Pharmacy Services

COMPOUND EVALUATION FORM					
Compound Name: LamoTRIgine	Container-closure system(s): tight light resistant container (i.e. amber plastic bottle)				
Strength: 1 mg/mL	Preservatives: Contained in Ora Blend				
Dosage Form: Oral Suspension	Beyond Use Date: 90 days				
Product Description: white suspension	Storage: Room Temperature				
Auxiliary Labels: Shake Well					
Quality control procedures (ex: pH test, etc.): Visual Inspection of final product					
Ingredients:					
LamoTRIgine 100 mg tablets Ora-BLEND (QS)					

lnst	truc	tions	for	pre	para	tion:

Directions:

- 1. Crush tablets in a mortar and triturate to a fine powder.
- 2. Wet powder with a minimal amount of vehicle and levigate to form a viscous, but smooth and uniform paste.
- 3. Continue adding vehicle geometrically, mixing well after each addition.
- 4. Transfer to a graduate.
- 5. Rinse mortar with vehicle, adding rinse to graduate, until almost final volume.
- 6. QS to final volume with vehicle. Stir well.

Calculations:

Ingredient Name/Strength/Form	QS	Quantity	Units
LamoTRIgine 100 mg tablets		2	tablets
Ora BLEND	X	200	mL

Equipment/ Supplies:

X Mortar and Pestle	X Stirring rod	☐ Ointment slab
X Graduated cylinder(s)	☐ Spatula	☐ Balance
☐ Weigh paper	X Amber (plastic/glass) bottle	☐ Ointment jar
X Counting tray	☐ Syringe	☐ Funnel

References:

Revised: TS 9/25



1. USP. Lamotrigine Compounded Oral Suspension. In: USP-NF. Rockville, MD: USP; June 2022

Revised: TS 9/25



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Revised: TS 9/25

