

Division: Pharmacy Policy	Subject: Prior Authorization Criteria
Original Development Date: Original Effective Date: Revision Date:	March 22, 2022  November 13, 2024

**Cresemba<sup>®</sup> (isavuconazonium sulfate)**

**LENGTH OF AUTHORIZATION:** Up to one year

**REVIEW CRITERIA:**

- Patient must be  $\geq 1$  year of age for Cresemba for injection or  $\geq 6$  years of age who weigh  $\geq 16$  kg for Cresemba capsules.
- Prescribed by or in consultation with an infectious disease specialist.
- Patient must have a diagnosis of invasive aspergillosis or invasive mucormycosis.
- Recent (within 60 days) fungal culture and sensitivity (C&S) results.

**DOSING AND ADMINISTRATION:**

- Refer to product labeling at <https://www.accessdata.fda.gov/scripts/cder/daf/>
- Available as the following:
  - Cresemba for injection: 372 mg of isavuconazonium sulfate lyophilized powder single-dose vial (equivalent to 200 mg of isavuconazole)
  - Cresemba capsules: 74.5 mg of isavuconazonium sulfate (equivalent to 40 mg of isavuconazole) and 186 mg of isavuconazonium sulfate (equivalent to 100 mg of isavuconazole)