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Drug Details

Drug Name(s)	AMELUZ
FDA Application No.	(NDA) 208081
Active Ingredient(s)	AMINOLEVULINIC ACID HYDROCHLORIDE
Company	BIOFRONTERA PHARMA AG
Original Approval or Tentative Approval Date	May 10, 2016
Chemical Type	3 New dosage form

- [There are no Therapeutic Equivalents](#)
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Products on Application (NDA) #208081

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

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLD	TE Code
AMELUZ	AMINOLEVULINIC ACID HYDROCHLORIDE	78MG	GEL;TOPICAL	Prescription	TBD  ¹¹	None

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