
STARTING TREATMENT WITH ERELZI® (ETANERCEPT)¹

HELPING YOU ACHIEVE
YOUR TREATMENT GOALS

AXIAL SPONDYLOARTHRITIS



This medicine is subject to additional monitoring. This will allow quick identification of new safety information. You can help by reporting any side effects you may get at <https://pvi1j.solutions.iqvia.com>

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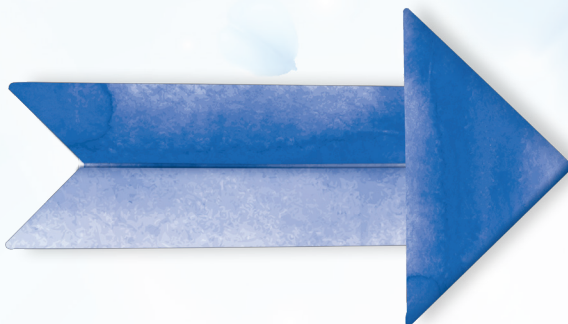
STARTING YOUR NEW TREATMENT



Your doctor has recommended that you start taking a biological medicine, known as a biologic.¹ This might be because your current medicine is not controlling your symptoms well enough or is causing side effects.¹

BIOLOGICS HAVE TRANSFORMED THE TREATMENT PARADIGM FOR AXIAL SPONDYLOARTHRITIS^{2,3}

Your doctor has recommended that you start treatment with Erelzi[®], which contains the active ingredient etanercept.^{1,4} You will start to use it as soon as you or your carer learn how to inject it.⁴



THIS GUIDE WILL HELP YOU UNDERSTAND
WHAT ERELZI[®] IS, WHY IT HAS BEEN PRESCRIBED
TO YOU, AND HOW IT CAN HELP YOU.

ABOUT BIOLOGIC MEDICINES



BIOLOGICS MAY BE DIFFERENT FROM ANY TREATMENT YOU HAVE TRIED BEFORE. THEY ARE THERAPIES THAT SPECIFICALLY TARGET INFLAMMATION.^{2,4}

Most drugs are made by combining chemical ingredients.⁶ Biologics are different because they contain proteins that are made by living cells.^{6,7} They are designed to act on a particular molecule, cell or other structure inside the body (a specific target).^{2,7}

Biologics are often administered by an injection through the skin, while some are given by intravenous infusion (a drip).⁷

BIOLOGICS HAVE MADE A BIG DIFFERENCE TO PEOPLE WITH AXIAL SPONDYLOARTHRITIS^{2,3}

Inflammation is a normal process and occurs when the body fights against bacteria and viruses.⁸ In autoinflammatory conditions such as rheumatoid arthritis,⁹ psoriatic arthritis,¹⁰ axial spondyloarthritis¹¹ and psoriasis,¹² the process is not controlled and the body targets its own tissue.⁸ A substance called tumour necrosis factor (TNF) is overproduced in patients with these conditions, and this can lead to inflammation and damage to joints, as well as to other organs and body systems.¹³⁻¹⁶

Several biologics have been developed to block the effects of TNF and control inflammation.¹⁷ As these biologics work against TNF, they are often referred to as 'anti-TNFs'.^{15,17} The medicine you have been prescribed is an anti-TNF.¹

TREATMENT WITH ERELZI®¹

HOW TO ADMINISTER ERELZI®



At the start of your treatment you will be trained on how to administer Erelzi®.⁴ Read all of the package leaflet information carefully before you start using Erelzi® because it contains important information for you. Erelzi® is injected under your skin (subcutaneous injection), using the device that was provided to you.⁴ Erelzi® is available either as a pre-filled syringe or SensoReady® pen.¹ Your doctor or nurse can help you decide which is best for you.

Rest assured that, throughout your entire treatment with Erelzi®, you will be followed up by your doctor on a regular basis.

HOW TO STORE ERELZI®

Erelzi® should be stored in the refrigerator (2–8°C) and must not be frozen.⁴ If you have a pack with several syringes or SensoReady® pens, please take one out of the pack for application and return the rest to the refrigerator straight away.⁴ If you are travelling, please make sure to include a cool pack with your medicine and return the medicine to a refrigerator as soon as possible.⁴

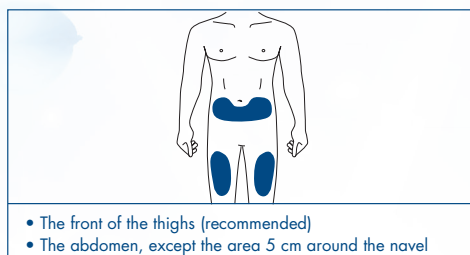
Erelzi® may be stored at temperatures up to a maximum of 25°C for a single period of up to four weeks, after which it should not be refrigerated again.⁴ Erelzi® should be discarded if not used within four weeks of removal from refrigeration.⁴

UNDERSTANDING YOUR DOSE

To treat axial spondyloarthritis, the recommended dose of Erelzi® is 25 mg twice a week or 50 mg once a week as an injection.¹ Your doctor will confirm the dose before you begin treatment.⁴ They will also decide how long you should take Erelzi® and whether retreatment is needed.⁴

WHERE TO INJECT

Before injection, let Erelzi® reach room temperature (take the blister pack containing pre-filled syringe or pen out of the refrigerator and leave unopened for 15 to 30 minutes).⁴ Erelzi® should be injected in one of these two sites:⁴

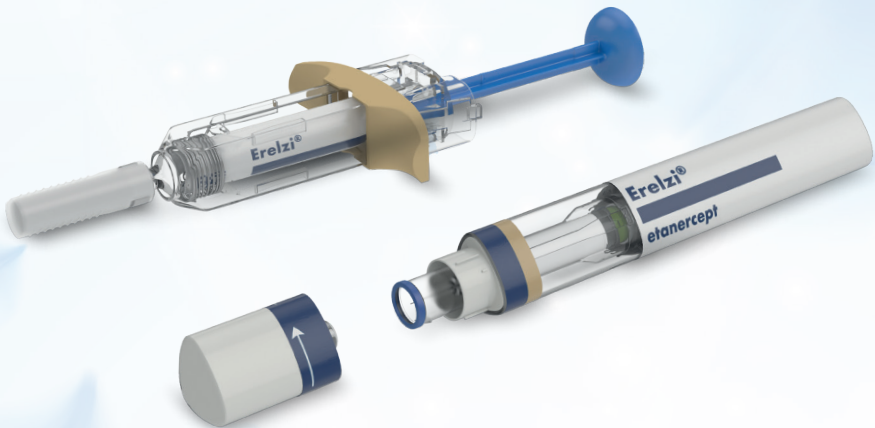


HOW TO DISPOSE

Dispose of the used syringe or pen in a sharps container (closable, puncture-resistant container).⁴ Do not throw away the used syringe with your household waste.⁴

GETTING READY TO START YOUR TREATMENT

You will be trained on how to administer your medicine, in order to get used to a new injection device.⁴



AFTER YOU OR YOUR CARER ARE COMFORTABLE
WITH ERELZI® ADMINISTRATION, YOU CAN
INJECT YOURSELF OR BE INJECTED AT HOME.⁴

This will give you the flexibility to inject when and where it suits your daily routine, and you may not have to go to the hospital or your doctor for the injection.

Your doctor will evaluate your past medical history, as well as any allergies or other conditions you may have and the medications you take, before you can start treatment.^{1,4}

You will also need to have some initial screening tests, which may include:

- Blood tests to evaluate if you have any ongoing infections¹
- Tests for tuberculosis, which may include a chest X-ray, as treatment with biologics may lead to reactivation of latent (inactive) tuberculosis, and must not be initiated in patients with active tuberculosis⁴

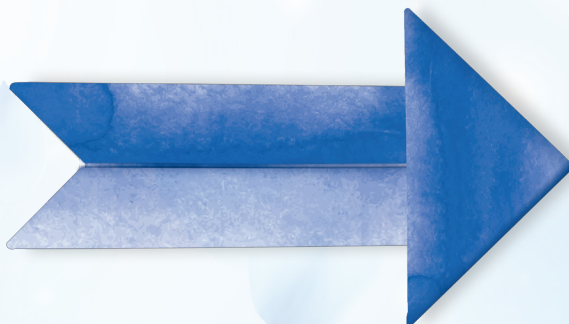


WHAT ARE THE NEXT STEPS?



You will receive training on how to use the injection device for Erelzi®.⁴ Ask your doctor if you have any questions about how to use it, as it is important you are comfortable with it as soon as possible.⁴ If you are unable to inject yourself, your family or carer may be trained to inject you with Erelzi®.⁴

If you feel unwell, contact your doctor immediately.⁴



YOU SHOULD ALWAYS USE ERELZI® EXACTLY
AS PRESCRIBED BY YOUR DOCTOR.⁴
IF YOU FEEL THAT THE TREATMENT
IS NOT RIGHT FOR YOU, TALK TO
YOUR DOCTOR TO DISCUSS POTENTIAL
OPTIONS REGARDING YOUR TREATMENT⁴

WHAT YOU CAN EXPECT FROM YOUR TREATMENT



If you've been given this leaflet, it is because your doctor thinks Erelzi® may help treat your condition.⁴ However, the benefits and side effects of treatment vary from person to person.⁴



TALK TO YOUR DOCTOR FOR MORE
INFORMATION ABOUT WHAT TO EXPECT
FROM YOUR TREATMENT WITH ERELZI®

POSSIBLE SIDE EFFECTS

Every medicine can have side effects, and over the course of your treatment, you may experience one or some of them, although not everybody gets them.⁴ It is not possible to predict which side effects you may have, and they may vary in severity and duration.^{1,4}

FREQUENT SIDE EFFECTS OF ERELZI® INCLUDE⁴

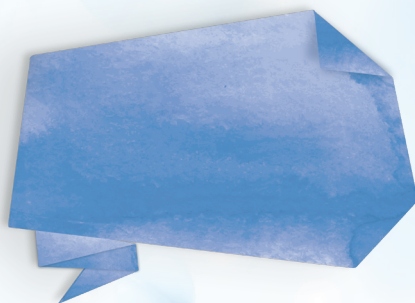
- Infections, including colds, sinusitis, bronchitis, urinary tract infections and skin infections
- Injection site reactions, including bleeding, bruising, redness, itching, pain, and swelling
- Headache
- Fever
- Itching, rash
- Allergic reactions
- Antibodies against normal tissue

Please see the Patient Information Leaflet for less frequent side effects which might occur when taking Erelzi®.⁴

No studies have been performed on effects on the ability to drive or use machines.⁴ You should not be treated with Erelzi® if you are pregnant or breastfeeding.⁴ Safety in pregnancy and lactation has not been established.⁴

YOU ARE NOT ALONE

For more information and support please contact your healthcare provider and/or patient support nurse. You can also email Sandoz at:



medical.za@sandoz.com

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References: 1. Sandoz SA (Pty) Ltd. ERELZI® Professional information. V3.0 (20/10/2022), approved 13 September 2022. 2. Biologic therapy. National Axial Spondyloarthritis Society. Accessed June 9, 2022. <https://nass.co.uk/managing-my-as/medication/biologic-therapy/>. 3. Agrawal P, Machado PM. Recent advances in managing axial spondyloarthritis. F1000Research. Accessed June 9, 2022. <https://f1000research.com/articles/9-697>. 4. Sandoz SA (Pty) Ltd. ERELZI® Patient information leaflet. V2.0 (15/11/2021), approved 26 January 2022. 5. Malaviya AP, Östör AJK. Rheumatoid arthritis and the era of biologic therapy. Inflammopharmacol. 2012;20(2):59-69. doi:10.1007/s10787-012-0123-y. 6. A patient's guide to understanding biosimilars. Global Healthy Living Foundation. 2018. Accessed June 3, 2022. 7. Morrow T. Defining the difference: what makes biologics unique? Biotechnology Healthcare. Accessed June 2, 2022. 8. Inflammation: what is it, causes, symptoms & treatment. Cleveland Clinic. Accessed June 6, 2022. <https://my.clevelandclinic.org/health/symptoms/21660-inflammation>. 9. Rheumatoid arthritis (RA). CDC. Accessed June 6, 2022. 10. Psoriatic arthritis - symptoms and causes. Mayo Clinic. Accessed June 6, 2022. <https://www.mayoclinic.org/diseases-conditions/psoriatic-arthritis/symptoms-causes/syc-20354076>. 11. Ankylosing spondylitis - symptoms and causes. Mayo Clinic. Accessed June 6, 2022. <https://www.mayoclinic.org/diseases-conditions/ankylosing-spondylitis/symptoms-causes/syc-20354808>. 12. Psoriasis - symptoms and causes. Mayo Clinic. Accessed June 6, 2022. <https://www.mayoclinic.org/diseases-conditions/psoriasis/symptoms-causes/syc-20355840>. 13. Yost J, Gudjonsson JE. The role of TNF inhibitors in psoriasis therapy; new implications for associated comorbidities. F1000 Med Rep. 2009;1. doi:10.3410/M1-30. 14. Manara M, Sinigaglia L. Bone and TNF in rheumatoid arthritis: clinical implications. RMD Open. 2015;1(Suppl 1):e000065. doi:10.1136/rmdopen-2015-000065. 15. Braun J, Sieper J, Breban M, et al. Anti-tumour necrosis factor therapy for ankylosing spondylitis: international experience. Annals of the Rheumatic Diseases. 2002;61(Supplement 3):51iii-60. doi:10.1136/ard.61.suppl_3.iii51. 16. Mease PJ. Tumour necrosis factor (TNF) in psoriatic arthritis: pathophysiology and treatment with TNF inhibitors. Annals of the Rheumatic Diseases. 2002;61(4):298-304. doi:10.1136/ard.61.4.298. 17. TNF inhibitors. American College of Rheumatology. Accessed June 6, 2022.

ERELZI® 25 mg (solution for injection). Reg. No.: 55/3.1/0392. Composition: Each pre-filled syringe contains 25 mg etanercept. ERELZI® 50 mg (solution for injection). Reg. No.: 55/3.1/0393. Composition: Each pre-filled syringe/pen contains 50 mg etanercept. Pharmacological Classification: A3.1 Anti-rheumatics (anti-inflammatory agents). For full prescribing information refer to the Sandoz Professional Information approved by the South African Health Products Regulatory Authority (SAHPRA).

Sandoz SA (Pty) Ltd, Reg. No. 1990/001979/07, Magwa Crescent West, Waterfall City, Jukskei View, Midrand, 2090. Tel: +27 (11) 347 6600. SANCAL Customer Call Centre: 0861 726 225. Reporting of AEs via the link: <https://pvi1j.solutions.iqvia.com> or the e-mail address, adverse.event.sac@sandoz.com ZA2407310741