

## 1. NAME OF THE MEDICINAL PRODUCT

GP-Betahistine Tablets 24mg

## 2. QUALITATIVE AND QUANTITATIVE COMPOSITION

Each tablet contains 24mg betahistine dihydrochloride equivalent to 15.63 mg betahistine.

Each tablet contains 24 mg of mannitol.

For the full list of excipients, see section 6.1

## 3. PHARMACEUTICAL FORM

Tablet

White colour, round (approx 10.00mm) & biconvex uncoated tablet scored on one side with the embossing "II" on either sides of the score and plain on other side.

The score line is only to facilitate breaking for ease of swallowing and not to divide into equal doses.

## 4. CLINICAL PARTICULARS

### 4.1 Therapeutic indications

Vertigo, tinnitus and hearing loss associated with Ménière's syndrome.

### 4.2 Posology and method of administration

*Posology*

Adults:

The dosage for adults is 24-48mg divided over the day. The 24mg strength can be taken 2 times daily. Daily dose should not exceed 48mg. The dosage should be individually adapted according to the response. Improvement can sometimes only be observed after a couple of weeks of treatment.

Paediatric population: not recommended for use in children below 18 years due to insufficient data on safety and efficacy.

Renal impairment: There are no specific clinical trials available in this patient group.

Hepatic impairment: There are no specific clinical trials available in this patient group.

*Method of Administration*

The score line is only to facilitate breaking for ease of swallowing and not to divide into equal doses.

### 4.3 Contraindications

Hypersensitivity to the active substance(s) or to any of the excipients listed in section 6.1.

Phaeochromocytoma.

### 4.4 Special warnings and precautions for use

Caution is advised in the treatment of patients with a history of peptic ulcer. Clinical intolerance to Betahistine in bronchial asthma patients has been shown in a relatively few patients. These patients need to be carefully monitored during the therapy.

Excipient Warnings

This product contains mannitol which may have a mild laxative effect.

### 4.5 Interaction with other medicinal products and other forms of interaction

No *in-vivo* interaction studies have been performed. Based on *in-vitro* data no *in-vivo* inhibition on Cytochrome P450 enzymes is expected.

*In vitro* data indicate an inhibition of betahistine metabolism by drugs that inhibit monoamino-oxidase (MAO) including MAO subtype B (e.g. selegiline). Caution is recommended when using betahistine and MAO inhibitors (including MAO-B selective) concomitantly.

As betahistine is an analogue of histamine, interaction of betahistine with antihistamines may in theory affect the efficacy of one of these drugs.

### 4.6 Fertility, Pregnancy and lactation

Pregnancy

There are no adequate data from the use of betahistine in pregnant women.

Animal studies do not indicate direct or indirect harmful effects with respect to reproductive toxicity at clinically relevant therapeutic exposure. As a precautionary measure, it is preferable to avoid the use of betahistine during pregnancy.

Lactation

It is not known whether betahistine is excreted in human milk.

Betahistine is excreted in rat milk. Effects seen post-partum in animal studies were limited to very high doses. The importance of the drug to the mother should be weighed against the benefits of nursing and the potential risks for the child.

Fertility

Animal studies did not show effects on fertility in rats.

### 4.7 Effects on ability to drive and use machines

Vertigo, tinnitus and hearing loss associated with Ménière's syndrome can negatively affect the ability to drive and use machines. In clinical studies specifically designed to investigate the ability to drive and use machines betahistine had no or negligible effects.

### 4.8 Undesirable effects

The following undesirable effects have been experienced with the below indicated frequencies in betahistine-treated patients in placebo-controlled clinical trials [very common ( $\geq 1/10$ ); common ( $\geq 1/100$  to  $< 1/10$ ); uncommon ( $\geq 1/1,000$  to  $< 1/100$ ); rare ( $\geq 1/10,000$  to  $< 1/1,000$ ); very rare ( $< 1/10,000$ )].

Immune System disorders

Hypersensitivity reactions e.g. anaphylaxis have been reported.

Nervous System disorders

Common: headache

In addition to those events reported during clinical trials, the following undesirable effects have been reported spontaneously during post-marketing use and in scientific literature. A frequency cannot be estimated from the available data and is therefore classified as "not known".

Gastrointestinal disorders

Common: nausea and dyspepsia

Mild gastric complaints (e.g. vomiting, gastrointestinal pain, abdominal distension and bloating) have been observed.

These can normally be dealt with by taking the dose during meals or by lowering the dose.

Skin and subcutaneous tissue disorders

Cutaneous and subcutaneous hypersensitivity reactions have been reported, in particular angioneurotic oedema, urticaria, rash, and pruritus.

### 4.9 Overdose

A few overdose cases have been reported. Some patients experienced mild to moderate symptoms with doses up to 640 mg (e.g. nausea, somnolence, abdominal pain). More serious complications (e.g. convulsion, pulmonary or cardiac complications) were observed in cases of intentional overdose of betahistine especially in combination with other overdosed drugs. Treatment of overdose should include standard supportive measures.

## 5. PHARMACOLOGICAL PROPERTIES

### 5.1 Pharmacodynamic properties

Pharmacotherapeutic group: Anti-vertigo preparations. ATC-Code: N07CA01

The mechanism of action of betahistine is only partly understood. There are several plausible hypotheses that are supported by animal studies and human data:

Betahistine affects the histaminergic system:

Betahistine acts both as a partial histamine H1-receptor agonist and histamine H3-receptor antagonist also in neuronal tissue, and has negligible H2-receptor activity. Betahistine increases histamine turnover and release by blocking presynaptic H3-receptors and inducing H3-receptor downregulation.

Betahistine may increase blood flow to the cochlear region as well as to the whole brain:

Pharmacological testing in animals has shown that the blood circulation in the striae vascularis of the inner ear improves, probably by means of a relaxation of the precapillary sphincters of the microcirculation of the inner ear. Betahistine was also shown to increase cerebral blood flow in humans.

#### Betahistine facilitates vestibular compensation:

Betahistine accelerates the vestibular recovery after unilateral neurectomy in animals, by promoting and facilitating central vestibular compensation; this effect characterized by an up-regulation of histamine turnover and release, is mediated via the H3 Receptor antagonism. In human subjects, recovery time after vestibular neurectomy was also reduced when treated with betahistine.

#### Betahistine alters neuronal firing in the vestibular nuclei:

Betahistine was also found to have a dose dependent inhibiting effect on spike generation of neurons in lateral and medial vestibular nuclei.

The pharmacodynamic properties as demonstrated in animals may contribute to the therapeutic benefit of betahistine in the vestibular system.

The efficacy of betahistine was shown in studies in patients with vestibular vertigo and with Ménière's disease as was demonstrated by improvements in severity and frequency of vertigo attacks.

### **5.2. Pharmacokinetic properties**

#### Absorption:

Orally administered betahistine is readily and almost completely absorbed from all parts of the gastro-intestinal tract. After absorption, the drug is rapidly and almost completely metabolized into 2-pyridylacetic acid. Plasma levels of betahistine are very low. Pharmacokinetic analyses are therefore based on 2-PAA measurements in plasma and urine.

Under fed conditions C<sub>max</sub> is lower compared to fasted conditions. However, total absorption of betahistine is similar under both conditions, indicating that food intake only slows down the absorption of betahistine.

#### Distribution:

The percentage of betahistine that is bound by blood plasma proteins is less than 5 %.

#### Biotransformation:

After absorption, betahistine is rapidly and almost completely metabolized into 2-PAA (which has no pharmacological activity).

After oral administration of betahistine the plasma (and urinary) concentration of 2-PAA reaches its maximum 1 hour after intake and declines with a half-life of about 3.5 hours.

#### Excretion:

2-PAA is readily excreted in the urine. In the dose range between 8 and 48 mg, about 85% of the original dose is recovered in the urine. Renal or faecal excretion of betahistine itself is of minor importance.

#### Linearity:

Recovery rates are constant over the oral dose range of 8 – 48 mg indicating that the pharmacokinetics of Betahistine are linear, and suggesting that the involved metabolic pathway is not saturated.

### **5.3. Preclinical safety data**

#### Chronic toxicity

Adverse effects in the nervous system were seen in dogs and baboons after intravenous doses at and above 120 mg/kg.

Chronic oral toxicity testing for 18 months in rats at a dose of 500 mg/kg and 6 months in dogs at a dose of 25 mg/kg showed betahistine to be well tolerated without changes in the clinical chemical and hematological parameters.

#### Mutagenic and carcinogenic potential

Betahistine does not have mutagenic potential.

In an 18 months chronic toxicity study in rats betahistine up to a dose of 500 mg/kg did not show any evidence for carcinogenic potential.

#### Reproduction toxicity

Effects in reproductive toxicity studies were observed only at exposures considered sufficiently in excess of the maximum human exposure. The relevance of changes noticed at higher dose to humans is unknown.

## **6. PHARMACEUTICAL PARTICULARS**

### **6.1. List of excipients**

Citric acid anhydrous  
Microcrystalline cellulose  
Mannitol (E421)  
Silica, colloidal anhydrous  
Purified talc

### **6.2. Incompatibilities**

Not applicable.

### **6.3. Shelf life**

36 months.

### **6.4. Storage condition**

Store below 30°C.

### **6.5. Nature and contents of container**

Alu/PVC /ACLAR blister packs containing 100 tablets.  
Alu/ PVC/ PVDC blister packs containing 100 tablets.  
Not all presentations are marketed.

### **6.6. Special precautions for disposal and other handling**

Any unused product or waste material should be disposed of in accordance with local requirements.

## **7. PRODUCT REGISTRANT**

Goldplus Universal Pte. Ltd  
103, Kallang Ave, #06-02,  
Singapore 339504

## **8. PRODUCT REGISTRATION NUMBER**

SINXXXXXP

## **9. DATE OF REVISION OF THE TEXT**

21/3/2023