

Review

# Potential Strategies to Control the Risk of Antifungal Resistance in Humans: A Comprehensive Review

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**Abstract:** Fungal infections are becoming one of the main causes of morbidity and mortality in people with weakened immune systems. Mycoses are becoming more common, despite greater knowledge and better treatment methods, due to the regular emergence of resistance to the antifungal medications used in clinical settings. Antifungal therapy is the mainstay of patient management for acute and chronic mycoses. However, the limited availability of antifungal drug classes limits the range of available treatments. Additionally, several drawbacks to treating mycoses include unfavourable side effects, a limited activity spectrum, a paucity of targets, and fungal resistance, all of which continue to be significant issues in developing antifungal drugs. The emergence of antifungal drug resistance has eliminated accessible drug classes as treatment choices, which significantly compromises the clinical management of fungal illnesses. In some situations, the emergence of strains resistant to many antifungal medications is a major concern. Although new medications have been developed to address this issue, antifungal drug resistance has grown more pronounced, particularly in patients who need long-term care or are undergoing antifungal prophylaxis. Moreover, the mechanisms that cause resistance must be well understood, including modifications in drug target affinities and abundances, along with biofilms and efflux pumps that diminish intracellular drug

levels, to find novel antifungal drugs and drug targets. In this review, different classes of antifungal agents, and their resistance mechanisms, have been discussed. The latter part of the review focuses on the strategies by which we can overcome this serious issue of antifungal resistance in humans.

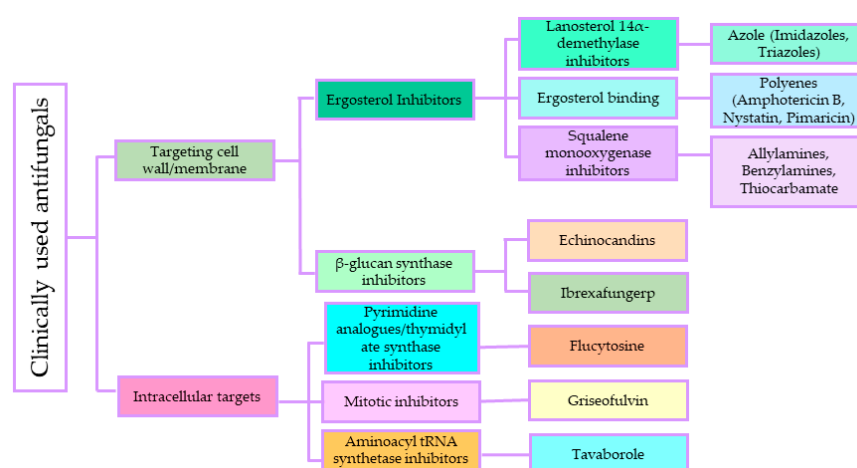
**Keywords:** antifungal; antifungal resistance; fungal infections; fungal treatments

## 1. Introduction

The human population has been plagued by various infectious diseases for a long time and they remain one of the leading causes of death [1]. Fungal infections are not considered a severe health issue, even though they affect over a billion people. Approximately 1 million individuals have invasive fungal infections (*Aspergillus*, *Candida*, *Cryptococcus*, and *Pneumocystis*) worldwide, resulting in the death of 1.7 million people annually [2,3]. The WHO has developed a priority list of fungi based on the severity of infection, the WHO FFPL. This is the first global effort to systematically classify fungal pathogens. Nineteen fungi were categorized into three priority groups (critical, high, and medium) in this list considering their unfulfilled requirements in research and development and their significant role in public health [4]. Almost every person is affected by a superficial fungal infection, which is easy to cure. However, individual infection by a specific invasive fungal pathogen can be fatal due to a lack of definitive diagnosis and treatment [5]. For instance, recent studies estimated ~ 300,000 cases of invasive *Aspergillosis*, 750,000 cases of *Candidiasis*, and more than 900,000 cases of *Mucormycosis* infection annually [6,7]. Additionally, the emergence of antifungal drug resistance in susceptible pathogens (the ubiquitous mould *Aspergillus fumigatus*) and novel fungal species resistance to multiple antifungal drugs (for example, the yeast *Candida auris*) are alarming critical concerns to human health worldwide [8–11]. The inaccessibility of efficient therapeutics against such fungal infections and the gain of resistance in fungi against antifungal agents is, therefore, contributing a hindrance to eradicating fungal infection while increasing the mortality rate [12]. In addition, an increasing number of diseased populations, such as older people and immunodeficient or seriously ill patients, such as AIDS, cancer, diabetic patients, and transplantation patients, are vulnerable to fungal disease, which provides a perfect niche for infection [13]. Additionally, fungal infections were connected with other bacterial and viral infections, including severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2); immunocompromised patients thereof are prone to invasive fungal infections [14]. For instance, the most unambiguous indication of co-infection was the appearance of mucormycosis, a fungal infection caused by Mucorales, in South Asian nations, particularly in India, following the first wave of the coronavirus diseases 19 (COVID-19) caused by SARS-CoV-2. [15,16]. However, it appears that severe immunocompromised hosts (such as those who have undergone organ transplantation, cancer, autoimmune disorders, immune system functional limitations, or neutropenia), hyperglycaemia or poorly controlled diabetes mellitus, and immunocompetence in post-traumatic cases are the leading causes of this infection [15].

Among the known fungal pathogens, drug-resistant pathogens in invasive fungal infections are responsible for causing more than 50% of the mortality rate; therefore, it is essential to understand the mechanism of drug resistance and the drawbacks of the currently available antifungal treatments for the clinical management of fungal infections [5]. Recent studies established that pathogenic fungi acquired various adaptive mechanisms to achieve antifungal drug resistance, such as alteration of the target drug site, target over-expression, upregulation of the multidrug transporters, biofilm formation, cell permeability, and stress response [17]. For instance, fungi gained primary resistance against the chemical action of antifungal drugs using intrinsic resistance parameters, including biofilm formation and cell wall impermeability. After continuous exposure to antifungal agents, fungi acquired secondary mechanisms to adopt resistance, termed acquired

resistance, via genetic alternations to modify the target binding site or overexpression of the targeted proteins, or stress response and stress response/epigenetic pathways alternation [18–21]. Additionally, fungi can also gain antifungal drug resistance via random mutations induced by the overuse of antifungal functional groups, such as azole and echinocandins [22]. Therefore, a systematic treatment for fungal infections is mainly based on four classes of antifungal drugs, i.e., polyenes, azoles, echinocandins, and antimetabolite agents (Figure 1) [19]. For example, heptaene amphotericin B is a class of polyene which binds to ergosterol, a significant component of the fungal cell membrane. It creates pores in the cell membrane to cause leakage of intracellular ions. Such drugs have been documented for the fungicidal or fungistatic activity against *Aspergillus fumigatus*, *A. flavus*, and *Candida* genera [23]. Likewise, azole drugs interfere with the ergosterol biosynthesis by inhibiting the ergosterol biosynthesis of the 14 $\alpha$ -lanosterol demethylase in fungi and are reported with potential activity against yeast and *Aspergillus* species [24]. Additionally, echinocandins (containing caspofungin, micafungin, and anidulafungin drugs) are known to inhibit the synthesis of  $\beta$ -d-glucans in fungi, which are essentially required for the formation of the fungal cell wall [25]. Similarly, flucytosine, an antimetabolite agent that inhibits fungal growth by alternating in fungal DNA and protein synthesis, is generally administered in conjunction with amphotericin B to treat refractory *Candida* infections and *Cryptococcal meningitis* infections [26]. Pathogenic fungi respond to these available antifungal agents effectively. Still, with prolonged usage of these drugs and due to some external factors, the pathogen attains resistance, resulting in no improvement in the infected individual even after treatment [27]. Other factors, such as a defective host immune system, poor antifungal activity, and specific fungal characteristics, such as antifungal tolerance and resistance, were also elucidated as critical factors in promoting antifungal drug resistance or the failure of antifungal treatment [28]. It is also important to mention that over-using antifungal drugs increases the chance of opportunistic pathogens attaining resistance [29]. So, it is essential to find specific and practical strategies to treat drug-resistant fungi by analysing the molecular mechanism leading to drug resistance, which can help identify new potential targets for developing new antifungal drugs [30]. Hence, the present review gives a brief understanding of the development of drug resistance in fungi against the commonly used antifungal agents and potential strategies adopted to overcome antifungal resistance, mainly in humans.



**Figure 1.** Flow chart depicting classification of antifungal drugs concerning the mode of action as well as names of drugs used in the management of fungal infections in humans.

## 2. Methodology

### 2.1. Literature Screening

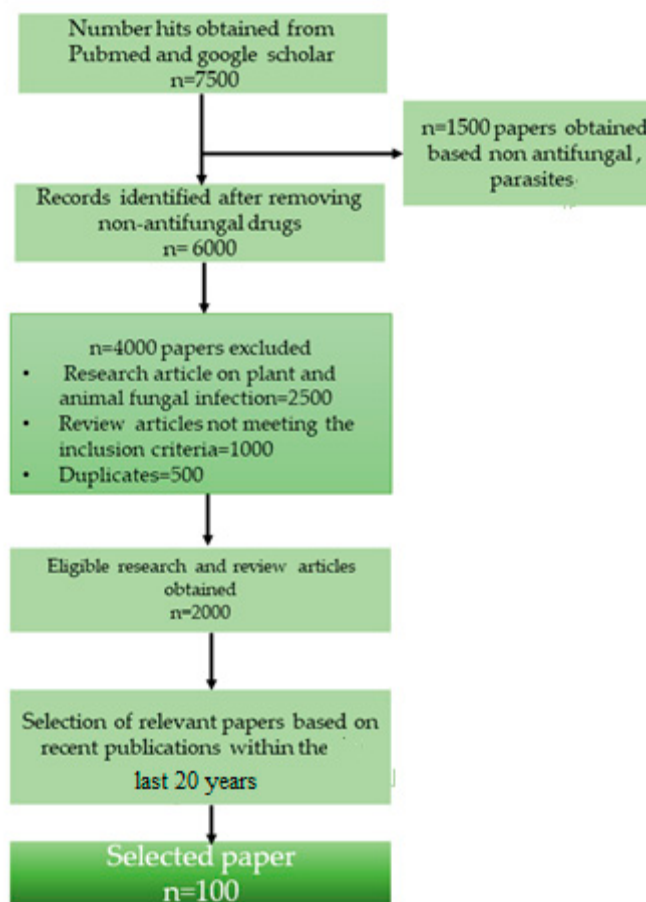
In this study, PubMed (<https://pubmed.ncbi.nlm.nih.gov/>, accessed on 20 June 2022) search engine was selected as they use the MEDLINE database to find relevant papers on life sciences and biomedicines. This search engine helps to screen the articles based on various criteria such as year of publication, type of article, text available, and other additional filters to make the screening of articles easier and accurate. Keywords ‘Antifungal’, ‘Antifungal resistance’, ‘Fungal infections’, ‘Fungal treatments’, and ‘Antifungal agents’ were used to search and collect the relevant literature from this database. Additionally, Clinical Trials (<https://clinicaltrials.gov/>, accessed on 20 June 2022) database was also considered as the secondary database to find the relevant information. Initially, the first search on the mentioned databases was performed on 26 June 2022, and relevant articles, both research and review papers, published between 2001 and 2022 were collected. Later, a second literature collection was conducted on 1 July 2022, to update (1 October 2022), and again, the most relevant publications were downloaded and considered in the present review. Following that, the collected literature was screened using the multiple keywords mentioned above to assemble the most pertinent publications.

### 2.2. Exclusion of Articles

Conclusively, the adopted methodology for the literature screening resulted in a collection of 7500 hits. This set of articles was further refined and articles containing information about the non-antifungal drugs (1500 articles), such as antiparasitic drugs and diseases, were excluded. Furthermore, from the available 6000 hits, another 4000 articles on antifungal agents related to plants and animals (2500 papers) were also removed from the set of selected papers. A set of articles containing incomplete information regarding antifungal resistance and published in languages other than English (1000 publications) were also excluded as they did not fulfil the selection criteria. Additionally, around 500 articles were found to be duplicates.

### 2.3. Collection of Data from Relevant Articles

From the remaining 2000 eligible research and review papers, only 100 relevant papers focusing on the inclusion criteria, such as evaluating antifungal resistance using in vitro and in vivo methods, various mechanisms of antifungal drug resistance shown by prominent fungal pathogens, and strategies used for the prevention of antifungal resistance, were sorted for the present systematic review. As the current paper concentrates on antifungal resistance in humans, 100 articles within the last 20 years were thus further sorted for a new collection that discussed antifungal resistance or antifungal medication in humans. To provide a comprehensive understanding of antifungal drug resistance, 60 articles were consulted from these 100 selected publications that describe the categorisation of antifungal medications, their mechanisms of action, the development of antifungal resistance, and strategies to overcome antifungal resistance in humans for a detailed discussion. In addition, some relevant articles were cited and discussed during the article revision. A systematic flow representing the adopted methodology for selecting and refining the literature is given in Figure 2.



**Figure 2.** Flow chart exhibiting selection criteria adopted for selecting research publications used in the design and compilation of the present review.

### 3. Results and Discussion

The overuse of existing antifungal medications in recent years has led to the evolution of antifungal-resistant fungi or the emergence of novel fungal species with solid antifungal tolerance [31]. As a result of its impact on human health, there is now a great deal of interest in understanding antifungal resistance from various viewpoints. The existing antifungal therapeutic agents are affected by other factors such as drug–drug interaction, toxicity, and limitation in routes of drug administration [32]. In the last two decades, no new antifungal classes have been available, and only a few new antifungal drugs of the existing class of drugs are being approved for treatment. Identification of novel antifungal pathways and drug targets is utilised for developing potential antifungal drugs with a novel mode of action or for a new formulation of existing antifungals [33,34]. A brief insight into the existing antifungal drugs and their resistance mechanisms will be helpful in the formulation of new antifungal agents.

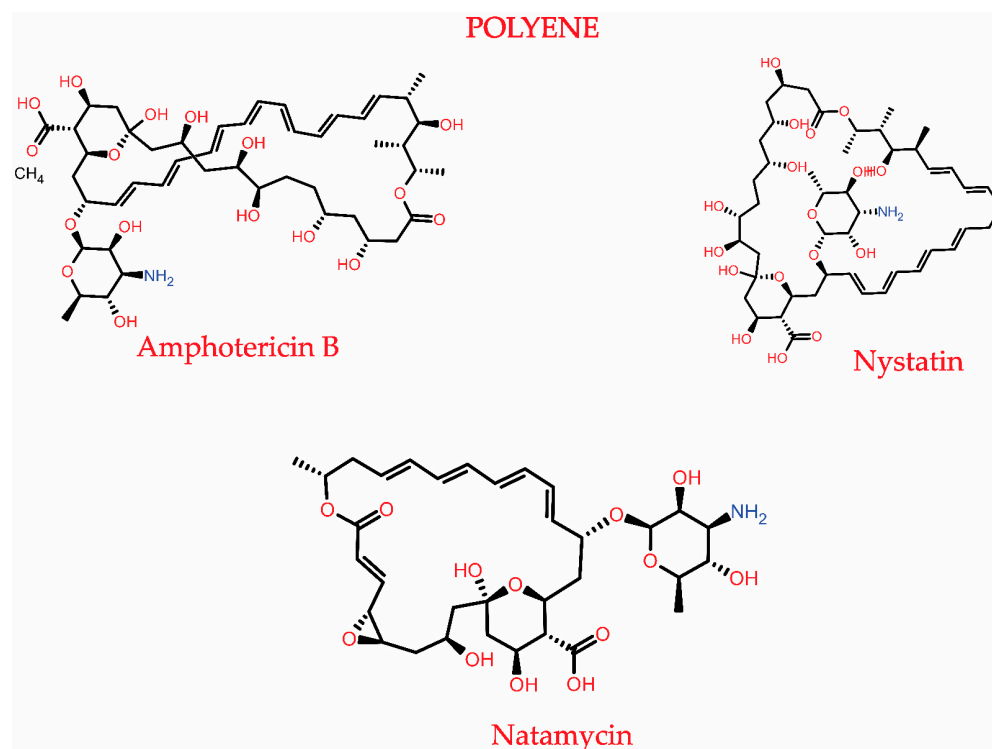
#### 3.1. Antifungal Agents and Their Mode of Action

Antifungal agents can be divided into nonspecific and specific agents based on the mode of action. Nonspecific antifungals mainly include disinfectants, antiseptics, and essential oils that can be applied to treat skin and mucous antifungal infections. Specific antifungals, also known as antimycotic drugs, have a particular mechanism of action [35]. For example, polyenes, antimetabolites agents, azoles, and echinocandins are the four classes of specific antifungal drugs approved by the Food and Drug Administration (FDA) for treating invasive fungal infections [36]. Only certain antifungals are taken into consideration in the current investigation because they are commonly used to treat invasive fungal infections.

### 3.1.1. Polyenes

The first antifungal polyene—fungicidin, later called nystatin—was discovered in 1949. More than 200 polyenes as fungal-specific antibiotics have been discovered and used to treat fungal infections in humans [36,37]. These antifungal agents show fungicidal activity against numerous species of *Aspergillus*, *Candida*, and *Cryptococcus* genera [17]. Generally, polyenes possess a cyclic heptaene or polyketide core macrolactone ring (20–40 carbon atoms including 3–8 conjugated double bonds) structure and are produced by the Gram-positive bacterium *Streptomyces nodosus*. Based on the conjugated double bonds, polyenes are characterised as trienes, tetraenes, pentaenes, hexaenes, heptaenes, etc. [38,39]. Approximately six polyenes were identified that could be used for antifungal therapy: amphotericin B, nystatin, natamycin (also called pimaricin), candicidin, trichomycin, and methyl partricin [37]. Amphotericin B (amphotericin B deoxycholate) is the most widely used and was first approved by the FDA to treat invasive fungal infections but, due to low therapeutic index and high nephrotoxicity, the usage of this drug has been limited [33,36]. To improve drug quality and reduce drug toxicity, liposomal amphotericin B was introduced, where the amphotericin B is induced into a small, unilamellar vesicle made up of a liposomal formulation. The liposomal amphotericin B is much safer and less toxic than the conventional amphotericin B and is used for the treatment of various fungal infections [40].

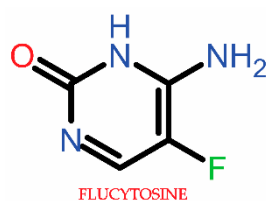
These antifungal agents exhibit two different modes of action against the fungi, i.e., (i) the polyenes incorporate into the fungal lipid bilayer and bind to the ergosterol molecule. This results in pore formation in the fungal cell wall and causes leakage of essential ions ( $K^+$ ,  $Mg^{2+}$ ,  $Ca^{2+}$ , and  $Cl^-$ ) and energy molecules (glucose), leading to the death of fungal cells; (ii) polyenes instigate reactive oxygen species (ROS) production and accumulation in fungi, which causes substantial damage to the fungal protein, mitochondria, cell membrane, and DNA [41]. Another mode of action of polyene is the extraction or ergosterol adsorption from the membrane leading to membrane destabilisation and membrane protein function disturbance [42] (Table 1, Figure 3).



**Figure 3.** Two-dimensional structural formula for the most common drugs under the polyene antifungal group. All the 2D structures were plotted using the academic version of Maestro Schrödinger suite 2022 (Schrödinger Release 2022-3: Maestro, Schrödinger, LLC, New York, NY, 2021).

### 3.1.2. Antimetabolite Agents (5-Fluorocytosine)

Antimetabolites are structurally characterized as analogues of essential metabolites but cannot be consumed by the human body [43]. The common classes of antimetabolites include purine antagonists (6-mercaptopurine) [44] and pyrimidine antagonists (5-fluorouracil) [45]. Some natural products, such as vinca alkaloids and taxol, have been reported as antimetabolites in disease management [46,47]. Some antimetabolites are also available for use against invasive fungal infections in humans. The best example is 5-fluorocytosine, a synthetic analogue of cytosine, which was first synthesized in 1957 as an antitumor drug and later approved by the FDA as an antifungal drug for humans in 1968 (Figure 4). Flucytosine effectively treats fungal infections caused by *Cryptococcus neoformans*, *Candida* spp., and *Cryptococcal meningitis* [48–50]. It is also used with amphotericin B to treat systemic mycoses, and this association also reduces the rate of nephrotoxicity in patients compared to treatment of amphotericin B alone [51,52]. Additionally, the combination of these drugs has a higher efficacy rate [53]. Typically, when the drug enters the fungus, the drug gets activated into 5-fluorouracil(5-FU) by the fungal cytosine deaminase enzyme and impedes DNA and RNA synthesis via intracytoplasmic conversion. For instance, 5-fluorouracil (5-FU) is converted into 5-fluorouridine monophosphate (5FUMP) with the help of uracil phosphoribosyltransferase (FUR1) and then 5-fluorouridine triphosphate to alter the RNA synthesis as well as protein synthesis [54–56]. Moreover, fluoro-deoxyuridylic acid, the modified form of 5-fluorouracil (5-FU), also inhibits DNA synthesis and causes DNA damage by the inhibition of thymidylate synthase [55] (Table 1).

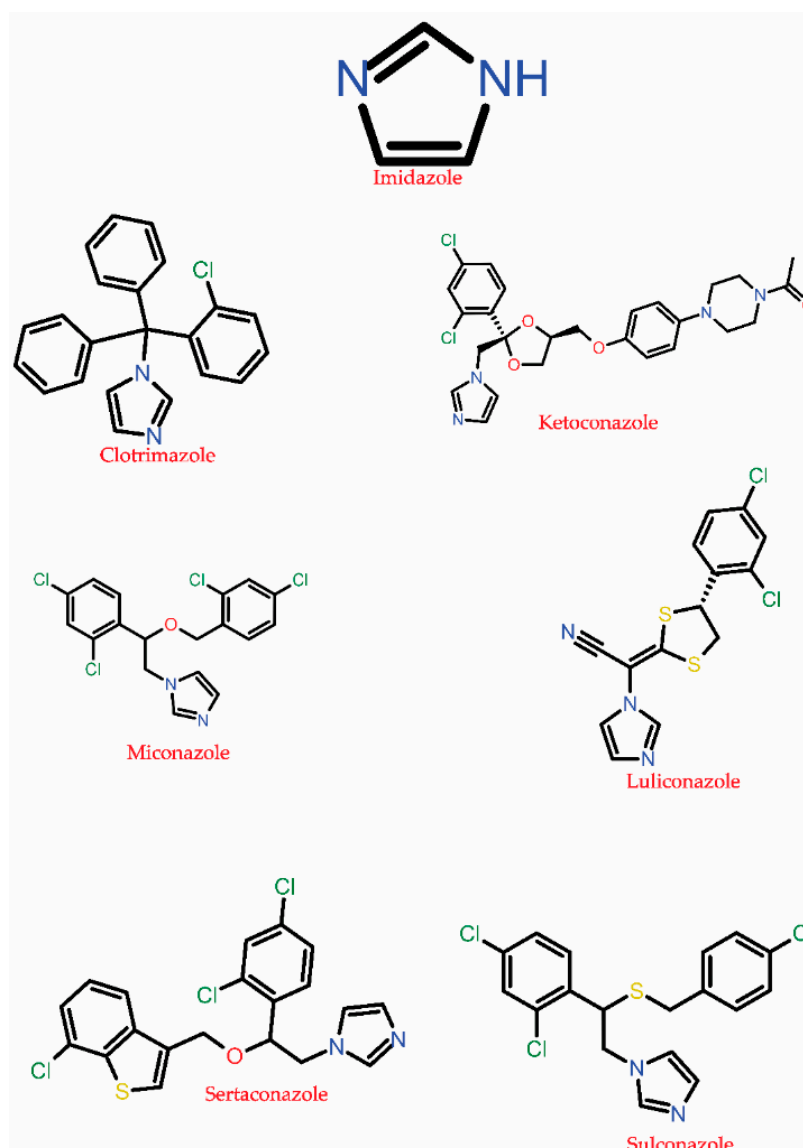


**Figure 4.** Two-dimensional structure of flucytosine used as an antimetabolite agent against invasive fungal infections.

### 3.1.3. Azole

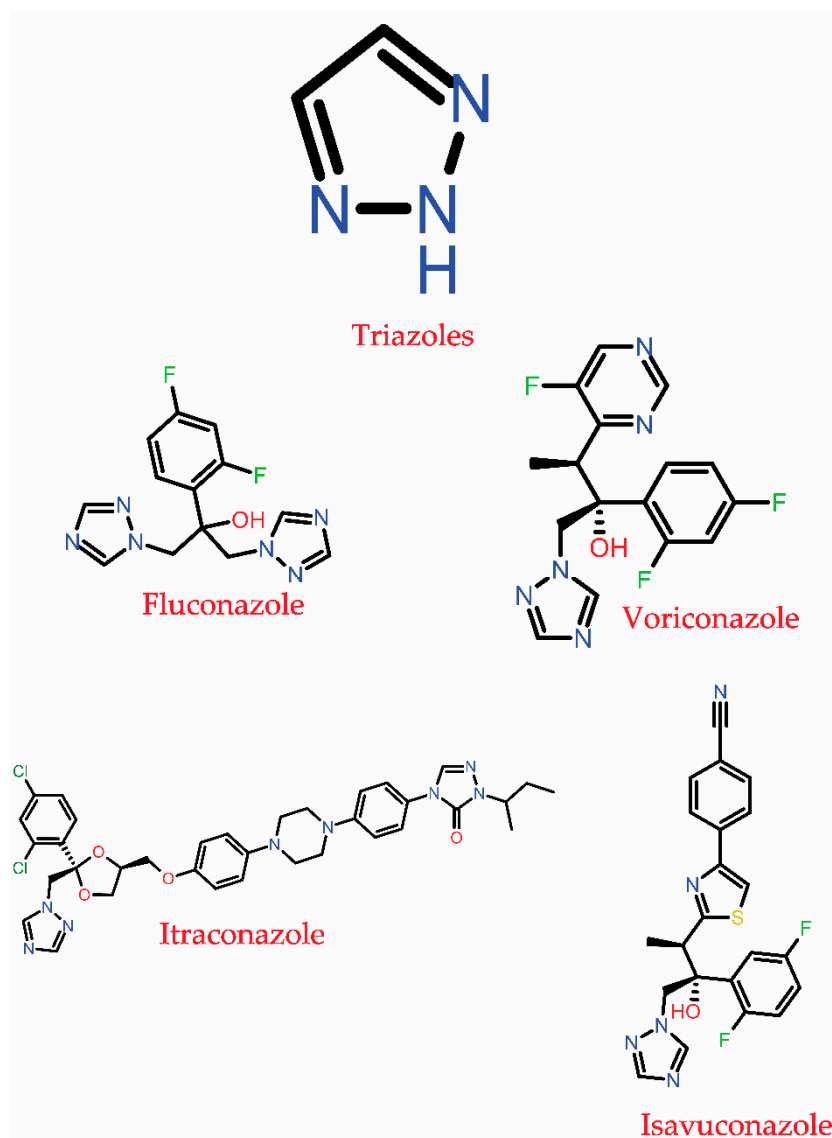
Azoles, a broad-spectrum class of antifungals, were discovered in 1944 and approved for human use in the 1950s [36]. Based on the number present in the aromatic ring of azoles, they are classified into three groups: imidazoles, triazoles, and tetrazoles. Imidazoles have two nitrogen atoms in the azole ring, and triazoles have three nitrogen atoms in the azole ring [57]. Miconazole, clotrimazole, econazole, ketoconazole, tioconazole, sulconazole, serconazole, and luliconazole represents the imidazole-based azole drugs and terconazole, fluconazole, isavuconazole (isavuconazonium—prodrug of isavuconazole), itraconazole, voriconazole, posaconazole, eficonazole, and albaconazole represents the triazole group of drugs [58]. Tetrazole, a recently developed compound under the azole group, shows broad spectrum activity against fungal species, but its usage is limited as most of the drugs are under trial stage and need to be approved by worldwide agencies [59,60]. Tetrazoles, which include one carbon and four nitrogen atoms, are a subclass of doubly unsaturated aromatic heterocycles with a five-membered ring. In nature, they do not exist [61]. Modified tetrazoles named quileseconazole (VT-1129) and oteseconazole (VT-1161) are the latest developed inhibitors of lanosterol 14 $\alpha$ -demethylase encoded by the *Cyp51* gene [54,62]. They are designed to overcome the problem of drug–drug interaction, a significant limitation of the azole class of antifungal drugs. These are developed by Mycovia Pharmaceuticals (previously known as Viamet Pharmaceuticals), Inc. (Durham, NC, USA) by replacing the triazole metal binding group with a tetrazole. This modification results in the development of more specific compounds in the inhibition of the *Cyp51* gene and shows minor drug–drug interaction [63,64]. The FDA approved

quileseconazole as an orphan drug for the treatment of cryptococcal meningitis, and oteseconazole is approved for the treatment of recurrent vulvovaginal candidiasis (RVVC) [54,65,66]. Quileseconazole and oteseconazole show antifungal activity against *Candida* and *Cryptococcus* species [67]. Drugs under the azole group inhibit the  $14\alpha$ -lanosterol demethylase enzymes, causing a disturbance in ergosterol biosynthesis, a significant cell membrane component. As a result, the fungal cells experience depletion of ergosterol and accumulation of toxic 14-methylated sterols, leading to cell lysis and death [68]. Azoles displays fungicidal and fungistatic activity against fungi from genera *Candida*, *Cryptococcus*, *Coccidioides*, *Aspergillus*, yeasts, and moulds. Voriconazole, isavuconazole, and itraconazole are the most preferred drugs for the treatment of invasive aspergillosis and fluconazole is one of the most safest and effective drugs used for the treatment of candida endophthalmitis [69–73] (Table 1) (Figures 5–7).

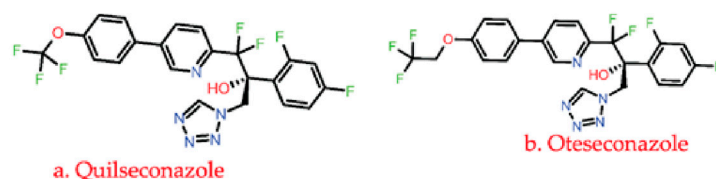


**Figure 5.** Two-dimensional structure of drugs under the imidazole group.





**Figure 6.** Two-dimensional structure of drugs under triazole group.

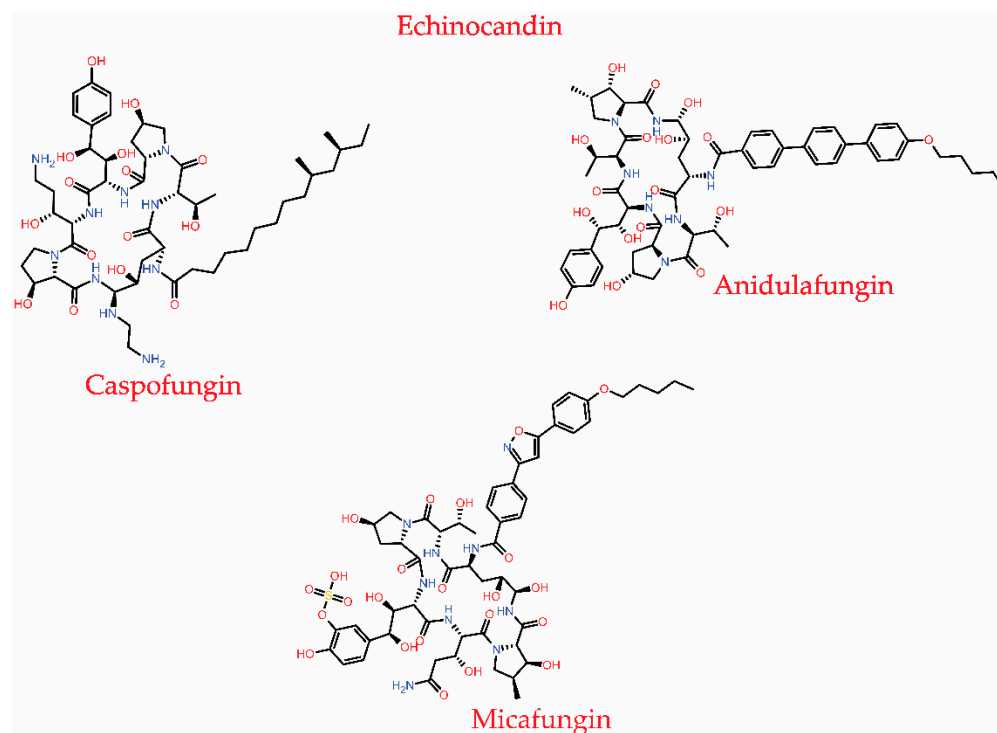


**Figure 7.** Two-dimensional structure of drugs under tetrazole group.

### 3.1.4. Echinocandins

Echinocandins, cyclic amphiphilic peptides with long lipophilic side chains, are the most recent class of antifungal drugs for the treatment of fungal infections; for example, caspofungin, micafungin, and anidulafungin belong to echinocandins, which are approved by the FDA and the European medicine agency for intravenous administration [74]. Echinocandins act as non-competitive inhibitors of  $\beta$ -1,3 glucan synthase, an essential enzyme complex responsible for cell wall synthesis in fungi [75]. This enzymatic disruption by echinocandins results in a leaky fungal cytoplasm and alters osmotic pressure, followed by fungal cell lysis, demonstrating fungicidal activity [76–79]. This fungicidal activity of echinocandins has been observed in the fungal species of *Candida* and *Saccharomyces*. Notably, due to the absence of cell walls in mammalian cells, it is less toxic and

has significantly less drug–drug interaction than other antifungal drugs, but they show low penetration into brain and CSF, due to which they are not preferred for the treatment of fungal infection in these tissues [76,80]. Echinocandins also demonstrated fungicidal activity against *Candida* spp. and fungistatic activity via inducing structural alterations in the fungus against *Aspergillus* spp. [81–84] (Table 1) (Figure 8).



**Figure 8.** Two-dimensional structure of drugs under the echinocandin antifungal group.

### 3.2. Resistance to Antifungal Agents

Antifungal resistance is an emerging worldwide issue due to new resistant variants of the existing fungal pathogens, for example, *Aspergillus fumigatus* and *Candida auris* [9,85]. To reduce toxicity in a host cell, the antifungal drug must act towards specific targets, which are not conserved between the fungi and the human host [17]. To overcome the issue of drug resistance, it is essential to understand the mechanism of drug resistance to antifungal drugs. Intrinsic and acquired resistance are two significant consequences of long-term treatment and a high range of prophylaxis [54]. The following sections briefly describe the mode of resistance gain in the fungi against four different classes of specific antifungal drugs.

#### 3.2.1. Polyene Resistance

Mechanisms of resistance to polyenes involve modification in the fungi membrane sterols, instigating antioxidant mechanisms to halt the damage caused by oxidative stress, and alternations in the ergosterol biosynthetic genes. Notably, twenty-five different genes are deciphered to play a significant role in regulating ergosterol biosynthesis (Table 1). For example, alternations (*ERG3*, *ERG5*, and *ERG11*), deletion (*ERG11*), and mutation (*ERG3*, *ERG5*, and *ERG11*) in the genes encoding respective enzymes were found to decrease the efficacy of amphotericin B in the treatment of infection caused by *Candida* species [86]. Mutation in the *MEC3*, a gene responsible for DNA damage homeostasis, also contributes to the high MIC of polyene in *C. auris* [87]. Moreover, overexpression of the molecular chaperones, such as the heat shock proteins (Hsp90 and Hsp70) family, further contributes to the development of intrinsic resistance in *Aspergillus terreus* against amphotericin B [88,89]. Alteration in the fungal cell wall composition, significantly increasing

the 1,3- $\alpha$ -glucan and 1,3- $\beta$ -glucan fraction, also contributes to amphotericin B resistance. This condition was seen in the amphotericin-B-resistant strains of *C. tropicalis*. The increase in the 1,3- $\beta$ -glucan is seen to be responsible for strengthening the immune response and survival of resistant strains [90].

### 3.2.2. Antimetabolite Agent Resistance

Flucytosine drugs make the pathogen easily resistant to the drug when used as monotherapy [91]. This is mainly used in combination therapy, along with amphotericin B and triazoles. It was found that some *Candida* spp. and *Saccharomyces cerevisiae* were resistant to this due to alternation in the *FCY2*, *FCY1*, and *FUR1* genes (Table 1), which are required for the uptake and conversion of flucytosine [92]. Another resistance mechanism reported is the upregulation of pyrimidines in de novo synthesis, leading to functional disturbance in uridine monophosphate pyrophosphorylase [91].

### 3.2.3. Azole Resistance

The most common mechanism in azole resistance is the alternation or overexpression of the *ERG11*, *Cyp51A*, and *Cyp51B* genes, which encode the lanosterol 14- $\alpha$ -demethylase enzyme belonging to the cytochrome P450 family [93]. For instance, overexpression of *ERG11* is due to the functional mutation in the Upc2 transcriptional activator in *C. albicans* [29]. Similarly, *A. fumigatus* becomes resistant to azole when alternation occurs in the *Srb* transcriptional factor, *HapE* modification, *Cyp51B* overexpression, and biofilm formation [18,94,95]. Upregulation of the *ABC* (adenosine triphosphate binding) cassette also causes acquired resistance to azole in *C. albicans* and *C. glabrata* [29]. The mutation of the *RAD50* gene in the double-strand break repair (DSBR) and the *MSH2* and *PMS1* genes of mismatch repair (MMR) of the DNA repair pathway are responsible for inducing fluconazole resistance in fungi. Additionally, cerebellar degeneration-related protein 1 and 2 (CDR1, CDR2) and SNQ2 are the transporters responsible for resistance to azole in fungi. For instance, when the SNQ2 transporter is removed from these groups, it was found to restore azole susceptibility, making it an essential factor in azole resistance [96]. Additionally, azole resistance was observed in *Candida dubliniensis*, a pathogenic yeast with phenotypic characteristics similar to *Candida albicans*, which mainly infects patients with HIV/AIDS. *C. parapsilosis*, *C. tropicalis*, and some non-*albicans* species were also found to be resistant against azoles and typically gained azole resistance by drug efflux, drug target modification, and alternation in the ergosterol biosynthesis [97,98] (Table 1).

### 3.2.4. Echinocandin Resistance

The echinocandin resistance mainly occurs due to the amino acid substitution or point mutation in the *FKS1* and *FKS2* subunits in glucan synthase [99]. In *C. albicans*, the change occurs in Ser<sup>641</sup> and Ser<sup>645</sup> and in *C. glabrata*, it is seen in Ser<sup>629</sup>, Phe<sup>659</sup>, and Ser<sup>663</sup>; both these changes lead to resistance [100]. Resistance mediated by *FKS2* is only seen in *C. glabrata* and can be reversed using the calcineurin inhibitor tacrolimus [18,29]. In *A. fumigatus*, in vitro resistance of echinocandin was found when induced with the substitution of Ser<sup>678</sup> in *FSK1* [94] (Table 1)

**Table 1.** List of all four groups of antifungal agents and their resistance mechanism.

Antifungal Agents	Mode of Action	Example of Anti-fungal Resistant Species	Resistance Mechanism	Reference
Polyenes • Amphotericin B • Nystatin • Natamycin	<ul style="list-style-type: none"> <li>○ Cell membrane disruption by binding to ergosterol.</li> <li>○ Excessive production and accumulation of reactive oxygen species (ROS) causing damage to the cell</li> </ul>	<i>Candida species</i> , and <i>Aspergillus terreus</i>	Alternation ( <i>ERG3</i> , <i>ERG5</i> , and <i>ERG11</i> ), deletion ( <i>ERG11</i> ), and mutation ( <i>ERG3</i> , <i>ERG5</i> , and <i>ERG11</i> ) genes encoding for C-5 sterol desaturase, C-22	[36,41,93]

	membrane, DNA, fungal protein, and mitochondria		sterol desaturase, and sterol 14-demethylase are responsible for the regulation of ergosterol biosynthesis.
<b>Antimetabolite agent</b>	Inhibits RNA and protein synthesis by binding 5-fluorouracil(5-FU) to the RNA strand.	<i>Candida</i> spp.	Alteration in the <i>FCY2</i> , <i>FCY1</i> , and <i>FUR1</i> genes that are responsible for the uptake and conversion of flucytosine. [54,55,92]
<b>Azole</b>			
• <b>Imidazole</b>			
○ Miconazole			
○ Clotrimazole			
○ Econazole			
○ Ketoconazole			
○ Tioconazole			
○ Sulconazole			
○ Sertaconazole			
○ Luliconazole			
○ Tinidazole			
○ Enilconazole			
○ Parconazole			
○ Eberconazole			
○ Lanoconazole			
○ Fenticonazole	Inhibition of fungal lanosterol 14 $\alpha$ -demethylase cytochrome P450 51,		
○ Bifonazole	P450 3A4, P450 2D6, P450 2C8, P450		
○ Sulconazole	2A6, P450 2E1 P450 2C9, etc., that	<i>C. albicans</i> , <i>C. glabrata</i> , <i>C. dubliniensis</i>	Alteration and overexpression of <i>ERG11</i> , <i>Cyp51A</i> , and [65,67,70,
○ Lombazole	disrupts the ergosterol biosynthesis leading to cell death and lysis. Some	<i>A. fumigatus</i> , yeast,	<i>Cyp51B</i> encodes for lanosterol 71,93,101
○ Sertaconazole	of these drugs also targets DNA, bile salt export pump, P-glycoprotein 1,	<i>Trichophyton rubrum</i>	14-alpha-demethylase enzyme. ,102]
○ Oxiconazole	Hydroxycarboxylic acid receptor 2,		
○ Butaconazole	oxygen-insensitive NADPH nitroreductase.		
○ Isoconazole			
○ Flutrimazole			
○ Ornidazole			
○ Metronizadole			
• <b>Triazoles</b>			
○ Itraconazole			
○ Fluconazole			
○ Voriconazole			
○ Posaconazole			
○ Isavuconazole			
○ Fosravuconazole			
○ Albaconazole			
○ Letrozole			
○ Anastrozole			
• <b>Tetrazole</b>			
○ Quilseconazole (VT-1129)			
○ Oteseconazole (VT-1161)			
<b>Echinocandin</b>	Inhibits $\beta$ -1,3 glucan synthase, an essential enzyme complex responsible for cell wall synthesis in fungi.	<i>C. glabrata</i> and <i>A. fumigatus</i>	Amino acid substitution or point mutation in the <i>FKS1</i> and <i>FKS2</i> genes present in the [75,99]
• Caspofungin			
• Micafungin			
• Anidulafungin			

### 3.2.5. Other Mechanisms Involved in Drug Resistance

#### (a) Biofilm formation

Biofilms are a surface-associated microbial communities attached by a self-made extracellular matrix [103]. The biofilm formation helps the organism withstand antimicrobial agents' attack and survive severe environmental conditions [104]. In the case of fungal pathogens, such as *C. albicans*, biofilm formation is involved in the pathogenicity and death of patients [105,106]. Researchers found that *C. auris* adheres to a polymeric surface that further develops into a biofilm and can resist various antifungal agents, especially caspofungin, which are very effective against *Candida* biofilms [107]. Fungal biofilms also show resistance to echinocandin and azole antifungal drugs [108,109].

#### (b) Modification of drug targets

The drug targets are modified by specific genetic changes that lead to a reduction in the affinity of drugs. For example, modification of echinocandin drug targets confers drug resistance [110]. Mutation in the *FKS* gene that encodes  $\beta$ -D-glucan synthase leads to substituting amino acids that result in the target site modification [111,112]. The drug affinity becomes reduced by several folds due to the change of targets. The mutation in the *ERG11* also results in amino acid substitutions that are responsible for azole resistance, and this further decreases the affinity of the azoles to the target site [113,114].

#### (c) Efflux pump

Fungal efflux is responsible for regulating the environment and removing toxic substances, including antifungal agents. The overexpression of various drug transporters leads to the efflux of the drug and also prevents the accumulation of antifungal medications in the cytosol [115]. Two major efflux pumps involved in drug resistance are the ATP-binding cassette (ABC) superfamily and the major facilitator superfamily (MFS). Among azole-resistant (AR) clinical isolates of *C. albicans*, *CDR1* and *CDR2* are two promiscuous ABC proteins that are upregulated [116,117].

### 3.3. Strategies to Overcome Resistance

A deeper understanding of antifungal resistance is important for developing effective counterstrategies to overcome it. With the emergence of antifungal resistance, the chances of survival of opportunistic pathogens have also increased, so it is important to find various practical approaches to eradicate drug resistance issues in pathogenic fungi.

#### 3.3.1. Development of a New Antifungal Drug

Developing new antifungal drugs is one of the most essential strategies to reduce the risk of antifungal resistance. Several drugs have been identified and synthesized by researchers, which are currently in the preclinical and clinical trial stages. Some of the compounds have similar properties but may have better efficacy than the existing compound and some of the compounds may have a new mechanism of action.

##### (a) SUBA-itraconazole and VT-1598

The super bioavailable itraconazole (SUBA-ITC) is approved by the FDA and used as a first-line antifungal agent for the treatment of allergic bronchopulmonary aspergillosis (ABPA) and other fungal infections in children [118]. This antifungal formulation was developed by Mayne Pharma L.td., mainly to enhance the bioavailability of itraconazole. It also ensures the bioavailability by solid dispersion in a pH-dependent matrix and interferes with the cytochrome P450 activity to reduce the rate of ergosterol synthesis. This antifungal formulation also showed broad-spectrum activity against *Blastomycosis* sp., *Histoplasmosis* sp., and *Aspergillosis* sp. [54,119].

Mycovia Pharmaceuticals has created VT-1598, the most recent tetrazole inhibitor of lanosterol 14-demethylase encoded by the *Cyp51* gene. VT-1598, which is still in Phase I of the clinical study, is effective against moulds, including *Aspergillus* species, *Coccidioides* species, and *Rhizopus arrhizus* [102,120] (Table 2) (Figure 9a).

## (b) Rezafungin and Ibrexafungerp

Rezafungin, developed by Cidara Therapeutics, belongs to the class of echinocandins that inhibits 1,3- $\beta$ -D-glucan synthesis and has been granted orphan drug designation by the FDA for the treatment of vulvovaginal candidiasis. It has been reported effective against *Candida* sp., *Aspergillus* sp., and *Pneumocystis* sp. [119,121]. Ibrexafungerp, an FDA-approved antifungal triterpenoid, was developed by SCYNEXIS Inc. and mainly inhibits 1,3- $\beta$ -D-glucan synthesis. Both rezafungin and ibrexafungerp have low toxicity, are highly bioavailable, and significantly inactivate the *Candida* sp., including *C. glabrata* and *C. auris* [122] (Table 2) (Figure 9d,e).

## (c) Olorofim

Another drug in development is olorofim, which comes under a new class of antifungal drugs called orotomides and is in the Phase II stage of a clinical trial. This class has a novel mode of action, where it inhibits the dihydroorotate dehydrogenase enzyme, an important enzyme in pyrimidine biosynthesis in fungi, to disturb the nucleic acid and cell wall synthesis. This class of antifungal has been reported with potential efficiency against *Aspergillus* sp. and *Scedosporium* sp. [32,123] (Table 2) (Figure 9c).

## (d) Amphotericin B Cochleate (CAMB)

The amphotericin B cochleate (CAMB), an oral formulation of amphotericin B, is a new drug derived from the polyene class and is under the Phase II stage of clinical trials, developed by Matinas BioPharma. CAMBs are formulated using phosphatidylserine along with phospholipid-calcium and are stable against degradation in the gastrointestinal tract. This formulation was successfully reported to treat *C. albicans* infection in a murine model [124] (Table 2).

## (e) MGCD290

MGCD290 is an orally administered drug that inhibits the Hos2 fungal histone deacetylase (HDAC) enzyme as well as affects the non-histone protein such as Hsp90. This formulation was developed by Mirati Therapeutics, Inc. and is under Phase II clinical trial against fungal infection. This drug is co-administrated with both azole and echinocandins, and is known for fungicidal activity against *Candida* sp. and *Aspergillus* sp. [119] (Table 2).

## (f) Fosmanogepix (APX001)

Fosmanogepix is a glycosylphosphatidylinositol inhibitor developed by Amplyx Pharmaceuticals which is currently in Phase II clinical trials. It is metabolized into its active form, manogepix, which targets the enzyme Gwt1, responsible for the glycosylphosphatidylinositol (GPI) anchor biosynthesis [125,126]. Fosmanogepix was also reported to actively inhibit the growth of yeast, moulds, *Candida* sp., *Cryptococcus* sp., *Coccidioides* sp., and *Aspergillus* sp. [127] (Table 2) (Figure 9b).

**Table 2.** List of new antifungal drugs.

New Antifungal Agents Developed	Mode of Action	Developer	Activity Against	Clinical Trial Stage	Reference
VT-1598	Inhibits lanosterol 14 $\alpha$ -demethylase	Mycovia Pharmaceuticals, North Carolina, USA	Moulds, <i>Aspergillus</i> spp., <i>Rhizopus arrhizus</i> , and <i>Coccidioides</i>	Phase I	[102,120]
SUBA-itraconazole	Enhance the bioavailability of itraconazole by its solid dispersion in a pH-dependent matrix.	Mayne Pharma Ltd. Salisbury South, Australia	<i>Aspergillus</i> spp., <i>Blastomyces dermatitidis</i>	FDA-approved	[118]
Rezafungin	Inhibition of 1,3- $\beta$ -D-glucan synthesis	Cidara Therapeutics, San Diego, California, USA	<i>Candida</i> spp., <i>Aspergillus</i> spp., and <i>Pneumocystis</i> spp.	FDA-approved as an orphan drug for the treatment	[119,121]

				of vulvovaginal candidiasis	
Ibrexafungin	Inhibition of 1,3-β-D-glucan synthesis	SCYNEXIS Inc., New Jersey, USA	<i>Candida</i> spp.	FDA-approved	[122]
Olorofim	Inhibition of dihydroorotate dehydrogenase responsible for pyrimidine biosynthesis.	Shionogi & Co., Ltd. And F2G Ltd., Osaka, Japan and Manchester, UK	<i>Aspergillus</i> and <i>Scedosporium</i> spp.	Phase II	[123]
Amphotericin B Cochleate (CAMB)	Cell wall disruption	Matinas Bio-Pharma, New Jersey, USA.	<i>C. albicans</i>	Phase II	[124]
MGCD290	Inhibits Hos2 fungal histone deacetylase (HDAC) and also affects non-histone protein Hsp90	Mirati Therapeutics, Inc., California, USA	<i>Candida</i> and <i>Aspergillus</i> spp.	Phase II	[119]
Fosmanogepix (APX001)	Inhibition of glycosylphosphatidylinositol.	Amplix Pharmaceuticals, California, USA	Yeast, moulds, <i>Candida</i> , <i>Cryptococcus</i> , <i>Coccidioides</i> , and <i>Aspergillus</i> spp.	Phase II	[125,126]
VL-2397	Unknown	Vical Pharmaceuticals, California, USA	<i>Aspergillus fumigatus</i>	Phase II	[128,129]
T-2307	Fungal mitochondrial disruption	Toyama Chemical Co, Tokyo, Japan.	<i>Candida</i> spp. <i>Cryptococcus</i> and <i>Aspergillus</i> spp.	Phase I	[130]

## (g) VL-2397

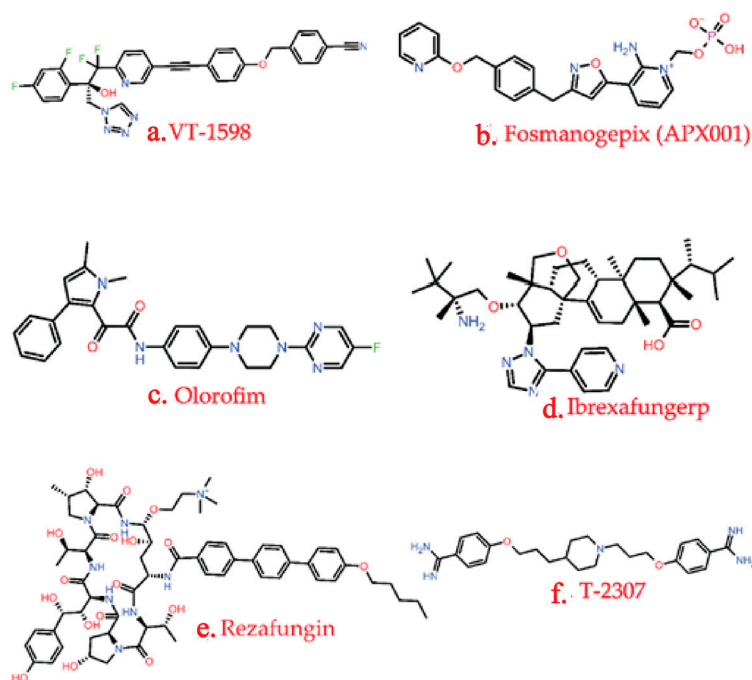
VL-2397, also known as ASP2397, was isolated from *Acremonium* sp. and formulated by Vical Pharmaceuticals. Fungi uptakes this drug through the siderophore iron transporter 1 (Sit1), leading to disruption of the intercellular process. It was reportedly effective against *Aspergillus fumigatus* under Phase II of clinical trials [121,128,129] (Table 2).

## (h) T-2307

This arylamide compound developed by Toyama Chemical Co., Toyama, Japan, is in Phase I of clinical trials. The structure of this drug is similar to that of aromatic diamidines [131]. It mainly disrupts fungal mitochondrial membranes and is effective against *Candida* sp., *Cryptococcus* sp., and *Aspergillus* sp. [130,132] (Table 2) (Figure 9f).

## (i) Retinoids and All-trans retinoic acid (ATRA)

Retinoids and all-trans retinoic acid (ATRA) are derived from Vitamin A. Studies have proved that these compounds have antifungal properties and can effectively treat various skin and systemic fungal infections. These compounds have shown a broad spectrum of activity against yeast, *Candida albicans*, and *Aspergillus fumigatus* [133,134]. Vitamin A serum is eligible for clinical practice to prevent skin and systemic fungal infection in psoriatic patients when treated with IL-17 inhibitor [135].



**Figure 9.** Two-dimensional structure of new antifungal agents developed.

### 3.3.2. Combination Therapy

Combination therapy is an effective approach to antifungal drug resistance and mainly helpful in extending the useful life of existing drugs. The combination of two drugs can be a much more effective way of killing the pathogen, reducing the pathogen population, and minimizing the chances of experiencing acquired resistance mutation [136,137]. Combination therapy also reduces the individual dosage of the drug, the length of treatment, and drug toxicity [136]. For instance, combination of amphotericin B and fluorocytosine is the best example of combination therapy for the treatment of persistent *Cryptococcal meningitis* and *Candida* fungal infections [138].

### 3.3.3. Antifungal Stewardship

Antifungal stewardship refers to the method in which the diagnostics method and the usage and dosage of antifungal drugs are combined to obtain a gained better clinical outcome, which can reduce the emergence of resistance in fungi [139,140]. As antifungal management is mainly based on the guidelines used for the treatment and diagnostic tests that are safe for the patient, the combination of health professionals and public health centres following the guidelines for proper care of patients formed a stewardship team to execute the program [141,142]. However, several health centres lack antifungal stewardship due to unavailability or lack of access to long-term diagnostic tools, and, hence, failed to manage and execute the program. Thus, it is recommended for each and every health care centre and institution to conduct antifungal stewardship programs that contain guidelines and diagnostic tests to guide patients about the treatment duration, as well as guidance from the specialists and pharmacists to adopt the dosage and mode of administration of drugs [143]. The Mycoses Study Group Education and Research Consortium (MSGERC) has designed a set of core recommendations that can uplift the practice of antifungal stewardship. Their core recommendation includes integration of antifungal stewardship goals into hospital strategic management policies with the proper guidance of a senior leader and a core team. The core team should have the knowledge of and clinical expertise in invasive fungal infection management. The team should be able to develop collaborative strategies with the help of clinical specialists for better antifungal therapies. They also recommend proper diagnostic testing for *Candida* and *Aspergillus* species



[140]. A study conducted by Valerio and team on antifungal stewardship in tertiary care institutes found that the annual antifungal drug expenditure was reduced within 2 years of implementation of an antifungal stewardship program. Along with that, the number of incidences was also reduced. This study helped to understand that the impact of antifungal stewardship can be an efficacious and cost-effective approach [144]. Considering all these factors, antifungal stewardship can significantly contribute to the prevention of drug resistance in fungi.

#### 3.3.4. Potential Drug Targets to Overcome Antifungal Resistance

A better understanding of the drug resistance mechanism can assist in finding new drug targets which can be targeted for the designing and development of novel antifungal drugs. For instance, Hsp 90 of fungi are one major drug target, as they contribute to the cellular stress response in fungi [145]. However, incompetence in the Hsp90 function leads to resistance development in fungi against the azole and echinocandin classes [146]. Thus, inhibition of Hsp90, such as that exhibited by geldanamycin, was adopted to reduce resistance to caspofungin in *A. fumigatus* [147]. Additionally, calcineurin is another drug target which is required for growth, virulence, and drug resistance development in fungi such as *C. albicans*, *A. fumigatus*, and *C. neoformans* [148,149]. Hence, such essential enzymes of the fungi can be targeted for the designing and development of new antifungal agents to escape the drug resistance in pathogenic fungi.

#### 3.3.5. One Health Approach in Combating Antifungal Resistance

The “One health” concept identifies that human, animals, and the environment are all indistinguishably connected and influential to one another [150]. This concept was first designed to combat the health of humans, plants, and animals during the emergence of the SARS-CoV and influenza virus in the year 2003–2004 and gained importance with the evolution of antimicrobial resistance in humans [151]. The number of patients who are highly susceptible to fungal infections is increasing as a result of changes in medical care, and this problem is made worse by the fact that our meagre supply of antifungal medications is being threatened (or compromised) by the emergence of drug-resistant fungi strains, which in some cases are linked to antifungal agents used in agriculture. The “One Health” idea seeks to educate scientists and decision-makers about these recognized and newly developing fungal dangers to global health by stressing the convergence of these dynamics. [150]. The American Academy of Microbiology organized a colloquium in October 2017 to address this issue, assembling a multidisciplinary and worldwide team of experts. This gathering took place ten years after the Academy hosted the first of its kind in 2007. One Health: Fungal Pathogens of Humans, Animals, and Plants, a new report, emphasizes the field’s extraordinary advancements and offers updated suggestions to deal with problems in public health and science in the modern world [150]. For instance, this concept has been used to closely analyse azole fungicide resistance in the environment and to reduce the burden of environmental resistance [152]. The “One Health” approach focuses on finding and prioritizing effective research methodologies that can be useful in developing various preventive guidelines and fill the major gaps in antifungal resistance conditions, as they can also originate from the environment, as well as from hospital facilities [8,153].

## 4. Conclusions

The rise in fungal infections has paradoxically advanced medicine. As a result, the frequent utilization of antifungal agents has also been transpired. In clinical setups, researchers and clinicians are facing the serious threat of antifungal resistance. Thus, along with the old strategies, some new methods are gaining popularity to tackle the growing drug resistance in fungi, as these new methods are more robust, and the accuracy lever is higher compared to the traditional ones. Additionally, new combinational therapies and

drug trials are coming to light which will lead us closer to winning the war against anti-fungal resistance with better drugs with lower toxicity, fewer cross-interactions with side proteins, and higher specificity to the target sites. Ultimately, elucidating strategies to combat fungal pathogens in humans may cast light on how fungal infections that pose a worldwide danger to biodiversity can be defeated.

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