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Drug Application (NDA): 021920

Company: BIONPHARMA INC

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For Important Information from FDA

<http://www.fda.gov/Drugs/DrugSafety/PostmarketDrugSafetyInformationforPatientsandProviders/ucm110423.htm>

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Products on NDA 021920 ▼

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Drug Name	Active Ingredients	Strength	Dosage Form/Route	Marketing Status
NAPROXEN SODIUM	NAPROXEN SODIUM	EQ 200MG BASE	CAPSULE;ORAL	Over-the-counter

Showing 1 to 1 of 1 entries

Approval Date(s) and History, Letters, Labels, Reviews for NDA 021920 ▲

Final Approvals or Tentative Approvals

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Approval Date	Submission	Action Type	Submission Classification	Review Priority; Orphan Status
7/2006	ORIG-1	Approval	Type 3 - New Dosage Form	STANDARD

Showing 1 to 1 of 1 entries

Supplements

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Submission Date	Submission	Supplement Categories or Approval Type	Letters, Reviews, Labels
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Submission Date	Submission	Supplement Categories or Approval Type	Letters, Reviews, Labels
6/2017	SUPPL-19	Labeling-Container/Carton Labels	Label (PDF) (https://www.accessdata.fda.gov/drugsatfda_docs/nda/2017/suppl/021920Orig1s01.pdf) Letter (PDF) (https://www.accessdata.fda.gov/drugsatfda_docs/nda/2017/suppl/021920Orig1s01.pdf)
1/2013	SUPPL-18	Labeling-Package Insert	Label (PDF) (https://www.accessdata.fda.gov/drugsatfda_docs/nda/2013/suppl/021820Orig1s01.pdf) Letter (PDF) (https://www.accessdata.fda.gov/drugsatfda_docs/nda/2013/suppl/021820Orig1s01.pdf)
3/2012	SUPPL-17	Labeling-Container/Carton Labels	Letter (PDF) (https://www.accessdata.fda.gov/drugsatfda_docs/nda/2012/suppl/021720Orig1s01.pdf)
2/2010	SUPPL-13	Labeling-Container/Carton Labels	Letter (PDF) (https://www.accessdata.fda.gov/drugsatfda_docs/nda/2010/suppl/021320Orig1s01.pdf)
4/2009	SUPPL-11	Labeling	Label (PDF) (https://www.accessdata.fda.gov/drugsatfda_docs/nda/2009/suppl/021120Orig1s01.pdf) Letter (PDF) (https://www.accessdata.fda.gov/drugsatfda_docs/nda/2009/suppl/021120Orig1s01.pdf)
5/2009	SUPPL-10	Labeling	Letter (PDF) (https://www.accessdata.fda.gov/drugsatfda_docs/nda/2009/suppl/021020Orig1s01.pdf)

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Supplements for NDA 021920 

OTC Drugs with the Same Active Ingredient, Strength and Dosage Form/Route 