

The Board of Pharmacy has received notice of the following product recall:

NDC	Name	Strength	Size	Lot	Expires
51862-858-01	Carbidopa and Levodopa	25mg/250mg tablets	100 count bottle	FG11514	05/2021

Mayne Pharma is conducting a recall of one lot, as referenced above, of Carbidopa and Levodopa, 25mg/250mg Tablets, to the retail level. The lot is being recalled due to a foreign tablet being found in Carbidopa/Levodopa Tablets USP 25mg/250mg. This lot was distributed in the US market between 14 October 2019 and 06 January 2020.

The Board of Pharmacy strongly encourages pharmacies to immediately review their quality assurance and recall policies and procedures to determine if any corrective action is required.