

# PATIENT INFORMATION ON MILNACIPRAN

[Mil-na-si-prin]

Brand name: Joncia

This information sheet has been produced by the Australian Rheumatology Association to help you understand milnacipran. It includes important information about:

- **how you should take your medication**
- **possible side (harmful) effects**
- **what tests you will have to monitor your condition and detect unwanted effects**
- **other precautions you should take while you are taking milnacipran.**

Please read it carefully and discuss it with your doctor.

## IMPORTANT THINGS TO REMEMBER

- While taking milnacipran you must see your rheumatologist regularly to ensure the treatment is working and minimise any possible side effects.
- You may need regular blood tests as suggested by your rheumatologist
- If you are worried about any side effects, you should contact your rheumatologist as soon as possible.

**For more information about fibromyalgia and other inflammatory conditions, see Arthritis Australia's website:**  
[www.arthritisaustralia.com.au](http://www.arthritisaustralia.com.au)

## What is milnacipran?

Milnacipran (brand name: Joncia®) belongs to a class of medicines called serotonin and noradrenaline reuptake inhibitors (SNRI). Other similar drugs from this group are used to treat depression and anxiety, but in Australia milnacipran is approved specifically to treat fibromyalgia. Although milnacipran is not listed on the Pharmaceutical Benefits Scheme (PBS), your rheumatologist may be able to obtain approval through the Special Access Scheme and prescribe it.

## What benefit can you expect from your treatment?

The purpose of taking medications in fibromyalgia is to control symptoms and improve quality of life. Medications in fibromyalgia should be used as part of a treatment plan together with non-medication approaches, such as physical activity.

Milnacipran has been studied in over 3000 fibromyalgia patients worldwide. In clinical studies that compared milnacipran to placebo, milnacipran improved the symptoms of pain, fatigue, sleep quality and quality of life.

Milnacipran can additionally have mild positive effects on disability and cognition (clarity of thinking).

## How is milnacipran taken?

Milnacipran is taken orally twice daily. It currently comes in capsules of two strengths: 25mg and 50mg.

To minimise side effects, it is recommended to commence with a single daily dose of 25mg for two days, then increase to 25mg twice daily for six days before increasing to the full dose of 50mg twice daily.

If after six days on 25mg twice daily side effects are occurring, it is recommended that the 25mg twice daily dosage be continued for another 7 days as there is a reasonable chance side effects will settle during this extra time on the lower dose.

In general, if no useful response occurs within 4 weeks of taking the full dose of milnacipran it should be ceased. Drug cessation should occur by weaning over at least 2 weeks. Do not stop milnacipran suddenly.

## Can other medications be taken with milnacipran?

As milnacipran increases serotonin levels within the brain, if it is taken together with other medications which also increase serotonin it may cause serotonin syndrome, a potentially serious medical condition. These medications include tramadol, triptan medications for migraine and most antidepressants, such as amitriptyline, escitalopram and St John's wort. In general,

milnacipran should not be taken concurrently with any of these.

Milnacipran works by an independent mechanism in the brain and so does not interact with pregabalin, allowing it to be safely taken at the same time. Similarly, pain relievers such as paracetamol and non-steroidal anti-inflammatories (such as Nurofen™) do not interact with milnacipran.

### How long is the treatment continued?

Treatment with milnacipran is continued indefinitely as long as it is effective and as long as no serious side effects occur. The beneficial effect does not tend to wear off.

If you stop milnacipran treatment for more than a few weeks there is a risk that your condition may worsen. Continue with your treatment unless advised by your doctor or unless side effects develop.

### Are there any side effects?

You might experience side effects with your treatment. Tell your doctor if you are concerned about possible side effects.

Most people with fibromyalgia experience at least one side effect from milnacipran, but only a minority of these people will stop the drug because of persistent side effects.

Although the effectiveness of milnacipran may be mildly improved by increasing the dose above 50mg twice daily to a maximum of 100mg twice daily, significant side effects occur more frequently.

A reduction in dose may minimise side effects so that you can continue to take the treatment. Your doctor will advise on any dose changes that are necessary.

### Most common possible side effects

Nausea, constipation, hot flushes, headaches and excessive sweating are the most common side effects of milnacipran.

### Less common or rare possible side effects

Less common side effects include dizziness, palpitations, high blood pressure and increased heart rate.

### More information about possible side effects

Information that comes with your milnacipran medicine describes in detail potential side effects that may occur with milnacipran.

### What precautions are necessary?

In general, milnacipran should not be used in people who have:

- Uncontrolled high blood pressure, cardiac rhythm irregularities, severe or unstable heart (coronary) disease or severe heart failure
- Closed-angle glaucoma
- Have an allergy to one of the ingredients in milnacipran (see medication information leaflet for list of ingredients).

In addition, before starting milnacipran, you should tell your doctor if you have any of the following:

- Kidney disease
- Epilepsy or seizure disorder
- Bipolar disorder or depression
- Enlargement of prostate or difficulty urination
- Bleeding problems.

If these conditions are well controlled, it may still be possible to cautiously take milnacipran.

Both before and during treatment with milnacipran, blood pressure and heart rate should be regularly monitored.

### Use with alcohol

Although there is no evidence of interaction with alcohol, as milnacipran has an effect on the brain, alcohol should only be consumed in moderation while taking it.

### Use in pregnancy and when breastfeeding

Milnacipran should not be used during pregnancy or breastfeeding.

### How to store milnacipran?

Store milnacipran in a cool, dry place, away from direct heat and light (e.g. not in the bathroom).

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.

