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FDA Approved Drug Products

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Drug Details

Drug Name(s)	ADEMPAS
FDA Application No.	(NDA) 204819
Active Ingredient(s)	RIOCIGUAT
Company	BAYER HLTHCARE
Original Approval or Tentative Approval Date	October 8, 2013
Chemical Type	1 New molecular entity (NME)
Review Classification	P Priority review drug O Orphan drug

- [There are no Therapeutic Equivalents](#)
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Products on Application (NDA) #204819

Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
ADEMPAS	RIOCIGUAT	0.5MG	TABLET;ORAL	Prescription	No	None
ADEMPAS	RIOCIGUAT	1MG	TABLET;ORAL	Prescription	No	None
ADEMPAS	RIOCIGUAT	1.5MG	TABLET;ORAL	Prescription	No	None
ADEMPAS	RIOCIGUAT	2MG	TABLET;ORAL	Prescription	No	None
ADEMPAS	RIOCIGUAT	2.5MG	TABLET;ORAL	Prescription	Yes	None

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