

State of Oklahoma Oklahoma Health Care Authority Bosulif® (Bosutinib) Prior Authorization Form

Member Name:	Date of Birth:	Member ID#:	
	Drug Information		
Pharmacy billing (NDC: Dose:) Start Date (or date of next dose): Regimen:		
	Billing Provider Inform	ation	
Provider NPI:	Provider Name:		
Provider Phone:	Provider Fa	Provider Fax:	
	Prescriber Informati	ion	
Prescriber NPI: Prescriber Name:			
Prescriber Phone:	Prescriber Fax:	Specialty:	
	Criteria		
B. Relapsed/re C. Bosutinib us D. Bosutinib us Yes E. E255K/V, F3 Chronic Myeloid Leu A. Chronic, accele B. Newly diagnose Yes Cother, please provide	Chromosome Positive (Ph+)? Yes fractory ALL? Yes No ed as a single-agent? Yes No led in combination with an induction re No 817L/VI/C, F359V/C/I, T315A, or Y253 likemia (CML) erated, or blast phase CML? Yes ed or resistant/intolerant to other Tyrose	egimen not previously given? 3H mutations? Yes No _ No sine Kinase Inhibitors (TKIs)?	
3. Has the member experience of the second o	idence of progressive disease while of the decision of the dec	bosutinib therapy? Yes No	
I certify that the indicated tre	atment is medically necessary and	Date: d all information is true and correct to the	
best of my knowledge. Please do not send in chart no	tes. Specific information will be reque	ested if necessary. Failure to complete this	

PLEASE PROVIDE THE INFORMATION REQUESTED AND RETURN TO:

form in full will result in processing delays.

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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