The medicine is dispensed with a doctor's prescription only

Hemangiol® **Oral Solution**

Active ingredient: The active ingredient is propranolol. Each 1 mL of solution contains 3.75 mg propranolol (equivalent to 4.28 mg propranolol hydrochloride).

Inactive and allergenic ingredients in the preparation: see section 2 "Important information about some of the ingredients of the medicine" and section 6 "Further information".

Read this leaflet carefully in its entirety before using 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 your child only. Do not pass it on to others. It may harm them, even if it seems to you that their illness is similar.

In addition to this leaflet, Hemangiol also has an information guide for the patient's parents. This guide contains important safety information, that you must know and follow before starting and during treatment with Hemangiol. Read both the information guide for the patient's parents and the patient leaflet before starting to use the medicine. Keep the information guide for further reference, if necessary

WHAT IS THE MEDICINE INTENDED FOR? The medicine is intended to treat proliferating

infantile hemangioma requiring systemic therapy:

- the hemangioma is life- or functionthreatening. the hemangioma is ulcerated and painful.
- and/or does not respond to simple wound
- the hemangioma is with a risk of permanent scarring or disfigurement. Hemangiol treatment is started in infants aged

5 weeks to 5 months. A hemangioma is a collection of extra blood

vessels that form a lump in or under the skin. A hemangioma can be superficial or deep. It is sometimes called a 'strawberry mark' because its surface resembles the shape of a strawberry.

Therapeutic group: The active ingredient propranolol belongs to a

group of medicines known as beta-blockers.

BEFORE USING THE MEDICINE

Do not use the medicine if: your child was born prematurely and

- has not reached the corrected age of 5 weeks (the corrected age being the age a premature baby would be if he/she had been born on their due date). your child is allergic (sensitive) to the active
- ingredient propranolol or to any of the other ingredients contained in this medicine (see section 2 "Important information about some of the ingredients of the medicine" and section 6 "Further information" in this leaflet). An allergic reaction can include a skin rash, itching or shortness of breath.
- your child has asthma or a history of breathing difficulties. your child has a slow heart rate for his age.
- Refer to the doctor if you are not sure. your child has a heart problem (such as heart rhythm disorders or heart failure).
- · your child has very low blood pressure. your child has a circulation problem which
- causes the toes and fingers to be numb and pale. your child is prone to low blood sugar level.
- your child has high blood pressure caused by a tumor in the adrenal gland. This tumor is called phaeochromocytoma. Do not give this medicine to your child if
- you are breastfeeding and taking medicines that must not be taken together with Hemangiol (see section "Breastfeeding" and section "Drug interactions"). Special warnings regarding use of the

medicine Before giving Hemangiol to your child, tell your child has liver or kidney problems. This medicine is not recommended in case of liver

- or kidney problems. your child has ever had an allergic reaction,
- regardless of its origin (e.g., from medicines or from an alimentary substance, etc.). An allergic reaction can include a rash, itching or shortness of breath.
- your child has psoriasis [a skin condition that causes the formation of red, dry plaques of thickened skin], as this medicine may worsen the symptoms of this condition. vour child has diabetes: in this case, monitor
- his blood sugar level more frequently. your child has PHACE syndrome (a condition
- combining hemangioma with vascular abnormalities including cerebral blood vessels), as this medicine may increase the risk of cerebral stroke. Important signs to pay attention to after administering Hemangiol to your child:

Risk of low blood sugar level

(hypoglycemia) This medicine may mask the warning signs of hypoglycemia (also known as low blood sugar level). The medicine may also aggravate

hypoglycemia in children, especially during a fasting period (e.g., low food intake, concomitant infection, vomiting) or when glucose demands are increased (cold, stress, infections), or in case of overdose. The warning signs may be: Minor: pallor, tiredness, sweating, shaking, palpitations, anxiety, hunger, difficulty waking

- · Major: sleeping more than usual, difficulty responding, poor feeding, decrease in body
- temperature, convulsions, brief pauses in breathing, loss of consciousness. The risk of hypoglycemia remains significant throughout the whole treatment period with the

medicine. To avoid risk of low blood sugar level

(hypoglycemia), give Hemangiol to your child during or straight after a meal and avoid giving the last dose close to night bedtime (see section 3 "How should you use the medicine?"). You must feed your child sufficiently and regularly during treatment. If your child is not eating enough or is vomiting, it is recommended to skip the dose. Do not give Hemangiol to your child until he/she starts eating normally If your child shows any signs of low blood sugar level (hypoglycemia) while taking

Hemangiol, stop treatment with the medicine and contact a doctor immediately or proceed to a hospital. If the child is conscious, give him/her a drink of liquid containing sugar. Risk of bronchospasm

Stop treatment and contact a doctor immediately

if after giving Hemangiol to your child, you observe any of the following symptoms that could be indicative of bronchospasm (breathing difficulties due to temporary narrowing of the airways in the lungs): cough, rapid breathing or difficulty breathing, or wheezing with or without bluish-colored skin. Stop treatment with the medicine and contact a doctor immediately if your child

shows cold-like symptoms associated with difficulty in breathing and/or wheezing during the course of treatment with Hemangiol.

Risk of hypotension and low heart rate (bradycardia) Hemangiol can decrease blood pressure and heart rate (bradycardia); therefore, your child

must be kept under close medical surveillance and heart rate monitoring for two hours after administration of the first dose or after the dose of the medicine is increased. Then, the doctor should regularly examine the child during treatment

Stop treatment with the medicine and contact a doctor immediately if your child has any signs of tiredness, coldness, pallor, bluish-colored skin, or fainting while using

Hemangiol.

Risk of high potassium level in the blood (hyperkalemia)

Hemangiol may increase blood potassium level (hyperkalemia). In case of a large ulcerated hemangioma, your child's blood potassium level should be monitored.

If your child is due to undergo general anesthesia

Tell the doctor that your child is receiving Hemangiol. This is because your child may suffer from low blood pressure if given certain anesthetics while taking this medicine (see section "Drug interactions"). It may be necessary to stop taking Hemangiól for at least 48 hours before anesthesia.

Tests and follow-up

- · If your child has diabetes, monitor his blood sugar level more frequently (see "Special warnings regarding use of the medicine").
- Hemangiol can cause a decrease in blood pressure and heart rate. Therefore, your child must be under close medical monitoring, with heart rate monitoring for 2 hours after administration of the first dose or after a dosage increase. Then, the doctor must regularly examine the child during the course of treatment. (see "Important signs to pay attention to after administering Hemangiol to your child").
- Hemangiol may increase blood potassium level. In case of a large ulcerated hemangioma, your child's blood potassium level should be monitored (see "Important signs to pay attention to after administering Hemangiol to your child").

Drug interactions

- If your child is taking, or has recently taken, other medicines, including nonprescription medicines and nutritional supplements, tell the doctor or pharmacist. This is because Hemangiol can change the way other medicines act, and some medicines can have an effect on the way Hemangiol acts.
- If you are breastfeeding your child, it is important to tell your doctor, pharmacist or the medical staff if you are taking any medicine, as it may pass into breast milk and influence your child's treatment. Your doctor will advise you on whether you should stop breastfeeding or not. Tell the doctor or pharmacist if your child is

taking or if you are breastfeeding and taking: Medicines to treat diabetes.

- · Medicines to treat heart or blood vessel
- problems, such as heart rate disturbances, chest pain or angina, high blood pressure, heart failure. Medicines to treat anxiety and depression, as well as medicines for more serious mental
- health problems and epilepsy. Medicines to treat tuberculosis.
- · Medicines to treat pain and inflammation.
- Medicines to lower blood lipid level. · Medicines for anesthesia.
- If you have further questions, consult your

doctor or pharmacist. Use of the medicine and food

The medicine should be given immediately after or while feeding the child. Breastfeeding

If you are breastfeeding your child Tell the doctor before giving this medicine to

- · Do not give this medicine to your child if
- you are taking medicines that must not be used with Hemangiol (see section "Drug interactions") Important information about some of the

ingredients of the medicine The medicine contains less than 1 mmol (23 mg) sodium per dose (essentially "sodium-free").

The medicine contains 2.08 mg propylene glycol/kg/day. If your baby is less than 4 weeks old, consult the doctor or pharmacist before giving this medicine, especially if the baby is taking other medicines that contain propylene glycol or alcohol. 3. HOW SHOULD YOU USE THE MEDICINE?

The treatment of your child is in accordance

with the recommendation of a doctor who has expertise in the diagnosis and treatment of infantile hemangioma. Always use the preparation according to the doctor's instructions. Check with your doctor or pharmacist if you are not sure about the dosage and treatment regimen of the preparation. The dosage and treatment regimen will be determined by the doctor only.

Never change the dosage given to your child

on your own. Every increase in dose or dose adjustment in accordance to your baby's weight must be done by the attending doctor · Dosage:

The dosage will be determined in accordance with your baby's weight, as per the following dosage regimen: Maaks Dose for Timing of

(daily dose)	oral intake	doses
First week (1 mg/kg/day)	0.5 mg/kg	one dose in the morning additional dose in late afternoon at an interval of at least 9 hours between the 2 doses
Second week (2 mg/kg/day)	1 mg/kg	
Third and following weeks (3 mg/kg/day)	1.5 mg/kg	
If necessary, you may mix the medicine with a small quantity of baby-milk or age-suitable		

apple and/or orange juice, and give it to your child in a baby bottle. Do not mix the medicine with a full bottle of milk or fruit juice. For children weighing up to 5 kg, you may mix the dose with one teaspoonful (approximately 5 mL) of baby-milk. For children weighing more

than 5 kg, the dose may be mixed with one tablespoonful (approximately 15 mL) of babymilk or age-suitable apple or orange juice. Use the mixture within 2 hours of preparation Do not exceed the recommended dose.

How to give Hemangiol to your child: Hemangiol is for oral use only.

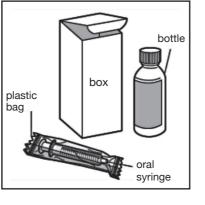
Give the medicine immediately after or while feeding the child.

- Always measure the dose by using the oral syringe supplied together with the bottle and do not replace it with any other syringe. If
- you lost the syringe, contact the registration Administer Hemangiol directly into your child's mouth using the oral syringe supplied with the bottle.
- Feed your child regularly to avoid a prolonged fast. If your child does not eat sufficiently or is vomiting, skip the dose.
- If your child spits up the dose or if you are uncertain whether he/she got the full dose, do not give another dose; wait until the next
- scheduled dose. Administration of Hemangiol and feeding the child must be done by the same person in order to avoid the risk of low blood sugar level (hypoglycemia) in your child. If different
- people are involved in treating the child, good communication between them is essential in order to ensure the safety of the child. Instructions for use: Step 1 - Remove the items from the box The box contains the following items that you will need to administer the medicine

1. A glass bottle containing 120 mL Hemangiol

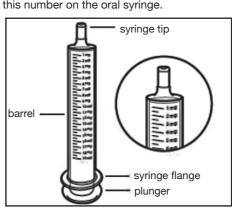
oral solution.

An oral measuring syringe, graduated in mg, provided with this medicine. Remove the bottle and the oral syringe from the box, and remove the syringe from the plastic bag



Step 2 - Check the dose

Check the Hemangiol dose in milligrams (mg) as prescribed for your child by the doctor. Locate this number on the oral syringe.

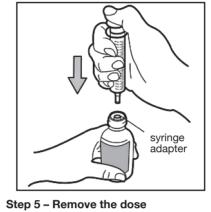


Step 3 - Open the bottle The bottle has a child-proof cap. Open the bottle

as follows: push the plastic cap downwards while turning the cap counter-clockwise (to the left). Do not shake the bottle before use

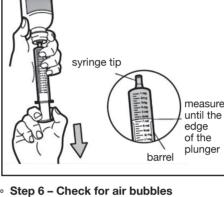


Insert tip of the oral syringe into the upright bottle and push the plunger of the syringe all the way down. Do not remove the syringe adapter from the neck of the bottle. Only use the oral syringe that is supplied with the medicine to measure and administer the dose. Do not use a spoon or any other measuring tool.



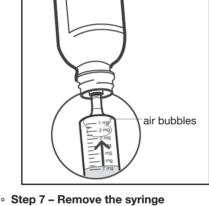
With the oral syringe in place, turn the bottle

upside down. Pull the plunger of the syringe up to the number of milligrams you need.



If you see air bubbles in the syringe, hold the

syringe upright, push the plunger upwards until all the large air bubbles have been pushed out and then readjust to the dose prescribed for your child by the doctor.

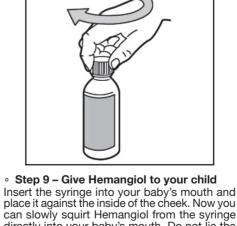


Turn the bottle upright and remove the entire syringe from the bottle. Be careful not to push

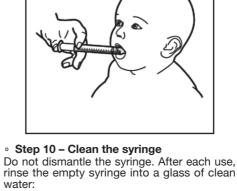
the plunger in during this step.



it clockwise (to the right).



directly into your baby's mouth. Do not lie the baby down immediately after administering the



1. Take a glass of clean water

2. Pull the plunger 3. Discard the water into the sink

4. Repeat this cleaning process 3 times

Do not use any soap or alcohol-based product for cleaning. Wipe the outside dry. Do not put the syringe in a sterilizer or a dishwasher. Until the next use, store the bottle and the syringe together in the medicine package, in a safe place where your child will not be able to see or reach it. Discard the syringe once the bottle is finished.

If you accidentally give your child a higher dosage of the medicine than required, contact the doctor immediately or proceed to a hospital emergency room, and bring the package of the medicine with you. If you forget to give the medicine to your child

Skip the forgotten dose and do not give a double dose to make up for the forgotten dose. Continue the treatment at the usual frequency: one dose in the morning and one dose in the late afternoon. Adhere to the treatment regimen recommended by the doctor. If you stop giving the medicine to your child

The medicine may be stopped at once at the end of the treatment, as decided by the doctor. Do not take medicines in the dark! Check

the label and the dose each time 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 FEFECTS

As with any medicine, use of Hemangiol can cause side effects in some users. Do not be alarmed by reading the list of side effects. Your child may not suffer from any of them. After administering Hemangiol, pay attention to important warning signs associated with the following potential side effects: low blood pressure, slow heart rate, low blood sugar level, and bronchospasm (breathing difficulties). See section 2 of this leaflet.

Very common side effects (effects that occur in more than 1 user in 10):

- · Inflammation of the bronchi (bronchitis) Sleep disorders (insomnia, poor quality of sleep and difficulties waking up)
- Diarrhea and vomiting

Common side effects (effects that occur in 1-10 in 100 users): Bronchospasm (breathing difficulties)

Bronchiolitis – inflammation of the small

- bronchi in the lungs, with breathing difficulties
- and wheezing in the chest, accompanied by cough and fever Decreased blood pressure
- · Decreased appetite Agitation, nightmares, irritability
- Somnolence

Skin redness

- Cold extremities Constipation, abdominal pain
- Diaper rash Uncommon side effects (effects that occur in 1-10 in 1,000 users):
- Heart conduction or heart rhythm disorders (slow or uneven heartbeats)
- Urticaria (allergic reaction of the skin), hair loss (alopecia)
- Decreased blood sugar levels · Reduction in the number of white blood cells
- Side effects of unknown frequency (effects whose frequency has not been determined yet):
- Convulsions linked to a low blood sugar level (hypoglycemia)
- · Bradycardia (abnormally slow heart rate) Low blood pressure Very low levels of white blood cells that fight
- infection · Circulation problems which cause pallor and
- numbness in the fingers and toes Elevated blood potassium level

If a side effect occurs, if one of the side effects worsens or if your child suffers from a side effect not mentioned in this leaflet, consult with the doctor.

Reporting 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:

In addition, you can report to Padagis via the following address: Padagis.co.il

https://sideeffects.health.gov.il

HOW SHOULD THE MEDICINE BE STORED? Avoid poisoning! This medicine and any other medicine should be kept in a closed place

- out of the reach and sight 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 package and the bottle label. The expiry date refers to the last day of that month. Storage conditions: Store below 25°C. Do not freeze. Store the

bottle in the original package to protect from

Store below 25°C

Between uses, store the bottle and oral syringe together in the original package. After first opening: may be used for two

months, but no later than the expiry date.

· Do not discard medicines into the wastewater

or household waste. Consult the pharmacist on how to dispose of medicines no longer used. These measures will help protect the environment. 6. FURTHER INFORMATION

Propylene glycol 92%, Ethyl butyrate, Vanillin, Undecalactone/Gamma, Ethyl acetate, Ethyl propionate, Maltol, Hydroxy-2-5-dimethyl-3(2h)furanone/4-, Hexyl alcohol, Hexen-1-Ol/Cis-3, Linalool, Isoamyl butyrate, Ethylhexanoate, Ethyl-2-methylbutyrate, Hexen-1-Yl acetate/ Cis-3-, Decalactone/gamma-, Methylbutyric acid/2-, Hydroxyphenyl)-2-butanone/4(para-, Methyl dihydrojasmonate, Methyl cinnamate, Ethyl isovalerate, Hexenal/trans-2-, Hexanoic acid, Diacetyl natural, Isovaleric acid, Ionone/ alpha, Ionone/beta, Ionone/gamma, Tocopherol/ alpha], Citric acid monohydrate, Purified water. For further information, see section 2 "Important information about some of the ingredients of the What the medicine looks like and the contents of the package: Hemangiol is a clear, colorless to slightly yellow,

the medicine contains a dark brown glass bottle containing 120 mL Hemangiol solution, which is closed with a child-proof cap. The package

contains an oral measuring syringe Registration Holder and Importer: Padagis Israel Agencies Ltd., 1 Rakefet St.,

National Drug Registry of the Ministry of Health: 15968.35287

In addition to the active ingredient, the medicine also contains: Hydroxyethylcellulose (250 grade), Vanilla flavor (IFF*SC011851) [contains Propylene glycol 75.2%, Vanillin, Water, Butyric acid, Ethyl butyrate, Piperonal], Saccharin sodium, Strawberry flavor (IFF*SN864761) [contains

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oral solution with a fruity odor. The package of