# ADALIMUMAB

[Add-a-LIM-you-mab]

Examples of brand names: Abrilada, Amgevita, Hadlima, Humira, Hyrimoz, Yuflyma

This information sheet is from the Australian Rheumatology Association to help you understand the medicine prescribed for you. It includes important information about:

- how to take your medicine
- possible side effects
- tests you <u>will</u> have to monitor your condition
- other precautions to take while you are taking adalimumab.

Please read it carefully and talk to your doctor.

### IMPORTANT THINGS TO REMEMBER

- You must see your rheumatologist regularly to make sure the treatment is working and check for possible side effects.
- You should have regular blood tests as suggested by your rheumatologist.
- It is important to tell your rheumatologist if you have a new serious illness such as a serious infection, cancer or heart failure.
- If you are worried about any side effects, you should contact your rheumatologist as soon as possible.
- If you stop adalimumab for any reason, you must contact your rheumatologist. Failure to do so may mean that your treatment may no longer be funded.
- If you plan to become pregnant, you must discuss the timing with your rheumatologist

For more information about inflammatory conditions associated with arthritis, see Arthritis Australia's website:

www.arthritisaustralia.com.au

## What is adalimumab?

Adalimumab belongs to a class of medicines called biological disease modifying antirheumatic drugs (biological DMARDs or bDMARDs). Specifically, it is a Tumour Necrosis Factor (TNF) inhibitor.

bDMARDs have now been given to over a million people worldwide since their first use in the late 1990s.

These medicines block substances called cytokines. These cytokines are found in excessive amounts in the blood and joints of people with conditions such as rheumatoid arthritis, psoriatic arthritis, juvenile arthritis and ankylosing spondylitis.

They cause inflammation, which results in symptoms of pain, joint swelling and stiffness, and can lead to joint damage. They also cause eye inflammation in people who have uveitis.

By blocking the cytokine called TNF, adalimumab lessens inflammation, pain symptoms and helps stop further joint damage.

# What benefit can you expect from your treatment?

Adalimumab can help people with inflammatory arthritis live longer with a better quality of life. Studies show that patients who take adalimumab have a lower risk of death and fewer flares.

Many patients with inflammatory arthritis experience reduced pain, swelling, and stiffness from taking adalimumab. It helps to slow or stop disease progression and prevent joint or organ damage. This can enhance mobility and reduce disability. Adalimumab usually takes several weeks to take effect. It is important to continue taking the medicine as prescribed to see full benefits.

### Stopping adalimumab

If you stop or delay your adalimumab treatment, you may worsen again. Continue your treatment, unless told by your rheumatologist to stop or unless side effects occur (see *Side effects*). If you stop adalimumab for any reason, you <u>must</u> contact your rheumatologist.

Failure to do so may mean that your treatment may no longer be funded.





#### **Brands of adalimumab**

There are several different brands or "biosimilar" adalimumab medicines. A biosimilar is a version of adalimumab that has been shown to have the same benefits and safety as the original product. It is preferable that switching between different adalimumab brands is minimised.

# How will you be checked while on adalimumab?

- Medicines like adalimumab are very expensive and highly subsidised by Medicare. Certain conditions must be met to receive it.
- Adalimumab will only be given if your disease is active and if standard treatments have not worked.
- It will only be kept going if it helps your condition. This must be checked between 12 and 16 weeks after the start of treatment.
- Blood tests are needed during your treatment to watch for side effects and decide if the treatment is working.
- How often you have blood tests will depend on what other medicines you are taking and what other illnesses you might have. Your rheumatologist will advise on this.

### How is adalimumab taken?

Adalimumab is injected under the skin of the abdomen or thigh. It comes in a pen or a syringe injection.

It can be injected by your doctor, nurse, carer or by you. If injecting yourself, be sure to follow the detailed instructions carefully to ensure the best response. It is important to change where you inject each time.

# What is the dosage?

The usual dose for adults with arthritis is 40mg once every two weeks.

# Can other medicines be taken with adalimumab?

Adalimumab may be used with other arthritis medicines including:

- other DMARDs such as methotrexate
- steroid medicines such as prednisolone or cortisone injections into the joint
- anti-inflammatory medicines (NSAIDs) such as naproxen (Naprosyn) or ibuprofen (Brufen, Nurofen)
- simple pain medicines such as paracetamol.

Adalimumab cannot be used with other bDMARDs.

There are separate information sheets for the medicines mentioned above.

## Are there any side effects?

You might experience side effects with your treatment. Contact your rheumatologist if you have any concerns about possible side effects. Many side effects go away when you stop taking adalimumab.

### Most common possible side effects

- Mild pain, swelling or itching at the site of the injection are very common (up to 20% of patients) but can be reduced by applying ice and antihistamine/steroid creams to the injection site and/or leaving the medicine out of the fridge for 30 minutes before injecting.
- Headaches, cough, stomach and bowel discomfort may also occur.
- As adalimumab affects the immune system, mild infections, mainly the upper respiratory tract (e.g. colds, sinusitis) may occur more often. Treatment with adalimumab may need to be briefly stopped for a serious infection so contact your rheumatologist for advice.

## Less common or rare possible side effects

- Serious infections such as tuberculosis (TB) are seen rarely, and screening for TB is needed before treatment begins.
- Rarely, adalimumab may cause an allergic reaction with itchy, red skin, a rash or a feeling of tightness in the chest and trouble breathing.
- Side effects involving the nerves, such as inflammation of the nerve to the eye, may also occur rarely, causing changes in vision or sensation.
- Very rarely 'drug-induced lupus' has occurred with symptoms of rash, fever and increased joint pain.
- Annual skin checks are suggested with any medicines that can suppress the immune system as there is a slight increase in risk in skin cancers. To date research and use over 20 years, have not shown an increase in risk of other cancers.

# What precautions are necessary? *Infections*

 If you have a current infection of any kind treatment with adalimumab should not be given until the infection is treated.

You will need some blood tests and a chest X-ray to exclude some chronic infections before your first bDMARD.

## Use with other medical conditions

Worsening may occur of the following conditions:

multiple sclerosis





- moderate to severe heart failure
- systemic lupus erythematosus (lupus/SLE).
   People with SLE are not often given adalimumab but each case will be assessed whether safe by your rheumatologist.

#### Use with other medicines

Adalimumab can interact with other medicines. You should tell all your doctors about all medicines you are taking or plan to take. This includes over the counter or herbal/naturopathic medicines.

#### **Vaccines**

- If you are taking adalimumab, you should not be immunised with 'live' vaccines such as: MMR (measles, mumps and rubella), Varicella vaccines (Chicken pox), OPV (oral polio virus), BCG (Bacillus Calmette Guerin), Japanese Encephalitis or Yellow Fever. Talk with your rheumatologist before receiving any vaccines.
- Pneumococcal vaccines and the yearly seasonal flu vaccinations are encouraged.

For more information on vaccination including the COVID-19 vaccination go to the ARA website; <a href="https://rheumatology.org.au/">https://rheumatology.org.au/</a> patients, medication information, vaccinations.

#### Surgery

 If you need surgery for any reason, adalimumab should be stopped before surgery. It can be started again after the operation at a time decided by your surgeon and rheumatologist (often once the wound has healed there is no infection).

#### Use with alcohol

 You may drink alcohol while taking adalimumab. If you are also taking methotrexate, you should be cautious about how much alcohol you drink.

#### Use in pregnancy and when breastfeeding

- It is important to discuss with you doctor if you are planning a pregnancy while on adalimumab.
- It may be used in pregnancy and in men trying to father a child.
- If adalimumab is kept going beyond 4
  months of pregnancy it may increase the risk
  of infection in the newborn when live
  vaccines may be due.
- The rotavirus vaccine should be given within the first six months of life. MMR may be given at 6 months.

 There is only limited information regarding adalimumab in breast milk and while small amounts may occur, it does not seem to be harmful.

More detailed information is available at: <a href="https://rheumatology.org.au/For-Patients/Pregnancy-Information">https://rheumatology.org.au/For-Patients/Pregnancy-Information</a>

#### How to store adalimumab

- Adalimumab should normally be kept refrigerated. If needed, for example when travelling, it may be stored below 25°C for up to 2 weeks and then used or discarded.
- Keep all medicines out of reach of children.

#### Questions?

If you have any questions or concerns write them down and discuss them with your doctor.

Your doctor's contact details

You should see your rheumatologist regularly to make sure the treatment is working and to minimise any possible side effects.

This Information Sheet has been prepared using materials obtained from various sources which have been reviewed by the Australian Rheumatology Association (ARA). It contains general information only and does not contain a complete or definitive statement of all possible uses, actions, precautions, side effects or interactions of the medicines referenced. This information is not intended as medical advice for individual conditions nor for making an individual assessment of the risks and benefits of taking a particular medicine. Decisions regarding the assessment and treatment of patients are the sole responsibility of the treating medical professional, exercising their own clinical judgment and taking into account all of the circumstances and the medical history of the individual patient.

ARA has used all reasonable endeavours to ensure the information on which this Information Sheet is based is accurate and up to date. However, the ARA accepts no responsibility or liability for the accuracy, currency, reliability and/or completeness of the information contained in this Information Sheet. To the maximum extent permitted by law, the ARA expressly disclaims any liability for any injury, loss, harm or damage arising from or in connection with use of and reliance on the information contained in this Information Sheet.

This information sheet is copyright and may be reproduced in its entirety but may not be altered without prior written permission from the ARA.

Please consider the environment before printing this resource.



