

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

### **Summary of risk management plan for Furosemide Medical Valley 20 mg and 40 mg tablets (Furosemide)**

This is a summary of the risk management plan (RMP) for Furosemide Medical Valley 20 mg and 40 mg tablets. The RMP details important risks of Furosemide Medical Valley 20 mg and 40 mg tablets, how these risks can be minimised, and how more information will be obtained about Furosemide Medical Valley 20 mg and 40 mg tablets risks and uncertainties (missing information).

Furosemide Medical Valley 20 mg and 40 mg tablets summary of product characteristics (SmPC) and its package leaflet give essential information to healthcare professionals and patients on how Furosemide Medical Valley 20 mg and 40 mg tablets should be used.

Important new concerns or changes to the current ones will be included in updates of Furosemide Medical Valley 20 mg and 40 mg tablets RMP.

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

Furosemide Medical Valley 20 mg and 40 mg tablets is authorised for:

1) Treatment of oedema associated with

- cardiac disease

- liver disease

- renal disease including nephrotic syndrome. In patients with nephrotic syndrome, therapy of the underlying disorder has priority.

Treatment of pulmonary oedema

2) Arterial hypertension

It contains furosemide as the active substance, and it is given by the oral route of administration.

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

Important risks of Furosemide Medical Valley 20 mg and 40 mg tablets, 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 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 Furosemide Medical Valley 20 mg and 40 mg tablets is not yet available, it is listed under 'missing information' below.

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

Important risks of Furosemide Medical Valley 20 mg and 40 mg tablets 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 Furosemide Medical Valley 20 mg and 40 mg tablets. 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).

<b>List of important risks and missing information</b>	
Important identified risks	None
Important potential risks	<ul style="list-style-type: none"> <li>• Medication errors</li> <li>• Lack of efficacy</li> </ul>
Missing information	<ul style="list-style-type: none"> <li>• Use in pregnant and lactating women</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 Furosemide Medical Valley 20 mg and 40 mg tablets.

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

There are no studies required for Furosemide Medical Valley 20 mg and 40 mg tablets.