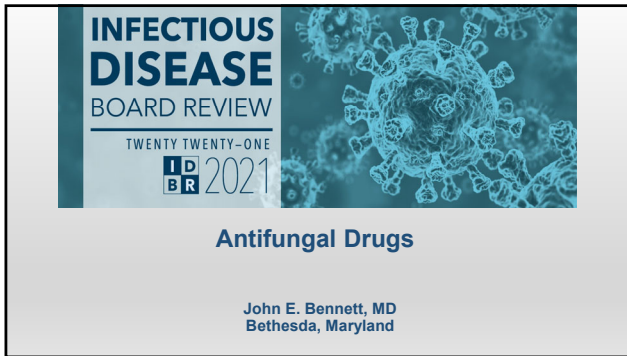


11 - Antifungal Drugs

Speaker: John Bennett, MD



Disclosures of Financial Relationships with Relevant Commercial Interests

- None

Disclosures of Off-Label Use

- Will be cited as discussed

Plan of the talk

- 1. review of antifungals
 - Key points are underlined
- 2. questions on antifungals with answers
- 3. Key points

Antifungal drugs

The diagram illustrates the mechanisms of various antifungal drugs on a cell. It shows a cell with a cell wall, cell membrane, nucleus, vacuole, and cytoplasm. Arrows indicate the following effects:

- Azoles:** stops sterol synthesis for the cell membrane.
- Echinocandins:** block cell wall synthesis (glucan fibers).
- Flucytosine:** blocks DNA synthesis in the nucleus.
- Amphotericin:** makes cell membrane leak.

DRUG RESISTANCE IN FUNGI: BLOCK TARGET ENZYME

1. ASPERGILLUS AND CANDIDA: AZOLE RESISTANCE IN CYP51A

- gene CYP51A ← modified CYP51A = drug resistance
- Lanosterol → C14-demethylase → ergosterol in cell membrane
- ↑ Azole

2. CANDIDA : ECHINOCANDIN RESISTANCE IN FKS1, FKS2

- genes FKS1 and FKS2 ← modified gene = drug resistance
- Substrates → glucan synthase → glucan fibers in cell wall
- ↑ Echinocandin

Antifungal resistant species ☹️

- **Amphotericin B resistant:** Scedosporium apiospermum (Pseudallescheria boydii), Aspergillus terreus, Variable in Candida lusitanae, C. auris +/-
- **Fluconazole resistant:** All moulds, Candida krusei, Candida auris, Candida haemulonii, some Candida glabrata
- **Voriconazole resistant:** mucormycosis, some cryptic Aspergillus species higher MIC's: (lentulus, ustus, calidoustus)
- **Posaconazole resistance:** like vori but more mucormycosis activity
- **Echinocandin resistance:** Cryptococcus, Trichosporon, Histoplasma, Blastomyces, Coccidioides, moulds other than Aspergillus.

11 - Antifungal Drugs

Speaker: John Bennett, MD

Azole antifungals

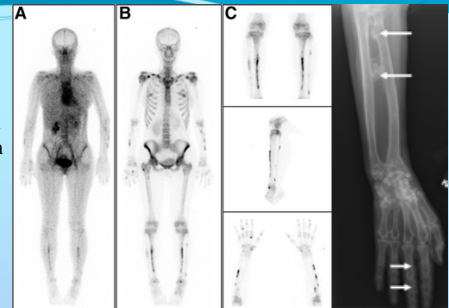
Voriconazole: the fundamentals

- Candida, Aspergillus, Scedosporium apiospermum, etc.
- Children are rapid metabolizers. Japanese 20% slow (2C19)
- Good CSF levels, none in urine.
- IV (sulfobutylcyclodextran=16x vori dose) accumulates in azotemia but not obviously toxic. Use oral in azotemia.
- Many drug interactions, Increases other drug levels: cyclosporine, tacrolimus, serolimus, steroids (budesonide, fluticasone), etc
- Side effects: hallucinations, hepatitis, photosensitivity, visual changes, peripheral neuropathy
- Many months of Rx: skin cancer, periostitis

Photosensitivity from voriconazole



- Voriconazole
Periostitis:
-Bone pain
-Months of Rx
-Alk phos high
-Plasma fluoride high (fluorosis)
-Bone scan
-Exostoses



Rossier, et al. Eur J Nuc Med Mol Imag 2011 Wermers, et al. CID 2011

Isavuconazonium/Isavuconazole

- Noninferior to vori in invasive aspergillosis.
- Use for mucor controversial
- Inferior to caspofungin for candidemia
- No good data on prophylaxis
- Pharma: like vori but long half life (5.4 days), no drug in CSF or urine. Fewer drug interactions than vori or posa. Teratogenic.
- Isavuconazonium 372mg=isavuconazole 200 mg
- Load with 200 mg q8h X6 then 200 mg qd, IV or PO
- No dose change for renal or moderate liver failure.

Posaconazole

- Approved for prophylaxis in GVHD or prolonged neutropenia.
- Extended release three 100 mg tablets twice first day then daily. IV same dose, has cyclodextran. 7-10 days for steady state. Check trough levels (usually 1-5 mcg/ml)
- Has been used in mucormycosis once patient has responded to amphotericin B
- Interactions with CYP3A4 increase some drug levels
- Well tolerated. Hypertension, hypokalemia

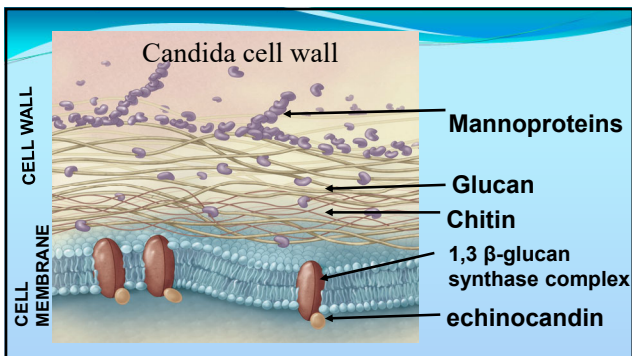
11 - Antifungal Drugs

Speaker: John Bennett, MD

FLUCONAZOLE

- FEW SIDE EFFECTS ,WIDE DOSAGE RANGE. DRY SKIN, ALOPECIA
- FOUND IN URINE, CSF. ACCUMULATES IN AZOTEMIA.
- DRUG-DRUG INTERACTIONS. TERATOGENIC
- CANDIDIASIS, COCCIDIOIDAL MENINGITIS, PROPHYLAXIS IN HSCT,
- VERY LOW BIRTHWEIGHT INFANTS, RINGWORM, OTHERS
- NO MOLD ACTIVITY

Echinocandins



Caspofungin, Micafungin, Anidulafungin

- All Candida (including C. auris and C. parapsilosis) susceptible but resistance can arise during long therapy. Mold activity: Aspergillus
- Cryptococcus, Trichosporon, endemic mycoses resistant
- IV once daily. Plasma half life: 10-15 hr.
- No drug in urine. Azotemia: same dose
- Protein binding high: poor penetration into CSF and vitreous humor of eye
- Drug interactions: none important

Clinical trials in deeply invasive candidiasis

- ☺ Treatment candidemia)
- ☺ Caspofungin, micafungin, anidulafungin effective
- ☹ Isavuconazole “not noninferior” to caspofungin in candidemia (don’t use)
- ☺ Prophylaxis for candidiasis: trials in micafungin (neutropenia), fluconazole (HSCT), posaconazole (HSCT)

Caspofungin and Micafungin in invasive aspergillosis

- ☹ IDSA Guidelines: “Primary therapy with an echinocandin is NOT recommended.”
- ☺ Prophylaxis for aspergillosis: micafungin best studied, most often used, not FDA approved

11 - Antifungal Drugs

Speaker: John Bennett, MD

Flucytosine

- Bioavailability 100%, good levels in CSF, eye, urine
- Accumulates in azotemia: bone marrow depression, hepatitis, colitis. Measure blood levels/dose adjust.
- Drug resistance arises during monotherapy.
- Used with ampho in cryptococcal meningitis

Now for a few questions



Question #1

A 47-year-old male with known HIV, poorly compliant with ARV, last CD4 20/mcl, presents with low grade fever and headache. Blood culture is growing a yeast, not yet identified. Starting micafungin would be a poor choice if the isolate is which of the following:

- A. *Candida parapsilosis*
- B. *Cryptococcus gattii*
- C. *Candida auris*
- D. *Candida krusei*
- E. *Candida glabrata*

Question #2

A 72 yr man with diabetes mellitus, renal failure and a central venous catheter developed fever and hypotension. Blood cultures grew *Candida lusitanae*. On day 5 of liposomal amphotericin B 5 mg/kg he remained febrile and his creatinine rose from 4.5 to 6.0 mg/dl.

Question #2 Continued

In addition to changing his IV catheter, which of the following would be most appropriate?:

- A. Itraconazole
- B. Micafungin
- C. Amphotericin B lipid complex
- D. IV Voriconazole
- E. Isavuconazole

Question #3

Echinocandin class of antifungals has which mechanism of action:

- A. inhibits synthesis of membrane sterols
- B. damages cytoplasmic membrane
- C. interferes with synthesis of fungal cell wall glucans
- D. inhibits fungal DNA synthesis
- E. interfere with synthesis of fungal cell wall chitin

11 - Antifungal Drugs

Speaker: John Bennett, MD

Question #4

A 37 yr female with diabetes mellitus is admitted for ketoacidosis, fever and sinus pain. Biopsy of a necrotic area of the middle turbinate shows wide, branching nonseptate hyphae. Serum creatinine is 2.5 mg/dl.

Question #4 Continued

Which of the following would be most appropriate?

- A. Voriconazole
- B. Anidulafungin
- C. Fluconazole
- D. Liposomal amphotericin B
- E. Itraconazole

Question #5

You are asked to advise your hem-onc colleagues as to what prophylactic antifungal agent might be useful in preventing aspergillosis in their patients with prolonged neutropenia or acute graft-vs-host disease .

Question #5 Continued

According to the IDSA guidelines and literature you recommend:

- A. itraconazole solution
- B. posaconazole
- C. micafungin
- D. voriconazole
- E. caspofungin

Question #6

45 yr old male 6 weeks post stem cell transplant for myelodysplasia, with a history of chronic hepatitis C was discharged home to Florida on cyclosporine, mycophenylate, prednisone , Bactrim (tmp/smz), citalopram and voriconazole. Diffuse nonpruritic erythema developed over his sun exposed skin.

Question #6 Continued

The most probable cause was:

- A. porphyria cutanea tarda
- B. graft versus host disease
- C. drug interaction
- D. voriconazole
- E. Bactrim allergy

11 - Antifungal Drugs

Speaker: John Bennett, MD

Question #7

A 66 yr old male with neutropenia following chemotherapy for lung cancer, serum creatinine 5 mg/dl, and congestive heart failure is found to have a *Scedosporium apiospermum* lung abscess.

Question #7 Continued

Which of the following would be preferred?

- A. Anidulafungin
- B. Itraconazole
- C. Micafungin
- D. Oral voriconazole
- E. Liposomal amphotericin B

Question #8

- 65 yr wm admitted with cryptococcal meningitis, seizures, diabetes mellitus and granulomatosis with polyangiitis. Given conventional amphotericin B, flucytosine, phenytoin, glipizide, prednisone and cyclophosphamide.
- By the end of the first week of treatment, his creatinine had risen from 1.6 to 3 mg/dl.
- By the end of the second week his WBC had fallen to 1.2K, platelets 60K and diarrhea began.

Question #8 Continued

The cause of his WBC falling to 1.2K, platelets 60K and copious diarrhea is most likely which of these drugs?

- A. flucytosine
- B. phenytoin
- C. glipizide
- D. cyclophosphamide
- E. cytomegalovirus

Take home messages

- Ampho: not *Scedosporium* (*Pseudallescheria boydii*), *Candida lusitanae*, *Asperillus terreus*
- Only ampho for mucormycosis
- Fluconazole: not *Candida krusei*, *Candida auris*,
- +/- *Candida glabrata*
- Echinocandins: not *Trichosporon* or *crypto*
- Know mechanisms of action: glucan, sterol, cell membrane, DNA synthesis
- Flucytosine WBC & plt fall, diarrhea, hepatitis

Take home, continued

- Voriconazole: **phototoxicity, periostitis, hallucinations**
- Azole interactions:
 - Increases other drug levels: cyclosporine, tacrolimus, serolimus, warfarin, midazolam, steroids, etc.
 - Decrease azole level: **phenytoin**, rifampin, etc

11 - Antifungal Drugs

Speaker: John Bennett, MD

The End

email

john_bennett@comcast.net