

## FDA Home<sup>3</sup> Drug Databases<sup>4</sup> Orange Book<sup>5</sup> Orange Book: Approved Drug Products with Therapeutic Equivalence Evaluations Search results from the "OB\_Rx" table for query on "203441."

Active Ingredient:	TEDUGLUTIDE RECOMBINANT				
Dosage Form;Route:	POWDER;SUBCUTANEOUS				
Proprietary Name:	GATTEX KIT				
Applicant:	NPS PHARMS INC				
Strength:	5MG/VIAL				
Application Number:	N203441				
Product Number:	001				
Approval Date:	Dec 21, 2012				
Reference Listed Drug	Yes				
RX/OTC/DISCN:	RX				
TE Code:					
Patent and Exclusivity Info for this product: View					

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FDA/Center for Drug Evaluation and Research Office of Generic Drugs Division of Labeling and Program Support Update Frequency: Orange Book Data - **Monthly** Generic Drug Product Information & Patent Information - **Daily** Orange Book Data Updated Through December 2015 Patent and Generic Drug Product Data Last Updated January 15, 2016

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#### Orange Book: Approved Drug Products with Therapeutic Equivalence Evaluations Patent and Exclusivity Search Results from query on Appl No 203441 Product 001 in the OB\_RX list.

## Patent Data

Appl No	Prod No	Patent No	Patent Expiration	Drug Substance Claim	Drug Product Claim	Patent Use Code	Delist Requested
N203441	001	5789379	Apr 14, 2016	Y	Y	U - 1320	
N203441	001	7056886	Sep 18, 2022		Y	U - 1320	
N203441	001	7847061	Nov 1, 2025			U - 1320	
N203441	001	9060992	Nov 1, 2025			U - 1320	

## **Exclusivity Data**

Appl No Prod No		Exclusivity Code	Exclusivity Expiration	
N203441	001	NCE	Dec 21, 2017	
N203441	001	ODE	Dec 21, 2019	

Additional information:

- 1. Patents are published upon receipt by the Orange Book Staff and may not reflect the official receipt date as described in 21 CFR 314.53(d)(5).
- 2. Patents listed prior to August 18, 2003 are flagged with method of use claims only as applicable and submitted by the sponsor. These patents may not be flagged with respect to other claims which may apply.

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