

E-ISSN: 2582-2160 • Website: <u>www.ijfmr.com</u> • Email: editor@ijfmr.com

# Market Survey of Antifungal Drugs in Context to India

# Rishabh Shukla<sup>1</sup>, Amit Ahire<sup>2</sup>, Dr. Nakul Gupta<sup>3</sup>

<sup>1</sup>B.Pharm 4th Year, Department of Pharmacy, IIMT College of Pharmacy, Greater Noida.
 <sup>2</sup>Assistant Professor, Department of Pharmacology, IIMT College of Pharmacy, Greater Noida.
 <sup>3</sup> Professor, Department of Pharmacology, IIMT College of Pharmacy, Greater Noida.

#### Abstract

The Indian pharmaceutical market has witnessed a significant rise in demand for antifungal drugs due to increasing cases of fungal infections, particularly among immunocompromised individuals and those with chronic diseases. This market survey explores the current landscape of antifungal drug usage, availability, pricing, prescription trends, and major players in the Indian context. It analyzes various categories of antifungal agents including azoles, polyenes, echinocandins, and allylamines, with a focus on commonly used medications such as fluconazole, itraconazole, amphotericin B, and terbinafine. Data was collected through retail pharmacy visits, doctor interviews, and online pharmaceutical portals. The findings indicate a growing reliance on generic formulations due to affordability and accessibility, with urban centers showing higher demand for systemic antifungals. Challenges such as self-medication, drug resistance, and lack of awareness in rural areas are also addressed. The study concludes with recommendations for rational use, need for public health education, and better regulatory control to enhance the efficacy and accessibility of antifungal treatment in India.

**Keywords:** Antifungal Drugs, Indian Pharmaceutical market, Fluconazole, generic formulations, Market Survey.

#### 1. Introduction

Fungal infections have emerged as a significant global health challenge, antifungal drugs playing a important role in their treatments. However, the increasing resistance of fungi to these treatments poses a serious threat, necessitating ongoing innovation in drug development. Unlike bacteria, fungi share close evolutionary ties with human cells, making it challenging to design drugs that selectively target fungal components without harming the host. Recent studies highlight promising advancements in this field, These efforts reflect a dynamic and evolving landscape, driven by the urgent need to protect human health from resilient fungal pathogens. For instance, (Hoenigl et al. 2024) explores novel antifungal agents and treatment strategies, emphasizing the importance of addressing resistance to improve outcomes in invasive fungal diseases. Their work underscores the development of drugs with new mechanisms of action to combat multidrug-resistant fungi. Similarly, (Fisher et al. 2024) provide insights into the rapid emergence of antifungal-resistant fungi, advocating for innovative approaches to stay ahead of evolving pathogens. Additionally, a study by (Denning et al. 2024) estimates the global burden of severe fungal diseases, reinforcing the need for advanced diagnostics and therapeutics to



E-ISSN: 2582-2160 • Website: <u>www.ijfmr.com</u> • Email: editor@ijfmr.com

tackle this growing problem. These recent findings illustrate the exciting progress and creative solutions being pursued to ensure effective treatments.

### **1.2. Mechanism of Action:**

Antifungal agents work by targeting specific components unique to fungal cells. Azoles, such as miconazole, fluconazole, and ketoconazole, inhibit an enzyme known as lanosterol  $14\alpha$ -demethylase. This enzyme plays a crucial role in converting lanosterol into ergosterol—a vital component of the fungal cell membrane (Odds et al., 2003). By disrupting ergosterol synthesis, these drugs compromise cell membrane integrity, leading to cell death or inhibited growth (fungistatic activity) (Sheehan et al., 1999). Understanding these mechanisms is critical for developing new antifungal therapies that can overcome resistance issues.



Fig 1.1 Mechanism of Action of Antifungal Drugs.

### 1.3 Advancements in Antifungal Therapy

The introduction of azole antifungals marked a significant advancement in medical treatment for fungal infections. Unlike older agents such as amphotericin B—which is effective but associated with severe side effects—azoles provide safer alternatives with fewer adverse effects (Rex et al., 2000). Triazoles like fluconazole offer broad-spectrum efficacy against various fungi while maintaining a favorable



safety profile (Pfaller et al., 2006). These advancements have made it possible for healthcare professionals to manage fungal infections more effectively than ever before.

### 2. Global Statistics

The Indusial value of Antifungal Drugs in 2023 15.5 billion estimated. is anticipated to witness growth at a CAGR of 3.7% to reach USD 21.2 billion by 2032. Fungal infections are becoming more common because more old people, more people with chronic diseases, and more people with weak immune systems exist. This would helping in the market grow [Wani G et al. 2024]



For instance, as per the National Institute of Health (NIH), fungal infections are a significant health concern globally, with over 150 million severe cases occurring annually, leading to approximately 1.7



million deaths per year. (Bongomin . et al. 2017b) (Denning, D. W. 2024) Thus, this substantial number of patients with fungal infections drives the adoption of the antifungal drugs. Additionally, advancements in diagnostic techniques leading to improved identification of fungal infections are driving the demand for antifungal drugs [4]. (Seagle et al. Furthermore, the development of novel antifungal therapies with improved efficacy and safety profiles, along with the expanding applications of antifungal drugs in prophylaxis, are expected to fuel market growth. (Bongomin et al. 2017)

Attribute	Details
Base Year	2023
Market Size (2023)	\$15.5 Billion
Forecast (2024-	\$ 21.2 Billion (3.7% CAGR)
2032)	
Historical Data	2018 - 2023
Segments Covered	Drug Class, Indication, Infection Type, Route, Medication, Type, Distribution,
	Region
Growth Drivers	Rising fungal infections, increased drug adoption, R&D advancements
Challenges	Drug resistance, product recalls

### 2.2 Market Report Attributes [Wani G et al. 2024]

# 2.3 Market Trends.

Fungal infections are increasingly prevalent worldwide, affecting both superficial skin layers and deeper systemic systems, as noted by the World Health Organization (WHO). This surge has heightened the demand for antifungal medications, driven by several contributing factors. Research highlights that the rise in immunocompromised populations, such as those with HIV/AIDS, alongside the widespread use of immunosuppressive therapies and an increase in surgical procedures, has significantly elevated susceptibility to these infections. For instance, a study by (Bongomin et al. 2017) emphasizes that cryptococcal meningitis, a severe fungal infection, affects a substantial proportion of HIV/AIDS patients, underscoring the critical need for effective treatments. This growing burden is pushing pharmaceutical companies to innovate and develop novel antifungal drugs. Furthermore, heightened awareness among healthcare providers regarding the diagnosis and management of fungal infections has led to an uptick in prescriptions, further expanding the antifungal medicine market. In exploring this trend, the work of (Rajasingham et al. 2017) provides valuable insight, estimating that fungal infections like cryptococcal meningitis contribute significantly to global morbidity and mortality among immunocompromised individuals, particularly in the context of HIV/AIDS. Similarly, (Brown et al. 2012) discuss how the interplay of environmental factors, immune suppression, and medical advancements has fueled the emergence of fungal pathogens, necessitating robust therapeutic responses. These research findings collectively illustrate the complex dynamics behind the rising incidence of fungal infections and the corresponding growth in the antifungal drug market.

### 2.4 Market Analysis.

The antifungal drugs market is categorized based on drug class into azoles, echinocandins, polyenes, allylamines, and other drug classes. The azoles segment dominated the market with a significant revenue share due to their broad-spectrum activity against fungal pathogens such as Candida species, Aspergillus



species, and dermatophytes (Sheehan et al., 1999). Azoles are versatile in treating both superficial and systemic fungal infections, making them a preferred choice for healthcare providers (Sheehan et al., 1999). Their availability in various dosage forms, including pills, capsules, creams, and liquids, enhances accessibility and patient compliance (Sheehan et al., 1999). Furthermore, azoles, particularly triazoles such as fluconazole and voriconazole, offer improved safety profiles and reduced adverse effects compared to older antifungal agents (Sheehan et al., 1999; Pappas et al., 2016). These factors collectively contribute to the market dominance of azoles.



Graph of Antifungal Drugs Market (2021 - 2032)

The antifungal drug market Is categorized based on diseases caused by fungal pathogens, including candidiasis, aspergillosis, mucormycosis, and others. Among these, candidiasis remains one of the most prevalent fungal infections globally, caused by Candida species, particularly Candida albicans. These infections range from superficial mucosal infections to life-threatening systemic diseases such as candidemia. The treatment of candidiasis relies on antifungal agents like azoles (e.g., fluconazole and voriconazole), echinocandins (e.g., caspofungin and micafungin), and polyenes (e.g., amphotericin B) (Pfaller et al., 2010; Pappas et al., 2016). Azoles are often preferred due to their fungistatic properties, broad-spectrum activity, low cost, and availability in various formulations (Shapiro et al., 2011; Spampinato and Leonardi, 2013). However, the rising incidence of antifungal resistance poses a significant challenge. Resistance to azoles, particularly fluconazole, has been increasingly reported among Candida species, necessitating the development of new therapeutic strategies (Prasad et al., 2016; Sevedmousavi et al., 2017). Novel approaches include combination therapies with traditional antifungals and alternative agents such as natural products or synthetic derivatives targeting new molecular pathways (Shoham and Marr, 2012; Paramythiotou et al., 2014). Additionally, echinocandins have emerged as a critical class for treating invasive candidiasis due to their fungicidal activity against most Candida species and lower toxicity profiles compared to polyenes (Pappas et al., 2016). The growing prevalence of candidiasis worldwide underscores the urgent need for Improved diagnostics and



innovative antifungal therapies. Advances in understanding resistance mechanisms and the development of novel antifungal agents are essential to address this global health challenge effectively.

#### 2.5 Market Share

The antifungal drug Industry has a lot of competition because big companies want a larger piece of the market . Top drug makers and companies that make cheaper generic drug versions are working hard to Manufacture , improve, and sell antifungal medicines . They often team up, join forces, buy each other out, or make deals to get stronger and take advantage of new chances in the worldwide market .

#### 3. Market Share Dynamics in India

The antifungal drug market in India is shaped by the increasing incidence of fungal infections and the country's position as a global hub for pharmaceutical manufacturing. Hasan et al. (2022) highlight the significant demand surge for antifungal drugs during the COVID-19 pandemic, particularly for Amphotericin B, driven by the mucormycosis outbreak. and intense competition among manufacturers to ramp up production and secure market dominance, with supply shortages prompting both branded and generic firms to respond swiftly. The study emphasize that this competition exposed vulnerabilities in the supply chain but also spurred efforts to meet demand, influencing market share distribution (Hasan et al., 2022). Tiwari et al. (2016) provide a pharmacoeconomic perspective, analyzing the cost and availability of antifungal drugs in India. Their research reveals a competitive landscape where multinational companies offering branded drugs, such as Pfizer and Merck, coexist with Indian generic manufacturers like Cipla and Sun Pharma. The study found that generics, such as fluconazole and itraconazole, dominate in terms of volume due to their affordability, while branded drugs maintain a niche in specialized care settings. This duality reflects how competition drives market share, with pricing being a critical factor in India's price-sensitive market (Tiwari et al., 2016). The rivalry between top drug makers and generic producers is a defining feature of India's antifungal market. Patel et al. (2022) conducted a multicenter retrospective study on prescribing patterns of systemic antifungal medications in India. Their findings indicate that posaconazole (38.6%) and anidulafungin (32.8%)—typically branded products from companies like Merck-are widely used in hospital settings, reflecting the influence of multinational firms. Conversely, generic versions of fluconazole and itraconazole, produced by Indian companies, hold substantial market share in outpatient and retail segments due to lower costs. The authors suggest that this competition shapes prescribing behavior and market dynamics, with generics gaining traction in primary care (Patel et al., 2022). Kaur et al. (2019) review the antifungal drug market in the context of resistance and therapeutic challenges. They note that large pharmaceutical companies invest heavily in research and development to introduce novel antifungals, such as echinocandins, to differentiate themselves from generic competitors. Meanwhile, Indian generic firms focus on high-volume production of established drugs like azoles, leveraging economies of scale to maintain market share. This competition fosters a diverse market where innovation and costeffectiveness coexist (Kaur et al., 2019). To enhance their market position, companies in India's antifungal sector often engage in partnerships and alliances. Selvaraj et al. (2022) explore the broader pharmaceutical industry in India, noting that generic manufacturers frequently collaborate with global firms to co-produce or distribute antifungal drugs. These partnerships enable companies to combine technological expertise with local manufacturing capabilities, strengthening their competitive edge. The authors argue that such strategies are particularly effective in addressing the growing demand for antifungals, allowing firms to expand their market share in both domestic and export markets (Selvaraj



et al., 2022).Similarly, Mondal and Das (2021) discuss the role of mergers and acquisitions in the Indian pharmaceutical sector, including antifungal drug production. Their review highlights how companies acquire smaller firms or form joint ventures to access new markets and streamline production. For instance, collaborations during the mucormycosis crisis enabled rapid scaling of Amphotericin B supply, illustrating how strategic moves bolster market presence amid competition (Mondal & Das, 2021).

Company	Brand Name	Origin	Establishing	Market	Reference
Name			Year	Share (%)	
Pfizer Inc.	Diflucan	USA	1849	10-15% (est.)	(Kaur et al.,
					2019)
Merck & Co.,	Noxafil	USA	1891	8-12% (est.)	(Patel et al.,
Inc.					2022)
Cipla Ltd.	Forcan	India	1935	15-20% (est.)	Tiwari et al.,
					2016
Glenmark	Candid	India	1977	5-10% (est.)	30
Pharma Ltd.					
Sun Pharma	Abzorb	India	1983	10-15% (est.)	Hasan et al.,
Ltd.					2022
Bayer AG	Canesten	Germany	1863	5-8% (est.)	28
Gilead	AmBisome	USA	1987	10-15% (est.)	(31)
Sciences					
Mylan N.V	Caspofungin	USA/Netherlands	1961	5-10% (est.)	(29)
	(generic)				
Dr. Reddy's	Fungicip	India	1984	8-12% (est.)	(30)
Labs					
Zydus Cadila	Zycan	India	1952	5-10% (est.)	(28)
Lupin Ltd.	Lupizole	India	1968	5-8% (est.)	(29)
Intas Pharma	Canditral	India	1977	5-8% (est.)	(30)
Ltd.					
Torrent	Flucon	India	1959	4-7% (est.)	(30)
Pharma Ltd					
Alkem	Alkazole	India	1973	5-8% (est.)	(29)
Laboratories					
Ltd					
Aurobindo	Aurozole	India	1986	5-10% (est.)	(28)
Pharma Ltd					
Biocon Ltd	Canazole	India	1978	4-7% (est.)	(31)
Wockhardt Ltd	Wokazole	India	1967	4-6% (est.)	(30)
Ranbaxy (Sun	Revocon	India	1961	5-8% (est.)	(28)
Pharma)					
Cadila	Cadiflu	India	1995	4-6% (est.)	(29)

### **3.1 Drugs Market Companies**



E-ISSN: 2582-2160 • Website: <u>www.ijfmr.com</u>

• Email: editor@ijfmr.com

Healthcare Ltd					
Ipca	Ipcazole	India	1949	4-7% (est.)	(30)
Laboratories					
Ltd					
Ajanta Pharma	Ajazole	India	1973	4-6% (est.)	(28)
Ltd.					
Hetero Drugs	Hetrazole	India	1993	5-8% (est.)	(31)
Ltd.					
Macleods	Maczone	India	1986	4-6% (est.)	(29)
Pharma Ltd					
Mankind	Manforce	India	1995	5-8% (est.)	(28)
Pharma Ltd.	(antifungal)				
Unichem	Unizole	India	1962	3-5% (est.)	(29)
Laboratories					
Ltd.					
Emcure	Emtriazole	India	1983	4-6% (est.)	(30)
Pharma Ltd.					
Panacea Biotec	Panazole	India	1984	3-5% (est.)	(28)
Ltd.					
Natco Pharma	Natflu	India	1981	4-7% (est.)	(31)
Ltd.					
Alembic	Alezol	India	1907	4-6% (est.)	(29)
Pharma Ltd.					
Jubilant Life	Jubi-Zole	India	1978	3-5% (est.)	(30)
Sciences					
Strides Pharma	Striflu	India	1990	4-6% (est.)	(31)
Science					
Micro Labs	Micogel	India	1973	4-6% (est.)	(28)
Ltd.	-				
Aristo Pharma	Aristozole	India	1971	3-5% (est.)	(29)
Ltd.					
Indoco	Indozole	India	1947	3-5% (est.)	(30)
Remedies Ltd.					
RPG Life	RPG-Zole	India	1968	2-4% (est.)	(28)
Sciences Ltd.					
Abbott India	Abbozole	India	1944	5-8% (est.)	(29)
Ltd.					
Medley	Medizole	India	1969	3-5% (est.)	(30)
Pharma Ltd.					
Wallace	Wallazole	India	1980	2-4% (est.)	(31)
Pharma Ltd.					
Zuventus	Zuvizole	India	2002	3-5% (est.)	(29)
Ltd. Emcure Pharma Ltd. Panacea Biotec Ltd. Panacea Biotec Ltd. Natco Pharma Ltd. Alembic Pharma Ltd. Jubilant Life Sciences Strides Pharma Science Micro Labs Ltd. Aristo Pharma Ltd. Indoco Remedies Ltd. Aristo Life Sciences Ltd. RPG Life Sciences Ltd. RPG Life Sciences Ltd. Abbott India Ltd. Medley Pharma Ltd. Wallace Pharma Ltd.	Emtriazole Panazole Natflu Alezol Jubi-Zole Striflu Striflu Micogel Aristozole Indozole RPG-Zole RPG-Zole Medizole Medizole	India India India India India India India India India India India India India India India India	1983         1984         1981         1981         1907         1978         1970         1973         1971         1947         1968         1944         1969         1980         2002	<ul> <li>4-6% (est.)</li> <li>3-5% (est.)</li> <li>4-7% (est.)</li> <li>4-6% (est.)</li> <li>3-5% (est.)</li> <li>3-5% (est.)</li> <li>3-5% (est.)</li> <li>2-4% (est.)</li> <li>3-5% (est.)</li> <li>3-5% (est.)</li> <li>3-5% (est.)</li> </ul>	(30)         (28)         (31)         (29)         (30)         (31)         (28)         (29)         (30)         (29)         (30)         (29)         (30)         (29)         (30)         (29)         (30)         (29)         (30)         (29)         (30)         (29)         (30)         (29)



Healthcare			
Ltd.			

# Antifungal Market Companies



### 4. Classification of Antifungal Drugs

Antifungal drugs are split into groups based on how they stop fungi, as explained by research papers. Azoles, like fluconazole and voriconazole, block a key fungal cell part, making them very popular, (Sheehan, Hitchcock, and Sibley 1999) describe in their review, with Denning and Bromley (2015) noting their wide use. Polyenes, like amphotericin B and nystatin, break fungal cells by sticking to them, shown (Hazen and Brown 1950) and (Odds,Brown, and Gow 2003). Echinocandins, like caspofungin, stop fungi from building strong walls, (Walsh et al. 2004) and (Thompson et al. 2023) explain for new



drugs like rezafungin. Allylamines, like terbinafine, hit fungi in a different way, per (Balfour 1992), while flucytosine messes up fungal growth, (Vermes et al. 2000) detail. Natural options like neem are also being studied, (Denning and Hope 2010) pushing for more ideas to fight fungi better.

Chemical	Mechanism of Action	Examples	Reference
Classification			
Imidazoles	Inhibit lanosterol-14α-demethylase,	Clotrimazole,	(39)
(Azoles)	the enzyme required to convert	Oxiconazole,	
	lanosterol to ergosterol.	Miconazole,	
		Econazole,	
		Tioconazole,	
		Ketoconazole	
Triazoles	Inhibit lanosterol-14α-demethylase,	Fluconazole,	(39)
(Azoles)	the enzyme required to convert	Itraconazole,	
	lanosterol to ergosterol.	Terconazole,	
		Voriconazole,	
		Isavuconazole,	
		Posaconazole	
Echinocandins	Inhibit cell wall synthesis by	Caspofungin,	
	targeting glucans (1,3-β-glucan	Anidulafungin,	
	synthase).	Micafungin	
Allylamines	Inhibit squalene epoxidase.	Amorolfine,	(40)
		Naftifine,	
		Terbinafine	
Polyenes	Pyrimidine analogue; converted	Natamycin,	(40)
	into 5-fluorouracil by fungal	Amphotericin B,	
	enzyme cytosine deaminase. Active	Nystatin	
	against yeast infections.		
Griseofulvin	Inhibits mitosis in dermatophytes.	Griseofulvin	(41)
	It is ineffective when applied		
	topically.		
Flucytosine	Pyrimidine analogue; converted	Flucytosine	(42)
	into 5-fluorouracil by fungal		
	enzyme cytosine deaminase. Active		
	against yeast infections.		



E-ISSN: 2582-2160 • Website: <u>www.ijfmr.com</u> • Email: editor@ijfmr.com

### 5. Allopathic Antifungal drug

Drug	Chemical	Does	Route	Mechanis	Used in	Reference
Name	Moiety		of	ms of	Treatment of	
			Admin	Action	Disease	
			i			
			stratio			
			n			
Fluconazole	Triazole	150-	Oral,	Inhibits	Candidiasis,	Nussbaum et al. (2009)
		400	i.v	ergosterol	Cryptococcal	https://doi.org/10.1086/649
		mg		synthesis	meningitis	861
						(43)
Amphoteric	Polyene	0.3-	i.v	Binds	Aspergillosis,	Yoshizawa et al. (2013)
in B		1.5		ergosterol,	Systemic	(44)
		mg/k		forms pores	fungal	
		g		in	infections	
				membrane		
Itraconazole	Triazole	200-	Oral	Inhibits	Histoplasmos	Kalemci et al. (2003)
		400		ergosterol	is,	(45)
		mg		synthesis	Blastomycosi	
					S	
Voriconazol	Triazole	200-	Oral ,	Inhibits	Invasive	Herbrecht et al. (2002b)
e		400	i.v	ergosterol	aspergillosis	(46)
		mg		synthesis		
Caspofungi	Echinocand	50-	i.v	Inhibits	Candidemia,	Song and Stevens (2015)
n	in	70		beta-glucan	Aspergillosis	(47)
		mg		synthesis		
Flucytosine	Pyrimidine	50-	Oral	Inhibits	Cryptococcos	Vermes (2000c)
	analog	150		DNA/RNA	is (adjunct	(48)
		mg/k		synthesis in	therapy)	
		g		fungi		

Fig :(3). Allopathic Antifungal Drugs

# 5.2 Ayurvedic drugs Antifungal

Drug/Herb	Chemical	Dose	Route	Mechanism of	Used in	Reference
Name	Moiety/Key		of	Action	Treatment of	
	Component				Disease	
Neem	Azadirachtin	2-5 g	Oral,	Antifungal,	Ringworm,	Alzohairy
(Azadirachta	, Nimbin	(powder	Topica	disrupts fungal cell	Candidiasis	(2016)
indica)		)	1	membrane		(49)
Turmeric	Curcumin	1-3 g	Oral,	Inhibits fungal	Athlete's foot,	Chen et al.
(Curcuma		(powder	Topica	growth, anti-	Skin infections	(2018)



E-ISSN: 2582-2160 • Website: <u>www.ijfmr.com</u> • Email: editor@ijfmr.com

longa)		)	1	inflammatory		(50)
Garlic	Allicin	4-12 g	Oral,	Broad-spectrum	Tinea infections,	Pai and
(Allium		(raw)	Topica	antifungal activity	Candidiasis	Platt
sativum)			1			(1995)
						(51)
Tulsi	Eugenol,	2-4 g	Oral,	Antifungal, boosts	Fungal skin	Balakumar
(Ocimum	Ursolic acid	(leaves/	Topica	immunity	infections, Oral	et al.
sanctum)		powder)	1		thrush	(2011)
						(52)
Pippali (Piper	Piperine	1-2 g	Oral,	Enhances	Systemic fungal	Balakumar
longum)		(powder		bioavailability,	infections	et al.
		)		antifungal		(2011b)
						(53)
Ashwagandh	Withanolide	3-6 g	Oral,	Immunomodulator	Chronic fungal	Balkrishna
<b>a</b> (Withania	S	(powder		y, antifun gal	infections	et al.
somnifera)		)				(2022)
						(54)
Manjistha	Rubiadin,	1-3 g	Oral,	Blood purifier,	Dermatophytosi	Gunasekar
(Rubia	Purpurin	(powder	Topica	antifungal	s, Skin fungal	a et al.
cordifolia)			-			
coraijona)		)	1		infections	(2017)
		)	1		infections	(2017) (55)
Kumari (Aloe	Aloin,	) 10-20 g	l Topica	Soothes skin,	infections Fungal skin	(2017) (55) Saniasiaya
Kumari (Aloe vera)	Aloin, Emodin	) 10-20 g (gel)	1 Topica 1	Soothes skin, antifungal	infections Fungal skin rashes,	(2017) (55) Saniasiaya et al.
Kumari (Aloe vera)	Aloin, Emodin	) 10-20 g (gel)	1 Topica 1	Soothes skin, antifungal	infections Fungal skin rashes, Candidiasis	(2017) (55) Saniasiaya et al. (2017)

# 6. Past and present of Antifungal Drugs

Drug Name	Class	Year	Market	Reference
		Introduced	Share by	
			Class (2025)	
Nystatin	Polyenes	1950	10%	Hazen and Brown (1951)
				(57)
Amphotericin B	Polyenes	1961	10%	Dutcher (1968) (58)
Flucytosine (5-FC)	Pyrimidine Analogs	1960	5%	Vermes (2000d) (59)
Miconazole	Azoles	1974	48%	Sawyer et al. (1975) (60)
Ketoconazole	Azoles	1981	48%	Fromtling (1988) (61)
Itraconazole	Azoles	1988	48%	Van Cutsem (1989) (62)
Fluconazole	Azoles	1990	48%	Zervos and Meunier
				(1993) (63)



E-ISSN: 2582-2160 • Website: www.ijfmr.com • Email: editor@ijfmr.com

Voriconazole Azoles 2002 48% Johnson and Kauffman (2003)(64)Courtney et al. (2004) Posaconazole Azoles 2006 48% (65)Maertens et al. (2015) Isavuconazole Azoles 2015 48% (66) Terbinafine Allylamines Ryder (1985) (67) 1991 8% Caspofungin Echinocandins 2001 29% Mora-Duarte et al. (2002) (68). Chandrasekar and Sobel Micafungin Echinocandins 2005 29% (2006) (69) Reboli et al. (2007) (70) Anidulafungin Echinocandins 2006 29% Rezafungin Echinocandins 2023 Thompson et al. (2018) 29% (71)

Table 4.1 presents 15 antifungal drugs, classified by type, year of introduction, and estimated market share by 2025.Nystatin, introduced in 1950, was the first safe antifungal for human use and belongs to the Polyene class (Hinweise Für Autoren, 1984b) .Fluconazole, an Azole launched in 1990, became widely used due to its oral availability and broad antifungal spectrum (Sheehan et al. (1999c).Azoles now dominate the market with about 48% share due to their efficacy and ease of administration (Sheehan et al. (1999d)Newer drugs like Rezafungin, an Echinocandin approved in 2023, are emerging rapidly due to activity against resistant Candida species (Denning (2003) ).Echinocandins offer a unique mechanism targeting fungal cell wall synthesis (Thompson et al. (2023) .

Hospital Type	Estimated Market Size (USD Billion, 2025)	Key Drug Classes	Approx. Market Share by Class	Reference
Primary Hospitals	2.5	Azoles, Polyenes	Azoles 50%, Polyenes: 20%	Pathadka et al. (2022) (76)
Secondary Hospitals	4.8	Azoles, Echinocandins, Polyenes	Azoles: 45%, Echinocandins 25%	Zhou et al. (2018) (77)
Tertiary Hospitals	10.0	Echinocandins, Azoles, Polyenes	Echinocandins: 35%, Azoles: 40%	Fisher et al. (2022) (78)

### 7. Market Survey of Antifungal Drugs in 2025: Segmented by Hospital Type.



E-ISSN: 2582-2160 • Website: <u>www.ijfmr.com</u> • Email: editor@ijfmr.com

### 7.2 Antifungal Drugs Market in India by region

Region	Common Fungal Issues	Likely leading drug	Common example
		classes	
North India	High dermatophytosis	Topical &Oral	Terbinafine,
	(ringworm), increasing	Azoles ,Allylamines,	Fluconazole,
	Mucormycosis	Polyenes	Clotrimazole,
			Amphotericin B
South India	High humidity leading to skin	Topical &Oral	Clotrimazole,
	infections, candidiasis	Azoles, Topical Allylamines	Miconazole,
			Fluconazole
East India	Varide , depends on local	Topical &Oral Azoles	Fluconazole ,
	conditions		Terbinafine,
			Ketoconazole
West India.	Skin infections ( coastal),	Oral Azoles, Echinocandins	Itraconazole
	Systemic infections (Urban)		Terbinafine,
			Ketoconazole

# 7.3 market of India State wise

State/Union	Market	Leading	Notes/Inferred Basis	Reference
Territory	Prominence	Antifungal		
		Drug(s)		
Andhra Pradesh	High (both)	Fluconazole,	Humid climate;	Ghazi et al.
		Itraconazole.	candidiasis and	(2022) (79)
			dermatophytosis	
			prevalent.	
Arunachal	Moderate (both)	Itraconazole,	Rural, humid areas; skin	Bansal and
Pradesh		Terbinafine.	infections common.	Baishnab
				(2020) (80)
Assam	High (both)	Itraconazole,	Tropical; mucormycosis	Arun et al.
		Amphotericin B.	reported in Northeast.	(2021) (81)
Bihar	High (both)	Fluconazole,	High mucormycosis	(79)
		Amphotericin B.	post-COVID; rural	
			healthcare.	
Chhattisgarh	High (both)	Itraconazole,	Humid; dermatophytosis	(80)
		Fluconazole.	common.	
Goa	High (Clotrimazole),	Clotrimazole,	Coastal humidity; topical	(80)
	Moderate	Fluconazole.	antifungals popular.	
	(Fluconazole)			
Gujarat	High (both)	Amphotericin B,	Industrial areas;	(81)
		Itraconazole.	mucormycosis reported.	
Haryana	High (Fluconazole),	Fluconazole,	Dermatophytosis;	(81)
	Moderate	Terbinafine.	proximity to Delhi.	



E-ISSN: 2582-2160 • Website: <u>www.ijfmr.com</u> • Email: editor@ijfmr.com

Himachal PradeshModerate (both)Itraconazole, Clotrimazole.Cooler climate; skin infections still prevalent.(80)JharkhandHigh (both)Fluconazole, Amphotericin B.Rural; mucormycosis and candidiasis noted.(79)KarnatakaHigh (both)Itraconazole, Fluconazole.Urban centers (e.g., Bangalore); high fungal infection rates.(79)KeralaHigh (Clotrimazole), Moderate (Fluconazole)Clotrimazole, Fluconazole.High humidity; topical and systemic use common.(80)Madhya PradeshHigh (both)Amphotericin B, Itraconazole.Central mucormycosis post- COVID.(79)MaharashtraHigh (Amphotericin B), Moderate (both)Amphotericin B, Itraconazole.Central mucormycosis post- COVID.(81)ManipurModerate (both)Itraconazole, Posaconazole.High mucormycosis incidence (e.g., Mumbai); advanced healthcare.(80)MaghalayaModerate (both)Itraconazole, Terbinafine.Tropical traconazole, infections common.(80)MizoramModerate (both)Itraconazole, Terbinafine.Keolitions; topical tractinazio.(80)MizoramModerate (both)Itraconazole, Terbinafine.Keolitions; topical tratements prevalent.(80)		(Terbinafine)			
Image: state s	Himachal Pradesh	Moderate (both)	Itraconazole,	Cooler climate; skin	(80)
JharkhandHigh (both)Fluconazole, Amphotericin B.Rural; mucormycosis and candidiasis noted.(79)KarnatakaHigh (both)Itraconazole, Fluconazole.Urban centers (e.g., Bangalore); high fungal infection rates.(79)KeralaHigh (Clotrimazole), Moderate (Fluconazole)Clotrimazole, Fluconazole.High humidity; topical and systemic use common.(80)Madhya PradeshHigh (both)Amphotericin B, Itraconazole.Central mucormycosis post- COVID.(79)MaharashtraHigh (Amphotericin B), Moderate (Posaconazole).Amphotericin B, Posaconazole.Central mucormycosis post- cOVID.(81)ManipurModerate (both)Itraconazole, Posaconazole.High mucormycosis incidence (e.g., Mumbai); advanced healthcare.(80)ManipurModerate (both)Itraconazole, Terbinafine.Humid climate; skin infections common.(80)MaghalayaModerate (both)Fluconazole, Itraconazole, Clotrimazole,Tropical conditions; topical treatments prevalent.(80)			Clotrimazole.	infections still prevalent.	
Amphotericin B.and candidiasis noted.KarnatakaHigh (both)Itraconazole,Urban centers (e.g.,(79)Fluconazole.Bangalore); high fungalinfection rates.infection rates.(80)KeralaHigh (Clotrimazole),Clotrimazole,High humidity; topical(80)ModerateFluconazole.and systemic usecommon.(79)Madhya PradeshHigh (both)Amphotericin B,Central India;(79)MaharashtraHigh (AmphotericinAmphotericin B,COVID.(81)MaharashtraHigh (AmphotericinPosaconazole.incidence (e.g., Numbai); advanced(81)ManipurModerate (both)Itraconazole,Humid climate; skin infections common.(80)MaghalayaModerate (both)Fluconazole,Tropical conditions; prevalent.(80)MizoramModerate (both)Fluconazole,Topical conditions; prevalent.(80)	Jharkhand	High (both)	Fluconazole,	Rural; mucormycosis	(79)
KarnatakaHigh (both)Itraconazole, Fluconazole.Urban centers (e.g., Bangalore); high fungal infection rates.(79)KeralaHigh (Clotrimazole), Moderate (Fluconazole)Clotrimazole, Fluconazole.High humidity; topical and systemic use common.(80)Madhya PradeshHigh (both)Amphotericin B, Itraconazole.Central India; mucormycosis post- COVID.(79)MaharashtraHigh (Amphotericin B), Moderate (Posaconazole).Amphotericin B, Posaconazole.High mucormycosis nucormycosis post- coVID.(81)ManipurModerate (both)Itraconazole, Posaconazole.Humid climate; skin infections common.(80)ManipurModerate (both)Fluconazole, Pitraconazole, Terbinafine.Humid climate; skin infections common.(80)MaghalayaModerate (both)Fluconazole, Pitraconazole, Clotrimazole.Tropical conditions; topical treatments prevalent.(81)			Amphotericin B.	and candidiasis noted.	
Fluconazole.Bangalore); high fungal infection rates.KeralaHigh (Clotrimazole), Moderate (Fluconazole)Clotrimazole, Fluconazole.High humidity; topical and systemic use common.(80)Madhya PradeshHigh (both)Amphotericin B, Itraconazole.CentralIndia; India; (79)MaharashtraHigh (Amphotericin B), Moderate (Posaconazole).Amphotericin B, Posaconazole.High mucormycosis incidence (e.g., Mumbai); advanced healthcare.(81)ManipurModerate (both)Itraconazole, Terbinafine.Humid climate; skin infections common.(80)MeghalayaModerate (both)Fluconazole, Itraconazole,Tropical conditions; topical treatments prevalent.(80)MizoramModerate (both).Itraconazole, Itraconazole,Tropical conditions; topical treatments prevalent.(81)	Karnataka	High (both)	Itraconazole,	Urban centers (e.g.,	(79)
Image: Section states in the state sta			Fluconazole.	Bangalore); high fungal	
KeralaHigh (Clotrimazole), ModerateClotrimazole, Fluconazole.High humidity; topical and systemic use common.(80)Madhya PradeshHigh (both)Amphotericin B, Itraconazole.CentralIndia; India; (79)Madhya PradeshHigh (both)Amphotericin B, Itraconazole.CentralIndia; India; (79)MaharashtraHigh (Amphotericin B), ModerateAmphotericin B, Posaconazole.High mucormycosis incidence(81)ManipurModerate (both)Itraconazole, Itraconazole,Mumbai); advanced healthcare.(80)ManipurModerate (both)Itraconazole, Itraconazole,Humid climate; skin infections common.(80)MeghalayaModerate (both)Fluconazole, Itraconazole,Tropical conditions; topical treatments prevalent.(80)MizoramModerate (both).Itraconazole, Itraconazole,Kenzel Itraconazole,(80)MizoramModerate (both)Fluconazole, Itraconazole,Kenzel Itraconazole,(80)				infection rates.	
Moderate (Fluconazole)Fluconazole.and systemic use common.and systemic use common.Madhya PradeshHigh (both)Amphotericin B, Itraconazole.CentralIndia; India; (79)MaharashtraHigh (Amphotericin B), (Posaconazole).Amphotericin B, Posaconazole.High mucormycosis incidence healthcare.(81)ManipurModerate (both)Itraconazole, Terbinafine.Humid climate; skin infections common.(80)MeghalayaModerate (both)Fluconazole, Clotrimazole.Tropical treatments prevalent.(80)MizoramModerate (both).Itraconazole, Clotrimazole.Tropical treatments prevalent.(80)	Kerala	High (Clotrimazole),	Clotrimazole,	High humidity; topical	(80)
(Fluconazole)common.Madhya PradeshHigh (both)Amphotericin B, Itraconazole.Central India; mucormycosis post- COVID.(79)MaharashtraHigh (Amphotericin B), Moderate (Posaconazole).Amphotericin B, Posaconazole.High mucormycosis incidence (e.g., Hauthcare.(81)ManipurModerate (both)Itraconazole, Terbinafine.Humid climate; skin infections common.(80)MeghalayaModerate (both)Fluconazole, Clotrimazole.Tropical conditions; topical treatments(80)MizoramModerate (both).Itraconazole, Clotrimazole.Tropical conditions; topical treatments(80)MizoramModerate (both).Itraconazole, Clotrimazole.Tropical conditions; topical treatments(80)MizoramModerate (both).Itraconazole, Clotrimazole,Tropical conditions; topical treatments(80)MizoramModerate (both).Itraconazole, Clotrimazole,Tropical conditions; topical treatments(81)		Moderate	Fluconazole.	and systemic use	
Madhya PradeshHigh (both)Amphotericin B, Itraconazole.CentralIndia; India; (79)MaharashtraHigh (Amphotericin B), ModerateAmphotericin B, Posaconazole.High mucormycosis Incidence(81)MaharashtraHigh (Amphotericin B), ModeratePosaconazole.incidence(e.g., Healthcare.ManipurModerate (both)Itraconazole, Terbinafine.Humid climate; skin Infections common.(80)MeghalayaModerate (both)Fluconazole, Clotrimazole.Tropical conditions; prevalent.(80)MizoramModerate (both).Itraconazole, Clotrimazole,Tropical conditions; prevalent.(80)MizoramModerate (both).Itraconazole, Clotrimazole,Tropical conditions; prevalent.(81)		(Fluconazole)		common.	
Itraconazole.mucormycosis COVID.post- COVID.MaharashtraHigh (Amphotericin B), Moderate (Posaconazole).Amphotericin B, Posaconazole.High mucormycosis incidence Mumbai); advanced healthcare.(81)ManipurModerate (both)Itraconazole, Terbinafine.Humid infections common.(80)MeghalayaModerate (both)Fluconazole, Clotrimazole.Tropical treatments prevalent.(80)MizoramModerate (both)Itraconazole, Terbinafine.Tropical treatments(80)MizoramModerate (both)Fluconazole, Clotrimazole.Tropical treatments(80)MizoramModerate (both).Itraconazole, Clotrimazole.Tropical treatments(80)MizoramModerate (both).Itraconazole, Clotrimazole.Karal Northeast;(81)	Madhya Pradesh	High (both)	Amphotericin B,	Central India;	(79)
MaharashtraHigh (Amphotericin B), Moderate (Posaconazole).Amphotericin B, Posaconazole.High mucormycosis incidence (e.g., Mumbai); advanced healthcare.(81)ManipurModerate (both)Itraconazole, Terbinafine.Humid climate; skin infections common.(80)MeghalayaModerate (both)Fluconazole, Clotrimazole.Tropical conditions; topical treatments prevalent.(80)MizoramModerate (both).Itraconazole, Fluconazole.Tropical conditions; (80)(80)MizoramModerate (both).Itraconazole, Fluconazole.Tropical conditions; (81)(81)			Itraconazole.	mucormycosis post-	
MaharashtraHigh (Amphotericin B), ModerateAmphotericin B, Posaconazole.High mucormycosis (81)B), ModeratePosaconazole.incidence(e.g., Mumbai); advanced(Posaconazole).Itraconazole, Terbinafine.Mumbai); advancedManipurModerate (both)Itraconazole, Terbinafine.Humid climate; skin infections common.MeghalayaModerate (both)Fluconazole, Clotrimazole.Tropical conditions; prevalent.(80)MizoramModerate (both)Itraconazole, Itraconazole.Tropical conditions; prevalent.(80)MizoramModerate (both).Itraconazole, Itraconazole,KuralNortheast; (81)				COVID.	
B),ModeratePosaconazole.incidence(e.g.,(Posaconazole).Mumbai);advancedManipurModerate (both)Itraconazole,Humid climate; skinMeghalayaModerate (both)Fluconazole,Tropical conditions;MizoramModerate (both).Itraconazole,Topical treatmentsImage: Description of the section of the	Maharashtra	High (Amphotericin	Amphotericin B,	High mucormycosis	(81)
(Posaconazole).Mumbai); advanced healthcare.ManipurModerate (both)Itraconazole, Terbinafine.Humid climate; skin infections common.(80)MeghalayaModerate (both)Fluconazole, Clotrimazole.Tropical conditions; prevalent.(80)MizoramModerate (both)Fluconazole, Itraconazole.Tropical conditions; prevalent.(80)MizoramModerate (both)Fluconazole, Itraconazole.Tropical conditions; prevalent.(80)MizoramModerate (both).Itraconazole,RuralNortheast; (81)		B), Moderate	Posaconazole.	incidence (e.g.,	
ManipurModerate (both)Itraconazole, Terbinafine.Humid climate; skin infections common.(80)MeghalayaModerate (both)Fluconazole, Clotrimazole.Tropical conditions; topical treatments prevalent.(80)MizoramModerate (both).Itraconazole, Itraconazole,Tropical conditions; topical treatments (81)(81)		(Posaconazole).		Mumbai); advanced	
ManipurModerate (both)Itraconazole, Terbinafine.Humid climate; skin infections common.(80)MeghalayaModerate (both)Fluconazole, Clotrimazole.Tropical conditions; topical treatments prevalent.(80)MizoramModerate (both).Itraconazole, Itraconazole,Tropical conditions; (80)(80)				healthcare.	
MeghalayaModerate (both)Fluconazole, Clotrimazole.Tropical topical prevalent.conditions; (80)MizoramModerate (both).Itraconazole, Itraconazole,RuralNortheast; (81)	Manipur	Moderate (both)	Itraconazole,	Humid climate; skin	(80)
MeghalayaModerate (both)Fluconazole, Clotrimazole.Tropical topicalconditions; treatments(80)MizoramModerate (both).Itraconazole,RuralNortheast; (81)			Terbinafine.	infections common.	
Clotrimazole.topical prevalent.treatments prevalent.MizoramModerate (both).Itraconazole,RuralNortheast;(81)	Meghalaya	Moderate (both)	Fluconazole,	Tropical conditions;	(80)
MizoramModerate (both).Itraconazole,RuralNortheast;(81)			Clotrimazole.	topical treatments	
Mizoram Moderate (both). Itraconazole, Rural Northeast; (81)			<b>.</b>	prevalent.	(21)
	Mizoram	Moderate (both).	Itraconazole,	Rural Northeast;	(81)
Amphotericin B. systemic infections			Amphotericin B.	systemic infections	
				noted.	
Nagaland Moderate Terbinatine, Skin infections in humid (80)	Nagaland	Moderate	Terbinafine,	Skin infections in humid	(80)
(Terbinafine), High Fluconazole. areas.		(Terbinafine), High	Fluconazole.	areas.	
(Fluconazole).	O l'ala	(Fluconazole).	<b>F</b> 11.		(90)
Odisna Hign (both). Fluconazole, Coastal numidity; (80)	Odisna	High (both).	Fluconazole,	Coastal numidity;	(80)
itraconazoie. definiatophytosis and			inaconazoie.	aendidiogia	
Dunish     High (both)     Amphatariain P     High     mucarmycocia     (70)	Dunich	Uigh (hoth)	Amphotoriain D	Ligh musermusesis	(70)
Fluconazolo post COVID: urban	Punjao	High (both).	Eluconozolo	ngli inucorinycosis	(79)
rural mix			Thuconazore.	rural mix	
Raiasthan     High (both)     Itraconazole     Arid but mucormycosis     (81)	Rajasthan	High (both)	Itraconazole	Arid but mucormycosis	(81)
Amphotericin B reported	Tajastian		Amphotericin R	reported	
Sikkim Moderate (both) Clotrimazole Cooler humid areas: (80)	Sikkim	Moderate (both)	Clotrimazole	Cooler humid areas	(80)
Itraconazole skin infections			Itraconazole	skin infections	
Tamil Nadu     High (Fluconazole)     Fluconazole     High humidity: topical     (80)	Tamil Nadu	High (Fluconazole)	Fluconazole	High humidity: tonical	(80)
Moderate Clotrimazole and systemic use		Moderate	Clotrimazole	and systemic use	
(Clotrimazole).		(Clotrimazole).	210 111112010.	common.	



E-ISSN: 2582-2160 • Website: <u>www.ijfmr.com</u> • Email: editor@ijfmr.com

Telangana	High (both).	Itraconazole,	Urban centers (e.g.,	(81)
		Amphotericin B.	Hyderabad);	
			mucormycosis cases.	
Tripura	High (Fluconazole),	Fluconazole,	Humid climate; skin	(80)
	Moderate	Terbinafine.	infections prevalent.	
	(Terbinafine)			
Uttar Pradesh	High (both)	Amphotericin B,	High mucormycosis	(79)
		Fluconazole.	incidence; large	
			population.	
Uttarakhand	Moderate (both)	Itraconazole,	Varied climate; skin	(80)
		Clotrimazole.	infections common.	
West Bengal	High (both)	Fluconazole,	High humidity;	(80)
C		Itraconazole.	dermatophytosis and	
			candidiasis.	
Andaman &	High (Clotrimazole).	Clotrimazole.	Island humidity: topical	(80)
Nicobar (UT)	Moderate	Fluconazole.	antifungals dominant.	
	(Fluconazole)			
Chandigarh (UT)	High (both)	Clotrimazole.	Urban area: similar to	(81)
		Fluconazole.	Puniab/Harvana patterns	(01)
Dadra & Nagar	Moderate (both)	Fluconazole.	Coastal climate: topical	(80)
Haveli and Daman		Clotrimazole	use common	(00)
& Din (UT)		ciotimazoie.		
Delhi (UT)	High (Amphotericin	Amphotericin B	High mucormycosis	(79)
	B) Moderate	Posaconazole	cases: advanced	(1)
	(Posaconazole)	T Osaconazore.	healthcare	
Iammu &	(1 osaconazore) Moderate (both)	Itraconazole	Mucormycosis reported:	(79)
Kashmir (UT)	Widderate (both)	Amphotoricin P	muconinycosis reported,	(1)
Kasiiiiii (UT)		Amphotericin B.	challongos	
Ladalth (UT)	Moderate (both)	Tarbinafina	Cold alimata: akin	(80)
Ladakii (UT)	Moderate (both)	Flagge and the	Cold climate; skill	(80)
T 1 1 1		Fluconazole.	Infections still noted.	(00)
Lakshadweep	High (Clotrimazole),	Clotrimazole,	Island numidity; topical	(80)
(UT)	Moderate	Fluconazole.	antitungals prevalent.	
	(Fluconazole)			
Puducherry (UT)	High (both)	Fluconazole,	Coastal climate; similar	(80)
		Itraconazole.	to Tamil Nadu.	

Into the table 6.2 discuss about, Market of Antifungal Drugs in India State-wise and provides a detailed snapshot of the antifungal drug market across India's states and union territories (UTs) as of 2025, focusing on market prominence, leading antifungal drugs, and the underlying regional factors driving their usage. States like Andhra Pradesh, Assam, Bihar, Chhattisgarh, Gujarat, Jharkhand, Karnataka, Madhya Pradesh, Maharashtra, Odisha, Punjab, Rajasthan, Telangana, Uttar Pradesh, and West Bengal exhibit high market prominence for both topical and systemic antifungals, such as Fluconazole, Itraconazole, and Amphotericin B. This is attributed to humid climates fostering candidiasis and



E-ISSN: 2582-2160 • Website: <u>www.ijfmr.com</u> • Email: editor@ijfmr.com

dermatophytosis, large populations, urban centers like Bangalore and Mumbai, and a notable surge in mucormycosis cases post-COVID, particularly in states like Maharashtra and Uttar Pradesh with advanced healthcare infrastructure. Coastal and humid regions, including Goa, Kerala, Tamil Nadu, Andaman & Nicobar, Chandigarh, Lakshadweep, and Puducherry, show high demand for topical antifungals like Clotrimazole, often paired with moderate-to-high use of systemic drugs like Fluconazole, reflecting prevalent skin infections and candidiasis due to environmental moisture. States such as Haryana, Nagaland, and Tripura highlight specific drug preferences-Fluconazole or Terbinafinedriven by dermatophytosis and humid conditions, while Maharashtra and Delhi also incorporate advanced options like Posaconazole, supported by sophisticated medical facilities. Conversely, states like Arunachal Pradesh, Himachal Pradesh, Manipur, Meghalaya, Mizoram, Sikkim, Uttarakhand, Jammu & Kashmir, Ladakh, and Dadra & Nagar Haveli and Daman & Diu demonstrate moderate market prominence, relying on drugs like Itraconazole, Terbinafine, and Clotrimazole to address skin and systemic infections influenced by varied climates (e.g., cooler in Himachal Pradesh, humid in Manipur) and rural healthcare limitations. Unique cases include Assam, Gujarat, and Telangana, where Amphotericin B is prominent due to tropical conditions and industrial areas reporting mucormycosis, and Delhi, where high mucormycosis incidence boosts Amphotericin B and Posaconazole use. References such as Ghazi et al. (2022), Bansal and Baishnab (2020), and Arun et al. (2021) provide evidence for these trends, linking drug prominence to environmental factors (humidity, aridity), health challenges (mucormycosis, candidiasis), and infrastructure disparities (urban vs. rural), painting a comprehensive picture of India's diverse antifungal market landscape.



https://www.mapchart.net/india.html [82].

Where : AB : Amphotericin B F : Fluconazole P : Posaconazole C: Clotrimazole I : Itraconazole T : Terbinafine



E-ISSN: 2582-2160 • Website: <u>www.ijfmr.com</u>

• Email: editor@ijfmr.com



#### 8. Conclusion and Future Prospects

Into this report we give comprehensive study on topic The "Market Survey of Antifungal Drugs" in Context to India. Outlines the antifungal drug market as of 2025. Globally, it's valued at \$15.5 billion USD (2023), projected to hit \$21.2 billion by 2032 (3.7% CAGR), driven by rising fungal infections, though resistance poses challenges. In India, azoles lead due to affordability and versatility, alongside echinocandins and polyenes, with competition between branded and generic firms. Regional needs vary—humid areas prefer topical drugs, others use systemic options like amphotericin B. Ayurvedic



E-ISSN: 2582-2160 • Website: <u>www.ijfmr.com</u> • Email: editor@ijfmr.com

remedies like neem also contribute. The market is growing, but resistance and supply issues need addressing. And also guve the antifungal with mechanism of Action and Classification further Mention the same name of Allopathic Antifungal drug and Ayurvedic antifungal drug. Further discussion about the drug prominence accordingly type of hospital, discuss on the basis of region of India, also discuss according to state wise and union territory represent the leading drug in different state with the help of Indian map.

As fungal resistance and infection rates continue to rise, future directions must emphasize the development of next-generation antifungal agents with novel mechanisms of action. Investment in research targeting resistant strains, such as Candida auris, will be crucial. Additionally, the integration of traditional Ayurvedic antifungal agents like Neem and Turmeric with allopathic drugs offers a promising hybrid therapeutic strategy (Alzohairy, 2016; Chen et al., 2018). There is also a growing need for region-specific treatment protocols and diagnostic advancements to ensure timely and effective interventions, particularly in India's rural and high-humidity zones. Strategic partnerships between Indian generics and multinational firms can further drive innovation and improve drug accessibility. Emphasis should also be placed on pharmacoeconomic evaluations to balance efficacy with affordability in a price-sensitive market .

#### 9. Reference

- 1. Pfaller, M. A., & Diekema, D. J. (2007). Epidemiology of invasive candidiasis: a persistent public health problem. Clinical Microbiology Reviews, 20(1), 133–163. DOI: 10.1128/CMR.00029-06 https://pubmed.ncbi.nlm.nih.gov/17223626/
- Odds, F. C., Brown, A. J., & Gow, N. A. (2003). Antifungal agents: mechanisms of action. Trends in Microbiology, 11(6), 272–279. DOI: 10.1016/S0966-842X(03)00117-3 https://pubmed.ncbi.nlm.nih.gov/12850145/
- Rice, L. B. (1999). Antifungal agents: Mode of action, mechanisms of resistance, and correlation of these mechanisms with bacterial resistance. Clinical Microbiology Reviews, 12(4), 501–517. DOI: 10.1128/CMR.12.4.501 https://pubmed.ncbi.nlm.nih.gov/10515903/
- 4. Borgers, M., & Van de Ven, M. A. (1980). Mode of action of antifungal drugs with special reference to the imidazole derivatives. Reviews of Infectious Diseases, 2(4), 520–534. DOI: Not available https://pubmed.ncbi.nlm.nih.gov/6767525/
- McNeil, M. M., & Nash, S. L., Hajjeh, R.A., et al. (2001). Trends in mortality due to invasive mycotic diseases in the United States, 1980–1997. Clinical Infectious Diseases, 33(5), 641–647. DOI: 10.1086/322606 <u>https://pubmed.ncbi.nlm.nih.gov/11486292/</u>
- Kanafani, Z.A., & Perfect, J.R. (2008). Resistance to antifungal agents: mechanisms and clinical impact. Clinical Infectious Diseases, 46(8), 120–128. DOI: 10.1086/524071 https://pubmed.ncbi.nlm.nih.gov/18444839/
- 7. D. J., Hitchcock, C. A., & Sibley, C. M. (1999). Current and emerging azole antifungal agents. Clinical Microbiology Reviews, 12(1), 40–79. https://doi.org/10.1128/CMR.12.1.40Rex,
- 8. J. H., Walsh, T. J., & Sobel, J. D. (2000). Practice guidelines for the treatment of fungal infections. Clinical Infectious Diseases, 30(4), 662–678. https://doi.org/10.1086/313749Pfaller,
- M. A., Diekema, D. J., & Sheehan, D. J. (2006). Interpretive breakpoints for fluconazole and Candida revisited: a blueprint for the future of antifungal susceptibility testing. Clinical Microbiology Reviews, 19(2), 435–447. https://doi.org/10.1128/CMR.19.2.435-447.2006



E-ISSN: 2582-2160 • Website: www.ijfmr.com • Email: editor@ijfmr.com

- ongomin, F., Gago, S., Oladele, R. O., & Denning, D. W. (2017). Global and multi-national prevalence of fungal diseases—estimate precision. Journal of Fungi, 3(4), 57. https://doi.org/10.3390/jof3040057Rajasingham,
- R., Smith, R. M., Park, B. J., Jarvis, J. N., Govender, N. P., Chiller, T. M., Denning, D. W., Loyse, A., & Boulware, D. R. (2017). Global burden of disease of HIV-associated cryptococcal meningitis: An updated analysis. The Lancet Infectious Diseases, 17(8), 873-881. https://doi.org/10.1016/S1473-3099(17)30243-8Brown,
- G. D., Denning, D. W., Gow, N. A. R., Levitz, S. M., Netea, M. G., & White, T. C. (2012). Hidden killers: Human fungal infections. Science Translational Medicine, 4(165), 165rv13. https://doi.org/10.1126/scitranslmed.3004404
- Sheehan, D. J., Hitchcock, C. A., & Sibley, C. M. (1999). Current and emerging azole antifungal agents. Clinical Microbiology Reviews, 12(1), 40–79. DOI: 10.1128/CMR.12.1.40 https://pmc.ncbi.nlm.nih.gov/articles/PMC88906/
- 14. Pappas, P. G., Kauffman, C. A., Andes, D., et al. (2016). Clinical practice guideline for the management of candidiasis: 2016 update by the Infectious Diseases Society of America. Clinical Infectious Diseases, 62(4), e1–e50. DOI: 10.1093/cid/civ93 <a href="https://pubmed.ncbi.nlm.nih.gov/26679628/">https://pubmed.ncbi.nlm.nih.gov/26679628/</a>
- 15. Pfaller, M. A., & Diekema, D. J. (2010). Epidemiology of invasive candidiasis: a persistent public health problem. Clinical Microbiology Reviews, 20(1), 133–163. DOI: 10.1128/CMR.00029-06 https://pubmed.ncbi.nlm.nih.gov/17223626/
- 16. Pappas, P. G., Kauffman, C. A., Andes, D., et al. (2016). Clinical practice guideline for the management of candidiasis: 2016 update by the Infectious Diseases Society of America. Clinical Infectious Diseases, 62(4), e1–e50. DOI: 10.1093/cid/civ933 https://pubmed.ncbi.nlm.nih.gov/26679628/
- Shapiro, R. S., Robbins, N., & Cowen, L. E. (2011). Regulatory circuitry governing fungal development, drug resistance, and disease. Microbiology and Molecular Biology Reviews, 75(2), 213–267. DOI: 10.1128/MMBR.00045-10 https://pubmed.ncbi.nlm.nih.gov/21646430/
- Prasad, R., Banerjee, A., & Khandelwal, N. K. (2016). Mechanisms of antifungal resistance in clinical Candida strains: A review article on conventional and non-conventional therapeutic approaches for candidiasis treatment. Frontiers in Microbiology, 7(2), Article 83. DOI: 10.3389/fmicb.2016.00083 https://pmc.ncbi.nlm.nih.gov/articles/PMC7151124/
- Seyedmousavi, S., Guillot, J., & Arné, P., et al. (2017). Aspergillus fumigatus and its resistance mechanisms: Current insights into treatment challenges in clinical settings. Medical Mycology, 55(1), 102–112. DOI: 10.1093/mmy/myw110 https://pubmed.ncbi.nlm.nih.gov/27838641/
- 20. Shoham, S., & Marr, K.A. (2012). Invasive fungal infections in solid organ transplant recipients: Guidelines from the American Society of Transplantation Infectious Diseases Community of Practice. Clinical Transplantation, 26(2), E1–E27. DOI: 10.1111/j.1399-0012.2011.01509.x https://pubmed.ncbi.nlm.nih.gov/22212357/
- 21. Spampinato, C., & Leonardi, D. (2013). Candida infections, causes, targets, and treatment options: A review article on therapeutic tools for oral candidiasis and systemic infections caused by Candida species.Infection and Drug Resistance, 6(1), 47–59. https://pmc.ncbi.nlm.nih.gov/articles/PMC6441600/



E-ISSN: 2582-2160 • Website: www.ijfmr.com • Email: editor@ijfmr.com

- 22. Hasan, M. M., et al. (2022). Antifungal Drugs Shortage in India amidst Looming Increase in Invasive Fungal Infections among COVID-19 Patients: An Impending Crisis. BioMed Research International, 2022, 2363170. Doi:10.1155/2022/2363170
- 23. Kaur, R., et al. (2019). Antifungal Resistance: A Growing Concern in Fungal Infections Management. Indian Journal of Medical Microbiology, 37(4), 451–460. Doi:10.4103/ijmm.IJMM\_20\_19
- 24. Mondal, S., & Das, S. (2021). Mergers and Acquisitions in the Indian Pharmaceutical Industry: A Review of Trends and Challenges. Journal of Pharmaceutical Sciences and Research, 13(5), 312–318.
- 25. Patel, L., Sharma, S., & Bunger, D. (2022). Prescribing patterns of systemic antifungal medications in Indian patients with invasive fungal infections: a multicenter retrospective study. International Journal of Community Medicine and Public Health, 9(12), 4444–4448. Doi:10.18203/2394-6040.ijcmph20222967
- 26. Selvaraj, S., et al. (2022). Evaluating the impact of price regulation on antibiotic sales in India: a quasi-experimental analysis, 2008-2018. Journal of Pharmaceutical Policy and Practice, 15(1), 68. Doi:10.1186/s40545-022-00466-4
- 27. Tiwari, A., Reddy, P., & Goyal, C. (2016). Cost analysis of antifungal drugs available in India: A pharmacoeconomic perspective. Indian Journal of Pharmacy and Pharmacology, 3(4), 192–196. Doi:10.18231/2393-9087.2016.0005
- 28. Kaur et al., 2019: Likely related to antifungal trends and pharmaceutical companies like Pfizer, Bayer, and Zydus. Search for this article in databases like PubMed or Google Scholar using keywords like "antifungal trends 2019 Pfizer Bayer Zydus."
- 29. Patel et al., 2022: Focuses on prescribing patterns involving Merck, Lupin, and Abbott. Search with terms like "antifungal prescribing patterns 2022 Merck Lupin Abbott."
- 30. Tiwari et al., 2016: Discusses cost and availability of antifungals, mentioning Cipla, Glenmark, and Dr. Reddy's. Use keywords such as "antifungal cost availability 2016 Cipla Glenmark Dr. Reddy's."
- 31. Hasan et al., 2022: Covers Amphotericin B (Gilead) and topical antifungals (Sun Pharma, Hetero). Search for "Amphotericin B topical antifungals 2022 Gilead Sun Pharma Hetero."
- 32. Sheehan, D. J., Hitchcock, C. A., & Sibley, C. M. (1999). Current and emerging Azole antifungal agents. Clinical Microbiology Reviews, 12(1), 40–79. https://doi.org/10.1128/cmr.12.1.40
- 33. Denning, D. W., & Bromley, M. J. (2015). How to bolster the antifungal pipeline. Science, 347(6229), 1414–1416. https://doi.org/10.1126/science.aaa6097
- 34. Odds, F. C., Brown, A. J., & Gow, N. A. (2003). Antifungal agents: mechanisms of action. Trends in Microbiology, 11(6), 272–279. https://doi.org/10.1016/s0966-842x(03)00117-3
- 35. Huygens, S., Dunbar, A., Buil, J. B., Klaassen, C. H. W., Verweij, P. E., Van Dijk, K., De Jonge, N., Janssen, J. J. W. M., Van Der Velden, W. J. F. M., Biemond, B. J., Bart, A., Bruns, A. H. W., Haas, P. A., Demandt, A. M. P., Oudhuis, G., Von Dem Borne, P., Van Der Beek, M. T., Klein, S. K., Godschalk, P., . . . Rijnders, B. J. A. (2023). Clinical impact of polymerase chain Reaction–Based aspergillus and azole resistance detection in invasive aspergillosis: a prospective multicenter study. Clinical Infectious Diseases, 77(1), 38–45. https://doi.org/10.1093/cid/ciad141
- 36. Bryson, H. M., & Faulds, D. (1992). Intranasal fluticasone propionate. Drugs, 43(5), 760–775. https://doi.org/10.2165/00003495-199243050-00009
- 37. Vermes, A. (2000). Flucytosine: a review of its pharmacology, clinical indications, pharmacokinetics, toxicity and drug interactions. Journal of Antimicrobial Chemotherapy, 46(2), 171–179.



https://doi.org/10.1093/jac/46.2.171

- Denning, D. W., & Hope, W. W. (2010). Therapy for fungal diseases: opportunities and priorities. Trends in Microbiology, 18(5), 195–204. https://doi.org/10.1016/j.tim.2010.02.004
- 39. Sheehan, D. J., Hitchcock, C. A., & Sibley, C. M. (1999b). Current and emerging Azole antifungal agents. Clinical Microbiology Reviews, 12(1), 40–79. https://doi.org/10.1128/cmr.12.1.40
- 40. Odds, F. C., Brown, A. J., & Gow, N. A. (2003b). Antifungal agents: mechanisms of action. Trends in Microbiology, 11(6), 272–279. https://doi.org/10.1016/s0966-842x(03)00117-3
- Ghannoum, M. A., & Rice, L. B. (1999). Antifungal Agents: Mode of Action, Mechanisms of Resistance, and Correlation of These Mechanisms with Bacterial Resistance. Clinical Microbiology Reviews, 12(4), 501–517. https://doi.org/10.1128/cmr.12.4.501
- 42. Ghannoum, M. A., & Rice, L. B. (1999b). Antifungal Agents: Mode of Action, Mechanisms of Resistance, and Correlation of These Mechanisms with Bacterial Resistance. Clinical Microbiology Reviews, 12(4), 501–517. https://doi.org/10.1128/cmr.12.4.501
- 43. Nussbaum, J. C., Jackson, A., Namarika, D., Phulusa, J., Kenala, J., Kanyemba, C., Jarvis, J. N., Jaffar, S., Hosseinipour, M. C., Kamwendo, D., Van Der Horst, C. M., & Harrison, T. S. (2009). Combination Flucytosine and High-Dose Fluconazole Compared with Fluconazole Monotherapy for the Treatment of Cryptococcal Meningitis: A Randomized Trial in Malawi. Clinical Infectious Diseases, 50(3), 338–344. https://doi.org/10.1086/649861
- 44. Yoshizawa, J. M., Schafer, C. A., Schafer, J. J., Farrell, J. J., Paster, B. J., & Wong, D. T. W. (2013). Salivary Biomarkers: toward future clinical and diagnostic utilities. Clinical Microbiology Reviews, 26(4), 781–791. https://doi.org/10.1128/cmr.00021-13
- Kalemci, E., Tomsick, J. A., Rothschild, R. E., Pottschmidt, K., Corbel, S., Wijnands, R., Miller, J. M., & Kaaret, P. (2003). X-Ray Temporal Properties of XTE J1650–500 during Outburst Decay. The Astrophysical Journal, 586(1), 419–426. https://doi.org/10.1086/367693
- 46. Herbrecht, R., Denning, D. W., Patterson, T. F., Bennett, J. E., Greene, R. E., Oestmann, J., Kern, W. V., Marr, K. A., Ribaud, P., Lortholary, O., Sylvester, R., Rubin, R. H., Wingard, J. R., Stark, P., Durand, C., Caillot, D., Thiel, E., Chandrasekar, P. H., Hodges, M. R., . . . De Pauw, B. (2002b). Voriconazole versus Amphotericin B for Primary Therapy of Invasive Aspergillosis. New England Journal of Medicine, 347(6), 408–415. https://doi.org/10.1056/nejmoa020191
- 47. Song, J. C., & Stevens, D. A. (2015). Caspofungin: Pharmacodynamics, pharmacokinetics, clinical uses and treatment outcomes. Critical Reviews in Microbiology, 42(5), 813–846. https://doi.org/10.3109/1040841x.2015.1068271
- 48. Vermes, A. (2000c). Flucytosine: a review of its pharmacology, clinical indications, pharmacokinetics, toxicity and drug interactions. Journal of Antimicrobial Chemotherapy, 46(2), 171–179. https://doi.org/10.1093/jac/46.2.171
- 49. Alzohairy, M. A. (2016). Therapeutics Role of Azadirachta indica(Neem) and Their Active Constituents in Diseases Prevention and Treatment. Evidence-based Complementary and Alternative Medicine, 2016(1). https://doi.org/10.1155/2016/7382506
- 50. Chen, C., Long, L., Zhang, F., Chen, Q., Chen, C., Yu, X., Liu, Q., Bao, J., & Long, Z. (2018). Antifungal activity, main active components and mechanism of Curcuma longa extract against Fusarium graminearum. PLoS ONE, 13(3), e0194284. https://doi.org/10.1371/journal.pone.0194284
- 51. Balakumar, S., Rajan, S., Thirunalasundari, T., & Jeeva, S. (2011). Antifungal activity of Ocimum sanctum Linn. (Lamiaceae) on clinically isolated dermatophytic fungi. Asian Pacific Journal of Tro-



pical Medicine, 4(8), 654–657. https://doi.org/10.1016/s1995-7645(11)60166-1

- 52. Balakumar, S., Rajan, S., Thirunalasundari, T., & Jeeva, S. (2011c). Antifungal activity of Ocimum sanctum Linn. (Lamiaceae) on clinically isolated dermatophytic fungi. Asian Pacific Journal of Tropical Medicine, 4(8), 654–657. https://doi.org/10.1016/s1995-7645(11)60166-1
- 53. Balakumar, S., Rajan, S., Thirunalasundari, T., & Jeeva, S. (2011b). Antifungal activity of Ocimum sanctum Linn. (Lamiaceae) on clinically isolated dermatophytic fungi. Asian Pacific Journal of Tropical Medicine, 4(8), 654–657. https://doi.org/10.1016/s1995-7645(11)60166-1
- 54. Balkrishna, A., Verma, S., Mulay, V. P., Gupta, A. K., Haldar, S., & Varshney, A. (2022b). Withania somnifera (L.) Dunal whole-plant extracts exhibited anti-sporotrichotic effects by destabilizing peripheral integrity of Sporothrix globosa yeast cells. PLoS Neglected Tropical Diseases, 16(6), e0010484. https://doi.org/10.1371/journal.pntd.0010484
- 55. Gunasekara, T., Radhika, N., Ragunathan, K., Gunathilaka, D., Weerasekera, M., Hewageegana, H., Arawwawala, L., & Fernando, S. (2017). Determination of antimicrobial potential of five herbs used in ayurveda practices against Candida albicans, Candida parapsilosis and methicillin resistant Staphylococcus aureus. Ancient Science of Life, 36(4), 187. https://doi.org/10.4103/asl.asl\_179\_16
- 56. Saniasiaya, J., Salim, R., Mohamad, I., & Harun, A. (2017). Antifungal Effect of Malaysian Aloe vera Leaf Extract on Selected Fungal Species of Pathogenic Otomycosis Species in In Vitro Culture Medium. Oman Medical Journal, 32(1), 41–46. https://doi.org/10.5001/omj.2017.08
- 57. Hazen, E. L., & Brown, R. (1951). Fungicidin, an antibiotic produced by a soil actinomycete. Experimental Biology and Medicine, 76(1), 93–97. https://doi.org/10.3181/00379727-76-18397
- 58. Dutcher, J. D. (1968). The discovery and development of Amphotericin B. Diseases of the Chest, 54, 296–298. https://doi.org/10.1378/chest.54.supplement\_1.296
- 59. Vermes, A. (2000e). Flucytosine: a review of its pharmacology, clinical indications, pharmacokinetics, toxicity and drug interactions. Journal of Antimicrobial Chemotherapy, 46(2), 171–179. https://doi.org/10.1093/jac/46.2.171
- 60. Sawyer, P. R., Brogden, R. N., Pinder, R. M., Speight, T. M., & Avery, G. S. (1975). Miconazole. Drugs, 9(6), 406–423. https://doi.org/10.2165/00003495-197509060-00002
- 61. Fromtling, R. A. (1988b). Overview of medically important antifungal azole derivatives. Clinical Microbiology Reviews, 1(2), 187–217. https://doi.org/10.1128/cmr.1.2.187
- 62. Van Cutsem, J. (1989c). The in-vitro antifungal spectrum of itraconazole. Mycoses, 32(s1), 7–13. https://doi.org/10.1111/j.1439-0507.1989.tb02290.x
- 63. Zervos, M., & Meunier, F. (1993). Fluconazole (Diflucan®): a review. International Journal of Antimicrobial Agents, 3(3), 147–170. https://doi.org/10.1016/0924-8579(93)90009-t
- 64. Johnson, L. B., & Kauffman, C. A. (2003). Voriconazole: a new triazole antifungal agent. Clinical Infectious Diseases, 36(5), 630–637. https://doi.org/10.1086/367933
- 65. Courtney, R., Radwanski, E., Lim, J., & Laughlin, M. (2004). Pharmacokinetics of Posaconazole Coadministered with Antacid in Fasting or Nonfasting Healthy Men. Antimicrobial Agents and Chemotherapy, 48(3), 804–808. https://doi.org/10.1128/aac.48.3.804-808.2004
- 66. Maertens, J. A., Raad, I. I., Marr, K. A., Patterson, T. F., Kontoyiannis, D. P., Cornely, O. A., Bow, E. J., Rahav, G., Neofytos, D., Aoun, M., Baddley, J. W., Giladi, M., Heinz, W. J., Herbrecht, R., Hope, W., Karthaus, M., Lee, D., Lortholary, O., Morrison, V. A., . . . Ullmann, A. J. (2015). Isavuconazole versus voriconazole for primary treatment of invasive mould disease caused by



E-ISSN: 2582-2160 • Website: <u>www.ijfmr.com</u> • Email: editor@ijfmr.com

Aspergillus and other filamentous fungi (SECURE): a phase 3, randomised-controlled, non-inferiority trial. The Lancet, 387(10020), 760–769. https://doi.org/10.1016/s0140-6736(15)01159-9

- 67. Ryder, N. S. (1985). Specific inhibition of fungal sterol biosynthesis by SF 86-327, a new allylamine antimycotic agent. Antimicrobial Agents and Chemotherapy, 27(2), 252–256. https://doi.org/10.1128/aac.27.2.252
- Mora-Duarte, J., Betts, R., Rotstein, C., Colombo, A. L., Thompson-Moya, L., Smietana, J., Lupinacci, R., Sable, C., Kartsonis, N., & Perfect, J. (2002). Comparison of caspofungin and amphotericin B for invasive candidiasis. New England Journal of Medicine, 347(25), 2020–2029. https://doi.org/10.1056/nejmoa021585
- 69. Chandrasekar, P. H., & Sobel, J. D. (2006). Micafungin: a new Echinocandin. Clinical Infectious Diseases, 42(8), 1171–1178. https://doi.org/10.1086/501020
- 70. Reboli, A. C., Rotstein, C., Pappas, P. G., Chapman, S. W., Kett, D. H., Kumar, D., Betts, R., Wible, M., Goldstein, B. P., Schranz, J., Krause, D. S., & Walsh, T. J. (2007). Anidulafungin versus Fluconazole for Invasive Candidiasis. New England Journal of Medicine, 356(24), 2472–2482. https://doi.org/10.1056/nejmoa066906
- 71. Thompson, G. R., Vazquez, J., Soriano, A., Skoutelis, A., Ostrosky-Zeichner, L., Mena, K., Navalta, L., Sandison, T., & Pappas, P. (2018). 1718. Rezafungin Clinical safety and efficacy in patients with candidemia and/or invasive candidiasis in the randomized, Double-Blind, multicenter, Phase 2 STRIVE study. Open Forum Infectious Diseases, 5(suppl\_1), S52. https://doi.org/10.1093/ofid/ofy209.124
- 72. Schaffner, C. P., et al. (1953). Nystatin, a new antifungal antibiotic. Antibiotics & Chemotherapy, 3(9), 545–548.DOI: 10.1159/000221631 https://www.karger.com/Article/Abstract/221631
- 73. Peyton, L. R., Gallagher, S., & Hashemzadeh, M. (2001). Current and Emerging Azole Antifungal Agents. Infectious Diseases in Obstetrics and Gynecology, 9(3), 133–144. DOI: 10.1155/S1064744901000236 https://www.ncbi.nlm.nih.gov/pmc/articles/PMC88906/
- 74. Denning, D. W. (2003). Echinocandin antifungal drugs. The Lancet, 362(9390), 1142–1151.DOI: 10.1016/S0140-6736(03)14472-8 https://www.thelancet.com/article/S0140-6736(03)14472-8/fulltext
- 75. Thompson III, G. R., et al. (2024). Efficacy and safety of rezafungin and caspofungin in candidaemia and invasive candidiasis. The Lancet Infectious Diseases, 24(3), 319–328.DOI: 10.1016/S1473-3099(23)00551-0https://www.thelancet.com/journals/laninf/article/PIIS1473-3099(23)00551-0/fulltext
- 76. Pathadka, S., Yan, V. K. C., Neoh, C. F., Al-Badriyeh, D., Kong, D. C. M., Slavin, M. A., Cowling, B. J., Hung, I. F. N., Wong, I. C. K., & Chan, E. W. (2022). Global consumption trend of antifungal agents in humans from 2008 to 2018: Data from 65 Middle- and High-Income countries. Drugs, 82(11), 1193–1205. https://doi.org/10.1007/s40265-022-01751-x
- 77. Zhou, W., Li, X., Osmundson, T., Shi, L., Ren, J., & Yan, H. (2018). WGS analysis of ST9-MRSA-XII isolates from live pigs in China provides insights into transmission among porcine, human and bovine hosts. Journal of Antimicrobial Chemotherapy, 73(10), 2652–2661. https://doi.org/10.1093/jac/dky245
- 78. Fisher, M. C., Alastruey-Izquierdo, A., Berman, J., Bicanic, T., Bignell, E. M., Bowyer, P., Bromley, M., Brüggemann, R., Garber, G., Cornely, O. A., Gurr, S. J., Harrison, T. S., Kuijper, E., Rhodes, J., Sheppard, D. C., Warris, A., White, P. L., Xu, J., Zwaan, B., & Verweij, P. E. (2022). Tackling the emerging threat of antifungal resistance to human health. Nature Reviews Microbiology, 20(9), 557–



571. https://doi.org/10.1038/s41579-022-00720-1

- 79. Ghazi, B. K., Zahid, U., Usman, M. A., Kazmi, Z., Hunain, R., Riaz, M. M. A., Elmahi, O. K. O., Essar, M. Y., & Hasan, M. M. (2022b). Antifungal Drugs Shortage in India amidst Looming Increase in Invasive Fungal Infections among COVID-19 Patients: An Impending Crisis. INNOVATIONS in Pharmacy, 13(2), 3. https://doi.org/10.24926/iip.v13i2.4480
- 80. Bansal, P., & Baishnab, S. (2020b). A pharmacoepidemiology study of local fungal infections in skin and venereal diseases outpatient department of a rural tertiary care hospital. International Journal of Basic & Clinical Pharmacology, 9(4), 616. https://doi.org/10.18203/2319-2003.ijbcp20201187
- Arun, A. B., Hasan, M. M., Rackimuthu, S., Ullah, I., Mir, T., & Saha, A. (2021b). Antifungal drug shortage in India amid an increase in invasive fungal functions during the coronavirus disease 2019 (COVID-19) pandemic. Infection Control and Hospital Epidemiology, 43(12), 1965–1966. https://doi.org/10.1017/ice.2021.426
- 82. India | Create a custom map | MapChart. (n.d.). MapChart. https://www.mapchart.net/india.html
- 83. Alzohairy, M. A. (2016b). Therapeutics Role of Azadirachta indica(Neem) and Their Active Constituents in Diseases Prevention and Treatment. Evidence-based Complementary and Alternative Medicine, 2016(1). https://doi.org/10.1155/2016/7382506
- 84. Subbarayan, E., & Chittoria, R. (2017). Innovative usage of accessory auricles as full-thickness skin graft. Journal of Cutaneous and Aesthetic Surgery, 10(3), 150. https://doi.org/10.4103/jcas.jcas\_70\_17