

Part VI: Summary of the risk management plan

Summary of risk management plan for DECADRON OFTALMICO (Dexamethasone sodium phosphate)

This is a summary of the risk management plan (RMP) for DECADRON OFTALMICO. The RMP details important risks of DECADRON OFTALMICO, and how more information will be obtained about DECADRON OFTALMICO's risks and uncertainties (missing information).

DECADRON OFTALMICO's summary of product characteristics (SmPC) and its package leaflet give essential information to healthcare professionals and patients on how DECADRON OFTALMICO should be used.

I. The medicine and what it is used for

DECADRON OFTALMICO is authorised for treatment of non-infectious inflammatory conditions affecting the anterior segment of the eye (see SmPC for the full indication). It contains dexamethasone sodium phosphate as the active substance and it is given by the ophthalmological route. DECADRON OFTALMICO is a sterile solution that does not contain a preservative. The solution from one individual single dose container is to be used immediately after opening for administration to the affected eye(s). Any remaining contents must be discarded immediately after administration.

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

Important risks of DECADRON OFTALMICO, together with measures to minimise such 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.

II.A List of important risks and missing information

Important risks of DECADRON OFTALMICO are risks that need special risk management activities to further investigate or minimise the risk, so that the medicinal product can be safely administered. 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 DECADRON OFTALMICO. 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);

List of important risks and missing information	
Important identified risks	None
Important potential risks	None
Missing information	None

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 DECADRON OFTALMICO.

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

There are no studies required for DECADRON OFTALMICO.