Summary
Vaccines contain an active component (the antigen) which induces the immune response. They may also contain additional components such as preservatives, additives, adjuvants and traces of other components. This fact sheet provides information about vaccine components including why they are present, and what, if any, risks these components may pose to vaccine recipients.

The following commonly asked questions are answered below. More general information on the vaccine components is also available by following the links in ‘Further reading’.

• What are the individual components in vaccines and why are they present?
  1. Active components
  2. Adjuvants
  3. Diluents
  4. Stabilisers
  5. Preservatives
  6. Trace components
• Do allergies to vaccines or vaccine components occur?
• Which vaccines contain animal-derived products and are there any alternatives?
• Which vaccines have used human tissue sources in their production?

What are the individual components in vaccines and why are they present?

1. Active components
The active component of a vaccine is known as the vaccine ‘antigen’. This is a modified or partial form of the virus, bacteria or the toxin that causes the disease against which the vaccine protects. The vaccine antigen is altered from its original form so it no longer causes disease but it can produce an immune response. There are a number of ways this is achieved:

Attenuated live viruses
Natural or ‘wild type’ viruses cause disease by reproducing themselves many millions of times in the body’s cells. In some vaccines where live virus is used, the virus has been treated and weakened (attenuated) in such a way that, when it is introduced to the body in the form of a vaccine, it induces an immune response without causing severe disease. The advantage of live, attenuated vaccines is that one or two doses usually provide lifelong immunity. Examples of attenuated live viral vaccines are the varicella, rotavirus and measles-mumps-rubella (MMR) vaccines.

Inactivated viruses
Some viruses or parts of viruses in vaccines are killed (inactivated) with a chemical such as formaldehyde. The killed virus cannot possibly reproduce itself or cause disease. The advantage of vaccines produced in this way is that the body still recognises the virus and produces an immune response. Because no viral replication occurs, these vaccines can be given to people with weakened immunity. The only disadvantage of these types of vaccines is that, generally, several doses must be given to achieve long-term immunity, but persons with weakened immunity may not respond to even multiple doses. Examples of inactivated vaccines are the inactivated poliomyelitis, influenza and hepatitis A vaccines.

Use part of the virus or bacterium
The hepatitis B, *Haemophilus influenzae* type b (Hib), and human papillomavirus (HPV) vaccines are examples of vaccines where only part of the virus or bacterium is used. The part of the virus or bacterium required to ‘induce immunity’ is identified and separated from the part which causes disease symptoms. In the case of hepatitis B, the vaccine is composed of a protein that resides on the surface of the virus. In the case of the *Haemophilus influenzae* type b (Hib) vaccine, only the outer coat, or polysaccharide, is used, joined on (conjugated) to a protein so that the immune system responds to it. These vaccines can be administered to people with weakened immunity, although, if the person’s immune system is too weak, they may not develop a satisfactory immune response.
Use a toxin produced by the bacteria
Some vaccines are manufactured by chemically
inactivating specific bacterial toxins. The inactivated
toxin is then referred to as a toxoid and used to produce a
vaccine, for example, diphtheria and tetanus-containing
vaccines. In the case of tetanus infection, exposure to very
little tetanus toxin is sufficient to cause disease, whereas
only a small amount of the tetanus toxoid in the vaccine
will induce a good immune response and cannot cause
disease. Having tetanus infection does not induce a long-
term immune response and non-immune individuals who
contract tetanus must be fully vaccinated to protect
against future exposure. The only way to be protected
against tetanus and diphtheria is to be vaccinated using
several doses of the appropriate vaccine.

2. Adjuvants
Adjuvants are used to enhance the immune response to a
vaccine. They include various aluminium salts such as
aluminium hydroxide, aluminium phosphate and
potassium aluminium sulphate (alum). One way adjuvants
are thought to improve the immune response is by
keeping the antigen(s) near the injection site so that they
can be readily accessed by cells of the immune system.
The use of aluminium adjuvants in vaccines generally
means that less antigen per dose of vaccine is required,
and, in some cases, fewer vaccine doses are needed. The
presence of adjuvants in vaccines can often be associated
with the local reactions that occur at the injection site
after vaccination.

Aluminium salts, in small amounts, have been added to
certain vaccines for about 60 years and a recent review of
all the available studies of aluminium-containing
diphtheria, tetanus and pertussis vaccines (either alone or
in combination) found that there was no evidence that
aluminium salts in vaccines cause any serious or long-
term adverse events. The exposure to aluminium from
vaccines is far less than that received from diet or
medications, such as some antacids. Although aluminium-
containing vaccines have been associated with local
reactions and, less often, with the development of
subcutaneous nodules at the injection site, other studies
have reported fewer reactions with aluminium-containing
vaccines than those without aluminium.

3. Diluents
A diluent is a liquid provided separately and used to dilute
a vaccine to the proper concentration prior to
administration. This is usually sterile saline or sterile
water.

4. Stabilisers
Additives are used as stabilisers and help maintain a
vaccine’s effectiveness by keeping the antigen and other
vaccine components stable during storage. Stabilisers
prevent the vaccine components adhering to the side of
the vaccine vial. Examples of additives include lactose
and sucrose (both sugars), glycerine and monosodium
glutamate (both of which are amino acids or salts of
amino acids), and human or bovine (cow) serum albumin
(both proteins). Gelatin, which is partially hydrolysed
collagen, usually of bovine (cow) or porcine (pig) origin,
is added to some vaccines as a stabiliser. Some members
of the Islamic and Jewish faiths object to vaccination on
the basis that some vaccines contain porcine-derived
products. However, these concerns have been addressed
by religious scholars (see ‘Which vaccines contain
animal-derived products and are there any alternatives?’
below). An anaphylactic allergy to gelatin is a
contra-indication to vaccination with certain vaccines (see
‘Do allergies to vaccines or vaccine components occur?’
below).

5. Preservatives
Preservatives are used to prevent fungal and/or bacterial
contamination of vaccines, and are present in some but
not all vaccines. Originally, preservatives were introduced
to prevent bacterial contamination of multi-dose vials.
However, multi-dose vials are no longer used routinely in
Australia. The preservatives used include thiomersal,
phenoxyethanol and phenol. Thiomersal (also known as
thimerosal) is a mercury-containing compound that is
discussed in detail in the National Centre for
Immunisation Research and Surveillance (NCIRS)
Thiomersal fact sheet:
http://www.ncirs.edu.au/immunisation/fact-
sheets/thiomersal-fact-sheet.pdf

Phenoxyethanol is an aromatic ether alcohol and is also
used as a preservative in many cosmetics. There has been
one case report suggesting that this preservative may be
associated with eczema. However, this link has not been
supported in other studies. Phenol is an aromatic alcohol
used as a preservative in very few vaccines. Preservatives
have been used in many vaccines and worldwide there
have been very few serious adverse events associated with
the use of these preservatives.

6. Trace components
Trace components are the remaining minute quantities of
substances that have been used in the early stages of the
production process of individual vaccines. Depending on
the manufacturing process used this may include trace amounts of cell culture fluids, egg proteins, yeast, antibiotics or inactivating agents. Usually, only minute traces of these substances are detected in the final vaccine product.

Antibiotics are sometimes used during the manufacturing process to ensure that bacterial contamination does not occur during the manufacturing process. Neomycin and/or polymyxin B are used in the manufacture of vaccines such as varicella (chickenpox) vaccines, some influenza vaccines, DTPa-combination vaccines and measles-mumps-rubella vaccine. Gentamicin is used in the manufacture of some influenza vaccines. No β-lactam or cephalosporin antibiotics are used in the manufacture of any vaccines currently used in Australia. Any individual with a severe allergy to any antibiotic or chemical who presents for vaccination should be appropriately assessed by the immunisation provider. The product information relating to each vaccine must be scrutinised for specific vaccine components before administering any vaccine to these individuals.

Inactivating agents are used during the manufacture of killed and toxoid vaccines. The bacteria, virus or toxin is inactivated during the manufacturing process but the antigenic components remain intact. The residual amount of these inactivating agents, for example formaldehyde or glutaraldehyde, in the final vaccine is very small.

Certain vaccines, such as influenza vaccines, may contain traces of egg proteins as the virus to be used for the vaccine is grown in actual chicken eggs before it is inactivated. Measles and mumps (but not rubella or varicella) vaccine viruses are grown in chick embryo tissue cultures and it is now recognised that MMR (and MMRV) vaccines contain negligible amounts of egg protein and can be safely given to children with egg allergy, even anaphylactic egg allergy. Other vaccines, such as the hepatitis B vaccines, hepatitis B-combination vaccines and human papillomavirus (HPV) vaccines, are manufactured using yeast. Production steps such as filtering and centrifugation greatly reduce the amounts of all of these products in the final vaccine; however, trace amounts may still be present.

**Do allergies to vaccines or vaccine components occur?**

Vaccines rarely produce allergy or anaphylaxis (a rapid and serious form of allergic reaction). Overall, the total risk of anaphylaxis in children and adolescents after one vaccination has been reported as <1 case per one million doses. Antibiotics, gelatin and egg proteins are the components most often implicated in these allergic reactions. Yeast has only rarely been associated with vaccine-related allergic reaction. In addition, people allergic to latex are potentially at risk, not from the vaccine itself but the presence of latex in the equipment used to hold the vaccine such as vaccine vial stoppers (bungs) and syringe plungers. Very few vaccine bungs contain natural latex. The product information sheet should be consulted to check for the presence of latex.

It is important that immunisation providers assess each individual for a history of allergies and previous reactions to vaccines prior to giving any dose of vaccine. However, depending on the allergy identified, there may often NOT be a contraindication to vaccination. For example, a history of an allergy to antibiotics most commonly relates to β-lactam or related antibiotics, and is not a contraindication to vaccines that contain neomycin, polymyxin B or gentamicin (see ‘Trace components’ above). Previous reactions to neomycin that just involved the skin are not considered a risk factor for anaphylaxis to vaccines manufactured with neomycin since there are only trace amounts of this antibiotic in the final product. Similarly, the measles and mumps components of MMR vaccine do not contain sufficient amounts of egg ovalbumin to contraindicate MMR vaccination of people with egg allergy (even anaphylaxis).

Where necessary, further advice should be sought from a medical practitioner with expertise in vaccination, the immunisation section within your State or Territory health authority, or your local Public Health Unit (see The Australian Immunisation Handbook Appendix 1, Contact details for Australian, State and Territory Government health authorities and communicable disease control). In addition, information about specialist immunisation clinics is available through state-based contacts listed in the most current edition of The Australian Immunisation Handbook or on the Immunise Australia Program website: [http://www.immunise.health.gov.au](http://www.immunise.health.gov.au). Information on how to access these clinics can be provided by your local health authority. Immunisation providers are trained to provide treatment in the rare event that a severe allergic response occurs immediately following a vaccine dose.

**Which vaccines contain animal-derived products and are there any alternatives?**

Community concerns around animal products in vaccines generally fall into two categories:
1) religious or faith-based concerns about the use of animal-derived products, and 2) concern about the possibility of animal diseases ‘crossing over’ to humans through use of vaccines.

1) Some vaccines utilise porcine (pork) products in the manufacturing process. This concern has been raised by some religious groups that have faith-based concerns about the consumption of pork. Scholars of these religions have various exceptions or rulings which allow the ‘ingestion’ of porcine or porcine-derived products in this context:

- For Muslims: Shariah law includes the principle of transformation in which unclean products can be made clean by extensive processing, thus making it permissible for observant Muslims to receive vaccines, even if the vaccines contain porcine gelatin (see ‘Stabilisers’ above).
- Judaism permits the use of non-edible forms of porcine products.
- Seventh-Day Adventists are not forbidden to use pork-derived medical products.

2) Bovine serum albumin or fetal calf serum is used in some vaccines and there were theoretical concerns that vaccines could be contaminated with variant Creutzfeldt-Jakob disease (vCJD). These theoretical concerns arose because the United Kingdom has documented rare cases of vCJD in humans following the ingestion of products from animals infected with bovine spongiform encephalopathy (BSE, also known as ‘mad cow disease’).

This question about vaccines and the use of bovine serum albumin has been addressed by the Australian Therapeutic Goods Administration (TGA), which requires that vaccine manufacturers adhere to strict standards and provide detailed information on the source of all materials used in the manufacturing process. These requirements include control on the source country of the animals used, the nature of the tissue used, and details of the manufacturing processes. In Australia, no case of vCJD from vaccines has been demonstrated and the TGA (after reports to the NHMRC Special Expert Committee on Transmissible Spongiform Encephalopathies) concluded that the vaccines used in Australia meet high safety standards and that any risk of transmission is theoretical only.

Internationally, advisory bodies also consistently state that the potential risk to vaccine recipients is essentially non-existent.

Which vaccines have used human tissue sources in their production?

Certain viruses grown for use in vaccines require the use of ‘cell lines’. These cell lines (called human diploid cell lines – WI-38 and MRC-5) were originally derived from human fetal tissue. The vaccines manufactured using cell lines originally derived from fetal tissue include: rubella vaccine and MMR vaccine, hepatitis A vaccines, varicella vaccines, rabies vaccine, and oral polio (Sabin) vaccine (no longer available in Australia).

Further reading


Additional web-based Information (accessed May 2013)

Australian Government Department of Health and Ageing, Therapeutic Goods Administration. BSE risk associated with the use of materials of bovine origin during the manufacture of vaccines. 2007. Available at:

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