

Immunisation

How vaccines work

Dr Fiona Ryan
Consultant in Public Health Medicine,
Department of Public Health
April 2015

Presentation Outline

- An understanding of the following principles
- Overview of immunity
- Different types of vaccines and vaccine contents
- Vaccine failures
- Time intervals between vaccine doses
- Vaccine overload
- Adverse reactions
- Herd immunity

Immunity

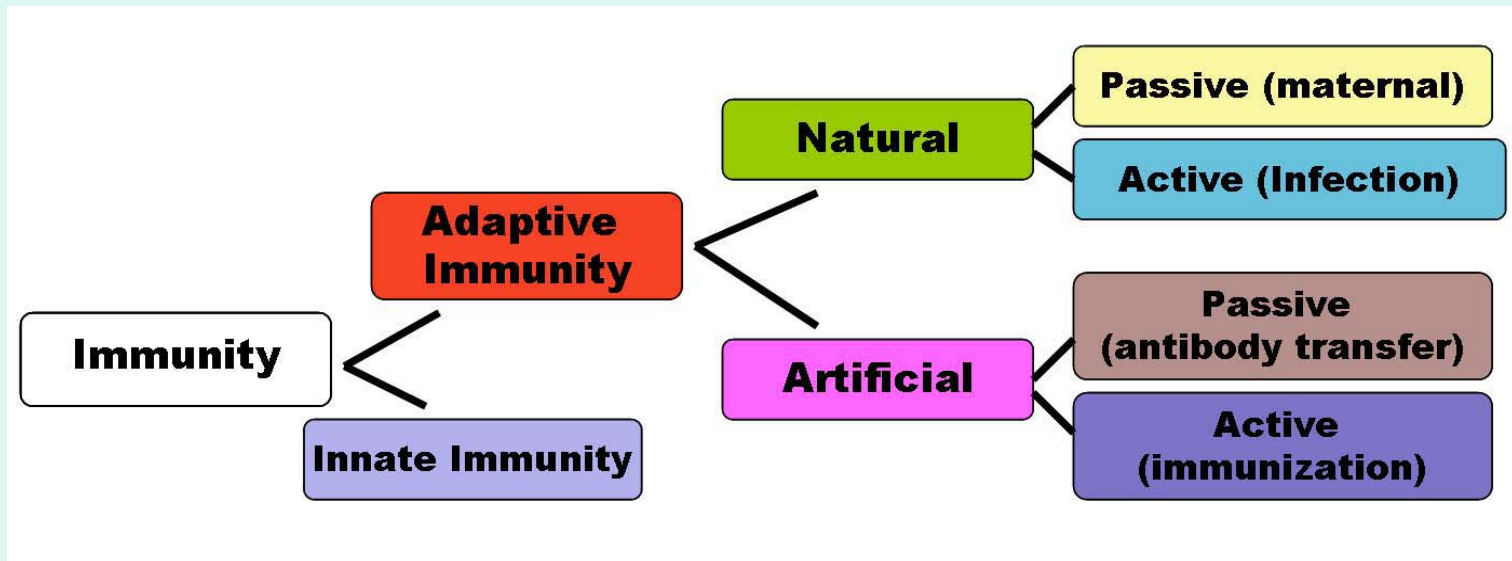
Immunity

- – The ability of the human body to protect itself from infectious disease

The immune system

- Cells with a protective function in the
 - bone marrow
 - thymus
 - lymphatic system of ducts and nodes
 - spleen
 - blood

Types of immunity

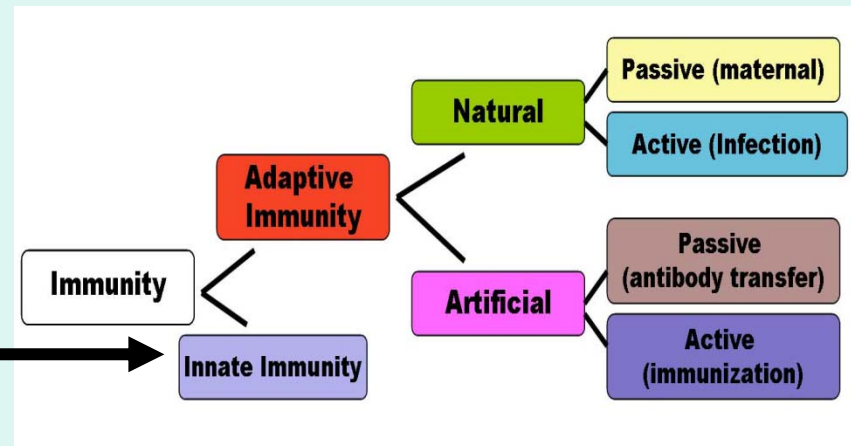


Source: http://en.wikipedia.org/wiki/Immunological_memory

Natural (innate) immunity

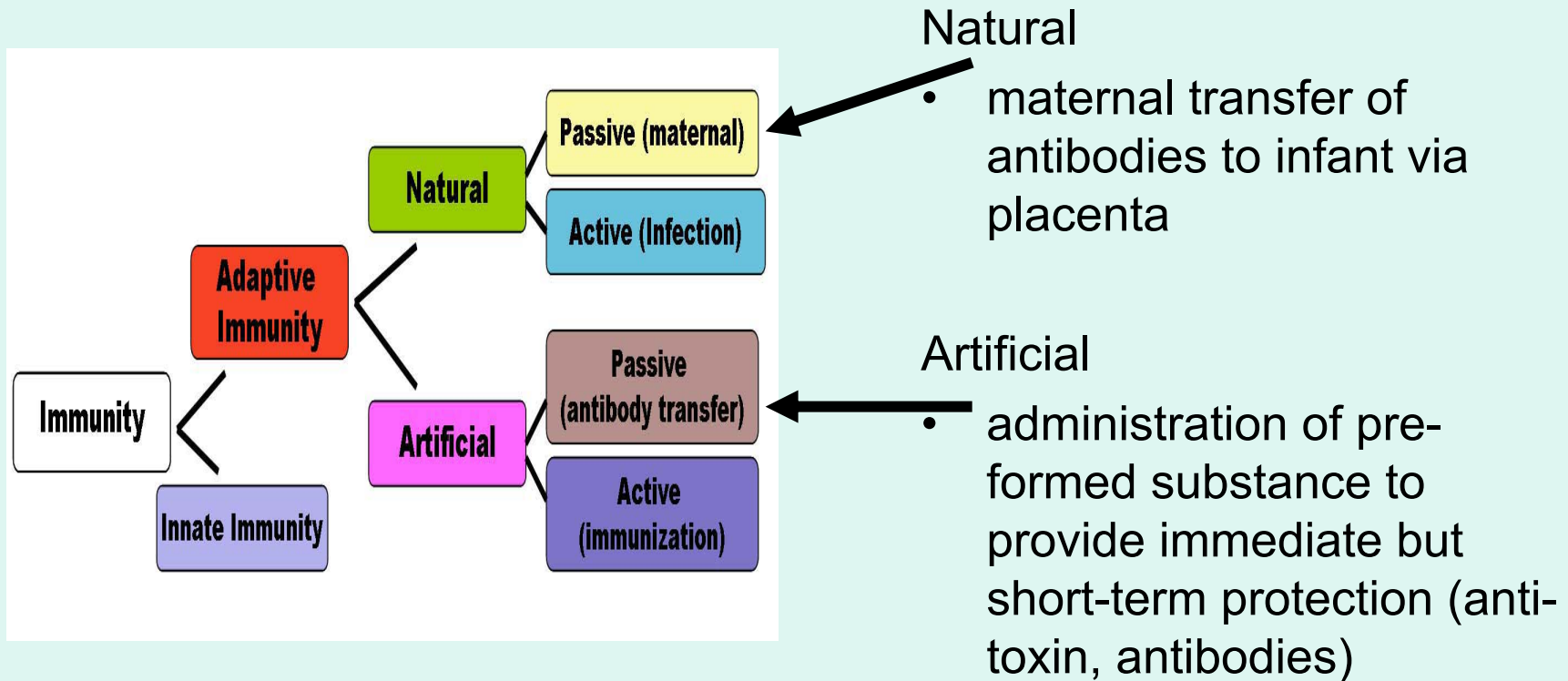
Non-specific mechanisms

- Physical barriers
 - skin and mucous membranes
- Chemical barriers
 - gastric and digestive enzymes
- Cellular and protein secretions
 - phagocytes, macrophages, complement system



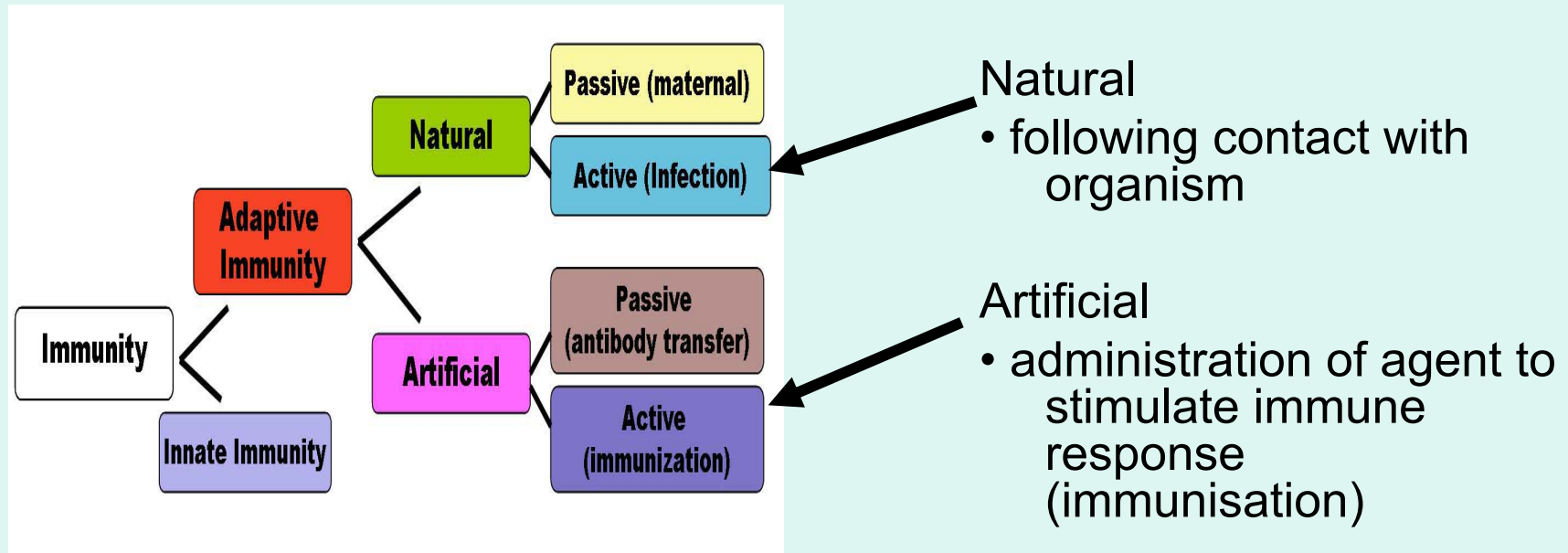
**** No “memory” of protection exists afterwards ****

Passive immunity – adaptive mechanisms



Protection is temporary and wanes with time (usually few months)

Active immunity – adaptive mechanisms



Acquired through contact with an micro-organism
Protection produced by individual's own immune system
Protection often life-long but may need boosting

How vaccines work

- Induce active immunity
 - Immunity and immunologic memory similar to natural infection but without risk of disease
- Immunological memory allows
 - Rapid recognition and response to pathogen
 - Prevent or modify effect of disease

Live attenuated vaccines

Weakened viruses /bacteria

- Achieved by growing numerous generations in laboratory
- Produces long lasting immune response after one or two doses
- Stimulates immune system to react as it does to natural infection
- Can cause mild form of the disease (e.g. mini measles which is non transmissible)
- CANNOT be given to immuno-compromised persons

e.g. BCG/ MMR/ Varicella/ Yellow fever

Inactivated vaccine and toxoids

- Cannot cause disease they are designed to protect against
- Doses
 - Two or more doses plus booster doses usually required
- Inactivated
 - e.g. pertussis, Hib, Pneumococcal, Men C, influenza
- Toxoids
 - e.g. tetanus, diphtheria

Vaccine Components

- Conjugating agents
 - – Carrier proteins which combine with antigens to improve immunogenicity
 - E.g. Men C, PCV, Hib
- Suspension fluid
 - Fluid (water, saline, tissue-culture mixture)
- Preservatives, stabilisers, antimicrobial agents
 - Trace amounts used to stabilise vaccine
 - May cause allergic reaction

Vaccine Components

- Adjuvants
 - Aluminium salt used to increase immunogenicity of vaccines containing inactivated micro-organisms or their products
- e.g.
 - Hepatitis B vaccine
 - Tetanus toxoid
 - Diphtheria toxoid

Vaccine Failure

- Primary
 - Inadequate immune response to vaccine (e.g.MMR1)
 - Infection possible any time post vaccination
- Secondary
 - Adequate antibody response immediately after vaccination
 - Level of antibodies decrease with time
 - Booster doses usually required
 - Feature of many inactivated vaccines

Time intervals between vaccine doses

Antigen combination	Recommended minimal interval between doses
≥2 killed antigens	No minimum interval
Killed and live antigens	No minimum interval
≥2 live antigens	Four-week minimum interval if not administered simultaneously

Time intervals between vaccine doses

- 2 Live vaccines - Minimum one month interval required
 - Allows **each immune response to develop**
 - Diminishes immune interference
- Interval between doses
 - Allows the next response to be a true secondary response – i.e. faster and bigger and with higher affinity IgG

Time Interval between vaccine doses

- Exceptions include
 - Yellow fever and MMR
 - Ideally separate by 1 month as may be suboptimal response to both if given together.
 - Rotavirus no interval needed
 - Anytime before or after other live vaccine
 - Influenza and PCV vaccine in those aged 12-23 months separate by at least 1 week due to slight increased risk of fever if given together

Can vaccines overload the immune system?

- The human body is composed of 10 trillion cells and contains 100 trillion bacteria
- On average there are
 - 1000 bacteria on each cm² of skin
 - 1,000,000 bacteria on each cm² of the scalp
 - 100,000,000 bacteria per gram of saliva
- The maximum number of antigens in a vaccine was ~3000 (DTwP, most from wP)
- With the new vaccines this number is much lower still
- No evidence that vaccines can overload the immune system

Adverse Events

Live vaccines:

- Frequency of adverse events falls with number of doses
- If antibody is made -> neutralises small amount of vaccine virus in any subsequent vaccine dose
- E.g. MMR

Inactivated vaccines

- Frequency of adverse events increases with number of doses
- Good antibody levels -> good secondary immune response
- May be inflammatory (i.e. produce a sore arm)
- E.g. tetanus, pertussis

Herd Immunity

Only applies to diseases which are passed from person to person

- For each disease
 - a certain level of immunity in the population which protects the whole population because the disease stops spreading in the community
- Provides indirect protection of unvaccinated as well as vaccinated individuals.
- May be the most important aspect of how vaccines work
 - MMR given to infants protects pregnant women from rubella.
 - Can eradicate disease even if some people remain susceptible

Vaccination: It works



REDPEN/
BLACK PEN

Hey guys - I don't even feel any rain. Why are we doing this again? Just put down the stupid umbrellas - they're bad for your arms anyway.