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**
- **Label Information**
- **Approval History, Letters, Reviews, and Related Documents**

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

Back to Top | Back to Previous Page | Back to Drugs@FDA Home

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