▼This medicinal product is subject to additional monitoring in Australia. This will allow quick identification of new safety information. Healthcare professionals are asked to report any suspected

adverse events at www.tga.gov.au/reporting-problems.

# Australian Product information

# KOZENIS (Tafenoquine) Tablets

# Name of the medicine

Tafenoquine succinate

# Qualitative and quantitative composition

Each tablet contains 188.2 mg of tafenoquine succinate (equivalent to 150 mg of tafenoquine).

Each tablet also contains 162.8 mg of the excipient mannitol. For a full list of excipients, see Section 6.1 LIST OF EXCIPIENTS.

# Pharmaceutical form

Film-coated tablets for oral administration.

Pink, film-coated, capsule-shaped tablets, plain on one side and debossed with ‘GS J11’ on the other side.

# Clinical particulars

## Therapeutic indications

Tafenoquine is indicated for the radical cure (prevention of relapse) of *Plasmodium vivax* (*P. vivax*) malaria in patients aged 16 years and older who are receiving appropriate antimalarial therapy for the acute *P. vivax* infection (see Section 4.2 Dose and method of administration).

## Dose and method of administration

All patients must be tested for glucose-6-phosphate dehydrogenase (G6PD) deficiency prior to prescribing tafenoquine (see Section 4.3 Contraindications and Section 4.4 Special warnings and precautions for use).

Pregnancy should be excluded prior to use the use of tafenoquine in females of child bearing potential (see Section 4.3 Contraindications and Section 4.6 Fertility, pregnancy and lactation).

Do not break or crush the tablets. Tafenoquine should be taken with food to increase systemic absorption and minimise gastrointestinal side effects (see Section 5.2 Pharmacokinetic properties). In the event of vomiting within 60 minutes after dosing, a repeat dose should be given. Re-dosing should not be attempted more than once.

Tafenoquine should be co-administered with chloroquine on the first or second day of the three days chloroquine administration for the treatment of acute *P.vivax* malaria.

Tafenoquine is NOT indicated for the treatment of acute *P. vivax* malaria. Consideration should be given to official guidance on the appropriate use of antimalarial medicinal products in areas where chloroquine is not recommended.

There are no data regarding co-administration of tafenoquine for the radical cure of *P.vivax* in conjunction with antimalarial agents other than chloroquine in the treatment of acute *P. vivax* infection (see Section 5.1 PHARMACODYNAMIC PROPERTIES, Clinical trials).

There are no data regarding the subsequent re-treatment of recurrent *P. vivax* infection with tafenoquine following initial dosing.

### Populations

### Adults and Adolescents (16 years and older)

A single 300 mg dose (two 150-mg tafenoquine tablets) is recommended to be given on Day 1 or Day 2 of the 3 day course of chloroquine (see Section 5.1 Pharmacodynamic properties).

### Children and Adolescents (up to 16 years of age)

The safety and efficacy of tafenoquine have not been established in children and adolescents less than 16 years of age.

### Elderly (65 years or older)

There are limited data available on the use of tafenoquine in patients aged 65 years and older.  However, there is no evidence that elderly patients require a different dose than younger adult patients (see Section 5.2 Pharmacokinetic properties).

### Renal Impairment

Tafenoquine has not been studied in patients with renal impairment. Dose adjustments in patients with renal impairment are unlikely to be required as tafenoquine is administered as a single one-time dose.

### Hepatic Impairment

Tafenoquine has not been studied in patients with hepatic impairment. Dose adjustments in patients with hepatic impairment are unlikely to be required as tafenoquine is administered as a single one-time dose.

## Contraindications

Tafenoquine is contraindicated in the following:

* G6PD deficiency (see Section 4.4 Special warnings and precautions for use).
* Pregnancy (see Section 4.6 Fertility, pregnancy and lactation).
* Breastfeeding an infant who is G6PD deficient or if the G6PD status of the infant is unknown (see Section 4.6 Fertility, pregnancy and lactation).
* Patients with known hypersensitivity to tafenoquine, other 8-aminoquinolines, or any component of the formulation.

## Special warnings and precautions for use

### Haemolytic anaemia and G6PD deficiency

Due to the risk of haemolytic anaemia in patients with G6PD deficiency, G6PD testing must be performed before prescribing tafenoquine (see Section 4.3 Contraindications). Withhold tafenoquine from patients with G6PD enzyme levels <70% of normal (see section 5.2 Pharmacokinetic properties). Monitor patients for clinical signs or symptoms of haemolytic anaemia. Advise patients to seek medical attention if signs of haemolytic anaemia occur.

### Methaemoglobinaemia

Asymptomatic elevations in methaemoglobin were observed in clinical studies (see Section 4.8 Adverse effects (UNDESIRABLE EFFECTS)). If signs or symptoms of methaemoglobinaemia occur, appropriate therapy should be instituted. Caution is advised in patients with nicotinamide adenine dinucleotide (NADH)-dependent methaemoglobin reductase deficiency.

### Psychiatric Effects

Mild to moderate, self-limiting psychiatric adverse reactions (e.g. anxiety, abnormal dreams) have been reported in clinical trials of tafenoquine (see Section 4.8 Adverse effects (UNDESIRABLE EFFECTS)). While there were no reports of serious psychiatric adverse reactions in clinical trials following a single 300 mg dose, cases of depression and psychosis have occurred following higher single doses (350 to 600 mg) of tafenoquine, mostly in subjects with a previous history of psychiatric disorders. Serious psychiatric disorders such as psychosis and depression have been associated with some quinoline anti-malarials. Caution is advised when administering tafenoquine to patients with a current or past history of serious psychiatric disorders. Individual patient risk-benefit should be assessed. Due to the long half-life of tafenoquine (15 days), psychiatric effects and hypersensitivity reactions may be delayed in onset and/or duration.

### Use in the elderly

### Refer to section DOSE AND METHOD OF ADMINISTRATION, Elderly (65 years or older).

### Paediatric use

### Refer to section DOSE AND METHOD OF ADMINISTRATION, Children and Adolescents (up to 16 years of age).

### Effects on laboratory tests

No data available.

## Interactions with other medicines and other forms of interactions

Tafenoquine is an inhibitor of human transporters organic cation transporter 2 (OCT2) and multidrug and toxin extrusion transporter (MATE) *in vitro*, potentially resulting in increased exposure to their substrates (e.g., dofetilide) (see Section 5.2 Pharmacokinetic properties). There is a small risk of lactic acidosis due to increased metformin exposure secondary to blockade of these transporters. Therefore, use with caution with metformin. Drugs with a narrow therapeutic index that are substrates of the renal transporters OCT2 and MATE should not be co-administered (e.g. phenformin, buformin, dofetilide, procainamide, and pilsicainide).

## Fertility, pregnancy and lactation

### Effects on fertility

In a rat fertility study, tafenoquine was given orally at 1.5, 5, and 15 mg/kg/day (up to about 0.5 times the human dose based on body surface area comparisons) to males for at least 67 days, including 29 days prior to mating, and to females from 15 days prior to mating through early pregnancy. Tafenoquine resulted in reduced number of viable foetuses, implantation sites, and corpora lutea at 15 mg/kg in the presence of maternal toxicity (mortality, piloerection, rough coat, and reduced body weight).

### Use in pregnancy (Category C):

Tafenoquine is contraindicated in pregnancy. There is a risk of haemolysis in patients with G6PD deficiency; and, even if a pregnant woman is not G6PD deficient, the foetus may be deficient in G6PD.

The effect of tafenoquine on human pregnancy is unknown. Tafenoquine resulted in dose related abortions when given orally to pregnant rabbits during organogenesis (GD 6 to 18), at doses of 7 mg/kg (about 0.4 times the clinical exposure based on body surface area comparisons) and above. Doses higher than 7 mg/kg were also associated with maternal toxicity (mortality and reduced body weight gain). In a similar study in rats, doses of 3, 10, or 30 mg/kg/day resulted in maternal toxicity (enlarged spleen, reduced body weight and reduced food intake) but no fetotoxicity, at the high-dose (equivalent to the clinical exposure based on body surface area comparisons). There was no evidence of malformations in either species. It is unknown if tafenoquine crosses the placenta.

Women of child-bearing potential should have a pregnancy test prior to starting treatment with tafenoquine and avoid becoming pregnant for 3 months after taking tafenoquine.

### Use in lactation.

It is not known whether tafenoquine is excreted in human milk. In a pre- and postnatal development study in rats, tafenoquine administered throughout pregnancy and lactation produced maternal toxicity and a reversible decrease in offspring body weight gain and decrease in motor activity; at 18 mg/kg/day, which is equivalent to about 0.6 times the clinical dose based on body surface area comparisons.

Tafenoquine should not be used during breastfeeding when the infant has G6PD deficiency or the status is unknown as haemolytic anaemia may occur (see Section 4.3 Contraindications).

Tafenoquine should only be used in a nursing mother if the expected benefit justifies the risk to an infant that is not G6PD deficient. Consideration should be given to the long half-life for tafenoquine as the drug may be present in the systemic circulation for 3 months following treatment with tafenoquine (see Section 5.2 Pharmacokinetic properties).

## Effects on ability to drive and use machines

There have been no studies to investigate the effect of tafenoquine on driving performance or the ability to operate machinery. A detrimental effect on such activities would not be anticipated from the pharmacology of tafenoquine. The clinical status of the patient and the adverse event profile of tafenoquineshould be borne in mind when considering the patient's ability to perform tasks that require judgement, motor or cognitive skills.

## Adverse effects (Undesirable effects)

### Clinical trial data

The adverse drug reaction profile was evaluated in 3 randomised, double-blind studies including a total 483 patients administered 300 mg tafenoquine in a single oral dose co-administered with chloroquine phosphate (600-mg free base on Days 1 and 2 with 300-mg free base on Day 3). Two of these studies were placebo-controlled and the third was an active-controlled study. The safety profile was also informed by supportive clinical studies, some of which included healthy volunteers who received the indicated dose. In the overall clinical development program, a total of 810 subjects received a single dose of tafenoquine 300 mg (>4,000 subjects received tafenoquine including other doses or regimens).

Adverse reactions are listed below by MedDRA body system organ class and by frequency.

The frequency categories used are:

Very common ≥ 1 in 10

Common ≥ 1 in 100 and < 1 in 10

Uncommon ≥ 1 in 1,000 and < 1 in 100

Rare < 1 in 1,000

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| **System organ class** | **Very**  **common** | **Common** | **Uncommon** | **Rare** |
| *Blood and lymphatic system disorders* |  | Haemoglobin decreased  Elevated methaemoglobin |  |  |
| *Immune system disorders* |  |  |  | Hypersensitivity reactions (e.g.,angioedema) |
| *Psychiatric disorders* |  | Insomnia | Anxiety | Abnormal dreams |
| *Nervous system disorders* |  | Headache  Dizziness | Somnolence |  |
| *Eye disorders* |  |  | Photophobia  Vortex keratopathy |  |
| *Gastrointestinal disorders* |  | Nausea  Vomiting |  |  |
| *Hepatobiliary disorders* |  |  | Alanine aminotransferase increased |  |
| *Renal and urinary disorders* |  | Blood creatinine increased |  |  |

**Reporting suspected adverse effects**

Reporting suspected adverse reactions after registration of the medicinal product is important. It allows continued monitoring of the benefit-risk balance of the medicinal product. Healthcare professionals are asked to report any suspected adverse reactions at [https://www.tga.gov.au/reporting-problems](http://www.tga.gov.au/reporting-problems).

## Overdose

Haemolytic anaemia in patients with G6PD deficiency and methaemoglobinaemia may be encountered in an overdose.

### Treatment

There is no specific treatment for an overdose with tafenoquine. If overdose occurs, the patient should be treated supportively with appropriate monitoring as necessary.

Further management should be as clinically indicated or as recommended by the national poisons centre.

For information on the management of overdose, contact the Poisons Information Centre on 13 11 26 (Australia).

# Pharmacological properties

## Pharmacodynamic properties

### Mechanism of action

Tafenoquine, an 8-aminoquinoline, eradicates *P. vivax* liver hypnozoites, preventing the relapse of malaria. The molecular target of tafenoquine is not known.

### Clinical trials

The DETECTIVE trial was a double-blind, randomised, controlled clinical trial of 522 adults positive for *P. vivax* in 3 regions (Asia, Africa, and Latin America). All patients received chloroquine phosphate (600-mg free base on Days 1 and 2 and 300-mg free base on Day 3) to treat the acute *P.vivax*  malaria and were randomised to one of the following: a one-time dose of tafenoquine (two 150 mg tablets) on Day 1 or Day 2 (N=260), primaquine 15 mg once daily for 14 days starting Day 2 (N=129), or placebo (N=133). Patients included in the study had a mean age of 35 (range 15-79 years), were primarily male (75%), and from the following regions: 70% South America (Brazil and Peru), 20% Southeast Asia (Thailand, Cambodia and the Philippines), and 11% Africa (Ethiopia). All patients enrolled in the study had a positive blood film for *P. vivax*. Those with mixed malaria infections were excluded.

The primary endpoint was recurrence-free efficacy 6 months post-dosing for tafenoquine added to chloroquine compared to chloroquine alone. Patients were considered recurrence-free if they demonstrated initial parasite clearance, took no anti-malarial medications, and were confirmed parasite-free at the final assessment (i.e., absence of relapse or new infection).

Due to the risk of haemolytic anaemia, patients were excluded from the study if they had a G6PD enzyme level <70% of the site median value for G6PD normals. An assay validation study determined G6PD eligibility requirements for the pivotal trials and found global median G6PD activity was 8.2 IU/gHb, with 70% of median at 5.7 IU/gHb (at 30ºC using Trinity® assay). Regional G6PD values (70% of median) were similar across the studied regions: 5.8 for South America, 5.6 for SE Asia, 5.7 for Africa). In this trial, the minimum G6PD enzyme level of any subject was 5.4IU/gHb.

The recurrence-free efficacy rates at 6 months amongst treatment groups are presented for the overall population in Table 1. The risk of recurrence for tafenoquine plus chloroquine was reduced by 70% compared to chloroquine alone. There was no evidence of an effect of host cytochrome P450 2D6 metabolizer class on treatment outcome with tafenoquine.

**Table 1. Recurrence-free efficacy at 6 months – Overall Populationa**

|  |  |  |  |
| --- | --- | --- | --- |
|  | **Tafenoquine/**  **Chloroquine**  **(n = 260)** | **Primaquine/**  **Chloroquined**  **(n = 129)** | **Chloroquine**  **(n = 133)** |
| Recurrence-free efficacyb  (95% CI)  HRc - (95% CI) difference from chloroquine  p value | 62%  (55, 69)  0.30 (0.22, 0.40)  <0.001 | 70%  (60, 77)  0.26 (0.18, 0.39)  <0.001 | 28%  (20, 36) |

1. Microbiologic intent to treat population; survival analysis
2. Kaplan-Meier Estimate
3. Hazards ratio of the risk of recurrence versus chloroquine alone obtained from a Cox’s proportional hazards model with treatment and region as covariates.
4. Statistical comparisons for efficacy cannot be made between tafenoquine/chloroquine and primaquine/chloroquine as the study was not powered for this comparison.

Cardiac Electrophysiology

At a cumulative dose of 1200 mg (400 mg/day for 3 days; 4 times the maximum recommended dose), tafenoquine did not prolong the QT interval to any clinically relevant extent.

## Pharmacokinetic properties

### Absorption

Maximum plasma concentrations were generally observed 12 to 15 hours following oral administration. Plasma AUC increased 41% and Cmax increased 31% for tafenoquine administration with a high fat meal compared to the fasted state.

### Distribution

Tafenoquine is highly plasma protein bound (>99.5%) and widely distributed (apparent oral volume of distribution >1,500 L). Following single and multiple oral dose administration, tafenoquine whole blood concentrations were on average 67% higher than corresponding plasma values, reflecting preferential partitioning of drug in the erythrocytes.

### Metabolism

Tafenoquine undergoes very slow metabolism, and drug-related material is excreted slowly, both unchanged and as metabolites. Tafenoquine is the principal circulating drug-related component and there are no major systemic metabolites in humans.

### Excretion

The clearance of oral tafenoquine is approximately 3 L/h based on plasma concentrations. The average terminal half-life is approximately 15 days. Definitive elimination data in humans has not been generated, although slow elimination of drug related material in urine is evident. In nonclinical species drug-related material is eliminated slowly in both urine and faeces (which includes some biliary secretion).

### Special Patient Populations

### Elderly patients (> 65 years old)

No formal studies have been conducted in elderly patients. In a population pharmacokinetic analysis in 675 subjects aged 15 to 79 years, there was no indication of an effect of age on the pharmacokinetics of tafenoquine.

### Renal impairment

No formal studies have been conducted to investigate the effect of renal impairment on the pharmacokinetics of tafenoquine.

### Hepatic impairment

No formal studies have been conducted to investigate the effect of hepatic impairment on the pharmacokinetics of tafenoquine.

### Drug Interaction Studies

Tafenoquine demonstrated *in vitro* inhibition of several CYPs including 1A2, 2A6, 2C8, 2C9 and 3A4 enzymes. Clinical studies have shown no clinically significant effects on the pharmacokinetics of substrates of CYP1A2 (caffeine), CYP2D6 (desipramine), CYP2C9 (flurbiprofen), or CYP3A4 (midazolam, chloroquine) following oral administration of tafenoquine.

Tafenoquine inhibited *in vitro* transport of metformin via human OCT2, MATE1 and MATE2-K transporters. Assessments based on systemic concentrations (unbound Cmax) of tafenoquine at therapeutic doses, compared with the IC50 values derived from *in vitro* transporter inhibition studies, were conducted and indicated a potential, but small, drug interaction risk with OCT2 and MATE substrates.

Concomitant administration of tafenoquine and chloroquine in man resulted in no clinically significant interaction.

Tafenoquine administered concomitantly with dihydroartemisinin-piperaquine (40 mg /320 mg as the tetrahydrate daily for 3 days) increased exposure of tafenoquine AUC0-inf 12% and Cmax 38%. This change was not considered clinically relevant. There was no significant change in dihydroartemisinin or piperaquine exposure.

Concomitant administration of tafenoquine with artemether-lumefantrine (20 mg/120 mg daily for day 1, followed by twice daily for 2 days) reduced the exposure of the dihydroartemisinin metabolite of artemether by 23% and 16% for AUC(0-tau) and Cmax respectively. This change was not considered clinically significant. There was no significant change in tafenoquine, lumefantrine or artemether exposure.

## Preclinical safety data

### Genotoxicity

Tafenoquine did not cause gene mutations or chromosomal damage in two definitive *in vitro* tests (bacterial mutation assay and mouse lymphoma L5178Y cell assay), or in an *in vivo* oral rat micronucleus test.

### Carcinogenicity

Two-year oral carcinogenicity studies were conducted in rats and mice. Tafenoquine was not carcinogenic in mice but was carcinogenic in rats inducing an increase in the incidence of renal cell tumours and hyperplasia in high dose (2 mg/kg/day) and mid dose (1 mg/kg/day) males compared with controls (normalised AUC0-8 weeks equivalent to 5.0 and 2.4 times the human dose per AUC0-∞ based on a single 300 mg dose, respectively). Given the single dose administration of tafenoquine, these findings are not considered to represent a carcinogenicity risk to humans.

### Animal toxicology and/or pharmacology

Tafenoquine has been evaluated in repeat dose toxicity studies of up to 13 weeks in duration in CD-1 mice, 26 weeks in Sprague Dawley rats, 52 weeks in beagle dogs and in a PK study in rhesus monkeys. Principal findings were haematological (e.g., decreased haemoglobin, increased methaemoglobin), pulmonary (e.g., increased numbers of foamy macrophages and the presence of eosinophilic material in alveoli), hepatic (e.g., increased liver weight, subacute inflammation), and renal toxicity (e.g., renal tubular lesions). The majority of these effects was both dose- and duration-dependent, and reversible upon cessation of treatment. The risk of clinically relevant toxicity outside of the known risk of haematologic effects associated with 8-aminoquinolines is low considering the single dose administration of tafenoquine.

### Microbiology

Tafenoquine has demonstrated schizontocidal activity against *Plasmodium vivax* in animal models.

# Pharmaceutical particulars

## List of excipients

Microcrystalline Cellulose

Mannitol

Magnesium Stearate

Hypromellose

Titanium Dioxide

Iron Oxide Red

Polyethylene Glycol

## Incompatibilities

No incompatibilities have been identified.

## Shelf life

In Australia, information on the shelf life can be found on the public summary of the Australian Register of Therapeutic Goods (ARTG). The expiry date can be found on the packaging.

## Special precautions for storage

Store in the original package to protect from moisture.

## Nature and contents of container

150 mg tablets supplied in:

* Child-resistant aluminium foil blister strip.

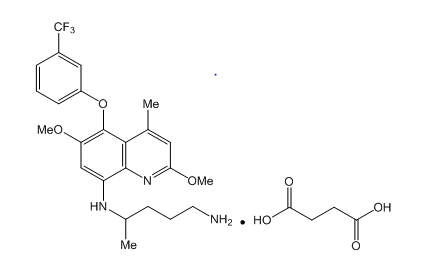
The pack contains two tablets.

## Special precautions for disposal

In Australia, any unused medicine or waste material should be disposed of by taking to your local pharmacy.

## Physicochemical properties

### Chemical structure



Molecular Formula: C24H28F3N3O3. C4H6O4

Molecular Weight: 581.58 as the succinate salt.

463.49 as free base.

Tafenoquine succinate is a pale green or pale orange to orange solid. The pKa values for tafenoquine succinate are 10.0 and 3.0, and it is sparingly soluble at pH 2 and practically insoluble at and above pH 6 in aqueous buffer. Tafenoquine succinate contains one chiral centre and is produced as a racemate.

### CAS number

CAS Registry Number: 106635-81-8

# Medicine schedule (Poisons Standard)

Schedule 4 – Prescription Only Medicine

# Sponsor

GlaxoSmithKline Australia Pty Ltd,

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Abbotsford, Victoria, 3067

# Date of first approval (ARTG entry)

13 September 2018

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