

B. PACKAGE LEAFLET

Package leaflet: Information for the user

Decitabine Accord 40 mg powder for concentrate for solution for infusion **Decitabine Accord 50 mg powder for concentrate for solution for infusion** decitabine

Read all of this leaflet carefully before you start using this medicine because it contains important information for you.

- Keep this leaflet. You may need to read it again.
- If you have any further questions, ask your doctor<,> <or> pharmacist < or nurse>.
- This medicine has been prescribed for you only. Do not pass it to others. It may harm them, even if their signs of illness are the same as yours.
- If you get any side effects, talk to your doctor, pharmacist or nurse. This includes any possible side effects not listed in this leaflet. See section 4.

What is in this leaflet

1. What Decitabine Accord is and what it is used for
2. What you need to know before you use Decitabine Accord
3. How to use Decitabine Accord
4. Possible side effects
5. How to store Decitabine Accord
6. Contents of the pack and other information

1. What Decitabine Accord is and what it is used for

What Decitabine Accord is

Decitabine Accord is an anti-cancer medicine. It contains the active substance ‘decitabine’.

What Decitabine Accord is used for

Decitabine Accord is used to treat a type of cancer called ‘acute myeloid leukaemia’ or ‘AML’. This is a type of cancer that affects your blood cells. You will be given Decitabine Accord when you are first diagnosed with AML. It is used in adults.

How Decitabine Accord works

Decitabine Accord works by stopping cancer cells from growing. It also kills cancer cells.

Talk to your doctor < or nurse> if you have any questions about how Decitabine Accord works or why this medicine has been prescribed for you.

2. What you need to know before you are given Decitabine Accord

Do not use Decitabine Accord

- if you are allergic to decitabine or any of the other ingredients of this medicine (listed in section 6).
- if you are breast-feeding.

If you are not sure if any of the above applies to you, talk to your doctor<,> <or> pharmacist < or nurse> before using Decitabine Accord.

Warnings and precautions

Talk to your doctor<,> <or> pharmacist < or nurse> before using Decitabine Accord if you have

- low numbers of platelets, red blood cells or white blood cells,

- an infection,
- liver disease,
- a serious kidney disorder,
- a heart disorder.

If you are not sure if any of the above applies to you, talk to your doctor<, > <or> pharmacist < or nurse> before using Decitabine Accord.

Decitabine Accord can cause a serious immune reaction called ‘differentiation syndrome’ (see section 4 ‘Possible side effects’).

Tests or checks

You will have blood tests before you start treatment with Decitabine Accord and at the start of each treatment cycle. These tests are to check that:

- you have enough blood cells, and
- your liver and kidneys are working properly.

Talk to your doctor about what your blood test results mean.

Children and adolescents

Decitabine Accord is not for use in children or adolescents under the age of 18.

Other medicines and Decitabine Accord

Tell your doctor<, nurse> or pharmacist if you are using, have recently used or might use any other medicines. This includes medicines obtained without a prescription and herbal medicines. This is because Decitabine Accord can affect the way some other medicines work. Also, some other medicines can affect the way Decitabine Accord works.

Pregnancy and breast-feeding

- If you are pregnant or breast-feeding, think you may be pregnant or are planning to have a baby, ask your doctor for advice before using this medicine.
- You should not use Decitabine Accord if you are pregnant as it may harm your baby. If you are able to become pregnant, your doctor will ask you to take a pregnancy test before you start treatment with Decitabine Accord. Tell your doctor immediately if you become pregnant during treatment with Decitabine Accord.
- Do not breast-feed if you are using Decitabine Accord. This is because it is not known if the medicine passes into the mother’s milk.

Male and female fertility and contraception

- Men should not father a child while using Decitabine Accord.
- Men should use effective contraception during treatment and for up to 3 months after treatment has stopped.
- Talk to your doctor if you wish to conserve your sperm before starting treatment.
- Women who are able to become pregnant must use effective contraception during treatment and for 6 months following completion of treatment.
- Talk to your doctor if you wish to freeze your eggs before starting treatment.

Driving and using machines

You may feel tired or weak after using Decitabine Accord. If this happens, do not drive or use any tools or machines.

Decitabine Accord contains potassium and sodium

40 mg decitabine per vial

- This medicine contains 0.4 mmol (15.6 mg) potassium in each vial. After preparing the medicine, it contains less than 1 mmol (39 mg) of potassium per dose, i.e. essentially ‘potassium- free’.

- This medicine contains 0.23 mmol (5.3 mg) sodium (main component of cooking/table salt) in each vial. After preparing the medicine, it contains between 5.3 mg-885 mg sodium per dose, equivalent to 0.3-44% of the recommended maximum daily dietary intake of sodium for an adult. Talk to your doctor if you are on a low salt diet.

50 mg decitabine per vial

- This medicine contains 0.5 mmol (19.5 mg) potassium in each vial. After preparing the medicine, it contains less than 1 mmol (39 mg) of potassium per dose, i.e. essentially 'potassium-free'.
- This medicine contains 0.29 mmol (6.7 mg) sodium (main component of cooking/table salt) in each vial. After preparing the medicine, it contains between 5.3 mg-885 mg sodium per dose, equivalent to 0.3-44% of the recommended maximum daily dietary intake of sodium for an adult. Talk to your doctor if you are on a low salt diet.

3. How to use Decitabine Accord

Decitabine Accord will be given to you by a doctor < or nurse > who is trained in giving this type of medicine.

How much to use

- Your doctor will work out your dose of Decitabine Accord. This depends on your height and weight (body surface area).
- The dose is 20 mg/m² body surface area.
- You will receive Decitabine Accord every day for 5 days, then 3 weeks without the medicine. This is called a 'treatment cycle' and it is repeated every 4 weeks. You will usually receive at least 4 treatment cycles.
- Your doctor may delay your dose and change the total number of cycles, depending on how you respond to the treatment.

How Decitabine Accord is given

The solution is given into a vein (as an infusion). This will take one hour.

If you are given more Decitabine Accord than you should

This medicine will be given by your doctor < or nurse >. In the unlikely event that you are given too much (an overdose) your doctor will check you for side effects and manage them accordingly.

If you forget your appointment to have Decitabine Accord

If you miss an appointment, make another one as soon as possible. This is because for this medicine to be as effective as possible, it is important to follow the dosing schedule.

If you have any further questions on the use of this medicine, ask your doctor < or nurse >.

4. Possible side effects

Like all medicines, this medicine can cause side effects, although not everybody gets them. The following side effects may happen with this medicine.

Tell your doctor or nurse immediately if you notice any of the following serious side effects

- Fever: this may be a sign of an infection caused by low levels of white blood cells (very common).
- Chest pain or shortness of breath (with or without fever or cough): these may be signs of an infection of the lung called "pneumonia" (very common) or inflamed lungs (interstitial lung disease [frequency not known]) or cardiomyopathy (heart muscle disease [uncommon]) which can be accompanied with swelling of ankles, hands, legs and feet.

- Bleeding: including blood in the stools. This may be a sign of bleeding in the stomach or gut (common).
- Difficulty with moving, speaking or understanding or seeing; sudden severe headache, seizure, numbness or weakness in any part of the body. These may be signs of bleeding inside your head (common).
- Difficulty breathing, swelling of the lips, itching or rash: This may be due to an allergic (hypersensitivity) reaction (common).
- Serious immune reaction (differentiation syndrome) that may cause fever, cough, difficulty breathing, rash, decreased urine, hypotension (low blood pressure), swelling of the arms or legs and rapid weight gain (not known).

Tell your doctor or nurse immediately if you notice any of the serious side effects above.

Other side effects of Decitabine Accord include

Very common (may affect more than 1 in 10 people)

- urine infection
- other infection in any part of the body, caused by bacteria, virus or fungi
- bleeding or bruising more easily - these may be signs of a drop in the number of blood platelets (thrombocytopenia)
- feeling tired or looking pale - these may be signs of a drop in the number of red blood cells (anaemia)
- high level of sugar in the blood
- headache
- nose bleeds
- diarrhoea
- vomiting
- nausea
- fever
- abnormal liver function

Common (may affect up to 1 in 10 people)

- an infection of the blood caused by bacteria - this may be a sign of a low level of white blood cells
- sore or runny nose, sore sinuses
- mouth or tongue ulcers
- high level of 'bilirubin' in the blood

Uncommon (may affect up to 1 in 100 people)

- a drop in the number of red blood cells, white blood cells and platelets (pancytopenia)
- heart muscle disease
- red, raised painful patches on the skin, fever, an increase in white blood cells - these may be signs of 'Acute Febrile Neutrophilic Dermatitis' or 'Sweet's Syndrome'

Not known (frequency cannot be estimated from the available data)

- inflamed gut (enterocolitis, colitis and caecitis), with symptoms of abdominal pain, bloating, or diarrhoea. Enterocolitis may lead to septic complications and may be associated with fatal outcome.

Reporting of side effects

If you get side effects, talk to your doctor, pharmacist or nurse. This includes any possible side effects not listed in this leaflet. You can also report side effects directly via

HPRA Pharmacovigilance

Website: www.hpra.ie

By reporting side effects you can help provide more information on the safety of this medicine.

5. How to store Decitabine Accord

- Your doctor<, nurse> or pharmacist is responsible for storing Decitabine Accord.
- Keep this medicine out of the sight and reach of children.
- Do not use this medicine after the expiry date which is stated on the carton and on the vial label after EXP. The expiry date refers to the last day of that month.
- After reconstitution, the concentrate must be further diluted within 15 minutes using cold infusion fluids. This prepared diluted solution can be stored refrigerated at 2°C - 8°C for up to a maximum of 4 hours, followed by up to 3 hours at room temperature (20°C - 25°C) before administration.
- Do not use the product, if reconstituted or diluted solution exhibits discoloration.
- Your doctor<, nurse> or pharmacist is responsible for disposing of any unused Decitabine Accord correctly.

6. Contents of the pack and other information

What Decitabine Accord contains

40 mg decitabine per vial

- The active substance is decitabine. Each vial of powder contains 40 mg decitabine. After reconstitution with 8 ml of water for injections, each ml of concentrate contains 5 mg of decitabine.
- The other ingredients are potassium dihydrogen phosphate (E340) and sodium hydroxide (E524) (for pH adjustment). See section 2.

50 mg decitabine per vial

- The active substance is decitabine. Each vial of powder contains 50 mg decitabine. After reconstitution with 10 ml of water for injections, each ml of concentrate contains 5 mg of decitabine.
- The other ingredients are potassium dihydrogen phosphate (E340) and sodium hydroxide (E524) (for pH adjustment). See section 2.

What Decitabine Accord looks like and contents of the pack

40 mg decitabine per vial

Decitabine Accord is a white to off-white powder for concentrate for solution for infusion. It is supplied in a 20 ml glass vial containing 40 mg decitabine. Each pack contains 1 vial.

50 mg decitabine per vial

Decitabine Accord is a white to off-white powder for concentrate for solution for infusion. It is supplied in a 20 ml glass vial containing 50 mg decitabine. Each pack contains 1 vial.

Marketing Authorisation Holder

Accord Healthcare Ireland Ltd
Euro House
Euro Business Park
Little Island
Cork T45 K857
Ireland

Manufacturer

Accord Healthcare Polska Sp.z.o.o.

ul. Lutomierska 50,
95-200, Pabianice,
Poland

Accord Healthcare Single Member S.A.
64th Km National Road Athens,
Lamia, 32009, Greece

Laboratori Fundació Dau
C/ C, 12-14 Pol. Ind.
Zona Franca, Barcelona, 08040, Spain

This medicine is authorised in the Member States of the European Economic Area under the following names:

Member States	Proposed (Invented) Names
Austria	Decitabin Accord 40 mg Pulver für ein Konzentrat zur Herstellung einer Infusionslösung Decitabin Accord 50 mg Pulver für ein Konzentrat zur Herstellung einer Infusionslösung
Germany	Decitabin Accord 40 mg Pulver für ein Konzentrat zur Herstellung einer Infusionslösung Decitabin Accord 50 mg Pulver für ein Konzentrat zur Herstellung einer Infusionslösung
Lithuania	Decitabine Accord 40 mg milteliai infuzinio tirpalo koncentratui Decitabine Accord 50 mg milteliai infuzinio tirpalo koncentratui
Norway	Decitabine Accord Decitabine Accord
Sweden	Decitabine Accord 40 mg Decitabine Accord 50 mg
Ireland	Decitabine Accord 40 mg powder for concentrate for solution for infusion Decitabine Accord 50 mg powder for concentrate for solution for infusion
Cyprus	Decitabine Accord 40 mg Κόνις για πυκνό σκεύασμα για παρασκευή διαλύματος προς έγχυση Decitabine Accord 50 mg Κόνις για πυκνό σκεύασμα για παρασκευή διαλύματος προς έγχυση
Italy	Decitabina Accord Decitabina Accord
Portugal	Decitabina Accord 40 mg Pó para concentrado para solução para perfusão Decitabina Accord 50 mg Pó para concentrado para solução para perfusão
Spain	Decitabina Accord 40 mg polvo para concentrado para solución para perfusión Decitabina Accord 50 mg polvo para concentrado para solución para perfusión EFG
Croatia	Decitabine Accord 40 mg prašak za koncentrat za otopinu za infuziju Decitabine Accord 50 mg prašak za koncentrat za otopinu za infuziju
Hungary	Decitabine Accord 40 mg por oldatos infúzióhoz való koncentrátumhoz Decitabine Accord 50 mg por oldatos infúzióhoz való koncentrátumhoz
Romania	Decitabină Accord 40 mg pulbere pentru concentrat pentru soluție perfuzabilă Decitabină Accord 50 mg pulbere pentru concentrat pentru soluție perfuzabilă
Slovenia	Decitabin Accord 40 mg prašek za koncentrat za raztopino za infundiranje Decitabin Accord 50 mg prašek za koncentrat za raztopino za infundiranje

This leaflet was last revised in 12/2025

The following information is intended for medical or healthcare professionals only:

1. RECONSTITUTION

Skin contact with the solution should be avoided and protective gloves must be worn. Standard procedures for dealing with cytotoxic medicinal products should be adopted.

40 mg decitabine per vial

The powder should be aseptically reconstituted with 8 ml of water for injections. Upon reconstitution, each ml contains approximately 5 mg of decitabine at pH 6.7 to 7.3. Within 15 minutes of reconstitution, the solution must be further diluted with cold (2°C - 8°C) infusion fluids (sodium chloride 9 mg/ml [0.9%] solution for injection or 5% glucose solution for injection) to a final concentration of 0.15 to 1.0 mg/ml.

50 mg decitabine per vial

The powder should be aseptically reconstituted with 10 ml of water for injections. Upon reconstitution, each ml contains approximately 5 mg of decitabine at pH 6.7 to 7.3. Within 15 minutes of reconstitution, the solution must be further diluted with cold (2°C - 8°C) infusion fluids (sodium chloride 9 mg/ml [0.9%] solution for injection or 5% glucose solution for injection) to a final concentration of 0.15 to 1.0 mg/ml.

For the shelf-life and the precautions for storage after reconstitution, see section 5 of the leaflet.

2. ADMINISTRATION

Infuse the reconstituted solution intravenously over 1 hour.

3. DISPOSAL

A vial is for single use only and any remaining solution must be discarded.

Any unused medicinal product or waste material should be disposed of in accordance with local requirements.