

Summary of risk management plan for Lidocaine Orion 20 mg/ml injection (lidocaine hydrochloride) Orion Corporation

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

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

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

I. The medicine and what it is used for

Lidocaine Orion is authorised to be used in cardiac arrhythmias; treatment of, and prophylaxis after, ventricular tachyarrhythmias including recurrence of ventricular fibrillation, especially in connection with myocardial infarction, mechanical myocardial irritation in connection with heart surgery, diagnostic procedures (e.g. cardiac catheterisation and angiocardiology), digitalis intoxication and intoxication with tricyclic antidepressants (see SmPC for the full indication). It contains lidocaine hydrochloride as the active substance and it is given by injection.

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

Important risks of Lidocaine Orion, together with measures to minimise such risks and the proposed studies for learning more about Lidocaine Orion'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*.

II.A List of important risks and missing information

This product has no safety concerns requiring additional pharmacovigilance actions or additional risk minimisation activities.

Safety concerns are adequately addressed in the product information.

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 Lidocaine Orion.

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

There are no studies required for Lidocaine Orion.