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

This medicine is dispensed without a doctor's prescription

liptipot Novimol®

Suspension Each 1 ml contains 100 mg paracetamol Inactive ingredients and allergens in the

preparation - see section 6 - "Additional information" and the "Important information about some ingredients of the medicine" section. Read the entire leaflet carefully before using the medicine. This leaflet contains concise information about the medicine. If you have any other questions, refer to the doctor or the pharmacist.

Use the preparation according to the instructions in the dosage section of this leaflet. Consult the pharmacist if you have further questions. Refer to the doctor if the fever persists for more than 3 days or if the symptoms do not resolve within

5 days despite use of the medicine. 1. What is the medicine intended for?

The medicine is intended to relieve pain and reduce fever. Therapeutic class: analgesic and

antipyretic. 2. Before using the medicine: Do not use this medicine if:

 There is a known sensitivity to paracetamol or to any of the other ingredients the medicine

contains (see section 6). Special warnings regarding the

use of the medicine:

 If you developed skin side effects in the past as a result of taking preparations containing paracetamol, do not take preparations containing paracetamol, so that severe skin side effects will not recur

· The preparation contains

paracetamol, which may cause liver damage when:

 Given at a higher dosage than recommended or for a prolonged period of time.

 Alcoholic beverages are consumed during the course of treatment.

Additional medicines which affect liver function are taken.

· In patients with glutathione deficiency, use of paracetamol may increase the risk for metabolic acidosis.

· Do not use this medicine frequently without consulting the doctor.

 Do not take additional antipyretics and analgesics or cold medicines without consulting a doctor or a pharmacist, to prevent paracetamol overdose or poisoning.

 Do not take other medicines from the same family and/or other preparations containing paracetamol.

 Avoid taking a high dosage (within the recommended limit) of this

medicine when fasting. If you are sensitive to any type of food or medicine, inform your doctor before starting treatment with this medicine.

 Consult a doctor regarding use of this medicine in children under 3 vears of age

 Consult a doctor about using this preparation or another preparation for pain relief and/or to reduce fever.

■ Before treatment with Tiptipot Novimol, inform the doctor if:

 You have or had liver disease or impaired liver function.

You have or had impaired kidney

· You have or had alcoholism

You have or had jaundice.

 You are pregnant or breastfeeding ■ Drug-drug interactions: If you are taking or have recently taken other medicines including non-prescription medicines and food supplements, tell the doctor or the pharmacist. Especially if you are taking:

 Anticoagulants, especially warfarin

 Preparations that stimulate production of enzymes in the liver (e.g. rifampicin, barbiturates) Medicines for epilepsy -

phenytoin, carbamazepine Non-steroidal anti-inflammatory

 Metoclopramide or domperidone (for treatment of nausea, vomiting and other digestive problems).

 Medicines that reduce the stomach emptying rate (e.g., propantheline, anticholinergic antidepressants and narcotic analgesics) will compromise the absorption of paracetamol.

Chloramphenicol (antibiotic)

 Probenecid (for treatment of Cholestyramine (for reduction of

excess blood lipids). Other analgesics or antipyretics

and anti-arthritis medicines. ■ Use of paracetamol and alcohol consumption:

During treatment with paracetamol do not consume alcohol due to increased risk of liver damage

■ Pregnancy, breastfeeding and fertility:

If you are pregnant or breastfeeding, consult a doctor before you start using the medicine.

■ Use in children:

Parents must report to the treating doctor all side effects and any additional medicine given to the child.

■ Important information about some ingredients of the medicine: This medicine contains 140 mg of sorbitol in each 1 ml. Sorbitol is a source of fructose. If there is a known intolerance to certain sugars or a diagnosis of hereditary fructose intolerance, consult with your doctor before taking this medicine. This medicine contains 1 mg of sodium benzoate in each 1 ml.

360 3.60 Up to 5 times 25 kg | 375 | 3.75 | Up to 5 times 26 kg 390 3.90 Up to 5 times 27 kg | 405 | 4.05 | Up to 5 times 28 kg | 420 | 4.20 | Up to 5 times 29 kg | 435 | 4.35 | Up to 5 times 30 kg 450 4.50 Up to 5 times

Age table

This medicine contains less than

23 mg of sodium per 1 ml, and is

3. How should you use the

if you are uncertain about the

dosage and how to use the

medicine?

preparation.

weight.

age only.

Weight table

child's body weight

8 kg | 120 | 1.20

10 kg | 150 | 1.50

11 kg | 165 | 1.65

12 kg | 180 | 1.80

13 kg | 195 | 1.95

14 kg | 210 | 2.10

15 kg | 225 | 2.25

16 kg | 240 | 2.40

17 kg 255 2.55

18 kg | 270 | 2.70

19 kg 285 2.85

20 kg | 300 | 3.00

21 kg 315 3.15

22 kg | 330 | 3.30

23 kg 345 3.45 Up to 5 times

135 1.35

Child's mg

weiaht

9 kg

following tables

therefore considered sodium-free.

Check with the doctor or pharmacist

The generally accepted dosage is:

Find the suitable dosage in the

the dosage in the weight table

indicating dosage according to

Only if the child's weight is not

determined according to age, as

shown in the age table indicating

dosage according to the child's

The dosage according to the child's

weight is calculated as 15 mg/kg of

the child's weight, per dose. In other

words. 0.15 ml for every kg of the

Dose

lin ml

Maximum

hours

number of

doses per 24

Up to 5 times

known - the dosage will be

If you know the child's weight - find

In case the child's weight is unknown:

Children of identical ages can be of significantly different weights. Therefore, an effort must be made to find out the child's weight and determine the dosage according to the weight table.

Only if it is impossible to find out the child's weight - the dosage can be determined according to the age table.

	Child's age	Dose in ml	Maximum number of doses per 24 hours
	1-2 years	1.20 ml	Up to 5 times
	2-3 years	1.60 ml	Up to 5 times
	4-5 years	2.40 ml	Up to 5 times

Doses should be taken/administered in intervals of at least 4 hours, as necessary, up to 5 doses in 24 hours.

Do not exceed the recommended

dose. Refer to the doctor if the fever persists for more than 3 days or if the symptoms do not resolve within 5 days despite use of the medicine.



Find the required amount in the dosage table, in the "Dose in ml" column.

Insert the syringe into the opening in the bottle's neck.

3. To fill the syringe, turn the bottle upside-down. With the syringe held in place, pull the plunger gently downwards to draw the medicine up to the appropriate mark on the syringe.

4. Turn the bottle upside-down again (right side up) and gently remove the syringe from the bottle's opening

5. Empty the contents of the syringe slowly into the child's mouth. towards the inner wall of the cheek.

6. Wash the two parts of the syringe well with lukewarm water. With liquid medicines, use the syringe or dropper intended for measuring the proper amount of medicine. If a measuring device is not included in the package, consult a pharmacist. Do not use a household teaspoon to measure the amount of medicine. Household teaspoons vary in size and it is likely you will not receive the correct amount of medicine. If you or your child took an

bring the package of the medicine

without an explicit instruction from

the doctor! Even if you feel well/the

child feels well, immediate treatment

developing severe liver damage.

vomiting, diarrhea, loss of appetite.

abdominal pain, swelling, increased

sweating, pain or tenderness in the

upper abdomen, and they may not

reflect the severity of liver damage.

Do not take medicines in the

dark! Check the label and the

dose every time you take the

medicine. Wear glasses if you

need them.

with you. Do not induce vomiting

is essential, due to the risk of

Side effects can be nausea and

overdose or if a child accidentally swallowed this medicine. immediately refer to a doctor or to a hospital emergency room and

Method of use:

Be sure to measure the dose with the included syringe Shake well before use.



If you have any other questions regarding use of the medicine. consult the doctor or the pharmacist.

4. Side effects

As with any medicine, use of Tiptipot Novimol may cause side effects. such as dizziness, in some users Do not be alarmed when reading the list of side effects. You may not experience any of them

Severe side effects: Stop using this medicine and refer to a doctor immediately if:

 Severe allergic reactions occur. such as rash, itching, swelling of the face, lips, tongue, throat and/ or extremities that can cause difficulty breathing or swallowing. Paracetamol may, in rare cases,

cause the appearance of severe skin diseases whose symptoms can be: redness, rash, blisters. widespread skin damage Severe skin side effects may appear even if you had no problems in the past taking preparations containing the active ingredient paracetamol. If skin side effects appear. discontinue treatment and refer to a

doctor immediately Signs of changes in the blood system occur, such as: bleeding bruises, development of infections

more easily. Additional very rare side effects side effects that occur in less than one out of 10 000 users: Bronchospasm, liver failure thrombocytopenia (low platelets). 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 vour doctor.

Side effects may be reported to the Ministry of Health by clicking on the link "Report side effects due to medicinal treatment" found on the Ministry of Health website homepage (www.health.gov.il). which will direct you to the online form for reporting side effects, or by clicking on the following link: https://sideeffects.health.gov.il/

5. How to store the medicine? Avoid poisoning! This medicine and any other medicine must be kept in a closed place out of the reach and sight of children and/or infants to avoid poisoning. Do not induce vomiting without an explicit instruction from the doctor Do not use the medicine after the expiry date (Exp.) appearing on the

Storage:

Store at a temperature lower than 25°C

package. The expiry date refers to

the last day of that month

The medicine may be used up to 6 months after the first opening. 6. Additional information In addition to the active ingredient. the medicine also contains: Sorbitol Solution 70%, Glycerol, Xanthan Gum, Sucralose Strawberry Cream Flavour, Sodium Benzoate, Citric acid, FD&C Red No. 40. Purified Water.

What does the medicine look like and what are the contents of the package:

A glass bottle containing 50 ml of a pink-red suspension with white particles, with a strawberry fragrance.

Manufacturer and license holder: CTS Chemical Industries Ltd 3 Hakidma st., Kiryat Malachi 83057

The format of this leaflet was determined by the Ministry of Health and its content was checked and approved by the Ministry of Health on 09/2013, and has been updated in accordance with the Ministry of Health instructions on 07/2020. Registration number of the medicine in the national drug registry of the Ministry of Health: 1346131137



