

**Artemether and Lumefantrine Dispersible Tablets 20/120 mg**

**ANNEX IV: MODEL STABILITY REPORT FOR FINISHED PHARMACEUTICAL  
PRODUCTS**

**ANNEX IV – Summary of Product Characteristic**

- Enclosed

## **SUMMARY OF PRODUCT CHARACTERISTICS**

## 1. NAME OF THE MEDICINAL PRODUCT

Artemether and Lumefantrine Cipla Dispersible Tablets 20mg/120mg\*

## 2. QUALITATIVE AND QUANTITATIVE COMPOSITION

Each dispersible tablet contains artemether 20mg and lumefantrine 120mg  
For a full list of excipients see section 6.1.

## 3. PHARMACEUTICAL FORM

Dispersible tablet

Yellow coloured, circular shaped, flat bevelled, uncoated tablets, debossed with 'CL' on one side and plain on the other side.  
No score line.

## 4. CLINICAL PARTICULARS

### 4.1 Therapeutic indications

Artemether and Lumefantrine Cipla Dispersible Tablets 20mg/120mg is indicated for the treatment of uncomplicated cases of malaria due to *Plasmodium falciparum* in adults, children and infants weighing 5 kg and above.

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 Artemether and Lumefantrine Cipla Dispersible Tablets 20mg/120mg.

Official guidance will normally include WHO  
(<http://www.who.int/malaria/publications/atoz/9789241549127/en/>)  
and local health authorities' guidelines (see also sections 4.4 and 5.1).

### 4.2 Posology and method of administration

Oral use

Treatment should be administered at the time of initial diagnosis or at the onset of symptoms. It is preferable that the patient has a positive diagnostic test before administration.

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\*Trade names are not prequalified by WHO. This is the national medicines regulatory authority's (NMRA) responsibility. Throughout this WHOPAR the proprietary name is given as an example only.

**Number of Artemether and Lumefantrine Cipla Dispersible Tablets 20mg/120mg tablets for treatment according to weight bands**

<b>Weight range</b>	<b><sup>st</sup> of treatment</b>	<b><sup>nd</sup> of treatment</b>	<b><sup>rd</sup> of treatment</b>
≥ 5kg to < 15kg	1 tablet twice daily (2 x 20mg/120mg A/L)	1 tablet twice daily (2 x 20mg/120mg A/L)	1 tablet twice daily (2 x 20mg/120mg A/L)
15kg to <25kg	2 tablets twice daily (2 x 40mg/240mg A/L)	2 tablets twice daily (2 x 40mg/240mg A/L)	2 tablets twice daily (2 x 40mg/240mg A/L)
25kg to <35kg	3 tablets twice daily (2 x 60mg/360mg A/L)	3 tablets twice daily (2 x 60mg/360mg A/L)	3 tablets twice daily (2 x 60mg/360mg A/L))
≥ 35kg	4 tablets twice daily (2 x 80mg/480mg A/L)	4 tablets twice daily (2 x 80mg/480mg A/L)	4 tablets twice daily (2 x 80mg/480mg A/L)

To increase absorption, Artemether and Lumefantrine Cipla Dispersible Tablets 20mg/120mg should be taken with food or a milky drink (see section 5.2). If a patient is unable to tolerate food, Artemether and Lumefantrine Cipla Dispersible Tablets 20mg/120mg should still be administered, but the systemic exposure may be reduced.

Patients who vomit within 1 hour of taking the medication should repeat the dose.

If a dose is missed, it should be taken as soon as realized and then the recommended regimen continued until the full course of treatment has been completed.

Method of administration for dispersible tablets

- 1. Take two (2) teaspoons (10 ml) of water in a small and clean container and add the required number of tablets.**
- 2. Swirl the container until tablet(s) disperses, and give the entire mixture immediately.**
- 3. Rinse the container with an additional 10 ml of water and get the child to drink this water.**

*Renal or hepatic impairment*

No dose adjustments are necessary in patients with renal or hepatic impairment. However, caution is advised when administering Artemether and Lumefantrine Cipla Dispersible Tablets 20mg/120mg to patients with severe renal or hepatic problems (see section 4.4).

*Paediatric patients weighing less than 5 kg:*

Artemether/lumefantrine is not recommended for use in children below 5 kg body weight due to a lack of data on safety and efficacy.

*Elderly*

No special precautions or dosage adjustments are necessary in such patients.

**4.3 Contraindications**

Artemether and Lumefantrine Cipla Dispersible Tablets 20mg/120mg is contraindicated in:

- patients with known hypersensitivity to artemether, lumefantrine or to any of the excipients.
- patients with severe malaria according to WHO definition.
- patients with a personal or family history of congenital prolongation of the QTc interval or sudden death, or with any other clinical condition known to prolong the QTc interval, such as patients with a history of symptomatic cardiac arrhythmias, clinically relevant bradycardia or severe cardiac diseases.
- patients taking drugs that are known to prolong QTc interval such as:
  - antiarrhythmics of classes IA and III

- certain antibiotics including some agents of the following classes: macrolides, fluoroquinolones, imidazole, and triazole antifungal agents
  - certain non-sedating antihistamines (terfenadine, astemizole)
  - cisapride
- patients with known disturbances of electrolyte balance e.g. hypokalaemia or hypomagnesaemia
  - patients taking any drug which is metabolized by the cytochrome enzyme CYP2D6 (e.g. flecainide, metoprolol, imipramine, amitriptyline, clomipramine)
  - patients taking drugs that are strong inducers of CYP3A4 such as rifampicin, carbamazepine, phenytoin, St John's wort.

#### 4.4 Special warnings and precautions for use

*Renal/hepatic dysfunction:* Artemether/lumefantrine has not been studied in patients with severe renal or hepatic impairment.

*Malaria prophylaxis:* Artemether/lumefantrine has not been evaluated for malaria prophylaxis.

*Malaria not caused by P. falciparum:* Artemether/lumefantrine has not been evaluated for the treatment of malaria due to *P. vivax*, *P. malariae*, *P. ovale* or *P. knowlesi* (see section 5.1).

Following treatment of mixed infections including *P. vivax*, follow-up treatment must be given in order to eradicate the exoerythrocytic forms of *P. vivax*.

##### *Other antimalarials:*

Unless there is no other treatment option, Artemether and Lumefantrine Cipla Dispersible Tablets 20mg/120mg should not be given concurrently with any other antimalarial agent due to limited data on safety and efficacy (see section 4.5).

If a patient deteriorates while taking Artemether and Lumefantrine Cipla Dispersible Tablets 20mg/120mg alternative treatment for malaria should be started without delay. In such cases, monitoring of the ECG is recommended and steps should be taken to correct any electrolyte disturbances.

Due to the potential of additive/synergistic QT-prolongation, close ECG-monitoring is advised when quinine is given after Artemether and Lumefantrine Cipla Dispersible Tablets 20mg/120mg (see section 5.1).

If Artemether and Lumefantrine Cipla Dispersible Tablets 20mg/120mg is given after mefloquine, close monitoring of food intake is advised (see section 4.5).

In patients previously treated with halofantrine, Artemether and Lumefantrine Cipla Dispersible Tablets 20mg/120mg should not be administered earlier than one month after the last halofantrine dose (see section 4.5).

*Hormonal contraceptives:* Artemether and Lumefantrine Cipla Dispersible Tablets 20mg/120mg may reduce the effectiveness of hormonal contraceptives. Patients should be advised to use an additional non-hormonal (i.e. barrier) method of birth control for one month after therapy with artemether/lumefantrine.

*Intake with food and drinks:* Patients who remain averse to food during treatment should be closely monitored, as the risk of recrudescence may be greater.

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

Artemether and Lumefantrine Cipla Dispersible Tablets 20mg/120mg should not be used in patients taking drugs that are known to prolong the QTc interval (see section 4.3), as effects may be additive and increase the risk of cardiac arrhythmia.

##### *Interaction with other antimalarials*

Artemether and Lumefantrine Cipla Dispersible Tablets 20mg/120mg should not be given concurrently

with any other antimalarial agent (see section 4.4). In addition, due to the propensity of some antimalarial agents to prolong the QTc interval, caution is advised when administering Artemether and Lumefantrine Cipla Dispersible Tablets 20mg/120mg to patients in whom there may still be detectable concentrations of these

Administration of a six-dose regimen of artemether/lumefantrine (over 60 hours) starting 12 hours after completion of a three-dose regimen of mefloquine or placebo in healthy volunteers showed no effect of mefloquine on plasma concentrations of artemether or the artemether/dihydroartemisinin ratio, but a 30-40% reduction in plasma levels of lumefantrine. These are possibly due to lower absorption secondary to a mefloquine-induced decrease in bile production. Patients that have been pretreated with mefloquine should be encouraged to eat at dosing times to compensate for the decrease in bioavailability. Plasma mefloquine concentrations from the time of addition of artemether/lumefantrine were not affected compared with a group that received mefloquine followed by placebo.

In patients previously treated with halofantrine, Artemether and Lumefantrine Cipla Dispersible Tablets 20mg/120mg should be dosed at least one month after the last halofantrine dose due to the long elimination half-life of halofantrine and the potential additive/synergistic effects on the QT-interval.

#### Interaction with CYP450 enzymes

Studies in humans have demonstrated that artemisinins have some capacity to induce CYP3A4 and CYP2C19 and inhibit CYP2D6 and CYP1A2. Although the magnitude of the changes was generally low it is possible that these effects could alter the therapeutic response or safety profile of drugs that are predominantly metabolised by these enzymes (see sections 4.3 and 5.2).

Lumefantrine was found to inhibit CYP2D6 in vitro. This may be of particular clinical relevance for compounds with a low therapeutic index (see section 4.3).

#### Interaction with CYP450 3A4 inhibitors

*Ketoconazole:* both artemether and lumefantrine are metabolised predominantly by the cytochrome enzyme CYP3A4, and do not inhibit this enzyme at therapeutic concentrations. The concurrent oral administration of ketoconazole with artemether/lumefantrine led to a modest increase (2 fold) in artemether, DHA, and lumefantrine exposure in healthy adult subjects. This increase in exposure to the antimalarial combination was not associated with increased side effects or changes in electrocardiographic parameters. Dose adjustment of Artemether and Lumefantrine Cipla Dispersible Tablets 20mg/120mg is not considered necessary when administered concomitantly with ketoconazole or other azole antifungals, but such combinations should be used with caution.

#### *HIV Treatment Medications*

##### HIV nucleoside and nucleotide reverse transcriptase inhibitors (NTRIs, e.g. abacavir, emtricitabine, lamivudine, tenofovir [TDF or TAF], zidovudine.)

Co-administration has not been studied but based on metabolism and clearance a clinically significant interaction is unlikely.

##### HIV non-nucleoside reverse transcriptase inhibitors (NNRTIs):

*Efavirenz:* Co-administration of efavirenz and artemether/lumefantrine lead to decreases in artemether exposure (51% and 79%), dihydroartemisinin exposure (46% and 75%) and lumefantrine exposure by (21% and 56%). Lumefantrine had no significant effect on efavirenz exposure in either study. Use with caution as decreased concentrations of artemether, dihydroartemisinin, or lumefantrine may result in a decrease of antimalarial efficacy.

*Nevirapine:* Lumefantrine is metabolised predominantly by CYP3A4. Upon co-administration with artemether/lumefantrine with nevirapine decreased the AUCs of artemether and dihydroartemisinin. In a cross-over study lumefantrine exposure was decreased by 20% and lumefantrine reduced nevirapine exposure by 46%. Use with caution.

*Rilpivirine:* Co-administration has not been studied but based on metabolism and clearance a pharmacokinetic interaction is unlikely. Rilpivirine should be used with caution when co-administered with a drug that has a potential risk to prolong the QT interval.

##### HIV Protease Inhibitors (PIs)

*Atazanavir*: Co-administration may increase plasma levels of artemisinins and lumefantrine. Both lumefantrine and atazanavir have been shown to prolong the QT interval.

*Darunavir*: Co-administration may increase plasma levels of artemisinins and lumefantrine.

*Lopinavir/ritonavir*: Data from clinical studies and population modelling suggest that co-administration of lopinavir/ritonavir and artemether decreases exposure of dihydroartemisinin (the biologically active metabolite) by ~40-60%. Lumefantrine AUC was significantly increased by 2.3-fold and there was trend towards increased C<sub>max</sub> (1.4-fold). The clinical meaning of these opposite effects on artemether and lumefantrine is not clear. Both lumefantrine and lopinavir have been shown to prolong the QT interval.

*Ritonavir*: Co-administration may increase plasma levels of artemisinins and lumefantrine, as both are metabolised by CYP3A4. Caution is recommended.

#### HIV Integrase Strand-Transfer Inhibitors (INSTIs)

*Dolutegravir, Raltegravir*: Co-administration has not been studied but based on metabolism/elimination and toxicity profiles there is little potential for interaction.

*Elvitegravir/cobicistat*: Co-administration has not been studied. Artemether and lumefantrine are metabolized by CYP3A4. Elvitegravir/cobicistat may increase concentrations of artemisinins and lumefantrine.

#### Pharmacokinetic Enhancer

*Cobicistat*: Co-administration has not been studied. Cobicistat may increase concentrations of artemisinins and lumefantrine by inhibition of CYP3A4.

#### *Antivirals against Hepatitis B or C*

Co-administration has not been studied. In many instances a clinically significant interaction appears unlikely. However, consult the summary of product characteristics of the desired medication.

## **4.6 Fertility, pregnancy and breast-feeding**

### *Pregnancy*

A moderate amount of data on pregnant women in their first trimester (more than 500 pregnancy outcomes) is available for artemether/lumefantrine. Data from a recent meta-analysis have shown that compared to quinine, artemether/lumefantrine treatment in the first trimester was not associated with an increased risk of miscarriage or stillbirth. While the data are limited, they indicate no difference in the prevalence of major congenital anomalies between treatment groups (for animal data see section 5.3).

A large amount of data on pregnant women in their second and third trimester (more than 4000 documented pregnancy outcomes) is available for artemisinin derivatives including artemether/lumefantrine. They indicate no fetal or neonatal toxicity.

Artemether and Lumefantrine Cipla Dispersible Tablets 20mg/120mg can be used during pregnancy.

### *Breast-feeding*

The amounts of artemether, dihydroartemisinin and lumefantrine in breast milk are small. Therefore, breast-feeding women can receive artemisinin-based combination therapies (including Artemether and Lumefantrine Cipla Dispersible Tablets 20mg/120mg) for malaria treatment.

### *Fertility*

There is no information on the effects of Artemether and Lumefantrine Cipla Dispersible Tablets 20mg/120mg on fertility in humans.

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

No studies on the effects on the ability to drive and use machines have been performed. Patients receiving Artemether and Lumefantrine Cipla Dispersible Tablets 20mg/120mg should be warned that dizziness, fatigue or asthenia may occur, in which case their ability to drive or operate machines may be impaired.

**4.8 Undesirable effects**

The safety of artemether/lumefantrine has been evaluated in adults, adolescents and children in clinical trials with more than 3500 patients.

Adverse reactions reported from clinical studies and post-marketing experience are listed below according to system organ class.

Adverse reactions are ranked under headings of frequency using the MedDRA frequency convention:

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

( $< 1/10,000$ )

Not known (cannot be estimated from available data).

**Table 1: Frequency of undesirable effects**

	Adults and adolescents above 12 years of age	Infants and children of 12 years of age and below (incidence estimates*)
<b>Cardiac disorders</b>		
Palpitations	Very common	Uncommon
Electrocardiogram QT prolonged	Uncommon	rare
<b>Nervous system disorders</b>		
Headache	Very common	Common
Dizziness	Very common	Common
Gait disturbance	uncommon	--
Ataxia, hypoaesthesia	Uncommon	--
Clonic movements	Common	Uncommon
Somnolence	uncommon	uncommon
<b>Respiratory, thoracic and mediastinal disorders</b>		
Cough	Common	Very common
<b>Gastrointestinal disorders</b>		
Vomiting	Very common	Very common
Abdominal pain	Very common	common
Nausea	Very common	Common
Decreased appetite	Very common	Very common
Diarrhoea	Common	Common
<b>Skin and subcutaneous tissue disorders</b>		
Rash	Common	Common
Pruritus	Common	Uncommon
Urticaria	Uncommon	Uncommon
Arthralgia	Very common	Common
Myalgia	Very common	Common
<b>General disorders and administration site conditions</b>		
Asthenia	Very common	Common
Fatigue	Very common	Common
<b>Immune system disorders</b>		
Hypersensitivity	Not known	Rare
<b>Blood and lymphatic system disorders</b>		
Delayed haemolytic anaemia*	Not known	Not known
<b>Hepatobiliary disorders</b>		
Liver function tests abnormal	Uncommon	Common
<b>Psychiatric disorders</b>		
Sleep disorders	Very common	uncommon

\* These adverse reactions were reported during post-marketing experience. Because these spontaneously reported events are from a population of uncertain size, it is difficult to estimate their frequency.

Reporting suspected adverse reactions after authorisation 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 to the marketing authorisation holder, or, if available, via the national reporting system.

#### 4.9 Overdose

Experience of overdosage with artemether and lumefantrine is limited. In cases of suspected overdosage symptomatic and supportive therapy should be given as appropriate, which should include monitoring of ECG and serum electrolytes.

### 5. PHARMACOLOGICAL PROPERTIES

#### 5.1 Pharmacodynamic properties

Pharmacotherapeutic group: Antimalarials, Artemisinin and derivatives, combinations, ATC code: P01BF01

##### Pharmacodynamic effects

Artemether and Lumefantrine Cipla Dispersible Tablets 20mg/120mg comprises a fixed ratio of 1:6 parts of artemether and lumefantrine, respectively. The site of antiparasitic action of both components is the food vacuole of the malarial parasite, where they are thought to interfere with the conversion of haem, a toxic intermediate produced during haemoglobin breakdown, to the nontoxic haemozoin, malaria pigment. Lumefantrine is thought to interfere with the polymerisation process, while artemether generates reactive metabolites as a result of the interaction between its peroxide bridge and haem iron. Both artemether and lumefantrine have a secondary action involving inhibition of nucleic acid- and protein synthesis within the malarial parasite.

##### Clinical efficacy

The efficacy of artemether/lumefantrine was evaluated for the treatment of acute, uncomplicated malaria (defined as symptomatic *P. falciparum* malaria without signs and symptoms of severe malaria or evidence of vital organ dysfunction) in five 6-dose regimen studies and one study comparing the 6-dose regimen with the 4-dose regimen. Baseline parasite density ranged from 500/μL - 200,000/μL (0.01% to 4% parasitaemia) in the majority of patients.

Studies were conducted in otherwise healthy, partially immune or non-immune adults and children ( $\geq 5$ kg body weight) with uncomplicated malaria in Thailand, sub-Saharan Africa, Europe, and South America.

Efficacy endpoints consisted of:

- 28-day cure rate, proportion of patients with clearance of asexual parasites within 7 days without recrudescence by day 28
- parasite clearance time (PCT), defined as time from first dose until first total and continued disappearance of asexual parasite which continues for a further 48 hours
- fever clearance time (FCT), defined as time from first dose until the first time body temperature fell below 37.5°C and remained below 37.5°C for at least a further 48 hours (only for patients with temperature  $>37.5^{\circ}\text{C}$  at baseline)

The modified intent to treat (mITT) population includes all patients with malaria diagnosis confirmation who received at least one dose of study drug. Evaluable patients generally are all patients who had a day 7 and a day 28 parasitological assessment or experienced treatment failure by day 28. The results are presented in the table below:

**Clinical efficacy results**

Study No.	Age	Polymerase chain reaction (PCR)-corrected 28-day cure rate <sup>1</sup> n/N (%) in evaluable patients	Median FCT <sup>2</sup> [25 <sup>th</sup> , 75 <sup>th</sup> percentile]	Median PCT <sup>2</sup> [25 <sup>th</sup> , 75 <sup>th</sup> percentile]	Year/ Study location
A025 <sup>4</sup>	3-62 years	93/96 (96.9)	n <sup>3</sup> =59 35 hours [20, 46]	n=118 44 hours [22, 47]	1996-97 Thailand
A026	2-63 years	130/133 (97.7)	n <sup>3</sup> =87 22 hours [19, 44]	NA	1997-98 Thailand
A028	12-71 years	148/154 (96.1)	n <sup>3</sup> =76 29 hours [8, 51]	n=164 29 hours [18, 40]	1998-99 Thailand
A2401	16-66 years	119/124 (96.0)	n <sup>3</sup> =100 37 hours [18, 44]	n=162 42 hours [34, 63]	2001-05 Europe, Columbia
A2403	2 months-9 years	289/299 (96.7)	n <sup>3</sup> =309 8 hours [8, 24]	n=310 24 hours [24, 36]	2002-03 3 countries in Africa
B2303 <sup>CT</sup>	3 months-12 years	403/419 (96.2)	n <sup>3</sup> =323 8 hours [8, 23]	n=452 35 hours [24, 36]	2006-07 5 countries in Africa
B2303 <sup>DT</sup>	3 months-12 years	394/416 (94.7)	n <sup>3</sup> =311 8 hours [8, 24]	n=446 34 hours [24, 36]	2006-07 5 countries in Africa

<sup>1</sup> Efficacy cure rate based on blood smear microscopy

<sup>2</sup> mITT population

<sup>3</sup> For patients who had a body temperature >37.5°C at baseline only

<sup>4</sup> Only the 6-dose regimen over 60 hours group data is presented

<sup>CT</sup> –Artemether/lumefantrine tablets administered as crushed tablets

<sup>DT</sup> –Artemether/lumefantrine Dispersible tablets

Artemether/lumefantrine is not indicated for, and has not been evaluated in, the treatment of malaria due to *P. vivax*, *P. malariae* or *P. ovale*, although some patients in clinical studies had co-infection with *P. falciparum* and *P. vivax* at baseline. Artemether/lumefantrine is active against blood stages of *Plasmodium vivax*, but is not active against hypnozoites.

**Resistance**

Strains of *P. falciparum* with a moderate decrease in susceptibility to artemether or lumefantrine alone can be selected in vitro or in vivo, but not maintained in the case of artemether.

Alterations in some genetic regions of *P. falciparum* [multidrug resistant 1 (pfmdr1), chloroquine resistance transporter (pfcrt), and kelch 13 (K13)] based on in vitro testing and/or identification of isolates in endemic areas where artemether/lumefantrine treatment was administered, have been reported. The clinical relevance of these findings is not known.

**QT/QTc Prolongation:**

For information on the risk of QT/QTc prolongation in patients see Contraindications, section 4.3.

In a healthy adult volunteer parallel group study including a placebo and moxifloxacin control group (n = 42 per group), the administration of the six dose regimen of artemether/lumefantrine with food was associated

with a moderate prolongation of QTcF (QT interval corrected by Fridericias formula). The mean changes from baseline at 68, 72, 96, and 108 hours post first dose were 7.45, 7.29, 6.12 and 6.84 msec, respectively. At 156 and 168 hours after first dose, the changes from baseline for QTcF had no difference from zero. No subject had a > 30 msec increase from baseline nor was an absolute increase to > 500 msec. Moxifloxacin control associated with a QTcF increase as compared to placebo for 12 hours after the single dose with a maximal change at 1 hour after dose of 14.1 msec.

## 5.2 Pharmacokinetic properties

### Artemether

#### *Absorption*

Artemether is absorbed fairly rapidly and dihydroartemisinin (DHA), the active metabolite of artemether, appears rapidly in the systemic circulation with peak plasma concentrations of both compounds reached about 2 hours after dosing. The absolute bioavailability is unknown.

Following single dose administration of 4 dispersible tablet(s) of Artemether and Lumefantrine Cipla Dispersible Tablets 20mg/120mg in healthy volunteers, the mean ( $\pm$  SD) artemether  $C_{max}$  value was 98 ( $\pm$  73) ng/ml, the corresponding value for AUC was 254 ( $\pm$  181) ng.h/ml, and the mean artemether  $t_{max}$  value was 2.68 ( $\pm$  0.87) hours.

The pharmacokinetic data for dihydroartemisinin were supportive and indicated a comparable bioavailability between Test and Reference.

In healthy volunteers the relative bioavailability of artemether was increased more than two-fold when taken with food.

#### *Distribution*

Artemether is 95.4% bound to human serum proteins in vitro. The active metabolite dihydroartemisinin (DHA) is also bound to human serum proteins (47-76%).

#### *Metabolism*

Artemether is rapidly and extensively metabolised with substantial first-pass metabolism.

Artemether is metabolised in the liver to the biologically active main metabolite DHA (demethylation), predominantly through the isoenzyme CYP3A4/5. The pharmacokinetics of artemether in adults is time-dependent. During repeated administration of artemether/lumefantrine, plasma artemether levels decreased significantly, while levels of the active metabolite (dihydroartemisinin) increased, although not to a statistically significant degree. The ratio of day 3/day 1 AUC for artemether was between 0.19 and 0.44, and was between 1.06 and 2.50 for dihydroartemisinin. This suggests that there was induction of the enzyme responsible for the metabolism of artemether. DHA is further converted to inactive metabolites, primarily by glucuronidation. In vivo data indicate that artemisinins have some capacity to induce cytochrome isoenzymes CYP2C19 and CYP3A4.

#### *Elimination*

Artemether and dihydroartemisinin are rapidly cleared from plasma with an elimination half-life of about 2 hours.

No urinary excretion data are available for humans. In rats and dogs unchanged artemether has not been detected in faeces and urine due to its rapid and high-first-pass metabolism, but several metabolites (unidentified) have been detected in both faeces and urine.

### Lumefantrine

#### *Absorption*

Absorption of lumefantrine, a highly lipophilic compound, starts after a lag-time of up to 2 hours, with peak plasma concentration about 6-8 hours after dosing. The absolute bioavailability is unknown.

Following single dose administration of 4 tablet(s) of Artemether and Lumefantrine Cipla Dispersible Tablets 20mg/120mg in healthy volunteers, the mean ( $\pm$  SD) lumefantrine  $C_{max}$  value was 3225 ( $\pm$  1302)  $\mu$ g/ml, the corresponding value for AUC 53859 ( $\pm$  28757)

In healthy volunteers the relative bioavailability of lumefantrine, when was taken after a high-fat meal, was increased sixteen-fold compared with fasted conditions. Food has also been shown to increase the absorption of lumefantrine in patients with malaria, although to a lesser extent (approximately two-fold), most probably due to the lower fat content of the food ingested by acutely ill patients. The food interaction data indicate that absorption of lumefantrine under fasted conditions is very poor. Patients should therefore be encouraged to take the medication with a normal diet as soon as food can be tolerated.

#### *Distribution*

Lumefantrine is 99.7% bound to human serum proteins in vitro.

#### *Metabolism*

Lumefantrine is N-debutylated, mainly by CYP3A4, in human liver microsomes. The systemic exposure to the metabolite desbutyl-lumefantrine, for which the in vitro antiparasitic effect is 5 to 8 fold higher than lumefantrine, was less than 1% of the exposure to the parent drug. Desbutyl-lumefantrine data is not available specifically for an African population. In vitro, lumefantrine significantly inhibits the activity of CYP2D6 at therapeutic plasma concentrations. In humans, the exposure to lumefantrine increases with repeated administration of artemether/lumefantrine over the 3-day treatment period, consistent with the slow elimination of the compound.

#### *Elimination*

Lumefantrine is eliminated very slowly with a terminal half-life of approximately 3 days. No urinary excretion data are available for humans. Lumefantrine is eliminated via the bile in rats and dogs, with excretion primarily in the faeces. After oral dosing in rats and dogs qualitative and quantitative recovery of metabolites in bile and faeces was relatively low, most of the dose being recovered as parent drug.

#### Pharmacokinetics in special patient populations

Specific pharmacokinetic studies have not been performed in patients with hepatic or renal insufficiency. No pharmacokinetic studies are available in elderly patients.

#### *Paediatric population*

In paediatric malaria patients, mean  $C_{max}$  (CV%) of artemether (observed after first dose) were 223 (139%), 198 (90%) and 174 ng/ml (83%) for body weight groups 5-<15, 15-<25 and 25-<35 kg, respectively, compared to 186 ng/ml (67%) in adult malaria patients. The associated mean  $C_{max}$  of DHA were 54.7 (108%), 79.8 (101%) and 65.3 ng/ml (36%), respectively compared to 101 ng/ml (57%) in adult malaria patients.

AUC of lumefantrine (population mean, covering the six doses of artemether/lumefantrine) were 577, 699 and 1150 µg.h/ml for paediatric malaria patients in body weight groups 5-<15, 15-<25 and 25-<35 kg, respectively, compared to a mean AUC of 758 µg.h/ml (87%) in adult malaria patients. The elimination half-lives of artemether and lumefantrine in children are unknown.

### **5.3 Preclinical safety data**

#### General toxicity

The main changes observed in repeat-dose toxicity studies were associated with the expected pharmacological action on erythrocytes, accompanied by responsive secondary haematopoiesis.

#### Mutagenicity

No evidence of mutagenicity was detected in in vitro or in vivo tests with an artemether:lumefantrine combination (consisting of 1 part artemether: 6 parts lumefantrine). In the micronucleus test myelotoxicity was seen at all dose levels (500, 1,000 and 2,000 mg/kg), but recovery was almost complete 48 hours after dosing.

#### Carcinogenicity

Carcinogenicity studies with the artemether/lumefantrine combination were not conducted.

### Reproductive toxicity studies

Reproductive toxicity studies performed with the artemether/lumefantrine combination caused maternal toxicity and increased post-implantation loss in rats and rabbits at doses 50 mg/kg/day (corresponding to approximately 7 mg/kg/day artemether) and 175 mg/kg/day (corresponding to 25 mg/kg/day artemether) respectively. These effects were not observed at lower doses.

Lumefantrine alone caused no sign of reproductive or development toxicity at doses up to 1,000 mg/kg/day in rats and rabbits.

Embryotoxicity has been observed in rat and rabbit reproductive toxicity studies conducted with artemether, a derivative of artemisinin. Artemisinins (e.g. artesunate) are known to be embryotoxic in animals.

Artemether caused increases in post-implantation loss and teratogenicity (characterised as a low incidence of cardiovascular and skeletal malformations) in rats at 19.4 mg/kg, and in rabbits at 30 mg/kg. Maternal toxicity was also observed in rabbits at 30 mg/kg/day. No other adverse effects were observed at lower doses in rabbits. The no observed effect dose was 3 mg/kg/day in rats and 25 mg/kg/day in rabbits.

The embryotoxic artemether dose, 20 mg/kg/day in the rat, yields artemether and dihydroartemisinin exposures similar to those achieved in humans.

Artesunate, a structurally related compound, also caused increases in post-implantation loss and teratogenicity (low incidence of cardiovascular and skeletal malformations) in rats at 6 mg/kg and in the lowest dose tested in the rabbits, 5 mg/kg/day (see section 4.6 for data in humans).

### Cardiovascular Pharmacology

In toxicity studies in dogs at doses  $\geq 600$  mg/kg/day only, there was some evidence of prolongation of the QTc interval, at higher doses than intended for use in man. In an in vitro assay of HERG channels, lumefantrine and the main metabolite desbutyl-lumefantrine showed some inhibitory potential in one of the currents responsible for cardiac repolarization. From the estimated IC<sub>50</sub> values, the order of potency of HERG current block was halofantrine (IC<sub>50</sub> = 0.04  $\mu$ M) > chloroquine (2.5  $\mu$ M)

>mefloquine (2.6  $\mu$ M) >desbutyl-lumefantrine (5.5  $\mu$ M) >lumefantrine (8.1  $\mu$ M). Clinical studies show, that prolongation of QTcF can occur with standard dosing of artemether/lumefantrine (see sections 4.4 and 5.1).

## **6. PHARMACEUTICAL PARTICULARS**

### **6.1 List of excipients**

Microcrystalline cellulose  
Croscarmellose sodium  
Crospovidone  
Hydroxypropyl methylcellulose  
Polysorbate 80  
Colloidal anhydrous silica  
Saccharin sodium  
Cherry flavor Permaseal (11035-31)  
Magnesium stearate

### **6.2 Incompatibilities**

Not applicable.

### **6.3 Shelf life**

36 months

### **6.4 Special precautions for storage**

Do not store above 30°C.

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

Tablets are packed in PVC/Aclar/PVC-Alu blister packs.

Pack size:                      Carton containing a blister of 6 tablets.  
   Carton containing a blister of 12 tablets.  
   Carton containing 30 blisters of 6 tablets each.  
   Carton containing 30 blisters of 12 tablets each.

### **6.6 Special precautions for disposal**

#### **No special requirements.**

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

## **7. SUPPLIER**

Cipla Ltd.  
Cipla House  
Peninsula Business Park  
Ganpatrao Kadam Marg  
Lower Parel  
Mumbai: 400013  
India  
Tel: +91 22 24826000

## **8. WHO REFERENCE NUMBER (PREQUALIFICATION PROGRAMME)**

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## **9. DATE OF FIRST PREQUALIFICATION/ RENEWAL OF PREQUALIFICATION**

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

March 2018  
Section 6 updated in June 2019.

Detailed information on this medicine is available on the World Health Organization (WHO) web site:  
<https://extranet.who.int/prequal>

## References

General reference sources for this SmPC include:

Riamet Summary of Product Characteristics, Novartis Pharmaceuticals UK Ltd, last updated May 2017. Available at: <http://www.medicines.org.uk/emc/medicine/9196/SPC/>

Coartem label, Novartis Pharmaceuticals Corp. USA; last updated January 2018. Available at: <https://www.accessdata.fda.gov/scripts/cder/daf/index.cfm?event=BasicSearch.process>

Guidelines for the treatment of malaria -- 3rd edition, April 2015. Available at: <http://www.who.int/malaria/publications/atoz/9789241549127/en/>

### Section 4.5

University of Liverpool, HIV and Hepatitis Drug Interactions websites. Available at: <https://www.hiv-druginteractions.org/>  
<https://www.hep-druginteractions.org/>

### Section 4.6 and others (information related to use in pregnancy)

Malaria Policy Advisory Committee Meeting, 16–18 September 2015, Geneva, Switzerland, Background document for Session 4; WHO/HTM/GMP/MPAC/2015.13; Malaria in pregnancy. Available at: <http://www.who.int/malaria/mpac/mpac-sept2015-erg-mip-report.pdf?ua=1>

Dellicour S, Sevene E, McGready R et al (2017). First-trimester artemisinin derivatives and quinine treatments and the risk of adverse pregnancy outcomes in Africa and Asia: A meta-analysis of observational studies. PLoS Med 14(5): e1002290: <https://doi.org/10.1371/journal.pmed.1002290>

*All weblinks were last accessed 13 January 2018*