FDA Home[3] Drug Databases[4] Drugs@FDA[5]

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FAQ[6] | Instructions[7] | Glossary[8] | Contact Us[9]

Email Link

Start Over

Back to Reports[10]

Drug Details

Drug Name(s)	**IBUPROFEN LYSINE**
FDA Application No.	**(ANDA) 202402**
Active Ingredient(s)	**IBUPROFEN LYSINE**
Company	**EXELA PHARMA SCIENCE**
Original Approval or Tentative Approval Date	**March 30, 2016**

- **Therapeutic Equivalents**
- **Approval History, Letters, Reviews, and Related Documents**
- **Labels are not available**

Products on Application (ANDA) #202402
Click on a column header to re-sort the table:

Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status	RLDTE Code
IBUPROFEN LYSINE	IBUPROFEN LYSINE	EQ 20MG BASE/2ML (EQ 10MG BASE/ML)	INJECTABLE;INTRAVENOUS	Prescription	No AP

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

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