

Tacrolimus 100 mg/100 mL (1 mg/mL) Oral Suspension

NDC	PRESENTATION	EXPIRATION (BUD)	CASE QUANTITY
85383-0121-1	Multi-dose Aqueous Oral Suspension	110 Days, Room Temp	5 Bottles



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Directions of use: For oral administration as directed by a licensed healthcare professional. Shake vigorously for 10 seconds before use. Dosing regimen may be individualized in accordance with applicable standards of care. Avoid grapefruit or grapefruit juice. Therapeutic drug monitoring is recommended to maintain appropriate trough levels.

Storage and Handling: Store at Controlled Room Temperature 20°C — 25°C (68°F — 77°F). Protect from light. Do not use if container closure is damaged. Follow applicable special handling and disposal procedures.

Active Ingredients: Tacrolimus USP 100 mg

Inactive Ingredients: SUSPENDIT[®] QS (a proprietary vehicle consisting of: Water, Amorphophallus Konjac Root Powder, Siraitia Grosvenorii (Monk) Fruit Extract, Xanthan Gum, Potassium Sorbate 0.25% (preservative), Sodium Benzoate 0.25% (preservative), Citric Acid, Disodium EDTA)

This is a compounded Drug. Hospital/office use only. Not for resale. Adverse events may be reported to the FDA at www.fda.gov/medwatch and 1-800-FDA-1088

"Verified uniform dosing, so you can have consistent outcomes for your patients."



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Formulated with SuspendIt[®]

A USP Verified Excipient oral suspending vehicle

Formulated Without:

- Sugar
- Gluten
- Casein
- Dairy
- Soy
- Egg
- Nuts
- Alcohol
- Parabens
- Propylene Glycol
- Modified Food Starch
- Carbomer

✓ Eliminates staff exposure & time, and hazardous drug compounding resources for a commonly compounded oral medication.