

Juvenile idiopathic arthritis - for adult patients with onset prior to age 18 - continuing authority application

Online PBS Authorities



You do not need to complete this form if you use the **Online PBS Authorities** system.

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 **continuing** PBS-subsidised biological medicines for an adult patient with a documented history of juvenile idiopathic arthritis with onset prior to the age of 18 years.

Important information

Continuing authority applications 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.

Applications for **balance of supply** 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.

Under no circumstances will phone approvals be granted for juvenile idiopathic arthritis for adult patients with onset prior to age 18 **continuing** authority applications.

Where the term 'biological medicine' appears, it refers to adalimumab, etanercept and tocilizumab.

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

Continuing treatment

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

The patient remains eligible to receive continuing treatment providing they continue to sustain a response to treatment.

Section 100 arrangements for tocilizumab i.v.

This item is available to a patient who is attending:

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

and is a:

- day admitted patient
- non-admitted patient, **or**
- 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 form.

Treatment specifics

The assessment of the patient's response to the course of treatment must be conducted within the time frame specified in the restriction. Where a demonstration of response is not conducted within the required time frame, the patient will be deemed to have failed treatment with that particular PBS-subsidised biological medicine.

A patient who has experienced a serious adverse reaction of a severity necessitating permanent treatment withdrawal is not considered to have failed treatment with that particular PBS-subsidised biological medicine.

For more information

Go to servicesaustralia.gov.au/healthprofessionals

or

an ESR reduced by at least 20% from ESR provided at baseline

Baseline ESR

Current ESR

Date of test (DD MM YYYY)

and/or

a C-reactive protein (CRP) level \leq 15mg/L

CRP

Date of test (DD MM YYYY)

or

an CRP reduced by at least 20% from CRP provided at baseline

Baseline CRP

Current CRP

Date of test (DD MM YYYY)

14 The patient has demonstrated an adequate response to treatment with this drug evidenced by:

an active joint count of < 10 active (swollen and tender) joints

Current active joint count

Date (DD MM YYYY)

or

an active joint count reduced by at least 50% from baseline

Baseline total active joint count

Current active joint count

Date (DD MM YYYY)

or

a major active joint count of \leq 2

Current major joint count

Date (DD MM YYYY)

or

at least a 50% reduction in the number of the following major active joints, down from at least 4: elbow, wrist, knee and/or ankle, and/or shoulder, cervical spine and/or hip

Baseline major active joint count

Current major joint count

Date (DD MM YYYY)

Where a patient has at least 4 active major joints and less than 20 total active joints at baseline, assessment of the major joints only will be used for all continuing applications.

Checklist

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

Details of the proposed prescription(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/privacypolicy

Prescriber's declaration

You do not need to **sign** the declaration if you complete this form using Adobe Acrobat Reader and return this form through Health Professional Online Services (HPOS) at servicesaustralia.gov.au/hpos

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 provided details of the proposed prescription(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.

I have read, understood and agree to the above.

Date (DD MM YYYY) (you **must** date this declaration)

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Prescriber's signature (**only** required if returning by post)



Returning this form

Return this form, details of the proposed prescription(s) and any relevant attachments:

- **online** (no signature required), upload through HPOS at servicesaustralia.gov.au/hpos
or
- by post (signature required) to:
Services Australia
Complex Drugs Programs
Reply Paid 9826
HOBART TAS 7001