

## australasian society of clinical immunology and allergy

23 January 2025

Chair, Pharmaceutical Benefits Advisory Committee (PBAC) Department of Health and Ageing GPO Box 9848 Canberra ACT 2601

Email: pbac@health.gov.au

Dear PBAC Chair,

Re: New PBS listing of lxifi® (infliximab)

On behalf of the Australasian Society of Clinical Immunology and Allergy (ASCIA) we are writing in support of the request by Pfizer Australia Pty Ltd for the Pharmaceutical Benefits Scheme (PBS) listing of Ixifi® (infliximab), as outlined in the table below, for review at the March 2025 PBAC meeting.

INFLIXIMAB		
Powder for I.V. infusion 100 mg	Severe active rheumatoid arthritis Ankylosing spondylitis Severe psoriatic arthritis	To request Section 100 (Highly Specialised Drugs Program)
lxifi®	Severe chronic plaque psoriasis Severe Crohn disease,	Authority Required listings of a new infliximab biosimilar under
PFIZER AUSTRALIA PTY LTD	Complex refractory fistulising Crohn Disease Moderate to severe ulcerative colitis	the same conditions as other biosimilar brands of infliximab.
(New PBS listing)		

The ASCIA membership includes clinical immunologists with expertise in managing rheumatoid arthritis and other autoimmune conditions.

ASCIA supports the ARA Rheumatoid Arthritis Clinical Care Standard which states that:

- Early diagnosis and treatment lead to better long-term clinical outcomes and a better quality of life for people with rheumatoid arthritis.
- Treatment with biological disease modifying antirheumatic drugs (biological DMARDs) such as infliximab should be started as soon as rheumatoid arthritis is diagnosed.

Furthermore, having multiple brands available of infliximab mitigates against the risk of supply issues.

Therefore, ASCIA supports the PBS listing of Ixifi® (infliximab) to facilitate early treatment of severe active rheumatoid arthritis and other relevant autoimmune conditions with biological DMARDs.

Yours sincerely.

Dr Michael O'Sullivan Jill Smith ASCIA President ASCIA CEO

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