Drugs@FDA: FDA Approved Drug Products

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New Drug Application (NDA): 209884
Company: NOVARTIS PHARMS CORP

- Medication Guide (https://www.accessdata.fda.gov/drugsatfda_docs/label/2019/209884s000lbl.pdf#page=22)

Products on NDA 209884

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Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	TE Code
MAYZENT	SIPONIMOD	0.25MG	TABLET;ORAL	Prescription	None
MAYZENT	SIPONIMOD	2MG	TABLET;ORAL	Prescription	None

Showing 1 to 2 of 2 entries

Approval Date(s) and History, Letters, Labels, Reviews for NDA 209884

Labels for NDA 209884