

Part VI: Summary of the risk management plan

Summary of risk management plan for Abavalex 10 mg/ml oral solution (brivaracetam)

This is a summary of the risk management plan (RMP) for *Abavalex 10 mg/ml oral solution* [*Abavalex oral solution*]. The RMP details important risks of *Abavalex oral solution*, how these risks can be minimised, and how more information will be obtained about *Abavalex oral solution's* risks and uncertainties (missing information).

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

Important new concerns or changes to the current ones will be included in updates of *Abavalex oral solution's* RMP.

I. The medicine and what it is used for

Abavalex oral solution is authorised as adjunctive therapy in the treatment of partial onset seizures with or without secondary generalisation in adults, adolescents and children from 2 years of age with epilepsy (see SmPC for the full indication). It contains Brivaracetam as the active substance and it is given by oral route of administration.

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

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

If important information that may affect the safe use of *Abavalex oral solution* is not yet available, it is listed under 'missing information' below.

II.A List of important risks and missing information

Important risks of *Abavalex oral solution* 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 *Abavalex oral solution*. 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	<ul style="list-style-type: none">• Suicidality (class label for anticonvulsant products)
Important potential risks	<ul style="list-style-type: none">• None
Missing information	<ul style="list-style-type: none">• Data during pregnancy and lactation• Long-term effects on growth, endocrine function or sexual maturation, neurodevelopment, cognitive and psychomotor development in paediatric patients.

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 *Abavalex oral solution*.

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

There are no studies required for *Abavalex oral solution*.