

Consumer Medicine Information

What is in this leaflet

This leaflet answers some common questions about MYOVIEV. It does not contain all the available information, nor does it take the place of talking to your doctor or pharmacist.

All medicines and diagnostic preparations have risks and benefits. Your doctor has weighed the risks of you being treated with MYOVIEV against the expected benefits.

If you have any concerns about being given this medicine, ask your doctor or pharmacist or treatment provider.

Keep this leaflet.

You may need to read it again.

What MYOVIEV is used for

MYOVIEV is used in the preparation of Technetium [^{99m}Tc] Tetrofosmin Injection, a radioactive medicinal product. Such substances are also known as radiopharmaceuticals.

Such radiopharmaceuticals are given in small amounts to find or rule out a disease. When used in this way the radiation your body receives is very low and is considered safe.

After the radioactive liquid is given to you, it is taken up by the organs of interest, especially the heart muscle or just passes through your body. The radiation is taken up by a special camera and pictures are prepared. These pictures allow the nuclear medicine doctor to detect any problems. MYOVIEV is used to help detect areas in the heart that may not be getting enough blood to work properly.

MYOVIEV is used as an aid in diagnosis only. It is not used to treat or cure the condition.

Before you are given MYOVIEV

When you must not be given it

MYOVIEV is not recommended for adolescents or children under 12 years of age.

You must not be given MYOVIEV if you are allergic to it or any of the ingredients listed at the end of this leaflet or have had an allergic reaction in a diagnostic procedure with the same or a similar preparation.

You must not be given MYOVIEV if you are pregnant.

Before you are given it

Your doctor must know about all of the following before you are given MYOVIEV. Tell your doctor if you:

1. are breastfeeding

MYOVIEV may be excreted in the breast milk. Therefore, you should stop breastfeeding and substitute formula feeds for at least 24 hours after you are given MYOVIEV.

- 2. are taking any medicines that affect your heart or blood pressure, such as**
- beta blockers**
- calcium antagonists**
- vasodilators like tablets or spray containing nitrates**

These medicines can affect the results of the diagnostic tests, so you should mention these to your doctor.

3. if you are taking any other medicines, including any that you have bought from your pharmacy, supermarket or health food shop.

Some medicines may affect the way other medicines work. Some can have an unfavourable influence on the diagnostic tests.

4. If you have allergies to:

- any other medicines
 - any other substances such as foods, substances to maintain food quality or freshness (preservatives) or colourants (dyes)
 - any of the ingredients of MYOVIEV listed at the end of this .
- 5. If you have kidney or liver problems**

If you have not told your doctor about any of the above, tell them before you are given any MYOVIEV.

How MYOVIEV is given

Your doctor may have special instructions for you to follow to get ready for your procedure. You should not eat the night before or have only a light breakfast on the morning of your test.

MYOVIEV is given as an injection into a vein. MYOVIEV must only be given by a doctor or nurse.

Your doctor will decide what dose of MYOVIEV you will receive.

MYOVIEV is usually given in two doses, one during exercise (or after you have got medicine to increase your heart rate) and one at rest about 4 hours after the first dose. Soon after it is given diagnostic images will be taken.

If you have any questions about taking MYOVIEV or about the diagnostic procedure, ask your doctor.

After being given MYOVIEV

MYOVIEV breaks down quickly after it is given and is largely removed from the body after about 48 hours. There are usually no special precautions to observe for radiopharmaceuticals like MYOVIEV because they are used in small amounts for diagnosis.

If you have any questions or concerns, be sure to discuss these with your doctor or pharmacist.

Side Effects

The following rare side effects have been reported after use of MYOVIEV:

- feeling warm
- symptoms of allergy/hypersensitivity
- headache
- shortness of breath
- low blood pressure
- swelling of the face
- dizziness
- hot flushes
- itchy or red rash

- vomiting
- metallic taste in the mouth
- disturbance in sense of smell
- mild burning sensation in the mouth
- increase in white blood cell count.

Occasionally other side effects not listed above may also occur in some patients. Tell your doctor if you notice anything else that is making you feel unwell.

Do not be alarmed by this list of side effects. You may not experience any of them.

Overdose

The dose of MYOVIEV you will receive is calculated by a qualified nuclear medicine doctor and given to you in a highly specialised setting by either the doctor or nuclear medicine technologist. Therefore the possibility of overdose is minimal.

Storage

MYOVIEV will be stored by the hospital or clinic.

The hospital or clinic will make sure that MYOVIEV is not used if the expiry date (EXP) printed on the pack has passed.

Further information

This is not all the information that is available on MYOVIEV. If you have any more questions or are not sure about anything, ask your doctor or treatment provider.

Product description

Ingredients

Each vial contains:
active ingredient -
tetrofosmin 0.23mg
other ingredients -
stannous chloride dihydrate
sodium bicarbonate
sodium gluconate
disodium sulfosalicylate
nitrogen gas in the headspace
Australian Registration Number:
AUST R 60998

Sponsor

MYOVIEV is supplied in Australia by:
GE Healthcare Australia Pty Ltd
ABN 32 001 408 402
241 O'Riordan St
Mascot NSW 2020

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