

Summary of Risk Management Plan for INTELENCE (Etravirine)

This is a summary of the risk management plan (RMP) for INTELENCE. The RMP details important risks of INTELENCE, how these risks can be minimized, and how more information will be obtained about INTELENCE's risks and uncertainties (missing information).

INTELENCE's Summary of Product Characteristics (SmPC) and its Package Leaflet (PL) give essential information to healthcare professionals and patients on how INTELENCE should be used.

This summary of the RMP for INTELENCE should be read in the context of all this information including the assessment report of the evaluation and its plain-language summary, all of which is part of the European Public Assessment Report (EPAR).

Important new concerns or changes to the current ones will be included in updates of INTELENCE's RMP.

I. The Medicine and What it is Used For

INTELENCE is authorized in combination with a boosted protease inhibitor (PI) and other antiretroviral (ARV) medicinal products for the treatment of human immunodeficiency virus (HIV)-1 infection in ARV treatment-experienced adult patients and in antiretroviral therapy (ART)-experienced pediatric patients from 2 years of age (see SmPC for the full indication). It contains etravirine (ETR or TMC125) as the active substance and it is given as an oral tablet (ETR 25 mg, 100 mg, 200 mg).

Further information about the evaluation of INTELENCE's benefits can be found in INTELENCE's EPAR, including in its plain-language summary, available on the European Medicines Agency website, under the medicine's webpage:

<https://www.ema.europa.eu/en/medicines/human/EPAR/intelence>

II. Risks Associated with the Medicine and Activities to Minimize or Further Characterize the Risks

Important risks of INTELENCE, together with measures to minimize such risks and the proposed studies for learning more about INTELENCE'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 PL and SmPC addressed to patients and healthcare professionals;
- Important advice on the medicine's packaging;
- The authorized 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 (eg, with or without prescription) can help to minimize 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, including Periodic Benefit-Risk Evaluation Report /Periodic Safety Update Report (PSUR) assessment 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 INTELENCE is not yet available, it is listed under ‘missing information’ below.

II.A. List of Important Risks and Missing Information

Important risks of INTELENCE 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 INTELENCE. 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 (eg, on the long-term use of the medicine).

List of Important Risks and Missing Information	
Important identified risks	None
Important potential risks	None
Missing information	None

II.B. Summary of Important Risks

Not applicable.

II.C. Postauthorization 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 INTELENCE.

II.C.2. Other Studies in Postauthorization Development Plan

There are no studies required for INTELENCE.