Part VI: Summary of the risk management plan

Summary of risk management plan for Aripiprazole, 300 & 400 mg powder and solvent for prolonged-release suspension for injection

This is a summary of the risk management plan (RMP) for Aripiprazole, 300 & 400 mg powder and solvent for prolonged-release suspension for injection. The RMP details important risks of Aripiprazole, 300 & 400 mg powder and solvent for prolonged-release suspension for injection, how these risks can be minimized, and how more information will be obtained about Aripiprazole, 300 & 400 mg powder and solvent for prolonged-release suspension for injection's risks and uncertainties (missing information).

Aripiprazole, 300 & 400 mg powder and solvent for prolonged-release suspension for injection's summary of product characteristics (SmPC) and its package leaflet give essential information to healthcare professionals and patients on how Aripiprazole, 300 & 400 mg powder and solvent for prolonged-release suspension for injection should be used.

Important new concerns or changes to the current ones will be included in updates of Aripiprazole, 300 & 400 mg powder and solvent for prolonged-release suspension for injection's RMP.

I. The medicine and what it is used for

Aripiprazole, 300 & 400 mg powder and solvent for prolonged-release suspension for injection is authorized for maintenance treatment of schizophrenia in adult patients stabilized with oral aripiprazole (see SmPC for the full indication). It contains Aripiprazole as the active substance, and it is given by intramuscular administration.

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

Important risks of Aripiprazole, 300 & 400 mg powder and solvent for prolonged-release suspension for injection, together with measures to minimise such risks and the proposed studies for learning more about Aripiprazole, 300 & 400 mg powder and solvent for prolonged-release suspension for injection's risks, are outlined below.

Measures to minimize 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 minimization* measures.

In addition to these measures, information about adverse reactions is collected continuously and regularly analyzed 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 Aripiprazole, 300 & 400 mg powder and solvent for prolonged-release suspension for injection is not yet available, it is listed under 'missing information' below.

II.A List of important risks and missing information

Important risks of Aripiprazole, 300 & 400 mg powder and solvent for prolonged-release suspension for injection are risks that need special risk management activities to further investigate or minimize 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 Aripiprazole, 300 & 400 mg powder and solvent for prolonged-release suspension for injection. 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 risk	Extrapyramidal Symptoms (EPS), including tardive dyskinesia.
Important potential risk	Orthostatic hypotension.
Missing information	Use in pregnancy and lactation. Use in Elderly Patients above 65 Years of Age.

II.B Summary of important risks

The safety information in the proposed Product Information is aligned to the reference medicinal product.

II.C Post-authorization development plan

II.C.1 Studies which are conditions of the marketing authorization

There are no studies which are conditions of the marketing authorization or specific obligation of Aripiprazole, 300 & 400 mg powder and solvent for prolonged-release suspension for injection.

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

There are no studies required for Aripiprazole, 300 & 400 mg powder and solvent for prolonged-release suspension for injection.