

Member Name:				
Member Date of Birth:				
Member UMI:				
Requesting Physician's Name:	NPI Number:			
Requesting Physician's Address:				
Office Contact: Phone #:	Fax #:			
Facility:	Facility NPI Number:			
Facility's Address:				
Date of Service:				
Diagnosis Code(s):				
DRUG INFORMATION (please select one)				
PREFERRED PRODUCTS	NON-PREFERRED**			
 Neulasta (J2506) Fulphila (Q5108) 	Udenyca (Q5111) Stimufend () Nyvepria (Q5120) Fylnetra () Rolvedon ()			
└	**A non-preferred product will be considered when the member has documented therapy failure after an adequate therapeutic trial of a preferred product, or the preferred product has not been tolerated or is contraindicated ** <u>Medicare members</u> currently established on a non- preferred therapy are not required to try a preferred option			
 Ziextenzo (Q5120) What is the member's cancer diagnosis and staging? 	member has documented therapy failure after an adequate therapeutic trial of a preferred product, or the preferred product has not been tolerated or is contraindicated ** <u>Medicare members</u> currently established on a non-			

Please verify member's eligibility and benefits through the health plan

pg. 1

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3.	What is the member's complete chemo regimen?	
4.	Is the member considered to be at low, intermediate, or high risk for febrile neutropenia?	🗆 Low 🗆 Intermediate 🗆 High
5.	Is the member at an increased risk for febrile neutropenia due to any of the following reasons?	 Persistent neutropenia (ANC of 1500/mm3 or less) History of febrile neutropenia Prior exposure to chemotherapy or radiation Bone marrow involvement by tumor Recent surgery and/or open wounds Liver or renal dysfunction Age > 65 years receiving full chemo dose intensity Comorbidities that can increase risk of serious infection Other:

Please attach all pertinent clinical information		
	Attached:	YES NO

Please verify member's eligibility and benefits through the health plan

pg. 2

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