

medicare



Fistulising Crohn's disease – continuing 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 **continuing** PBS-subsidised biological medicines for patients with complex refractory fistulising Crohn's disease.

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 complex refractory fistulising Crohn's disease **continuing** 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 continuing treatment.

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.

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 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 particular PBS-subsidised biological medicine.

For more information

Go to servicesaustralia.gov.au/healthprofessionals

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Online PBS Authorities			Hospital details		
	You do not need to complete this form if you use the Online PBS Authorities system.	8	Hospital name		
	Go to servicesaustralia.gov.au/hppbsauthorities				
			This hospital is a:		
o a	tient's details		public hospital		
	_		private hospital		
1	Medicare card number	9	Hospital provider number		
	Ref no.				
	Or				
	Department of Veterans' Affairs card number	Co	nditions and criteria		
		To	qualify for PBS authority approval, the following conditions		
2	Family name		ust be met.		
		10	The patient is being treated by a:		
	First given name	10	gastroenterologist		
	The grown hame		consultant physician specialising in gastroenterology (either		
			internal or general medicine)		
3	Date of birth (DD MM YYYY)	11	This application is for the:		
			first continuing treatment with:		
4	Patient's weight		adalimumab		
	kg		infliximab i.v. (at a dose of 5mg/kg)		
			ustekinumab s.c.		
Pr	escriber's details		or		
-			subsequent continuing treatment with:		
5	Prescriber number		adalimumab		
			infliximab i.v. (at a dose of 5mg/kg)		
3	Family name		ustekinumab s.c.		
		12			
		12	Has the patient previously received PBS-subsidised treatment with this drug (regardless of formulation) for this condition?		
	First given name		Yes		
			No .		
7	Business phone number (including area code)	13	Has the assessment of response been conducted within the		
		.0	time frame specified in the PBS restriction?		
	Alternative phone number (including area code)		Yes		
			No 🗌		



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14	The patient has demonstrated an adequate response to treatment with this drug for this condition evidenced by:						
		ase in the number of open drainir % from baseline	ng fistula	e of at			
	and/or						
		ed reduction in drainage of all fist e, together with less pain and indo atient.	. ,				
Response assessment must be conducted following a minimum of or up to 12 weeks of treatment, or betwee and 16 weeks of therapy depending on the biological medicine requested. This assessment must be submitted later than 4 weeks from the cessation of treatment.							
Provide details of current Fistula assessment Date of assessment (no more than 1 month o				/IM YYYY)			
	tulae						
	-	nptom grading table parameter in this table must be as	ssigned a	a value			
	Symptom	Descriptions	Score	Subtotal			
	Discharge	no discharge	0				
		minimal mucous discharge	1				
		moderate mucous or purulent	2				

Symptom	otom Descriptions		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 induration	no induration	0	
	minimal induration	1	
	moderate induration	2	
	substantial induration	3	
	gross fluctuance/abscess	4	

Fistulae symptom grading total score

Checklist

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The relevant attachments need to be provided with this form.

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.

Lunderstand that

i understand that:		
$\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