

Iwifin™ (eflornithine) Prior Authorization Form

Member Name: _____ **Date of Birth:** _____ **Member ID#:** _____

Drug Information

Pharmacy Billing (NDC: _____ **) Start Date (or date of next dose):** _____

Dose: _____ **Regimen:** _____

Pharmacy Information

Pharmacy NPI: _____ **Pharmacy Name:** _____

Pharmacy Phone: _____ **Pharmacy Fax:** _____

Prescriber Information

Prescriber NPI: _____ **Prescriber Name:** _____

Prescriber Phone: _____ **Prescriber Fax:** _____ **Specialty:** _____

Criteria

For Initial Authorization:

1. Please indicate the diagnosis and information:

Neuroblastoma

A. Is diagnosis high-risk neuroblastoma? Yes ___ No ___

B. Has member had at least a partial response to prior multiagent, multimodality therapy including anti-GD2 immunotherapy? Yes ___ No ___

C. Will eflornithine be used as a single agent to reduce the risk of relapse for a maximum of 2 years? Yes ___ No ___

D. Member's body surface area (BSA): _____ Date taken: _____

Other _____

Additional Information: _____

For Continued Authorization:

1. Date of last dose: _____

2. Does member have any evidence of progressive disease while on eflornithine? Yes ___ No ___

3. Has the member experienced any adverse drug reactions related to eflornithine therapy? Yes ___ No ___

If yes, please specify adverse reactions: _____

Additional Information: _____

Prescriber Signature: _____ **Date:** _____

I certify that the indicated treatment is medically necessary and all information is true and correct to the best of my knowledge. Please do not send in chart notes. Specific information will be requested if necessary. Failure to complete this form in full will result in processing delays.

PLEASE PROVIDE THE INFORMATION REQUESTED AND RETURN TO:

University of Oklahoma College of Pharmacy
Pharmacy Management Consultants
Product Based Prior Authorization Unit

Fax: 1-800-224-4014
Phone: 1-800-522-0114 Option 4

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