

# ADT<sup>TM</sup> Booster *(ay-dee-tee boo-ster)*

## Diphtheria and Tetanus Vaccine, Adsorbed

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### Consumer Medicine Information

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#### What is in this leaflet

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This leaflet answers some common questions about **ADT<sup>TM</sup> Booster**.

It does not contain all the available information.

It does not take the place of talking to your doctor or pharmacist.

All medicines, including vaccines, have risks and benefits. Your doctor has weighed the risks of you or your child (of five years or older) having **ADT<sup>TM</sup> Booster** against the benefits they expect it will have.

**If you have any concerns about this vaccine, talk to your doctor, nurse or pharmacist.**

**Keep this leaflet.**

You might need to read it again.

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#### What ADT<sup>TM</sup> Booster is used for

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**ADT<sup>TM</sup> Booster** is a “combination” vaccine. It helps prevent **two** diseases, each caused by a different infection. The diseases are

- diphtheria *and*
- tetanus.

Both of these infections are serious and can be life-threatening.

**ADT<sup>TM</sup> Booster** is used to vaccinate children ( $\geq 5$  years of age) and adults who have previously received at least three doses of a vaccine for primary immunisation against diphtheria and tetanus.

**ADT<sup>TM</sup> Booster** is not intended for primary immunisation against diphtheria and tetanus.

**ADT<sup>TM</sup> Booster** is given as one **additional** dose (**booster** dose) with intervals according to national recommendations.

#### **How ADT<sup>TM</sup> Booster works**

**ADT<sup>TM</sup> Booster** works by getting your body to produce its own protection against the two types of bacteria (germs). The germs are those that cause two different and serious infections

- diphtheria *and*
- tetanus.

The vaccine does not contain live germs and cannot give you these illnesses.

After you have **ADT<sup>TM</sup> Booster**, your body makes substances called antibodies. These antibodies fight both the diphtheria and the tetanus germs. When you come into contact with these germs, your body is usually ready to destroy them.

Most people who receive the booster dose (suitable only if in the past they have had the full primary course against diphtheria and tetanus) will produce enough antibodies to protect against both the diphtheria and tetanus diseases. However, as with all vaccines, 100% protection cannot be guaranteed.

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## **Before you are given ADT™ Booster**

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### ***When you or your child must NOT be given ADT™ Booster***

**Do not give ADT™ Booster to a child under 5 years of age.**  
ADT™ Booster is not suitable for children under 5 years of age.

**Do not use ADT™ Booster after the expiry date printed on the pack.**

**Do not use ADT™ Booster if the packaging is torn, shows signs of tampering, or does not look quite right.**

**If you are not sure whether you or your child should have ADT™ Booster, talk to your doctor or pharmacist.**

### ***Before you or your child are given ADT™ Booster***

**Tell your doctor if you or your child have allergies to:**

- **ADT™ Booster, Tetanus Vaccine, Diphtheria Vaccine** or any of the ingredients listed at the end of this leaflet
- any other medicines
- any other substances, such as foods, preservatives or dyes.

As for all vaccines, medical supervision and treatment should be available in case there is a severe allergic reaction.

**Tell your doctor if you or the person to be immunised are pregnant or intend to become pregnant.**

Your doctor will discuss the possible risks and benefits of having **ADT™ Booster** during pregnancy.

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## **How ADT™ Booster is given**

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**ADT™ Booster** is given by a trained health professional, as an injection into the muscle.

### ***How much is given and when***

For the **booster dose**, one dose of 0.5 mL is given.

**Ask your doctor or pharmacist to answer any questions you may have.**

### ***If you are given too much (overdose)***

Because each ADT™ Booster contains only one dose, overdosage is unlikely.

If you think you or anyone else may have been given too much of this medicine

- consult your doctor immediately or
- telephone the Poisons Information Centre (telephone 13 11 26 in Australia or 0800 POISON (0800 764 766) in New Zealand) for advice, or
- go to Accident and Emergency at your nearest hospital.

Do this even if there are no signs of discomfort or poisoning. Urgent medical attention may be required.

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## After having ADT™ Booster

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### ***Things you must do***

Keep an updated record of your vaccinations or your child's vaccinations.

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## Side effects

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**Tell your doctor or pharmacist as soon as possible if you or your child feel unwell after having ADT™ Booster.**

All medicines, including vaccines, can have side effects. **ADT™ Booster** may have unwanted side effects in some people. Sometimes they are serious, most of the time they are not. You or your child may need medical treatment if you get some of the side effects.

**Ask your doctor or pharmacist to answer any questions you have.**

**Tell your doctor or pharmacist if you notice any of the following and they worry you:**

- reaction at the injection site such as temporary redness, tenderness or swelling
- a small lump at the injection site; sometimes this may last for a few weeks
- fever, general malaise, eczema and inflammation of the skin.

### ***Allergic reaction:***

As with all vaccines given by injection, there is a very small risk of a severe allergic reaction.

**If any of the following happen, consult your doctor or pharmacist immediately, or go to Accident and Emergency at your nearest hospital.**

- sudden signs of allergy such as rash, itching or hives on the skin, swelling of the face, lips, tongue or other parts of the body
- shortness of breath
- collapse.

These are very serious side effects. If you or your child have them, you may have had a severe allergic reaction to **ADT™ Booster**. You or your child may need urgent medical attention or hospitalisation.

This type of side effect mostly occurs within the first few hours of being given the vaccine.

**Other side effects not listed above might occur in some people. Tell your doctor or pharmacist if you notice anything that is making you or your child feel unwell.**

**Do not be alarmed by this list of possible side effects.**

You or your child may not experience any of them.

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## Storing ADT™ Booster

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

- **Keep it where children cannot reach it.**
- **Keep it in the original pack until it is time for it to be injected.**
- **Keep it in the refrigerator, between 2°C and 8°C. DO NOT FREEZE ADT™ Booster.** Freezing destroys the vaccine.

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## Product description

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### ***What ADT™ Booster looks like***

ADT™ **Booster** is supplied as a single dose (0.5 mL) in a needle-less pre-filled glass syringe or vial. The vaccine should appear as white and grey particles suspended in a colourless fluid.

### ***Ingredients***

Active ingredients:

- Diphtheria Toxoid: at least 2 International Units (IU)
- Tetanus Toxoid: at least 20 IU.

Other ingredients:

- Aluminium hydroxide
- Sodium chloride
- Sodium hydroxide
- Water for injection.

### **ADT™ Booster does NOT contain:**

- lactose
- sucrose
- gluten
- tartrazine *or*
- any other azo dyes
- preservatives.

The manufacture of this product includes exposure to bovine derived materials. No evidence exists that any case of vCJD (considered to be the human form of bovine spongiform encephalitis) has resulted from the administration of any vaccine product.

### ***Manufacturer/Distributor/ Supplier***

#### ***Manufacturer***

**ADT™ Booster is made in Denmark by:**

AJ Vaccines A/S  
5, Artillerivej  
DK-2300 Copenhagen S  
Denmark

#### ***Distributor***

**ADT™ Booster is distributed in Australia by:**

Seqirus (Australia) Pty Ltd  
63 Poplar Road  
Parkville Victoria 3052  
Australia

**ADT™ Booster is distributed in New Zealand by:**

Seqirus (NZ) Ltd  
PO Box 62 590  
Greenlane,  
Auckland 1546  
New Zealand

Telephone: 0800 502 757

***Registration number***

Australia  
AUST R 130906  
AUST R 130919

***Date of preparation***

28 April 2017

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