

State of Oklahoma Oklahoma Health Care Authority

Intravenous Iron Therapy Prior Authorization Request

Member Name:	Date of Birti	h: Member ID#:
	Drug Info	rmation
Medication Name:		Strength:
Dose:	Regimen:	Start Date:
HCPCS code:B		
Billing Provider Information		
Provider NPI:	Provider N	Name:
Provider Phone:	Provider F	Fax:
	Prescriber Ir	nformation
Prescriber NPI:	Prescriber	Name:
Prescriber Phone:	Prescriber	· Fax:
Criteria		
 Please indicate the diagnosis for lron Deficiency Anemia lron Deficiency Anemia who other: If member has Chronic Kidney I a. Stage of Chronic Kidney I b. Is the member on di Please submit laboratory results Has the member had a trial of one a. If "Yes", please proving it. Dates of the ora ii. Member's response b. If "No", please proving the member: 	vith Chronic Kidney Disease Disease, please provide the form of the provide the form of the provide the form of the provide the following: all iron therapy trial: onse to oral iron therapy: de a patient-specific, clinically	ollowing information: emia
6. Has the member had a trial of In a. If "Yes", please proving it. Dates of the Iron ii. Member's response b. If "No", please proving member:	ron Dextran? Yes No ride the following: on Dextran trial: onse to Iron Dextran: de a patient-specific, clinically	(Iron Dextran is available without prior authorization) significant reason why Iron Dextran is not appropriate for the
Prescriber Signature: I certify that the indicated treatment is medically necessary and all information is true and correct to the best of my knowledge. Pease do not send in chart notes. Specific information will be requested if necessary. Failure to complete this form in full will result in processing delays.		

PLEASE PROVIDE THE INFORMATION REQUESTED AND RETURN TO:

University of Oklahoma College of Pharmacy Pharmacy Management Consultants Product Based Prior Authorization Unit

> Fax: 1-800-224-4014 Phone: 1-800-522-0114 Option 4

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