



Guilan University  
of Medical Sciences



Drug Information Leaflet (2)

## “Caspofungin”

**Amirmomenin Hospital**  
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NOVEMBER  
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## “Caspofungin”

### **Introduction**

Caspofungin is one of the licensed agents in Echinocandins class. Echinocandins are the newest class of antifungal agents to be developed. This agent is active against *Candida* and *Aspergillus*, but not *C. neoformans* or the agents of zygomycosis and mucormycosis.

### **Indication**

Aspergillosis, invasive (salvage therapy)  
Candidemia and other *Candida* infections  
*Candida* infection, prophylaxis in neutropenic cancer patients at substantial risk (off-label use)  
Candidiasis, chronic disseminated (hepatosplenic) (off-label use)  
Candidiasis, empiric therapy (non-neutropenic ICU patients) (off-label use)  
Candidiasis, esophageal  
Candidiasis, esophageal, in HIV-infected patients (alternative agent) (off-label use)  
Candidiasis, intravascular infections (native or prosthetic valve endocarditis, infection of implantable cardiac devices, suppurative thrombophlebitis) (off-label use)  
Candidiasis, osteoarticular infections (osteomyelitis or septic arthritis) (alternative therapy) (off-label use)  
Candidiasis, prophylaxis against invasive candidiasis (high-risk ICU patients in units with a high rate of invasive candidiasis) (alternative therapy; off-label use)  
Candidiasis, oropharyngeal (refractory disease) (alternative therapy) (off-label use)  
Fungal infections, empiric therapy (neutropenic patients)

**Dosing: Adult** (Duration of caspofungin treatment should be determined by patient status and clinical response).

### **Aspergillosis, invasive (salvage therapy)**

IV: Initial dose: 70 mg on day 1; subsequent dosing: 50 mg once daily. Duration of therapy should be a minimum of 6 to 12 weeks and depends on site of infection, extent of disease, and level/duration of immunosuppression.



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## **“Amphotericin B deoxycholate (conventional)”**

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### **Candidemia and other *Candida* infections**

IV: Initial dose: 70 mg on day 1; subsequent dosing: 50 mg once daily; generally continue for at least 14 days after the last positive culture or longer if neutropenia warrants. Higher doses (150 mg once daily infused over ~2 hours) compared to the standard adult dosing regimen (50 mg once daily) have not demonstrated additional benefit or toxicity in patients with invasive candidiasis (Betts 2009).

**Note:** IDSA Candidiasis guidelines recommend transition to fluconazole (usually after 5 to 7 days in non-neutropenic patients) in clinically stable patients with fluconazole-susceptible isolates and negative repeat cultures.

### ***Candida* infection, prophylaxis in neutropenic cancer patients at substantial risk (off-label use)**

IV: 50 mg once daily.

### **Candidiasis, chronic disseminated (hepatosplenic) (off-label use)**

IV: Initial dose: 70 mg on day 1; subsequent dosing: 50 mg daily for several weeks, followed by oral fluconazole therapy.

### **Candidiasis, empiric therapy (non-neutropenic ICU patients) (off-label use)**

IV: Initial dose: 70 mg on day 1; subsequent dosing: 50 mg once daily. Consider discontinuing after 4 to 5 days in patients with no clinical response; continue treatment for 2 weeks in patients who improve on antifungal therapy.

### **Candidiasis, esophageal**

IV: Manufacturer's labeling: 50 mg once daily; continue for 7 to 14 days after symptom resolution.

**Note:** The majority of patients studied for this indication also had oropharyngeal involvement.

Alternate recommendations: Initial dose: 70 mg on day 1; subsequent dosing: 50 mg daily; may transition to oral fluconazole therapy once oral intake tolerable. In patients with fluconazole-refractory disease, continue caspofungin for 14 to 21 days.

### **Candidiasis, esophageal, in HIV-infected patients (alternative agent) (off-label use)**

IV: 50 mg once daily; continue for 14 to 21 days. **Note:** A higher rate of relapse has been reported with echinocandins than with fluconazole.



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**Candidiasis, intravascular infections (native or prosthetic valve endocarditis, infection of implantable cardiac devices, suppurative thrombophlebitis) (off-label use):** IV: 150 mg daily. For native or prosthetic valve endocarditis, therapy should continue for at least 6 weeks after valve replacement surgery (longer durations in patients with abscesses or other complications); for patients with implantable cardiac devices, therapy should continue for 4 to 6 weeks after surgery (4 weeks for infections limited to generator pockets and at least 6 weeks for infections involving the wires); for suppurative thrombophlebitis, continue for at least 2 weeks after candidemia has cleared.

**Note:** Step-down to fluconazole therapy is recommended in clinically stable patients with fluconazole-susceptible isolates and negative repeat cultures.

**Candidiasis, osteoarticular infections (osteomyelitis or septic arthritis) (alternative therapy) (off-label use)**  
IV: 50 to 70 mg daily for at least 14 days, followed by fluconazole.

**Candidiasis, prophylaxis against invasive candidiasis (high-risk ICU patients in units with a high rate of invasive candidiasis) (alternative therapy; off-label use)**  
IV: Loading dose: 70 mg on day 1, then 50 mg daily.

**Candidiasis, oropharyngeal (refractory disease) (alternative therapy) (off-label use)**  
IV: Initial dose: 70 mg on day 1; subsequent doses: 50 mg once daily.

### **Fungal infections, empiric therapy (neutropenic patients)**

IV: Initial dose: 70 mg on day 1; subsequent dosing: 50 mg once daily; continue until resolution of neutropenia; if fungal infection confirmed, continue for a minimum of 14 days (continue for at least 7 days after resolution of both neutropenia and clinical symptoms); if clinical response inadequate, may increase up to 70 mg once daily if tolerated.

### **Products**

- Caspofungin acetate is available in vials of 50 and 70 mg of drug.
- Equilibrate the refrigerated vials to room temperature.
- Reconstitute both the 50- and 70-mg vials with 10.5 mL of sterile water for injection, sodium chloride 0.9%, or bacteriostatic water for injection and mix gently until dissolved.
- **Do NOT use dextrose-containing solutions.**
- Withdrawing 10 mL of the reconstituted solution will provide the full 50 or 70 mg as a clear solution.
- Do not use hazy, precipitated, or discolored solutions.
- The reconstituted solution should be withdrawn within 1 hour after reconstitution for preparation of the IV infusion.



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## **Administration**

- Caspofungin acetate is administered by IV infusion over a period of 1 hour.
- For a 70-mg dose, a solution volume of 250 mL should be used.
- For a 50- or 35-mg dose, a solution volume of either 250 or 100 mL may be used.

## **Stability & Compatibility Information**

- Intact vials of caspofungin acetate should be stored between 2 and 8 °C.
- **Intact vials exposed to ambient room temperature for longer than 48 hrs should be discarded.**
- The reconstituted solution may be stored for up to 1 hour after reconstitution at room temperature up to 25 °C but **should be withdrawn within 1 hour after reconstitution** to prepare the IV infusion solution.
- Caspofungin acetate may be diluted in **sodium chloride 0.9, 0.45, or 0.225% or Ringer's injection, lactated** for administration. The drug diluted in these solutions is stable for up to 24 hrs at room temperature up to 25 °C and up to 48 hrs refrigerated.
- Caspofungin acetate is **unstable in dextrose-containing solutions**; such solutions should not be used for reconstitution or dilution of this drug.

## **Other Drugs**

The manufacturer recommends that caspofungin acetate **not be administered with any other drugs.**

solution	compatible	incompatible
Sodium chloride 0.9%	*	

## **References**

- caspofungin: Drug information available from Lexicomp® Drug
- caspofungin: Patient drug information available from <https://www.uptodate.com/contents/caspofungin-patient-drug-information>