

Evolan Pharma AB	Risk Management Plan	
Name of the medicinal product: Prednisolone 31.25 mg rectal solution	Version number: 0.1	1.8.2
Prednisolone 31.25 mg rectal solution		

PART VI: SUMMARY OF THE RISK MANAGEMENT PLAN

Summary of risk management plan for Prednisolone 31.25 mg rectal solution (prednisolone)

This is a summary of the risk management plan (RMP) for Prednisolone 31.25 mg rectal solution (hereafter referred as Prednisolone rectal solution). The RMP details important risks of Prednisolone rectal solution, how these risks can be minimised, and how more information will be obtained about Prednisolone rectal solution's risks and uncertainties (missing information).

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

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

I. The medicine and what it is used for

Prednisolone rectal solution is authorised for the treatment of ulcerative proctosigmoiditis (see SmPC for the full indication). It contains prednisolone sodium phosphate as the active substance and it is for rectal use.

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

Important risks of Prednisolone rectal solution, 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

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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 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 Prednisolone rectal solution 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 Prednisolone rectal solution. 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.

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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 Prednisolone rectal solution.

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

There are no studies required for Prednisolone rectal solution.