PATIENT PACKAGE INSERT IN ACCORDANCE WITH THE PHARMACISTS' REGULATIONS (PREPARATIONS) - 1986

The medicine is dispensed with a doctor's prescription only

Puri-Nethol Tablets 50 mg

Each tablet contains mercaptopurine 50 mg Inactive ingredients and allergens in the preparation: see section 2 "Important information about some of the ingredients of the medicine" and section 6 in the leaflet "Further information".

Read the leaflet carefully in its entirety before using the medicine. This leaflet contains concise information about the medicine. This leaflet contains concise information about the

medicine. If you have further questions, refer to the doctor or pharmacist.

This medicine has been prescribed for the treatment of your ailment. Do not pass it on to others. It may harm them, even if it seems to you that their ailment is similar.

1. WHAT IS THE MEDICINE INTENDED FOR?

Puri-Nethol is used to treat various types of leukemia.

Therapeutic group: Cytotoxic immunosuppressive medicines.

BEFORE USING THE MEDICINE

- Do not use the medicine:

 if you are sensitive (allergic) to the active ingredient or to any of the additional ingredients contained in the medicine
- any of the additional ingredients contained in the medicine (see section 6). if you are breastfeeding. you should not be vaccinated with the yellow fever vaccine while taking Puri-Nethol, as it may be lethal.

- While taking Puri-Nethol, as it may be lethal.

 Special warnings regarding use of the medicine
 Before treatment with Puri-Nethol, tell the doctor if:

 You were recently vaccinated or are about to be vaccinated. If you have been vaccinated with the yellow fever vaccine.

 You take Puri-Nethol, you should not have a live organism vaccines (such as:flu, measles and BCG vaccines, ect.) until the doctor confirms that this is safe for you. This is because certain vaccines may cause infections during treatment with Puri-Nethol.

 You have kidney or liver problems, since your doctor will have to check that they are functioning properly.

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 You have a condition in which your body produces amounts of an enzyme called TPMT (thiopurine methyltransferase) that are too low, as your doctor may have to adjust the dosage. You are planning to have a baby. This warning applies to both men and women. Puri-Nethol may harm your sperm or eggs (see "Pregnancy, breastfeeding and fertility" section in the leaflet).

 You have a sensitivity (allergy) to a medicine called Imuran You have a sensitivity (allergy) to a medicine called Imuran
- (azathioprine) (also used to treat cancer).
 You have, or have not, had chickenpox, shingles or hepatitis B (a liver disease caused by a virus).

- nepatitis b (a liver disease caused by a virus).

 You have a genetic disorder called Lesch-Nyhan Syndrome. If you are receiving treatment to suppress the immune system, taking Puri-Nethol may increase your risk for:

 Tumors, including skin cancer. Therefore, during treatment with the medicine, avoid prolonged exposure to sunlight, wear protective clothing and use sunscreen with a high protection factor. Lymphoproliferative disorders:

 Treatment with Puri-Nethol increases your risk for a certain
- - type of cancer called lymphoproliferative disorder. When the treatment includes a number of immunosuppressive medicines (including thiopurines), this condition may lead
- medicines (including thiopurines), unis conductions, to death.

 A combination of multiple immunosuppressive medicines, given concomitantly, increases the risk for lymphatic system disorders due to a viral infection [Epstein-Barr virus (EBV)-associated lymphoproliferative disorders].

 Taking Puri-Nethol may increase your risk for:

 Developing a severe condition called Macrophage Activation Syndrome (hyperactivation of white blood cells related to inflammation processes) which usually occurs in people suffering from certain types of arthritis.

- Inflammation processes) which usually occurs in people suffering from certain types of arthritis.
 Some patients with inflammatory bowel disease who received Puri-Nethol developed a rare and aggressive type of cancer called Hepatosplenic T-cell Lymphoma (see section 4 "Side effects").
 Use of Puri-Nethol for the treatment of inflammatory bowel disease (IBD) is an unlicensed indication.

Infections
When you are treated with Puri-Nethol, the risk of a viral, fungal

and bacterial infections is increased, and the infections can be more serious. See also section 4 "Side effects".
Tell your doctor before starting treatment whether or not you

have had chickenpox, shingles or hepatitis B (a liver disease caused by a virus).

Children and adolescents

Low blood sugar values (sweating more than usual, nausea, dizziness, confusion, etc.) have sometimes been observed in children, primarily in children under 6 years of age or who have

a low body weight.
Talk to your child's doctor if this occurs
Blood tests

Blood tests
Treatment with Puri-Nethol may affect your bone marrow. This means that you may have a lower number of white blood cells, platelets and red blood cells (less common) in your blood. At the beginning of the treatment, the doctor will refer you to do daily blood tests, and at least a weekly test during the treatment, this is in order to monitor the levels of these cells in your blood. If you stop the treatment early enough, the blood cell levels will return to normal values.
Liver function
Puri-Nethol has a toxic effect on your liver. Therefore, the doctor

Liver function

Puri-Nethol has a toxic effect on your liver. Therefore, the doctor will refer you for weekly liver function tests during treatment with Puri-Nethol. If you already have a liver disease or if you are taking other medicines that can have an effect on your liver, the frequency of the tests will increase. If you notice yellowing of the whites of the eyes or of the skin (jaundice), report this to the doctor immediately, as it may be necessary to immediately stop treatment with Puri-Nethol.

Sun and UV light

Sun and UV light

While you are taking Puri-Nethol, you are more sensitive to the sun and UV light. You must make sure to limit your exposure to the sunlight and UV light, wear protective clothing and use a sunscreen with a high sun protection factor.

If you are not sure whether the above applies to you, refer to the doctor or pharmacist before taking Puri-Nethol.

Drug interactions
If you are taking, or have recently taken other medicines, including non-prescription medicines and nutritional supplements, tell the doctor or pharmacist. Especially if you are taking: you are taking:

• Ribavirin (to treat viral infections).

- Other cytotoxic medicines (chemotherapy to treat cancer). Allopurinol, thiopurinol, oxypurinol or febuxostat (to treat
- gout).
- intestinal problem called ulcerative colitis) Sulfasalazine (to treat rheumatoid arthritis or ulcerative
- colitis) Methotrexate (to treat rheumatoid arthritis or severe
- psoriasis). Infliximab (to treat Crohn's disease and ulcerative colitis, rheumatoid arthritis, ankylosing spondylitis or severe psoriasis). Warfarin or acenocoumarol – anticoagulants (to thin the
- blood)
- Anti-epileptic medicines such as phenytoin, carbamazepine.

Blood levels of anti-epileptic medicines may need to be monitored and their dosages adjusted, if necessary.

Use of vaccines while taking Puri-Nethol If you are due to be vaccinated, consult first with the doctor or nurse. When you are taking Puri-Nethol, do not get vaccinated

nurse. When you are taking Puri-Nethol, do not get vaccinated with live vaccines (e.g., polio, measles, mumps, rubella) until the doctor confirms that this is safe for you, as certain vaccines may cause infections during treatment with Puri-Nethol.

Use of the medicine and food
See section 3 "How should you use the medicine?".

Pregnancy, breastfeeding and fertility
If you are planning to have a baby, do not take Puri-Nethol without first consulting your doctor. This warning applies to both men and women. Puri-Nethol may harm your sperm or eggs.

Reliable contraception must be used to avoid pregnancy whilst

you or your partner are taking Puri-Nethol. Both men and women should continue to use effective contraception for at least 3 months after stopping treatment. If you are already pregnant, you must talk to your doctor before taking Puri-Nethol. Women who are pregnant or who are planning to be pregnant or who are breastfeeding should not come into contact with Puri-Nethol. preparation. Do not breastfeed while taking Puri-Nethol. Consult your doctor, pharmacist or midwife.

Driving and operating machinery

Puri-Nethol is not expected to affect your ability to drive or operate machinery, but no studies have been performed to confirm this

Important information about some of the ingredients of

Important information about some of the ingredients of the medicine
Puri-Nethol tablets contain lactose. If the doctor has told you that you are sensitive to certain sugars, refer to the doctor before using the medicine.

3. HOW SHOULD YOU USE THE MEDICINE?

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Puri-Nethol should only be given to you on the instruction of a doctor who specializes in treating blood diseases.
Always use the preparation according to the doctor's instructions. Check with the doctor or pharmacist if you are uncertain about the dosage and treatment regimen of the preparation.
The dosage and treatment regimen will be determined by the doctor only. The dosage will be determined individually and depends on other treatments being given to the patient. It is important to be sure to take the medicine at designated times.
Take the medicine at least one hour before or 3 hours after food or drinking milk.
Swallow the medicine whole with a glass of water.

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 The score line on the tablet is not intended for halving the tablet. It is not recommended to halve, crush or chew the tablets

Tests and follow-up
During the course of treatment with Puri-Nethol, the doctor will
perform routine blood tests. This is to check your blood cell

count and liver function.

The doctor may perform other blood and urine tests to monitor uric acid level, which can rise during treatment with Puri-Nethol. The doctor may change the dosage during treatment in accordance with the test results.

- In adult patients the dosage may be reduced in accordance with the results of kidney and liver functions. In patients with kidney or liver problems the dosage may
- be reduced.
- If you have a TPMT enzyme deficiency the dosage may be reduced. Do not exceed the recommended dose

If you accidentally take a higher dosage
If you have taken an overdose or if a child or anyone else has accidentally swallowed the medicine, refer immediately to the doctor or proceed to a hospital emergency room and bring the package of the medicine with you.

If you forget to take the medicine
If you forget to take the medicine
If you forget to take the medicine with you.

If you forgot to take this medicine at the designated time, tell the attending doctor. **Do not take a double dose to compensate** for a forgotten dose.

If you stop taking the medicine
Adhere to the treatment regimen as recommended by the doctor. Even if there is an improvement in your health, do not stop the treatment with the medicine without consulting the doctor.

stop the treatment with the medicine without consulting the doctor.

Do not take medicines in the dark! Check the label and the dose <u>each time</u> you take medicine. Wear glasses if you need them. If you have further questions regarding use of the medicine, consult the doctor or pharmacist.

4. SIDE EFFECTS

As with any medicine use of Puri-Nethol may cause side effects.

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As with any medicine, use of Puri-Nethol may cause side effects in some users. Do not be alarmed when reading the list of side effects. You may not suffer from any of them.

Refer immediately to your specialist doctor or the hospital if any of the following symptoms appears:

An allergic reaction – signs may include:

Skin rashes

High fever

Joint pain

Swollen face

Any signs of fever or infection (sore threat sore mouth or

- Swollen face
 Any signs of fever or infection (sore throat, sore mouth or urinary problems).
 Any unexpected bruising or bleeding, as this could indicate low production of blood cells of a particular type. If you suddenly feel unwell (even if your body temperature is normal) and have abdominal pain and nausea, as this cauld be a claim of an inflamed parceas.

s normal and nave abuninal pain and haused, as this could be a sign of an inflamed pancreas.

• Yellowing of the whites of the eyes or skin (jaundice).

Consult the doctor if you have one of the following side effects:

Very common side effects – occur in more than one in ten users

A deap in the number of white blood cells and platelate (more A drop in the number of white blood cells and platelets (may

show up in blood tests)

Common side effects – occur in 1-10 in 100 users

Nausea or vomiting
 Liver damage, may show up in blood tests
 A drop in the number of red blood cells which may make you

tired, weak or breathless (called anemia)

- Loss of appetite
 Inflammation of the mouth (stomatitis)
 Uncommon side effects occur in 1-10 in 1,000 users
- Mouth ulcers Inflammation of the pancreas (pancreatitis)
- Joint pain

- Skin rash
 Fever
 Irreversible damage to the liver (hepatic necrosis)
 Rare side effects occur in 1-10 in 10,000 users
 Hair loss
 In most temperary law anomy count.
- In men: temporary low sperm count
- Swollen face Various types of cancer including blood, lymph and skin
- cancer
- Very rare side effects occur in less than one in 10,000 users
- 0,000 users
 A different type of leukemia than that which is being treated
 Cancer of the spleen and liver (in patients suffering from
 inflammatory bowel disease [IBD])
 Ulcers in the intestines; symptoms may include abdominal

octes in the intestines, symptoms may include addominar pain and bleeding
Side effects with unknown frequency (effects whose frequency have not been determined yet)
A rare type of cancer – Hepatosplenic T-cell Lymphoma (see section 2 "Special warnings regarding use of the

medicine") Increased sensitivity to sunlight and UV light causing skin reactions

Additional side effects in children

Additional side effects in children
Low blood sugar levels (sweating more than usual, nausea,
dizziness, confusion, etc.) have been reported in some children
taking Puri-Nethol. The frequency of this effect is not known;
however, most of the children were under the age of 6 years
and had a low body weight.
If a side effect occurs, if one of the side effects worsens
or if you suffer from a side effect not mentioned in this
leaflet, consult the doctor.
Reporting of side effects:
Side effects can be reported to the Ministry of Health by clicking
on the link "Report Side Effects of Drug Treatment" found on the
Ministry of Health homepage (www.health.gov.il) that directs
you to the online form for reporting side effects, or by entering
the link:

https://sideeffects.health.gov.il
Additionally, you can report to the company via the following address

- Padagis.co.il
 HOW SHOULD THE MEDICINE BE STORED?
 Avoid poisoning! This medicine, and any other medicine, should be stored in a safe place out of the reach and sight the stored in the of children and/or infants in order to avoid poisoning. Do not induce vomiting unless explicitly instructed to do so by the
- doctor. Do not use the medicine after the expiry date (exp. date) that appears on the carton package and the bottle label. The expiry date refers to the last day of that month.
 Store in the original package below 25°C, in a dry place. Keep the bottle tightly closed.
- If your doctor instructs you to stop treatment, it is important to return all remaining medicine to the pharmacist so that he/she can dispose of it according to the guidelines for disposal of dangerous substances. Only keep the remainder of the medicine with you upon specific instruction from the
- doctor.
 FURTHER INFORMATION In addition to the active ingredient, the medicine also contains:

Lactose monohydrate, maize starch, modified maize starch, magnesium stearate, stearic acid.

What the medicine looks like and contents of the

package:
Round and convex, pale yellow tablets, with a line (which is not used for halving) on one side, where PT is marked above the line and 50 is marked below it, in a glass bottle containing 25 tablets

- Registration holder: Padagis Israel Agencies Ltd., 1 Rakefet Shoham Manufacturer and address: Aspen Pharma Trading Ltd.
- Dublin, Ireland. Revised in November 2022 according to MOH guidelines.
 Registration number of the medicine in the National Medicines Registry of the Ministry of Health: 3344.22532