

Multiple myeloma newly diagnosed monotherapy – lenalidomide – initial authority application

Online services



Requesting PBS Authorities online provides an immediate assessment in real time.

For more information and how to access the **Online PBS Authorities** system, go to servicesaustralia.gov.au/hppbsauthorities

When to use this form

Use this form to apply for **initial** PBS-subsidised lenalidomide as monotherapy treatment for patients with newly diagnosed multiple myeloma.

Important information

Initial applications to start PBS-subsidised treatment can be made in real time using the **Online PBS Authorities** system or in writing and must include sufficient information to determine the patient's eligibility according to the PBS criteria.

Under no circumstances will phone approvals be granted for multiple myeloma **initial** authority applications.

The information in this form is correct at the time of publishing and may be subject to change.

Continuing treatment

This form is **ONLY** for **initial** treatment.

After an authority application for initial treatment has been approved, applications for **continuing** treatment can be made in real time using the **Online PBS Authorities** system or by phone. Call 1800 700 270 Monday to Friday, 8 am to 5 pm local time.

Call charges may apply.

Section 100 arrangements for lenalidomide

This item is available to a patient who is attending:

- an approved private hospital
- a public participating hospital, **or**
- a public hospital

and is:

- a day admitted patient
- a non-admitted patient, **or**
- a patient on discharge.

This item is not available as a PBS benefit for in-patients of a public hospital.

The hospital name and provider number must be included in this authority form.

Treatment specifics

Patients receiving lenalidomide under the PBS listing must be registered in the risk management program relevant for the brand of lenalidomide being prescribed and dispensed.

For more information

Go to servicesaustralia.gov.au/healthprofessionals

Current diagnostic reports

- 14** Nomination of which disease activity parameters will be used to assess response:

Results for (a) or (b) or (c) should be provided for all patients. Where the patient has oligo-secretory or non-secretory multiple myeloma, either (c) or (d) or if relevant (e), (f) or (g) evidence should be provided.

- a) the level of serum M protein
Provide the current level of serum M protein

Date of current diagnostic report (DD MM YYYY)

Unique identifying number/code or provider number
- b) Bence-Jones proteinuria
Provide the current results of 24-hour urinary light chain M protein excretion

Date of current diagnostic report (DD MM YYYY)

Unique identifying number/code or provider number
- c) the serum level of free kappa and lambda light chains
Provide the current results of the serum level

Date of current diagnostic report (DD MM YYYY)

Unique identifying number/code or provider number
- d) a bone marrow aspirate or trephine
Provide the current percentage of plasma cells in a bone marrow aspirate or on biopsy

Date of current diagnostic report (DD MM YYYY)

Unique identifying number/code or provider number
- e) if present, the size and location of lytic bone lesions
Provide details

- f) if present, the size and location of all soft tissue plasmacytomas by clinical or radiographic examination (that is, MRI or CT-scan)

Provide details

If applicable, provide the date of current radiographic report (DD MM YYYY)

Unique identifying number/code or provider number

- g) if present, the level of hypercalcaemia, corrected for albumin concentration.

Provide the current calcium corrected for albumin concentration

Date of current diagnostic report (DD MM YYYY)

Unique identifying number/code or provider number

Checklist

- 15**  The relevant attachments need to be provided with this form.

- The completed authority prescription form(s).

Privacy notice

- 16** Personal information is protected by law (including the *Privacy Act 1988*) and is collected by Services Australia for the purposes of assessing and processing this authority application. Personal information may be used by Services Australia, or given to other parties where the individual has agreed to this, or where it is required or authorised by law (including for the purpose of research or conducting investigations). More information about the way in which Services Australia manages personal information, including our privacy policy, can be found at servicessaustralia.gov.au/privacy

Prescriber's declaration

17 I declare that:

- I am aware that this patient must meet the criteria listed in the current Schedule of Pharmaceutical Benefits to be eligible for this medicine.
- I have informed the patient that their personal information (including health information) will be disclosed to Services Australia for the purposes of assessing and processing this authority application.
- I have attached the completed authority prescription form(s) and the relevant attachments as specified in the Pharmaceutical Benefits Scheme restriction.
- the information I have provided in this form is complete and correct.

I understand that:

- giving false or misleading information is a serious offence.

Prescriber's signature

Date (DD MM YYYY)

<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>	<input type="text"/>
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Returning this form

Return this form and any supporting documents:

- **online**, upload this form, the authority prescription form(s) and any relevant attachments through Health Professional Online Services (HPOS) at **servicesaustralia.gov.au/hpos**
- **or**
- by post, send this form, the authority prescription form(s) and any relevant attachments to:

Services Australia
Complex Drugs Programs
Reply Paid 9826
HOBART TAS 7001