

DIASUNATE TABLETS

NAME OF THE MEDICINAL PRODUCT

Diasunate® 25mg/67.5mg tablets Artesunate, amodiaquine

Diasunate® 50mg/135mg tablets Artesunate, amodiaquine

Diasunate® 100mg/270mg tablets Artesunate, amodiaquin

QUALITATIVE AND QUANTITATIVE COMPOSITION

Diasunate is a fixed dose combination of amodiaquine and artesunate.

Each DIASUNATE 25mg/67.5mg tablet contains 25 milligrams of artesunate 67.5 milligrams of amodiaquine (as hydrochloride)

Each DIASUNATE 50mg/135mg tablet contains 50 milligrams of artesunate and 135 milligrams of amodiaquine (as hydrochloride).

Each DIASUNATE 100mg/270mg tablet contains 100 milligrams of artesunate and 270 milligrams of amodiaquine (as hydrochloride).

PHARMACEUTICAL FORM

Tablets :

DIASUNATE 25mg/67.5mg is a round bilayer tablet : one layer is yellow coloured, the other one is white to slightly yellow, with score line, engraved on one side "AS" and on the other side "25".

DIASUNATE 50mg/135mg is a round bilayer tablet : one layer is yellow coloured, the other one is white to slightly yellow, with score line, engraved on one side "AS" and on the other side "50".

DIASUNATE 100mg/270mg is a round bilayer tablet : one layer is yellow coloured, the other one is white to slightly yellow, with score line, engraved on one side "AS" and on the other side "100".

The tablets may be mottled but it does not alter the safety and effectiveness of the product.

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

CLINICAL PARTICULARS

Therapeutic Indication

DIASUNATE is indicated for the treatment of uncomplicated cases of malaria due to *Plasmodium falciparum* strains which are susceptible to amodiaquine as well as to artesunate.

The most recent official guidelines on the appropriate use of antimalarial agents and local information

on the prevalence of resistance to antimalarial drugs must be taken into consideration for deciding on

the appropriateness of therapy with DIASUNATE.

Official guidance will normally include WHO

(<http://www.who.int/malaria/docs/TreatmentGuidelines2006.pdf>) and public health authorities guidelines (see also sections "Special warnings and precautions for use" and "Pharmacodynamic

properties”).

DIASUNATE should not be used in regions where amodiaquine resistance is widespread. (See also Sections “special warnings and precautions for use” and pharmacokinetic properties”).

Posology and method of administration

Oral use

The dosage of artesunate and amodiaquine is:

- 4 mg/kg (range 2 to 10 mg/kg) body weight of artesunate and
- 10 mg/kg (range 7.5 to 15 mg/kg) body weight of amodiaquine base once daily for 3 days.

Weight range (approximate age range)

**1st day
of treatment**

**2nd day
of treatment**

**3rd day
of treatment**

≥ 4.5kg to < 9 kg
(2 to 11 months)*

25 mg AS
67.5 mg AQ

25 mg AS
67.5 mg AQ

25 mg AS
67.5 mg AQ

≥9kg to <18kg
(1 to 5 years)*

50 mg AS
135 mg AQ

50 mg AS
135 mg AQ

50 mg AS
135 mg AQ

≥18kg to <36kg
(6 to 13 years)*

100 mg AS
270 mg AQ

100 mg AS
270 mg AQ

100 mg AS
270 mg AQ

≥ 36kg
(14 years and
above)*

200 mg AS
540 mg AQ

200 mg AS
540 mg AQ

200 mg AS

540 mg AQ

* if a weight-age mismatch occurs, dosing should be weight-based.

AS: artesunate

AQ: amodiaquine

DIASUNATE should not be taken with a high-fat meal (see section “pharmacokinetic properties”).

The tablets should be swallowed with water.

For patients unable to swallow the tablets whole, e.g. very young children, the tablets can be dissolved

in water before administration. The tablets can also be crushed and administered with water.

Should vomiting occur within half an hour after dosing, a repeated dose of ARTESUNATE AMODIAQUINE WINTHROP is to be taken. In case of further vomiting, treatment for severe malaria should be considered.

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Renal/hepatic impairment:

No data are available on dosing in hepatically or renally impaired patients (see section “special warnings and precautions for use”).

Contraindications

☒ Hypersensitivity to the active substances or to any of the excipients,

☒ History of liver injury during treatment with amodiaquine,

☒ Previous haematological event during treatment with amodiaquine,

☒ Retinopathy (in case of frequent treatment).

DIASUNATE must not be used for malaria prophylaxis, since it may result in agranulocytosis and severe hepatotoxicity (see section “special warnings and precautions for use”).

Special warnings and precautions for use

DIASUNATE should not be used in regions where amodiaquine resistance is widespread, as the treatment with the combination under such conditions may mean effectively a treatment with artesunate alone with an insufficient duration and decreased plasma concentrations as compared to artesunate alone (see section “Interaction with other medicinal products”). As a result, the risk of development of resistance of *P.falciparum* to artesunate increases significantly.

Amodiaquine is effective against some chloroquine-resistant strains of *P.falciparum*, although there is cross-resistance.

DIASUNATE has not been evaluated for the treatment of complicated malaria and is therefore not recommended.

DIASUNATE has not been evaluated in the treatment of malaria due to *Plasmodium vivax*, *Plasmodium malariae* or *Plasmodium ovale* and is therefore not recommended.

DIASUNATE has not been evaluated for malaria prophylaxis. The use of amodiaquine for prophylaxis results in an unacceptably high risk of agranulocytosis and liver toxicity and is contraindicated. Therefore, the combination of amodiaquine and artesunate is also contraindicated for malaria prophylaxis (see section “Contraindications”).

It is not known, whether the toxicity of amodiaquine, observed with prophylactic use (i.e. agranulocytosis, hepatotoxicity), may also develop after repeated cycles of curative treatment.

DIASUNATE has not been studied specifically in patients with thalassaemia, sickle cell anaemia or G6PD deficiency.

In the absence of specific clinical studies, caution should be exercised in patients with renal or hepatic impairment.

Symptoms suggestive of the following diseases should be carefully monitored:

- ☑ Hepatitis, pre-icteric phase and especially when jaundice has developed,
- ☑ Agranulocytosis (as suggested, for instance, by a clinical condition including fever and/or tonsillitis and/or mouth ulcers).

When these symptoms develop or exacerbate during the course of therapy with ARTESUNATE AMODIAQUINE WINTHROP, laboratory tests for liver function and/or blood cell counts should be performed at once. Immediate discontinuation of treatment may be required.

In such cases, continuation of treatment with amodiaquine increases the risk of death.

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Cardiovascular effects have been reported with other amino-4-quinoline derivatives during high dose

treatment. There is no evidence that an overdose of amodiaquine causes any of the life-threatening cardiovascular complications often seen after an overdose of chloroquine. However, by chemical class

analogy, caution should be exercised, especially with patients who have recently taken other antimalarial drug with cardiovascular side effects (quinine, quinidine, halofantrine, lumefantrine, mefloquine) or those who are under treatment with cardiovascular drugs or other drugs with the potential to prolong the QT interval (see section “overdosage”).

The combination of artesunate and amodiaquine may induce neutropenia (see section “Undesirable effects”) and increase the risk of infection.

Caution is advised when combining DIASUNATE tablets with drugs inhibiting, inducing or competing for CYP2C8 (see section “Interaction with other medicinal products”).

Co-administration of DIASUNATE and efavirenz should be avoided, since this combination has been noted to cause marked hepatotoxicity.

Interactions with other medicinal products and other forms of interaction

Interactions with drugs used for treatment of HIV and/or tuberculosis may occur, though little clinical

data is available. Prescribers should be vigilant for adverse events potentially related to such interactions, including liver toxicity and neutropenia.

In the absence of clinical data, DIASUNATE is not recommended to be administered concomitantly with drugs known to inhibit the liver enzymes cytochrome (CYP) 2A6 (e.g. methoxsalen, pilocarpine, tranylcypromine) and/or CYP2C8 (e.g. trimethoprim, ketoconazole, ritonavir, saquinavir, lopinavir, gemfibrozil, montelukast,) (see section “Pharmacokinetic properties”).

No pharmacokinetic interactions of artesunate with other antimalarial drugs of importance have been

identified. However, concomitant administration of DIASUNATE with other antimalarial treatments is not recommended, as no data on efficacy and safety are available.

A statistically significant decrease in dihydroartemisinin (DHA), the main active metabolite of artesunate, occurs with concomitant use of artesunate and amodiaquine (C_{max} decreased 47%, AUC_{0-inf} decreased 17%).

Agranulocytosis and hepatitis have been reported following the use of amodiaquine in long term prophylaxis treatments (see section “Undesirable effects”). Therefore, caution should be observed when prescribing amodiaquine-containing products, such as ARTESUNATE AMODIAQUINE

WINTHROP, concurrently with other drugs with a potential for liver and/or haematological toxicity. Though no pharmacokinetic interactions have been documented, amodiaquine and desethylamodiaquine inhibit CYP 2D6 in vitro and may cause clinically significant interactions with some β -blockers, antidepressants, and antipsychotics drugs. Caution should be exercised when coadministration of these agents with DIASUNATE is deemed necessary.

Pregnancy and lactation

Pregnancy

Malaria is known to be particularly hazardous during pregnancy. The benefits and risks of therapy with DIASUNATE to mother and foetus must be assessed by the prescriber.

The safety of amodiaquine in pregnant women has not been conclusively established, although many

years of experience with the drug have not indicated any teratogenicity.

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Data on a limited number of exposed pregnant women do not indicate any adverse effect of artemisinin on

pregnancy or on the health of the foetus/newborn child. Animal data indicate a limited embryotoxic effect

at doses of 6 mg/kg/day or more (see section “preclinical safety data”).

During 1st trimester of pregnancy, DIASUNATE should not be used

unless clearly necessary e.g. if treatment is life-saving for the mother, and if another antimalarial is not

suitable or not tolerated.

During 2nd or 3rd trimesters of pregnancy, DIASUNATE may be used

with caution, only if other antimalarials are unsuitable.

Lactation

The amounts of antimalarials in breast milk are small. Therefore, lactating women can receive artemisinin-based combination therapies (including DIASUNATE) for malaria treatment.

Effects on the ability to drive and use machines

Patients receiving DIASUNATE should be warned that

somnolence, dizziness or asthenia may occur, in which case they should not drive or use machines.

Undesirable effects

The tolerability to the fixed combination DIASUNATE has been

evaluated through two studies involving 1003 patients treated with the fixed dose combination: one

conducted in Burkina-Faso, and another one conducted in Senegal, Cameroon, Mali, and Madagascar.

The tolerability was evaluated as comparable to reference treatments.

About 30% of treated patients experienced adverse reactions. Most of the reported adverse reactions

were similar to symptoms usually seen during a malaria attack.

The most frequent adverse reactions observed were:

anorexia, abdominal pain, nausea, asthenia, somnolence, insomnia and cough (see hereafter).

The most serious adverse reactions observed were:

asthenia, anaemia and vertigo.

The adverse events considered at least possibly related to the treatment are listed hereafter by body

system, organ class and absolute frequency.

The adverse reactions are ranked under body-system and frequency using the following convention:

very common: $\geq 1/10$; common: $\geq 1/100$ to $< 1/10$; uncommon: $\geq 1/1000$ to $< 1/100$; rare : $\geq 1/10,000$

to $< 1/1000$; very rare : $< 1/10,000$; not known : cannot be estimated from the available data.

The type and frequencies of all adverse reactions observed from the two pivotal studies are summarised hereafter:

Class-organ Frequency Adverse events

Infections and infestations Uncommon Bronchitis acute, gastroenteritis, oral

candidiasis

Blood and lymphatic system

disorders

Uncommon Anaemia

Metabolism and nutrition

disorders

Uncommon Hypoglycaemia

Psychiatric disorders Common Anorexia, insomnia

Uncommon Hallucination

Nervous system disorders Common Somnolence

Uncommon Paraesthesia

Eye disorders Uncommon Ocular icterus

Ear and labyrinth disorders Uncommon Vertigo

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Class-organ Frequency Adverse events

Cardiac disorders Uncommon Arrhythmia, bradycardia

Respiratory, thoracic, and
mediastinal disorders

Common Cough

Gastro-intestinal disorders Common Nausea, abdominal pain

Uncommon Diarrhoea, vomiting

Skin and subcutaneous tissue
disorders

Uncommon Pruritus, rash, face oedema, skin disorders

Musculoskeletal and
connective tissue disorders

Uncommon Arthralgia

General disorders and
administration site conditions

Common

Uncommon

Asthenia

Oedema peripheral, pyrexia

The following adverse reactions have been reported with amodiaquine, especially at higher doses and/or during prolonged treatment:

☑ Blood and lymphatic system disorders: cases of leucopenia and neutropenia (agranulocytosis)

☑ Nervous system disorders: rare neuromyopathy

☑ Eye disorders, varying in type and severity: transient accommodation disorders, corneal opacifications regressive once treatment is stopped, very rarely, irreversible retinopathy justifying specialist ophthalmic attention

- ☒ Hepato-biliary disorders: severe and sometimes fatal hepatitis
- ☒ Skin and subcutaneous disorders: slate-grey pigmentation, notably affecting the fingers and mucous membranes.

Post-marketing experience

In published literature data, generated mostly during post-approval use of amodiaquine and/or artesunate, additional types of events have been reported. Since frequency estimates are highly variable across the studies, no frequencies are given for these events. For some of these events, it is unclear whether they are related to amodiaquine/artesunate or occur as a result of the underlying disease process:

- headache, dizziness
- cold, flu, rhinitis, shivering, sore throat
- convulsion
- splenomegaly, jaundice
- allergic reaction.

If any of the side effects is serious or unexpected, you should inform the supplier (see section “supplier”) and/or health authority, as per local regulation.

Overdose

In cases of suspected overdose, the patient should be urgently transferred to a specialized unit where appropriate monitoring and symptomatic and supportive therapy should be applied.

Amodiaquine

☒ The dangerous dose of amodiaquine cannot be stated precisely because of the low number of known cases; by analogy with chloroquine, it can be estimated at around 2 grams as a single administration in adults,

☒ Symptoms: headache, dizziness, visual disorders, cardiovascular collapse and convulsions, followed by early respiratory and cardiac arrest.

Artesunate

No cases of overdose have been reported to date.

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PHARMACOLOGICAL PROPERTIES

Pharmacodynamic properties

Artesunate Amodiaquine Combination (ATC code: P01BF03)

DIASUNATE is an artemisinin-based combination therapy which consists of two blood schizonticides, with independent modes of action and different intraparasitic biochemical targets.

DIASUNATE is indicated in areas where parasite resistance rate to amodiaquine remains below the threshold defined by WHO.

Efficacy and safety of DIASUNATE in uncomplicated *P.*

falciparum malaria have been demonstrated in clinical trials in West and Central Africa and in Madagascar. Inconsistent results have been seen in some areas where combinations of artesunate and

amodiaquine have been studied, probably due to a higher prevalence of amodiaquine resistance.

Artesunate: Antimalarial (ATC code: P01BE03).

Artesunate is a hemisuccinate derivative of dihydroartemisinin, which is obtained by the reduction of

artemisinin, a sesquiterpene lactone endoperoxide extracted from a plant used in traditional Chinese

medicine, known as sweet or annual wormwood (*Artemisia annua*).

The chemical mechanism of action of artesunate has been widely studied and appears well established.

The artesunate endoperoxide bridge is split by haeme within the infected erythrocyte, generating singlet oxygen. Parasite proteins, particularly in membranous structures, are thus alkylated, leading to parasite death.

In-vitro experiments in *P. falciparum* have shown that artemisinin derivatives are active against a broad spectrum of the life cycle of the parasite, from the relatively inactive ring stage to late schizonts.

The schizonticidal and gametocytocidal activities of artesunate, administered orally have been demonstrated in vivo on chloroquine-sensitive strains of Plasmodium (*P. berghei* in mice and *P. knowlesi* in monkeys) and on chloroquine-resistant strains (*P. berghei* in mice).

In-vitro, artesunate appears to be inactive against extra-erythrocyte forms, sporozoites, liver schizontes or merozoites.

When administered orally, artesunate consistently acts more quickly than orally administered chloroquine and intravenous quinine in all animal models studied, regardless of the strain or dose tested. In macaques (the animal model most similar to humans) infected with a chloroquine-resistant

strain of *P. knowlesi*, cure was obtained with the same dose of artesunate and quinine.

Amodiaquine: Antimalarial (ATC code: P01BA06).

Amodiaquine is a synthetic 4-aminoquinoline antimalarial. Its activity is characterized by a schizonticidal action on *Plasmodium falciparum*, *Plasmodium vivax*, *Plasmodium ovale* and *Plasmodium malariae* by destroying intraerythrocytic forms.

The mechanism of action of 4-aminoquinoline derivatives against plasmodium is not yet completely known. It is nonetheless accepted that these derivatives, one of which is amodiaquine, penetrate the

infected red blood cells and prevent the parasite from polymerizing haeme into an insoluble product

called haemozoin, leading to parasite death.

Strains of *Plasmodium falciparum* resistant to 4-aminoquinolines (chloroquine, amodiaquine) are present in many areas, and their geographical distribution is constantly changing. However, amodiaquine remains active against some chloroquine-resistant *P. falciparum* strains.

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Pharmacokinetic properties

Artesunate

Absorption

After oral administration, absorption is rapid. Most of the artesunate is promptly biotransformed, mainly through plasma esterases, into the active metabolite dihydroartemisinin (DHA).

After administration of two DIASUNATE 100mg/270mg tablets

(i.e total dose of 540 mg amodiaquine and 200 mg artesunate) in healthy volunteers (n=32), the mean

(CV) artesunate C_{max} value was 162.9 ng/ml (75%), and the corresponding value for AUC was 89.9 ng.h/ml (51%). The median (range) artesunate t_{max} value was 0.25 hours (0.25-1.33 h).

The mean (CV) DHA C_{max} value was 460.4 ng/ml (3 %), and the corresponding value for AUC was 712.2 ng.h/ml (36%). The median (range) DHA t_{max} value was 0.75 hours (0.5-1.33 h).

Distribution

DHA has been shown to substantially accumulate in *P. falciparum*-infected erythrocytes. Artesunate is

not significantly protein-bound.

Metabolism

Artesunate is extensively hydrolysed by plasma esterases and perhaps also by CYP2A6. Its main

metabolite, DHA is presumed to account for most of the in vivo antimalarial activity. DHA is further metabolised through glucuronidation prior to excretion.

Elimination

Artesunate has a plasma half-life of 3-29 minutes. The active metabolite DHA has a plasma half-life of

40 to 95 minutes. The modes of excretion of DHA have not been fully elucidated.

Amodiaquine

Absorption

After oral administration in healthy subjects, amodiaquine is quickly absorbed and biotransformed into

its main active form, desethylamodiaquine. The absolute bioavailability of amodiaquine is not known.

After administration of two DIASUNATE 100mg/270mg tablets

(i.e total dose of 540 mg amodiaquine and 200 mg artesunate) in healthy volunteers (n=32), the mean

(CV) amodiaquine C_{max} value was 9.2 ng/ml (33%), and the corresponding value for AUC was 65.7 ng.h/ml (45%). The median (range) amodiaquine t_{max} value was 0.79 hours (0.48-8 h).

The mean (CV) desethylamodiaquine C_{max} value was 147.9 ng/ml (41%), and the corresponding value for AUC was 9947.8 ng.h/ml (43%). The median (range) desethylamodiaquine t_{max} value was 2

hours (1.33- 8 h).

Distribution

The volume of distribution of amodiaquine is estimated at 20 to 40 l/kg.

Desethylamodiaquine, the main metabolite of amodiaquine, is assumed to be the main active form after oral administration. It is mainly found in blood, at much higher concentrations than unchanged

amodiaquine. Its concentration in whole blood is 4-6 times higher than in plasma.

Metabolism

The hepatic first pass metabolism of amodiaquine is high, with formation of the active metabolite, desethylamodiaquine, presumably via the CYP2C8 isoenzyme. Further metabolism includes oxidation

and glucuronoconjugation.

Elimination

Amodiaquine is eliminated principally through biotransformation with only around 2% excreted unchanged in urine. Desethylamodiaquine is slowly eliminated with a terminal half-life of 9-18 days.

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Diasunate

Artesunate and amodiaquine interaction

Single dose data have shown that the co-administration of artesunate and amodiaquine leads to a 47%

decrease in the C_{max} of dihydroartemisinin, and a 17% decrease of its AUC_{0-inf}, relative to what is seen when artesunate is administered alone. If DIASUNATE is used in the presence of amodiaquine resistance, this might further compromise the antimalarial activity of

DIASUNATE (see also sections “Therapeutic indication”, “special warnings and precautions for use” and “pharmacodynamic properties”).

Special populations

For the combined use of artesunate and amodiaquine, no pharmacokinetic data are available for patients with impaired renal or hepatic function.

Food effect

When DIASUNATE was taken with a high fat meal in healthy volunteers, the C_{max} and AUC(0-t) of amodiaquine increased 23% and 58% respectively, compared to fasting. The C_{max} and AUC(0-t) of the active metabolite desethylamodiaquine (DeAQ) increased 18% and 12% respectively with a high-fat meal, compared to fasting.

Conversely, when DIASUNATE was taken with a high fat meal in healthy volunteers, the C_{max} and AUC(0-t) of artesunate decreased 66% and 13% respectively, compared to fasting. The C_{max} and AUC(0-t) of the active metabolite dihydroartemisinin (DHA) decreased 48% and 5% respectively with a high-fat meal, compared to fasting.

Preclinical safety data

General toxicity

Artesunate presents low acute toxicity. After repeated administration of 50 mg/kg/day in rats and 82.5 mg/kg/day in dogs, i.e. 5 and 8.25 times the proposed maximal therapeutic dose in man it is potentially toxic to the haematopoietic organs, the immune system and response, the liver and kidneys.

For *amodiaquine* histopathological changes (pigmentation) were seen in the heart at 30 mg/kg/day in

rats. The statistically significant effects seen *in vitro* on ion channels in the heart at 0.1 µM in the hERG current (expressed in Human Embryonic Kidney cells) as well as the increase in QRS complex and QT interval durations at concentrations higher than 0.1µM in the isolated rabbit Purkinje fibres appeared to be due to a non-specific multi-ion channel blockade. Pigmentations were also seen in liver, kidney and thyroid glands in rats as well as in kidneys, liver and lymph nodes in dogs (at doses

of 25mg/kg/day). Also an increase in haemosiderosis in the spleen and bone marrow as well as thymus

lymphoid depletion were observed.

The toxicity after acute and chronic administration of the combination artesunate/amodiaquine was

similar to that of artesunate and amodiaquine, when administered alone. In repeated dose toxicity studies, the incidence and the severity of lesions were generally related to the dose levels.

Amodiaquine given alone at 30 mg/kg/day induced effects very similar to those of the 12/30 mg/kg/day artesunate amodiaquine combination.

Genotoxicity:

Artesunate did not show any mutagenic or clastogenic potential in *in vitro* and *in vivo* tests (Ames, mouse micronucleus). Although amodiaquine, like chloroquine, has shown both mutagenic and clastogenic potential, studies with the artesunate amodiaquine combination in the Ames test and micronucleus in rat did not demonstrate any evidence of genotoxicity.

Carcinogenesis:

No studies of the carcinogenic potential of the combination of artesunate and amodiaquine or the individual agents have been conducted.

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Toxicity to Reproduction :

Reproductive toxicology studies, conducted in rats and rabbits, confirmed the known embryotoxic and

teratogenic potential of artesunate and the maternal toxicity associated with amodiaquine. The combination did not demonstrate any particular effects on fertility or associated parameters. In the peri-postnatal study, the offspring from the F1 generation did not show any effect on sexual development, and despite an early slowing of bodyweight increases with some effect on testicular and

epididymal weights, no sequelae were noted on reproductive capacity.

No new toxicity was induced through the administration of the two substances in combination.

Safety pharmacology studies:

Slight sedative effect, a decrease in body temperature, a slight natriuretic effect and a decrease in endogenous creatinine clearance were observed with *artesanate* after single intravenous doses of 200 mg/kg (mice), 450 mg (rats, rabbits and dogs) and after single oral doses of 180 mg/kg in male rats. In conscious telemetered dogs, atrio-ventricular blocks and depressant effects on smooth muscles

were reported from 10 mg/kg (single oral dose). Since these effects were observed only in female animals, at a low incidence and without relation to dose, the relationship to artesunate administration

remains to be confirmed. Neither neurotoxicity nor prolongation of QT(c) interval were shown. Amodiaquine is likely to induce cardiovascular adverse effects, particularly transient prolongation of

QT interval duration at 30 mg/kg administered orally. This dose level corresponds to approximately

2-fold the maximum recommended therapeutic dose. At the dose level of 100 mg/kg administered orally (about 6.7 fold the maximum recommended therapeutic dose), also slight respiratory depressant

and natriuretic effects were noted.

Oral administration of both agents, amodiaquine followed by artesunate, was safe for the CNS, the cardiovascular and respiratory systems at dose levels of artesunate/amodiaquine corresponding to approximately 1.67 / 1.81 fold the maximum therapeutic dose levels (15/5.5 mg/kg amodiaquine/artesunate). The observed natriuretic effect on the kidney was very slight and transient.

PHARMACEUTICAL PARTICULARS

List of excipients

Croscarmellose sodium, Povidone K30, Silicia colloidal anhydrous, Microcrystalline cellulose, Magnesium stearate, Calcium carbonate DC CS90 (calcium carbonate and maize starch)

Incompatibilities

Not applicable.

Shelf life

The shelf-life of the product as packaged for sale is 36 months.

Special precautions for storage

The product should be stored below 30°C in the original package.

Nature and contents of container

***DIASUNATE* 25mg/67.5mg tablets**

3 tablets packaged in an aluminium/aluminium blister pack.

Box containing 1 or 25 blisters per pack.

***DIASUNATE* 50mg/135mg tablets**

3 tablets packaged in an aluminium/aluminium blister pack.

Box containing 1 or 25 blisters per pack.

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***DIASUNATE* 100mg/270mg tablets**

3 tablets packaged in an aluminium/aluminium blister pack.

6 tablets packaged in an aluminium/aluminium blister pack

Box containing 1 or 25 blisters per pack.

Instructions for use and handling and disposal

Not applicable

DATE OF REVISION OF THE TEXT:

NA

Marketing authorisation holder

Emzor Pharmaceutical Industries Limited

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