

Part VI: Summary of the risk management plan

Summary of risk management plan for Efrinter (Ivermectin)

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

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

Important new concerns or changes to the current ones will be included in updates of Efrinter's RMP.

I. The medicine and what it is used for

Efrinter is authorised for the following indication:

- for the topical treatment of inflammatory lesions of rosacea (papulopustular) in adult patients. (see SmPC for the full indication).

It contains ivermectin as the active substance and it is given by cutaneous use only.

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

Important risks of Efrinter, together with measures to minimise such risks and the proposed studies for learning more about Efrinter'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, so that immediate action can be taken as necessary. These measures constitute *routine pharmacovigilance activities*.

If important information that may affect the safe use of Efrinter is not yet available, it is listed under 'missing information' below.

II.A List of important risks and missing information

Important risks of Efrinter 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 Efrinter. 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	• Contact dermatitis (allergic or irritant)
Important potential risks	• Systemic Allergic Reactions • Accidental oral ingestion •
Missing information	• Exposure during pregnancy • Exposure during lactation • Use longer than one year • Off-label use • Use with concomitant topical treatments • Use with laser or UV radiation

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 that are conditions of the marketing authorization or specific obligation of the Applicant's Efrinter.

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

There are no studies required for the Applicant's Efrinter.