

All required fields are purple and bolded

PATIENT	First Name	Middle Initial	Last Name	Suffix
	Date of Birth (mm/dd/yyyy)	Gender <input type="checkbox"/> Male <input type="checkbox"/> Female <input type="checkbox"/> Other		
	Address			Floor/Suite/ Unit
	City		State	ZIP Code
	Primary Phone	Mobile Phone <input type="checkbox"/> (same as primary)	Email	
	Preferred Method of Contact <input type="checkbox"/> Primary Phone <input type="checkbox"/> Mobile Phone <input type="checkbox"/> Email		Preferred Language: <input type="checkbox"/> English <input type="checkbox"/> Spanish <input type="checkbox"/> Other language (please specify)	
	Authorized Representative Name (if applicable)			Relationship to Patient
	Phone		Email	

PRESCRIBER	First Name		Last Name	
	Specialty		NPI Number	
	State License Number		Medicaid Number	
	Tax ID		Name of Institution/Practice	
	Address			Floor/Suite/Unit
	City		State	ZIP Code
	Phone	Fax	Email	
	Preferred Method of Contact <input type="checkbox"/> Phone <input type="checkbox"/> Fax <input type="checkbox"/> Email			
Primary Contact Name (if different from prescriber)				
Phone		Fax		
Email				

INSURANCE	Provide copies of all medical and prescription cards — front and back			
	<input type="checkbox"/> Patient has no insurance			
	Primary Medical Insurance Name			Insurance Phone
	Subscriber Name		Relationship to Patient	
	Member ID	Group	Plan Code	
	Prescription (PBM) Insurance Name			Insurance Phone
	Subscriber Name			
	Member ID	RxBIN	RxPCN	RxGROUP

Patient's Full Name	Date of birth (mm/dd/yyyy)
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CLINICAL AND LAB RESULTS	ICD Code:		
	<input type="checkbox"/> D66.0 Hereditary factor VIII deficiency (please specify below) <ul style="list-style-type: none"> <input type="checkbox"/> Classic hemophilia <input type="checkbox"/> Deficiency factor VIII (with functional defect) <input type="checkbox"/> Hemophilia NOS <input type="checkbox"/> Hemophilia A <input type="checkbox"/> Other diagnosis (Please specify) _____		
	REQUIRED LAB ELIGIBILITY RESULTS DOCUMENTATION		
	Before administration of ROCTAVIAN, the following baseline assessments for AAV5 antibodies and liver health are required and results may be requested by the patient's insurance provider. Please call BioMarin RareConnections at 1.833.762.8284, or your BioMarin representative, if you have questions about these tests including the required companion diagnostic (CDx)		
	Please Confirm Test Status:		
	AAV5 Antibody Test: AAV5 DetectCDx™	<input type="checkbox"/> Completed	
Liver Function Test: Alanine transaminase (ALT) Note to prescriber: Blood tests for liver function are included in the liver fibrosis assessment listed below	<input type="checkbox"/> Completed As part of liver fibrosis assessment blood draw	<input type="checkbox"/> Completed Via independent blood draw	<input type="checkbox"/> Not completed
Liver Fibrosis Assessment: via blood test (e.g., FibroTest®, FibroSURE® or similar) OR via liver elastography ultrasound (e.g., FibroScan®) Note to prescriber: Blood tests for liver fibrosis include liver function (ALT) results	<input type="checkbox"/> Completed Via blood draw	<input type="checkbox"/> Completed Via ultrasound	<input type="checkbox"/> Not completed
Patient allergies			
<input type="checkbox"/> NKDA <input type="checkbox"/> Yes (please list) _____ Concurrent medications _____			

INFUSION SITE	<input type="checkbox"/> Information provided in Prescriber section on first page		
	Infusion Site Name		
	Address		Floor/Suite/Unit
	City		State ZIP Code
	Infusion Site NPI		Infusion Site Contact (if available)
	Phone	Fax	Email

PRESCRIPTION	Current weight (kg)	Date weight measured (mm/dd/yyyy)
	ROCTAVIAN™ (valoctocogene roxaparvovec-rvox) is provided in 10 mL vials containing an extractable volume of no less than 8 mL (6 x 10 ¹³ vg). Dose volume is based on body weight. To calculate a patient's dose in milliliters (mL), multiply body weight in kg by 3. The multiplication factor 3 represents the per-kilogram dose (6 x 10 ¹³ vg/kg) divided by the amount of vector genomes per mL of the ROCTAVIAN solution (2 x 10 ¹³ vg/mL). To calculate number of vials to be thawed, divide patient's dose volume in mL by 8 and round up to the next whole number of vials.	
	$\frac{\text{Patient's weight (kg)}}{\text{Dose (mL)}} \times 3 = \text{number of vials required} / 8 =$	
	Directions: Administer _____ ml as a single intravenous infusion per manufacturer product labeling Dose: _____ vg Dispense (number of vials): _____	
		Refills: None
		NDC #: 68135-927-48

PRODUCT COORDINATION	<input type="checkbox"/> Ship-to-site for product (if different from infusion site) <input type="checkbox"/> (select if same as infusion site)		
	Ship-to-site Name		
	Address		Floor/Suite/Unit
	City		State ZIP Code
	Ship-to-site Contact Name		Phone Fax
	Email	Shipping Instructions	

PRESCRIBER DECLARATION	Prescriber Declaration: By signing below, I, as the prescribing physician, certify that the information provided on this form was completed by me or at my direction. I understand and agree that, as the Prescriber, I will comply with my state-specific prescription requirements such as e-prescribing, state-specific prescription form, fax language, etc. Non-compliance with state-specific requirements could result in outreach to me, as the Prescriber.	
	I verify that the patient and prescriber information contained in this enrollment form is complete and accurate to the best of my knowledge and that I have prescribed ROCTAVIAN based on my professional judgment of medical necessity. I have informed my patient of the resources available in the BioMarin RareConnections program and have confirmed my patient's (or their respective caregiver's) consent to enroll in the program. I have obtained all required patient permissions and have complied with all federal and state laws with respect to disclosures and release of the provided information to BioMarin Pharmaceutical Inc., BioMarin RareConnections, and its affiliates, agents, and contractors (collectively, "BioMarin", as well as to or between other service providers such as laboratories and pharmacies, and for the purposes described herein by any means allowed under applicable law.	
	I understand that the information provided herein will be used for the purposes of BioMarin to investigate and verify patient's insurance and coverage benefits, to contact this patient to help obtain a signed patient consent form and/or to refer the patient to or contact the patient for purposes of enrollment in a patient education program, verify patient's insurance coverage benefits for ROCTAVIAN and any related services, to coordinate the dispensing and delivery of ROCTAVIAN (including transmitting the prescription to the appropriate pharmacies) utilizing the patient's benefit plan, assist in initiating or continuing therapy, provide prior authorization and appeals information, verify eligibility for a co-pay program, and identify additional financial resources, provide me and my patient with other education and support associated with ROCTAVIAN, and for BioMarin internal business purposes such as conducting quality control, data analysis, and gathering feedback to improve patient support and resources.	
Prescriber's Signature. Please make a selection		
Prescriber's Signature/Dispense as Written (no stamps or initials) Date _____		
Prescriber's Signature/Substitution Permitted (no stamps or initials) Date _____		

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