

# ANTIFUNGAL DRUGS

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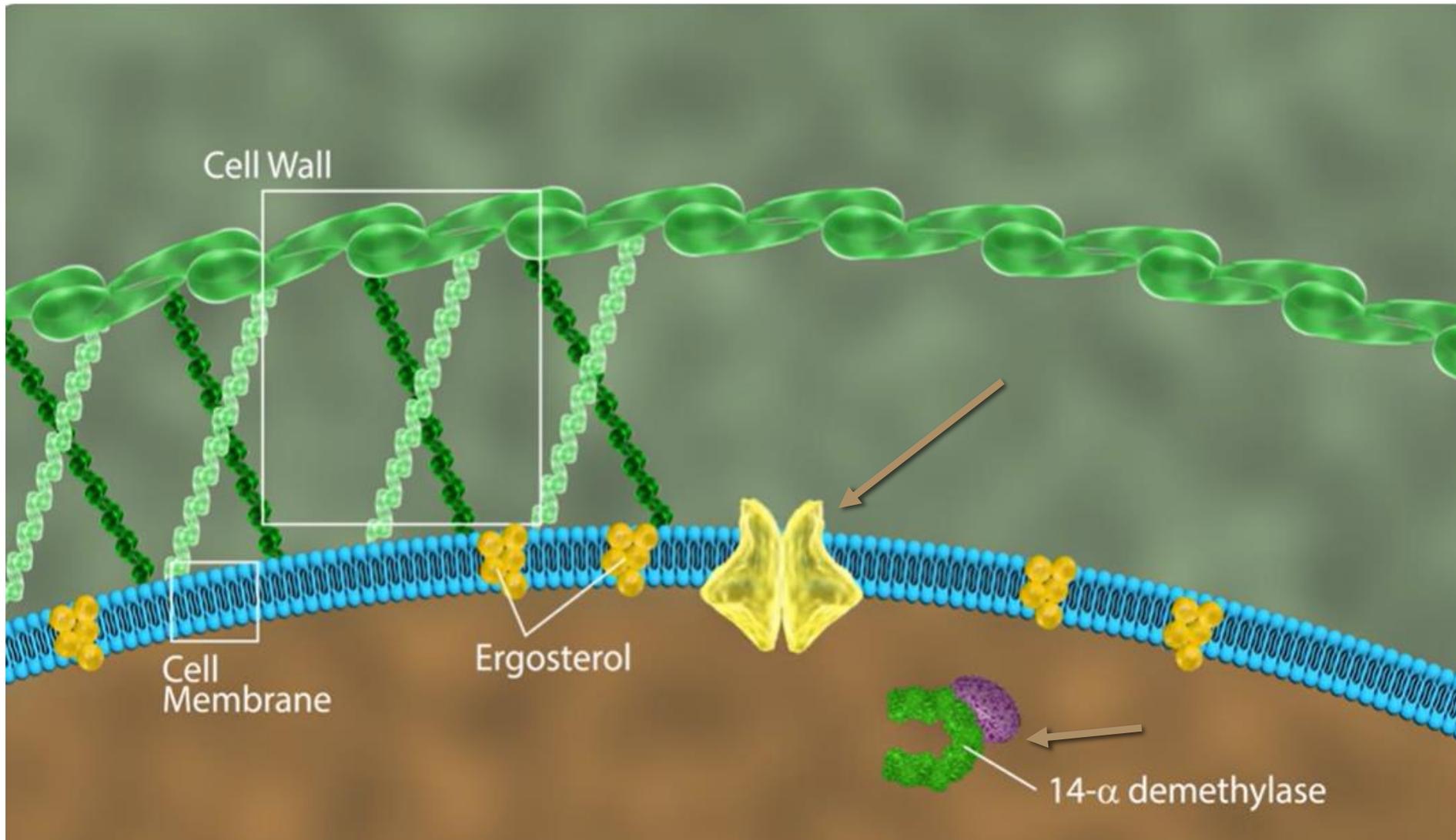
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# FUNGAL CELL WALL STRUCTURE



# Antifungal therapy

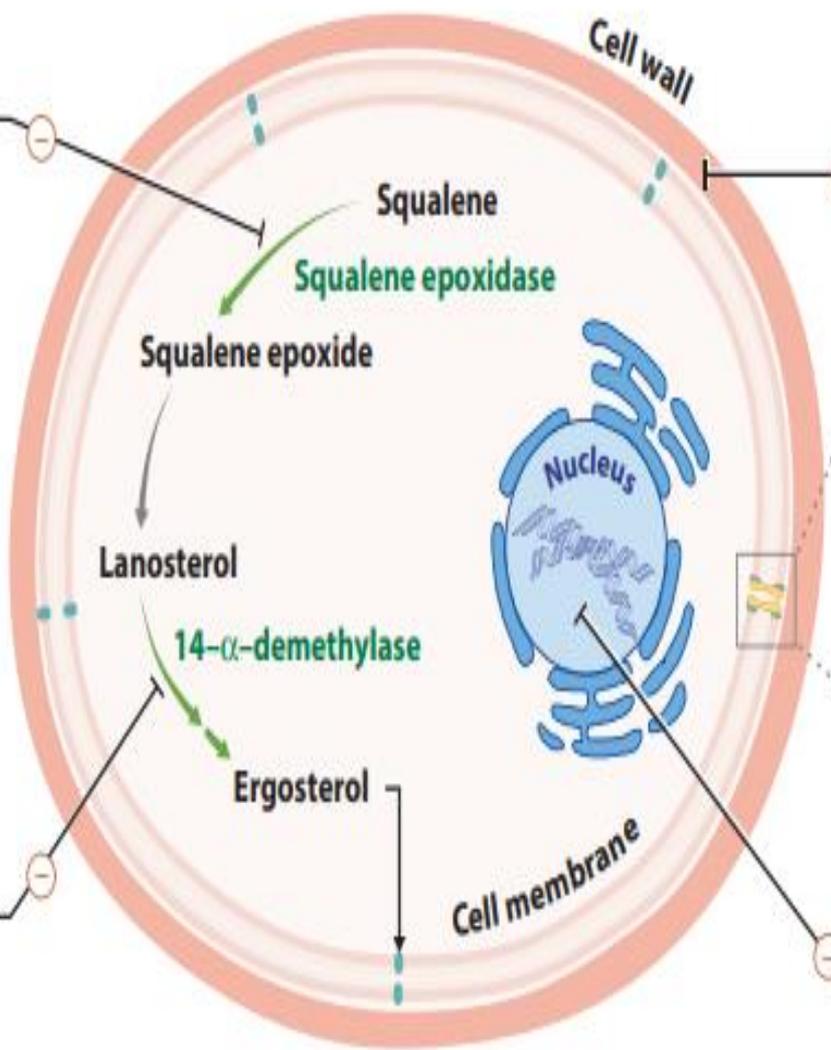
## LANOSTEROL SYNTHESIS

Terbinafine

## ERGOSTEROL SYNTHESIS

Azoles  
Clotrimazole  
Fluconazole  
Itraconazole  
Ketoconazole  
Miconazole  
Voriconazole

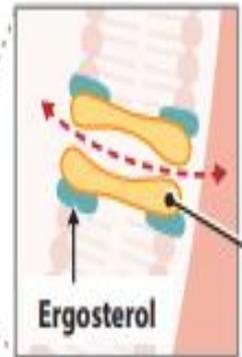
## FUNGAL CELL



## CELL WALL SYNTHESIS

Echinocandins  
Anidulafungin  
Caspofungin  
Micafungin

## CELL MEMBRANE INTEGRITY



Polyenes  
Amphotericin B  
Nystatin

## NUCLEIC ACID SYNTHESIS

Flucytosine

# Antifungals

Cell wall inhibitors

Antimetabolites

Terbinafine

Amphotericin B

Nystatin

Echinocandins

**Azoles**

Griseofulvin

Flucytosine

# Terbinafine

## **Mechanism:** *fungicidal*

- Inhibition of squalene epoxidase enzyme which is essential for ergosterol synthesis of cell membrane.

## **Indications:**

Systemic (oral) & topical for dermatophytes (more effective than griseofulvin).

## **Side effects:**

GIT and taste disturbances, hepatotoxicity, headache.

# Advantages over Azoles:

1. Squalene epoxidase enzyme is not present in human (more selective toxicity).
2. No inhibition of cytochrome P<sub>450</sub> (no serious adverse effect of azoles).

# Azoles

## **Mechanism of action: fungicidal**

inhibit ergosterol synthesis of cell membrane by inhibiting fungal cytochrome P<sub>450</sub> leading to membrane dysfunction.

## **Members :**

**1- Ketoconazole**

**2- Itraconazole**

**3- Fluconazole**

**4- Posaconazole**

# Ketoconazole:

1<sup>st</sup> oral broad spectrum antifungal.

## It is used for:

- Deep fungal infections (mild - non meningeal). 2<sup>nd</sup> line to amphotericin
- Candida infection.
- Dermatophytes resistant to griseofulvin & terbinafine (oral and topical).

## Avoid combination with:

- ❑ Antacids or H<sub>2</sub> blockers → decrease gastric acidity → decrease ketoconazole absorption.
- ❑ Amphotericin B: ketoconazole → decrease amphotericin effect by decreasing ergosterol

## Adverse effects:

1. Nausea - vomiting - rash (common).
2. Hepatotoxic (serious).
3. **Inhibition of human cytochrome P<sub>450</sub>**

## **Inhibition of human cytochrome P<sub>450</sub>:**

**it inhibits:**

### **Steroid synthesis which depends on cytochrome P450:**

- ❖ Corticosteroids → adrenal suppression (used in Cushing's disease).
- ❖ Testosterone → gynecomastia & impotence (used in cancer prostate).
- ❖ Female sex hormones → menstrual irregularities & infertility

### **Metabolism of drugs → drug interactions:**

- ❖ Increased level of astemizole & terfenadine → arrhythmia.
- ❖ Increased level of oral anticoagulants & antiepileptics.

## Itraconazole and fluconazole

- ❖ These drugs are azoles that are more specific to fungal cytochrome P<sub>450</sub> than to human cytochrome P<sub>450</sub> compared to ketoconazole.
- ❖ Less toxic (less effect on human cytochrome P<sub>450</sub>): less hepatotoxic, less adrenal suppression & less drug interactions.
- ❖ More effective.

# Fluconazole:

**Drug of choice in esophageal and oropharyngeal candidiasis.**

- Drug of choice in treatment and secondary prophylaxis against cryptococcal meningitis.
- Equivalent to amphotericin B in systemic candidiasis

# *Posaconazole*

- The **broadest-spectrum** azole.
- The only azole with activity against **mucormycosis**.
- It is used for prophylaxis of fungal infections during **cancer chemotherapy**.
- Inhibitor of **CYP3A4** → increasing the levels of cyclosporine and tacrolimus

# Amphotericin B

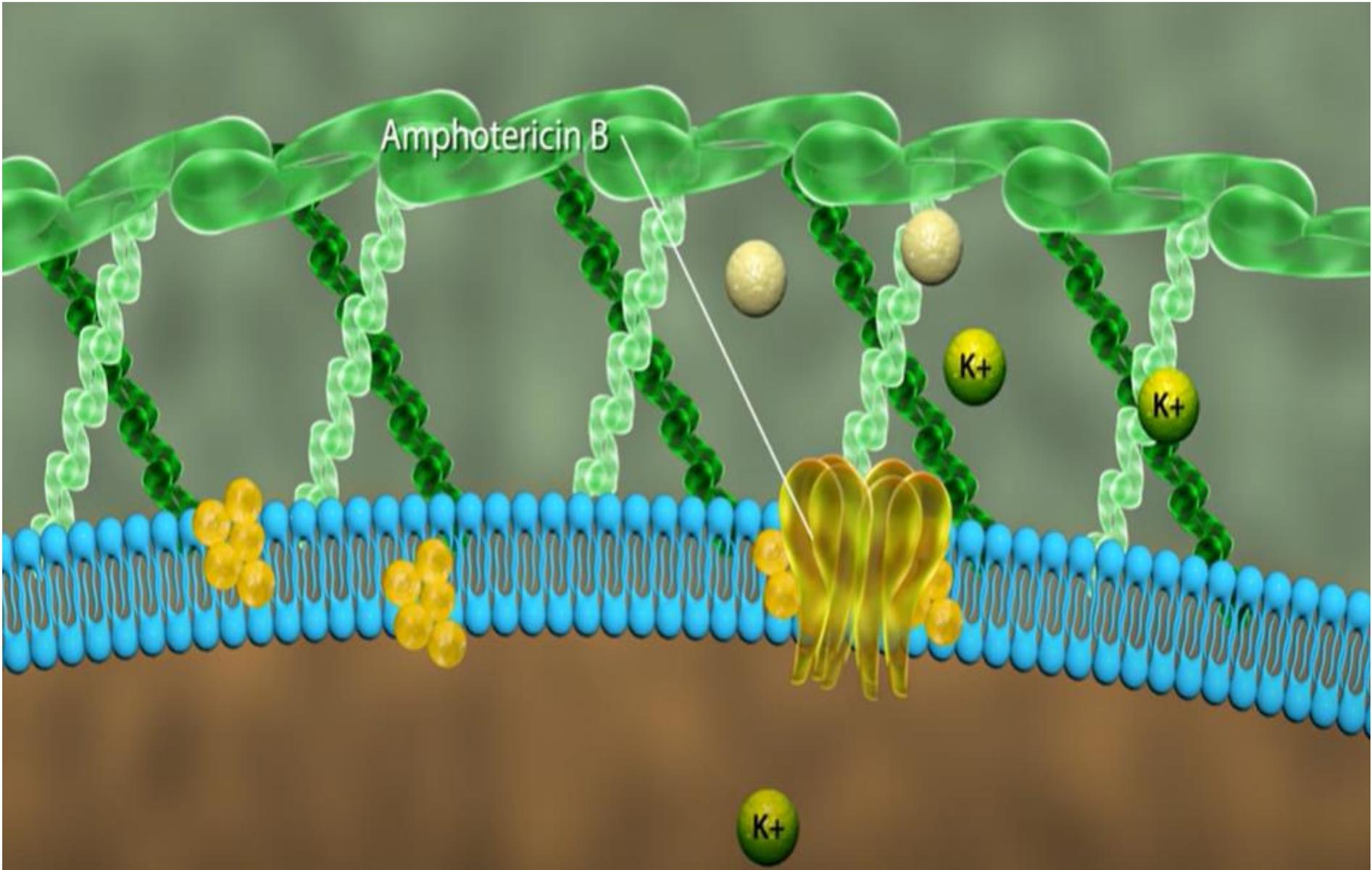
## **Mechanism of action: fungicidal**

- Binds to ergosterol of cell membrane → formation of artificial pores → leakage of important cell constituents' → cell death.

## **Indications: deep infections especially:**

- Severe life threatening (I.V - not absorbed orally).
- Meningitis (intrathecal- does not reach CSF after I.V.I).

Amphotericin B



- Side effects & toxicity:

- **Infusion related:** Fever, rigors, vomiting, hypotension & shock after I.V infusion.

**Can be avoided by:** Slow infusion rate and pretreatment with antihistamines, antipyretics.

- **Dose-related:** nephrotoxicity. **Can be decreased by:** dose reduction.

- **Convulsion.**

# Nystatin

## **Mechanism:**

Binds to ergosterol of fungal cell membrane → formation of artificial pores—» damage of membrane → leakage of important cell constituents → cell death.

# Indications: (too toxic for systemic use).

## Used locally in:

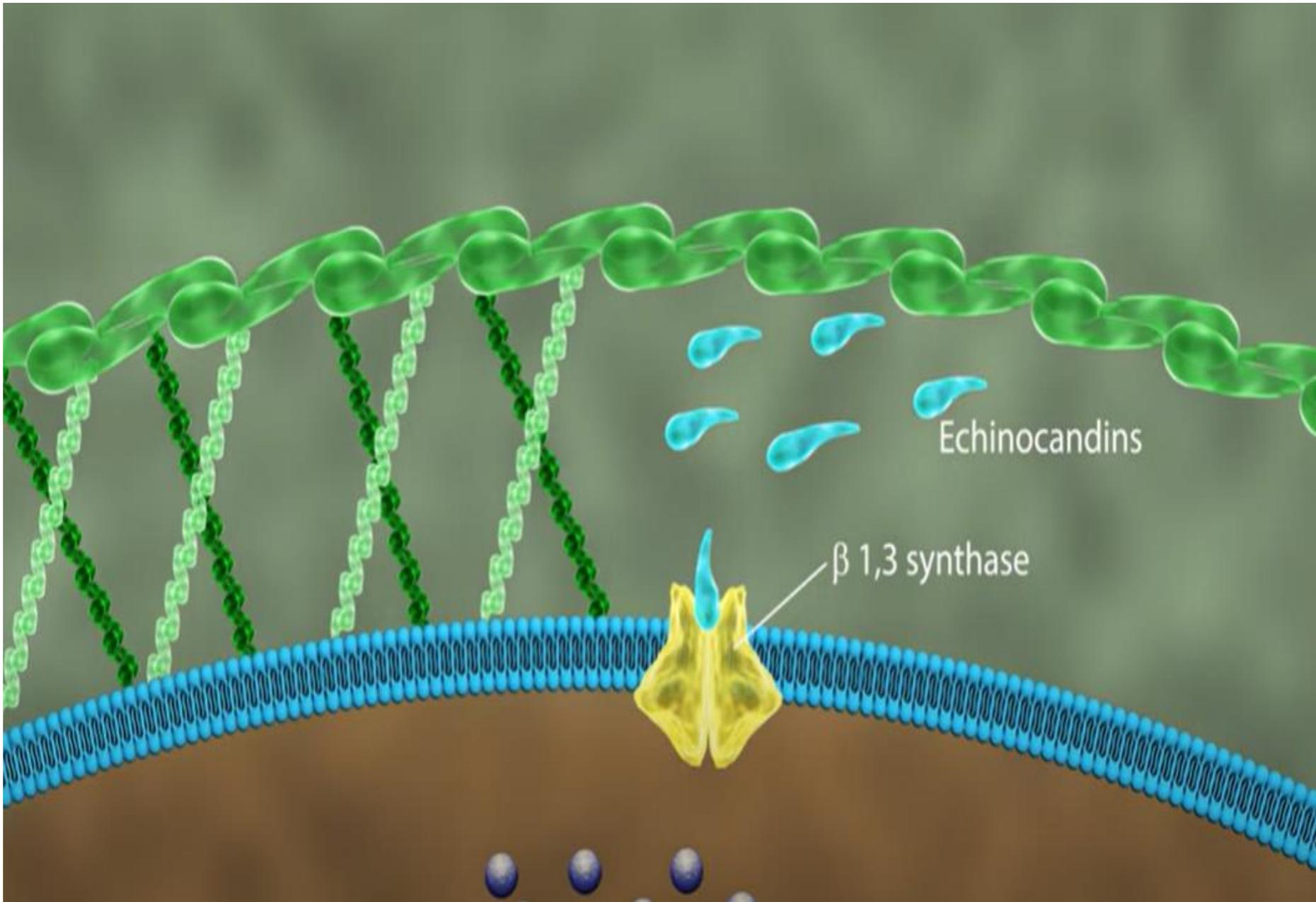
1. Oropharyngeal and GI **Candida**: oral (**not absorbed**).
2. Cutaneous **Candida**: topical (**non irritant- rarely causes allergy**).
3. Vaginal **Candida**: It is given **both topically and orally** because quite often vaginal **Candida** is associated with gastrointestinal **Candida** which acts as a source of reinfection of vagina.

# Echinocandins

## Caspofungin – Micafungin

- Mechanism:

Inhibits synthesis of a glucose polymer that is necessary for maintaining structure of fungal cell wall → loss of cell wall integrity → lysis & death.



- **Uses:** (IV)

**Caspofungin:** candidiasis & invasive aspergillosis refractory to amphotericin.

**Micafungin:** mucocutaneous candidiasis and for prophylaxis of *Candida* infections in bone marrow transplant patients

- **Adverse Effects:**

***Infusion-related:*** GIT upset, headache, fever & flushing (histamine release).

# Flucytosine

## Mechanism of action:

- Cytotoxic, transformed to 5-fluorouracil (5-FU) → inhibits nucleic acid synthesis.
- **Selective toxicity** occurs because mammalian cells cannot transform flucytosine into 5-FU.

## ▪ Indications:

*Given orally with amphotericin or azoles in Cryptococcal infections.*

## Adverse effects:

1. Bone marrow depression (reversible).
2. Hair loss.
3. Hepatotoxic.

## Advantages of combination of flucytosine with amphotericin B:

1. **Decrease resistance** to amphotericin B.
2. Decrease amphotericin **nephrotoxicity** (lower doses of amphotericin are used).

# Griseofulvin

## **Mechanism:** *Fungistatic*

Concentrated in newly formed keratin (e.g nails) preventing its infection by:

- Interfering with microtubular function → interfere with mitosis.
- Inhibiting nucleic acid synthesis.

## **Indications:** not active topically

- Dermatophyte infections (given orally: decreased absorption by high fat diet).
- Largely replaced by terbinafine & azoles

## Adverse effects :

1. Nausea-vomiting.
2. Headache - mental confusion.
3. Hepatotoxic.
4. Enzyme inducer → decrease warfarin level.
5. Teratogenic , Carcinogenic

## *Systemic therapy is used in:*

- 1- Resistance to topical therapy.
- 2- Wide or inaccessible areas.
- 3- Severe infections.
- 4- Low immunity of patient.

N.B: Superficial fungal infections are **treated first with topical** agents

# Antifungals

*Drugs for systemic (deep) fungal infections*

Amphotericin B

Echinocandins

**Azoles**

Flucytosine

*Drugs for superficial infections*

Nystatin

Terbinafine

**Azoles**

Griseofulvin

THANK YOU