

# **EU Risk Management Plan**

**for**

## **Midazolam Accord 1 mg/ml solution for injection/infusion in pre-filled syringe (Midazolam)**

### **Part VI: Summary of the risk management plan**

#### **Summary of risk management plan for Midazolam Accord 1 mg/ml solution for injection/infusion in pre-filled syringe (Midazolam)**

This is a summary of the risk management plan (RMP) for Midazolam Accord 1 mg/ml solution for injection/infusion in pre-filled syringe. The RMP details important risks of Midazolam Accord 1 mg/ml solution for injection/infusion in pre-filled syringe, how these risks can be minimised, and how more information will be obtained about Midazolam Accord 1 mg/ml solution for injection/infusion in pre-filled syringe's risks and uncertainties (missing information).

Midazolam Accord 1 mg/ml solution for injection/infusion in pre-filled syringe's summary of product characteristics (SmPC) and its package leaflet give essential information to healthcare professionals and patients on how Midazolam Accord 1 mg/ml solution for injection/infusion in pre-filled syringe should be used.

Important new concerns or changes to the current ones will be included in updates of Midazolam Accord 1 mg/ml solution for injection/infusion in pre-filled syringe's RMP.

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

Midazolam Accord 1 mg/ml solution for injection/infusion in pre-filled syringe is a short acting sleep-inducing active substance that is indicated:

##### In adults:

- Conscious sedation before and during diagnostic or therapeutic procedures with or without local anaesthesia.
- Anaesthesia
  - Premedication before induction of anaesthesia
  - Induction of anaesthesia
  - As a sedative component in combined anaesthesia

- Sedation in Intensive Care Units

In children ( $\geq 12$  years age):

- Conscious sedation before and during diagnostic or therapeutic procedures with or without local anaesthesia.
- Anaesthesia
  - Premedication before induction of anaesthesia
- Sedation in Intensive Care Units

It contains midazolam as the active substance and it is given by intravenous route.

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

Important risks of Midazolam Accord 1 mg/ml solution for injection/infusion in pre-filled syringe together with measures to minimise such risks and the proposed studies for learning more about Midazolam Accord 1 mg/ml solution for injection/infusion in pre-filled syringe'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.

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

Important risks of Midazolam Accord 1 mg/ml solution for injection/infusion in pre-filled syringe 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 Midazolam Accord 1 mg/ml solution for injection/infusion in pre-filled syringe. 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>• None</li> </ul> |
| Important potential risks  | <ul style="list-style-type: none"> <li>• None</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 Midazolam Accord 1 mg/ml solution for injection/infusion in pre-filled syringe.

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

There are no studies required for Midazolam Accord 1 mg/ml solution for injection/infusion in pre-filled syringe.

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