# CHILDHOOD IMMUNIZATION: INTRODUCTION

# Prophylaxis

# Introduction

- Vaccination:
  - The act of giving a vaccine (antigen)
- Immunization:
  - Induction of an immune response following exposure to an antigen
- The ultimate goals are eradication and control of disease.
- The immediate goal is prevention of disease.

This is Ben. He is immunocompromised and cannot be vaccinated. But thanks to community immunity, he is protected from major diseases.



By vaccinating, you are not only protecting yourself and your children, but also people unable to be vaccinated.

# **Active vs Passive immunity**

ACTIVE	PASSIVE
Immunogenic antigen is given then the body forms its own protective antibodies.	<b>Ready-made immune globulin</b> (antibodies) from human or animal sources are given to the body.
Long term protection (Sometimes life long)	Temporary immunity that decreases with time (turnover of the administered immunoglobulin)
Examples: Natural: Infection	<b>Examples:</b> Natural: Mother's Ig to infant (transplacental/breast milk) effective for about 6 months.
Artificial: Vaccination	<b>Artificial</b> : Adminestration of antibodies (e.g: Hepatitis B IG, Varicella IG)

### Table 1.1. Comparison of 20th Century Annual Morbidityand Current Morbidity: Vaccine-Preventable Diseases<sup>a</sup>

	20th Century	2010 Reported	Percent
Disease	Annual Morbidity <sup>8</sup>	Cases	Decrease
Smallpox	29 005	0	100
Diphtheria	21 053	0	100
Measles	530 217	63	>99
Mumps	162 344	2612	98
Pertussis	200 752	27 550	86
Polio (paralytic)	16 316	0	100
Rubella	47 745	5	>99
Congenital rubella syndrome	152	0	100
Tetanus	580	26	96
Haemophilus influenzae	20 000	246 <sup>d</sup>	00

\*National Center for Immunization and Respiratory Diseases. Historical Comparisons of Vaccine-Preventable Disease

Morbidity in the U.S. Atlanta, GA: Centers for Disease Control and Prevention

<sup>b</sup>Roush SW, Murphy TV, Vaccine-Preventable Disease Table Working Group. Historical comparisons of morbidity and mortality for vaccine-preventable diseases in the United States. *JAMA*. 2007;298(18):2155-2163

<sup>e</sup>Centers for Disease Control and Prevention. Notice to readers: final 2010 reports of nationally notifiable infectious diseases. MMWR Morb Mortal Wkly Rep. 2011;60(32):1088-1101

<sup>d</sup>23 type b and 223 unknown serotype (<5 years of age).



# Smallpox

• Eradicated in 1980, case-fatality rate 30-50% according to type and age!

# Poliomyelitis!

- Rapid asymmetric acute flaccid paralysis (paralytic poliomyelitis) caused by poliovirus
- Proximal muscles > distal
- Areflexia.
- Cranial nerve (bulbar poliomyelitis)
- Paralysis of the diaphragm may lead to impaired respiration.



# Tetanus (Lockjaw)!

- Caused by Clostridium Tetanus that excretes Neurotoxin in a contaminated wound.
- Generalized tetanus (lockjaw): trismus and severe painful generalized muscular spasms.
- Autonomic dysfunction: diaphoresis, tachycardia, blood pressure, and arrhythmias.



# Pertussis (Whooping cough)

- Bordetella Pertussis
- Catarrhal stage, paroxysmal stage and convalescent stage (6 to 10 weeks).
- Complications: syncope, sleep disturbance, incontinence, rib fractures, pneumonia, conjunctival bleeding, hernia, hypoxia, seizures (2%), encephalopathy, and death.
- <6 months can be atypical: gasping, bradycardia, or apnea; absence of whoop.



# **Diphtheria!**

- Membranous nasopharyngitis or obstructive laryngotracheitis caused by diphtheria toxin.
- Extensive neck swelling with cervical lymphadenitis (bull neck) is a sign of severe disease.
- Complications:
  - upper airway obstruction;
  - myocarditis with heart block;
  - cranial and peripheral neuropathies.
- Case fatality rates up to 10%, sometimes > 20% in older adults.



Live attenuated	Inactivated vaccine
<b>Live organisms</b> >Lost ability to induce the disease >Retain capacity to grow but slowly and locally only	<b>Killed organisms</b> <ul> <li>(by heat/chemicals [formaldehyde]).</li> </ul>
Can produce antigens continuously: > More potent > 1 Dose is sufficient > Longer immunity	Cannot keep producing antigens: >Less potent >Need booster doses >Shorter immunity
Induce humoral & cellular immunity	Induce humoral immunity only > Antibody titers fall with time
<b>Risk of infection</b> Can mutate back to its virulent form	No risk of infection
<ul> <li>Not safe for immunocompromised people 1-immunocompromised individuals.</li> <li>2-People being treated for certain chronic illnesses (steroids used)</li> <li>3-Pregnant women unless absolutely necessary.</li> </ul>	Safe

### **Types Of Vaccines**



### **Recombinant Vaccines**

 They are made by inserting viral genes that code for important antigens into common baker's yeast. The yeast then produces the antigens, which are collected and purified for use in the vaccine

**Production of Recombinant HB Vaccine** 



# **Polysaccharide-based Vaccines**

#### • <u>Pure</u> polysaccharide vaccines:

- They induce a T-cell INDEPENDENT immune response. This means that they stimulate B cells without the need for T helper cells (humoral immunity only)
- This causes them to have weaknesses:
  - 1. Not immunogenic in children younger than 2 years of age (underdeveloped immune system. B cells must be activated through T cells)
  - 2. No booster response in adults (antibody titers don't increase)
- How was this problem solved?
  - By joining the polysaccharide molecule to a protein molecule and making a **conjugated** polysaccharide vaccines.
- This way the polysaccharide vaccine will stimulate a T cell DEPENDENT immune response.
- Conjugated vaccines:
  - 1. Induce an immune response in children younger than 2 years
  - 2. have a booster response.

Images of phagocytosis Streptococcus pneumoniae and capsule



*licrobiology: An Evolving Science,* Third Edition Figure 23.28d opyright © 2014 W. W. Norton & Company, Inc.

# **Types of vaccines**

- Live-attenuated: <u>BCG, MMR, OPV, Rota, Varicella</u>, (oral typhoid, yellow fever), Nasal Influenza virus vaccine.
- **Inactivated:** <u>DTaP</u> (toxoids and inactivated components) (Tdap, Td, DTP), IPV, Hib (polysaccharide conjugate), Hepatitis A (inactivated), Meningococcal, pneumococcal (polysaccharide conjugate or polysaccharide), **Influenza virus** (inactivated)
- Genetically engineered (recombinants antigens): Hepatitis B, HPV
- Live-attenuated vaccines are <u>contraindicated</u> in cases of cell-mediated immune defects and pregnancy.
- OPV is the only vaccine contraindicated when household contains an immunocompromised member.

### **Examples For Each Type Of Vaccines**



# Administration

• Most: IM (Ant-lat thigh or deltoid).

- Intradermal (ID): BCG (or SC)
- SC: MMR, Varicella, Polysaccharide vaccines, IPV (or IM)
- Intranasal: Nasal influenza vaccine
- Oral: OPV, Rota



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### **Site of Administration**

• You must avoid vascular, neural or tissue injury when injecting the vaccine.

- The preferred sites for IM administration are:
  - Anterolateral aspect of the thigh in infants
  - The deltoid region in children and adults .



# Administration

- Combined vaccines, concurrent administration
- Catch up vaccines.
- Interval for live-vaccines administration after: Chemotherapy, high dose steroids, blood products.
- **Recommended Minimum Interval** Antigen Combination Between Doses 2 or more inactivated<sup>a</sup> May be administered simultaneously or at any interval between doses Inactivated plus live May be administered simultaneously or at any interval between doses 2 or more live<sup>b</sup> 28-day minimum interval if not administered simultaneously

<sup>a</sup> See text for exceptions.

# **Immunization in Pregnancy**

- Influenza Vaccine Each Influenza Season.
- Tdap With Each Pregnancy.
- Pregnancy is generally a contraindication for live-virus vaccines
- Killed Vaccines not Indicated and not contraindicated.

# **Possible side effects to all vaccines**

- In general they occur early within 24-48 hours of vaccination and are self-limited.
- However, reactions following live vaccines (e.g. MMR) may be delayed and resemble a mild version of the disease.
- 1. Local reactions to injectable vaccines
- 2. Anaphylaxis to the vaccine or one of it's components (contraindications for further similar doses)
- 3. Syncope
- 4. Fever

# Not a contraindication!

- The followings are NOT a contraindications to vaccine administration:
- 1. Mild illness with or without fever
- 2. Breast feeding
- 3. Local rxns or fever after previous vaccine
- 4. Preterm birth
- 5. Penicillin allergy
- 6. Concurrent antibiotics use
- 7. Family history of seizure, controlled seizures

#### Vaccines included in the national immunization program (NIP)

- Viral (6): Measles, mumps, rubella (MMR); poliovirus (IPV, OPV) hepatitis A and B viruses, Rotavirus
- **Bacterial (5)**: Haemophilus influenzae type b (Hib), Tuberculosis (BCG), Diphtheria, Tetanus, Pertussis (DTP, DTaP, Td)
- Other vaccines in the American NIP: Tdap, varicella, meningococcal, pneumococcal, human papilloma virus, influenza virus.
- Other vaccines for travelers or exposure: Typhoid, yellow fever, rabies

#### برنامج التطعيم الوطنهي للأطفال (قبل سن دخول المدرسة)

المطعوم	العمر والجرعة
التـــدرن BCG	أقرب وقت بعد الولادة
	( أول مراجعة للمركز الصحي )
<ul> <li>مطعوم شلل الأطفال ١٩٧</li> </ul>	على عمر شهرين (61 يومر) يعطى الطفل الجرعة
<ul> <li>المطعوم الثلاق DaPT</li> </ul>	الأولى من:
(ضد الدفتيريا والسعال الديكي اللاخلوي والكزاز)	
<ul> <li>مطعوم المستدعية النزلية نوع (ب)</li> </ul>	
<ul> <li>مطعوم التهاب الكيد نوع + (ب)</li> </ul>	
<ul> <li>مطعوم الروتا فيروس</li> </ul>	
<ul> <li>مطعوم شلل الأطفال IPV</li> </ul>	على عمر 3 شهور (91 يوم )
<ul> <li>المطعوم الثلاق DaPT</li> </ul>	بعط ، الطفا ، الحرعة الثانية من:
(ضد الدفتيريا والسعال الديكى اللاخلوي والكراز)	0.0.0.0.0.0
<ul> <li>مطعوم المستدمية التزلية نوع (ب)</li> </ul>	
<ul> <li>مطعوم التهاب الكيد نوع + (ب) مطعوم الشلل القموى + (OPV)</li> </ul>	
<ul> <li>مطعوم الروتا فيروس</li> </ul>	
<ul> <li>مطعوم شلل الأطفال ١٩٧</li> </ul>	على عمر 4 شهور (121 يومر) يعطى الطفل.
<ul> <li>المطعوم الثلاق DaPT</li> </ul>	الجرعة الثالثة من:
(ضد الدفتيريا والسعال الديكي اللاخلوي والكزاز)	
<ul> <li>مطعوم المستدمية النزلية توع (ب)</li> </ul>	
<ul> <li>مطعوم التهاب الكبد نوع + (ب) مطعوم الشلل القموي + ( OPV)</li> </ul>	
<ul> <li>مطعوم الروتا فيروس</li> </ul>	
<ul> <li>مطعوم الحصبة</li> </ul>	علی عمر 9 شهور
<ul> <li>مطعوم الشلل الفموي OPV</li> </ul>	
.Vit. A 100000 I.U -	
<ul> <li>الجرعة الأولى من مطعوم الثلاق الفيروسي (MMR)</li> </ul>	عند بلوغ الطفل عامه الأول يعطى الطفل:
ضد الحصبة والحصبة الألمانية والنكاف	
<ul> <li>الجرعة المدعمة من مطعوم الشلل ومطعوم (DPT)</li> </ul>	على عمر 18 شهر يعطى الطفل؛
<ul> <li>الجرعة الثانية من مطعوم التلاثي الفيروسي (MMR)</li> </ul>	
.Vit. A 200000 LU -	

lotherhood & More		Vaccination Sc	طاعيم hedule	جدول الم		
The No.						
	الجرعة المدعمة	الجرعة الرابعة IV	الجرعة الثالثة III	الجرعة الثانية II	الجرعة الأولى I	اسم المطعوم
					CUEN/N	التدرن BCG
						شلل الأطفال المقتول IPV
	clop rein	114phs	C.Ke/1-11	C1/2/7		شلل الأطفال الفموي OPV
	C20/10/N					الثلاثي البكتيري DPT
			C.18/1.10	C18/9/7	Non/o	DPT IPV +Hib الخماسي المحسن
						DPT+BV +Hib الخماسي العادي
		1. 1. 1. 1. 1. 1. 1. 1. 1. 1. 1. 1. 1. 1	al. 132	CA8/9 4	aleNo	التهاب الكبد الوبائي HBV
						Hib المستدمية النزلية
					c. 10/11/1	الحصبية Measles فيتامين ((١٠٠ الف وحدة دولية)
				cidiely	0/1-610	الثلاثي الفيروسي MMR فيتامين أ(١٢٠٠ألف وحدة دولية)
						مطعوم الروتافيروس *
						Others li
					نعم 🗌 لا 🗌	هل تم أخذ عينة المسح الطبي للتحري عن الأمراض الوراثية

\* ملاحظة:عدد الجرعات يعتمد على الشركة المصنعة للمطعوم

#### Table 1 Recommended Child and Adolescent Immunization Schedule for ages 18 years or younger, United States, 2021

These recommendations must be read with the notes that follow. For those who fall behind or start late, provide catch-up vaccination at the earliest opportunity as indicated by the green bars. To determine minimum intervals between doses, see the catch-up schedule (Table 2). School entry and adolescent vaccine age groups are shaded in gray.

Vaccine	Birth	1 mo	2 mos	4 mos	6 mos	9 mos	12 mos	15 mos	18 mos	19–23 mos	2-3 yrs	4-6 yrs	7-10 yrs	11–12 yrs	13–15 yrs	16 yrs	17–18 yrs
Hepatitis B (HepB)	1ª dose	۹ 2 <sup>rd</sup> c	iose•		•		3 <sup>st</sup> dose										
Rotavirus (RV): RV1 (2-dose series), RV5 (3-dose series)			1ª dose	2 <sup>nd</sup> dose	See Notes												
Diphtheria, tetanus, acellular pertussis (DTaP <7 yrs)			1ª dose	2 <sup>rd</sup> dose	3 <sup>st</sup> dose			<b>∢</b> 4≏ d	oseÞ			5ª dose					
Haemophilus influenzae type b (Hib)			1ª dose	2 <sup>rd</sup> dose	See Notes		3 <sup>st</sup> or 4 See I	* dose									
Pneumococcal conjugate (PCV13)			1ª dose	2 <sup>rd</sup> dose	3 <sup>rd</sup> dose		<b>∢</b> 4 <sup>∞</sup> c	iose									
Inactivated poliovirus (IPV <18 yrs)			1 <sup>e</sup> dose	2 <sup>rd</sup> dose	•		- 3ª dose -					4 <sup>e,</sup> dose					
Influenza (IIV)							A	nnual vacci	nation 1 or	2 doses				Annua	vaccination	n 1 dose on	ly .
Influenza (LAIV4)											Annual 1 o	l vaccinatio r 2 doses		Annual	vaccination	1 dose on	ly .
Measles, mumps, rubella (MMR)					See N	lotes	<b>∢</b> 1° d	ioseÞ				2 <sup>nd</sup> dose					
Varicella (VAR)							<b>∢</b> 1= d	ioseÞ				2 <sup>rd</sup> dose					
Hepatitis A (HepA)					See N	lotes	3	2-dose serie	s, See Note	s							
Tetanus, diphtheria, acellular pertussis (Tdap ≥7 yrs)														Tdap			
Human papillomavirus (HPV)													•	See Notes			
Meningococcal (MenACWY-D ≥9 mos, MenACWY-CRM ≥2 mos, MenACWY-TT ≥2years)								See Notes						1ª dose		2 <sup>rd</sup> dose	
Meningococcal B															See Not	es	
Pneumococcal polysaccharide (PP5V23)														See Notes			

# Schedule of the NIP and UNRWA in Jordan

Time of vaccination	Vaccine (s)	Comments
Within the first month of life	BCG	Only 1 dose
2 months of age (60+ days)	(DTaP, IPV, Hib: الخماسي), HepB, RotaV	Not in NIP at this age: PCV 13
3 months (90+ days)	(DTaP, IPV, Hib), HepB, RotaV, OPV	
4 months (120+ days)	(DTaP, IPV, Hib), HepB, RotaV, OPV	Final doses of Hib, HepB and RotaV. Not in NIP at 6mo: FluV
9 months	Measles, OPV	Monovalent measles
12 months	MMR, HepA	Not in NIP : Varicella
18 months	MMR, OPV <sup>b</sup> , DTP <sup>b</sup> , , HepA	Final MMR. <sup>b</sup> indicates booster
6 years, first grade	OPV <sup>b</sup> , Td	Reduced diphtheria vaccine
10 <sup>th</sup> grade	Td	Not in NIP: At 11 y: HPV, MCV4

# Vaccines: BCG

- Bacillus Calmette–Guérin (the only bacterial live attenuated )
- Reduce disseminated and life-threatening manifestations of TB in young children (meningitis and miliary TB, 80% efficacy)
- Specific adverse events: generally not serious
  - 1%, localized abscess and lymphadenopathy.
  - Osteitis, as long as several years after BCG.
  - Disseminated (2 per 1 million).
  - Anti-tuberculosis recommended for osteitis and disseminated.
- Live vaccine contraindications.

# Vaccines: HepB

#### • To whom?

- 1. Infants
- 2. Adults at high risk.
- 3. Special circumstances (infant born to a HepB +ve mother)
- For infants born to HBsAg + mothers: HepB vaccine and 0.5 mL of hepatitis B immune globulin (HBIG) within 12 hours of birth
- Efficacy 90-95%. Protection for 20 years or longer

# Vaccines: Hib

- Typable H. influenzae include 6 serotypes from A to F including H. influenzae B.
- Haemophilus influenzae type B (Hib) was the most common cause of <u>childhood bacterial meningitis</u>.
- It is given to individuals at increased risk for invasive Hib:
  - 1. Splenic dysfunction,
  - 2. immuncompromised
  - 3. younger than 5 years of age
- Efficacy 95-100%
- Adverse events and contraindications

Haemophilus influenzae infections



# Vaccines: IPV/OPV

- The IPV covers all the 3 serotypes of poliovirus while OPV covers serotypes 1 and 3.
- Efficacy 99-100%

#### IPV

- Type of vaccine?
- Inactivated/whole cell
- Route of administration?
- = IM
- To whom?
- Can be given to immunocompromised people
- Can be given to immunocompromised contacts of an infected/recently vaccinated person

#### OPV

- Type of vaccine?
- Live attenuated
- Route of administration?
- Oral

#### • To whom?

 Only to immunocompetent people and who don't have an immunocompromised household.

#### IPV

#### Advantages?

- No vaccine associated paralytic polio
- Can be given to immunocompromised patients
- Disadvantages?
- No mucosal immunity
- No contact immunity
- Harder to administer
- Costly

#### OPV

#### Advantages?

- Mucosal immunity
- Contact immunity
- Use in outbreaks
- Gives herd immunity
- Easy to administer
- Cheaper

#### Disadvantages?

 Vaccine associated paralytic polio

#### Contraindications:

- Immunodeficiency including antibody disorders,
- household of immunocompromised child.

# Vaccines: DTaP

#### • Type of vaccine and components?

- Inactivated/ Whole cell (DTP) (newer versions are inactivated acellular <u>DTaP</u>; currently in use; it is a combined purified antigens; fewer side effects, more prolonged immune response and can be given to adults)
- 2. Tetanus and diphtheria are toxoid vaccines.
- There are several different types of vaccines that can safely prevent diphtheria, tetanus, and pertussis:
  - 1. **DTaP** (diphtheria, tetanus, and acellular pertussis) vaccine, which is given to children <7 yrs
  - 2. **DT** (diphtheria and tetanus) vaccine, which is given to children
  - 3. **Tdap** (combined tetanus, diphtheria and acellular pertussis) vaccine, which is given to adolescents and adults (including pregnant)
  - 4. Td (tetanus and diphtheria) vaccine, which is given to adolescents and adults (including pregnant)
- Tetanus and Diphtheria booster every 10 years.
- Efficacy: 98-100% after 5 doses

# **Side effects of the DTP vaccination**

#### **OMild** (Common):

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•Fever, Redness, swelling, Soreness (1 in 4)
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-Fussiness, Tiredness or poor appetite and Vomiting (1 in 50).

These problems occur more often after the 4<sup>th</sup> and 5<sup>th</sup> doses of the DTP series than after earlier doses.

#### •Moderate Problems (Uncommon):

- Seizure (1 in 14,000),
- Non-stop crying for 3 hours or more (1 in 1,000)
- High fever (1in 16,000)

#### • Severe Problems (Very Rare) :

- -Serious allergic reaction (1 in a million dose)
- -Long-term seizures, coma, or lowered consciousness
- -Permanent brain damage.

# Vaccines: DTaP

#### Absolute Contraindications:

- 1. Encephalopathy (e.g., coma, decreased level of consciousness; prolonged seizures) within 7 days of previous dose (to pertussis vaccine)
- 2. Anaphylactic reaction to a previous dose.

#### <u>Relative</u> contraindications:

- 1. Progressive neurologic disorder (infantile spasms/ uncontrolled epilepsy/ progressive encephalopathy )
  - vaccine postponed until symptoms are controlled
- 2. Temperature of 40.5° C or higher within 48 hours after vaccination with a previous dose of DTP/DTaP
- 3. Collapse or shock-like state (i.e., hypotonic hyporesponsive episode) within 48 hours after receiving a previous dose of DTP/DTaP
- 4. Seizure within 3 days after receiving a previous dose of DTP/DTaP
- Persistent, inconsolable crying lasting 3 or more hours within 48 hours after receiving a previous dose of DTP/DTaP

# Vaccines: Rota V

- Live attenuated; given orally.
- Contraindications:
- Severe Combined Immune Deficiency.History of intussusception
- Precautions:
- Severe illness including gastroenteritis
- Other types of immunodeficiency
- Chronic GI illness



### **Latest Rotavirus Vaccines Recommendations**

Rotarix	RotaTeq
(RV1)	(RV5)
2	3
6 wks	6 wks
20 wks	12 wks
24 wks	32 wks

	ACIP Reco
	2009
Doses	
Min age	б wks
Max age- 1st dose	14 wks 6 days*
Max age- any dose	8 mon 0 days*

# Vaccines: MMR

- Efficacy after 1 dose 95%, 2 doses 99%
- Live vaccine contraindications
- Side effects:
- 1. Measles-like rash (rash and fever) within 7 days
- 2. Arthralgia or arthritis (rubella component) (7-21 days after the vaccine)
- 3. Mild form of mumps (swelling of the cheek and neck): 3-4 weeks after vaccine.
- 4. Febrile seizures
- 5. Orchitis, parotitis (mumps component)
- 6. Thrombocytopenia (measles component)

Vitamin A enhanced the antibody response to measles vaccine given at 9 months of age significantly. And decrease the severity of measles infection.



# Hepatitis A (HepA)

- Inactivated virus.
- Newly introduced in Jordanian National Program.
- Given IM × 2 doses (1<sup>st</sup> dose at 12 months with 6-month interval for the 2° dose).

# VACCINES NOT PART OF THE JORDANIAN VACCINATION PROGRAM

### **Pneumococcal vaccine**

#### There are 2 types of pneumococcal vaccines against Strep. pneumonia:

- A. Pneumococcal conjugated vaccine(PCV13): Given for children <2 years old and can be given for children older than 2 years?
  - 1. Cystic fibrosis or chronic lung disease.
  - 2. Cochlear implants
  - 3. Patients with Splenectomy / Asplenia
  - 4. Immunocompromised patients (increased risk of infection)
  - 5. Nephrotic syndrome
  - 6. leaks of cerebrospinal fluid
  - 7. Sickle cell disease
- B. Pneumococcal polysaccharide vaccine (PPV)/ Pneumovax: protects against 23 different types of pneumococcus bacteria. Given to older children and adults.

# **Meningococcal vaccine**

- What are the two types of meningococcal vaccine available in the US?:
- 1. Tetravalent polysaccharide vaccine (MPSV4): For ages 2 years and up
- 2. Tetravalent polysaccharide-protein <u>conjugate</u> vaccine (MCV4): Less than 2 years
- Both protect against serogroups <u>A, C, Y, and W-135</u> of Neisseria Meningitis
- There is new vaccine currently available against serogroup b, which is actually the most prevalent and lethal one.
- Whom is the vaccine given to?
- 1. Impaired immunity: Nephrotic syndrome (immunoglobulin loss), Splenectomy and Complement deficiencies.
- 2. Travelers to areas endemic or epidemic with meningococcal infection.

# Influenza vaccine

- Type of vaccine?
- Inactivated/Whole cell
- Each injected seasonal influenza vaccine contains three killed influenza viruses:
- 1. one influenza type A subtype H3N2 virus strain
- 2. one influenza type A subtype H1N1 (swine flu) virus strain
- 3. one influenza type B virus strain
- Vaccination before the start of influenza seasons annually is usually recommended for any individual above 6 months of age.
- Route of administration?
- $IM \rightarrow Killed$
- Nasal spray  $\rightarrow$  live attenuated

### Varicella vaccine

- Live attenuated
- First dose: 12- 15 months of age
- Second dose: up to 4 years of age

# Thanks