

Jemperli Endometrial Cancer / Solid Tumor Jemperli (Dostarlimab-gxly) J9272 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.

	Date Requested							
	Requesto	r	Clinic name: _		Phone		/ Fax	
MEMBER INFORMATION								
*Name:			*	*ID#:		*DOB:		
PRESCRIBER INFORMATION								
*Name:				□MD □FNP □DO □NP □PA *Phone:				
*Address:				*Fax:				
DISPENSING PROVIDER / ADMINISTRATION INFORMATION								
*Name: Phone:								
	lress:			Fax:				
PROCEDURE / PRODUCT INFORMATION								
нс	PC Code	Name of Drug	□ Self-administered	Dose (Wt: _	kg Ht:)	Frequency	End Date if known
Chart notes attached. Other important information:								
Diagnosis: ICD10: Description:								
\Box 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) Provider has reviewed the attached "Criteria for Continuation" and attests the member meets ALL required PA Continuation criteria. 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):								

For questions or assistance, please contact Customer Service at 1-877-672-8620, daily, 8am – 8pm (PST) (TTY users should call 1-800-735-2900).



Prior Authorization Group – Jemperli PA

Drug Name(s): JEMPERLI

DOSTARLIMAB-GXLY

Criteria for approval of Non-Formulary/Preferred Drug:

- 1. Prescribed for an approved FDA diagnosis (as listed below):
- 2. Drug is being used appropriately per NCCN GUIDELINES, MCG GUIDELINES, CMS recognized compendia, authoritative medical literature, evidence-based guidelines and/or accepted standards of medical practice.
- 3. 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.

Exclusion Criteria:

• N/A

Prescriber Restrictions:

• Oncology or related specialty

Coverage Duration:

Approval will be for 6 months

FDA Indications:

- Endometrial cancer, Recurrent or advanced, dMMR disease, that has progressed on or following prior treatment with a platinum-containing regimen in any setting, not candidates for curative surgery or radiation
- Solid tumor, Recurrent or advanced, dMMR disease, that have progressed on or following prior treatment and who
 have no satisfactory alternative options

Off-Label Uses:

• N/A

Age Restrictions:

Safety and effectiveness have not been established in pediatric patients

Other Clinical Consideration:

- Per NCCN Guidelines, Jemperli may be appropriate for the following diagnoses when used in patients whose cancer is progressing on or following prior treatment and who have no satisfactory alternative treatment options:
 - o Ampullary Adenocarcinoma
 - o Breast Cancer
 - Colorectal Cancer
 - o Esophageal, esophagogastric Junction and Gastric cancer
 - o Occult Primary Cancer
 - o Ovarian Cancer
 - Small Bowel Adenocarcinoma

Resouces:

https://www.micromedexsolutions.com/micromedex2/librarian/CS/982351/ND_PR/evidencexpert/ND_P/evidencexpert/DUPLICATIONSHIELDSYNC/5C61D3/ND_PG/ evidencexpert/ND_B/evidencexpert/ND_AppProduct/evidencexpert/ND_T/evidencexpert/PFActionId/evidencexpert.GoToDashboard?docId=933289&contentSetId= 100&title=Dostarlimab-gxly&servicesTitle=Dostarlimab-gxly&brandName=Jemperli&UserMdxSearchTerm=Jemperli&=null#

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