

Consumer Medicine Information (CMI) summary

The [full CMI](#) on the next page has more details. If you are worried about your child receiving this vaccine, speak to your doctor or pharmacist.

1. Why is my child receiving Hexaxim?

Hexaxim contains the active ingredients diphtheria, tetanus, pertussis (acellular, component), hepatitis B (rDNA), poliomyelitis (inactivated) and haemophilus influenzae type b conjugate vaccine (adsorbed). Hexaxim is a vaccine used to help protect your child against diphtheria, tetanus, pertussis, hepatitis B, poliomyelitis and serious diseases caused by Haemophilus influenzae type b.

For more information, see Section [1. Why is my child receiving Hexaxim?](#) in the full CMI.

2. What should I know before my child receives Hexaxim?

Your child should not receive Hexaxim if they have ever had an allergic reaction to Hexaxim or any of the ingredients listed at the end of the CMI.

Talk to your doctor your child has any other medical conditions or takes any other medicines.

For more information, see Section [2. What should I know before my child receives Hexaxim?](#) in the full CMI.

3. What if my child is taking other medicines?

Some medicines may interfere with Hexaxim and affect how it works. A list of these medicines is in Section [3. What if my child is taking other medicines?](#) in the full CMI.

4. How does my child receive Hexaxim?

- Hexaxim is injected into the muscle in the upper part of your child's leg or upper arm, by a doctor or nurse.
- Your child will receive two injections given at an interval of at least eight weeks apart, or three injections given at an interval of at least four weeks apart. This vaccine should be used according to the local vaccination program.

More instructions can be found in Section [4. How does my child receive Hexaxim?](#) in the full CMI.

5. What should I know after my child receives Hexaxim?

Things you should do	• Keep an updated record of your child's vaccination history. • Attend any appointments made by your doctor or nurse. • Report any side effects to your doctor.
Looking after your medicine	• Hexaxim is usually stored in the doctor's surgery or clinic. However, if you need to store Hexaxim, keep it in its original packaging in the refrigerator, between 2°C and 8°C. Do not freeze Hexaxim.

For more information, see Section [5. What should I know after my child receives Hexaxim?](#) in the full CMI.

6. Are there any side effects?

The most common side effects of Hexaxim are pain, redness or swelling at the injection site, loss of appetite, crying, sleepiness, vomiting, irritability, fever (temperature 38°C or higher), abnormal crying (prolonged crying), diarrhoea and injection site hardness.

Serious side effects can include episodes when your child goes into a shock-like state or is pale, floppy and unresponsive for a period of time (hypotonic reactions or hypotonic hyporesponsive episodes HHE), serious allergic reaction (anaphylactic reaction) and fits (convulsions) with or without fever.

For more information, including what to do if your child has any side effects, see Section [6. Are there any side effects?](#) in the full CMI.

HEXAXIM

Active ingredients: *acellular, component*), hepatitis B (rDNA), poliomyelitis (inactivated) and haemophilus influenzae type b conjugate vaccine (adsorbed)

Consumer Medicine Information (CMI)

This leaflet provides important information about receiving Hexaxim. **You should also speak to your doctor or pharmacist if you would like further information or if you have any concerns or questions about your child receiving Hexaxim.**

Where to find information in this leaflet:

1. [Why is my child receiving Hexaxim?](#)
2. [What should I know before my child receives Hexaxim?](#)
3. [What if my child is taking other medicines?](#)
4. [How does my child receive Hexaxim?](#)
5. [What should I know after my child receives Hexaxim?](#)
6. [Are there any side effects?](#)
7. [Product details](#)

1. Why is my child receiving Hexaxim?

Hexaxim contains the active ingredients diphtheria, tetanus, pertussis (acellular, component), hepatitis B (rDNA), poliomyelitis (inactivated) and haemophilus influenzae type b conjugate vaccine (adsorbed). Hexaxim is a vaccine used to protect against infectious diseases.

Hexaxim is used to help protect infants against diphtheria, tetanus, pertussis, hepatitis B, poliomyelitis and serious diseases caused by haemophilus influenzae type b. Hexaxim can be given to children from six weeks of age.

The vaccine works by causing the body to produce its own protection (antibodies) against the bacteria and viruses that cause these different infections:

- Diphtheria is an infectious disease that usually first affects the throat. In the throat, the infection causes pain and swelling which can lead to suffocation. The bacteria that cause the disease also make a toxin (poison) that can damage the heart, kidneys and nerves.
- Tetanus (often called lock jaw) is usually caused by the tetanus bacteria entering a deep wound. The bacteria make a toxin (poison) that causes spasms of the muscles, leading to inability to breathe and the possibility of suffocation.
- Pertussis (often called whooping cough) is a highly infectious illness that affects the airways. It causes severe coughing that may lead to problems with breathing. The coughing often has a "whooping" sound. The cough may last for one to two months or longer. Whooping cough can also cause ear infections, chest infections (bronchitis) which may last a long time, lung infections (pneumonia), fits, brain damage and even death.

- Hepatitis B is caused by the hepatitis B virus. It causes the liver to become swollen (inflamed). In some people, the virus can stay in the body for a long time, and can eventually lead to serious liver problems, including liver cancer
- Poliomyelitis (often just called polio) is caused by viruses that affect the nerves. It can lead to paralysis or muscle weakness most commonly of the legs. Paralysis of the muscles that control breathing and swallowing can be fatal.
- Haemophilus influenzae type b infections (often just called Hib) are serious bacterial infections and can cause meningitis (inflammation of the outer covering of the brain), which can lead to brain damage, deafness, epilepsy, or partial blindness. Infection can also cause inflammation and swelling of the throat, leading to difficulties in swallowing and breathing, and infection can affect other parts of the body such as the blood, lungs, skin, bones, and joints.

Important information about the protection provided

Hexaxim will only help to prevent these diseases if they are caused by the bacteria or viruses targeted by the vaccine. Your child could get diseases with similar symptoms if they are caused by other bacteria or viruses.

The vaccine does not contain any live bacteria or viruses and it cannot cause any of the infectious diseases against which it protects.

This vaccine does not protect against infections caused by other types of Haemophilus influenzae nor against meningitis due to other micro-organisms.

Hexaxim will not protect against hepatitis infection caused by other agents such as hepatitis A, hepatitis C and hepatitis E.

Because symptoms of hepatitis B take a long time to develop, it is possible for unrecognised hepatitis B infection to be present at the time of vaccination. The vaccine may not prevent hepatitis B infection in such cases.

Remember that no vaccine can provide complete, lifelong protection in all people who are vaccinated.

2. What should I know before my child receives Hexaxim?

Warnings

Your child should not receive Hexaxim if:

- they have had a previous allergic reaction to Hexaxim, or any of the ingredients listed at the end of this leaflet.
- Always check the ingredients to make sure you can use this medicine.

- has had respiratory disorder or swelling of the face (anaphylactic reaction) after administration of Hexaxim.
- has had an allergic reaction to any other diphtheria, tetanus, pertussis, poliomyelitis, hepatitis B or Hib containing vaccines.
- suffered from a severe reaction affecting the brain (encephalopathy) within 7 days of a prior dose of a pertussis vaccine (acellular or whole cell pertussis).
- has an uncontrolled condition or severe illness affecting the brain and nervous system (uncontrolled neurologic disorder) or uncontrolled epilepsy.

Check with your doctor if your child:

- has any other medical conditions.
- has a moderate or high temperature or an acute illness (e.g. fever, sore throat, cough, cold or flu). Vaccination with Hexaxim may need to be delayed until your child is better.
- has had any of the following events after receiving a pertussis vaccine, as the decision to give further doses of pertussis containing vaccine will need to be carefully considered:
 - o fever of 40°C or above within 48 hours not due to another identifiable cause.
 - o collapse or shock-like state with hypotonic-hyporesponsive episode (drop in energy) within 48 hours of vaccination.
 - o persistent, inconsolable crying lasting 3 hours or more, occurring within 48 hours of vaccination.
 - o fits (convulsions) with or without fever, occurring within 3 days of vaccination.
- previously had Guillain-Barre syndrome (temporary inflammation of nerves causing pain, paralysis and sensitivity disorders) or brachial neuritis (severe pain and decreased mobility of arm and shoulder) after being given a vaccine containing tetanus toxoid (an inactivated form of tetanus toxin). In this case, the decision to give any further vaccine containing tetanus toxoid should be evaluated by your doctor.
- is having a treatment that suppresses her/his immune system (the body's natural defences) or has any disease that causes the weakness of the immune system. In these cases, the immune response to the vaccine may be decreased. It is normally recommended to wait until the end of the treatment or disease before vaccinating. However, children with long standing problems with their immune system such as HIV infection (AIDS) may still be given Hexaxim but the protection may not be as good as in children whose immune system is healthy.
- is born prematurely. Lower responses to the vaccine may be observed in relation with immaturity of the immune system. However, according to national recommendations, vaccination should not be delayed. In addition, longer gaps than normal between breaths may occur for 2-3 days after vaccination.
- suffers from an acute or chronic illness including chronic renal insufficiency or failure (inability of the kidneys to work properly).

- suffers from any undiagnosed illness of the brain or epilepsy which is not controlled. Your doctor will assess the potential benefit offered by vaccination.
- has any problems with the blood that cause easy bruising or bleeding for a long time after minor cuts. Your doctor will advise you whether your child should have Hexaxim.
- has fainted when having a previous injection. Fainting can occur before or following needle injection.
- take any medicines for any other condition.

During treatment, your child may be at risk of developing certain side effects. It is important you understand these risks and how to monitor for them. See additional information under Section [6. Are there any side effects?](#)

3. What if my child is taking other medicines?

Tell your doctor or pharmacist if your child is taking any other medicines, including any medicines, vitamins or supplements that you buy without a prescription from your pharmacy, supermarket or health food shop.

Some medicines may interfere with Hexaxim and affect how it works.

Having other vaccines

Your doctor will advise you if Hexaxim is to be given with another vaccine.

Check with your doctor or pharmacist if you are not sure about what medicines, vitamins or supplements your child is taking and if these affect Hexaxim.

4. How does my child receive Hexaxim?

How is Hexaxim given

- Hexaxim is administered to your child by your doctor or nurse
- Hexaxim is injected into the muscle in the upper part of your child's leg or upper arm

When is Hexaxim given

First course of vaccination (primary vaccination)

- Your child will receive two injections given at an interval of at least eight weeks apart, or three injections given at an interval of at least four weeks apart.
- This vaccine should be used according to the local vaccination programme.

Additional injections (booster)

- After the first course of injections, your child may require a booster dose, in accordance with local recommendations.
- Your doctor will advise you if your child requires a booster dose.

If you forget a dose of Hexaxim

If your child misses a scheduled injection, it is important that you discuss with your doctor or nurse who will decide when to give the missed dose.

It is important to follow the instructions from the doctor or nurse so that your child completes the course of injections. If not, your child may not be fully protected against the diseases.

If you have any further questions on the use of this vaccine, ask your doctor, pharmacist or nurse.

If you use too much Hexaxim

Overdose is unlikely as your doctor is giving you the injection.

If you think that your child has been given too much Hexaxim, your child may need urgent medical attention.

You should immediately:

- phone the Poisons Information Centre (by calling 13 11 26), or
- contact your doctor or nurse, or
- go to the Emergency Department at your nearest hospital.

You should do this even if there are no signs of discomfort or poisoning.

5. What should I know after my child receives Hexaxim?

Things you should do

- Keep an updated record of your child's vaccination history.
- Attend any appointments made by your doctor or nurse.
- Report any side effects to your doctor.

Looking after your medicine

Follow the instructions in the carton on how to take care of your medicine properly.

Hexaxim is usually stored in the doctor's surgery or clinic, or at the pharmacy. However, if you need to store Hexaxim:

- Keep out of reach and sight of children.
- Keep Hexaxim in the original pack until it is time for it to be given.
- Keep it in the refrigerator, store at 2°C to 8°C. Do not freeze Hexaxim.
- Do not use Hexaxim if the packaging is torn or shows signs of tampering.

Keep it where young children cannot reach it.

Getting rid of any unwanted medicine

If you no longer need to use this medicine or it is out of date, take it to any pharmacy for safe disposal.

Medicines should not be disposed of via wastewater or household waste. Ask your pharmacist how to dispose of

medicines no longer required. These measures will help to protect the environment.

Do not use this medicine after the expiry date.

6. Are there any side effects?

All medicines can have side effects. If you do experience any side effects, most of them are minor and temporary. However, some side effects may need medical attention.

See the information below and, if you need to, ask your doctor or pharmacist if you have any further questions about side effects.

Less serious side effects

Less serious side effects	What to do
Gastrointestinal-related: • Vomiting • diarrhoea General and administration site-related: • pain, redness or swelling at the injection site • sleepiness • irritability • fever (temperature 38°C or higher) or high fever (temperature 39.6°C or higher) • injection site hardness • large reactions at the injection site (larger than 5 cm), including extensive limb swelling from the injection site beyond one or both joints. These reactions start within 24-72 hours after vaccination, may be associated with redness, warmth, tenderness or pain at the injection site, and get better within 3-5 days without the need for treatment Metabolism and nutrition-related: • loss of appetite Nervous system-related: • crying • abnormal crying (prolonged crying) Skin and subcutaneous tissue-related: • rash	Speak to your doctor if you have any of these less serious side effects and they worry you.

Serious side effects

Serious side effects	What to do
Allergy-related: • allergic reaction	Call your doctor straight

Serious side effects	What to do
- serious allergic reaction (anaphylactic reaction) - difficulty in breathing - blueness of the tongue or lips - a rash - swelling of the face or throat - low blood pressure causing dizziness or collapse Serious allergic reactions are a very rare possibility (may affect up to 1 in 10,000 people) after receiving any vaccine. Signs and symptoms of serious allergic reactions usually develop quickly after the injection is given and while the child is still in the clinic or doctor's surgery. Nervous system-related: - episodes when your child goes into a shock-like state or is pale, floppy and unresponsive for a period of time (hypotonic reactions or hypotonic hyporesponsive episodes HHE). - fits (convulsions) with or without fever	away, or go straight to the Emergency Department at your nearest hospital if you notice any of these serious side effects.

Tell your doctor or pharmacist if you notice anything else that may be making you feel unwell.

Other side effects not listed here may occur in some people.

Other side effects not listed above have been reported occasionally with other diphtheria, tetanus, pertussis, poliomyelitis, hepatitis B or Hib containing vaccines and not directly with Hexaxim:

- temporary inflammation of nerves causing pain, paralysis and sensitivity disorders (Guillain-Barré syndrome) and severe pain and decreased mobility of arm and shoulder (brachial neuritis) have been reported after administration of a tetanus containing vaccine.
- inflammation of several nerves causing sensory disorders or weakness of limbs (polyradiculoneuritis), facial paralysis, visual disturbances, sudden dimming or loss of vision (optic neuritis), inflammatory disease of brain and spinal cord (central nervous system demyelination, multiple sclerosis) have been reported after administration of a hepatitis B antigen containing vaccine.
- swelling or inflammation of the brain (encephalopathy/encephalitis).
- in babies born very prematurely (at or before 28 weeks of gestation) longer gaps than normal between breaths may occur for 2 -3 days after vaccination.
- swelling of one or both feet and lower limbs which may occur along with bluish discolouration of the skin, redness, small areas of bleeding under the skin and

severe crying following vaccination with Haemophilus influenzae type b containing vaccines. If this reaction occurs, it is mainly after first injections and within the first few hours following vaccination. All symptoms should disappear completely within 24 hours without the need for treatment.

If your child gets any side effects, talk to your doctor, pharmacist or nurse. This includes any possible side effects not listed in this leaflet.

Reporting side effects

After you have received medical advice for any side effects your child experiences, you can report side effects to the Therapeutic Goods Administration online at www.tga.gov.au/reporting-problems in Australia or to the Medsafe online at <https://pophealth.my.site.com/carmreportnz/s/> in New Zealand. By reporting side effects, you can help provide more information on the safety of this medicine.

Always make sure you speak to your doctor or pharmacist before you decide to stop taking any of your medicines.

7. Product details

This medicine is only available with a doctor's prescription.

What Hexaxim contains

Active ingredients (main ingredients)	Each 0.5 ml dose of Hexaxim contains: At least 20 IU (30 Lf) of diphtheria toxoid At least 40 IU (10 Lf) of tetanus toxoid 25 micrograms of pertussis toxoid and 25 micrograms of pertussis filamentous haemagglutinin 10 micrograms of hepatitis B surface antigen 29 D antigen Units of poliovirus Type 1, 7 D antigen Units of poliovirus Type 2, 26 D antigen Units of poliovirus Type 3* 12 micrograms of Haemophilus type B polysaccharide conjugated to 22- 36 micrograms of tetanus protein *These quantities are the same as those previously expressed when measured by a suitable method.
Other ingredients (inactive ingredients)	dibasic sodium phosphate mono basic potassium phosphate trometamol sucrose

	essential amino acids (cystine, tyrosine, arginine hydrochloride, histidine, isoleucine, leucine, lysine hydrochloride, methionine, phenylalanine, threonine, tryptophan and valine) water for injections
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Do not take this medicine if you are allergic to any of these ingredients.

The vaccine may contain traces of glutaral, formaldehyde, neomycin, streptomycin and polymyxin B. The manufacture of this product includes exposure to bovine materials. No evidence exists that any case of vCJD (considered to be the human form of bovine spongiform encephalopathy) has resulted from the administration of any vaccine product.

What Hexaxim looks like

Hexaxim is provided as a fully liquid suspension for injection in pre-filled syringe (0.5 mL).

Hexaxim is available in a pack containing 1 pre-filled syringe with 1 or 2 needles.

Hexaxim is available in a pack containing 10 pre-filled syringes with 10 or 20 needles.

Not all pack sizes may be marketed.

After shaking, the normal appearance of the vaccine is a whitish cloudy suspension (Aust R 215536).

Who distributes Hexaxim

Distributed in Australia by:

sanofi-aventis australia Pty Ltd

12-24 Talavera Road

Macquarie Park NSW 2113

Freecall: 1800 818 806

Email: medinfo.australia@sanofi.com

Distributed in New Zealand by:

Pharmacy Retailing (NZ) Ltd t/a Healthcare Logistics

PO Box 62027

Sylvia Park Auckland 1644

Freecall: 0800 283 684

Email: medinfo.australia@sanofi.com

This leaflet was prepared in October 2025.

hexa-ccdsV12-14-cmiv5-07oct25