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\checkmark	fuse@fuseinfusion.com

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Cana	kinuma	b (I	laris)
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Canakinumab (Ilaris) Provider Order Form	Date:				
PATIENT INFORMATION					
Name: ICD-10 code (required): □ NKDA Allergies:	DOB: SEX: M				
REFERRA	AL STATUS				
□New Referral □Referral Renewal □Medication/Order C	hange Benefits Verification Only Discontinuation Order				
PHYSICIA	N INFORMATION				
Referral Coordinator Name:	Referral Coordinator Email:				
Ordering Provider:	Provider NPI:				
Referring Practice Name:	Phone: Fax:				
Practice Address:	City: State: Zip Code:				
OBSERVATION (PLEASE SELECT BELOW) Patient is required to stay for 30 minutes observation period Patient is NOT required to stay for observation time Other: SPECIAL INSTRUCTIONS	THERAPY ADMINISTRATION Canakinumab (Ilaris) For Stills Disease including Adult Onset Stills Disease and Systemic Juvenile Idiopathic Arthritis. □ 4mg/kg (with a max of 300mg) for patients with a body weight greater than or equal to 7.5kg subcutaneous every 4 weeks For Cryopyrin-Associated Periodic Syndromes (CAPS) □ 150mg for patients with body weight greater than 40kg subcutaneous every 8 weeks □ 2mg/kg for patients with body weight greater than or equal to 15kg and less than or equal to 40kg subcutaneous every 8 wks For Tumor Necrosis Factor Receptor Associated Periodic Syndrome, Hyperimmunoglobulin D Syndrome/Mevalonate Kinase Deficiency, Familial Mediterranean Fever Body weight less than or equal to 40kg □ 2mg/kg subcutaneous every 4 weeks □ 4mg/kg subcutaneous every 4 weeks □ 4mg/kg subcutaneous every 4 weeks □ 500mg subcutaneous every 4 weeks □ 300mg subcutaneous every 4 weeks □ 300mg subcutaneous every 4 weeks □ 300mg subcutaneous every 4 weeks - consider if clinical response not adequate. Refills:□ Zero / □ for 12 months / □ (if not indicated order will expire one year from date signed)				
NOTES/ADDITIONAL COMMENTS:					
ORDERING PROVIDER Signature X	Date				

Provider _____ Phone ____ Fax _____