NDC	DRUG PROD DESC	INTRODUCED TO MARKET DATE
83457-0107-01	 Hyrimoz® (adalimuab-adcz) 80 mg/0.8 mL Prefilled Pen	 2/1/2024
83457-0108-01	Hyrimoz® (adalimuab-adcz) 20 mg/0.2 mL Prefilled Syringe	2/1/2024

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	Hyrimoz® (adalimuab-adcz)	<u> </u>
	Starter Pack 80 mg/0.8 ml-40	! !
83457-0112-01	mg/0.4 mL Prefilled Pen	2/1/2024
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	Hyrimog@ (adalimush adas) 00	
	Hyrimoz® (adalimuab-adcz) 80	1
83457-0113-01	mg/0.8 mL Prefilled Pen	2/1/2024
10 or 11 Digit NDC	This will include: Drug Trade Name,	Date Product was commercially
-	A .: Y 1: . C: .: 1 1 C:	-

Active Ingredient, Strength, and Size

Submissions Made To AGO.highcostprescriptiondrugs@vermont.gov

Key Used for Full Report Due 30 Days After Initial Notice

available

Cordavis Hyrimoz has being developed as a co-manufacturing biosimilar initiative. Hyrimoz exists on the market to-day under other manufaturer NDCs. There is currently no specific marketing and/or advertising budget or plan associated with this NDC. Cordavis is introducing the Hyrimoz NDC to create compention in the Biosimilar market and help drive down the overall cost of drugs. Cordavis Hyrimoz pricing strategy was to bring a Biosimilar to market at a price point that was at least 80% below the list price of the brand equivalent. \$ 2,600.00 Quivalent. N Cordavis Hyrimoz has being developed as a co-manufacturing biosimilar initiative. Hyrimoz exists on the market to-day lunder other manufacturer NDCs. There is currently no specific marketing and/or advertising budget or plan associated with this NDC. Cordavis is introducing the Hyrimoz NDC to create compention in the Biosimilar market and help drive down the overall cost of drugs. Cordavis Hyrimoz Pricing strategy was to bring a Biosimilar to market at a	WAC AT INTRODUCTION	MARKETING PRICING PLAN	MARKETING PRICING
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\$ 3,900.00	equivalent.	N

WAC on the date the Product was commercially available

A narrative description of the marketing Please respond with Yes or No and pricing plans used in the launch of the new prescription drug in the United States and internationally

ESTIMATED PATIENTS	BREAKTHROUGH THERAPY INDICATOR	PRIORITY REVIEW INDICATOR
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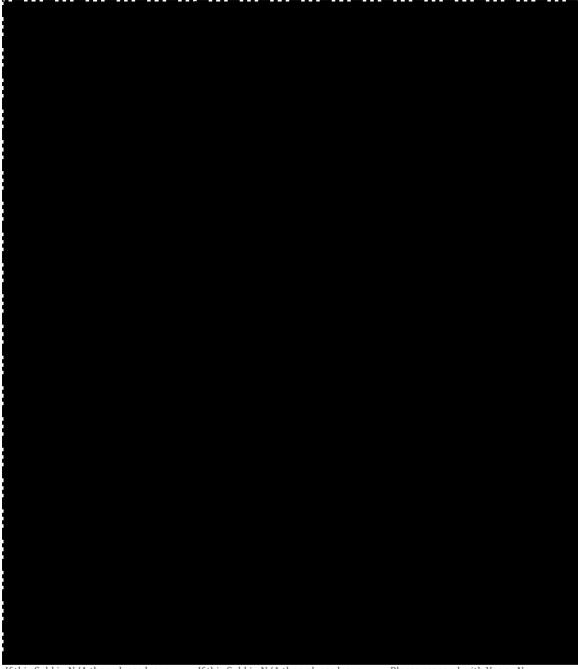
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e estimated number of patients in the nited States with a condition for which	Indicate whether the drug was granted breakthrough therapy designation or	Indicate whether the drug was granted breakthrough therapy designation or

the new prescription drug may be prescribed

breakthrough therapy designation or priority review by the federal Food and Drug Administration prior to approval

breakthrough therapy designation or priority review by the federal Food and Drug Administration prior to approval

ACQUISITION DATE	ACQUISITION PRICE	ACQUISITION PRICE NONPUBLIC

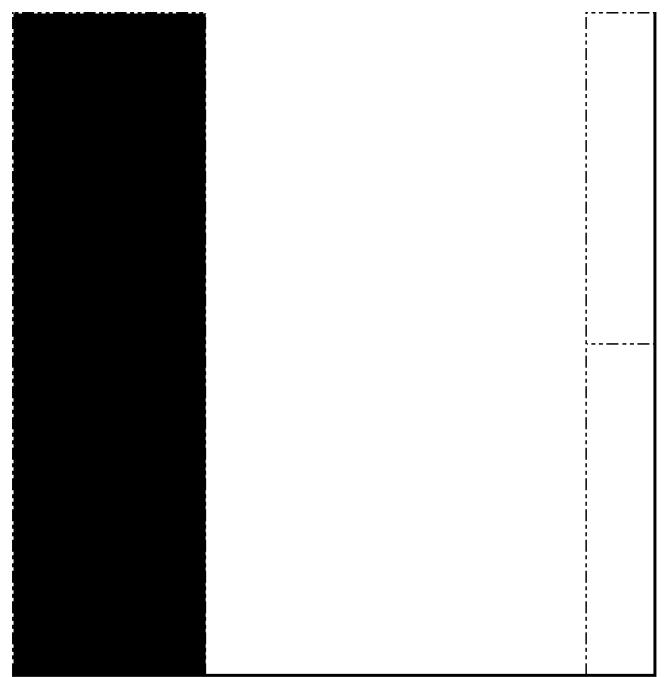


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Please respond with Yes or No

ACQUISITION PRICE COMMENT	GENERAL COMMENTS	



If this field in N/A then please leave blank