

## Part VI: Summary of the risk management plan

### Summary of risk management plan for Hydroxychloroquine Accord 200mg Film-coated Tablets (Hydroxychloroquine sulphate)

This is a summary of the risk management plan (RMP) for Hydroxychloroquine Accord 200mg Film-coated Tablets. The RMP details important risks of Hydroxychloroquine Accord 200mg Film-coated Tablets, how these risks can be minimised, and how more information will be obtained about Hydroxychloroquine Accord 200mg Film-coated Tablets's risks and uncertainties (missing information).

Hydroxychloroquine Accord 200mg Film-coated Tablets's summary of product characteristics (SmPC) and its package leaflet give essential information to healthcare professionals and patients on how Hydroxychloroquine Accord 200mg Film-coated Tablets should be used.

Important new concerns or changes to the current ones will be included in updates of Hydroxychloroquine Accord 200mg Film-coated Tablets's RMP.

#### I. The medicine and what it is used for

Hydroxychloroquine Accord 200mg Film-coated Tablets is authorized for following indications:

##### Adults

Hydroxychloroquine Accord tablets are recommended for the treatment of rheumatoid arthritis, discoid and systemic lupus erythematosus, and dermatological conditions caused or aggravated by sunlight.

This product is also indicated in adults for prevention and treatment of uncomplicated malaria caused by *Plasmodium vivax*, *P. ovale*, *P. malariae* and chloroquine sensitive *P. falciparum*.

##### Paediatric Population

Treatment of juvenile idiopathic arthritis (in combination with other therapies), discoid and systemic lupus erythematosus.

Also indicated for prevention and treatment of uncomplicated malaria, caused by *Plasmodium vivax*, *P. malariae*, *P. ovale* and chloroquine-sensitive *P. falciparum*.

It contains hydroxychloroquine sulphate as the active substance and it is given orally.

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## II. Risks associated with the medicine and activities to minimise or further characterise the risks

Important risks of Hydroxychloroquine Accord 200mg Film-coated Tablets together with measures to minimise such risks and the proposed studies for learning more about Hydroxychloroquine Accord 200mg Film-coated Tablets's risks, are outlined below.

Measures to minimise the risks identified for medicinal products can be:

- Specific information, such as warnings, precautions, and advice on correct use, in the package leaflet and SmPC addressed to patients and healthcare professionals;
- Important advice on the medicine's packaging;
- The authorised pack size - the amount of medicine in a pack is chosen so to ensure that the medicine is used correctly;
- The medicine's legal status - the way a medicine is supplied to the patient (e.g. with or without prescription) can help to minimise its risks.

Together, these measures constitute routine risk minimisation measures.

In addition to these measures, information about adverse reactions is collected continuously and regularly analysed, including PSUR assessment and signal management activity, so that immediate action can be taken as necessary. These measures constitute routine pharmacovigilance activities.

### II.A List of important risks and missing information

Important risks of Hydroxychloroquine Accord 200mg Film-coated Tablets are risks that need special risk management activities to further investigate or minimise the risk, so that the medicinal product can be safely taken. Important risks can be regarded as identified or potential. Identified risks are concerns for which there is sufficient proof of a link with the use of Hydroxychloroquine Accord 200mg Film-coated Tablets. Potential risks are concerns for which an association with the use of this medicine is possible based on available data, but this association has not been established yet and needs further evaluation. Missing information refers to information on the safety of the medicinal product that is currently missing and needs to be collected (e.g. on the long-term use of the medicine).

Important identified risks	<ul style="list-style-type: none"> <li>• Overdose</li> </ul>
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	<ul style="list-style-type: none"> <li>• Severe hypoglycaemia</li> <li>• Visual disturbance</li> <li>• Gastrointestinal effects</li> </ul>
Important potential risks	<ul style="list-style-type: none"> <li>• Patients with hepatic and renal insufficiency</li> <li>• Cardiac conduction disorders</li> <li>• Haematological effects</li> <li>• Musculoskeletal effects</li> <li>• Dermatitis</li> <li>• Use in pregnancy</li> <li>• Use in breastfeeding</li> </ul>
Missing information	<ul style="list-style-type: none"> <li>• None</li> </ul>

## II.B Summary of important risks

The safety information in the proposed Product Information is aligned to the reference medicinal product.

## II.C Post-authorisation development plan

### II.C.1 Studies which are conditions of the marketing authorisation

There are no studies which are conditions of the marketing authorisation or specific obligation of Hydroxychloroquine Accord 200mg Film-coated Tablets.

### II.C.2 Other studies in post-authorisation development plan

There are no studies required for Hydroxychloroquine Accord 200mg Film-coated Tablets.

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