



# Part B Prior Authorization Step Therapy Guidelines

Castleman's Disease

Sylvant (siltuximab) J2860 Prior Authorization Request

Medicare Part B Form

Instructions: \* Indicates required information – Form may be returned if required information is not provided. Please fax this request to the appropriate fax number listed at the bottom of the page.

□ Standard Request– (72 Hours)				<b>Urgent Request</b> (standard time frame could place the member's life, health or ability in serious jeopardy)							
	Date Requested										
	Requestor Clinic name:			Phone / Fax					/ Fax		
MEMBER INFORMATION											
*Nar	me:	* [	D#:*DOB:								
PRESCRIBER INFORMATION											
*Name: DMD DFNP							□PA	*Phone	9:		
*Address:*F							*Fax:				
DISPENSING PROVIDER / ADMINISTRATION INFORMATION											
*Name: Phone:											
*Address: Fax:											
PROCEDURE / PRODUCT INFORMATION											
нс	PC Code	Name of Drug	Dos	e (	(Wt:	kg	Ht:	)	Frequency	End Date if known	
										KIIOWII	
□ Self-administered □ Provider-administered □ Home Infusion											
Chart notes attached. Other important information:											
Diagnosis: ICD10: Description:											
□ Provider attests the diagnosis provided is an FDA-Approved indication for this drug											
CLINICAL INFORMATION											
<ul> <li>New Start or Initial Request: (Clinical documentation required for all requests)</li> <li>Provider has reviewed the attached "Criteria for Approval" and attests the member meets         ALL required PA criteria.         If not, please provide clinical rationale for formulary exception:     </li> </ul>											
□ Continuation Requests: (Clinical documentation required for all requests)											
Patient had an <u>adequate response</u> or <u>significant improvement</u> while on this medication. If not, please provide clinical rationale for continuing this medication:											
ACKNOWLEDGEMENT											
Request By (Signature Required):											



## Prior Authorization Group - Orphan Drug: Castleman's Disease PA

Drug Name(s): SYLVANT SILTUXIMAB

Criteria for approval of Non-Formulary/Preferred Drug:

- 1. Prescribed for an approved FDA diagnosis (as listed below):
- 2. Member does not have any clinically relevant contraindications, or CMS/Plan exclusions, to the requested drug.
- If the member meets all these criteria, they may be approved by the Plan for the requested drug.
- Quantity limits and Tiering will be determined by the Plan.
- Continuation Requests: Provider must verify continued clinical benefit in confirmatory trial(s).

Exclusion Criteria: N/A

Prescriber Restrictions: N/A

Coverage Duration: Approval will be for 12 months

#### FDA Indications:

Sylvant Multicentric Castleman's disease, In patients known to be HIV-negative and human herpesvirus-8-negative

Off-Label Uses: N/A

Age Restrictions: Safety and efficacy not established in pediatric patients

### **Other Clinical Consideration:**

Concomitant use: Do not administer live vaccines

#### **Resources:**

https://www.micromedexsolutions.com/micromedex2/librarian/CS/280669/ND\_PR/evidencexpert/ND\_P/evidencexpert/ /DUPLICATIONSHIELDSYNC/7AC27D/ND\_PG/evidencexpert/ND\_B/evidencexpert/ND\_AppProduct/evidencexpert/ND\_T /evidencexpert/PFActionId/evidencexpert.GoToDashboard?docId=930940&contentSetId=100&title=Siltuximab&service sTitle=Siltuximab&brandName=Sylvant&UserMdxSearchTerm=sylvant&=null#