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New Drug Application (NDA): 209884

Company: NOVARTIS PHARMS CORP

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- [Medication Guide \(https://www.accessdata.fda.gov/drugsatfda_docs/label/2019/209884s0001b1.pdf#page=22\)](https://www.accessdata.fda.gov/drugsatfda_docs/label/2019/209884s0001b1.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

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

Labels for NDA 209884 ▾