



Trodelvy™ (sacituzumab govitecan-hziy) Medication Precertification Request

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(All fields must be completed and legible for precertification review.)

Aetna Precertification Notification

Phone: 1-866-752-7021

FAX: 1-888-267-3277

For Medicare Advantage Part B:

Phone: 1-866-503-0857

FAX: 1-844-268-7263

Please indicate: Start of treatment: Start date ____ / ____ / ____
 Continuation of therapy, Date of last treatment ____ / ____ / ____

Precertification Requested By: _____ **Phone:** _____ **Fax:** _____

A. PATIENT INFORMATION

| | | | | | |
|--|--|---|-------|--------------------|-------------|
| First Name: | | Last Name: | | DOB: | |
| Address: | | | City: | | State: ZIP: |
| Home Phone: | | Work Phone: | | Cell Phone: Email: | |
| Patient Current Weight: _____ lbs or _____ kgs | | Patient Height: _____ inches or _____ cms | | Allergies: | |

B. INSURANCE INFORMATION

| | | | |
|--|--|--|--|
| Aetna Member ID #: _____ | | Does patient have other coverage? <input type="checkbox"/> Yes <input type="checkbox"/> No | |
| Group #: _____ | | If yes, provide ID#: _____ Carrier Name: _____ | |
| Insured: _____ | | Insured: _____ | |
| Medicare: <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, provide ID #: | | Medicaid: <input type="checkbox"/> Yes <input type="checkbox"/> No If yes, provide ID #: | |

C. PRESCRIBER INFORMATION

| | | | | | |
|--|--|------------|----------------------|--|-------------|
| First Name: | | Last Name: | | (Check One): <input type="checkbox"/> M.D. <input type="checkbox"/> D.O. <input type="checkbox"/> N.P. <input type="checkbox"/> P.A. | |
| Address: | | | City: | | State: ZIP: |
| Phone: | | Fax: | | St Lic #: NPI #: DEA #: UPIN: | |
| Provider Email: | | | Office Contact Name: | | Phone: |
| Specialty (Check one): <input type="checkbox"/> Oncologist <input type="checkbox"/> Other: _____ | | | | | |

D. DISPENSING PROVIDER/ADMINISTRATION INFORMATION

| | | | |
|--|---|--|--|
| Place of Administration: | | Dispensing Provider/Pharmacy: Patient Selected choice | |
| <input type="checkbox"/> Self-administered | <input type="checkbox"/> Physician's Office | <input type="checkbox"/> Physician's Office | <input type="checkbox"/> Retail Pharmacy |
| <input type="checkbox"/> Outpatient Infusion Center | Phone: _____ | <input type="checkbox"/> Specialty Pharmacy | <input type="checkbox"/> Other: _____ |
| Center Name: _____ | | Name: _____ | |
| <input type="checkbox"/> Home Infusion Center | Phone: _____ | Address: _____ | |
| Agency Name: _____ | | Phone: _____ Fax: _____ | |
| <input type="checkbox"/> Administration code(s) (CPT): _____ | Address: _____ | TIN: _____ PIN: _____ | |

E. PRODUCT INFORMATION

Request is for: Trodelvy (sacituzumab govitecan-hziy) Dose: _____ **Frequency:** _____

F. DIAGNOSIS INFORMATION - Please indicate primary ICD code and specify any other where applicable.

Primary ICD Code: _____ **Secondary ICD Code:** _____ **Other ICD Code:** _____

G. CLINICAL INFORMATION - Required clinical information must be completed in its entirety for all precertification requests.

For Initiation Requests (clinical documentation required for all requests):

Breast cancer
Please indicate the clinical setting the requested drug will be used: Recurrent disease Metastatic disease
 Unresectable locally advanced disease Other

Yes No Has the patient received at least two prior therapies, at least one of them for metastatic disease?
 Yes No Unknown Does the patient have a diagnosis of triple-negative breast cancer confirmed by the breast cancer cells testing negative for ALL of the following receptors: A) human epidermal growth factor receptor 2 (HER2), B) estrogen, and C) progesterone?

Urothelial cancer (UC)
Please indicate the clinical setting the requested drug will be used: Locally advanced disease Metastatic disease Other

Yes No Has the patient received a platinum-containing chemotherapy (e.g., cisplatin, carboplatin)?
 Yes No Has the patient received either a programmed death receptor-1 (PD-1) or a programmed death-ligand 1 (PD-L1) inhibitor?
Please explain:
 a programmed death receptor-1 (PD-1) inhibitor (e.g., Keytruda, Opdivo)
 a programmed death-ligand 1 (PD-L1) inhibitor (e.g., Bavencio, Tecentriq)

For Continuation Requests (clinical documentation required for all requests):

Yes No Is there evidence of unacceptable toxicity or disease progression on the current regimen?

H. ACKNOWLEDGEMENT

Request Completed By (Signature Required): _____ **Date:** ____ / ____ / ____

Any person who knowingly files a request for authorization of coverage of a medical procedure or service with the intent to injure, defraud or deceive any insurance company by providing materially false information or conceals material information for the purpose of misleading, commits a fraudulent insurance act, which is a crime and subjects such person to criminal and civil penalties.

The plan may request additional information or clarification, if needed, to evaluate requests.