

Summary of risk management plan for benzydamine/cetylpyridinium chloride oralmucosal spray, solution

This is a summary of the risk management plan (RMP) for Benzydamine hydrochloride/ Cetylpyridinium Billev. The RMP details important risks of Benzydamine hydrochloride/ Cetylpyridinium Billev and how more information will be obtained about Benzydamine hydrochloride/ Cetylpyridinium Billev's risks and uncertainties (missing information).

Benzydamine hydrochloride/ Cetylpyridinium Billev's summary of product characteristics (SmPC) and its package leaflet give essential information to healthcare professionals and patients on how Benzydamine hydrochloride/ Cetylpyridinium Billev should be used.

Important new concerns or changes to the current ones will be included in updates of Benzydamine hydrochloride/ Cetylpyridinium Billev's RMP.

I. The medicine and what it is used for

Benzydamine hydrochloride/ Cetylpyridinium Billev is indicated in adults and children for anti-inflammatory, analgesic and antiseptic treatment of irritations in the throat, mouth and gums (see SmPC for the full indication). It contains benzydamine/cetylpyridinium chloride as the active substance and it is given by orally.

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

Important risks of Benzydamine hydrochloride/ Cetylpyridinium Billev together with measures to minimise such risks and the proposed studies for learning more about benzydamine/cetylpyridinium chloride '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

Important risks of Benzydamine hydrochloride/ Cetylpyridinium Billev 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 Benzydamine hydrochloride/ Cetylpyridinium Billev. 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 | Not applicable |
| Important potential risks | Not applicable |
| Missing information | Not applicable |

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 Benzydamine hydrochloride/ Cetylpyridinium Billev.

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

There are no studies required for Benzydamine hydrochloride/ Cetylpyridinium Billev.