

Perjeta (pertuzumab) J9306

Please provide the information below, please print your answer, attach supporting documentation, sign, date, and return to our office as soon as possible to expedite this request.

Without this information, we may deny the request in 30 days.

Date of request	Patient	Date of birth	ProviderOne client ID		
Prescriber	Billing provider NPI number	Telephone number	FAX number		
Drug/strength/dose (Procedu	ure/HCPC Code: 19306)				
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Does patient have HER2-Positive disease?					
2. What is patient's diagnos Metastatic breast can					
Operable early stage, locally advanced, or inflammatory breast cancer					
Other (please specify):					
Must provide justification for off-label use					
3. What is prescribed regimen:					
Perjeta + trastuzumab (Herceptin) + docetaxel Perjeta + trastuzumab (Herceptin)					
Other (please specify):					
4. Is Perjeta being used as a first line treatment?					
If no, mark all that apply:					
Patient received prior anti-HER2 therapy. Indicate which therapy:					
_					
Patient received prior chemotherapy for metastatic disease. Indicate which therapy:					
Patient has been previously treated with chemotherapy and trastuzumab.					
5. Please provide dosing schedule (select all that apply):					
☐ Metastatic breast cancer: 840mg initial dose, followed by 420mg every 3 weeks until disease					
progression or intolerable	e side effects.				
☐ Neoadjuvant treatment of locally advanced, inflammatory, or early stage breast cancer:					
840mg initial dose, followed by 420mg every 3 weeks for total of 6 doses					
Other: Dosing schedule:					
Must provide justification and documentation for off-label dosing					

6. For patients who have already been taking Perjeta:

Metastic disease:

- Has there been any evidence of disease progression while on Perjeta?
- Has the patient had any intolerable side effects?

Neoadjuvant use:

- How many cycles has the patient received?
 Dates they were received:
- Has the patient had any intolerable side effects?

Chart notes/progress notes required with request

Prescriber's signature	Prescriber's specialty	Date

A typed and completed *General Authorization for Information* form (13-835) must be attached to your request.

Fax to: 1-866-668-1214