

medicare



Fistulising Crohn's disease – initial or recommencement (treatment break greater than 5 years) authority application

Online PBS Authorities



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** or **recommencing** PBS-subsidised biological medicines for patients with complex refractory fistulising Crohn's disease who are either:

- initiating PBS-subsidised treatment
- recommencing PBS-subsidised treatment after a treatment break > 5 years.

Important information

Authority applications to start or recommence 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.

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 complex refractory fistulising Crohn's disease **initial** or **recommencement** authority applications.

Where the term 'biological medicine' appears, it refers to adalimumab, infliximab or ustekinumab.

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 or **recommencing** treatment after a treatment break greater than 5 years.

After an authority application for the **first continuing** treatment has been approved, **subsequent continuing** treatments with PBS-subsidised biosimilar brands of biological medicines are **Authority Required (STREAMLINED)** and do not require authority approval from Services Australia for the listed quantity and repeats.

Continuing treatment with PBS-subsidised **infliximab s.c.** is **Authority Required (STREAMLINED)** and does not require authority approval from Services Australia for the listed quantity and repeats.

Section 100 arrangements for infliximab i.v. and ustekinumab i.v.

These items are 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.

These items are 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

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 PBS-subsidised biological medicine.

For more information

Go to servicesaustralia.gov.au/healthprofessionals

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0	nline PBS Authorities
	You do not need to complete this form if you use the Online PBS Authorities system. Go to servicesaustralia.gov.au/hppbsauthorities
Pa	tient's details
1	Medicare card number Ref no. Department of Veterans' Affairs card number
2	Family name
	First given name
3	Date of birth (DD MM YYYY)
4	Patient's weight kg
Pr	escriber's details
5	Prescriber number
6	Family name
	First given name
7	Business phone number (including area code) Alternative phone number (including area code)

HO	spital details
8	Hospital name
	This hospital is a:
	public hospital
_	private hospital
9	Hospital provider number
Co	nditions and criteria
	qualify for PBS authority approval, the following conditions ust be met.
10	The patient is being treated by a:
	gastroenterologist
	consultant physician specialising in gastroenterology (either internal or general medicine)
11	This application is for:
	adalimumab
	ustekinumab s.c with i.v. loading
	Go to 14
	infliximab i.v.
	(at a dose of 5 mg/kg) • Go to 13
	infliximab s.c. with i.v. loading (at a dose of 5 mg/kg) Go to 12
12	Does the patient have a concurrent PBS authority application for the IV form of infliximab that is approved or to be approved? Yes Go to 14 No
13	Will the treatment exceed a total of 3 doses to be administered at weeks 0, 2 and 6? Yes No



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14	The patient has:
	confirmed Crohn disease, defined by standard clinical, endoscopic and/or imaging features, including histologica evidence, with the diagnosis confirmed by a gastroenterologist or a consultant physician
	andan externally draining enterocutaneous or rectovaginal fistula.
15	Provide details of current Fistula assessment
	Date of assessment (no more than 1 month old) (DD MM YYYY) Number of externally draining complex fistulae

Fistulae symptom grading table

Note: Each parameter in this table must be assigned a value

Symptom	Descriptions	Score	Subtotal
Discharge	no discharge	0	
	minimal mucous discharge	1	
	moderate mucous or purulent discharge	2	
	substantial discharge	3	
	gross faecal soiling	4	
Pain	no pain	0	
	mild discomfort	1	
	moderate discomfort	2	
	marked discomfort	3	
	severe pain	4	
Degree of	no induration	0	
induration	minimal induration	1	
	moderate induration	2	
	substantial induration	3	
	gross fluctuance/abscess	4	

Checklist

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The relevant attachments need to be provided with this form.

Fistulae symptom grading total score

Details of the proposed prescription(s).

Privacy notice

17 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 **servicesaustralia.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

18 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.

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$\bullet $ 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)		
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