



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)				Urgent Request (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											
 New Start or Initial Request: (Clinical documentation required for all requests) 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: 											
□ 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#