Childhood Immunization

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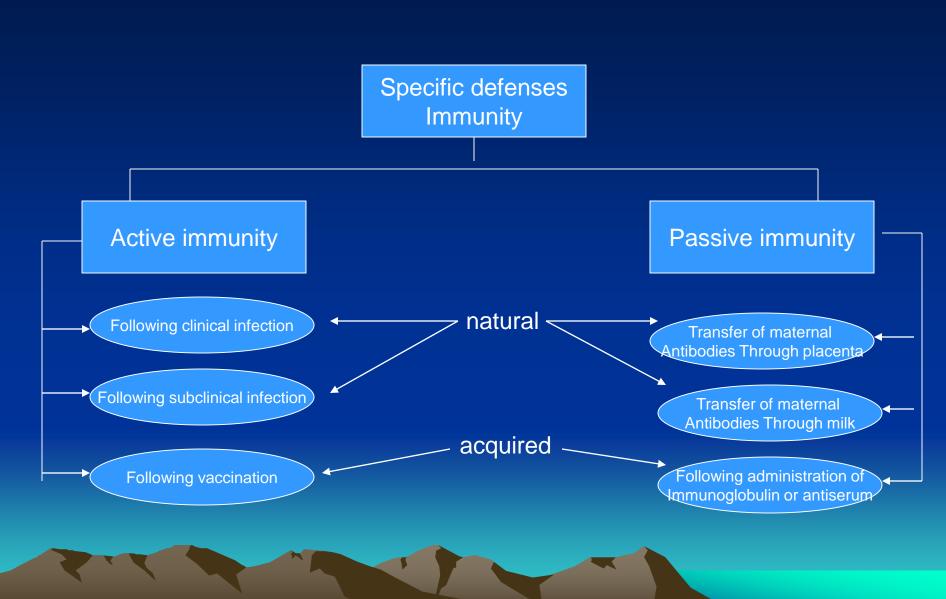
Childhood Immunization

- Definitions & General Concepts
- Safety of Immunization
- Standards for immunization practices.
- Cold chain and hazards of immunization

Definitions

- Vaccination : means having a vaccine that is actually getting the injection
- Immunization: The process of inducing immunity artificially. means both receiving a vaccine and becoming immune to a disease, as a result of being vaccinated

Immunity



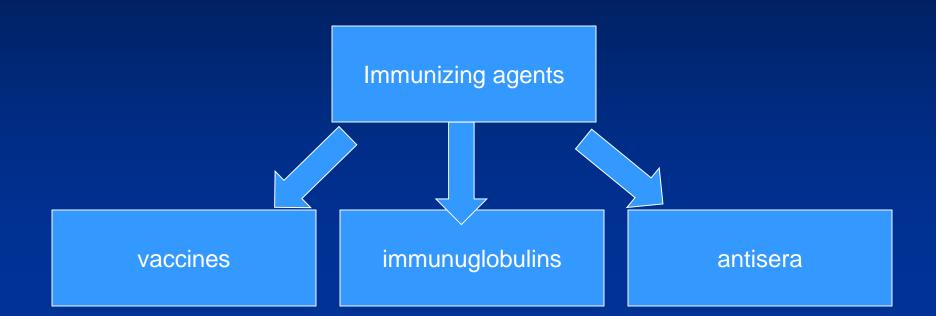
Active immunity

 Resistance developed in response to stimulus by an antigen (infecting agent or vaccine) and is characterized by the production of antibodies & cellular elements by the host.

Passive immunity

 Immunity conferred by an antibody produced in another host. It may be acquired naturally (Transplacental transmission) or artificially (through an antibody-containing preparation).

Immunizing agents



Immunizing Agents:

 Vaccine: A preparation of proteins, polysaccharides, or nucleic acids that are delivered to the immune system as single entities, part of a complex, or by live attenuated agents to induce specific responses that inactivate, destroy or suppress the pathogen.

Immunizing Agents

 Toxoid : A modified bacterial toxin that has been made non-toxic but retains the capacity to stimulate the formation of antitoxin like DT vaccines.

Antitoxin : An antibody derived from the serum of humans or animals after stimulation with specific antigens ,used to provide passive immunity like Diphtheria ,Tetanus

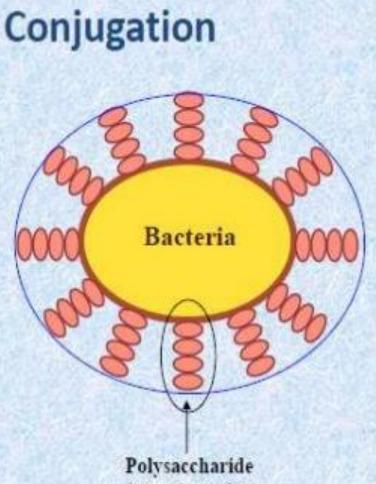
Immunoglobulins

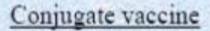
- There are 5 major classes: IgM, IgA, IgG, IgE, IgD.
- Two types of human immunoglobulin preparations are available for passive immunization:
 - Normal human immunoglobulin e.g. hepatitis A , measles Ig (IM) , and Botulism (IV)
 - Specific (hyper-immune) human Ig i e.g. hepatitis B and Varicella

What is the other use of Igs?

Types of vaccines

- Attenuated live vaccines
- Inactivated (killed vaccines)
- Toxoids
- Polysaccharide and polypeptide (cellular fraction) vaccines
- Surface antigen (recombinant) vaccines.
- Conjugate vaccines







Polysaccharide linked to carrier protein

(sugar) coating

Conjugation is the process of attaching (linking) the polysaccharide antigen to a protein carrier (e.g. diphtheria or tetanus) that the infant's immune system already recognises in order to provoke an immune response

Types of vaccines

 BCG Hepatitis A Typhoid Typhoid Typhoid Cholera Plague Pertussis Oral polio Plague 	Live Attenuated vaccines	Killed Inactivated vaccines	Toxoids	Cellular fraction vaccines	Recombina nt vaccines	Conjugate vaccines
 Yellow Yellow Salk polio Intra- Measles Mumps Influenza Varicella Intranasal Influenza 	 Typhoid oral Plague Oral polio Rota v Yellow fever Measles Mumps Rubella Varicella Intranasal 	 Typhoid Cholera Pertussis Plague Rabies Salk polio Intra- muscular influenza Japanise 	-	 polysaccharide vaccine Pneumococcal polysaccharide vaccine Hepatitis B polypeptide 		Influenza b, pneumococcal, meningococcal vaccines

Safety of Immunization:

Vaccine administration:

- Disposable syringes vs.reusable glass syringes.
- ✤70% alcohol solution
- All vaccines containing an adjuvant should be given IM to avoid granuloma or necrosis.
- Given in the antlat. Of thigh < 18 months, deltoid or triceps in older children
- Aqueous vaccines may be given IM, SQ, ID

Vaccine factors

It is safe to administer many combinations of vaccines simultaneously.

- Inactivated vaccines can be given together or at any time after different vaccines.
- Live-virus vaccines, if not on the same day, should be given at least 30 days apart.
- Ig does not interfere with killed vaccines.

Ig can interfere with the immune response to measles & varicella vaccines, it should be administered at least 2 wk after measles vaccine.

Time intervals between vaccine doses

- 2 or more killed vaccines No minimum interval. Doses of same killed vaccine 4 weeks apart, 8 weeks for PCV
- Killed and live vaccines No minimum interval . 2 or more live vaccines 4 week minimum interval if not administered simultaneously
- Immunoglobulin and live vaccines 3 months or more

Host Factors:

Healthy Children:

Minor acute illnesses, with or without fever, are not contraindications to vaccination.

- Moderate to severe febrile illness may be a reason to postpone vaccination can be giving with precaution
- Routine P/E and Taking Temp are not necessary in healthy children.

Catch-up vaccination:

- vaccine series does not need to be restarted, regardless of the time that has elapsed
- Depend on type of vaccine, age and the level of acquired protection
- If level of protection is inadequate following HBV a series of 3 doses should be repeated

Children with Chronic Illnesses

- Most chronic diseases are not contraindications to vaccination
- Premature Infants should be immunized according to their chronological age ,not gestational age. (except birth dose of hepatitis B, if mother is HB -ve & infant < 2 kg at birth defer until 30 days of age)
- Vaccine doses should not be reduced for preterm or LBW infants.
- Pertussis v. can be giving with precaution in those with a progressive CNS disorder.

Immunodeficient Children

- Congenital Immunodeficient children: may not be vaccinated with live vaccines
- Other vaccines may be safe, yet they may fail to evoke a proper immune response.
- Children with cancer, on steroids ,or Immunosuppressive agents are not to receive live vaccines.
- HIV : MMR are recommended except in severe immunodeficiency ,OPV & BCG are not (? Rota v), other vaccinations in Libyan schedule should be giving.

Periods of maintained immunity due to vaccines

- Short period (months): cholera vaccine
- Three to five years: DPT vaccine
- Five or more years: BCG vaccine
- Ten years : yellow fever vaccine , hep. A
- Solid immunity: measles, mumps, rubella.

Levels of effectiveness

- Absolutely protective(100%): yellow fever vaccine
- Almost absolutely protective (99%): measles, mumps, rubella, hep. A, diphtheria and tetanus toxoids.
- Highly protective (80-95%): polio, BCG, Hepatitis B, and pertussis vaccines.
- Moderately protective (40-60%) cholera vaccine, and influenza killed vaccine.

previous Vaccination schedule of Libyan children

Visit No.	Age	Vaccines (dose)
First	At birth	BCG HBV polio
Second	At two months	 OPV DPT+ Hib v + HBV (penta)
Third	At four months	 OPV DPT+ Hib v + HBV (penta)
Fourth	At sixth months	 OPV DPT+ Hib v + HBV (penta)
Fifth	At 12 months	• MMR
Sixth	At 18 months	 DPT (Booster1) OPV (Booster1) MMR
Seventh	At school entry (6 years)	 DT (Booster2) OPV (Booster2) ACYW135 (meningococcal v)
eighth	12 years	•OPV
Ninths	15 years	·Td

The DPT + Hib + HBV vaccine is a penta vaccine and is given in one dose.

New Vaccination schedule of Libyan children

		-
Visit No.	Age	Vaccines (dose)
First	At birth	BCG HBV OPV
Second	At two months	IPV + HBV + DTaP+ Hib + Rota + PCV
Third	At four months	IPV + HBV + DTaP+ Hib + Rota + PCV
Fourth	At sixth months	IPV + HBV + DTaP+ Hib + Rota
Fifth	At nine months	OPV A,C,Y,W135 (meningococcal)
Sixth	At 12 months	MMR PCV A,C,Y,W135 (meningoc.)
Seventh	At 18 months	DTaP OPV MMR
Eighth	At school entry (6 years)	Td + OPV + meningococcal v
Ninths	15 years	Td + OPV+ HPV(3 doses)

The DTaP + Hib + HBV-JPV vaccine is a Hexa vaccine and is given in one dose. Haemophilus Influenza b, pneumococcal, and meningococcal vaccines are conjugated vaccines

Routes of administration

- Deep subcutaneous or intramuscular route (most vaccines)
- Oral route (sabine vaccine), Rota v
- Intradermal route (BCG vaccine)
- Intranasal route (live attenuated influenza vaccine)

Certain available vaccines and their routes of administration

Vaccine	Туре	Route
BCG	Live-attenuated mycobacterial	Intradermal
DTaP	D&T = Toxoids aP = detoxified components from <i>Bordetella</i> pertussis	Intramuscular
Hep. B (HBV) , hep A	Inactivated viral antigen	Intramuscular
Hib , peumococcus & meningiococcus	Polysaccharide	Intramuscular
MMR & varicella	Live attenuated viruses	Subcutaneous
OPV & Rota v	Live attenuated virus	Oral

BCG = Bacillus Calmette – Guerin vaccine (tuberculosis).
 DTap = Diphtheria, tetanus and acellular pertussis vaccine.
 MMR = Live measles, mumps and rubella viruses in a combined vaccine.
 OPV = Oral Poliovirus vaccines containing attenuated poliovirus types 1,2 and 3
 Hib = Haemophilus Influenza b

Contraindications to vaccinations:

•Absolute

Temporary

Contraindications to vaccines <u>Absolute</u>

- 1- general contraindication for all vaccines is anaphylactic reaction to a prior dose.
- 2-Contraindication of subsequent pertussis vaccines
- are: encephalopathy (e.g., coma, decreased level of
- consciousness, prolonged seizures) not attributable
- to another identifiable cause within 7 days of
- administration of a previous dose pertussis v.

Contraindications to vaccines <u>Temporary:</u>

- Severe illness that needs hospitalization, and usually deferred in children with moderate to severe acute illnesses
- 2. Some Immunosuppression disorder.
- 3. Recent receipt of blood or immunoglobulin.

4. Pregnancy.

The Cold Chain

- The "cold chain" is a system of storage and transport of vaccines at low temperature from the manufacturer to the actual vaccination site.
- The cold chain system is necessary because vaccine failure may occur due to failure to store and transport under strict temperature controls.

Cold Chain

\cap	Vaccine Manufacturer
X	Vaccine*
Y	Distributor
Ŷ	Vaccine*
P	Provider's Office
Q	Vaccine*
U	Patient
B	

Vaccines must be stored properly from the time they are manufactured until they are administered to your patients

> Manufacturer to distributor Distributor to office Office to Child

Cold Chain Equipment

The recommended equipment typically used for vaccine storage are :

cold rooms,
 refrigerators and
 freezers.

For transporting vaccines equipment such as

 cold boxes,
 vaccine carriers and
 international containers are commonly used.

Cold chain equipment "For vaccine storage"



refrigerators



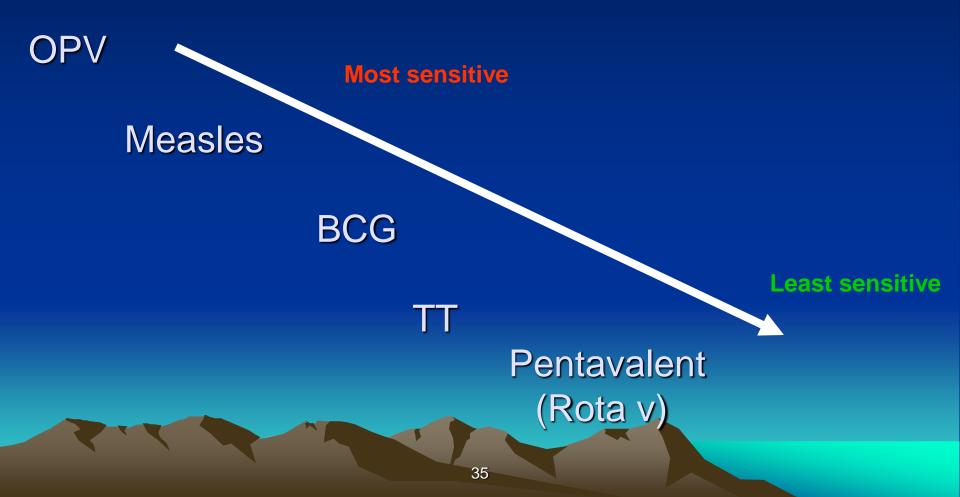
Cold room

Freezer

Cold chain equipment "for transporting vaccine"



Vaccines sensitive to heat



HAZARDS OF IMMUNIZATION

- No immune response is entirely free from the risk of adverse reactions or remote squeal. The adverse reactions include:
- 1. Reactions inherent to inoculation
- 2. Reactions due to faulty techniques
- 3. Reactions due to hypersensitivity

1. Reactions inherent to inoculation:

- local reactions: pain, swelling, redness, tenderness and development of a small nodule or sterile abscess at the site of injection.
- General reactions :fever, malaise, headache and other constitutional symptoms. Most killed bacterial vaccines (e.g., typhoid) cause some local and general reactions. Diphtheria and tetanus toxoids and live polio vaccine cause little reaction.

• 2. Reactions due to faulty techniques:

Faulty techniques may relate to

- faulty production of vaccine (e.g. inadequate inactivation of the microbe, inadequate detoxication),
- Unproper dose,
- improper immunization site or route,
- vaccine reconstituted with incorrect diluents,
- wrong amount of dilute used,
- vaccine prepared incorrectly for use (e.g., an adsorbed vaccine not shaken properly before use),
- vaccine or dliluent contaminated,
- vaccine stored incorrectly,
- contraindications ignored (e.g. a child who experienced a severe reaction (like anapmylaqxis) after a previous dose of vaccine is immunized with the same vaccine),
- reconstituted vaccine of one session of immunization used again at the subsequent session.

- 3. Reactions due to hypersensitivity:
- Administration of antisera (e.g., ATS) may occasionally give rise to anaphylactic shock and serum sickness.
- The symptoms may appear within a few minutes of injection or may be delayed up to 2 hours.

Immunization : WHY IT IS IMPORTANT ??

- Immunization is a remarkably successful & very cost effective means of preventing infectious diseases.
- A leading achievement of public health
- It is either to prevent primary infection or secondary consequences of infection



