis in lungs. The allergic bronchopulmonary aspergillosis (ABPA) mainly occurs in patients having severe asthma and they immediately develop allergic response against the conidia of *Aspergillus*. The development of IA in patients is also due to lack of proper awareness and poor availability of microbiological tests to diagnose the infections (Garcia rubio et al., 2017).

**TRIAZOLE RESISTANCE IN *A. FUMIGATUS***

The azole resistance has become a global health threat as spores of *A. fumigatus* can disperse very easily by air currents. It has been shown that 90% of the azole resistant *A. fumigatus* isolates contain TR34-L98H (Tandem repeat-TR) mutation in cyp51A. This resistant genotype is also isolated from the azole naïve patients, who were never given any azole drug treatment, but might have received sufficient pressure exerted by the azole drugs. However, patients suffering from acute or chronic aspergillosis, do not exhibit aspherical stages of spores in lungs. In these patients, infection generally proceeds through the help of invasive hyphae.

Though, it is possible to develop resistance mutation in hyphae but it would have a very little effect on the phenotype of fungus because hyphae are surrounded by large number of wild type nuclei.

2) The environmental route: The isolation of azole resistant isolates from the azole naïve patients shows that they can procure azole resistant fungal infection from the environment. These isolates generally contain a tandem repeat (TR) in the promoter region along with point mutation in the coding region of cyp51A. There are mainly two reasons which justify the substantial use of azole fungicides in agriculture, first these compounds are not very expensive and secondly, they show a broad spectrum activity against various fungi. Generally all azoles have fungicidal effect rather than fungicidal effect (Azevedo et al., 2015). However azole compounds are very stable in nature and can persist in the environment for several months. Compost plays an important role in the development of azole resistance. It acts as an ecological niche for the various spores of fungi along with azole residues in it. Therefore, compost present in garden and agricultural field can act as a route for the emergence of resistant isolates (Snelders et al., 2009).

In *A. fumigatus* different patterns of azole resistance are identified ranging from multi azole resistance to pan azole resistance and these are found to be more common than resistance to single triazole. *A. fumigatus* contains two copies of target genes cyp51 (=ERG11 in yeast) - cyp51A and cyp51B, but each gene encodes different structural protein. The cyp51A gene encodes lanosterol 14a-demethylase required for ergosterol synthesis while cyp51B is required for the alternative functions in a cell, concerned with growth. Furthermore, till now no mutation has been detected in cyp51B gene that could be credited for azole resistance (Snelders et al., 2015). The triazole class of antifungal compounds inhibits the lanosterol 14a-demethylase enzyme. The enzyme plays a key role in the biosynthesis of ergosterol. The nitrogen atom of five membered triazole-ring binds to iron atom of heme group present in the active site of enzyme and inhibits the demethylation of lanosterol at C-14 thereby leading to the accumulation of methylated sterols in a cell. The accumulation of methylated toxic sterols in a cell makes plasma membrane leaky and impairs the growth of fungi (Bossche et al., 1995). There are many factors which determine the emergence of resistant isolates such as population size of fungi, its mode of reproduction, concentration of antifungal used and pharmacokinetic/pharmacodynamic (PK/PD) properties of the drug used (Bull et al., 2019). A clear understanding on emergence of triazole resistance in this fungus can be gained by understanding the basic structure of CYP51A enzyme, its role in the cell and how mutations in the cyp51A gene obliterates the azole binding to the active site of the enzyme.

**Structure of CYP51A Protein**

It has been studied that wild type CYP51A protein contain two ligand entry channels called as channel 1 and 2. The cyp51A sequence of *Mycobacterium tuberculosis* shares about 28% sequence identity with the cyp51A sequence of *A. fumigatus* (Snelders et al., 2010). The structure of wild type CYP51A protein of *A. fumigatus* is based on the protein homology model and X-ray crystallography structure of *Mycobacterium tuberculosis* protein (Chowdhary et al., 2014; Gollapudy et al., 2004; Xiao et al., 2004). The ligand entry channels helps in the binding of lipophilic sterol molecules and allow the docking of azole compounds at the active site of the protein. The presence of single nucleotide polymorphisms (SNPs) in cyp51A gene can cause structural alterations in CYP51A protein (Abdolrasool et al., 2015). The resistance to azole class of antifungal compounds depends upon the structural properties of the protein and its interaction with the compounds. The amino acids in codons G138, G54, M220 are located at the opening of the ligand entry channels. The G138 is present at the helix of channel 1, while G54 and M220 are located at the opening of channel 2 (Figure 1). All these residues make direct interactions with the azole that are required for binding at the heme group of the protein. The channel 1 is located parallel to the plane of heme group while channel 2 is perpendicular to the plane of heme group (Snelders et al., 2010).

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1) The patient route/ azole therapy: Treatment of patients for a longer period of time with azole drugs results in the selection of resistant mutant fungal isolates (Arendrup et al., 2010). Asexual sporulation in *A. fumigatus* facilitates the transfer of resistance mutation through the dispersal of spores (Chowdhary and Meis, 2018; Resendiz sharpe et al., 2018). These spores are generally formed in the lungs of patients suffering from caviary aspergillosis. The formation of asexual spores in lungs would allow the escape of resistant uninculturated spores from the heterokaryotic mycelium that germinate and form the entire colony that expresses the resistance mutations (Zhang et al., 2015; Verweij et al., 2009b). The formation of multiple spores also enables the fungi to adapt to the selective pressure exerted by the azole drugs. However, patients suffering from acute or chronic aspergillosis, do not exhibit asexual stages of spores in lungs. In these patients, infection generally proceeds through the help of invasive hyphae.

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Figure 1 CYP51A protein model of A. fumigatus. This model shows the important amino acids associated with resistance to azole compounds. The two ligand channels are indicated by arrows. The heme cofactor is represented black in color (Eveline Snelders et al., Antimicrob. Agents Chemother. 2010; doi:10.1128/AAC.01599-09) (Permission for Re-Use by ASM, License Number-497882805448)

The amino acid leucine at codon 98 in A. fumigatus is highly conserved and very important for maintaining the structural integrity of the CYP51A protein. It also plays a pivotal role in the docking of azole. The L98 residue forms an arch or gate like structure. This residue is neither located at substrate binding cleft nor at the heme group but present in a conserved loop like structure that connects two helices (B and B') in the protein structure. In wild type CYP51A protein the hydrophobic leucine residue forms an interaction with the proline 124 located on adjacent loop (BC) in the tertiary structure of the protein (Snelders et al., 2011).

Point Mutation in cyp51A Gene

Non-synonymous mutation at G54 and M220 leads to the substitution of smaller hydrophobic residues with the larger ones that will cause the blocking of channel 2 and inhibit the docking of azole compound. However there are two ligand entry channels present, but mutation in any one of the channel can inhibit the binding of azole compounds (Chowdhary et al., 2014). Majority of the mutations, mostly amino acid substitutions, are associated with channel 2 in CYP51A protein. These will cause a structural distortion in the CYP51A protein that will further inhibit the docking of azole molecules. Besides, the position and type of amino acid substitution in CYP51A protein also determines the pattern of azole cross resistance (Howard et al., 2009). Isolates having mutation at G54 residue will show cross resistance to itraconazole (ITC) and posaconazole (POS). Mutation at G54 includes 4 types of amino acid substitution and these are characterized as G54E, G54V, G54R, and G54W. The mutation G54W, for example, results in the ITC and POS resistant phenotype but it shows some susceptibility toward the voriconazole (VRC). In this type of mutation, glycine is replaced with the larger and hydrophobic tryptophan amino acid, so it will block the channel 2 and disturb the docking of azole molecule (Snelders et al., 2010). Both ITC and POS contain a long side chain and have ring like structure at the tail. Presence of two rings in the long side chains of ITC and POS will help them to bind G54 and M220 at ligand entry channel for the stable docking at heme cofactor. However, VRC is a compact molecule and lacks a long side chain; therefore it is not able to make any kind of interaction with these residues (Fraczek et al., 2011). The absence of cross resistance between the ITC and VRC is mainly due to the presence of differences in their chemical structures (Meneau and Sanglard, 2005). Resistance to ITC drug is generally associated with reduced susceptibility to POS as well, because both these drugs have similar structures. Therefore, isolates resistant to ITC also possess elevated Minimum Inhibitory concentration (MIC) to POS (Howard et al. 2009; Xiao et al. 2004). Therefore, mutation at G54 and M220 will affect the docking of only long chain azole compound such as ITC and POS (Fraczek et al., 2011).

The second type of point mutation in CYP51A occurs at M220 residue with amino acid substitution include M220V, M220K, M220T, M220I that shows resistance to ITC with decreased susceptibility to VRC and POS (Table 1); while isolates carrying amino acid substitution G138C show pan azole resistant phenotype (Mellado et al., 2004). The G448 residue is present at the opposite side of the heme group and it has the ability to alter the position of heme group. The G448S mutation shows resistance to voriconazole (VRC) with a reduced susceptibility to itraconazole and posaconazole (Snelders et al., 2010). This mutation acts by reducing the ability of binding of azole compound to heme group (Fraczek et al., 2011).

Table 1 Azole Resistance due to mutation in cyp51A

<table>
<thead>
<tr>
<th>Point Mutation in cyp51A at position</th>
<th>Amino Acid Substitution</th>
<th>Phenotype of the mutant</th>
</tr>
</thead>
<tbody>
<tr>
<td>54</td>
<td>G54E, G54V, G54R, G54W</td>
<td>Resistant to ITC</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Resistant to POS</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Sensitive to VRC</td>
</tr>
<tr>
<td>220</td>
<td>M220V, M220T, M220I</td>
<td>Resistant to ITC</td>
</tr>
<tr>
<td></td>
<td></td>
<td>High MIC/Resistant to POS</td>
</tr>
<tr>
<td></td>
<td></td>
<td>High MIC/Variable resistance to VRC</td>
</tr>
<tr>
<td>448</td>
<td>G448S</td>
<td>High MIC/Resistant to ITC</td>
</tr>
<tr>
<td></td>
<td></td>
<td>High MIC/Variable resistance to POS</td>
</tr>
<tr>
<td></td>
<td></td>
<td>Resistant to VRC</td>
</tr>
<tr>
<td>98</td>
<td>L98H</td>
<td>Multiple Azole resistance</td>
</tr>
</tbody>
</table>

(ITC-Itraconazole, POS-Posaconazole, VRC-Voriconazole) (Garcia rubio et al., 2017; Fraczek et al., 2011)

The amino acid substitution L98H also obstructs the binding of azoles (Snelders et al., 2011). The amino acid substitution from leucine to histidine results in the change of hydrophobicity at the surface of protein, as leucine is hydrophobic in nature while histidine is a polar residue. L98 is the only hydrophobic residue present on the hydrophilic surface of CYP51A protein. This results in narrowing of diameter of channel 1 and inhibits the entry of the ligand. Additionally, mutation L98 residue will re-localize the tyrosine residues at position 107 and 121 which further leads to blocking of channel 2 and result in the multiple azole resistant (MAR) phenotypes (Snelders et al., 2010, 2011). The tyrosine residue at 107 and 121 makes hydrogen bond with the two carboxyl group of the heme. The negatively charged carboxyl group will attract the histidine towards itself to neutralize negative charge, thereby blocking the entry of ligand though channel 2.

Tandem Repeats in cyp51A Gene

One of the most common resistance mechanism associated with azole compounds in both clinical and environmental isolates includes a 34 base pair tandem repeat (TR34) in the promoter region of cyp51A along with point mutation (L98H) in coding region. Changes in the promoter region of cyp51A will result in overexpression of gene (Chowdhary et al., 2017). The site directed mutagenesis studies show that both 34 base pair tandem repeats and L98H mutation are essential for the emergence of ARAF (Snelders et al., 2011).

Another mechanism of resistance mutation includes a 46 base pair tandem repeat in promoter region of cyp51A gene along with amino acid substitution of tyrosine for phenylalanine and threonine for alanine at positions 121 and 289 respectively (TR46/Y121F/T289A). The isolates carrying TR34/L98H mutation will show ITC resistant phenotype while isolates having TR46/Y121F/T289A phenotype show resistance to VRC (Lestrade et al., 2019).

Presence of tandem repeats at the 5’ end of cyp51A gene is an important mechanism for the development of azole resistance. The presence of different types of TR variants (TR34, TR46, TR53, and TR46) in isolates result in the emergence of pan azole resistant phenotypes. Over the past few years, new TR variants are reported that exhibit variation in the number (duplication in TR34 and TR46) and length of duplicated region such as TR120 (Bull et al., 2019; Hare et al., 2019).

The presence of a specific mutation in cyp51A gene also corresponds to development of specific azole resistance in fungal isolates. For example, the single nucleotide changes in cyp51A gene are thought to occur in patients that are having long term azole therapy (Lestrade et al., 2019). The tandem repeats in promoter region are mainly found in plant pathogenic molds and they generally acquire resistance mutation during the exposure ofazole fungicides in environment. These azole resistant phenotypes contain mutation at the transcriptional promoter or enhancer region (Verweij et al., 2016a,b). The incorporation of tandem repeats or duplication of the region mainly occurs during the sexual reproduction of the mold (Verweij et al., 2009b).

Analogous Binding of Medical Triazoles and Agricultural Fungicides to CYP51A

The use of sterol demethylase inhibitors (DMIs) as fungicides was authorized between 1990 and 1996 whereas the emergence of first TR34/L98H fungal isolate occurred in 1997 (Snelders et al., 2012). Both clinical and environmental isolates carry this mutations and show cross resistance to medical and agricultural azoles. The agricultural use of azole fungicides is thought to be a major route responsible for the emergence and spread of TR34/L98H and TR46/Y121F/T289A mutants.

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The hydrophobic residues (Y121, F115, V120, L128) present at the active site of CYP51A protein makes vanderwaals interactions with the di-halogenated phenyl group of triazoles, while the nitrogen atom of the five membered ring of triazole makes interactions with the heme group of CYP51A protein. In addition to this, the D-proponate ring (C₆H₄COO⁻) of the heme moiety forms hydrogen bond with hydroxyl group of triazoles (Chowdhary et al., 2013; Snelders et al., 2011) (Figure 2). The DMI fungicides also bind to the hydrophobic pocket created by the same residues in the target enzyme of A. fumigatus. The Y121 residue is very essential for the maintenance of the integrity and stability of heme group at the active site of a CYP51A protein. This residue forms a hydrophobic pocket and is used by all types ofazole compounds (ITC, POS, VRC and DMI fungicides) for the interaction. Amino acid substitution Y121F therefore disrupts the formation of hydrogen bond between the heme group and hydroxyl group of tyrosine (Snelders et al., 2015).

Non cyp51A Mutations
Mutations that do not involve cyp51A are also reported in clinical isolates conferring azole resistance (Sharma et al., 2019). These mutations are classified into different types-a) overexpression of efflux pumps, b) upregulation of sterol regulatory element binding protein (SrbA), c) mutation in HapE and (d) mutation in hmg1 gene

a. Overexpression of efflux pumps
One of the most important mechanism of resistance includes the over expression of efflux transporters that reduces the intracellular concentration of azole in the cell. Molecular characterization of the isolates shows that the A. fumigatus contain wild type cyp51A gene but still show azole resistance, which is due to the overexpression of cdr1B gene encoding the ABC transporter (Paul et al., 2013). Overexpression of ATP binding cassette (ABC) transporters and major facilitator superfamily (MFS) transporters has been found in azole resistant isolates. The ABC transporter contains two membrane spanning domain and two nucleotide binding domain that catalyze the ATP hydrolysis for the drug transport. On the other hand, MFS transporters are transmembrane proteins and they utilize the electrochemical proton motive force for the drug efflux (Sanglard, 2016). Cdr1B is the only known ABC transporter that mediates azole resistance and AftMDR3 is a MFS transporter which is found to be overexpressed in itraconazole (ITC) resistant isolates (Slaven et al., 2002; Frazeck et al., 2013). A potential transcriptional regulator AtuR regulates the expression of gene encoding Cdr1B transporter. Mutation in this transcription factor will result in overexpression of the gene encoding Cdr1B transporter.

b. Upregulation of Sterol regulatory element binding protein (SrbA)
Sterols are defined to be the major component of the fungal cell membrane. They are required for the maintenance of fluidity and permeability of the membrane. The mammals contain cholesterol as a principal sterol in membrane while fungi contain ergosterol in its membrane. The sterol regulatory element binding proteins (SREBP)s acts as a regulator for the biosynthesis of ergosterol. It generally binds to the sterol regulatory element (SRE) present at the promoter region of the gene and controls the transcription of genes involved in biosynthesis of sterols. The A. fumigatus contain SREBP homolog called as SrbA. It is defined as a transcription factor that contains basic helix loop helix (bHLH) as a DNA binding domain (Gsaller et al., 2016; Dhingra and Cramer, 2017). The SrbA binds to SRE motif in the promoter region and directs the transcription of several genes such as cyp51AB that are involved in the synthesis of ergosterol in A. fumigatus. The presence of tandem repeats in TR34 isolates results in the duplication of SRE motifs in promoter of cyp51A gene (Meis et al., 2016; Gsaller et al., 2016). Thus, binding of SrbA to SRE in TR region leads to increased expression of cyp51A gene. Therefore, SrbA plays crucial role in maintaining sterol profile of cell membranes of the fungi. The srbA null mutant shows hyper susceptibility toward the azole compounds (Wilger et al., 2008).

c. Mutation in HapE
The repeat sequence in TR34 is bound by both SrbA and CCAAT binding complex (CBC). It has been identified that CCAAT motif present in the 5’transcribed region of cyp51A gene of A. fumigatus plays an important role in the expression of cyp51A gene. CBC is a multi-subunit transcription factor and contains three subunits (HapB/HapC/HapE). The CBC recognizes a CGAAT and CCAAT motif promoter regions of cyp51A gene. It binds to the HapE, an ATP-dependent helicase, that acts as a negative regulator of ergosterol biosynthesis. Inactivation of CBC through mutation results in the increased expression of several enzymes of the ergosterol biosynthetic pathways through transcriptional activation and confers resistance to azole compounds. Mutation in HapE subunit at P88L impairs the binding affinity of CBC at CCAAT (Cumps et al., 2012a,b). Therefore, mutation in HapE subunit will result in the overexpression of sterol encoding gene by increasing the binding of SrbA (Gsaller et al., 2016).

d. Mutations in hmg1 gene Confers Resistance to Triazole in A. fumigatus
The biosynthesis of ergosterol starts with the acetyl-CoA and requires large amount of ATP for its formation. The enzyme HMG-CoA reductase catalyzes conversion of mevalonate from HMG-CoA. It is defined as a rate determining step in the biosynthesis of sterol in eukaryotes (Hagiwara et al., 2018). The sterol sensing domain of HMG-CoA reductase is very essential for the regulation of sterol biosynthesis. The isolate having mutation S269F, S269Y in the hmg1 gene (that encodes HMG-CoA reductase) exhibits reduced susceptibility towards the azole antifungal. The mutation S269F and S269Y alter the activity of HMG-CoA reductase enzyme by diminishing its sterol sensing ability (Hagiwara et al., 2018; Rybak et al., 2019). Presence of large amount of ergosterol in a cell will require more azole drug to inhibit the growth of fungus. Thus, the content of ergosterol in the cell membrane is a primary determinant involved in development of azole resistance.

ASSSESSMENT OF ANTIFUNGAL DRUG RESISTANCE
The Clinical Laboratory Standards Institute (CLSI) and the European Committee on Antimicrobial Susceptibility Testing (EUCAST) have developed standard protocols to determine antifungal susceptibility in yeasts and molds. The methods developed by them are used to determine MIC values (expressed in microgram per milliliter). CLSI and EUCAST have defined the epidemiological cut off values (ECV or ECOFF) and clinical breakpoints (based on MIC distribution, PK/PD of azoles, dose of drug, resistance mechanisms and outcome of treatment) for the study of triazole resistance in A. fumigatus (Sanglard, 2016; Verweij et al., 2009a). ECOFF values helps to identify non-wild type isolates that exhibit resistance to antifungal. They have also proposed the methods to identify in vitro resistance against the triazole in fungi. The resistant isolates generally show decreased MIC as compared to wild type isolates. However, MIC testing of various azole mold are not available, therefore these isolates are generally sent to the clinical mycology laboratories which can cause further delay in treatment. Therefore, therapeutic failure often occurs because patient is not able to respond to drug at a standard dose.

LABORATORY METHODS TO DETECT ASPERGILLUS RELATED INFECTION
In order to eliminate infections caused by fungi, it is very important to diagnose the infection at very early stage. Failure to detect azole resistance mutation can increase the mortality rate among the azole treated patients. To overcome these problems, scientists have developed novel diagnostic methods that are mainly used for the identification of the species involved in infection. Most of the microbiological laboratories in developing countries depend upon the conventional methods to identify Aspergillus species. The conventional method includes a direct microscopic examination, histopathological analysis, morphological character identification and standard microbiological cultures. The microscopic examination of tissue having invasion by fungi may provide the diagnostic clue for IA. Gomori Methenamine Silver stain (GMS) and Periodic Acid-Schiff (PAS) can be used to identify IA in tissue sections (Ullmann et al., 2018). Most of these stains are expensive and can be used to study various specimens such as Bronchoalveolar Lavage (BAL), Cerebrospinal fluid (CSF), sputum and tissue sections. However, these methods do not provide the accurate identification of the causative organism at species level (Arvanitis et al., 2014). The culture based methods are still used to detect triazole resistance but these methods are time consuming and have very less sensitivity. It has been studied that most of the patients suffering from aspergillosis or any other Aspergillus-related disease are found to be culture
negative. Therefore, analysis of the resistance mutations in culture negative patients is very difficult. Thus, early diagnosis of IA and initiation of antifungal therapy may improve the chances of survival in patients. But it is very difficult to recognize IA at very early stages.

There are different types of serological tests that are available such as immunodiffusion, ELISA and western blotting. ELISA assay targets the Galactomannan (GM) antigen. GM is a polysaccharide made up of mannann backbone and contains galactofuranosyl as side chain. This antigen is released by the Aspergillus during its early stage of infection. The level of fungal markers increases with the severity of the fungal infection. The GM that is released can be measured by an optical density index (ODI). However, this assay generally fails to determine the causative organism involved in infection and its susceptibility towards antifungal (Ullmann et al., 2018).

Many assays are available for the antifungal susceptibility testing such as disk diffusion, Etest, MIC strip kits, Sensititre YeastOne colorimetric assay and MTT assays. It is always suggested to perform an antifungal susceptibility test whenever the patient is tested positive for the A. fumigatus culture. The problem with this approach is that MIC determination is not routinely done in microbiological laboratories. Recently, the matrix assisted laser desorption ionization time of flight mass spectrometry (MALDI-TOF-MS) is being used for the accurate identification of the isolates. MALDI-TOF-MS is based on mass spectrometry and identifies the protein fingerprints of the microorganisms (Cassagne et al., 2016). It is possible to identify organism at species, genus and even strain level by comparing the spectral data of causative organisms with the reference spectra provided at the database in computer software (Arvanitakis et al., 2014).

**MOLECULAR METHODS TO DIAGNOSE AZOLE RESISTANCE IN A. FUMIGATUS**

Invasive fungal infections (IFIs) constitute a serious threat to immunocompromised patients. Species identification is a first step to detect the resistance mutation as it will help in the study of intrinsically resistant species (Lestrade et al., 2019; Ullmann et al., 2018). High mortality rate associated with the azole resistant isolates underlies the need of non-culture based methods to detect the species involved in infection and triazole resistance in clinical samples (Jenkins et al., 2019; Bull et al., 2018). The molecular method detects the resistance mutation in CYP51A protein and ensures the amplification of only fungal DNA and avoid any kind of cross reactivity with human DNA. A multiplex real-time PCR assay is available that detects one (TR34/L98H; Mycogenie developed by Ademtech, Pessac, France) or two (TR34/L98H and TR46/Y121F/T298A; AsperGenius, developed with PathoNostics, Maastricht, Netherlands) resistance mutation that is associated with the environmental route of transmission of azole resistance (Rath and Steinmann, 2018). Both these assays have advantages and disadvantages. The major benefit of AsperGenius assay includes its ability to detect the mixed infections and resistance mutation in culture negative patients. The AsperGenius assay detects mutation by the help of melting curve analysis (Meis et al., 2016; White et al., 2015). This assay shows its diagnostic effect in BAL fluid and serum samples while MycoGenie PCR system is generally recommended for the testing of biopsy, respiratory and serum samples. MycoGenie PCR system shows a very high sensitivity to detect the 28S rDNA gene of A. fumigatus (Jenkins et al., 2019; Rath and Steinmann, 2018). The important drawback includes the low sensitivity of the assays because cyp51A gene is a single copy gene present in each Aspergillus cell so that detection of resistance markers in serum might not be successful (Verweij et al., 2015). The problem associated with these commercial test kits is that they are only able to identify TR associated mutations. Therefore a negative test by these methods does not mean that the patient is not suffering from azole resistant aspergillus (Lestrade et al., 2019; Bull et al., 2019; Meis et al., 2016).

Microsatellite analysis is a best approach to study the genetic diversity between the fungal isolates. The next generation sequencing techniques such as illumina sequencing, amplicon sequencing of ITS regions or ribosomal gene, helps in determining the severity of fungal infections. The amplicon sequencing of ITS regions and whole genome sequencing of fungal isolates helps in identifying the species involved in infection (Zolfi et al., 2016).

Presence of point mutations in the cyp51A gene is a critical feature in the resistance of azole resistant isolates. Molecular approaches that targets the mutation associated with the coding region of cyp51A gene (G54, M220, G138, G448) are still lagging behind. Pyrosequencing is an important DNA sequencing technique based on ‘sequencing by synthesis’ principle that can rapidly screen out point mutations associated with theazole resistance (Van der torre et al., 2016; Meis et al., 2016).

Nucleic acid amplification provides a rapid, sensitive and simple technique that can be used in clinical laboratories to identify early stage infections. Most of the PCR based techniques relies on the use of specific primer targeting the internal transcribed sequence (ITS) of the rDNA which helps in identifying the species. Oligonucleotides are used to study Aspergillus species by amplifying the benA and calmodulin (caM) gene that can act as a secondary identification markers for the identification of causative organism involved in infection. Taxonomic tool such as ribosome gene and inter transcribed spacer regions are generally used for inter section level identification in A. fumigatus while the β-tubulin locus is used for the identification of individual species belonging to various Aspergillus sections (Chowdhary et al., 2016; Zakaria et al., 2020).

**EPIDEMIOLOGY OF CLINICALLY ISOLATED ASPERGILLUS FUNGI**

As per the epidemiological data, A. fumigatus is the predominant etiological agent of the invasive pulmonary aspergillosis in immunocompromised patients. Prospective cohort studies of transplant-associated fungal infections in the United States carried out by Transplant-Associated Infection Surveillance Network (TRANSNET) have also reported that A. fumigatus is the leading cause of invasive aspergillosis followed by A. flavus, A. niger, A. terreus (Kontoyiannis et al., 2010). By analyzing this data isolated from the posttransplant patients, Amphotericin B (AmpB) in the polyene and shows very poor oral absorption therefore it is solubilized with deoxycholate for the intravenous administration. AmB-deoxycholate is a potent antifungal drug with a broad spectrum activity, but it shows nephrotoxicity and infusion reactions (Stone et al., 2016). These disadvantages promoted the development of less toxic lipid based formulations of AmB such as amphotericin B lipid complex (ABLC). The new formulation includes AmB nanoparticle suspensions and AmB arabinogalactan or AmB-PEG (AmB-Polyethylene glycol) that shows high efficacy with little toxicity. The primary mechanism of AmB action includes its ability to form ion channels in the plasma membrane leading to leakage of ions from the cell. The second mechanism of action involves the oxidative damage of a cell by producing ROS. Lipid based formulation are generally recommended to avoid any kind of AmB related nephrotoxicity. The available lipid formulations are ABLC (Abeleum), AmB colloidal dispersion (ABCD, amphotec, Amphotec) and liposomal AmB (AmBisome) from the Pfizer. The difference in pathogenicity from those of A. fumigatus. The growing incidence of azole resistant species demands the accurate identification and antifungal susceptibility of the isolates (Pappas et al., 2010; Hagiwara et al., 2016).

**TREATMENT OPTIONS**

It has been successfully demonstrated that azole resistance is the predominant cause of treatment failure in most cases. The elevated MIC shown by azole resistant isolates reduces the efficiency of azole monotherapy in treatment (Lepak et al., 2013). CD101 (blastfungin) is a novel echinocandin that has been found to be effective in mouse model suffering from IA. ITC is mainly used in the treatment of CPA while VRC is used for IA. Amphotericin B (AmpB) is a polyene and shows very poor oral absorption therefore it is solubilized with deoxycholate for the intravenous administration. AmB-deoxycholate is a potent antifungal drug with a broad spectrum activity, but it shows nephrotoxicity and infusion reactions (Stone et al., 2016). These disadvantages promoted the development of less toxic lipid based formulations of AmB such as amphotericin B lipid complex (ABLC). The new formulation includes AmB nanoparticle suspensions and AmB arabinogalactan or AmB-PEG (AmB-Polyethylene glycol) that shows high efficacy with little toxicity. The primary mechanism of AmB action includes its ability to form ion channels in the plasma membrane leading to leakage of ions from the cell. The second mechanism of action involves the oxidative damage of a cell by producing ROS. Lipid based formulation are generally recommended to avoid any kind of AmB related nephrotoxicity. The available lipid formulations are ABLC (Abeleum), AmB colloidal dispersion (ABCD, amphotec, Amphotec) and liposomal AmB (AmBisome) from the Pfizer. The difference in pathogenicity from those of A. fumigatus. The growing incidence of azole resistant species demands the accurate identification and antifungal susceptibility of the isolates (Pappas et al., 2010; Hagiwara et al., 2016).
contain the binding site for ATP followed by a middle domain and C-terminal domain that contain binding site for the various downstream effector proteins and co-chaperons (Lamoth et al., 2014). The Hsp90 inhibitor geldanamycin and its derivatives 17-allylamino-17-demethoxygeldanamycin (17-AAG) and 17-demethoxygeldanamycin (17-DAMG) shows in vitro antifungal activity against the A. fumigatus. Geldanamycin restricts the paradoxical effect of caspofungin and potentiates its antifungal activity. The echinocandin (caspofungin, anidulafungin and micafungin) class of antifungal inhibits the synthesis of β-1,3-D-Glucan. The paradoxical effect refers to the dye shifting of hsp90 at higher concentrations. This effect occurs in A. fumigatus due to the activation of calcineurin signaling pathways which increases the chitin in cell wall. Hsp90 is one of the client proteins that activate this pathway. Therefore, use of Hsp90 inhibitors such as geldanamycin and its derivatives can attenuates the paradoxical effect. Targeting of Hsp90 and calcineurin signaling pathways can result in increased susceptibility towards the antifungal.

The anti-calcineurin drug FK506 (tacrolimus and cyclosporin), Hsp90 inhibitor (Geldanamycin and its derivatives 17-AAG, 17-DAMG) and histone deacetylase (HDAC) inhibitor such as Trichostatin A exhibit in vitro antifungal activity against various mold species. Trichostatin A inhibitor further shows synergistic interactions with caspofungin (Lamoth et al., 2015a,b).

CONCLUSION

The emergence of azole resistance in A. fumigatus is a major global health issue. The article highlights importance of fungal cyp51A gene and the associated mutation in the gene that forms part of a primary pathway of emergent azole resistant A. fumigatus mutants. The fact that these strains show cross-resistance to the three important medical triazoles and also to agricultural triazole fungicides, pointed to the spread of resistant isolates from environment to the clinics. There is therefore growing need for the development of novel approach methods to identify resistance mutations in order to initiate appropriate antifungal therapy in patients. Several new antifungals need to be developed that will show more advantage as compared to current drugs in terms of overcoming drug resistance and could offer less toxic effects. Environmental survey should be conducted to determine the prevalence of resistant isolates in agricultural fields. Furthermore, the indiscriminate usage of fungicides should be avoided in agricultural fields to minimize the emergence of resistant isolates. The increasing number of affected patients emphasizes the need for an elaborate study of fungal epidemiology in different countries and a thorough assessment of azole resistant mutations and its associated treatment failure. The future research should focus mainly on effectiveness of new pharmaceutical agents that can be used in alone or synergistically to avoid the emergence of resistant isolates.

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