

medicare



Severe chronic plaque psoriasis – initial authority application

When to use this form	Use this form to apply for initial PBS-subsidised biological medicines for patients 18 years or over with severe chronic plaque psoriasis.
Important information	Initial applications to start PBS-subsidised treatment must be 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 severe chronic plaque psoriasis initial authority applications.
	Where the term 'biological medicine' appears, it refers to adalimumab, bimekizumab, etanercept, guselkumab, infliximab, ixekizumab, risankizumab, secukinumab, tildrakizumab and ustekinumab. Patients are eligible for PBS-subsidised treatment with only 1 biological medicine at any time.
	A copy of the PASI calculation sheets is provided for your convenience.
	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 a written 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.
Section 100 arrangements for infliximab i.v. only	 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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Severe chronic plaque psoriasis – initial authority application

Patient's details Hospital details for infliximab i.v. only 1 Medicare card number 8 Hospital name Ref no. This hospital is a: or Department of Veterans' Affairs card number public hospital private hospital Т 9 Hospital provider number 2 Dr Mr Mrs Miss Ms Other Family name **Conditions and criteria** First given name To qualify for PBS authority approval, the following conditions must be met. 3 Date of birth (DD MM YYYY) **10** Is the patient, 18 years or over, being treated by a dermatologist? No 4 Patient's weight Yes kg **11** This application is for: adalimumab **Prescriber's details** bimekizumab 5 Prescriber number etanercept guselkumab infliximab i.v. 6 Dr Mr Mrs Miss Ms Other infliximab s.c. with i.v. loading Family name (and an authority prescription for at least 2 i.v. doses at weeks 0 and 2 is attached) ixekizumab First given name risankizumab secukinumab 7 Business phone number (including area code) tildrakizumab ustekinumab Alternative phone number (including area code) 1 1 1



12	The patient:			and/or					
			ie psoriasis (whole body) where	acitretin at a dose of at least 0.4 mg/k	g/day				
		ave been prese itial diagnosis	nt for at least 6 months from the	Dose	mg				
Г	or			From (DD MM YYYY)					
	hand or s	sole of a foot v	le psoriasis of the face, palm of /here the plaque or plaques have						
	been pres diagnosis		6 months from the time of initia	PASI score, if applicable					
	and			Date of assessment (DD MM YYYY)	Date of assessment (DD MM YYYY)				
			bsidised treatment with a						
	-	medicine for the	his condition	and/or					
	and	vo trootmont …	th this highering as	apremilast at a dose of 30 mg twice a	day				
			th this biological medicine as other than methotrexate).	Dose	mg				
	The patient has failed to achieve an adequate response, as indicated by a Psoriasis Area and Severity Index (PASI)			From (DD MM YYYY)					
	assessment, f	•	mum of 6 weeks treatment to	To (DD MM YYYY)					
		•	JVA) – a minimum of 3 treatme	PASI score, if applicable					
	per week		,	Date of assessment (DD MM YYYY)	Date of assessment (DD MM YYYY)				
	From (DD	MM YYYY)							
	To (DD MI	M YYYY)		and/or and/cr deucravacitinib at a dose of 6 mg once daily					
	PASI scor	e, if applicable		Dose	mg				
		ssessment (DD	MM YYYY)						
				From (DD MM YYYY)					
	and/or								
	methotre	exate at a dose	of at least 10 mg weekly	PASI score, if applicable					
	Dose		mg	Date of assessment (DD MM YYYY)	Date of assessment (DD MM YYYY)				
	From (DD	MM YYYY)							
	To (DD MI	M YYYY)		A PASI assessment must be completed for each prior treatment course preferably whilst still on treatment, but					
	PASI scor	e, if applicable		no later than 4 weeks following cessation	of treatment.				
		ssessment (DD	MM YYYY)	14 Provide details of contraindications or intole	rances to any of th				
				prior therapies including the degree of toxic	prior therapies including the degree of toxicity.				
	and/or			For details of the toxicity criteria, go to					
	ciclosporin at a dose of at least 2 mg/kg/day			servicesaustralia.gov.au/healthprofessio Intolerance must be of a severity to necessi					
	Dose		mg	treatment withdrawal.					
	Erom (DD			Prior therapy contraindication or toxicity and	l grade.				
	From (DD To (DD MI	MM YYYY) M YYYY)		Phototherapy					
		e, if applicable							
	Date of as	ssessment (DD		Methotrexate					
				Methotrexate					

 \square

Ciclosporin

Aci	tretin
Apr	emilast
Deı	icravacitinib
	patient has failed to achieve an adequate response to pr
trea	tment as demonstrated by:
	a whole body PASI score > 15
	PASI score
	Date of assessment (DD MM YYYY)
or	
	chronic plaque psoriasis classified as severe due to a
	plaque or plaques on the face, palm of a hand or sole of

foot where: **at least 2** of the **3** PASI symptom subscores for erythema, thickness and scaling are rated as severe or very severe

the skin affected $is \ge 30\%$ of the face, palm of a hand or sole of a foot.

Date of assessment (DD MM YYYY)

The PASI assessment must not be **older than 4 weeks** at the time of application.

Checklist

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The relevant attachments need to be provided with this form.

Details of the proposed prescription(s).

The completed PASI calculation sheet(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.

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)

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

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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



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PASI calculation and body diagram – face, hand and foot

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		Body regio	n				
Indicate the degree of	FACE	RIGHT PALM	LEFT PALM	RIGHT SOLE	LEFT SOLE		
involvement of the body region surface as a percentage	%	%	%	%	%		
OR							
Clearly indicate the plaque characteristics for each body region by circling the number which best corresponds to the patient's skin condition (circle one number in each box)							
	0 = none						
	1 = slight						
Erythema	2 = moderate						
	3 = severe						
	4 = very severe						
	0 = none						
	1 = slight						
Thickness	2 = moderate						
	3 = severe						
	4 = very severe						
	0 = none						
	1 = slight						
Scaling	2 = moderate						
	3 = severe						
	4 = very severe						
Mark clearly on the diagrams the extent of the affected area(s)		R	L				

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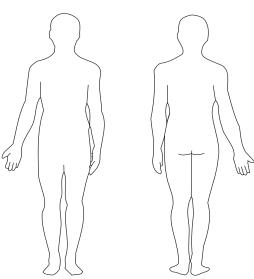
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PASI calculation and body diagram – whole body

	Deting coord	Body region (and weighting factor)				
Plaque characteristic	Rating score	Head	Upper Limbs	Trunk	Lower Limbs	
Erythema	0 = None 1 = Slight					
Thickness	2 = Moderate					
Scaling	3 = Severe 4 = Very severe				-	
Add together each of the 3 scores for each of the body regions to give 4 separate sub totals.						
	Sub Totals	A1=	A2=	A3=	A4=	
Multiply each sub total by the amount of body surface area represented by that region i.e. A1 x 0.1 for head, A2 x 0.2 for upper limbs, A3 x 0.3 for trunk, A4 x 0.4 for lower limbs to give a value B1, B2, B3 and B4 for each body region respectively						
		A1 x 0.1 = B1	A2 x $0.2 = B2$	A3 x 0.3 = B3	A4 x $0.4 = B4$	
		B1=	B2=	B3=	B4=	
Degree of involvement as % for each body region affected (score each region with score between 0–6)	0 = None 1 = 1-9% 2 = 10-29% 3 = 30-49% 4 = 50-69% 5 = 70-89% 6 = 90-100%					
For each body region multiply sub total B1, B2, B3 and B4 by the score (0–6) of the % of body region involved to give 4 subtotals C1, C2, C3 and C4						
		B1 x score = C1	B2 x score = C2	B3 x score = C3	B4 x score = C4	
		C1=	C2=	C3=	C4=	
The patient's PASI score is the su	m of C1+C2+C3+C4			PASI=		

Shade in the affected areas



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