



MEDICARE FORM

Signifor LAR (pasireotide) Medication Precertification Request

Page 1 of 2

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

For Michigan MMP:

FAX: 1-844-241-2495

PHONE: 1-855-676-5772

For other lines of business:

Please use other form

Note: Signifor LAR is non-preferred for acromegaly. The preferred products are Sandostatin LAR and Somatuline Depot.

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"/> Endocrinologist <input type="checkbox"/> Other: _____ | | | | | |

D. DISPENSING PROVIDER/ADMINISTRATION INFORMATION

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

E. PRODUCT INFORMATION

Request is for: Signifor LAR (pasireotide) 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):

Note: Signifor LAR is non-preferred for acromegaly. The preferred products are Sandostatin LAR and Somatuline Depot.

- Yes No Has the patient had prior therapy with Signifor LAR within the last 365 days?
 Yes No Has the patient had a trial and failure, intolerance, or contraindication to any of the following? (select all that apply)
 Sandostatin LAR (octreotide acetate) Somatuline Depot (lanreotide)

Please explain if there are any other medical reason(s) that the patient cannot use any of the following preferred products when indicated for the patient's diagnosis? (select all that apply)

- Sandostatin LAR (octreotide acetate) Somatuline Depot (lanreotide)

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| | | | |
|--------------------|-------------------|---------------|-------------|
| Patient First Name | Patient Last Name | Patient Phone | Patient DOB |
|--------------------|-------------------|---------------|-------------|

G. CLINICAL INFORMATION (continued) – Required clinical information must be completed in its entirety for all precertification requests.

Acromegaly

Please indicate the patient's pretreatment IGF-1 (insulin-like growth factor 1) level compared to the laboratory's reference normal range based on age and/or gender: IGF-1 level is higher than the laboratory's normal range IGF-1 level is lower than the laboratory's normal range
 IGF-1 level falls within the laboratory's normal range

Yes No Has the patient had an inadequate or partial response to surgery?
 ↳ Yes No Is there a clinical reason why the patient has not had surgery?

Cushing's syndrome/disease

Yes No Did the patient have surgery that was not curative?
 ↳ Yes No Is the patient a candidate for surgery?

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

Acromegaly only:

Please indicate how the patient's IGF-1 (insulin-like growth factor 1) level changed since initiation of therapy:
 IGF-1 level has increased IGF-1 level has decreased or normalized IGF-1 level has not changed

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.