

Summary of risk management plan Ticagrelor Pharmascience (ticagrelor)

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

Ticagrelor Pharmascience's summary of product characteristics (SmPC) and its package leaflet give essential information to healthcare professionals and patients on how Tarelik should be used.

I. The medicine and what it is used for

Ticagrelor Pharmascience is co-administered with acetylsalicylic acid (ASA), is indicated for the prevention of atherothrombotic events in adult patients with

- acute coronary syndromes (ACS) or
- a history of myocardial infarction (MI) and a high risk of developing an atherothrombotic event (see SmPC for the full indication). It contains ticagrelor as the active substance and it is given by oral administration.

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

Important risks of Ticagrelor Pharmascience, together with measures to minimise such risks and the proposed studies for learning more about Ticagrelor Pharmascience'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 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 Ticagrelor Pharmascience is not yet available, it is listed under 'missing information' below.

II.A List of important risks and missing information

Important risks of Ticagrelor Pharmascience 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 Ticagrelor Pharmascience. 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"> • Increased risk of bleeding
Important potential risks	<ul style="list-style-type: none"> • None
Missing information	<ul style="list-style-type: none"> • Long-term use in patients with prior ischaemic stroke

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

Not applicable.

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

Not applicable.

Summary of risk management plan Ticotromb (ticagrelor)

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

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

I. The medicine and what it is used for

Ticotromb is co-administered with acetylsalicylic acid (ASA), is indicated for the prevention of atherothrombotic events in adult patients with

- acute coronary syndromes (ACS) or
- a history of myocardial infarction (MI) and a high risk of developing an atherothrombotic event (see SmPC for the full indication). It contains ticagrelor as the active substance and it is given by oral administration.

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

Important risks of Ticotromb, together with measures to minimise such risks and the proposed studies for learning more about Ticotromb'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 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 Ticotromb is not yet available, it is listed under 'missing information' below.

II.A List of important risks and missing information

Important risks of Ticotromb 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 Ticotromb. 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">• Increased risk of bleeding
Important potential risks	<ul style="list-style-type: none">• None
Missing information	<ul style="list-style-type: none">• Long-term use in patients with prior ischaemic stroke

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

Not applicable.

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

Not applicable.