Isavuconazonium sulfate (Cresemba®)

IV and PO Only

Use requires formal ID Consult

Activity: Coverage against most strains of the following microorganisms, both in vitro and in clinical infections: Aspergillus flavus, Aspergillus fumigatus, Aspergillus niger, and Mucorales such as Rhizopus oryzae and Mucormycetes species

Criteria for Use:

- Treatment of invasive aspergillosis
- Treatment of invasive mucormycosis

Unacceptable Uses:

- Treatment for other fungal infections (Blastomyces, Histoplasma, etc.)
- Contraindicated with known hypersensitivity to isavuconazole. Caution in use in patients with hypersensitivity to other azoles
- Contraindicated in patients with familial short QT syndrome (shortens the QTc interval in a concentration-related manner)

Dosing in Adults:

- Standard dose: 372 mg IV/PO Q8H x 6 doses (48 hours; load), then 372 mg Q24H starting 12-24 hours after last loading dose
- No renal or hepatic dose adjustment
- IV must be administered via an infusion set with in-line filter (pore size 0.2-1.2 micron) and should be infused over a minimum of 1 hour
- No dose adjustment is necessary when changing from IV to PO

Monitoring:

AST/ALT/bilirubin at baseline and every 1-2 weeks after

Considerations for Use:

- Elevated LFTs have been reported in clinical trials. Elevations generally reversible and do not require discontinuation
- May cause fetal harm when administered to a pregnant woman

Important note regarding drug interactions:

- Isavuconazole is a substrate/inhibitor of CYP3A4 and has multiple drug interactions that may affect its levels and/or those of co-administered drugs. Dose adjustment may be necessary
- Some drugs with interactions of major significance include:
 - Carbamazepine
 - Rifampin/rifabutin
 - Ritonavir

Warfarin
Tacrolimus

Cyclosporine

- lacrolimus

Sirolimus

– Phenytoin

ALT= Alanine aminotransferase; AST= Aspartate aminotransferase; H= hour(s); ID= infectious diseases; IV= Intravenous; LFT's= Liver Function Tests; PO= by mouth; Q= every