10. 01. 174	New Hampshire Medicaid Fee-for-Service ProgramPrior Authorization Drug Approval FormRoctavian™ (valoctocogene roxaparvovec-rvox)DATE OF MEDICATION REQUEST:																						
SE	SECTION I: PATIENT INFORMATION AND MEDICATION REQUESTED																						
LAST NAME:								FIRST NAME:															
MEDICAID ID NUMBER:								DATE OF BIRTH:															
										_			] _										
GENDER: Male Female																							
Drug Name:								Strength:															
Do	Dosing Directions:									Length of Therapy:													
SECTION II: PRESCRIBER INFORMATION																							
		AME:										FIRST		ME:									
																				<u> </u>			
SPECIALTY:								NPI NUMBER:															
PH	ION		BER:								_	FAX NUMBER:											
			_				_								_				] _				
SE	стіс	)N III: (		CAL	HIST	ORY						I			1		•			L			
1.	SECTION III: CLINICAL HISTORY  1. Does the patient have severe congenital factor VIII deficiency, confirmed by factor VIII activity <1 Yes No IU/dL testing?																						
2.	Have other bleeding disorders been ruled out?									No													
3.	. Is the patient AAV serotype 5 (AAV5) antibody negative as determined by an FDA-approved or 🛛 🗌 Yes 🗌								No														
	CLIA-compliant test?																						
4.											No												
5.	Does the patient have significant hepatic fibrosis (stage 3 or 4) or cirrhosis?										No												
6.	ls t	Is the patient on a stable dose of exogenous factor VIII for prevention of bleeding episodes?										No											
	a.	-							-			-										<u> </u>	
	a.	Negill	icii d	nu st	urt u	are.																	

(Form continued on next page.)

Fax to DHHS; medication is administered in inpatient setting: Phone: 1-603-271-9384 Fax: 1-603-314-8101





## New Hampshire Medicaid Fee-for-Service Program Prior Authorization Drug Approval Form

Roctavian<sup>™</sup> (valoctocogene roxaparvovec-rvox)

PATIENT LAST NAME:	PATIENT FIRST NAME:											
SECTION III: CLINICAL HISTORY												
7. Does the patient have a hypersensitivity to mannitol?	Yes No											
Has the patient received prior hemophilia adeno-associated virus vector-based gene therapy? 🛛 🗌 Yes 🗌 N												
9. Is the patient negative for factor VIII inhibitor titers on initial test or re-test?												
10. Is the patient received a bypass agent (e.g. Feiba)?												
11. Will the liver function be assessed after Roctavian™ dose according to a facility protocol?												
a. Attach copy of baseline liver function tests.												
12. Does the patient have any of the following:	Yes No											
<ul> <li>Hepatitis B</li> <li>Hepatitis C</li> <li>Non-alcoholic fatty liver disease</li> </ul>												
<ul> <li>Chronic alcohol consumption</li> <li>Non-alcoholic steatohepatitis</li> <li>Advanced age</li> </ul>												
13. If yes to question 12, will the patient have regular live fetoprotein elevation?	r ultrasounds and testing for alpha- Yes No											
14. Attach protocol for post-Roctavian™ monitoring.												

Please provide any additional information that would help in the decision-making process. If additional space is needed, please use a separate sheet.

I certify that the information provided is accurate and complete to the best of my knowledge and I understand that any falsification, omission, or concealment of material fact may subject me to civil or criminal liability.

PRESCRIBER'S SIGNATURE:	DATE:					
Facility where infusion to be provided:						
Medicaid Provider Number of Facility:						
Fax to DHHS; medication is administered in inpatient setting: Phone: 1-603-271-9384						
Fax: 1-603-314-8101	THERAPEUTICS					