AZERBAIJAN MEDICAL UNIVERSITY
DEPARTMENT OF MEDICAL MICROBIOLOGY and IMMUNOLOGY

LESSON 14.

INTRODUCTION TO BASIC VIROLOGY. MICROBIOLOGY DIAGNOSIS OF ACUTE RESPIRATORY VIRAL INFECTIONS (FAMİLİES OF *ORTHOMYXOVİRİDAE* AND *PARAMYXOVİRİDAE*)

FACULTY: ***General Medicine***
SUBJECT: ***Medical microbiology - 2***

Discussed questions:

1. Introduction to specific virology
2. Collection of examination materials during various viral diseases.
3. Microbiological diagnostic methods of viral infections: express, virology, serology

a) Detection of the virus or its components from the examination material taken from the patient (express diagnostics - IFR, ELISA, RIM, PCR, etc.).

b) Virological method - cultivation of pathological material in various biological objects (bodies of laboratory animals, chicken embryos and tissue cultures) and subsequent indication and identification

• Virus indication methods (hemagglutination reaction (HAR), hemadsorption phenomenon, cytopathic effect (CPE), intracellular inclusions, "negative colonies", "color test", interference phenomenon, CFT)

• Virus identification methods (BNR, CFT, HALR, hemadsorption retardation reaction, PHAR, immunodiffusion reactions, PHAR, RIM, IFR, ELISA, immunoelectron microscopy)

c) Serological method - serodiagnosis of viral infections, taking double sera, establishing serological reactions (KBR, BNR, HALR, IFR, RIM, ELISA).

4. Viruses that cause acute respiratory infections
5. *Orthomyxoviridae* family. General features, classification.

- Influenza viruses. Virion structure, structural features, cultivation, resistance. Influenza virus antigens. Classification of influenza A viruses by neuraminidase and hemagglutinin, antigen variability, ecology. Pathogenesis of influenza. Complicating effects of bacterial flora, immunity, microbiological diagnostics. Principles of specific treatment and prevention (vaccines, immunoglobulin, interferon, chemicals)

6. *Paramyxoviridae* family. General features, classification. Structure and chemical composition of virion, cultivation. Hemolysis, hemagglutination and hemadsorption properties. Resistance.

- Parainfluenza viruses, their role in human pathology, features of immunity
- Mumps virus, cultivation. Pathogenicity features. Immunity. Specific prevention.
- Respiratory syncytial viruses, cultivation. Pathogenicity features. Immunity.
- Morbillivirus genus. Measles virus. Pathogenicity features. Immunity. Semi-acute sclerosing panencephalitis. Microbiological diagnosis, specific prophylaxis.

Purpose of the lesson:

- To acquaint students with the tasks of special virology, the collection of examination materials during viral diseases and the methods of laboratory diagnosis of these diseases. To acquaint them with the morpho-biological features of influenza, parainfluenza, respiratory syncytial virus, epidemic parotitis and measles viruses and to provide information about the methods of laboratory diagnosis of diseases caused by these viruses.

SPECIAL VIROLOGY

- ***Special virology*** – studies the morpho-biological characteristics of clinically important viral pathogens, the pathogenesis, diagnosis and treatment of the infections caused by them.
- Viruses are divided into the following groups according to their structural characteristics:
- Simple DNA viruses
- Complex DNA viruses
- Simple RNA viruses
- Complex RNA viruses

The study of some viruses is based not on their structure, but on their biological properties:

- Arboviruses
- Latent viral infections, etc.

Specimens Appropriate for Laboratory Diagnosis of Various Clinical Syndromes

Syndrome	Specimen
Respiratory	Nasal or throat swab; nasopharyngeal aspirate; sputum
Enteric	Feces
Genital	Genital swab, urine
Eye	Conjunctival (and/or corneal) swab
Skin	Vesicle fluid/swab/scraping; biopsy solid lesion
Central nervous system	Cerebrospinal fluid; feces (enteroviruses)
Generalized	Throat swab[a]; feces[a]; blood leukocytes[a]
Autopsy/biopsy	Relevant organ
Any	Blood for serology[b]

METHODS OF MICROBIOLOGICAL DIAGNOSIS OF VIRAL INFECTIONS

Rapid (express) methods

Direct detection of antigens or nucleic acid of the pathogen in clinical material obtained from the patient.

Virological method

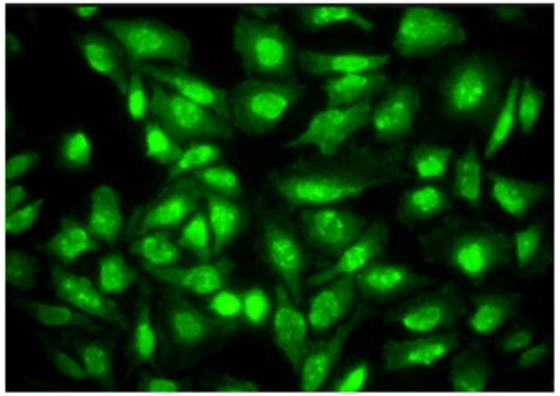
Isolation of virus from clinical material and its indication and identification.

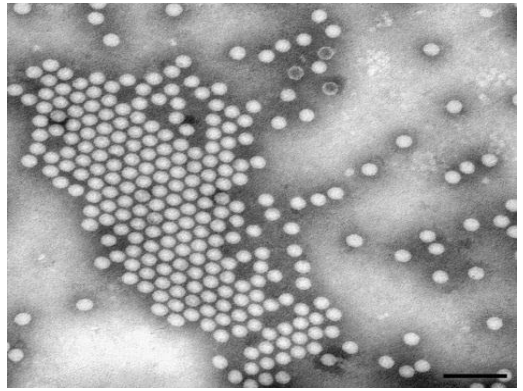
Serological method

Determination of the titer of antibodies against the virus in the paired sera of the patient.

RAPID (EXPRESS) METHODS

- **Rapid detection** is the identification of the virus and its antigens in bio substrates (biopsy samples, sediment epithelium, leukocytes, histological sections, etc.).
- **Rapid methods include:**
- **serological method -** determination of viral antigen using diagnostic antiviral sera: immunofluorescence, enzyme-linked immunosorbent assay (ELISA), radioimmunoassay, immunoelectrophoresis, immunoelectron microscopy, direct and indirect hemagglutination reaction, reversed indirect hemagglutination inhibition (RIHAI) test;
- **microscopic method -** detection of inclusions formed by viruses in cells using light, luminescent or electron microscopy:
- **molecular-genetic method -** molecular hybridization, PCR





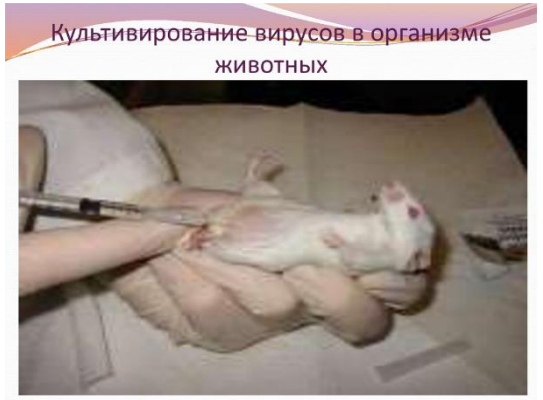
VIROLOGICAL METHOD

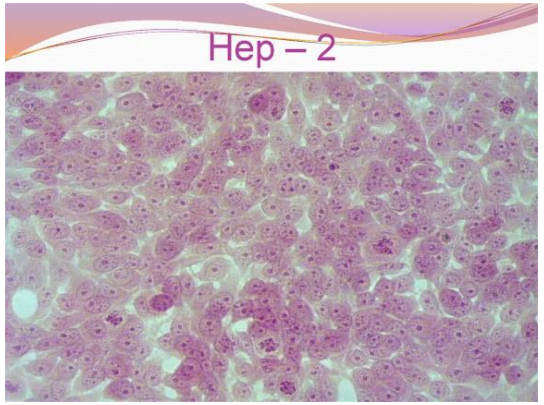
The virological method is based on the cultivation of viruses in sensitive biological objects (cell cultures, chicken embryos, laboratory animals).

It consists of the following stages:

- *Collection of pathological material*
- *Selection of a sensitive test system.*
- *Injection based on the principle of cytotropism.*
- *Indication (detection) of the virus.*
- *Virus identification (type determination).*

- **Indication** of viruses, that is, non-specific detection of infection, is based on the determination of the biological properties of viruses and their interaction with sensitive cells.
- **Identification** means identifying the type and variant of the virus.





CULTİVATİON OF VİRUSES

LABORATORY ANİMALS

CELL (TISSUE) CULTURES

Monolayer

- **Primary**
- **Diploid cell strain**

- **Continuous cell lines**

EMBRYONATED CHICKEN EGGS

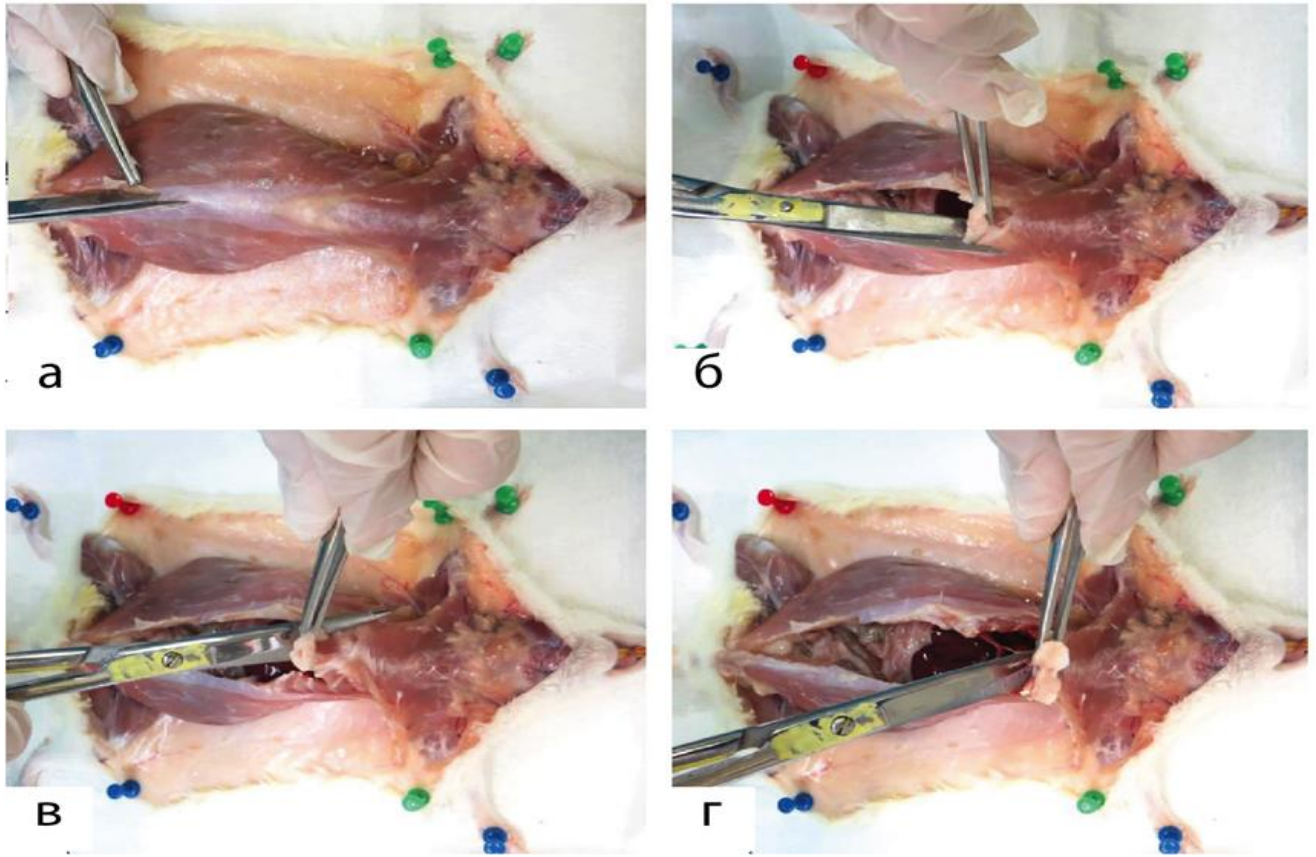
Cell Line - Types

- Primary cell lines – 5-10 times can divide
  - Monkey Kidney cell lines – Myxoviruses, Enteroviruses, Adenoviruses
  - Human Amnion cell line
  - Chick embryo cell line
- Secondary or diploid cell lines – 10 to 50 times can divide
  - Human Fibroblast cell line – for recovery of CMV
  - Human embryonic lung cell strain - MRC-5, WI-38 – for Vaccines, growth of virus
- Continuous cell lines – cancerous , immortal, haploid
  - HeLa cell line – from cervix
  - HEp-2 cell line - from larynx
  - KB cell line
  - McCoy
  - Vero cell line
  - BHK

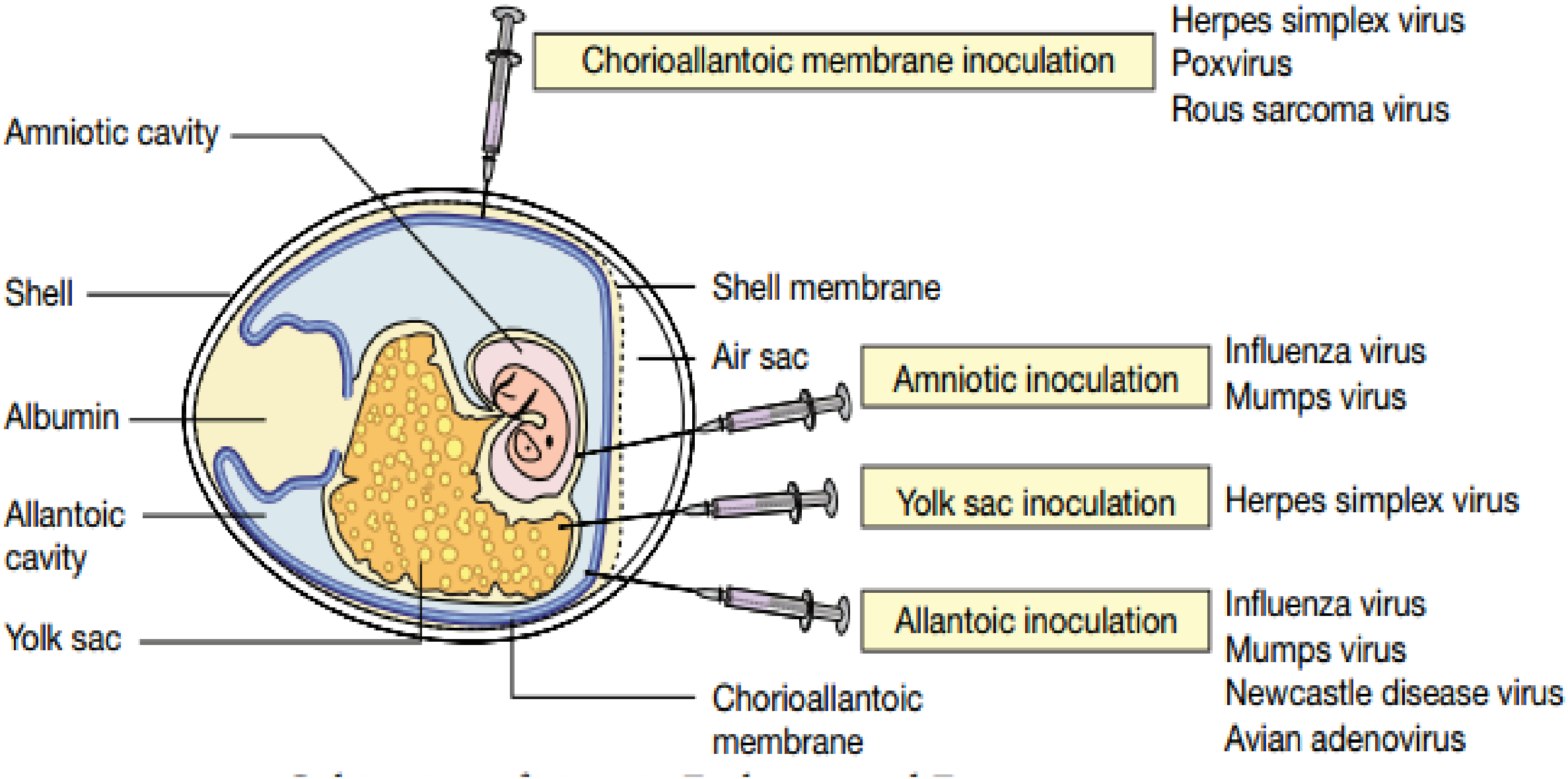
INDICATION OF VIRUSES IN THE BODY OF LABORATORY ANIMALS

The indication is based on the following characteristics

- *typical symptoms of the disease*
- *pathomorphological changes in animal organs and tissues*



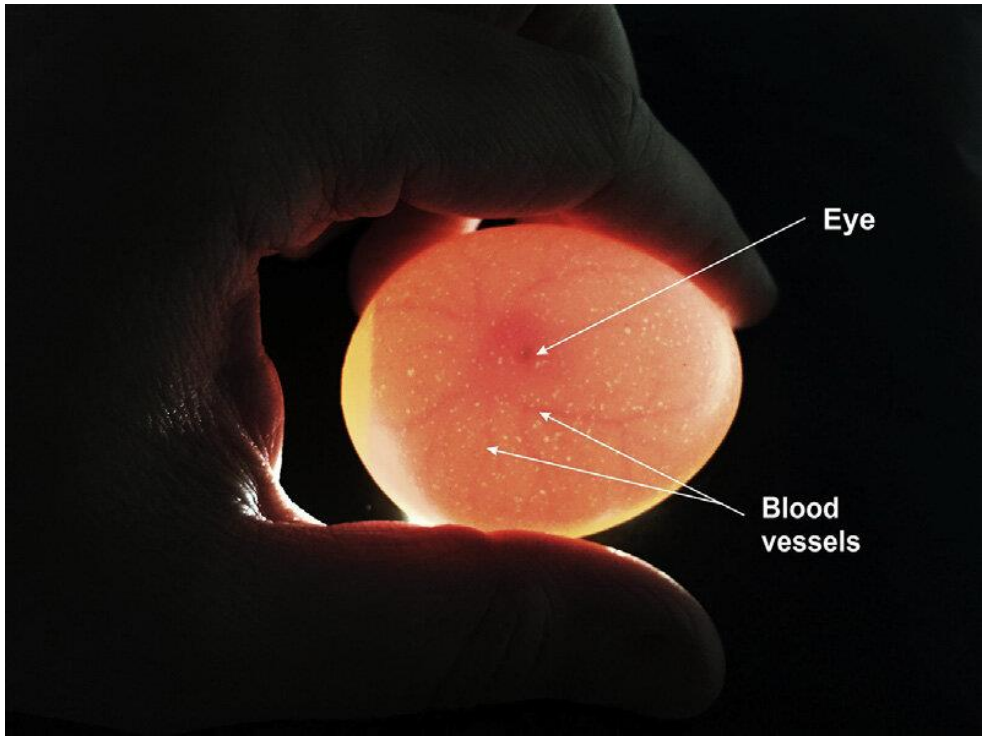
CULTIVATION OF VIRUSES IN EMBRYONATED EGG

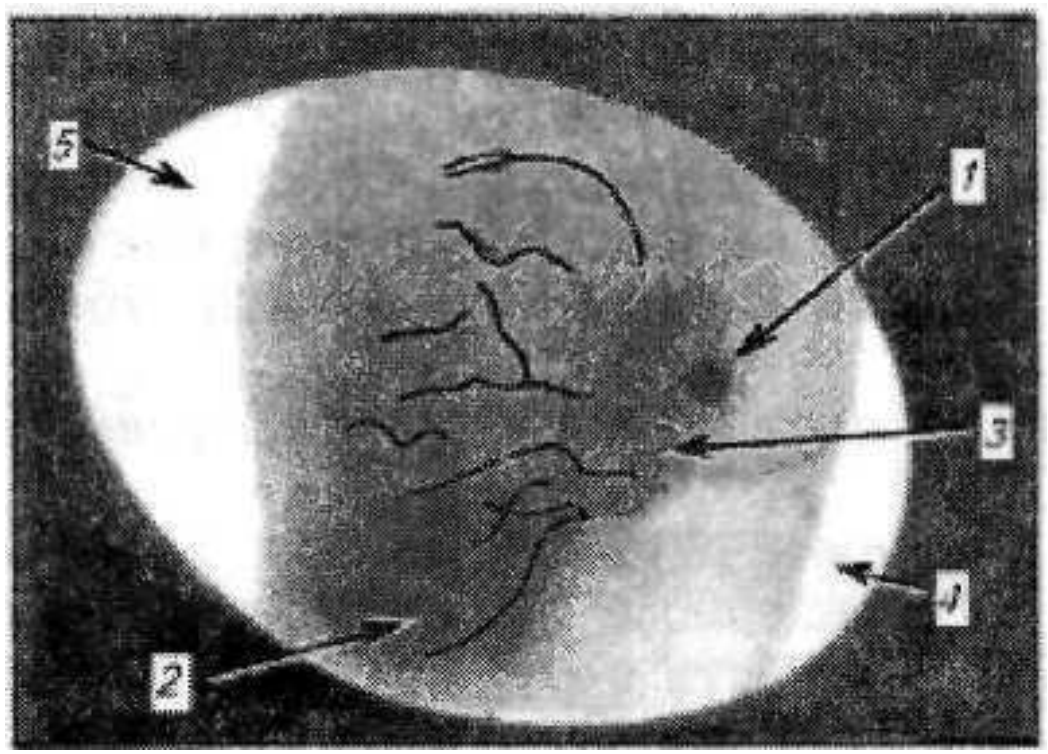


INDICATION OF VIRUSES IN INFECTED CHICKEN EMBRYOS

The development of viruses in an infected chicken embryo is determined by the following:

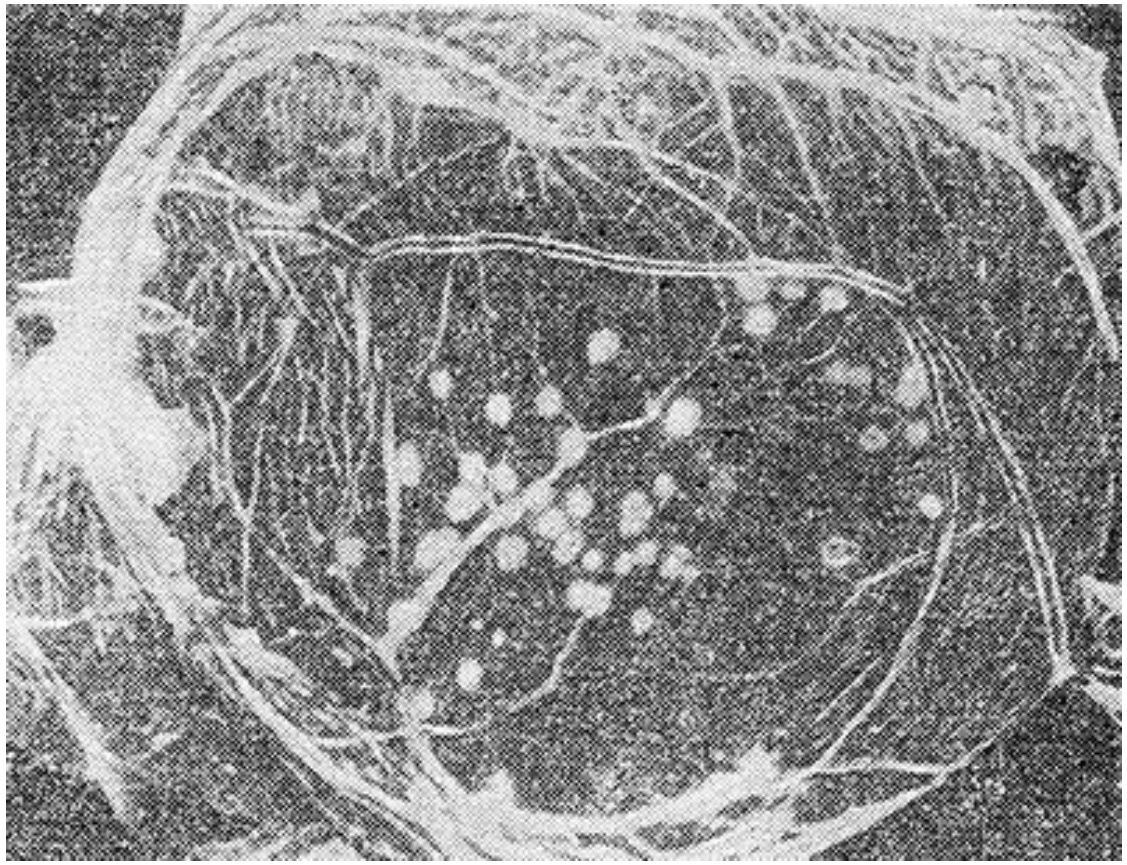
- *the death of the embryo,*
- *areas of necrosis caused by some viruses in the chorion-allantois membrane,*
- *hemagglutination reaction with amniotic and allantoic fluids,*





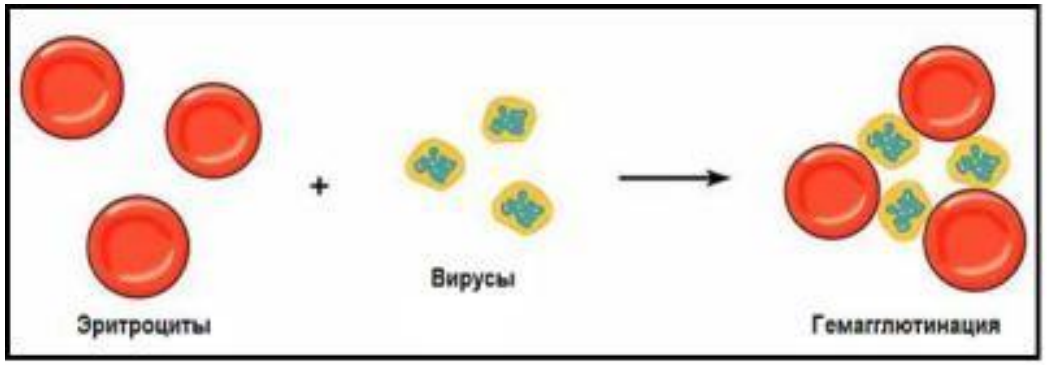
CHANGES IN THE CHORION-ALLANTOIS MEMBRANE

- During the study of changes in the chorionic-allantois membrane, it is cut with scissors and its contents are poured into a Petri dish.
- The chorion-allantois membrane remains inside the shell. It is removed with tweezers, placed in a Petri dish, washed with physiological solution, and ***the characteristics of focal lesions*** is studied on a dark background.



HEMAGGLUTINATION REACTION WITH AMNIOTIC AND ALLANTOIC FLUIDS

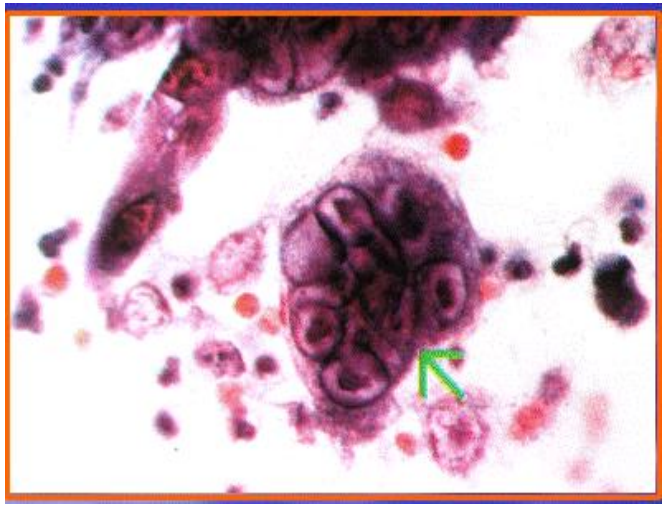
- The presence of the virus in the allantois and amniotic fluids of an infected embryo is determined by the hemagglutination reaction.



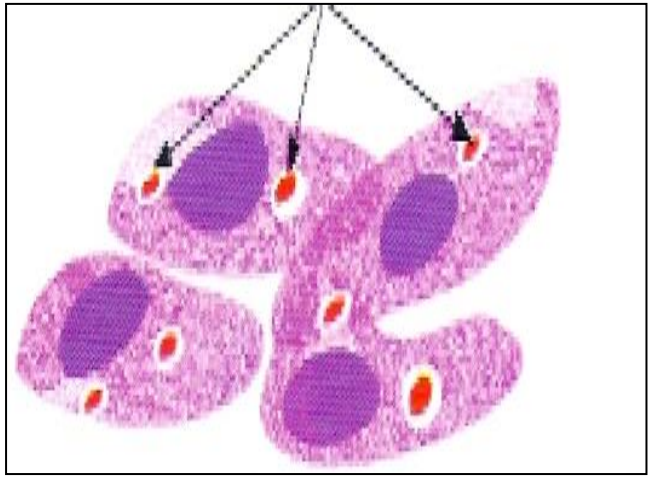
- This reaction is based on the ability of the hemagglutinin antigens of some viruses to agglutinate the erythrocytes of various animals and is used in the ***indication of viruses.***

METHODS OF INDICATING VIRUSES IN CELL CULTURES:

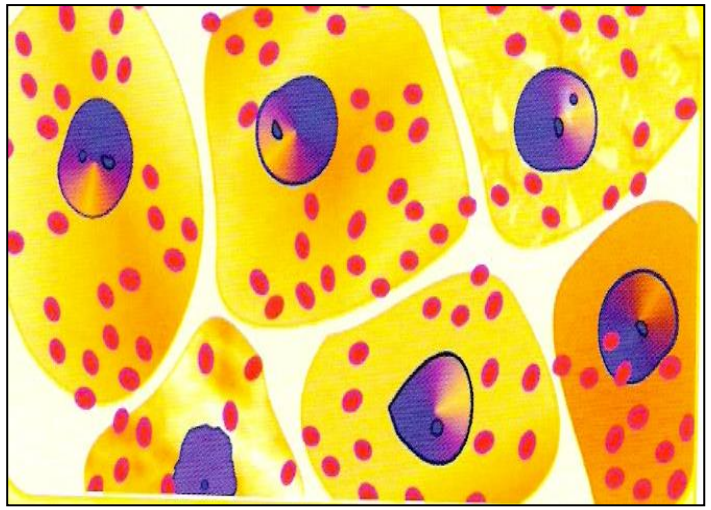
1. cytopathic effect (CPE)
2. intracellular inclusions
3. the formation of plaques
4. hemadsorption phenomenon
5. "color" test



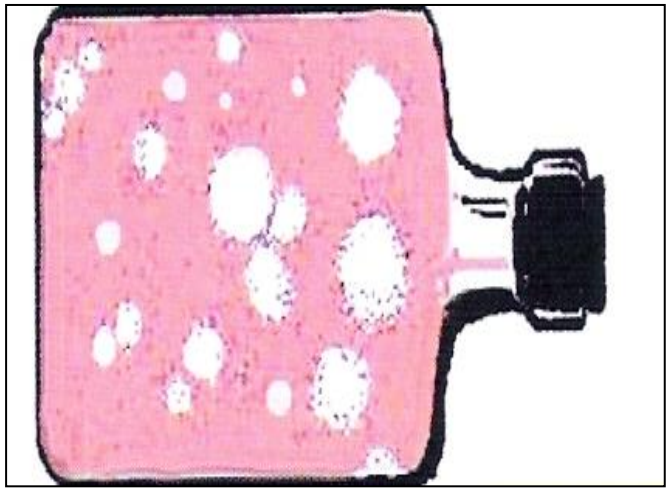
cytopathic effect (CPE)



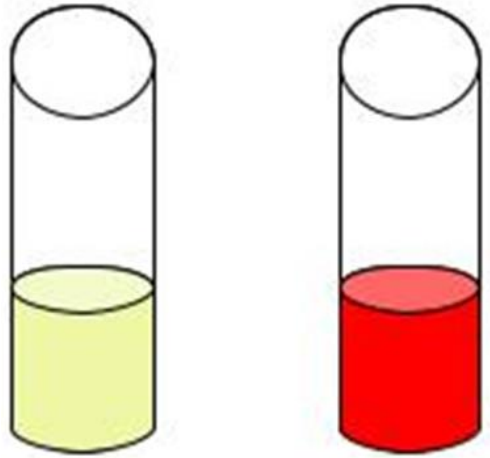
intracellular inclusions



hemadsorption phenomenon



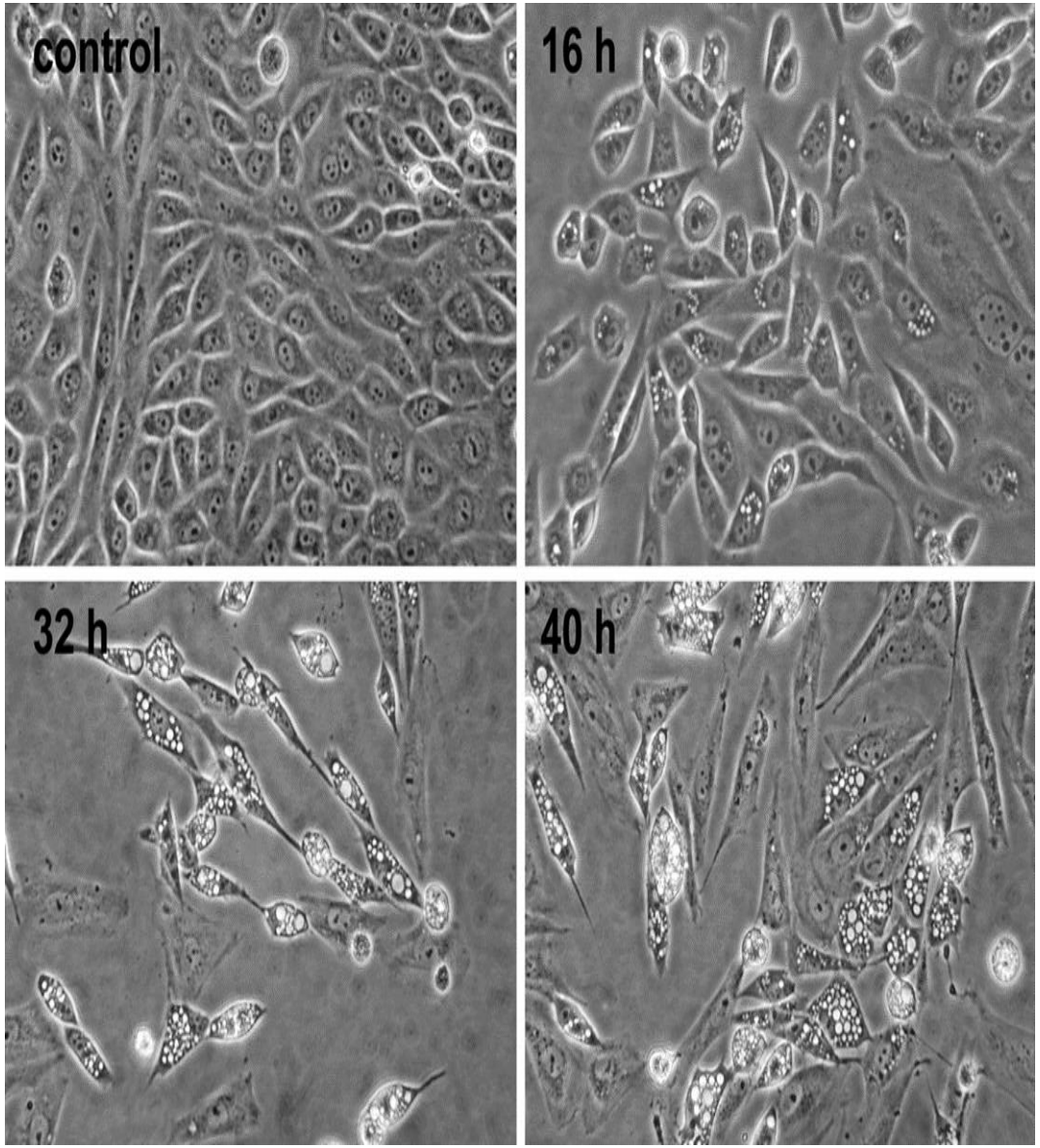
Negative colonies



«Color test»

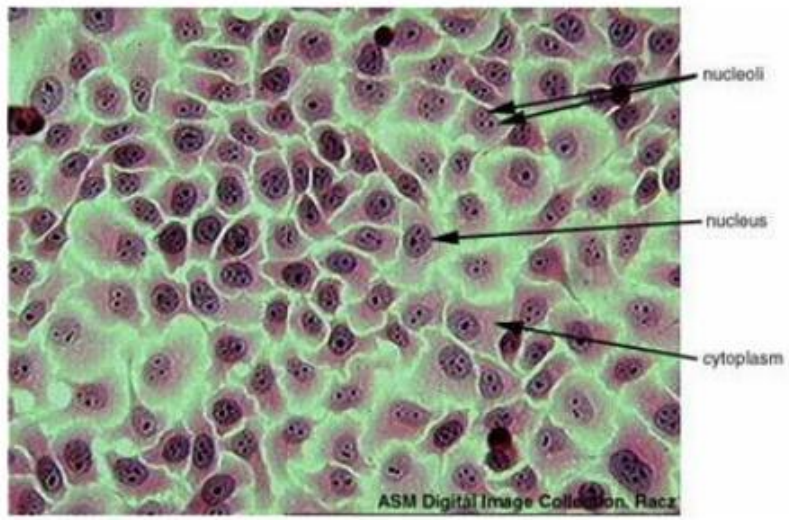
CYTOPATHIC EFFECT (CPE)

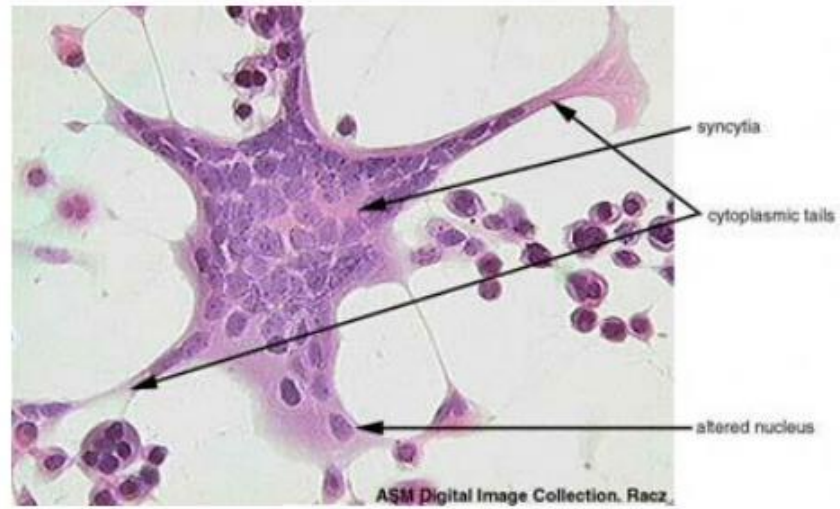
- During reproduction in cell culture, some viruses cause their degeneration, that is, cytopathic effect (CPE).
- After CPE virus infection, the tissue culture is evaluated in dynamics by studying it under a microscope at different times. CPE detection is one of the methods of virus **indication and identification.**
- Some viruses can be detected and identified by the **inclusions** they form in the cytoplasm and nucleus of infected cells.
- The shape of the inclusions varies, and their sizes vary from 0.25 μm to 25 μm.
- They represent the places of accumulation of virus particles and are detected by the Giemsa method and in preparations stained with fluorochrome.
- ***Different CPE are specific for different viruses.***



Examples of Cytopathic Effects of Viral Infection

- Nuclear shrinking (pyknosis)
- Proliferation of nuclear membrane
- Vacuoles in cytoplasm
- Syncytia (cell fusion)
- Margination and breaking of chromosomes
- Rounding up and detachment of cultured cells
- Inclusion bodies



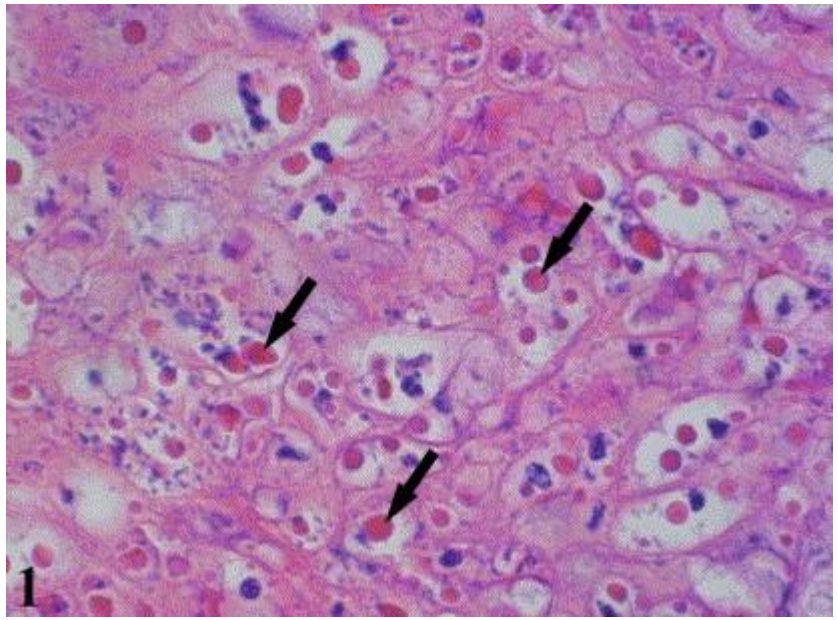


Cytopathic Effect (CPE)

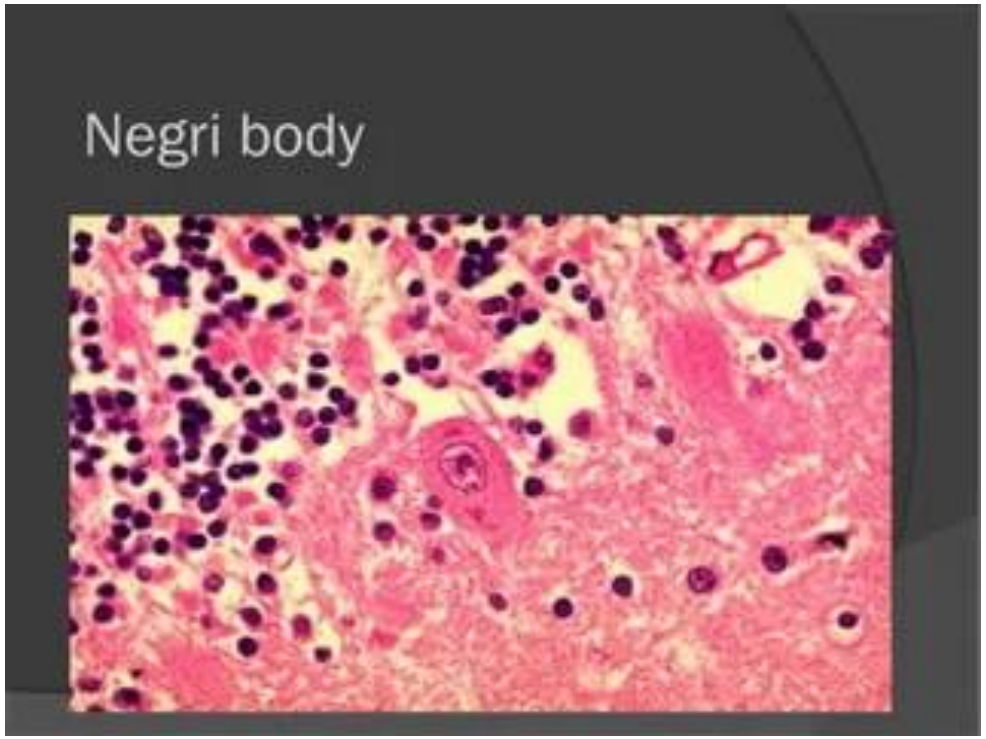
Types of Cytopathic elfect (CPE)	Virus
Rapid crenation(leaf like) and degeneration of the entire cell sheet	Enteroviruses – Eg. Polio,
Syncytium or multinucleated giant cell formation	Measles, RSV, HSV
Diffuse roundening and ballooning of the cell line	HSV
Cytoplasmic vacuolations	SV 40 (Simian vacuolating virus-40)
Large granular clumps resembling bunches of grapes	Adeno virus

INTRACELLULAR İNCLUSİONS (BODIES)

- Some viruses can be detected and identified by the inclusions they form in the cytoplasm and nucleus of infected cells.
- The shape of the inclusions varies, and their sizes vary from 0.25 µm to 25 µm.
- They represent the places of accumulation of virus particles and are detected by the Giemsa method and in preparations stained with fluorochrome.

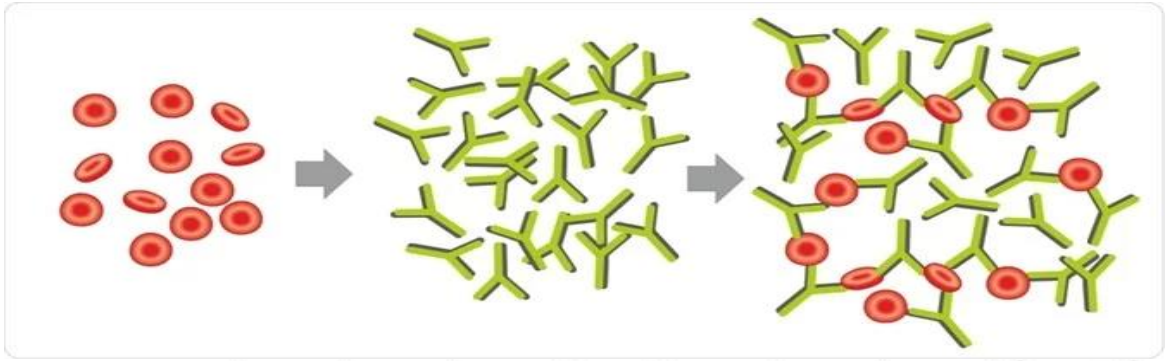


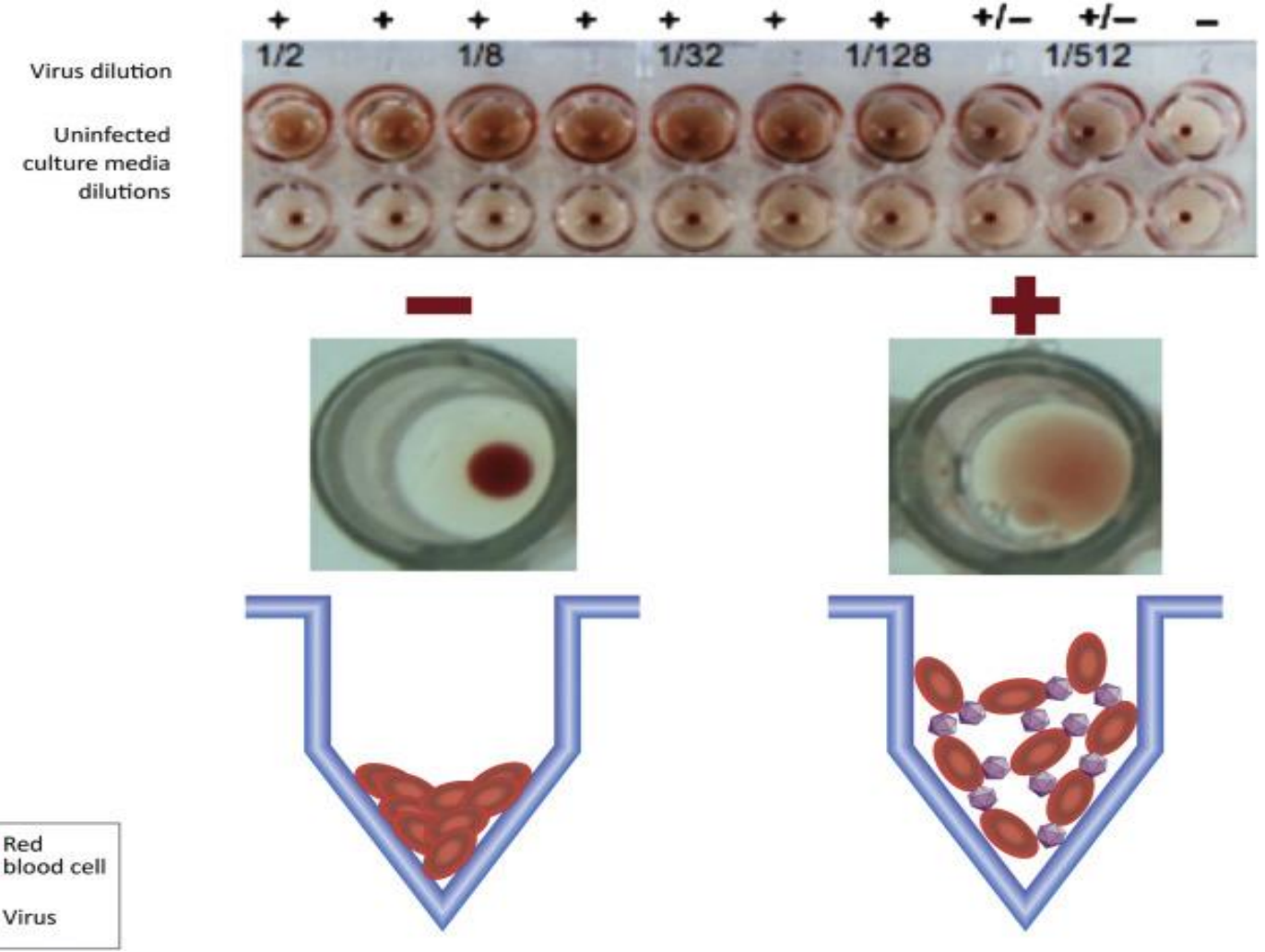
Guarnieri bodies



Inclusion body	Virus
Intracytoplasmic inclusion bodies	
Negri bodies	Rabies virus
Molluscum bodies	Molluscum contagiosum virus
Guarnieri bodies	Vaccinia virus
Bollinger bodies	Fowl pox virus
Perinuclear cytoplasmic acidophilic bodies	Reovirus
Intranuclear inclusion bodies	
Owl's eye inclusion bodies	Cytomegalovirus
Cowdry type A inclusion bodies	Herpes simplex virus and measles virus
Intranuclear basophilic	Adenovirus
Acidophilic inclusion bodies	Papovavirus

- **HEMAGGLUTINATION** is a reaction that causes clumping of red blood cells in presence of some **enveloped** viruses, such as the influenza virus. A glycoprotein on the viral surface, namely hemagglutinin, interacts with red blood cells, leading to the clumping of red blood cells and the formation of a lattice.
- In **absence of an enveloped** virus, red blood cells precipitate at the bottom of a well, forming a red-colored dot. However, in presence of a virus, red blood cell clumps are dispersed, forming no red-colored dot. This is the basic principle of a hemagglutination assay.

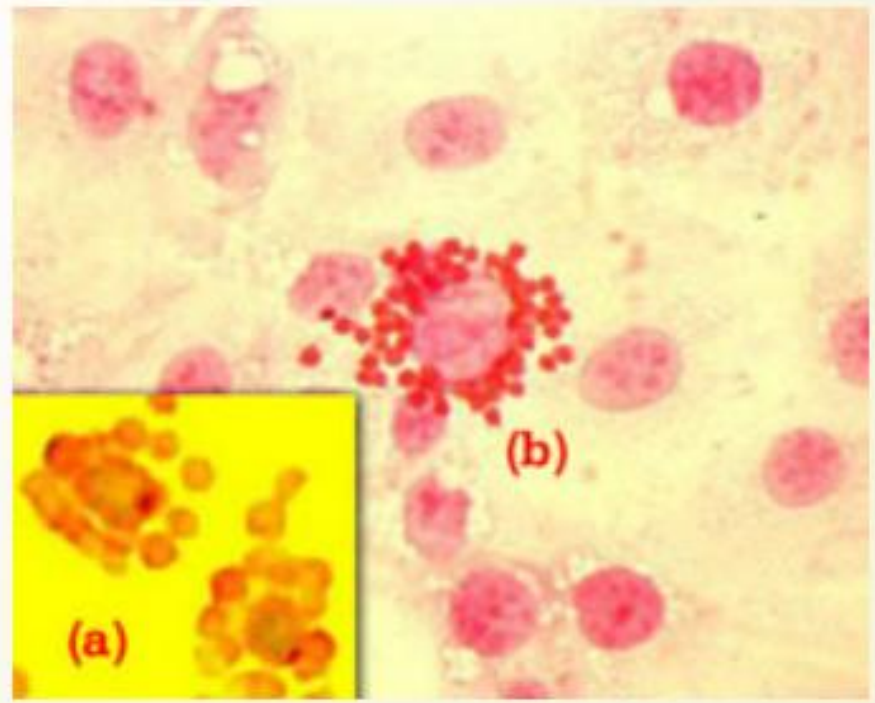




Hemadsorption

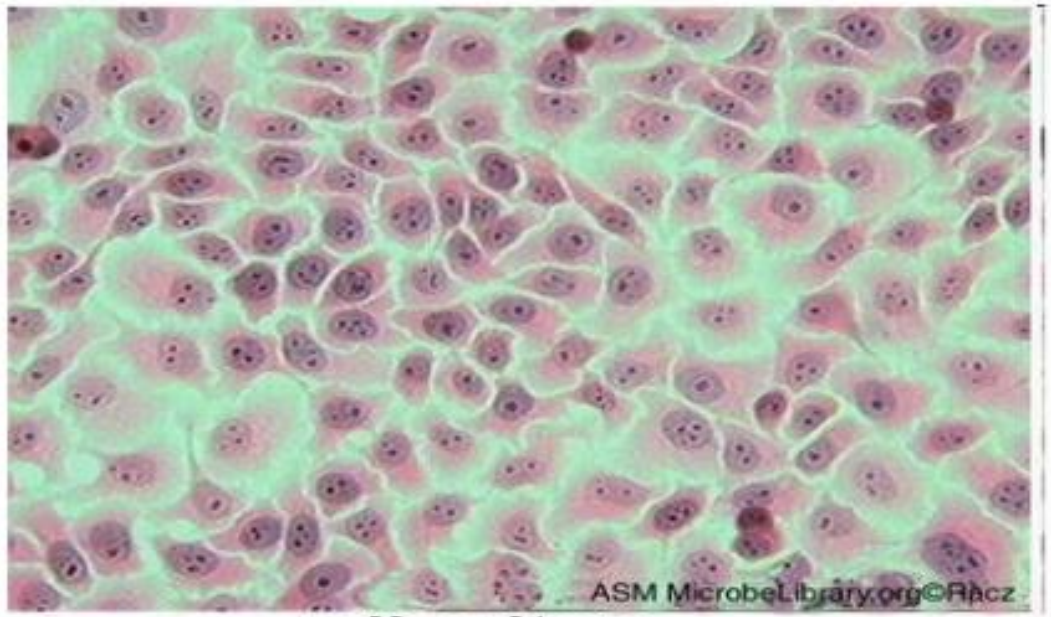
- Infective cell produce some proteins which have the ability to adsorb erythrocytes phenomenon known as *Haemadsorption* .
- Incorporation of viral glycoprotein peplomers into the plasma membrane of infected cells where they serve as receptors for ligands on the surface of erythrocytes.

Eg. orthomyxoviruses,
paramyxoviruses, and togaviruses,

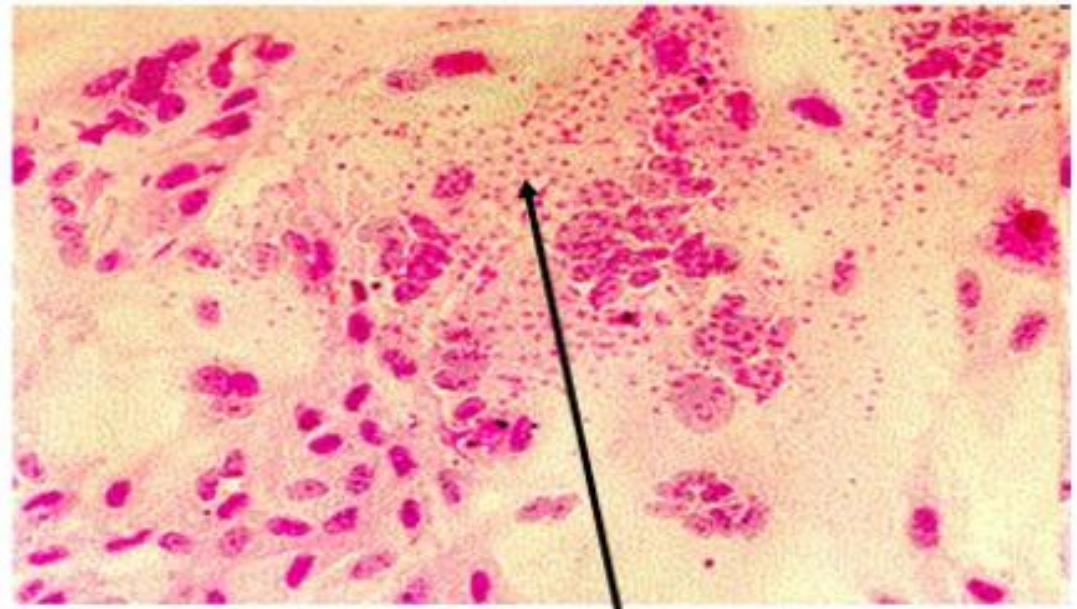


Hemadsorption (Hads)

- Virus growth in cell cultures is detected by testing for hemadsorption: red cells are added to the culture and adhere to virus budding from infected cells.
- If the culture tests positive, hemadsorption inhibition test with specific antisera is used to identify the virus.



cell culture



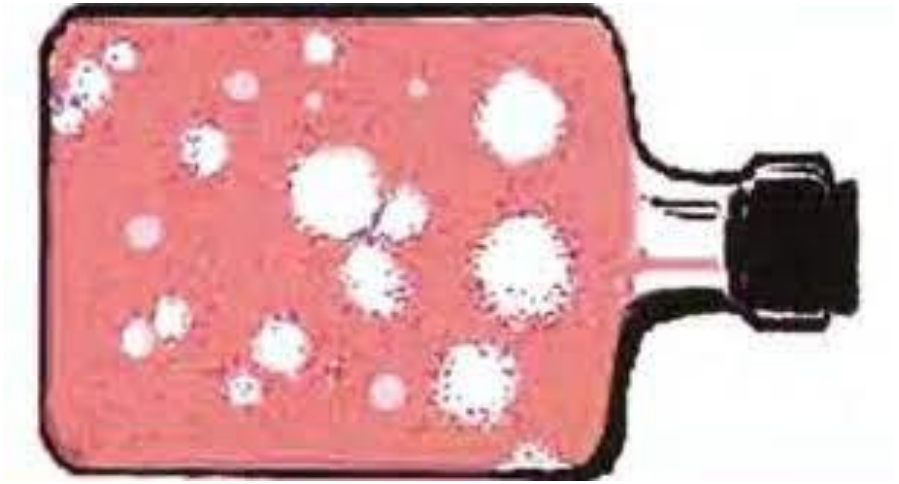
positive Hads

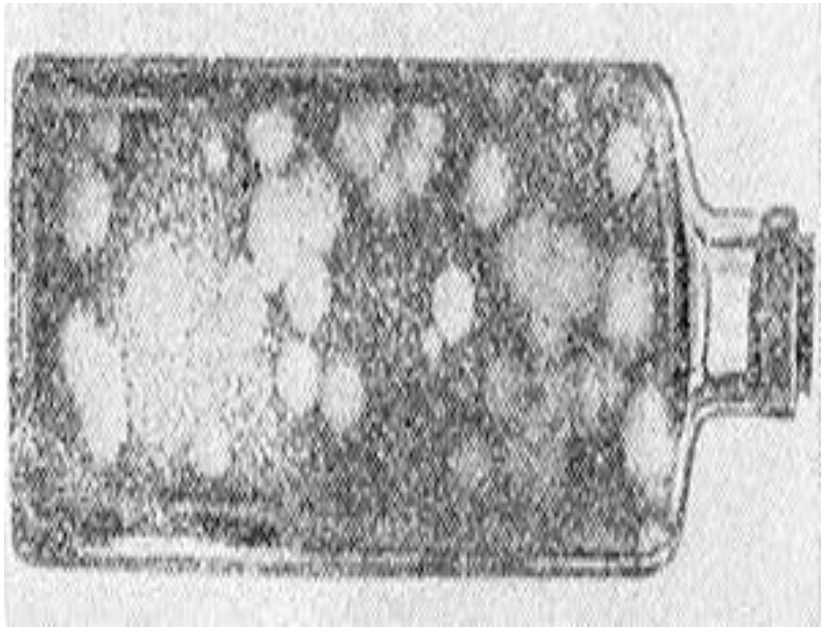
"NEGATİVE COLONİES"

•The growth of some viruses in cell cultures results in the destruction of cells in the corresponding region, which can be used to identify the viruses by revealing these areas ("negative colonies").

•After infecting the cell culture, adding an agar layer on top of it limits the areas of virus reproduction.

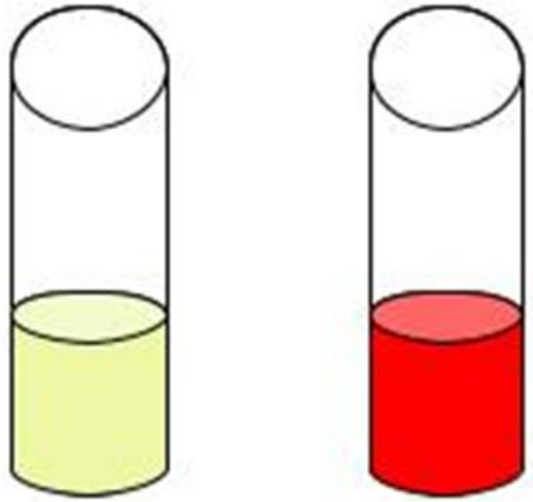
•As a result, the necrosis areas formed by them are isolated from each other.

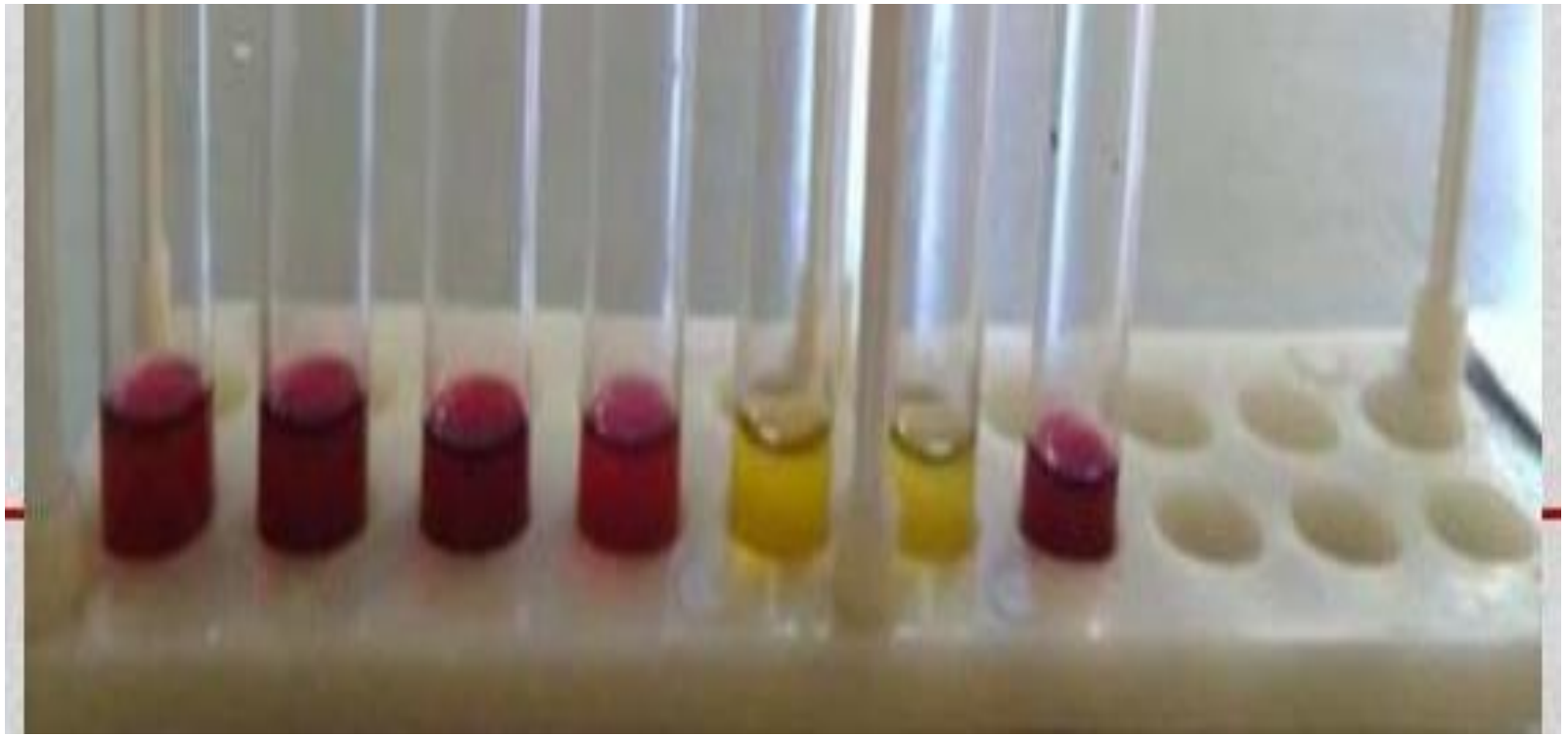




«COLOR TEST»

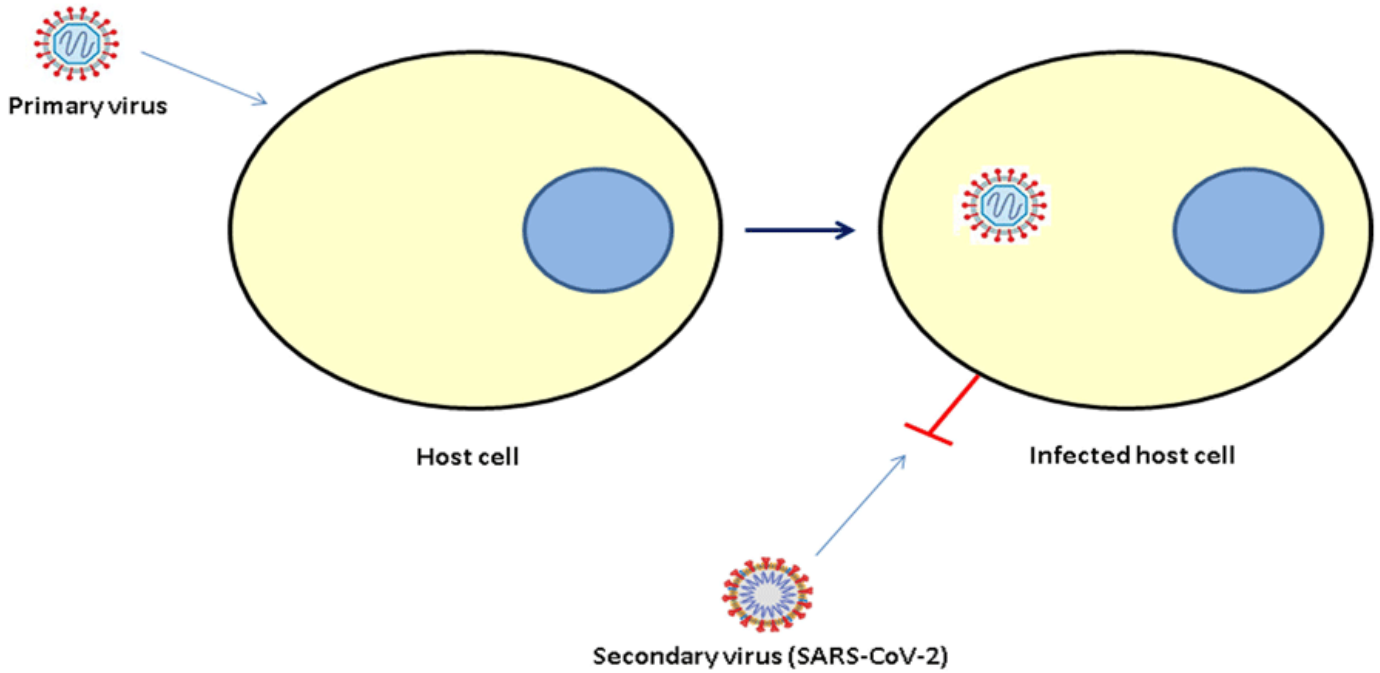
- The growth of viruses in cell cultures can be indicated by means of a "color test". For this, cell culture cultivated in a nutrient medium with an indicator (e.g., methyl red) is used.
- As the virus growth, the cells are destroyed, so the original color **(red)** of the medium remains unchanged.
- If the virus does not growth, a change in the color of the medium **(yellow)** is observed as a result of the effect of the metabolic products of the cells.





INTERFERENCE PHENOMENON

- In some cases, the phenomenon of interference is used to indicate viruses that do not cause CPE, especially those that are cultivated. The essence of interference is that a cell infected with one type of virus becomes resistant to other viruses.
- For example, rubella virus does not cause CPE despite cultivation in different cell cultures. In primary cell cultures, this virus can be detected due to the interference phenomenon.
- For this, the cell culture infected with the rubella virus is also infected with an indicator virus that produces CPE, for example, with the vesicular stomatitis virus. CPE is not observed because the growth of rubella virus in cell culture inhibits the replication of the indicator virus. However, when the rubella virus does not growth in cell culture, the indicator virus begins to multiply, and this is manifested by CPE.

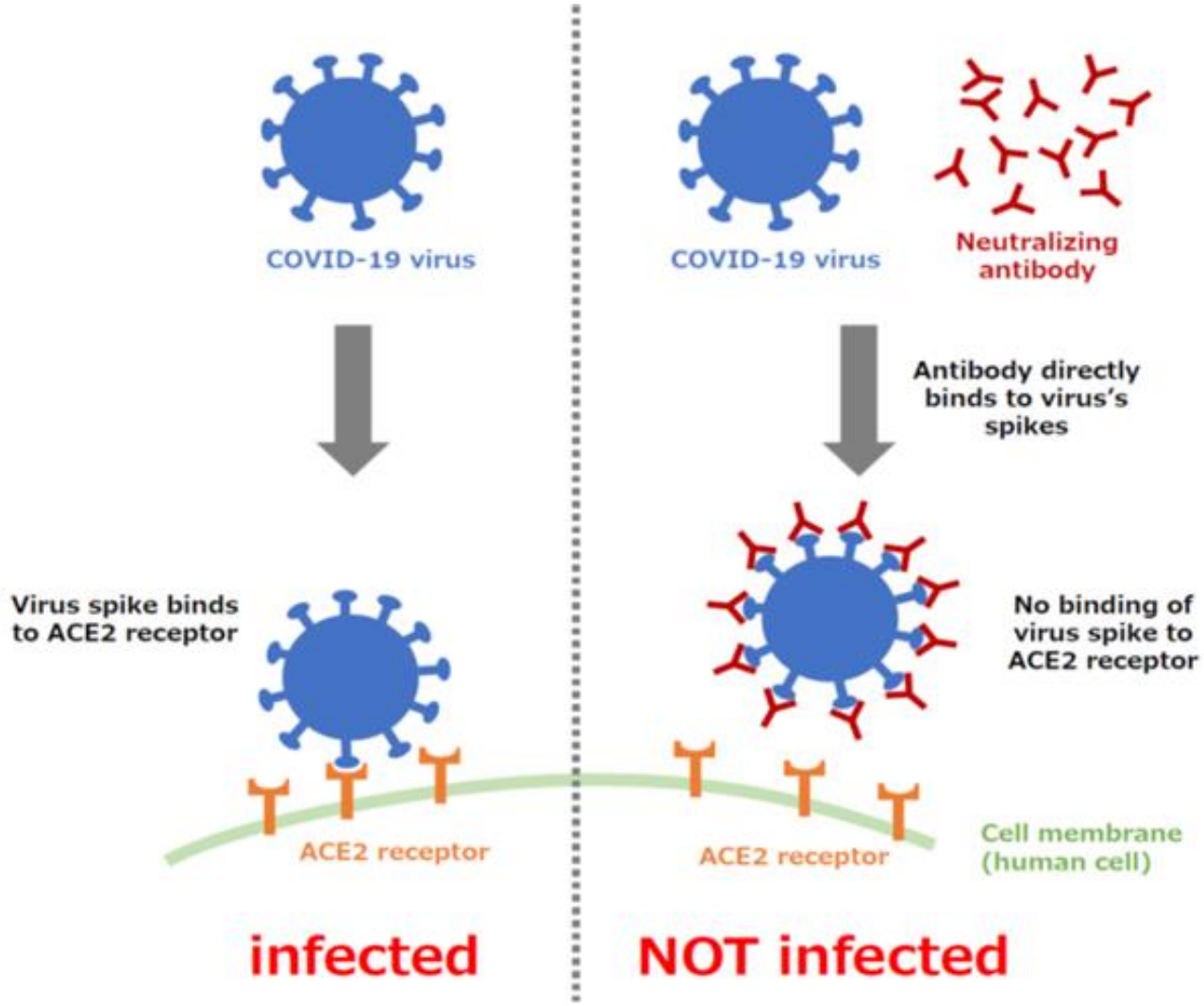


IDENTİFİCATİON OF VİRUSES

- Identification of viruses is the determination of their **variant, species, genus and family affiliation.**
- Virus identification is based on this principle: identifying the **unknown based on the known.**
- For the identification of viruses, serological reactions are performed using a known component - specific antiviral sera.
- These reactions include: neutralization reaction (**NR**), hemagglutination inhibition reaction (**HIR**), (hemadsorption inhibition reaction (**HAdsIR**), passive hemagglutination reaction (**PHAR**), complement fixation test (**CFT**), immunofluorescence reaction (**IFR**), enzyme-linked immunosorbent assay (**ELISA**)) and so on.
- These sera containing specific antivirus antibodies are called **diagnostic**.

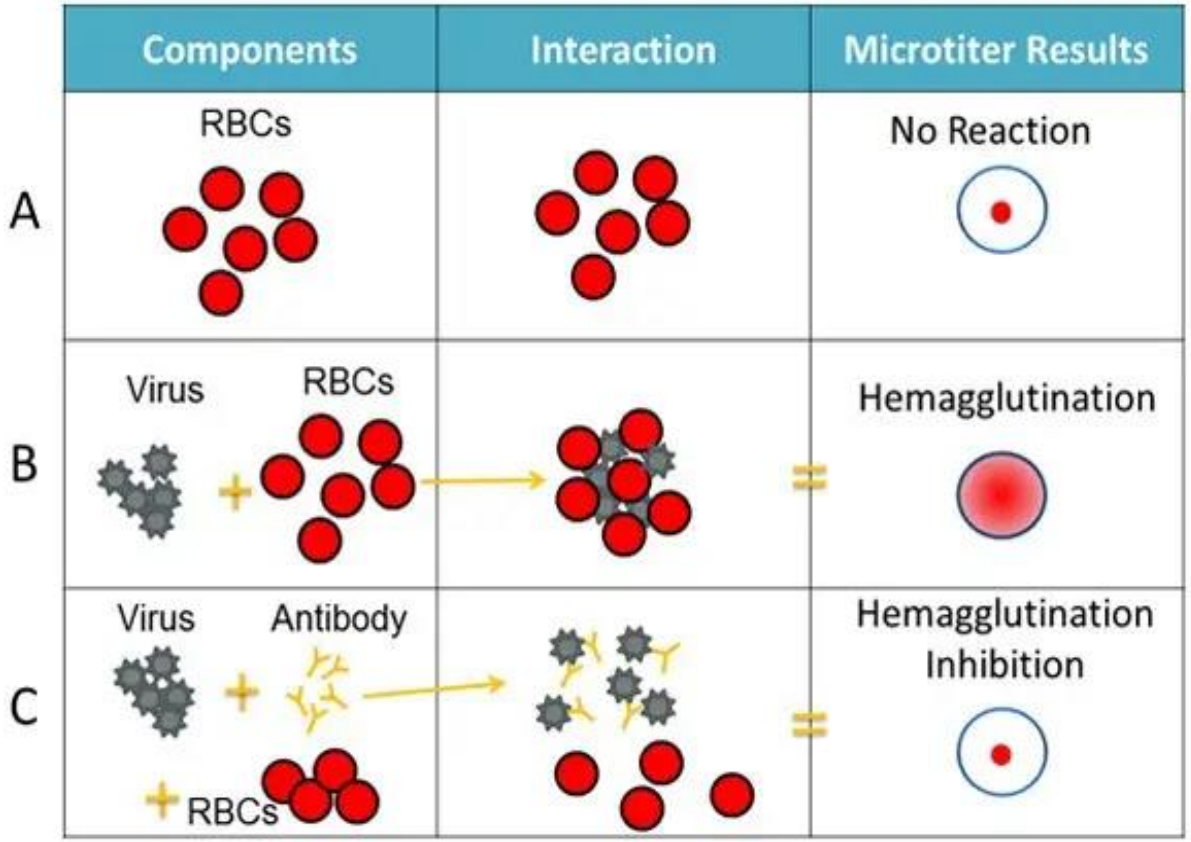
NEUTRALİZATİON REACTİON OF VİRUSES

- Virus neutralization reaction (biological neutralization reaction) allows identification of viruses.
- Due to the effect of appropriate antibodies, viruses do not cause disease in sensitive laboratory animals, do not have a cytopathic effect on cell and tissue cultures, and do not multiply in chicken embryos.



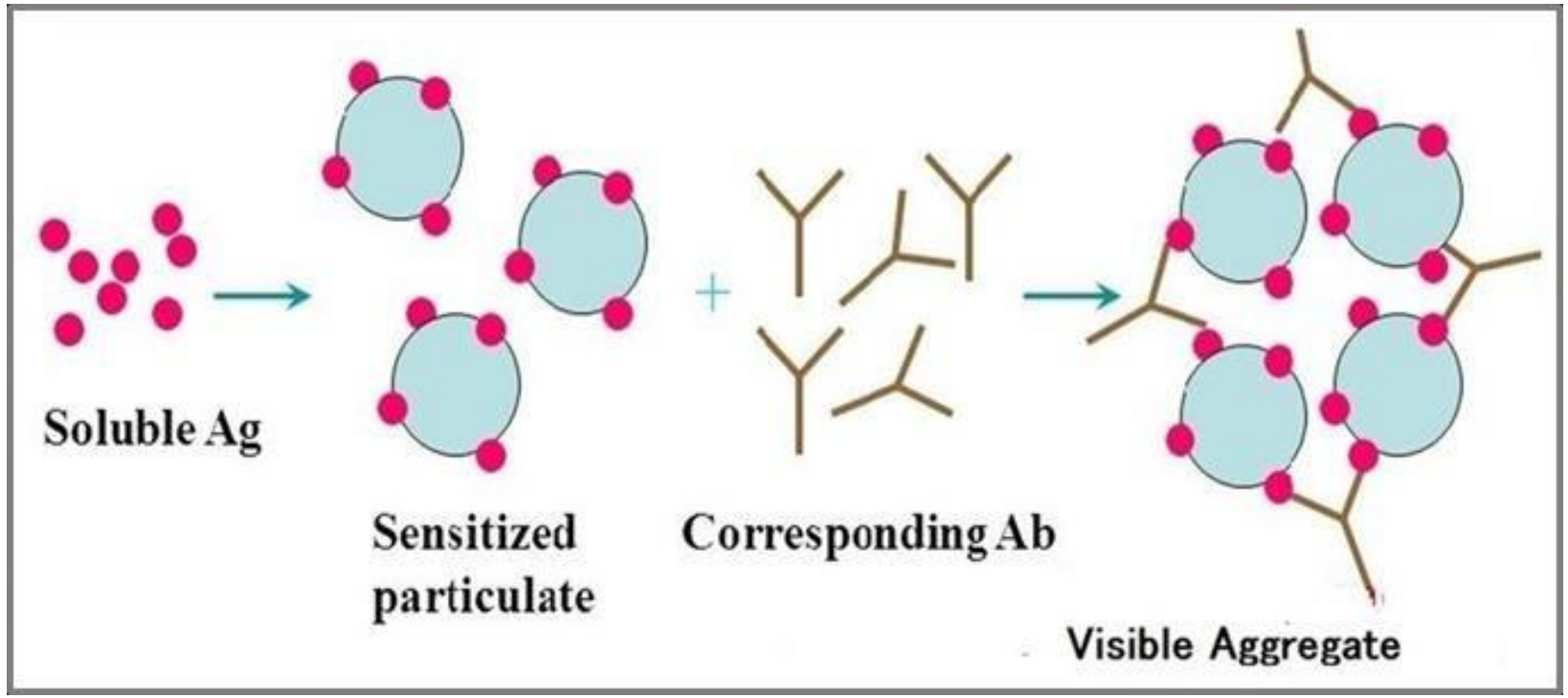
HEMAGGLUTINATION-INHIBITION (HI) ASSAY

- This reaction is used to identify some viruses (influenza, measles, tick-borne encephalitis, etc.).
- To determine the type of viruses in the examined material, serum containing antibodies against certain viruses is added to it.
- If there is a corresponding virus in the material, due to the effect of their antibodies, they lose their ability to agglutinate erythrocytes and the titer of the reaction decreases significantly.



PASSİVE HEMAGGLUTİNATİON REACTİON (PHAR)

The erythrocytes with adsorbed antigens come into contact with the corresponding antibodies in the blood serum, which causes the erythrocytes to stick together and settle to the bottom of the test tube or well in the form of sediment.



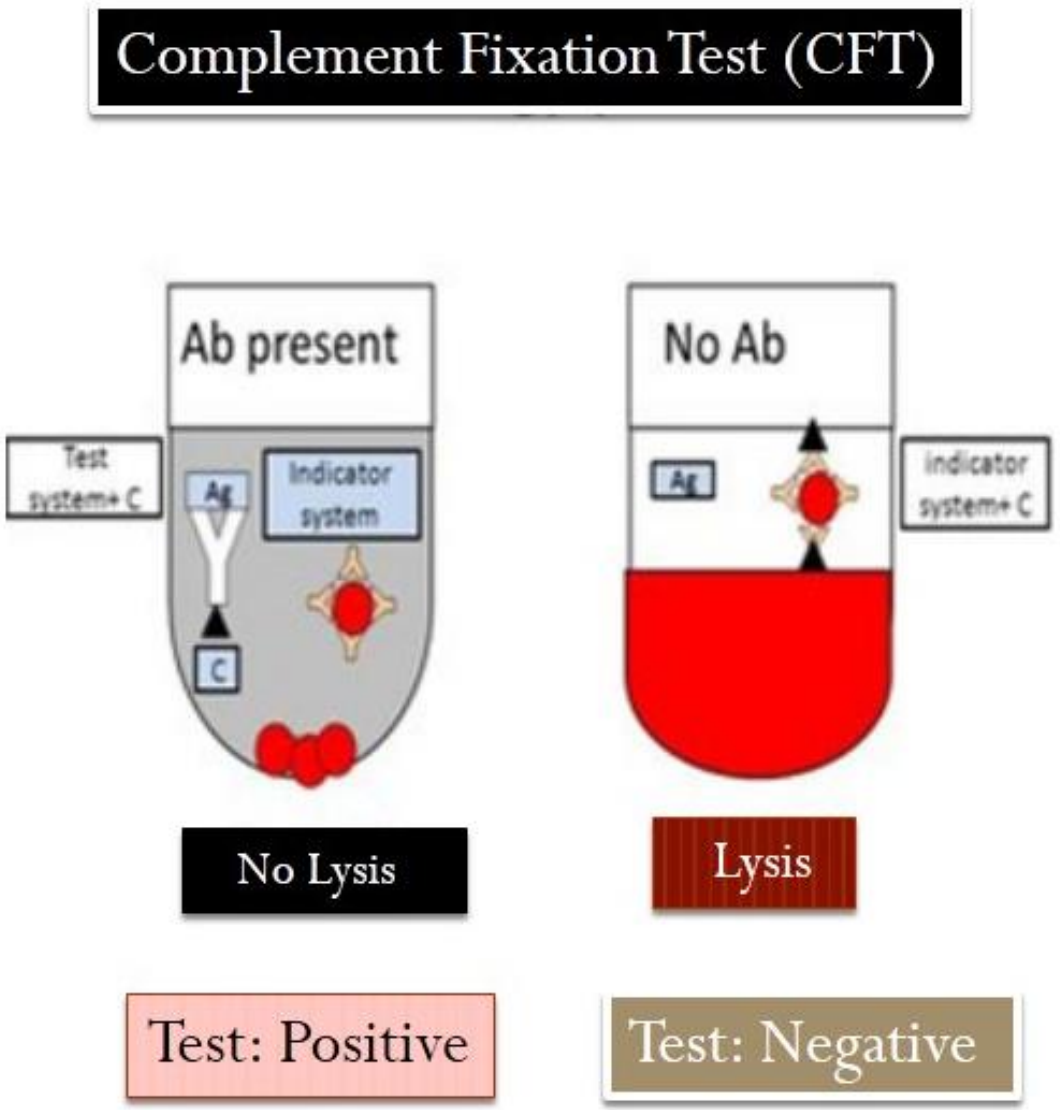
COMPLEMENT FIXATION TEST (CFT)

Complement fixation test (CFT) - when antigens and antibodies are compatible with each other, they form an immune complex, complement binds to it and a complement-antigen-antibody complex is formed. If the antigen-antibody complex is not formed, then the complement remains free.

CFT consists of two stages:

Phase 1st - incubation of the mixture containing antigen + antibody + complement,

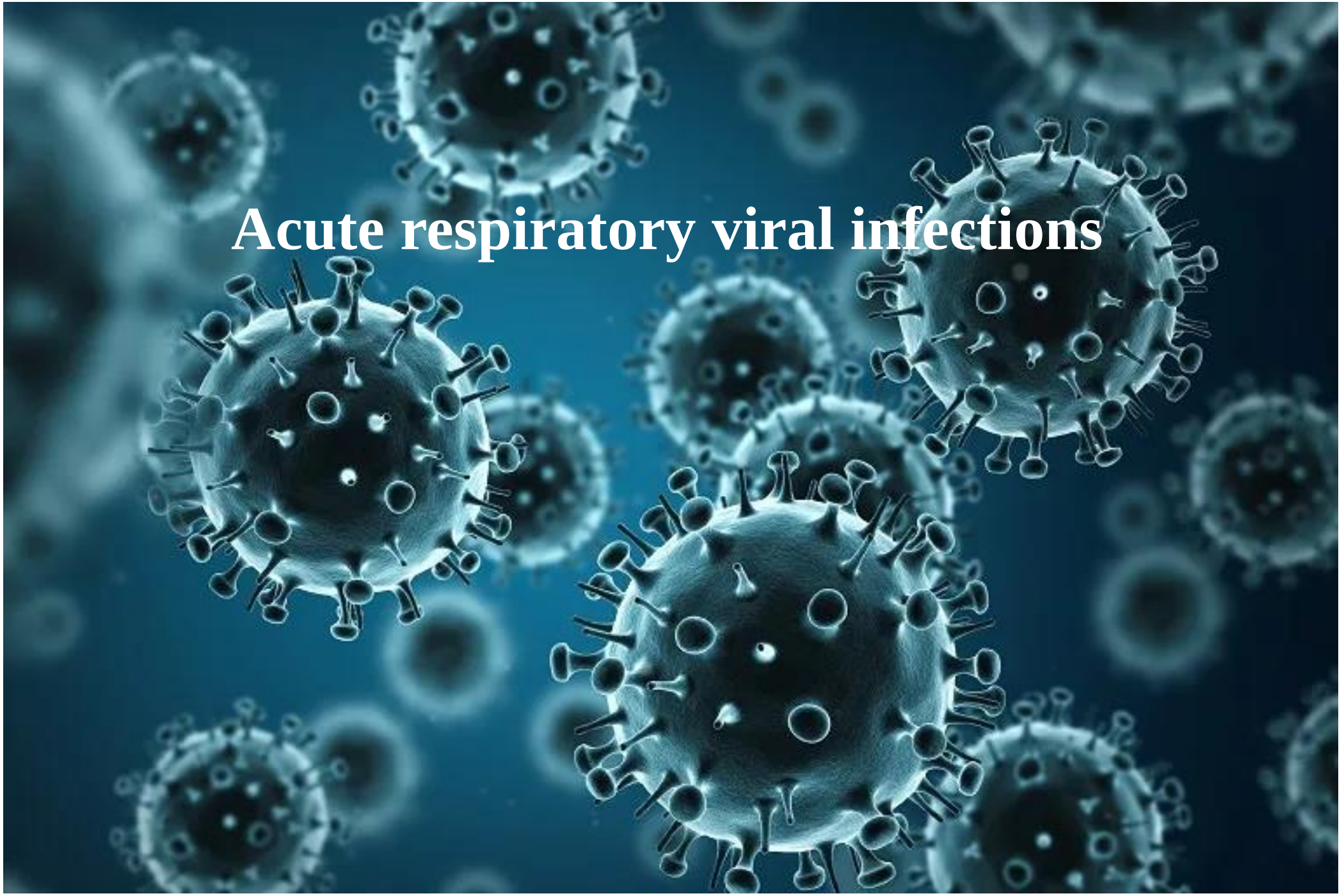
Phase 2nd (indicator) - detection of free complement in the mixture by adding a hemolytic system consisting of sheep erythrocytes and hemolytic serum. In the 1st phase of the reaction, when an antigen-antibody complex is formed, the combination of complement occurs, then in the 2nd phase, hemolysis of erythrocytes sensitized by antibodies will not occur (the reaction is **positive**). If the antigen and antibody do not match (no antigen or antibody in the test sample), the complement remains free and binds to the erythrocyte-anti-erythrocyte antibody complex in phase 2, causing hemolysis (**negative** reaction).



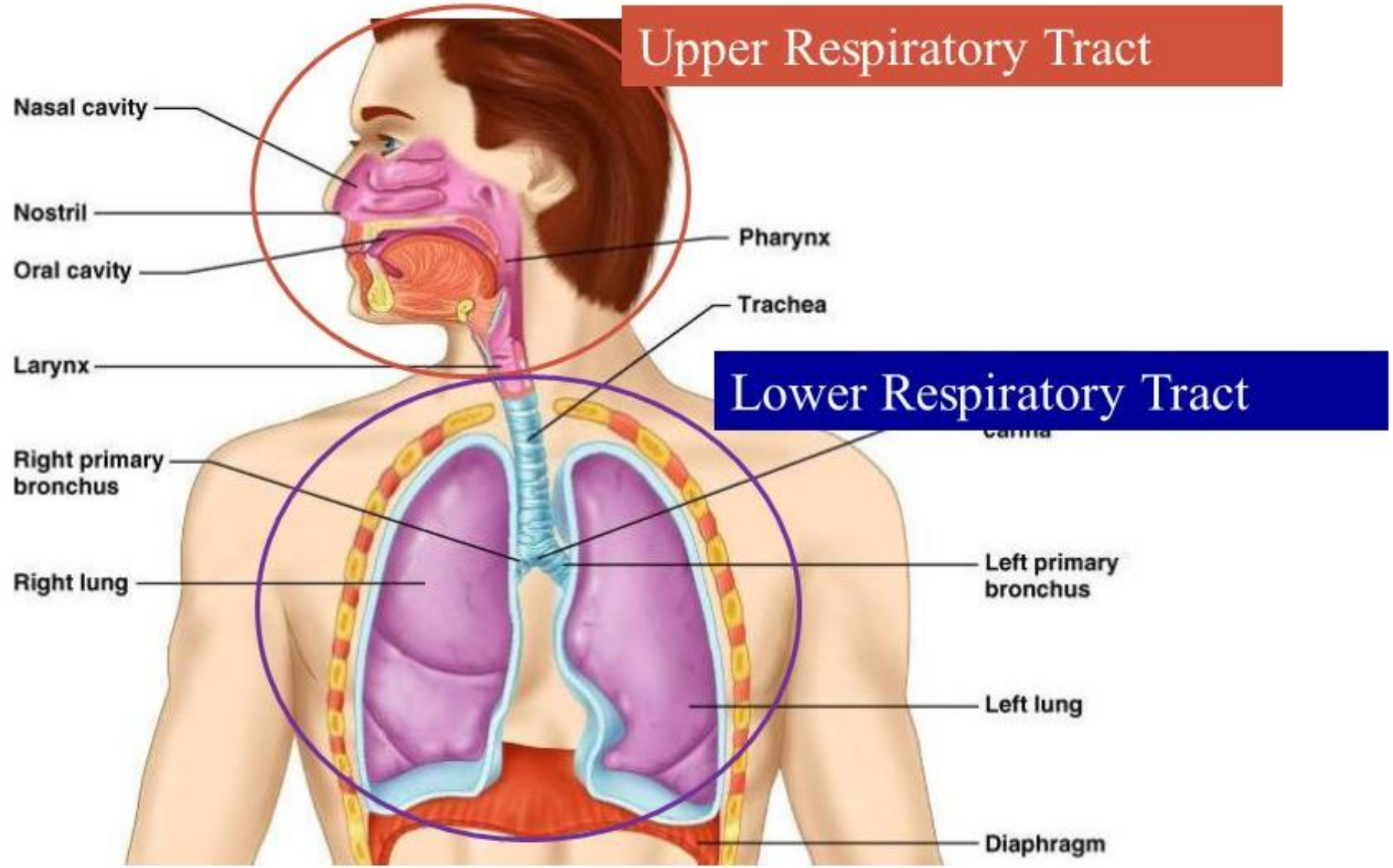
SEROLOGICAL METHOD

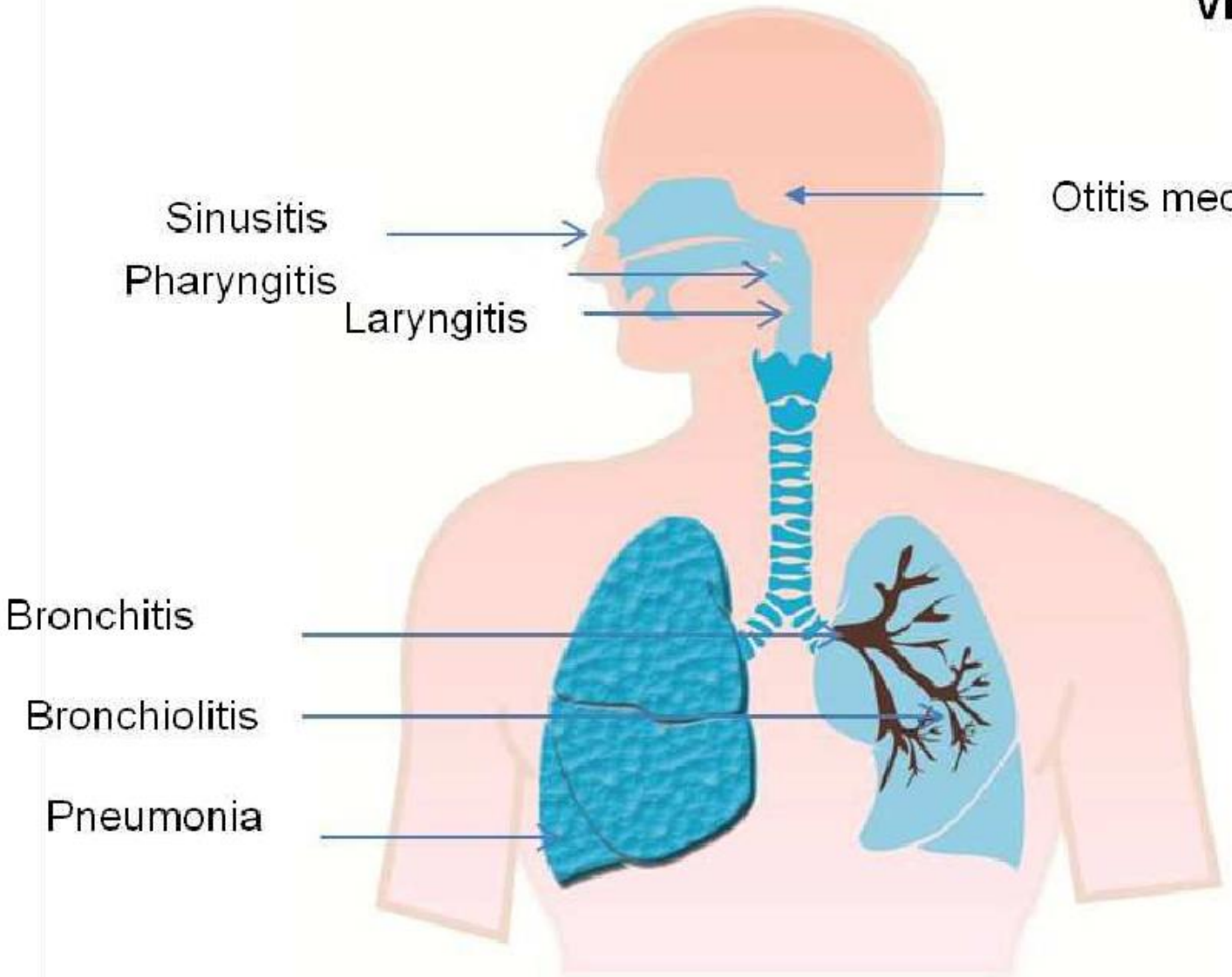
- Serological method - determination of antibodies in the blood of sick or recovered people. At this time, an increase in the titer of antibodies in the paired serum of the patient using viral diagnostics is considered as a positive result.
- Paired sera - two sera taken from the patient at the beginning of the disease and after 1-4 weeks.
- Serological reactions (PHAR, CFT, HIR, NR, ELISA, etc.) are performed with both sera to determine and compare antibody titers. The presence of IgM in the serum is determined for early diagnosis of the disease.


CONTINUE
LET GO


Acute respiratory viral infections

Upper and Lower Respiratory Tracts





Viruses that infect the upper respiratory tract

Rhinovirus
Coronavirus
Influenza virus
Parainfluenza virus
Respiratory Syncytial virus
Herpesvirus
Adenovirus
Bocavirus
Coxsackivirus

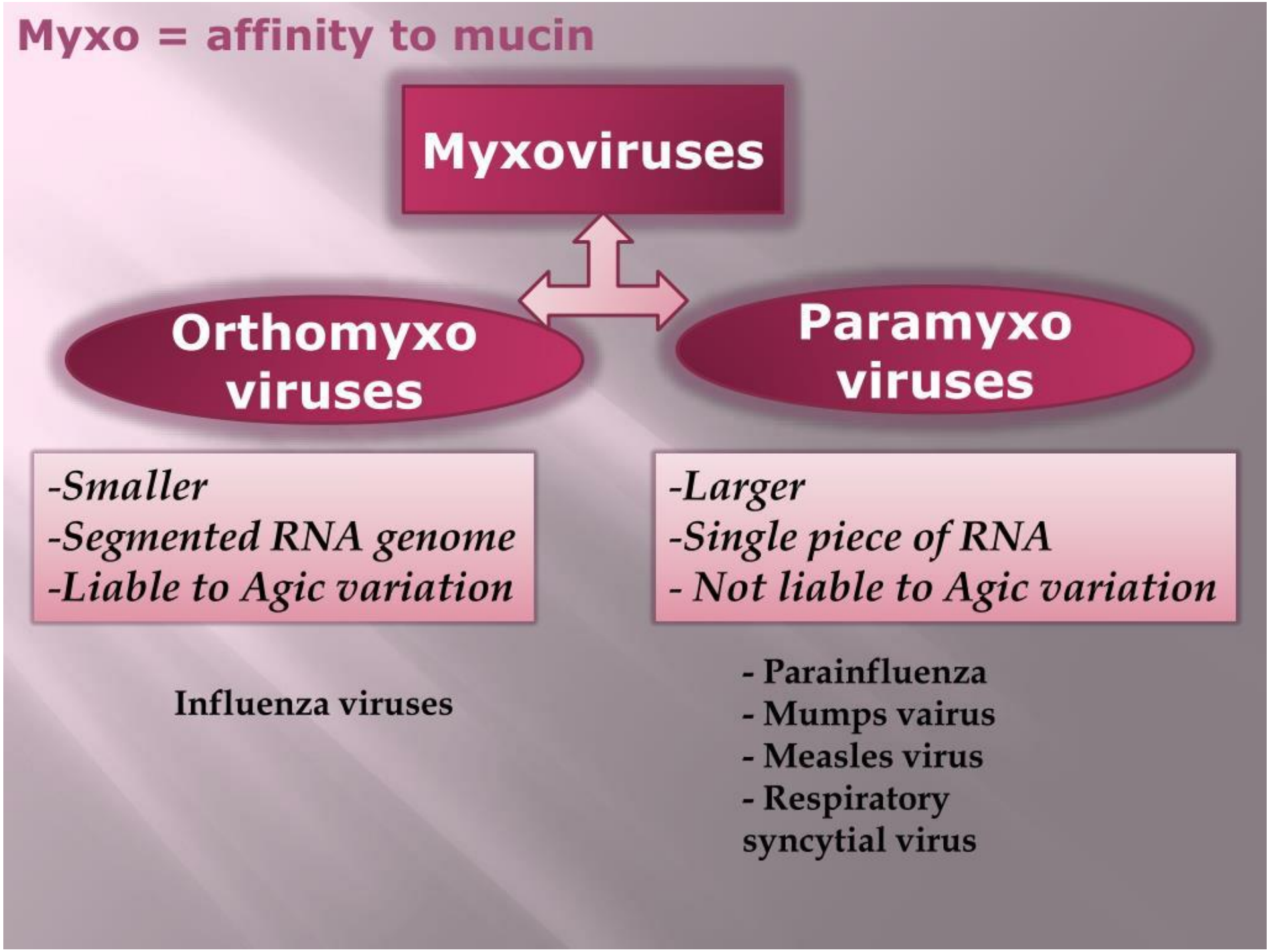
Viruses that infect the lower respiratory tract

Influenza virus
Parainfluenza virus
Respiratory Syncytial virus
Adenovirus
Bocavirus
Metapneumovirus

Viruses that initiate infection via respiratory tract

Site of infection	Family	Viruses
Local respiratory infection	Orthomyxoviridae	Influenza A and B viruses
	Paramyxoviridae	Parainfluenza viruses (4 types), respiratory-syncytial virus RSV (3 types)
	Picornaviridae	Rinoviruses (113 types)
	Reoviridae	Reoviruses (3 types)
	Coronaviridae	Types 1-4
	Adenoviridae	Types 1-7, 14, 21
Generalized diseases, usually with initial respiratory symptoms	Herpesviridae	Varicella virus, Epstain-Barr virus (EBV), cytomegalovirus
	Paramyxoviridae	Mumps and measles viruses
	Togaviridae	Rubella virus
	Picornaviridae	Some enteroviruses
	Bunyaviridae	Hantaviruses
	Arenaviridae	Lassa fever virus

Myxo = affinity to mucin



Orthomyxovirus (Influenza) Family

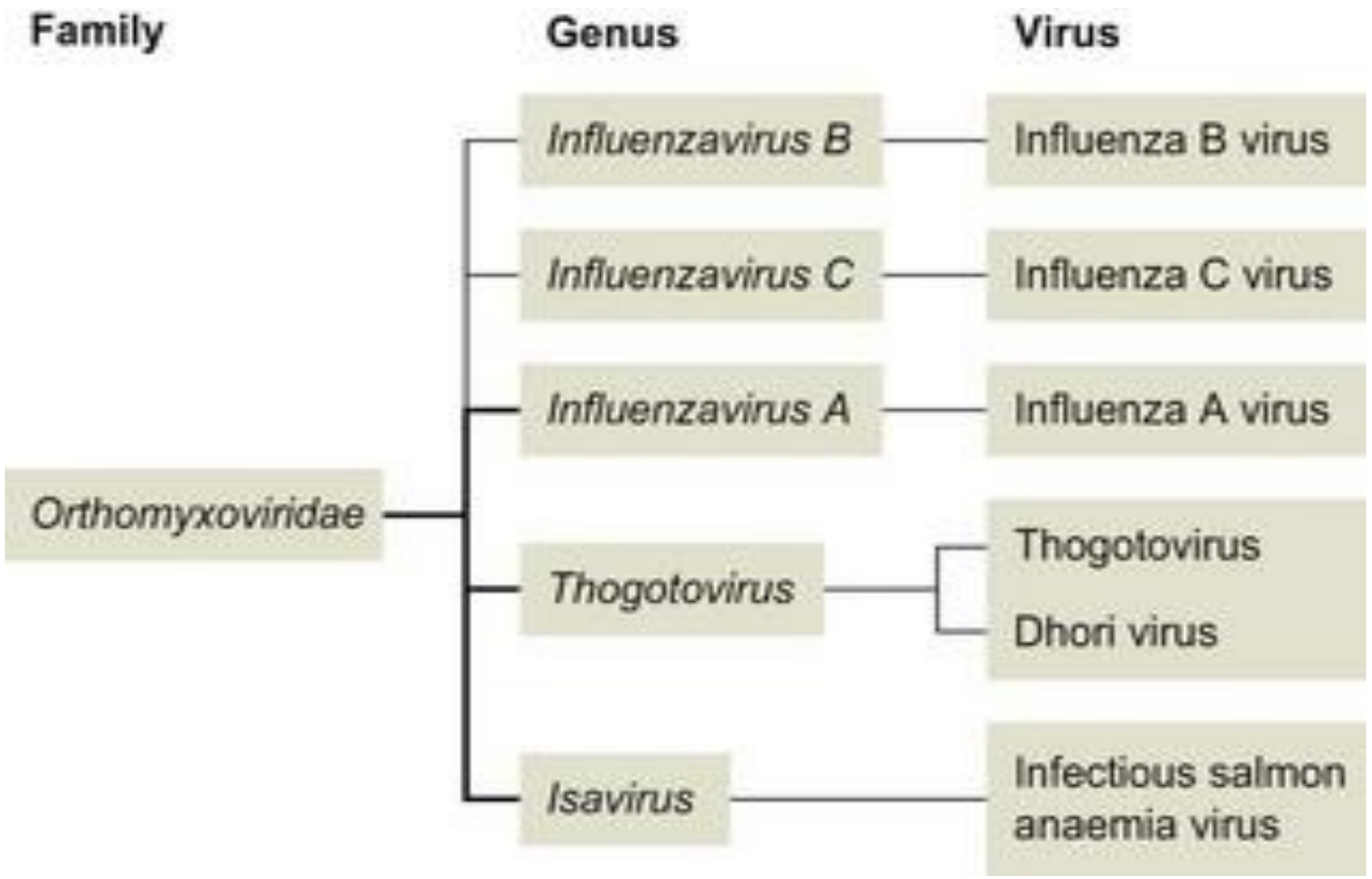
The name myxovirus was originally applied to influenza viruses. It meant virus with an affinity for mucins. Now there are 2 main groups – the orthomyxoviruses and the paramyxoviruses

Differences between orthomyxoviruses and paramyxoviruses

Feature	Orthomyxoviruses	Paramyxoviruses
Viruses and diseases	Influenza A,B,C	Mumps, measles, respiratory syncytial, parainfluenza
Genome	Single-stranded RNA in 8 pieces, MW 2-4x10^6	Single-stranded RNA in single piece, MW 5-8x10^6
Inner ribonucleo-protein helix	9-nm diameter	18-nm diameter

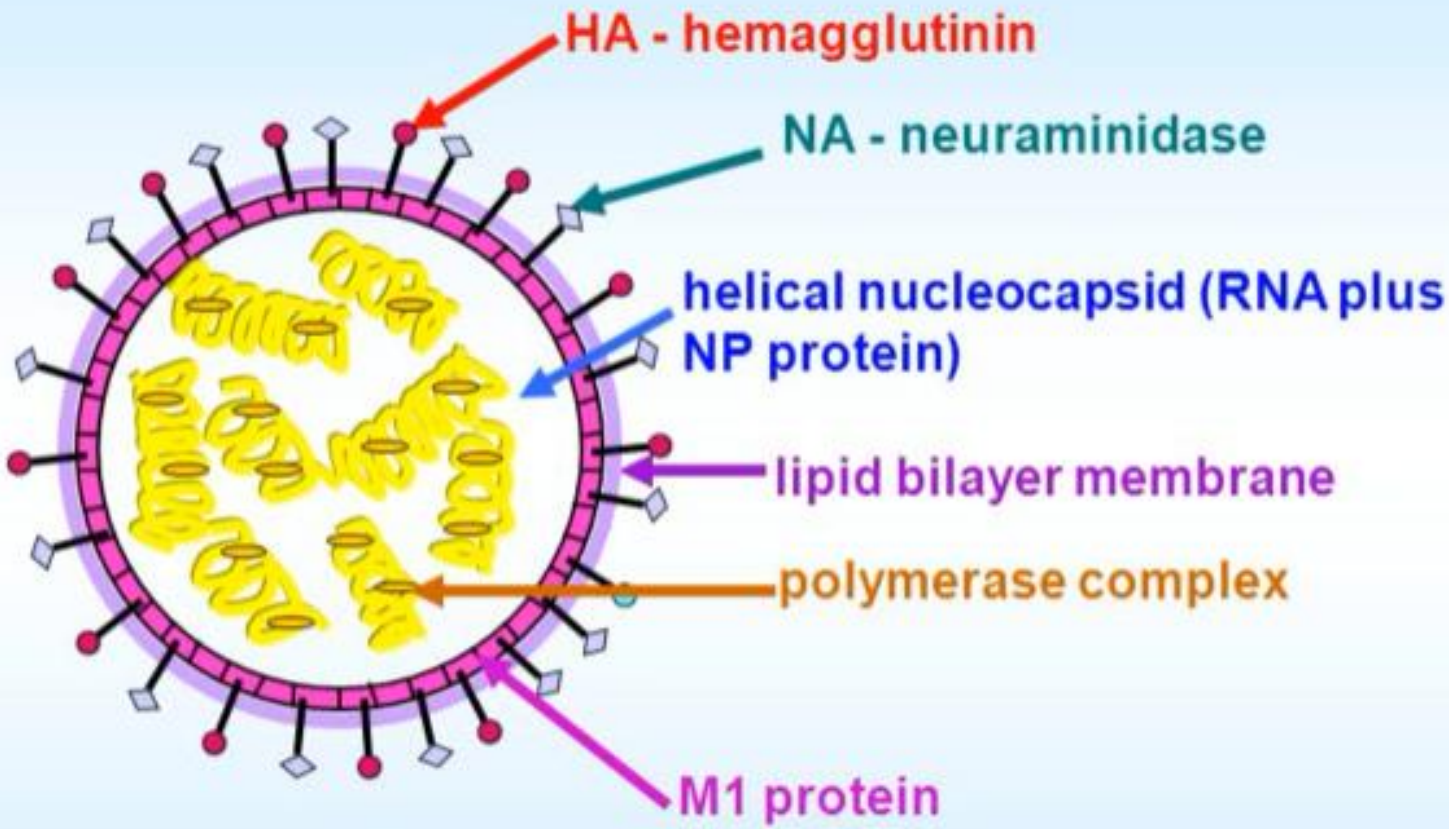
Ortomyxoviridae - Taxonomy

- Kingdom: Orthornavirae
- Phylum: Negarnaviricota
- Class: Insthoviricetes
- Order: Articulavirales
- Family: Orthomyxoviridae
- Genus:
  - Alphainfluenzavirus
  - Betainfluenzavirus
  - Gammainfluenzavirus
  - Deltainfluenzavirus
  - Isavirus
  - Quaranjavirus
  - Thogotovirus



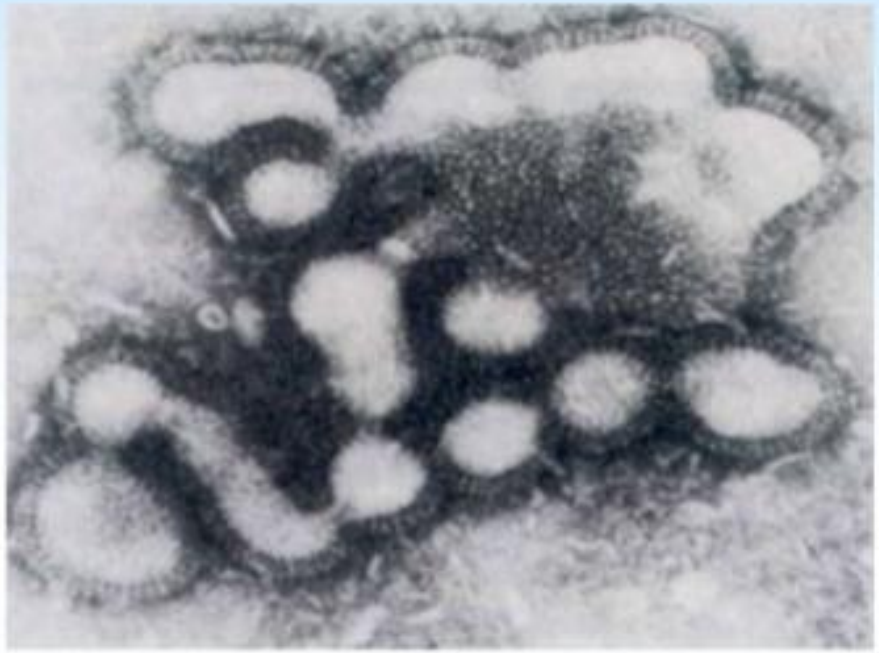
- **Influenzavirus D** - primarily affect cattle and are not known to infect or cause illness in people. (Medical Microbiology, 9th Edition, 2020, Patrick Murray, Ken Rosenthal, Michael Pfaller)

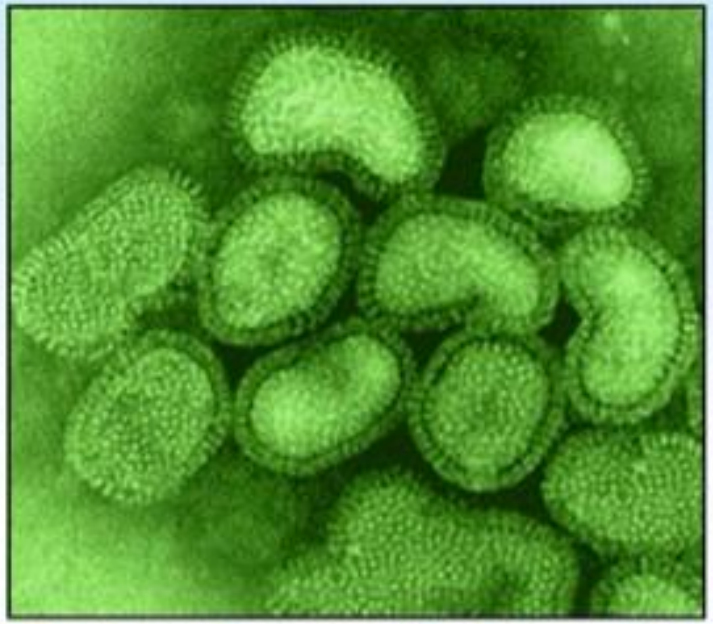
ORTHOMYXOVIRUSES



type A, B, C : **NP**, **M1** protein
sub-types: **HA** or **NA** protein

Influenza virus A

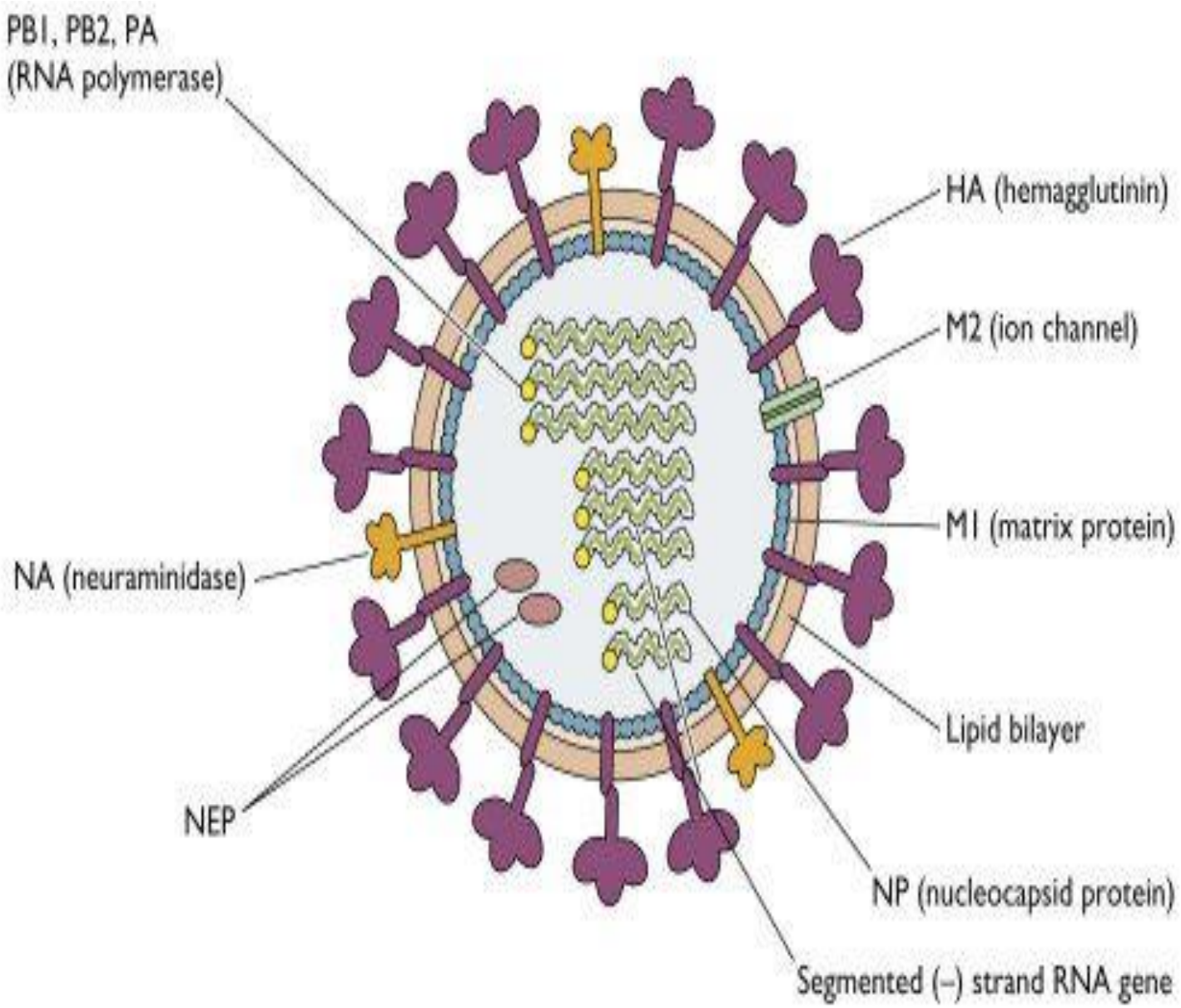




Orthomyxoviruses: medium-sized, enveloped, (-) sense that vary in shape from spherical to helical. Their genome is segmented into eight pieces

INFLUENZA VIRUS (STRUCTURE)

- The virion is **polymorphic**, mostly **spherical**, but sometimes rod-shaped. Influenza viruses, which vary in size in a wide range, are approximately **100 nm** in diameter.
- A nucleocapsid with **helical symmetry** is located in the center of the complex virion. In addition to ribonucleoprotein, nucleocapsid also includes three proteins of enzyme nature **(P1, P2 and P3).**
- The genome consists of **single-stranded segmented negative-RNA strand.** Influenza A and B viruses have **8** segments, and C virus has **7** segments.
- Nucleocapsid is surrounded by **matrix (M1)** and membrane or **ion channel (M2)** proteins.



INFLUENZA VIRUS (STRUCTURE)

The virion is surrounded by a lipoprotein membrane from the outside.
There are glycoprotein spikes on its surface. These spikes are composed of two complex glycoproteins: **hemagglutinin (H) and neuraminidase (N).**
Influenza type C virus does not have **neuraminidase.**

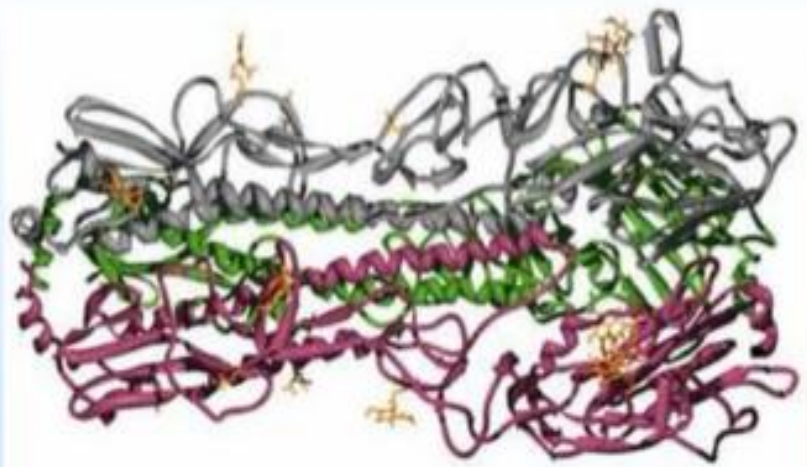
Haemagglutinin (HA)

Encoded by **RNA** segment # 4

Can agglutinate red blood cells - hence the nomenclature

Cleavage by host-cell protease is required (resulting in **HA1** and **HA2**) for infection to occur

Hemagglutinin glycoprotein is the viral attachment protein and fusion protein, and it elicits neutralizing, protective antibody responses



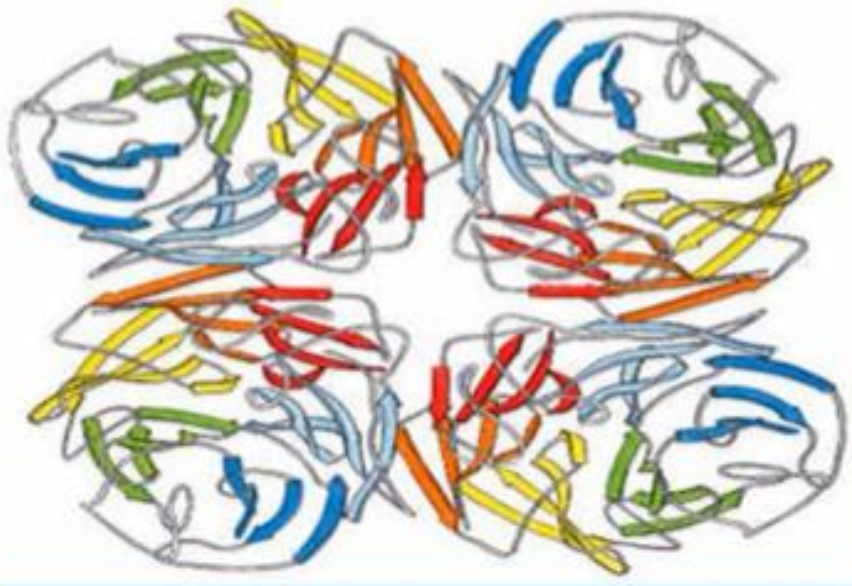
Neuraminadase (NA)

Encoded by **RNA** segment # 6

Removes neuraminic (sialic) acid from cell and permits dissemination of viruses

Important in releasing mature virus from cells

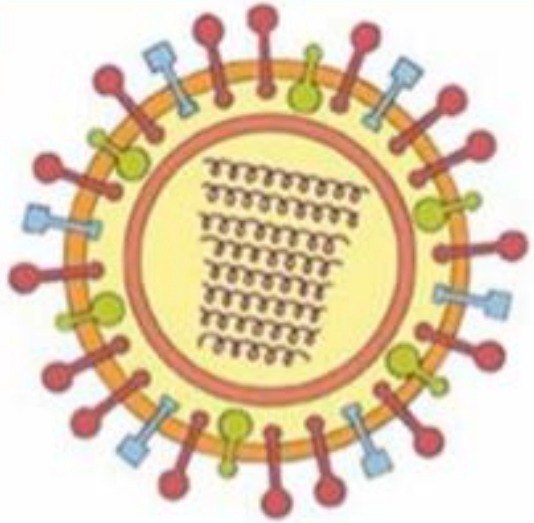
Stimulates production of protective antibodies



Orthomyxoviruses. Nomenclature

Human influenza virus

Influenza A/Bangkok/1/79(H3N2)
Influenza A/Singapore/1/57(H2N2)
Influenza B/Ann Arbor/1/86

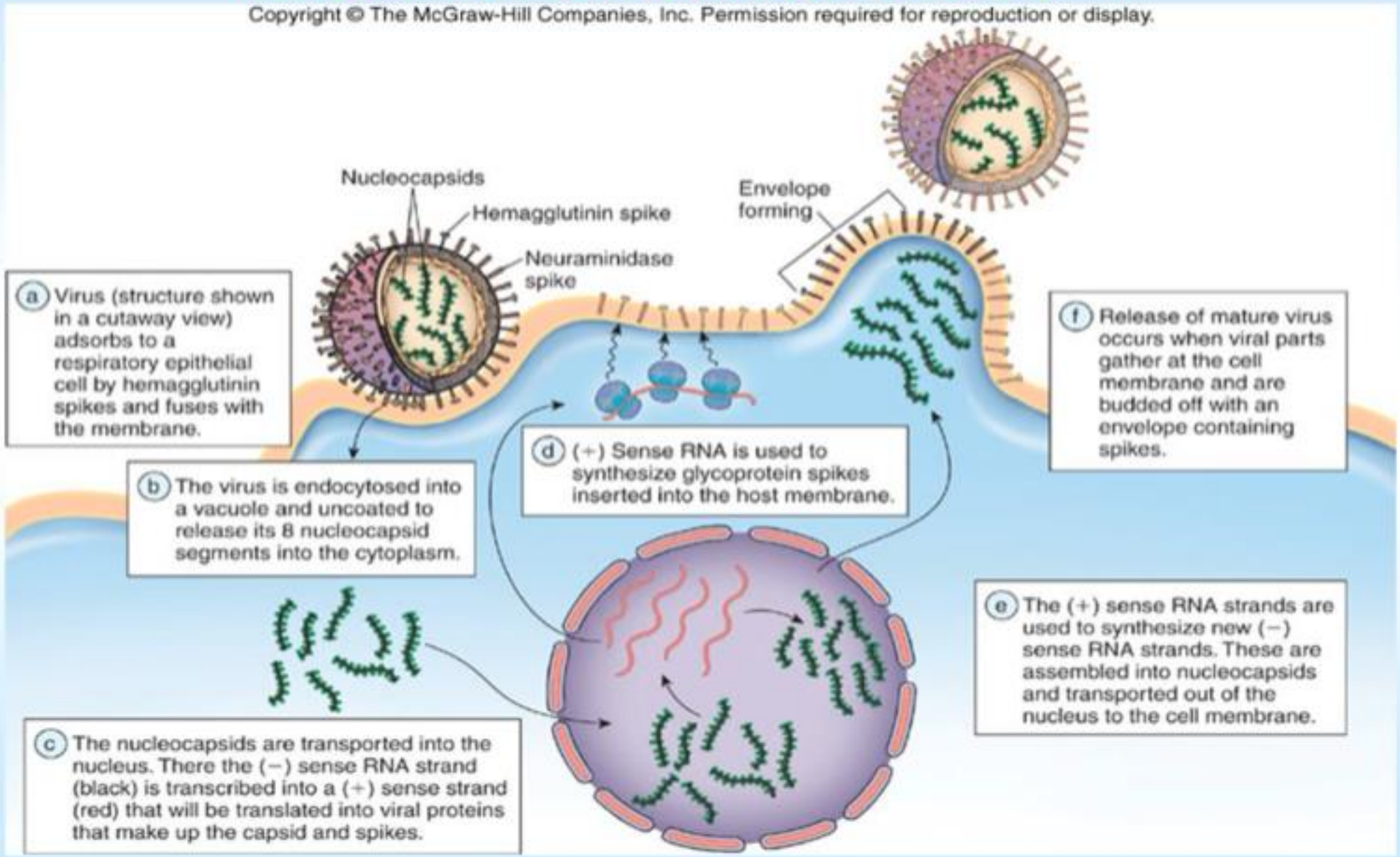


Influenza type — Year of isolation — Hemagglutinin subtype

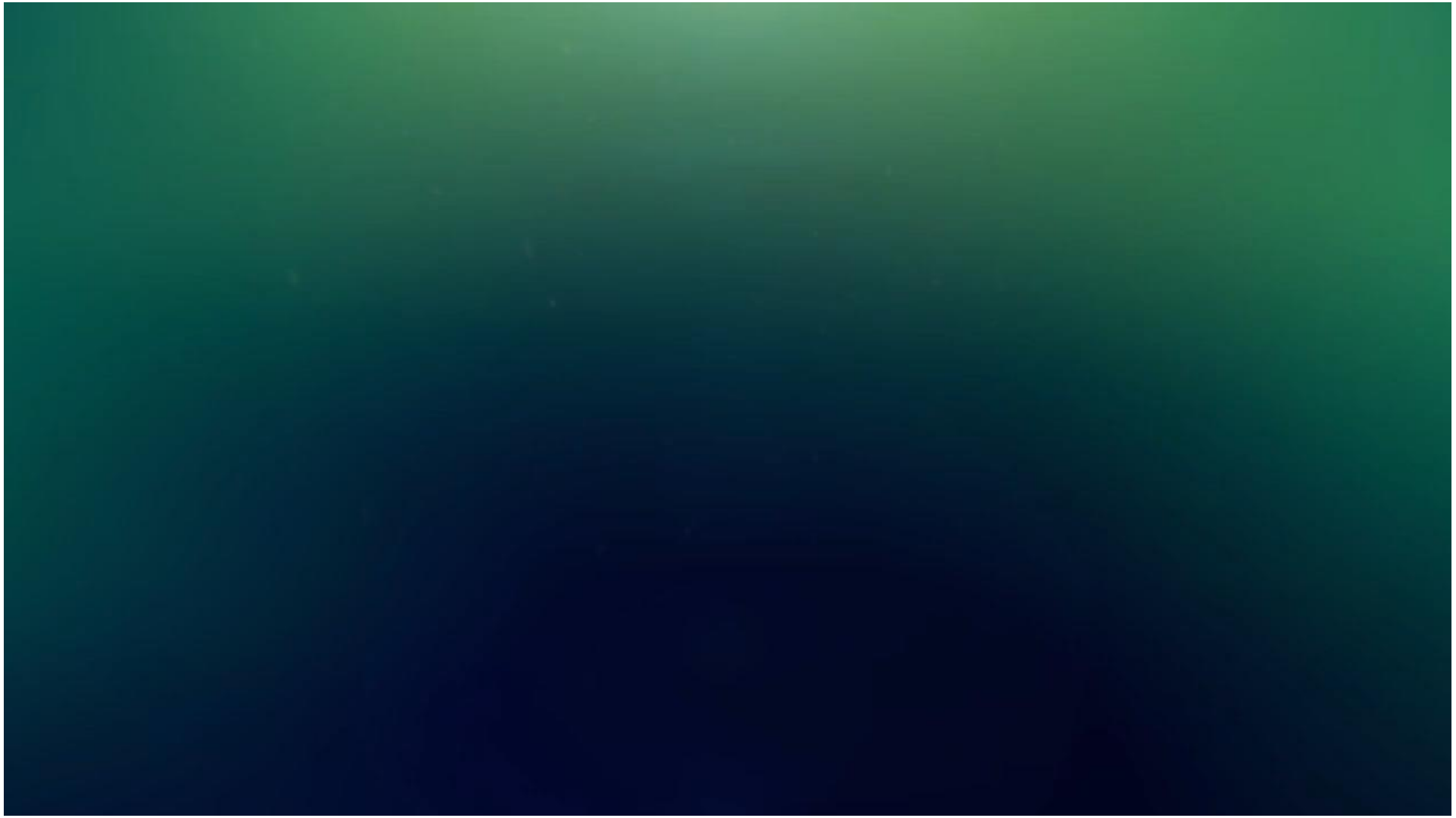
A/Sydney/5/97 (H3N2)

Geographic source — Isolate number — Neuraminidase subtype

Influenza virus reproduction

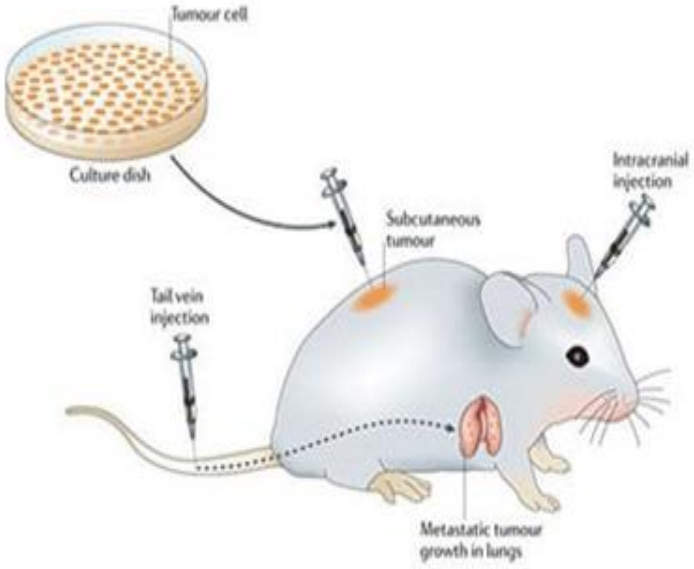


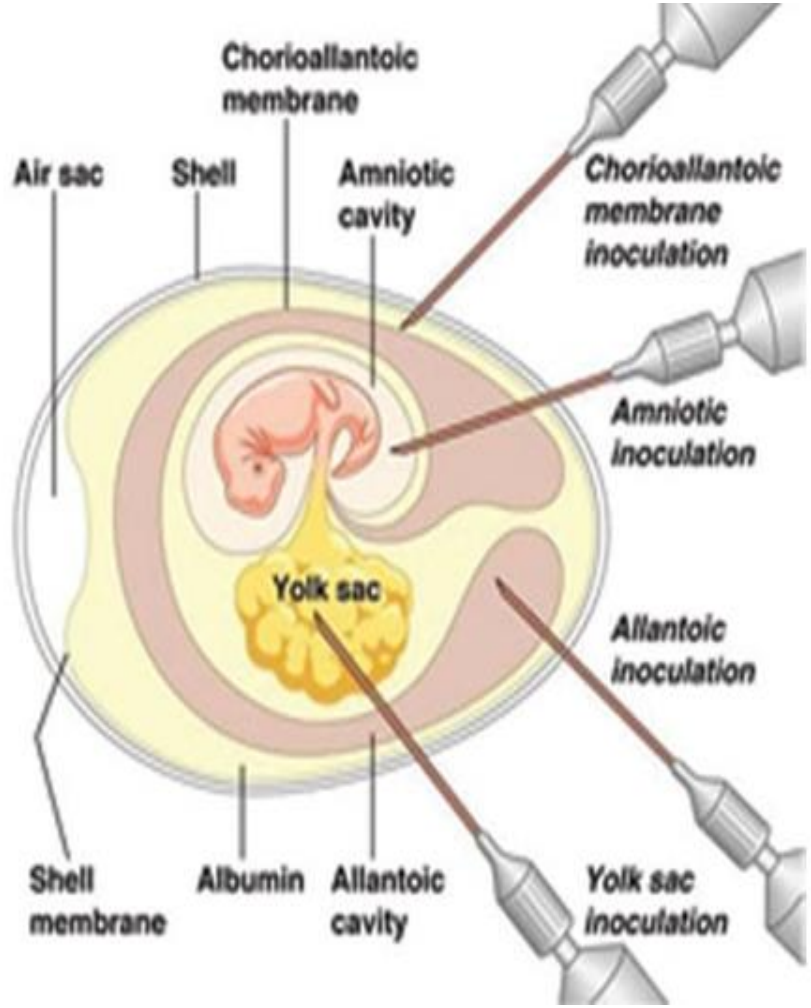
INFLUENZA VIRUS (REPRODUCTİON)



INFLUENZA VİRUS (CULTİVATİON)

- Chicken embryos are the optimal laboratory model for the cultivation of most strains of influenza viruses.
- Viruses can also be cultivated in **cell cultures** (primary cultures of monkey and dog kidney cells) and in **laboratory animals.**





Antigen

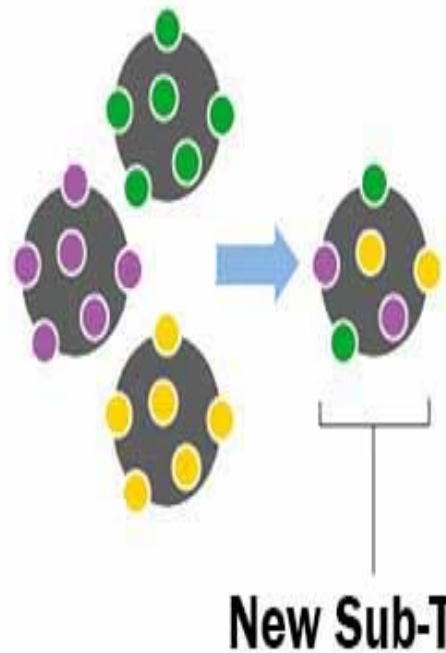
Influenza viruses are divided into 3 groups determined by the **ribonucleoprotein (RNP)** antigen and **M antigen**

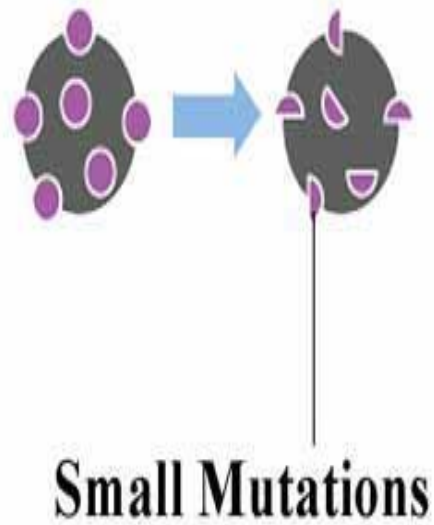
- Soluble antigens: include **ribonucleoprotein** and **M protein** which are much stable in antigenicity.
- Surface antigens: include **HA** and **NA** which are much variable in antigenicity.

Antigenic Shift	Antigenic Drift
Major change in genom	Minor change in genom
Gene reassortment	Point mutation
Happen accidently	Happen gradually
Only in A viruses	Both in A and B viruses
Result in new subtype	Result in new strain

Differences Between

Antigenic shift & Antigenic drift



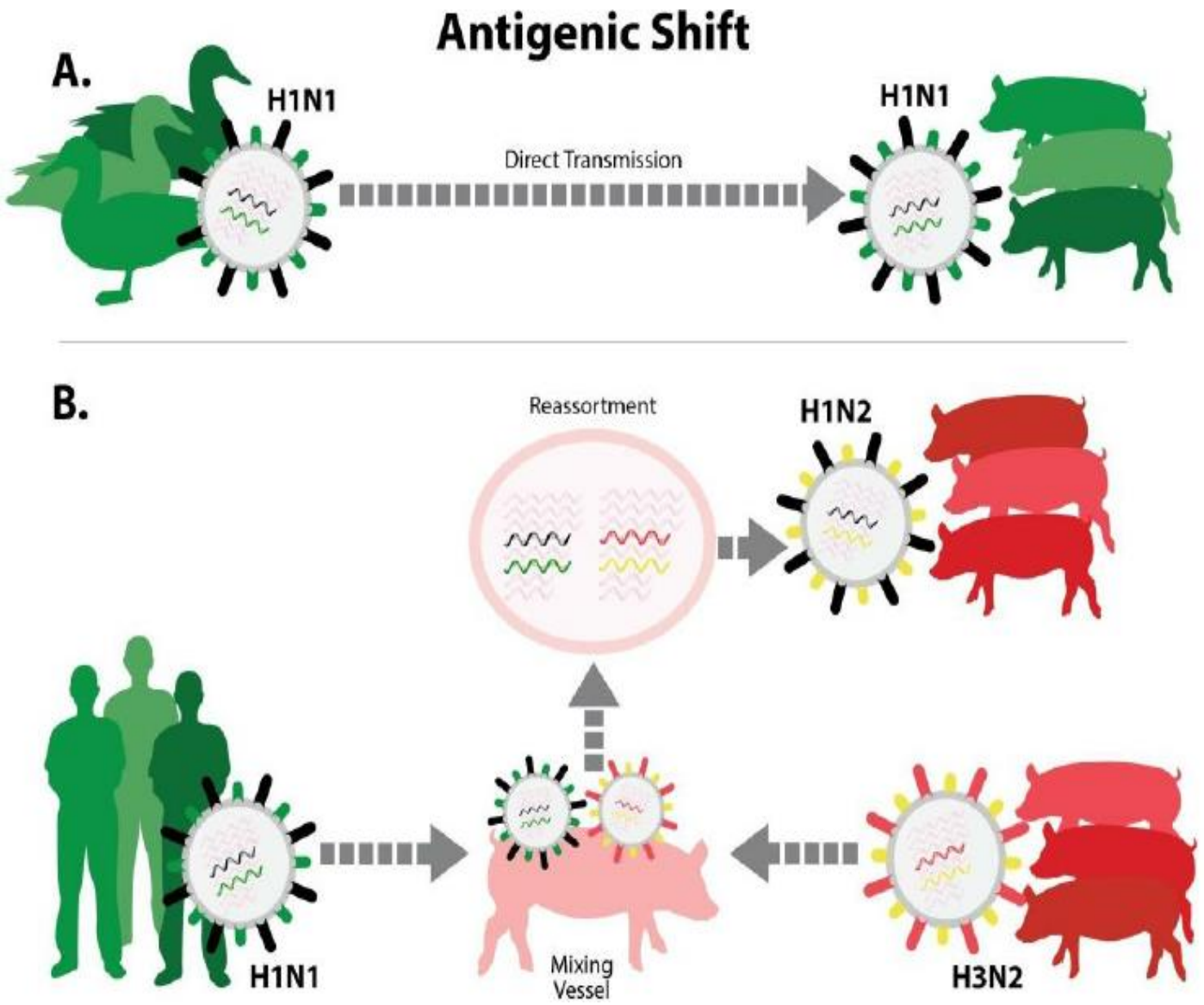


ANTİGENIC SHIFT

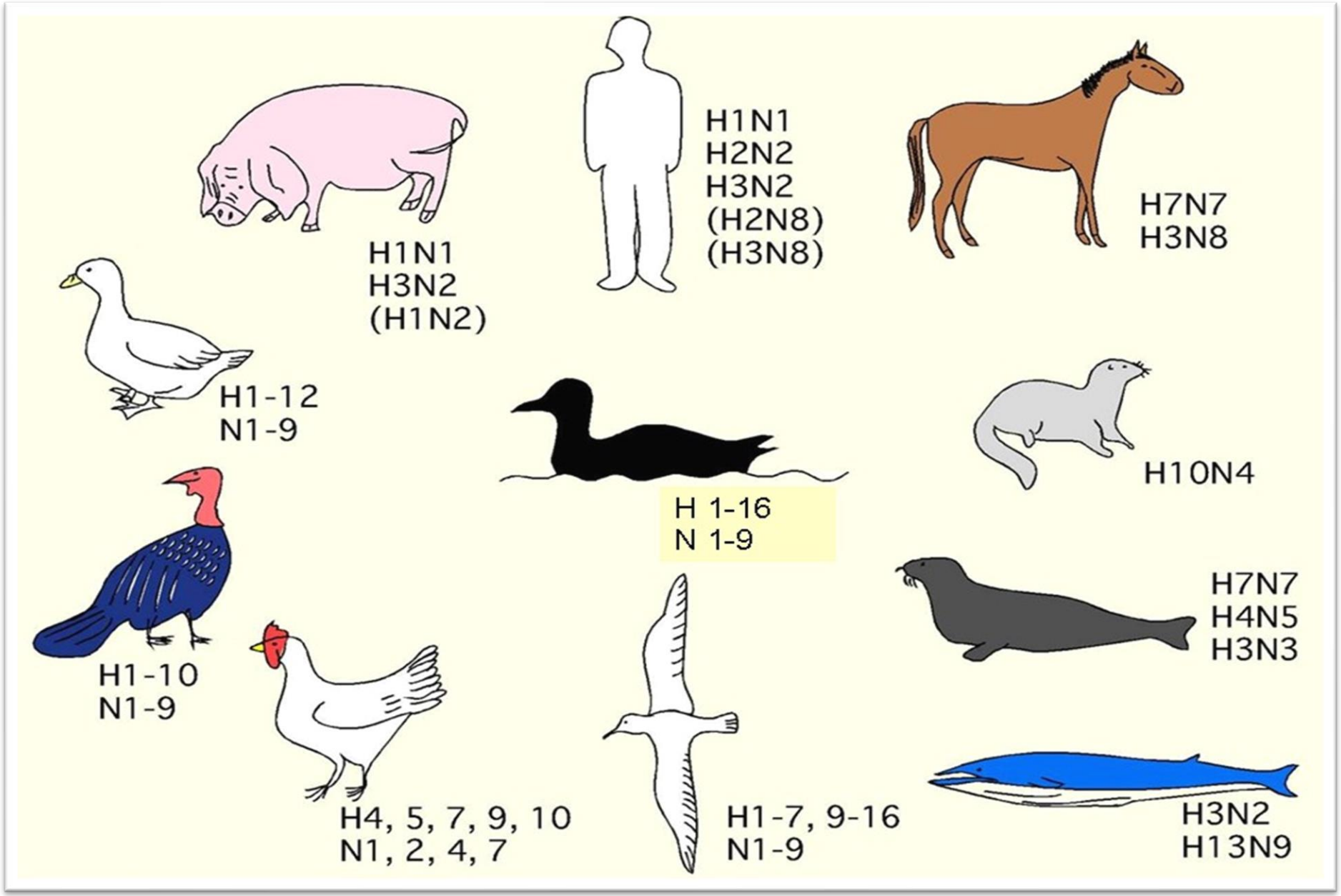
There are two ways that an influenza virus with new antigenic properties may enter the pig population.

(A) Virus that was previously adapted to another animal host, such as avian species, enters pigs and adapts to circulate efficiently in swine. The diagram portrays the inter-species transmission of an avian H1N1 virus, which became established in European swine populations;

(B) Virus previously adapted to another host, such as birds or humans, co-infects a pig along with a common swine-adapted strain. This can lead to gene reassortment, producing a new "reassortant" virus that contains an HA and/or NA antigenically different from those that previously circulated in swine. The diagram portrays reassortment between human seasonal H1N1 and swine H3N2 viruses. In both (A) and (B), the swine population lacks antibodies to important surface proteins of the new virus.

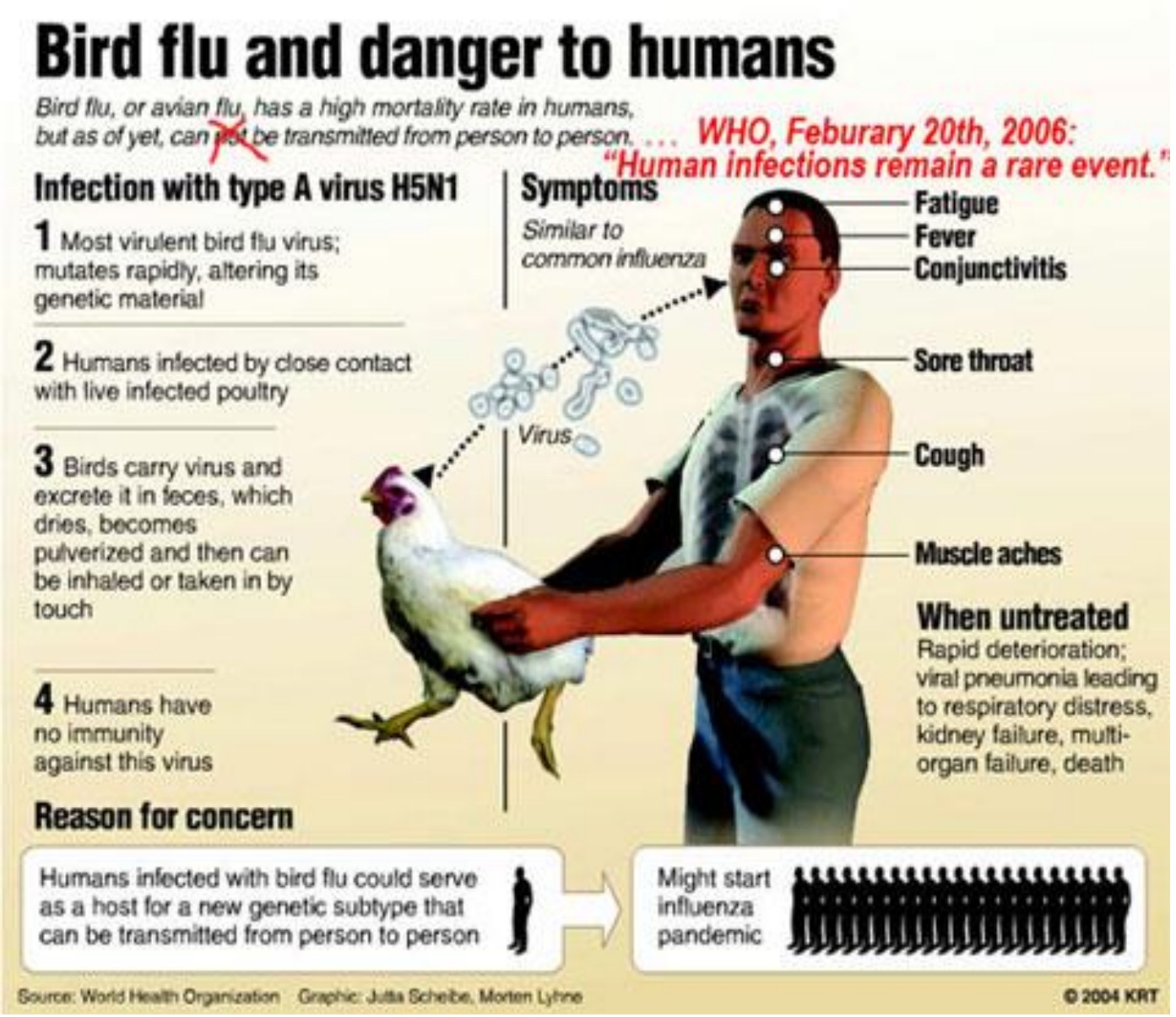


Occurrence of influenza virus subtypes in different organisms



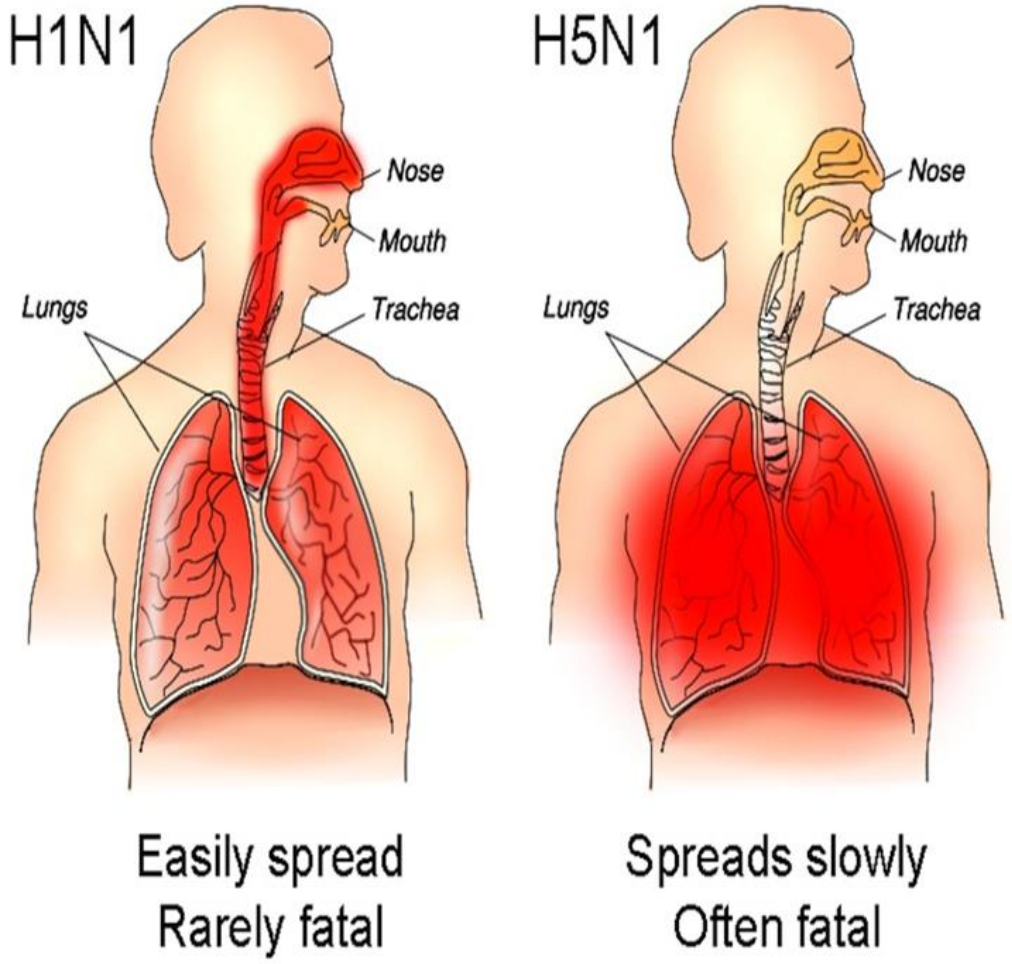
BIRD FLU (H5N1)

- In 1997, the first case of bird flu virus (H5N1 subtype of influenza A virus) was registered in Hong Kong. The source of infection was domestic birds. Avian influenza causes diseases of varying severity in some birds, from asymptomatic infections to lethal infections.
- Infection in ducks is usually asymptomatic. Influenza viruses in their bodies multiply in the intestinal epithelium, fall into the water in high concentrations with feces, and remain viable there for weeks. In this way, influenza viruses infect poultry and pigs.
- So far, pandemic influenza virus strains have emerged as a result of genetic sorting of avian and human influenza viruses. It is assumed that the genetic sorting of avian and human influenza viruses occurs in the body of pigs, since the body of pigs has receptors against both avian and human viruses.



SWINE FLU (H1N1)

♦ In 2009, in California - the first case of H1N1 subtype of influenza A virus (swine flu virus) was registered;
♦ the source of infection was pigs;
♦ swine flu - which quickly spread across most of the world's continents and caused many deaths - has become a pandemic;
♦ due to its antigenic structure - this virus, which is not different from the *«Spanish»* virus, differs in its high pneumotropism and causes high mortality rates mainly in elderly or weakened persons;
♦ unlike common flu viruses - "swine flu" virus is highly pathogenic and kills white mice quickly.

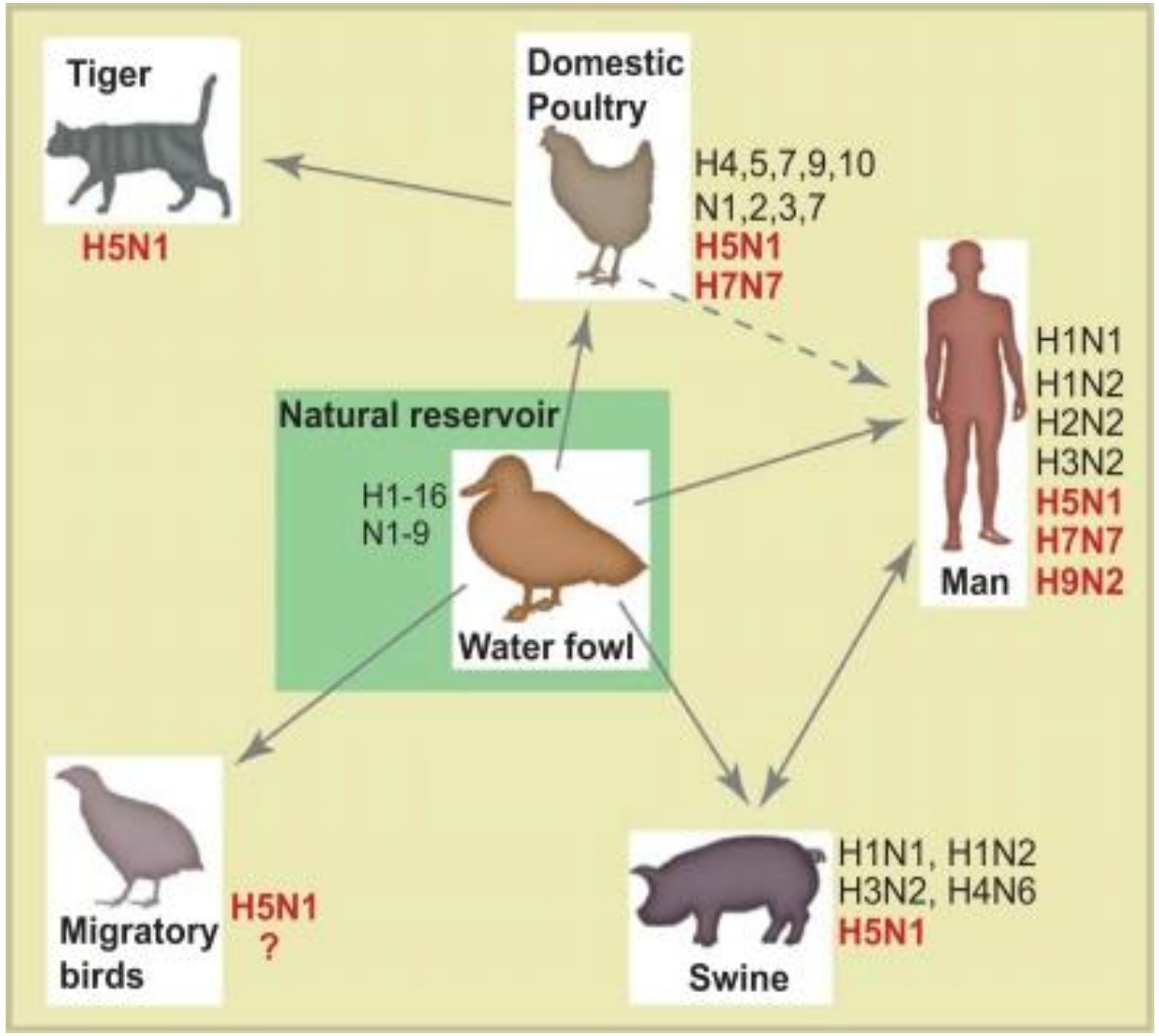


Resistance of Virus

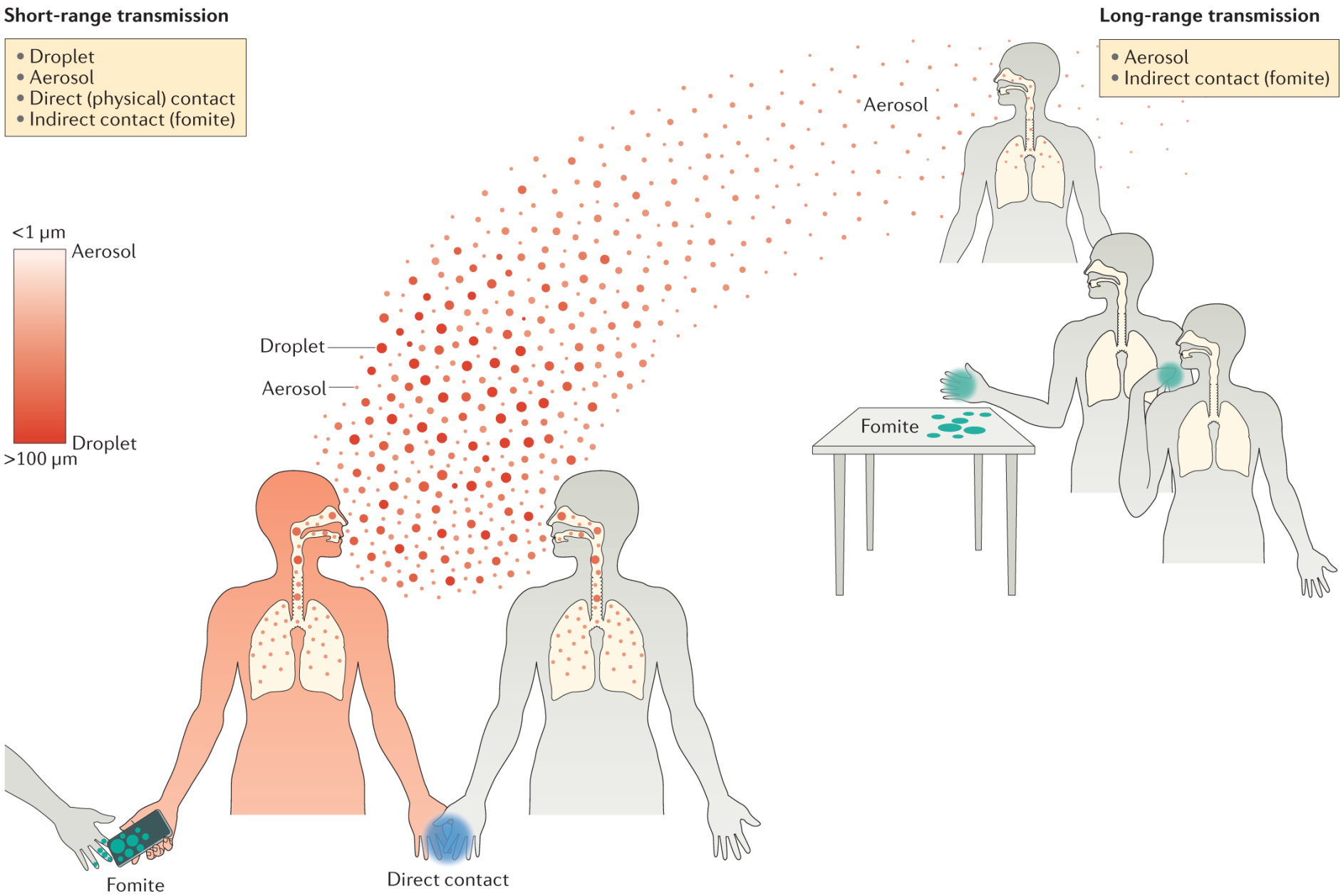
- Inactivated by heating at 50^0c for 30 mt
- Survive for 1 week at 0 – 4^0c for 1 week
- Virus preserved at – 70^0c
- **Survive in the blankets for 2 weeks**
- Ether, formaldehyde, Phenol destroy the virus

SOURCE OF INFECTION AND MODE OF TRANSMISSION:

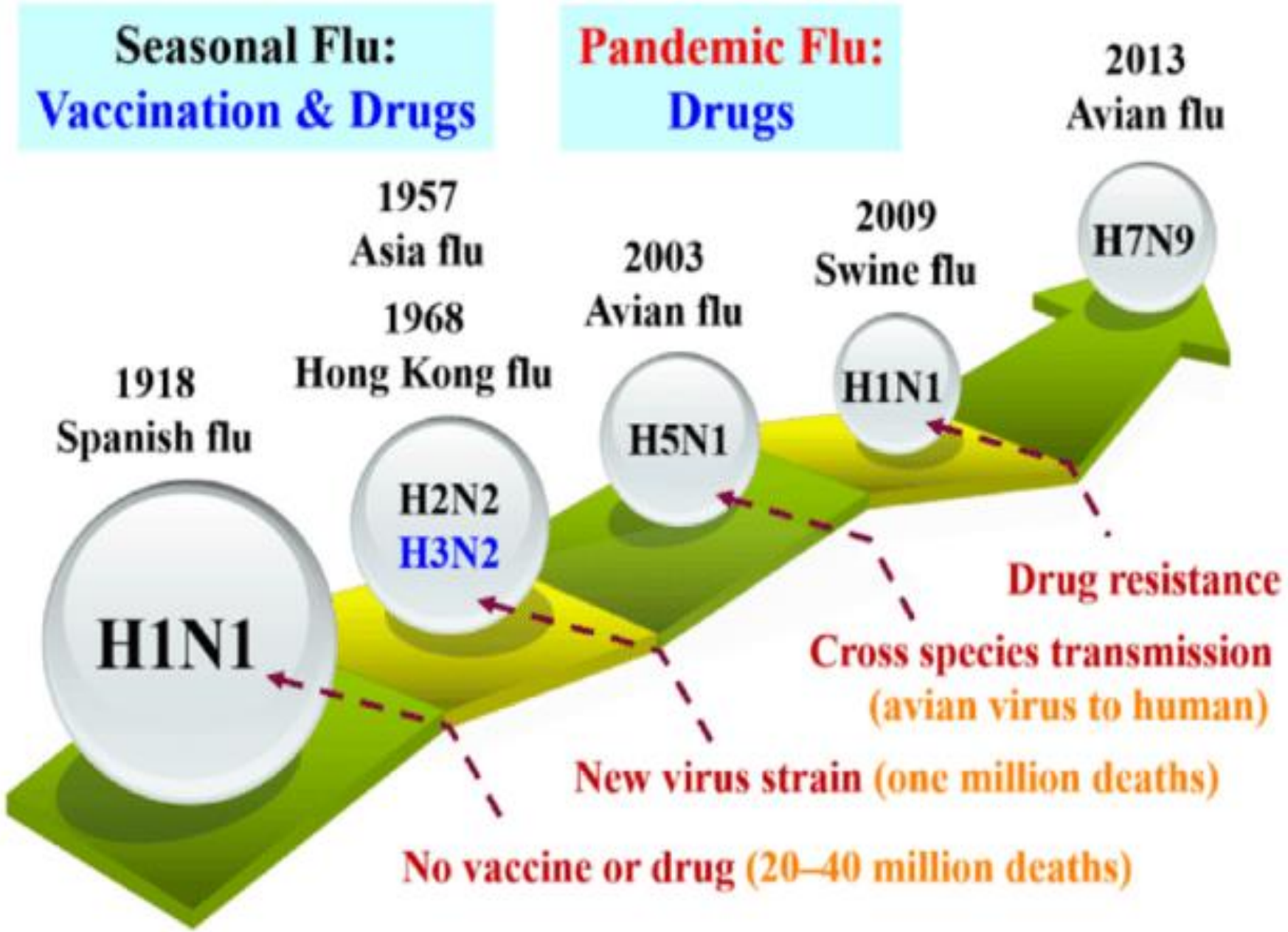
- The source of infection is **sick people,** sometimes **birds and animals**. People are very susceptible to the flu virus. Infection occurs mainly through **air droplets** (when coughing, sneezing, talking).
- Type A influenza virus periodically causes pandemics. Southeast Asia (China) is the epicenter of the emergence of new pandemic strains of influenza A viruses. Here, high compact population, close contact with domestic animals and birds create conditions for the recombination of human and animal viruses.
- Since the end of 2005, the "bird flu" caused by the H5N1 subtype of influenza A virus, and the "swine flu" caused by the H1N1 subtype have started in the world since 2009.



TRANSMISSION



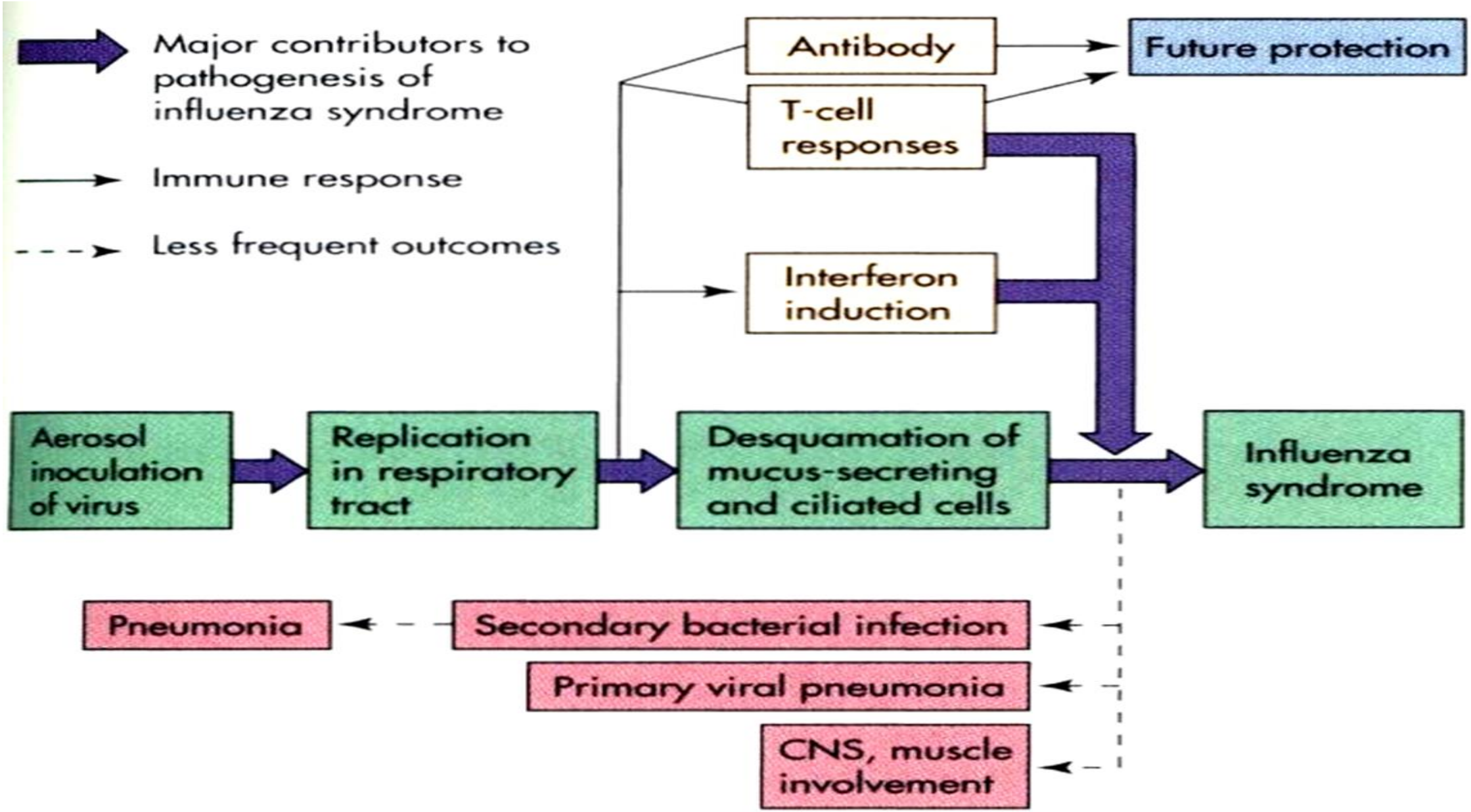
EPİDEMİOLOGY



Pathogenesis of influenza:

- The primary reproduction of the influenza virus that has entered the body occurs in the **epithelial cells** of the **upper respiratory tract** (sometimes lung alveoli). As a result of the destruction and desquamation of the surface epithelium, inflammation and edema develop here, but the basal membrane of the epithelium is not damaged.
- Some subtypes of influenza viruses can immediately enter the alveoli and cause the development of primary acute pneumonia (atypical pneumonia). This is one of the main causes of death in high-risk patients
- **The virus is rarely transmitted to the blood.**
- Damage to the epithelium of the respiratory tract creates conditions for the development of secondary bacterial infections, especially caused by staphylococci, streptococci (pneumococcus) and hemophilic bacteria.
- During the flu, transient secondary immunodeficiency develops, which creates conditions for the development of secondary bacterial infections.
- Secondary bacterial **pneumonias** are also one of the main causes of death.

Pathogenesis



Clinical manifestations of influenza:

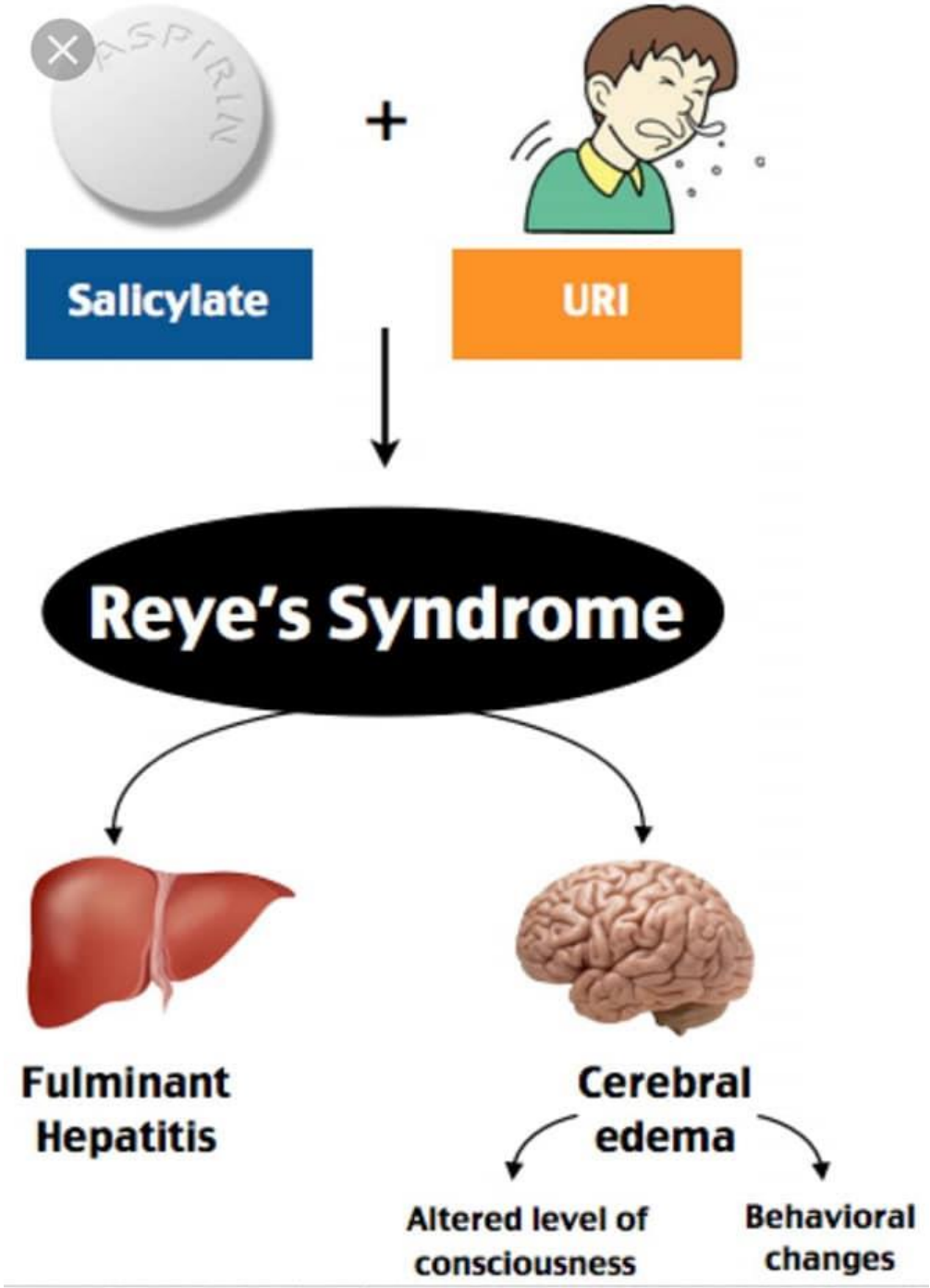
•The **latent period** of influenza can last from **1 to 4 days**.

•**Uncomplicated flu.** The disease begins acutely, usually with high fever, headaches, joint and muscle pain felt throughout the body, and weakness. Catarrh of the upper respiratory tract - cough, back pain, rhinitis and rhinorrhea develop.

Complications of the flu

•**Pneumonia**, the most serious complication of influenza in the elderly and debilitated with chronic diseases, as well as in pregnant women, is one of the main causes of death during influenza. Viral pneumonia, secondary bacterial pneumonias, as well as mixed viral-bacterial pneumonias are possible during influenza. Bacterial pneumonias are most commonly caused by *S.aureus*, *S.pneumoniae and H.influenzae*.

•**Reye's syndrome** is an acute encephalopathy that occurs in children and teenager aged 2-16 years and is rarely observed. Mortality is approximately 10-40%. The higher occurrence of Reye's syndrome after administration of salicylates in these infections suggests a possible relationship between salicylates (**aspirin**, etc.) and this syndrome.



Immunity in Influenza

Immunity to an influenza virus is type-specific and lasts for many years.
Recurrent cases of influenza are caused primarily by antigenically different strains.

- Antibody to HA - >protective
- Antibody to NA - > decrease severeity
- Serum antibody - > years
- Secretory antibody - > months

Influenza Diagnosis

- Clinical and epidemiological characteristics
- Isolation of influenza virus from clinical specimen (e.g., nasopharynx, throat, sputum)
- Significant rise in influenza IgG by serologic assay
- Direct antigen testing for type A virus

Microbiological diagnosis of influenza

Examination material - during the first three days of the disease, it is possible to obtain viruses as a result of the examination of the materials taken from the nose, or pharynx, as well as swabs from the sputum. Sometimes, nasal mucosal impression smear are studied.

Virological

Materials intended for virological examination should be stored at $+4^0$C until examination. Freezing reduces the possibility of detecting the influenza virus, if the examination is to be carried out later than 5 days, the material is stored frozen at -70^0C. Chicken embryos are mostly used for cultivation. In cell cultures, the virus can be indicated by the hemadsorption test 3-5 days after inoculation, and in the culture fluid after 5-7 days by the hemagglutination (**HAR**) reaction. The subtype of the acquired virus is determined by the hemagglutination inhibition reaction (**HIR**), and the type is determined by **CFT**.

Express diagnostics

Virus antigens can be detected in the examined material by **IFR** (direct and indirect variant), but it has poor sensitivity compared to the virological method.

It is possible to determine the viral genome in the material by **PCR**. In 2006, Real-time reverse transcriptase PCR was proposed to detect the "bird flu" virus (A/H5N1).

Serological

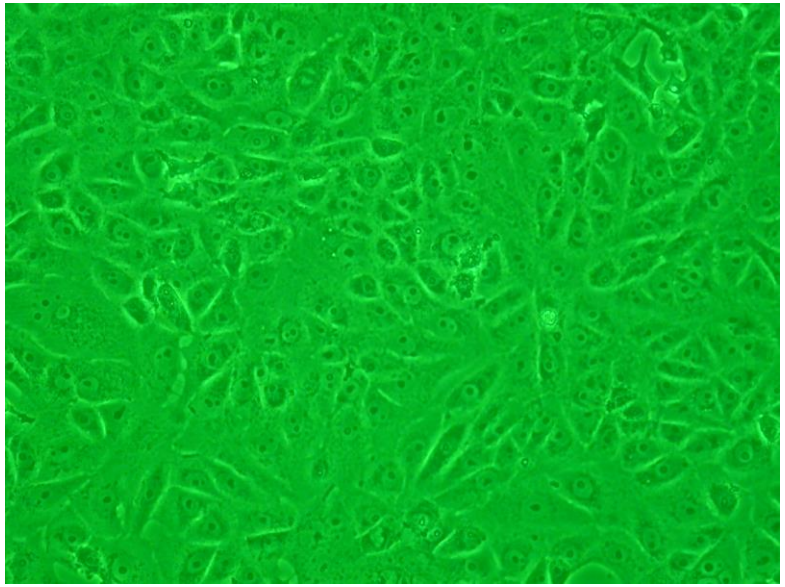
Preliminary serodiagnosis of influenza is carried out by **HIR** and **ELISA**. Since the blood serum of healthy people can contain antibodies against influenza viruses, paired blood sera of the patient (taken at an interval of 10-14 days during the acute period of the disease and during the convalescence period) are studied.

A four-fold increase in the titer of antibodies in the blood serum confirms the diagnosis. The serological method is often used for retrospective diagnosis.

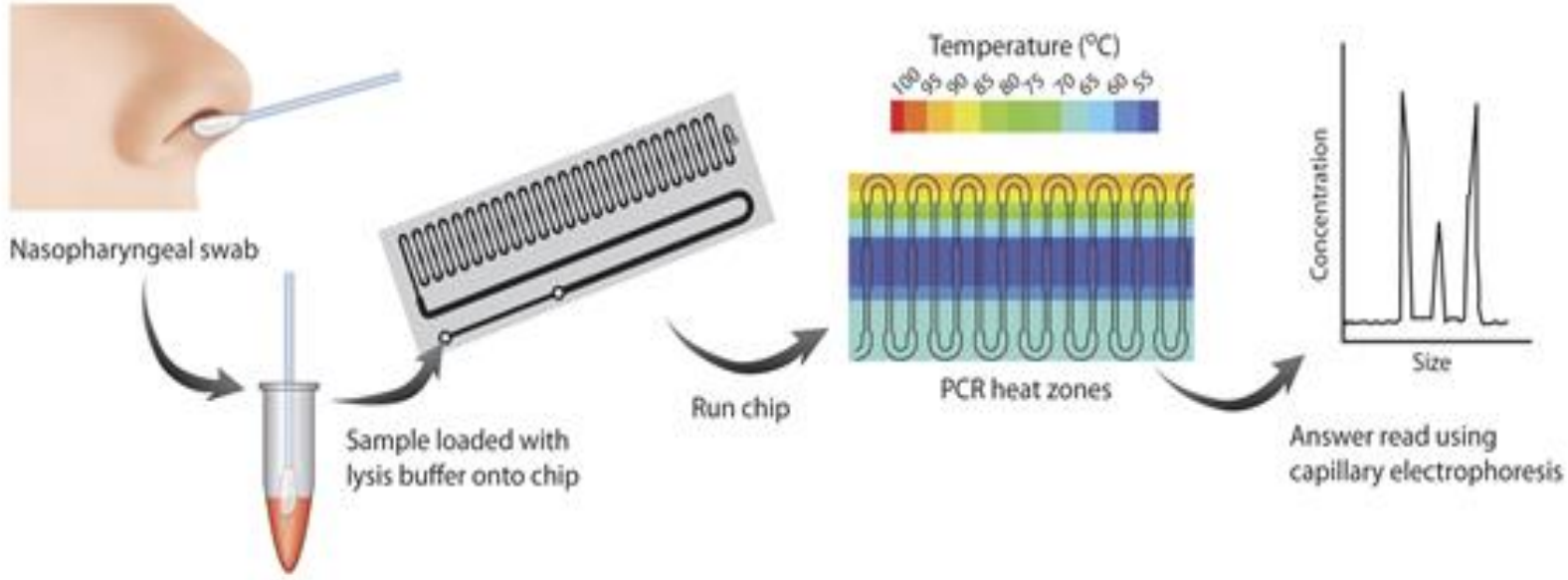
Microbiological diagnosis of influenza

Examination material - during the first three days of the disease, it is possible to obtain viruses as a result of the examination of the materials taken from the nose, or pharynx, as well as swabs from the sputum. Sometimes, nasal mucosal impression smear are studied.

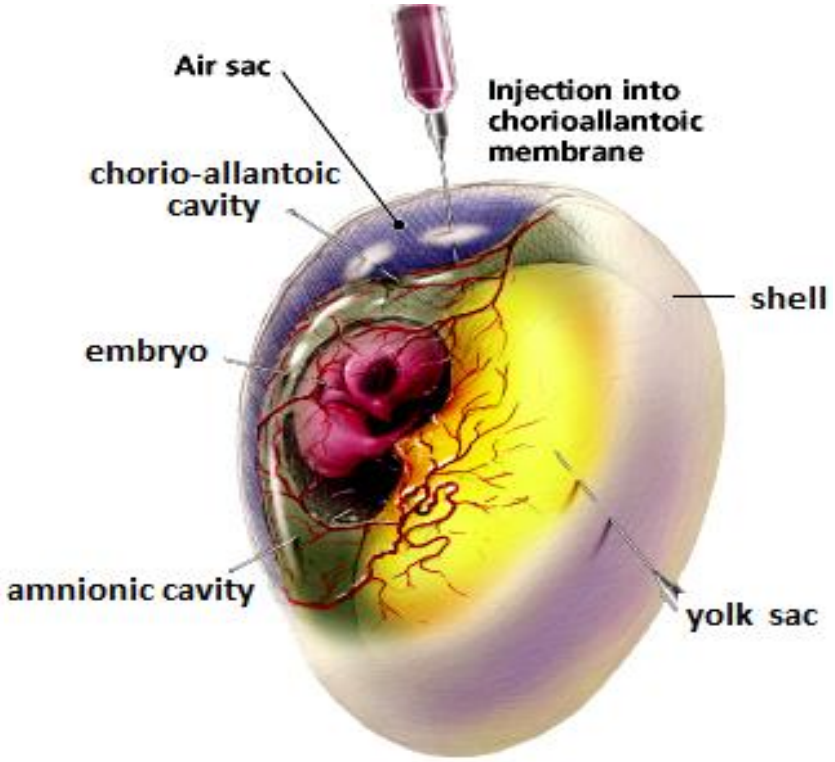
Virological

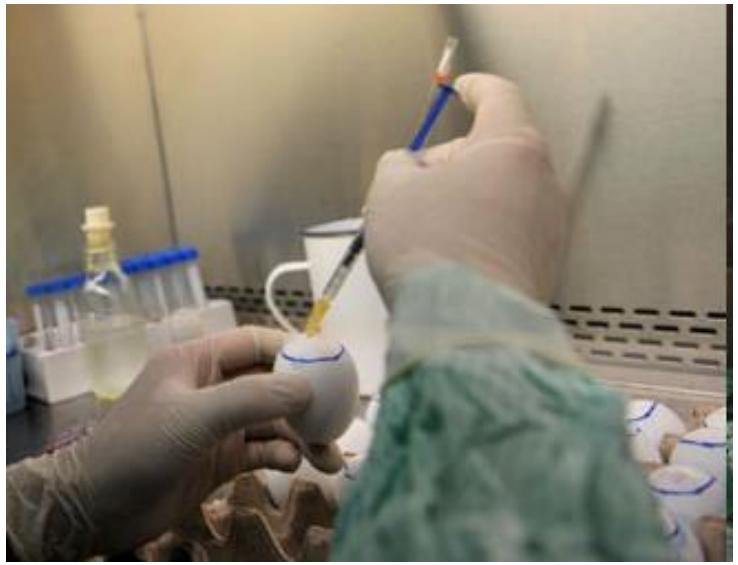


Express diagnostics

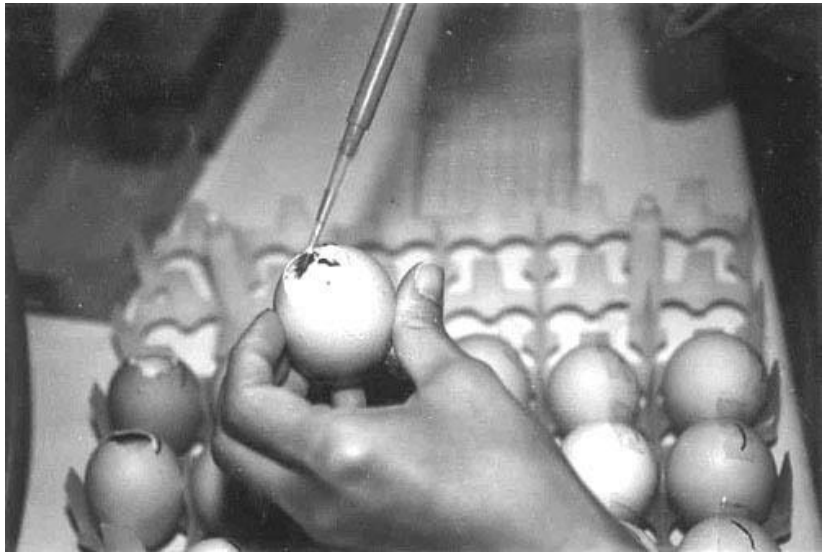


Chick embryo culture method





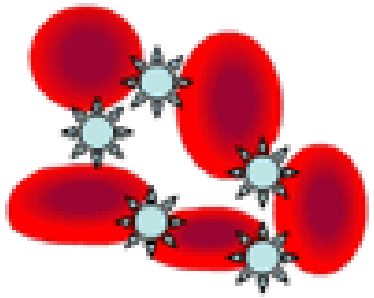
Inoculation of chick embryo



Removing allantoic fluid

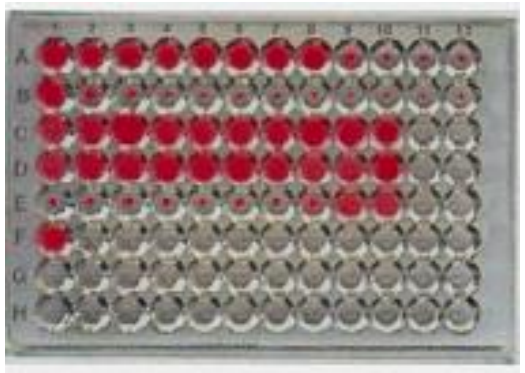
- Fluid from the amniotic or allantoic cavity of chick embryos is tested for the presence of newly formed viruses by haemagglutination test;
- the virus in positive fluids is then identified by haemagglutination inhibition test with specific antisera.

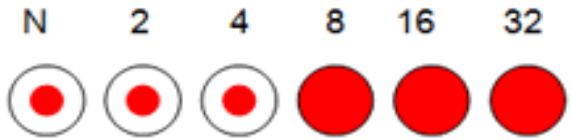
Haemagglutination inhibition test (HAI)



Influenza viruses bind to red blood cells using the haemagglutinin causing the formation of a lattice.

HA: two-fold serial dilutions of a virus are prepared, mixed with red blood cells, and added to the wells of a plastic plate. The red blood cells that are attached to virus particles form a lattice that coats the well. The red blood cells that are not bound by virus sink to the bottom of a well and form a button.

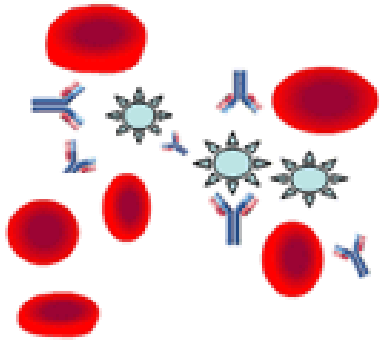


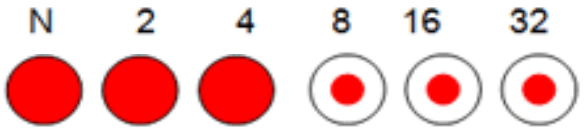


negative HA positive HA

The basis of the HAI assay is that antibodies to influenza virus will prevent attachment of the virus to red blood cells.

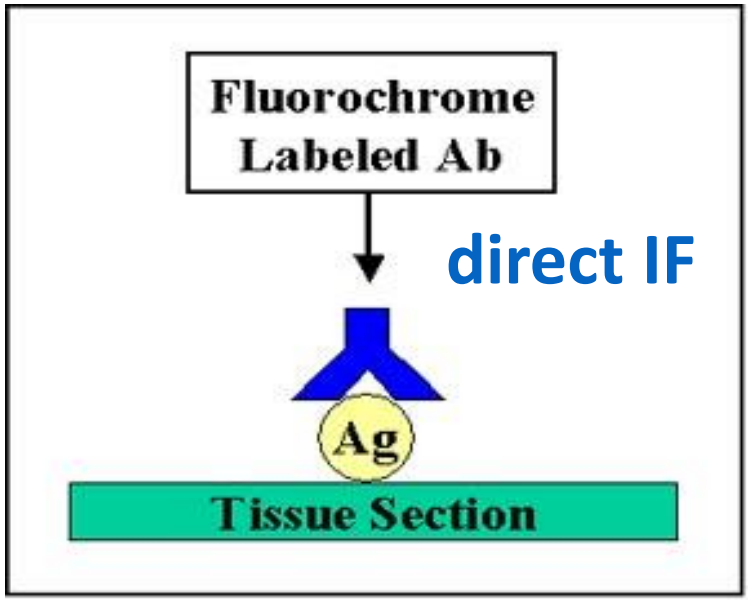
By adding specific antibodies to the virus it is possible to block this interaction and detect the virus. If antibodies to the virus are specific, hemagglutination will not be observed.

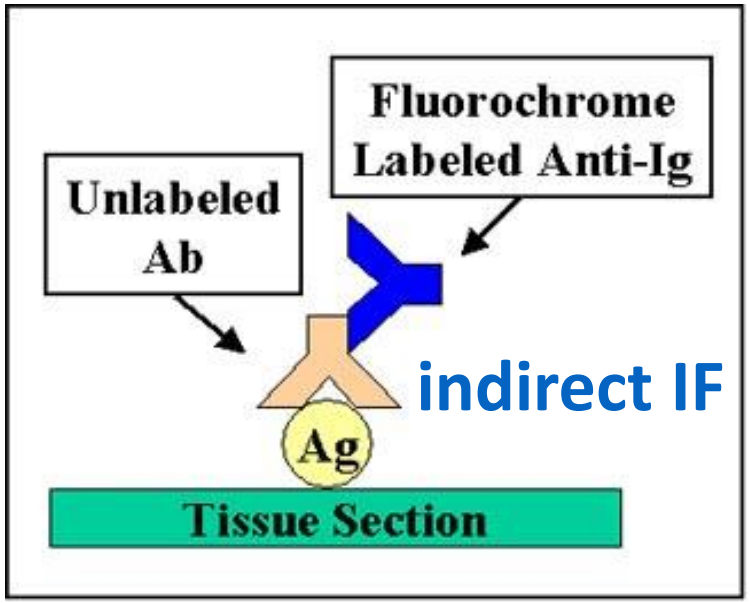


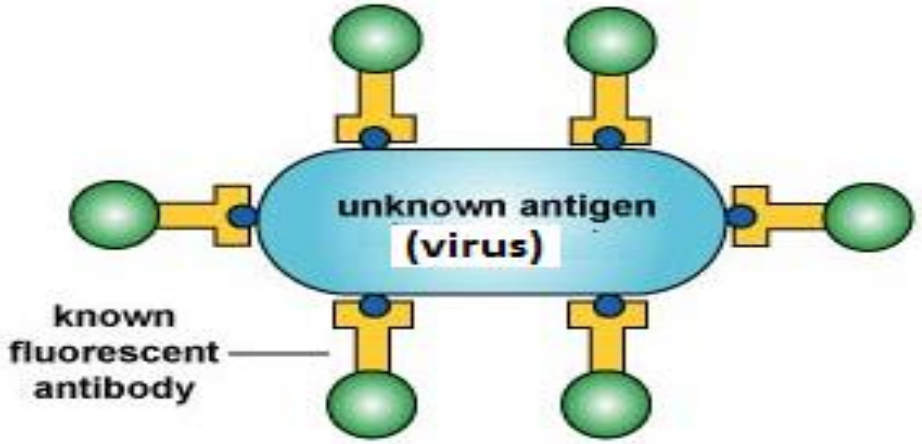


negative HAI positive HAI

Express diagnostics
Immunofluorescence (IF) to detect virus into host cells







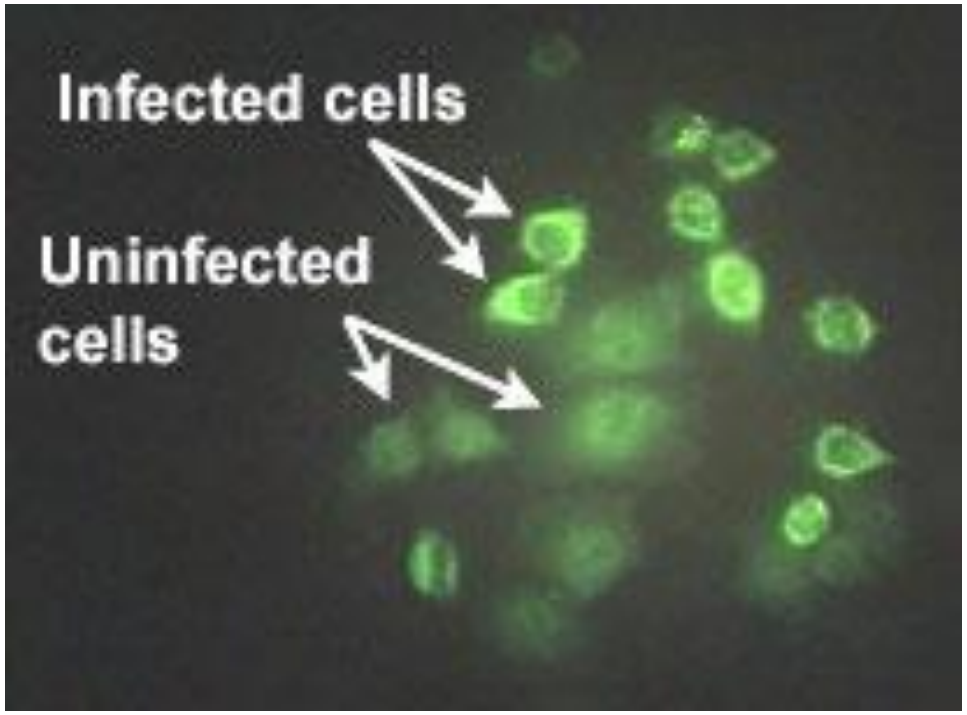


TABLE 61-6 Laboratory diagnosis of influenza virus infection

Method	Test	Detection
Direct antigen detection	IF, ELISA	Influenza virus antigen in respiratory secretions
Virus isolation		
Cell culture	Primary monkey kidney cell, Madin-Darby canine kidney cell, and hemadsorption to infected cells	Limited cytopathic effects Presence of HA protein on cell surface
Chick embryo	Allantoic and amniotic cavity	Fluid is tested for hemagglutination of virus
Serology	Hemagglutination inhibition, Hemadsorption inhibition, ELISA, CFT, and IF	Demonstration of a rise in serum antibody titer
Molecular diagnosis	RT-PCR	Viral nucleic acid in the nasopharyngeal cells

ELISA, enzyme-linked immunosorbent assay; CFT, complement fixation test; IF, immunofluorescence; RT-PCR, reverse transcriptase-polymerase chain reaction.

TREATMENT

- **RIMANTADINE** (blocks the M2 ion channel) (M2)
  - type A only, needs to be given early
- **AMANTADINE** (blocks the M2 ion channel) (M2)
  - type A only, needs to be given early
- **ZANAMIVIR** (neuraminidase inhibitors) (NA)
  - types A and B, needs to be given early
- **OSELTAMIVIR** (neuraminidase inhibitors) (NA)
  - types A and B, needs to be given early

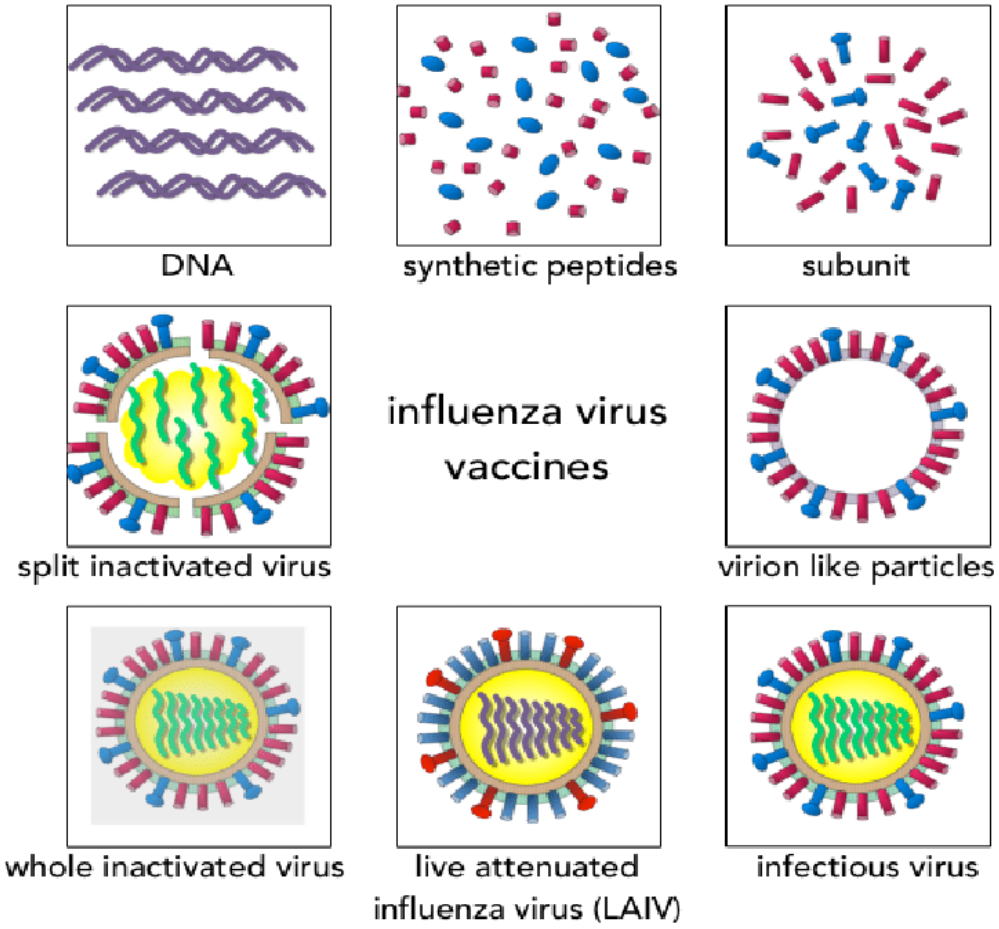
Prophylaxis
Masks and Hand Washing

- *To be Continued…*

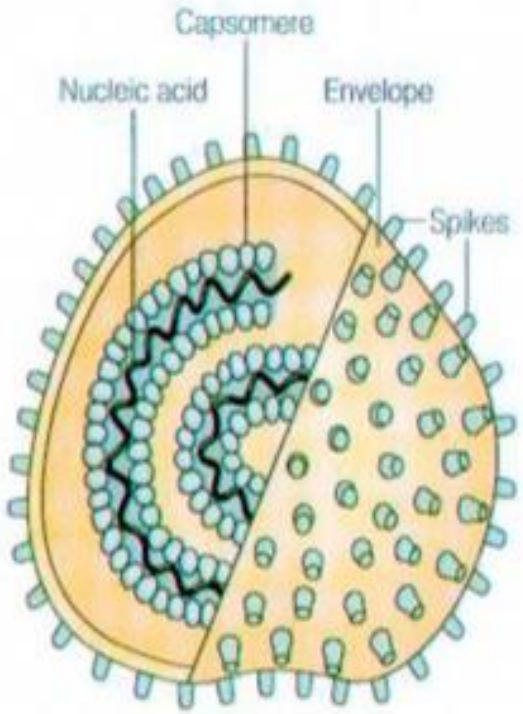
- Hand washing
  - Generally perceived to be useful
  - No studies specifically performed for influenza
  - Easy to recommend
- Masks
  - Effectiveness not shown for influenza
  - However, could reduce transmission associated with large droplets

VACCINES

- Rimantadine can be used for emergency **chemical prevention** during flu epidemics.
- Various vaccines are used for **specific prevention**. Inactivated (killed) and live vaccines are available.
- **Whole virion vaccines** consist of inactivated viruses
- **Subvirion vaccines** - consist of virus particles that have been broken down by detergents.
- Vaccines made from **surface antigens** include purified H and N glycoproteins.
- **Live (live, attenuated) vaccines**. Thermolabile mutants of the influenza virus have recently been obtained that can growth at 25^0C but cannot growth at body temperature (37^0C). Such viruses can replicate in the nasopharynx, where the temperature is relatively low (33^0C), but cannot replicate in the lower respiratory tract. Live vaccines made from such viruses are administered as an intranasal spray.



Influenzavirus B



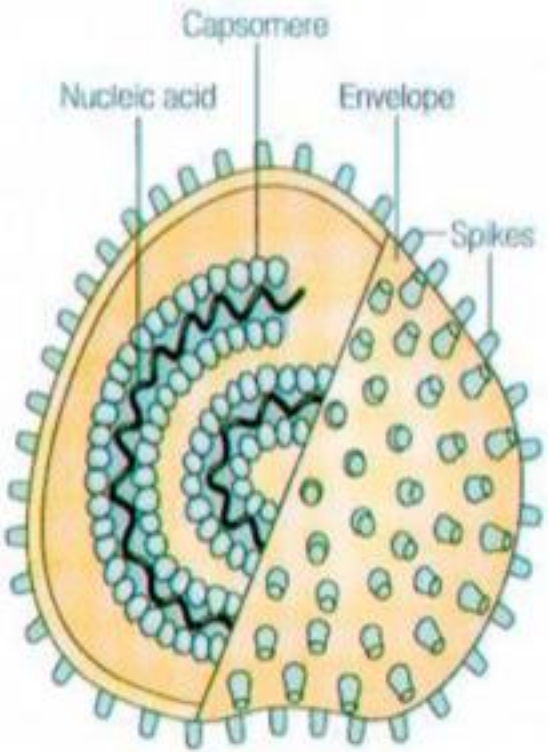
(a) An enveloped helical virus

(b) An *Influenzavirus*

- **Virions enveloped**
- **About 500 spikes**
- **Nucleocapsid enclosed within lipoprotein membrane**
- **Virions contain 8 segments of linear negative-sense single stranded RNA**
- **Total genome length is 13588 nt**
- **The largest segment 2341 nt**

- Infect much man and birds.
- Cause human disease but generally not as severe as A types.
- Believed to be epidemiologically important - reassortment with type A leads to epidemics.

Influenzavirus C



(a) An enveloped helical virus

(b) An *Influenzavirus*

- Virions enveloped
- Many spikes
- Nucleocapsid enclosed within lipoprotein membrane
- Virions contain 7 segments of linear negative-sense single stranded RNA
- Total genome length is 12900 nt
- Glycoprotein
- -hemagglutinin esterase fusion (HEF)
- esterase -> receptor destroying enzyme


return
L
Continue

Family Paramyxoviridae

Subfamily Paramyxovirinae:

Genera:

- Morbillivirus – measles virus,
- Respirovirus (earlier Paramyxovirus) – parainfluenza virus serotypes 1 and 3
- Rubulavirus - parainfluenza virus serotypes 2, 4a, 4b, mumps virus
- Henivirus – Australian Hendra-virus (diseases of human and horses), Malasian Nipah-virus (diseases of human and swine)

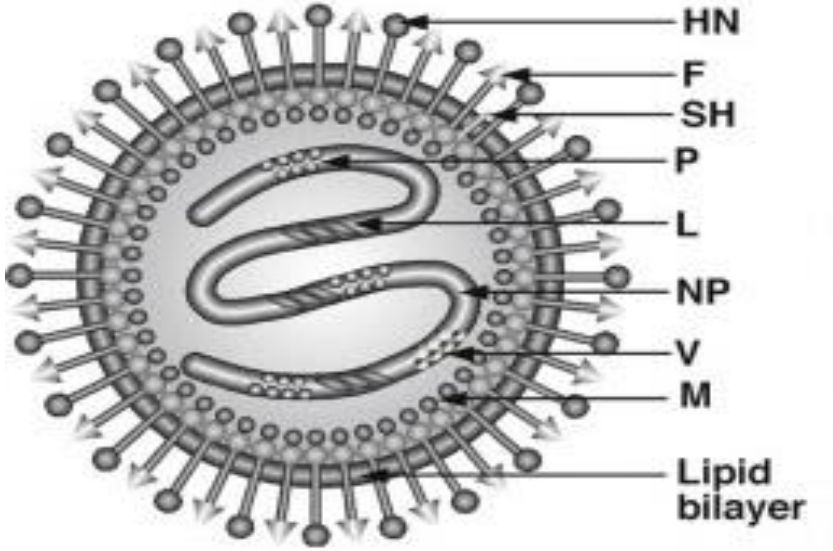
Subfamily Pneumovirinae

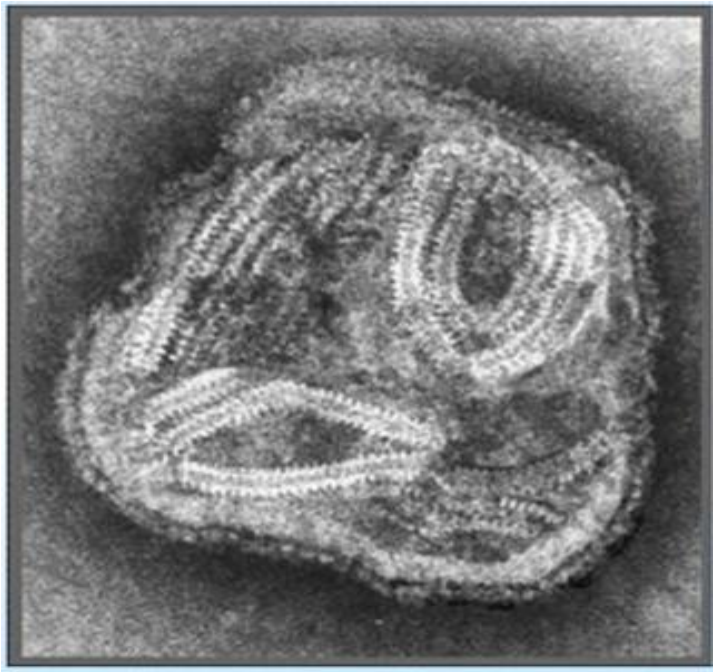
Genera:

- Pneumovirus – RS-virus
- Metapneumovirus – human metapmeumovirus (diseases in children)

Paramyxoviruses:

- **Structure**. The virions of *Paramyxoviruses* are **enveloped**, **polymorphic**, 150 nm and larger (sometimes 700 nm).
- The genome of the virus consists of **linear single-stranded RNA**, which combines with a number of proteins to form a nucleocapsid with **helical symmetry.**
- The lipid virion membrane has two types of transmembrane glycoprotein spikes: one of them consists of three glycoproteins **(HN, or H, or G)** with hemagglutinin and/or neuraminidase activity.
- The activity of these glycoproteins, which ensures the virus's connection with the host cell, allows different types of the family to be differentiated**. It is found in HN - parainfluenza and mumps, H - measles virus, G - respiratory syncytial virus.**
- Another glycoprotein consists of **fusion protein F-protein** (in English, fusion), connects cell membranes and has hemolytic activity.

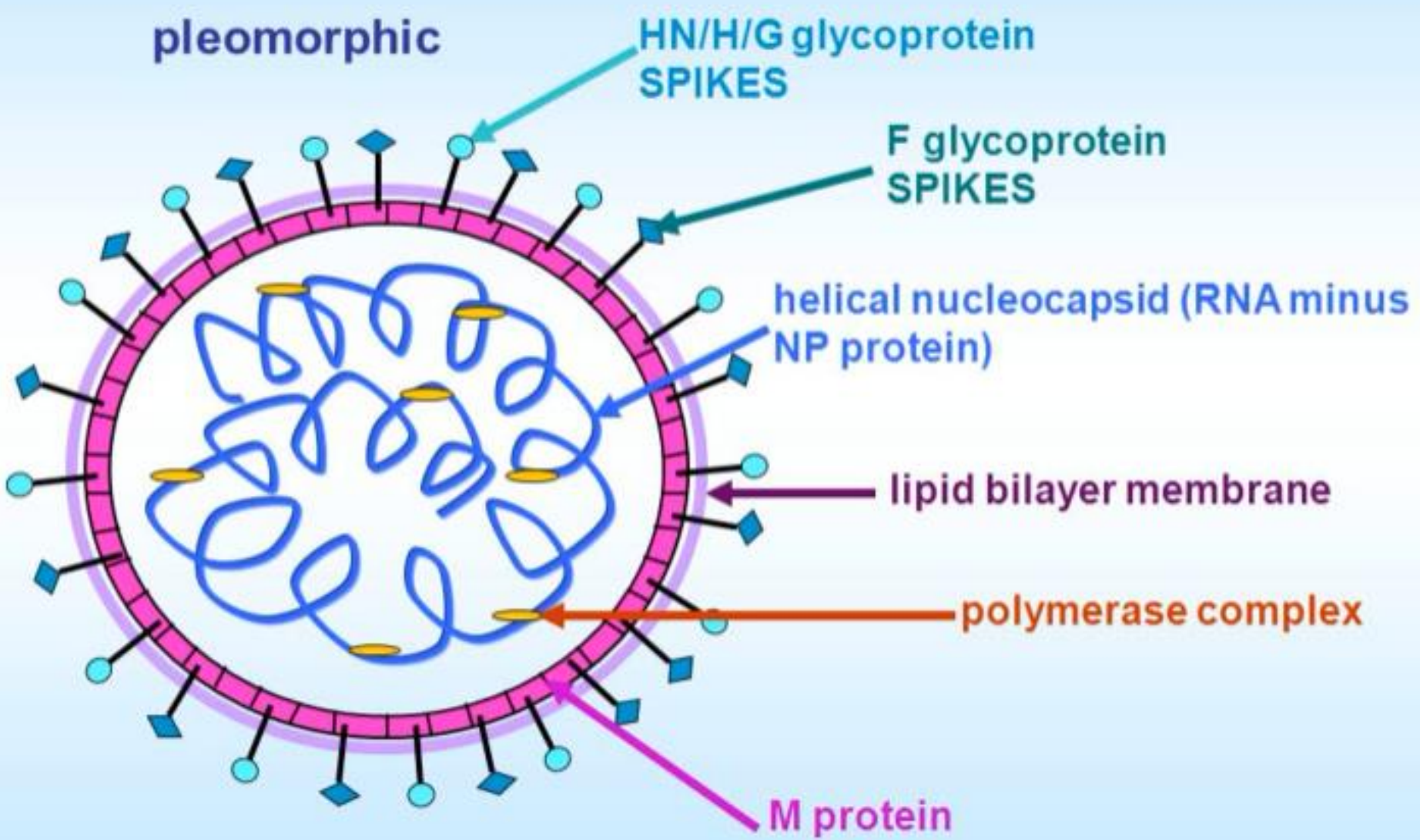




PARAMYXOVIRUS FAMILY
properties of attachment protein

GENUS	GLYCOPROTEINS	TYPICAL MEMBERS
Paramyxovirus genus	HN, F	HPIV1, HPIV3
Rubulavirus Genus	HN, F	HPIV2, HPIV4 mumps virus
Morbillivirus genus	H, F	measles virus
Pneumovirus genus	G, F	respiratory syncytial virus

PARAMYXOVIRUSES



Reproduction of Paramyxoviruses:

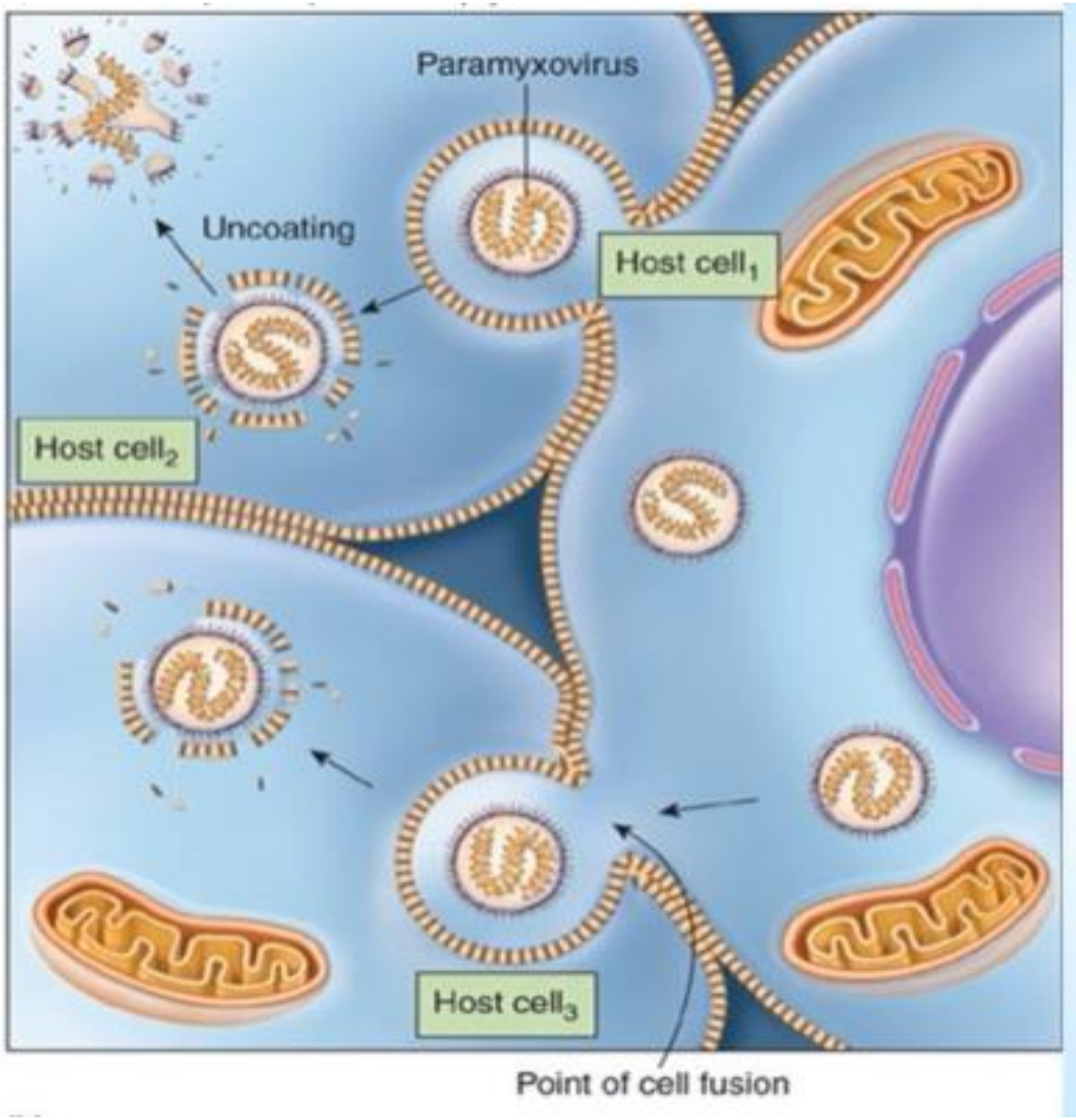
•***Paramyxoviruses*** are adsorbed to sialic acid receptors on the surface of the host cell through **HN-, H- or G-proteins** in the membrane.

•The virion enters the cell directly without endosome formation.

•Genome transcription, replication and protein synthesis take place in the cytoplasm of the host cell.

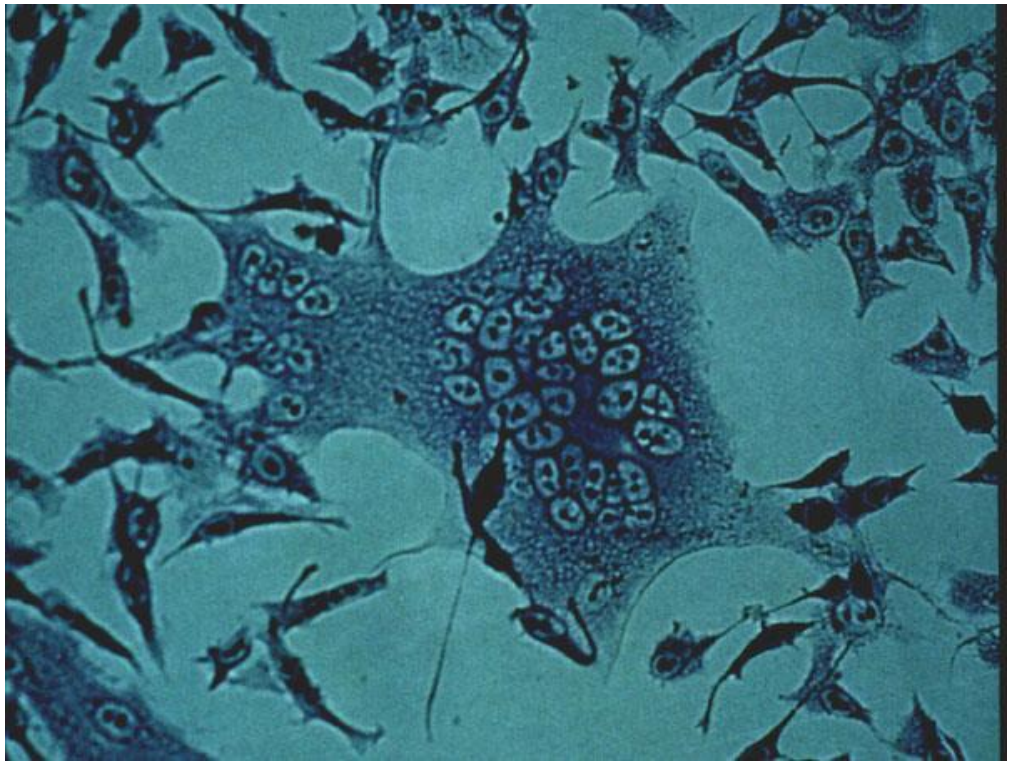
•The virion exits the cell by budding.

•F_0-glycoproteins formed as a result of proteolytic cleavage of F-glycoproteins in the cell membrane under the influence of appropriate proteases of the host cell form **syncyts** by connecting the membranes of neighboring cells.



Paramyxoviridae cultivation

- Cultivation of *Paramyxoviruses* are carried out in primary and continuous cell cultures.
- The **cytopathic effect** is manifested by the formation of **syncytium** (polykaryons) and acidophilic inclusions in the cytoplasm.

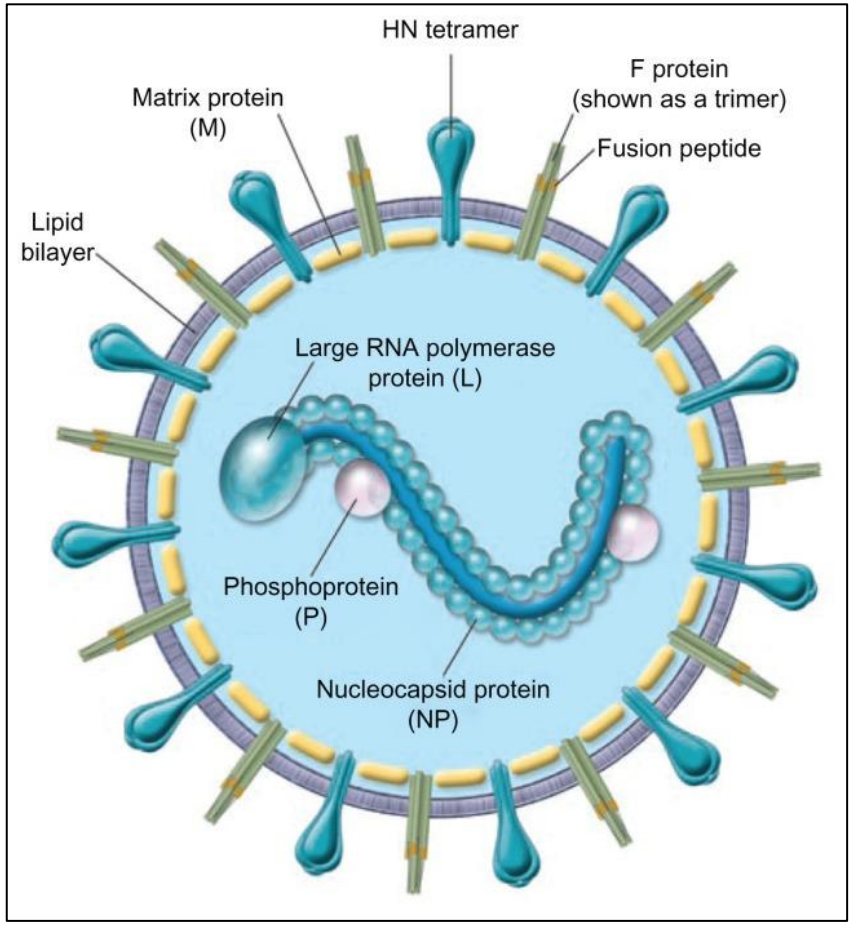


Resistance to environmental factors:

- *Paramyxoviruses* are among the most **persistent** viruses in the environment.
- They are **sensitive** to temperatures above 50^0C, detergents, disinfectants and other factors.
- They have higher resistance to low temperature.

Parainfluenza virus

- Human *Parainfluenza (HPIV)* virus serotypes 1 and 3 belong to the *Respirovirus* genus, and serotypes 2, 4a and 4b belong to the *Rubulavirus* genus.
- 4 main serotypes of *Parainfluenza* viruses are distinguished according to the antigens of glycoprotein spikes in the membrane - **HN-**, **NP-** and **F-proteins**.
- Viruses of serotypes 1, 2, 3 have common antigens with epidemic mumps viruses.

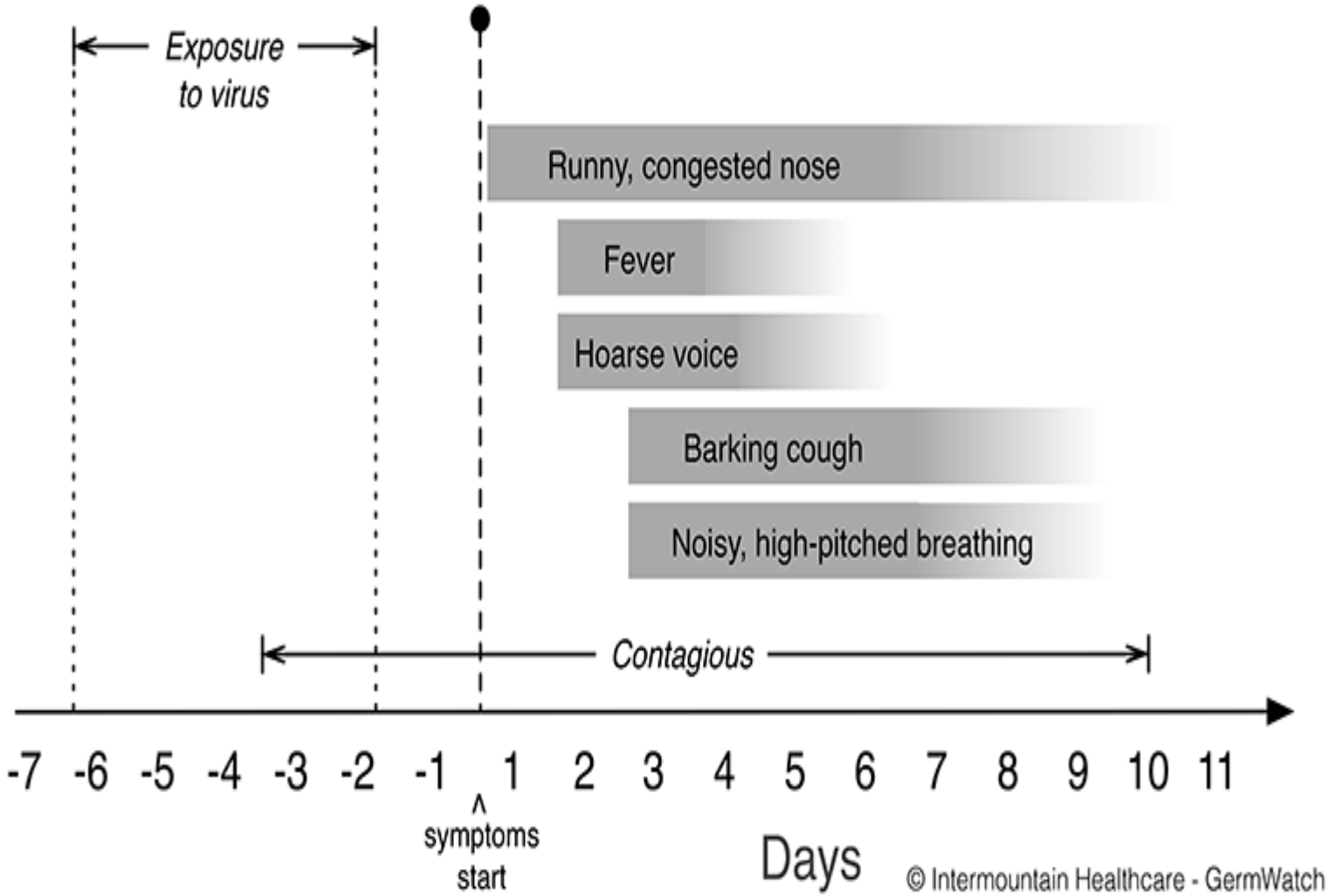


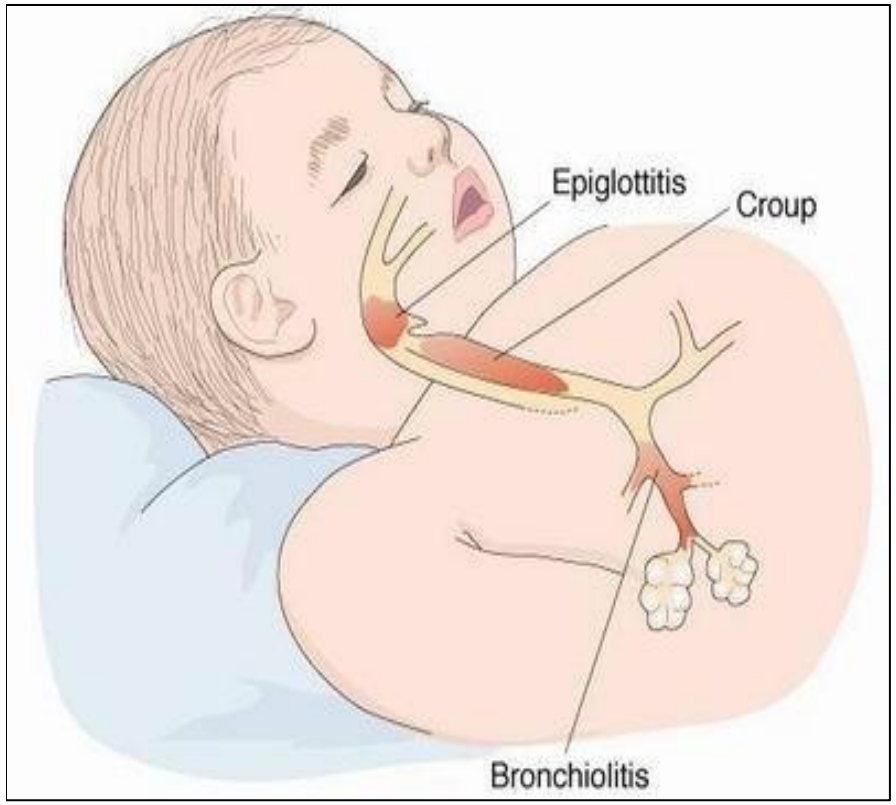
Clinical features of parainfluenza (PIV)

- The **source** of infection is **patient**. Infection occurs mainly through **airborne droplets.**
- Incubation period is 2 to 6 days.

- **Clinical symptoms:**
- Rhinitis, pharyngitis, cough, fever, croup (laryngotracheobronchitis), bronchiolitis, and pneumonia.
- **Croup** - the subglottic region becomes narrower and results in difficulty with breathing, a seal bark-like cough and hoarseness.
- **There is clinical variation between the different PIV types.**
- **PIV-1 and 2:** croup in children ages 2-6 years in autumn/early winter.
- **PIV-3:** bronchiolitis and pneumonia, and croup sporadically, without a particular seasonal occurrence.
- **PIV-4:** mild upper respiratory infections.

Parainfluenza Virus Infection Timeline





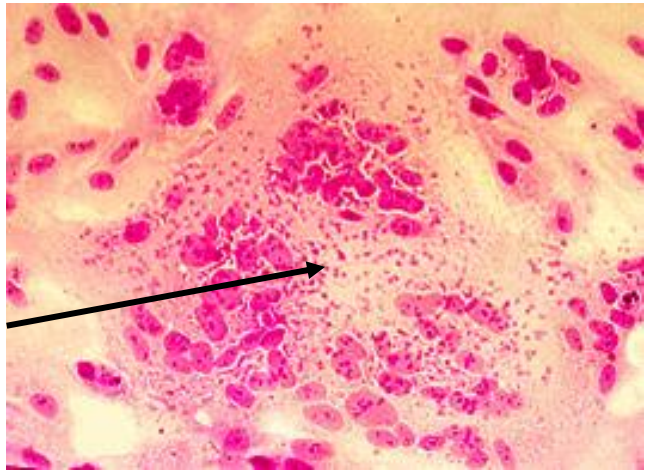
LABORATORY DIAGNOSIS OF PARAINFLUENZA

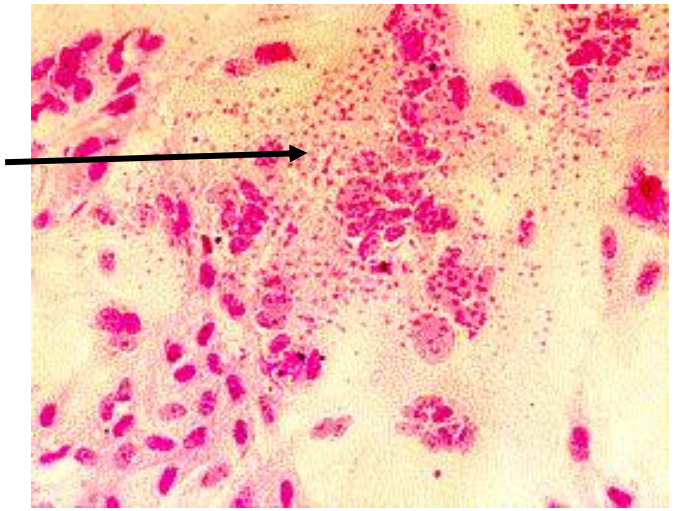
1. **Detection of antigen** from nasopharyngeal aspirates and throat swab **by IF and PCR**.

2. **The virus isolation** in cell culture.

Indication: Haemadsorption of erythrocytes on the surface of cells infected with virus.

Identification: **HadsI**, **HAI, NT, CFT.**

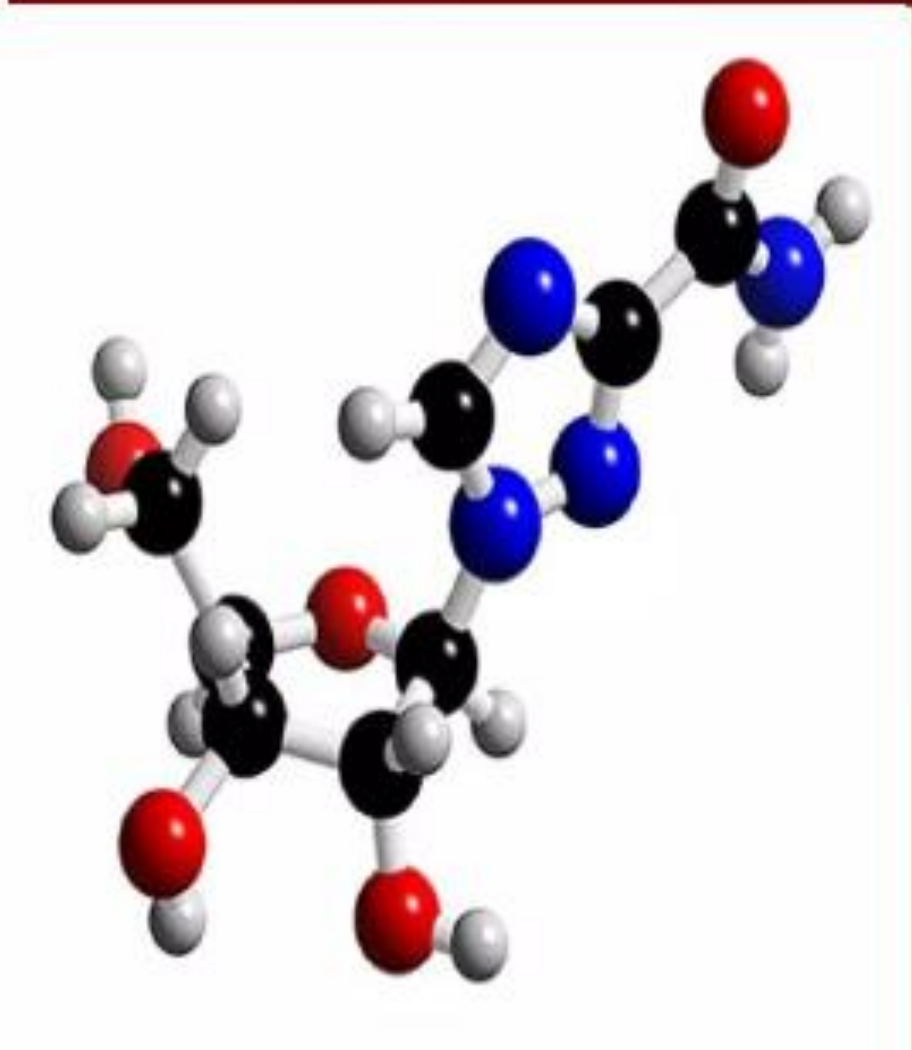




3. **Serology** – detection of rise in titer of IgG in paired sera:

NT, ELISA, CFT, **HAI.**

Treatment and Prophylaxis.

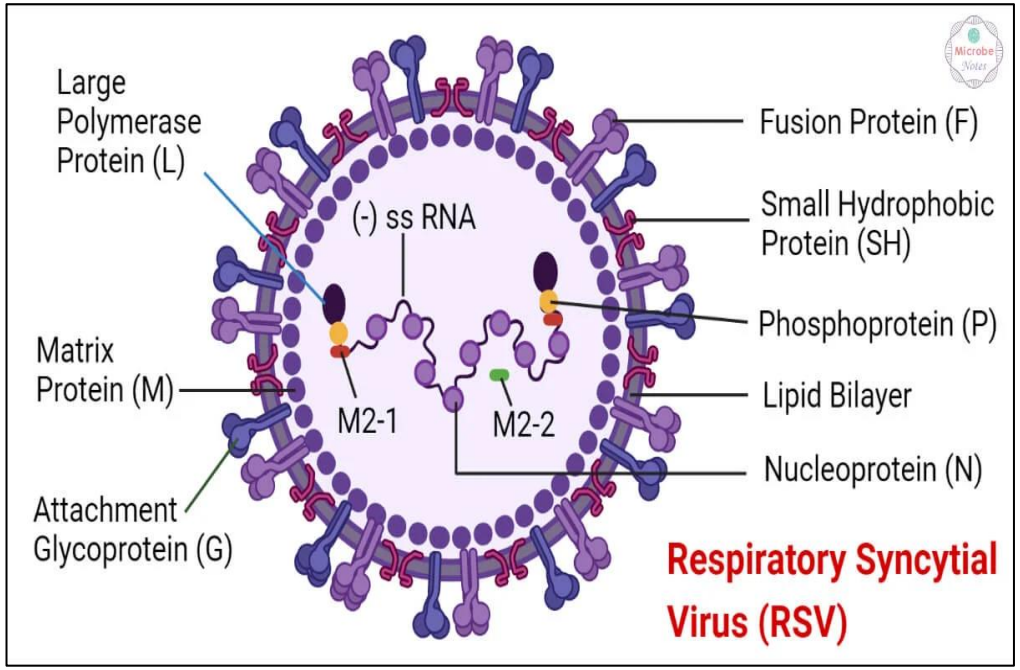


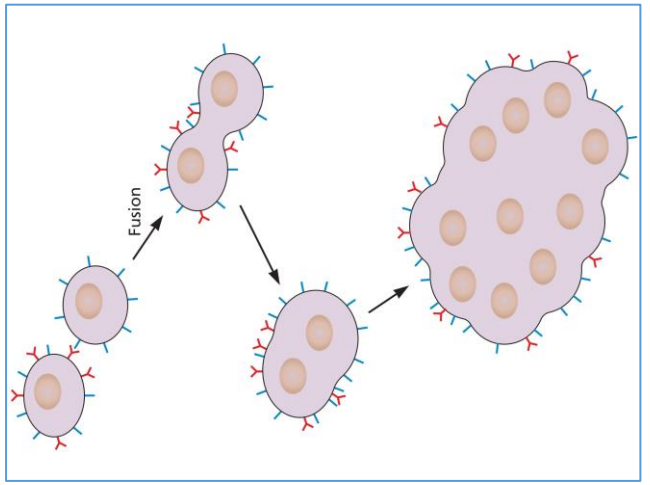
- Treatment with **Ribavirin**
- **No Vaccine to date.**


R S V
Respiratory Syncytial Virus

Respiratory syncytial virus (RS-virus)

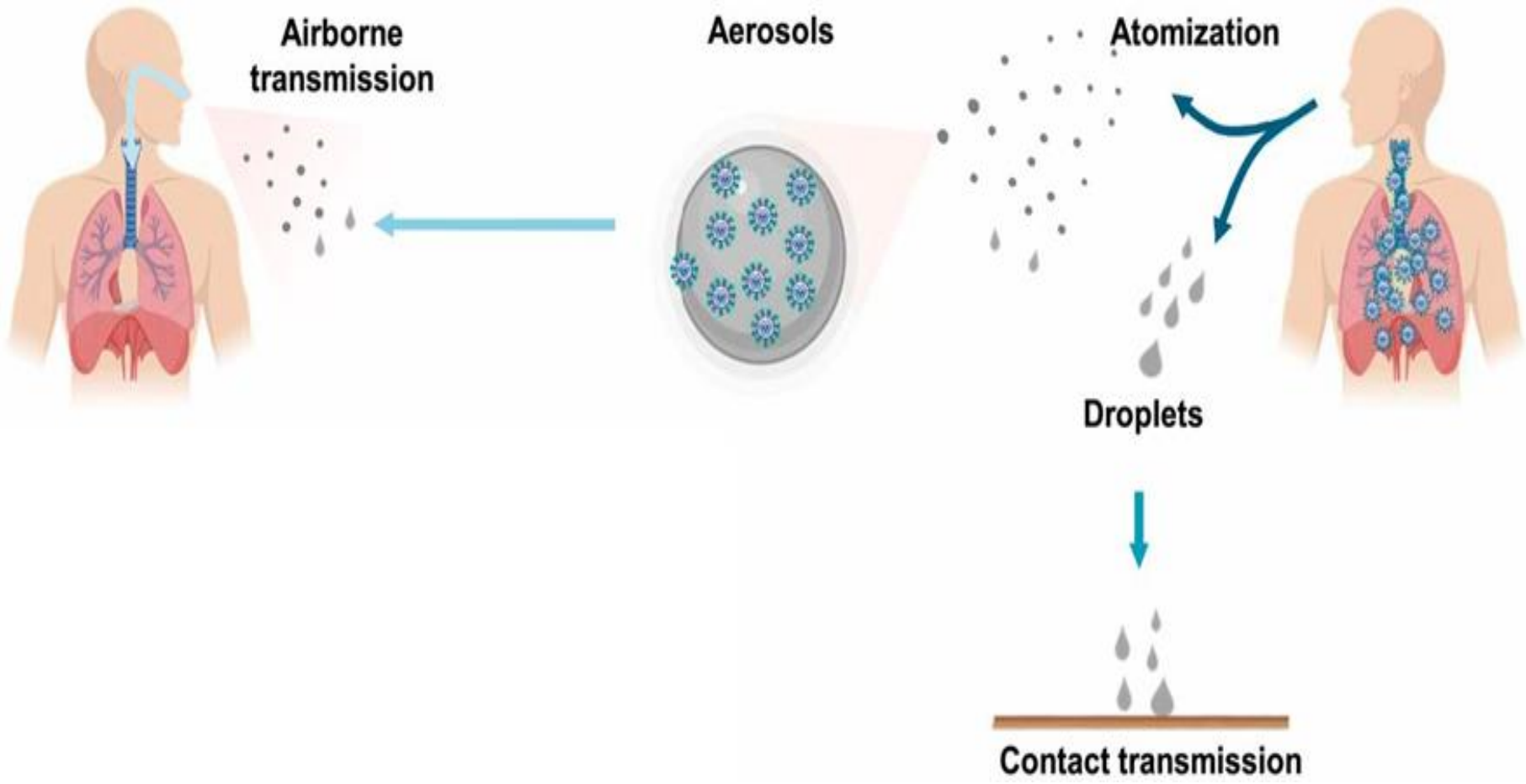
- RS-virus belongs to the ***Pneumovirus*** genus of the ***Paramyxoviridae*** family.
- RS-virus is distinguished by its **polymorphism**: in addition to the usual spherical forms, thread-shaped forms are also found. Large glycoprotein spikes in the lipoprotein membrane have no hemagglutinating and neuraminidase activity, so they are called **G glycoproteins**. This glycoprotein binds the virus to host cell receptors.
- F glycoproteins characteristic of RS-virus connect the membranes of neighboring cells, resulting in the formation of syncytium.
- RS-virus got its name due to the nature of cytopathic effect in cell culture - formation of **symplast** and **syncytium**.





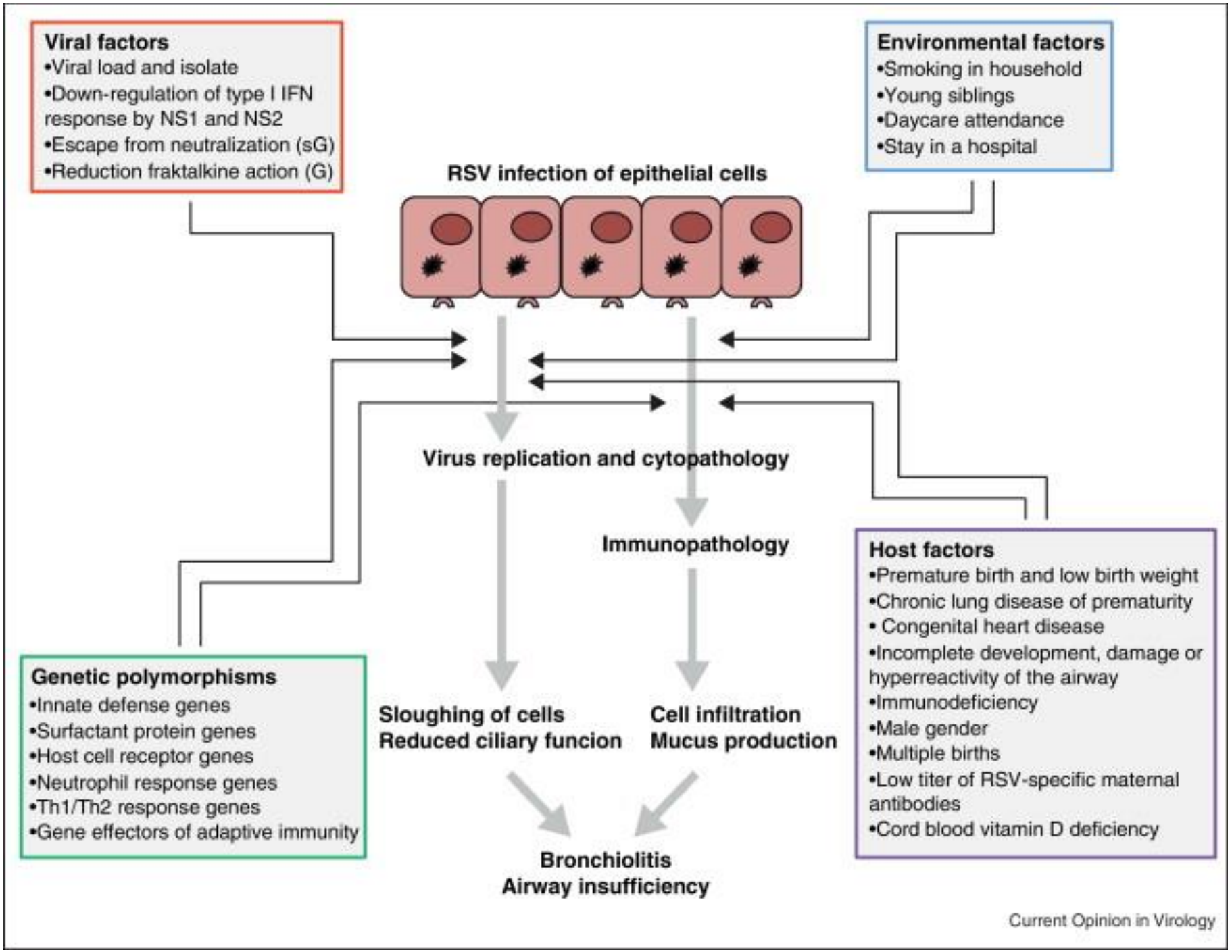
Mode of transmission of RS-virus infections:

- Infection occurs through **air-droplet**, as well as contact-household ways. The portal of the entry of the virus occurs in the mucous membranes of the upper respiratory tract - nasopharynx.



PATHOGENESIS

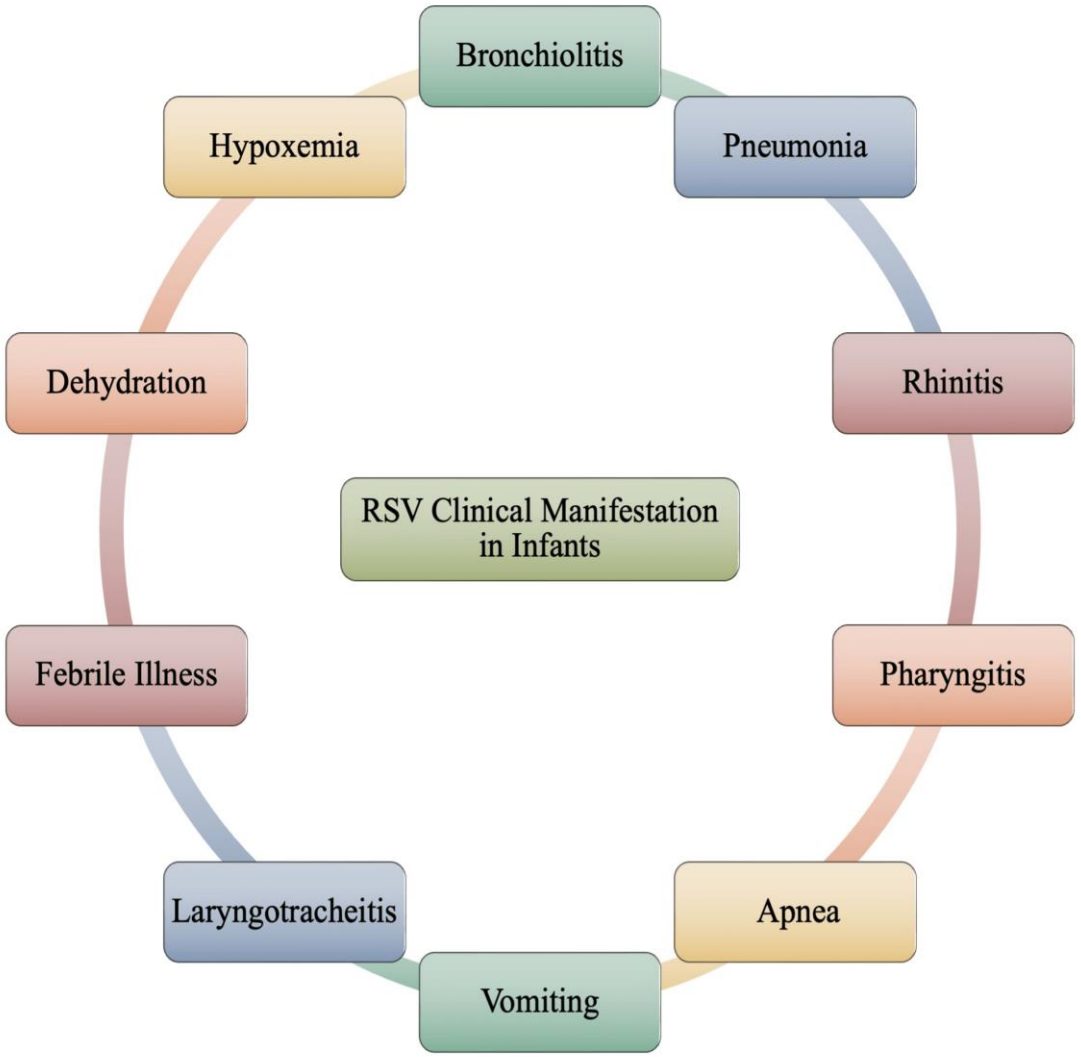
- The virus multiplies inside the epithelial cells, causing their destruction. The pathological process quickly spreads to the lower respiratory tract, causing bronchiolitis and pneumonia.
- Viremia is rarely observed.

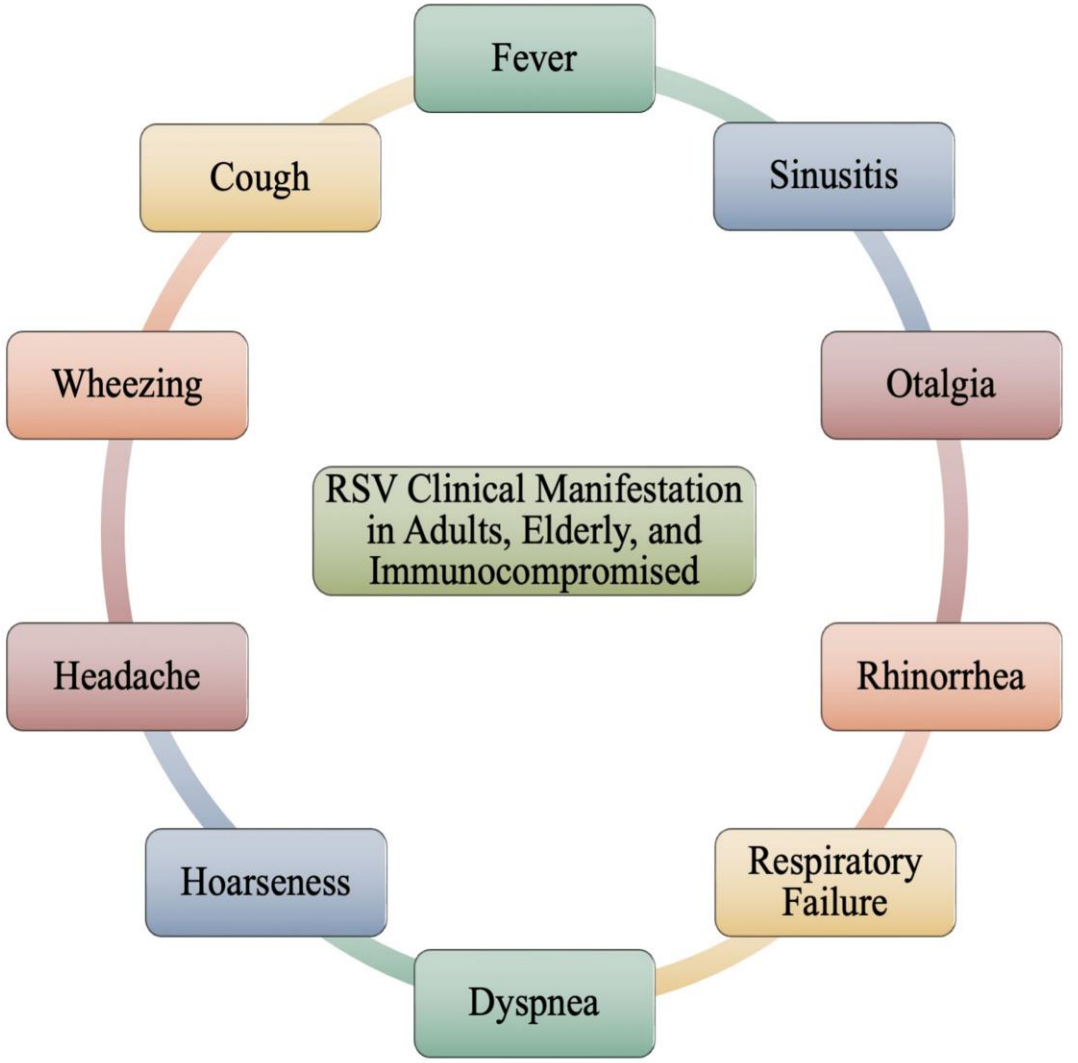


Clinical manifestations of RS-virus infections:

- RS-virus is one of the most common causes of lower respiratory tract infections in infants and children. This virus is the leader among the microorganisms that cause **bronchiolitis** and **pneumonia** in children under one year of age.
- Clinical manifestations of RS-virus infections vary widely, from mild cold symptoms to pneumonia in infants and bronchiolitis in young children. After a latent period lasting 3-5 days, acute catarrhal symptoms of the upper respiratory tract first appear, followed by bronchiolitis and pneumonia. RS-virus is more dangerous for children up to 6 months old - they develop severe bronchiolitis and pneumonia.
- RS-virus infections can be complicated by inflammation of the middle ear. About half of **otitis** in infants are accompanied by RS-virus infections.

Clinical manifestations of RS-virus infections:





Microbiological diagnosis of RS-virus infections

Examination material - *nasopharyngeal lavage or nasopharyngeal swabs*

Virological

RS-virus can be obtained from nasal mucus of patients by virological method - by infecting **HeLa and Hep-2** cell cultures for this purpose. After 10 days of incubation in cell cultures, RS-virus induces a cytopathic effect by producing giant cells and **syncytium**.

Exspress diagnostics

It is possible to detect the RS-virus in materials taken from the nasopharynx with a swab by **IFR, ELISA** and **PCR**. Detection of the virus indicates current illness, as RS-virus is never found in healthy individuals.

Serological

A serological method based on the detection of specific antibodies in blood serum by **IFR, ELISA** and **NT** can be applied, but serological tests have little diagnostic value, they are mostly used in **epidemiological** studies.

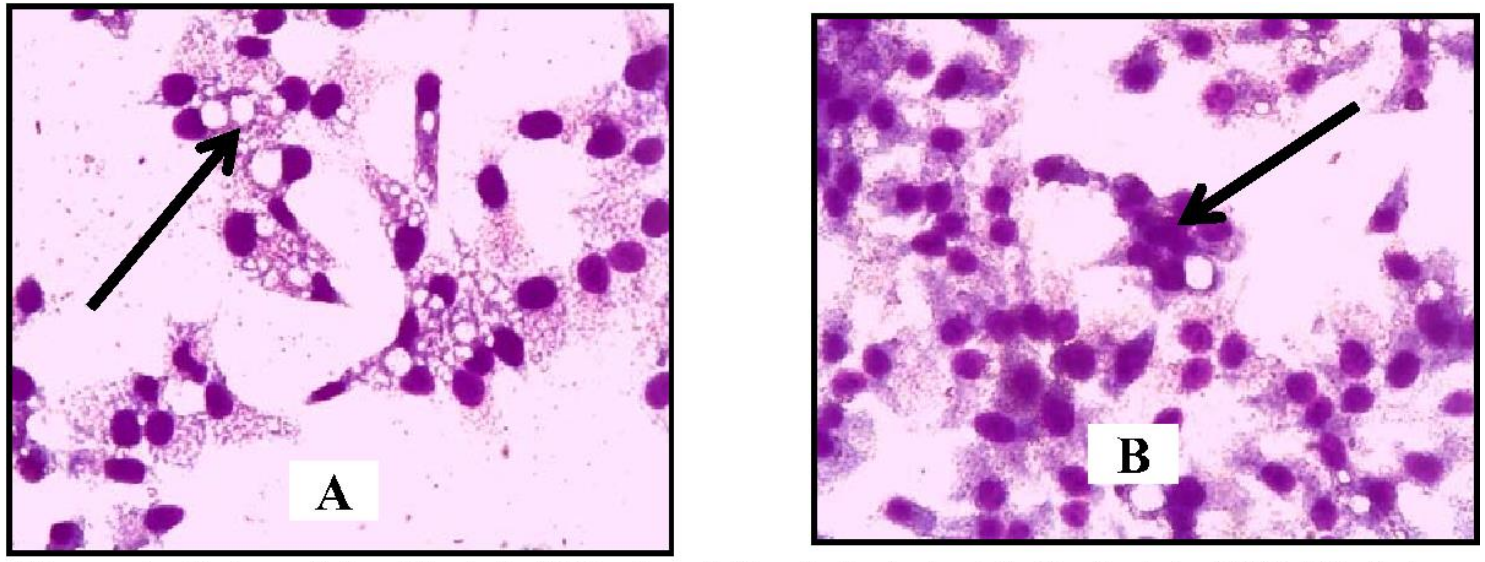


Figure 1: Cytopathic effects in HEp-2 cell line infected with the isolated HRSV, 5 days

TREATMENT

- A supportive management with tube feeding in cases of difficulty in suckling
- Use of oxygen if indicated.
- **Ribavirin** is a specific antiviral drug, proved to effective when given as a small particle aerosol although it is apparently not effective intravenous infusion.

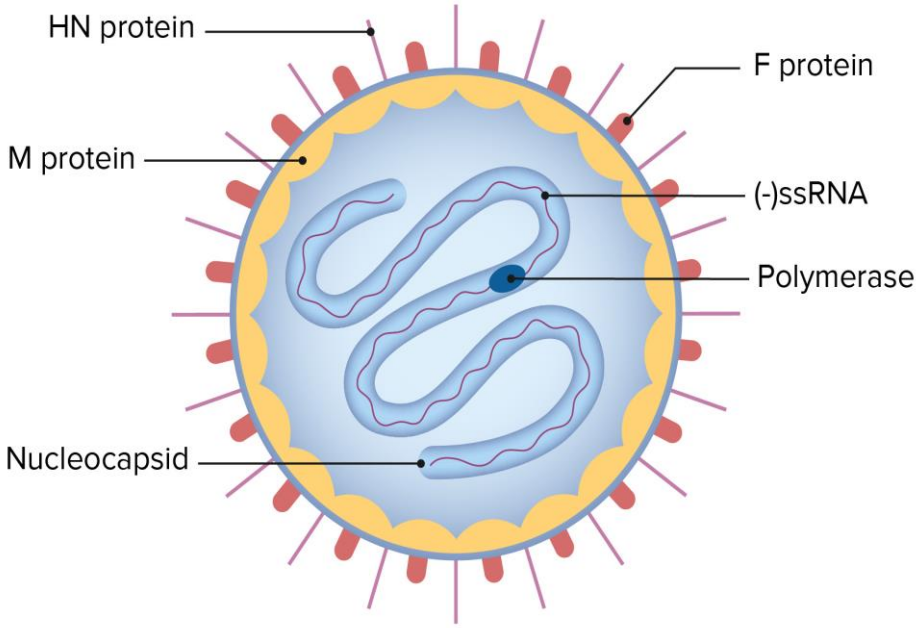
VACCINE - FAILURES

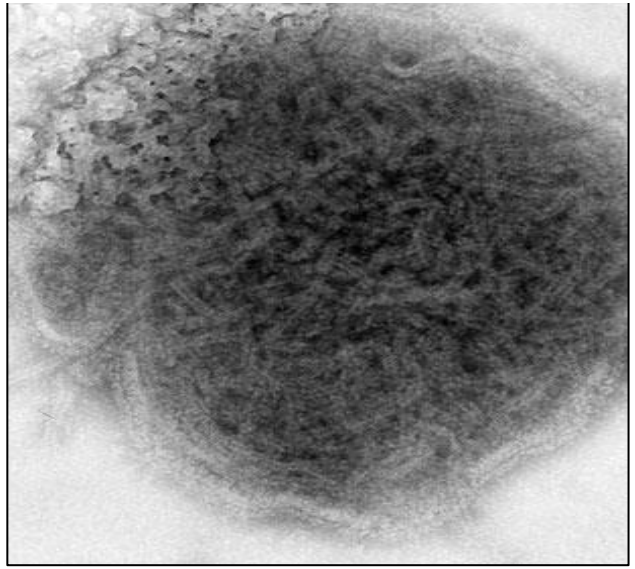
- **A formalin inactivated crude, whole virus vaccine was tried in 1960, but failed to produce immunity in the recipients**
- The difficulties in preparing safe vaccine for RSV lie with young and immunologically immature recipients.
- **Yet to date there is no safe vaccine available for universal use**

MUMPS

MUMPS VİRUS

- Mumps virus belongs to the ***Rubulavirus*** genus of the ***Paramyxoviridae*** family.
- The outer membrane of the virus contains **HN-** and **F-**glycoproteins. Therefore, the virus, having hemagglutinating activity, can be used in chicken, guinea pig, etc. causes erythrocytes to agglutinate. F-glycoproteins bind the membranes of host cells, that is, they have symplast formation and also hemolytic activity.
- Mumps virus can be **cultivated** in monkey kidney cell culture. The cytopathic effect is manifested by the formation of convoluted **giant cells**



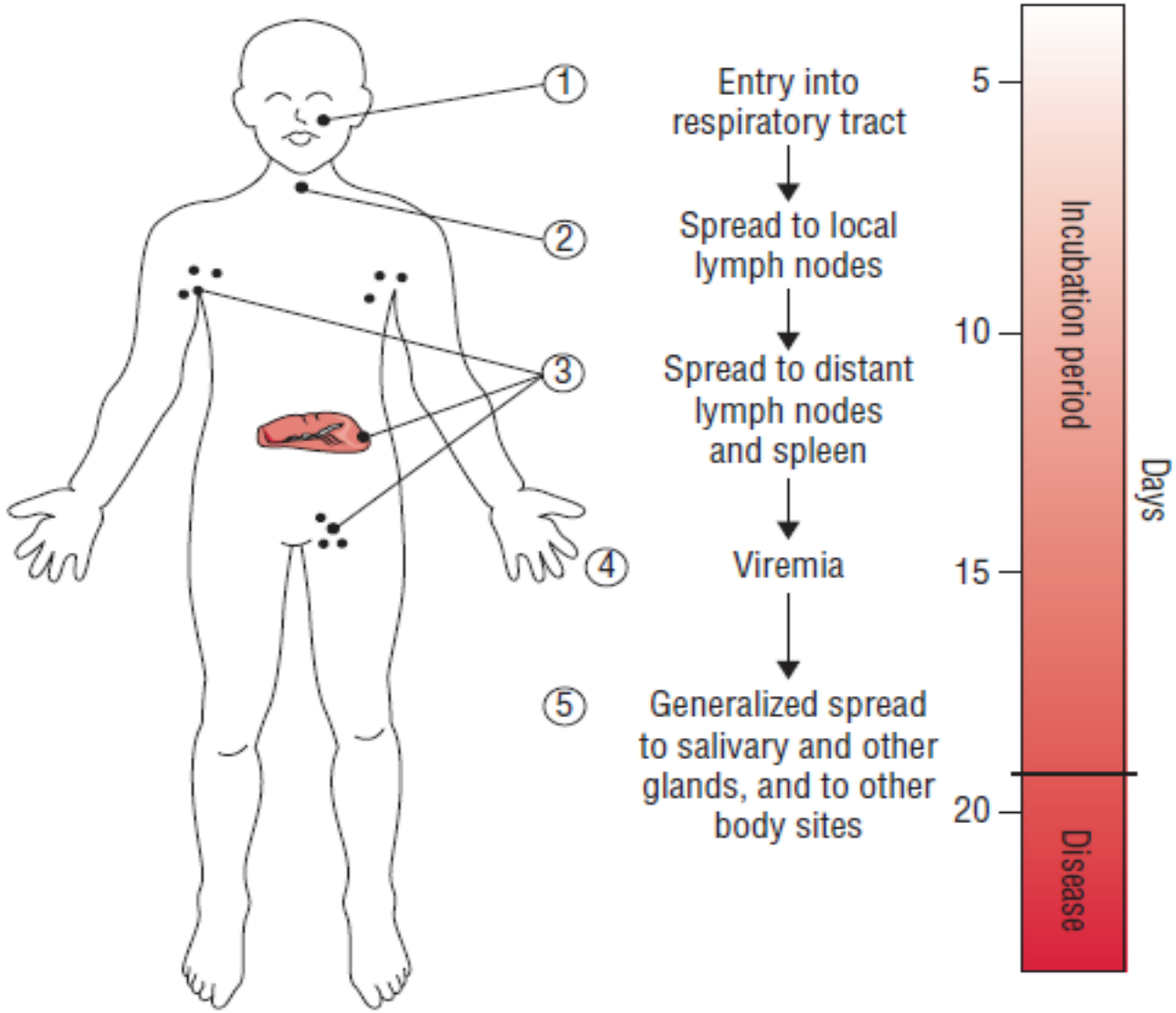


PATHOGENESİS

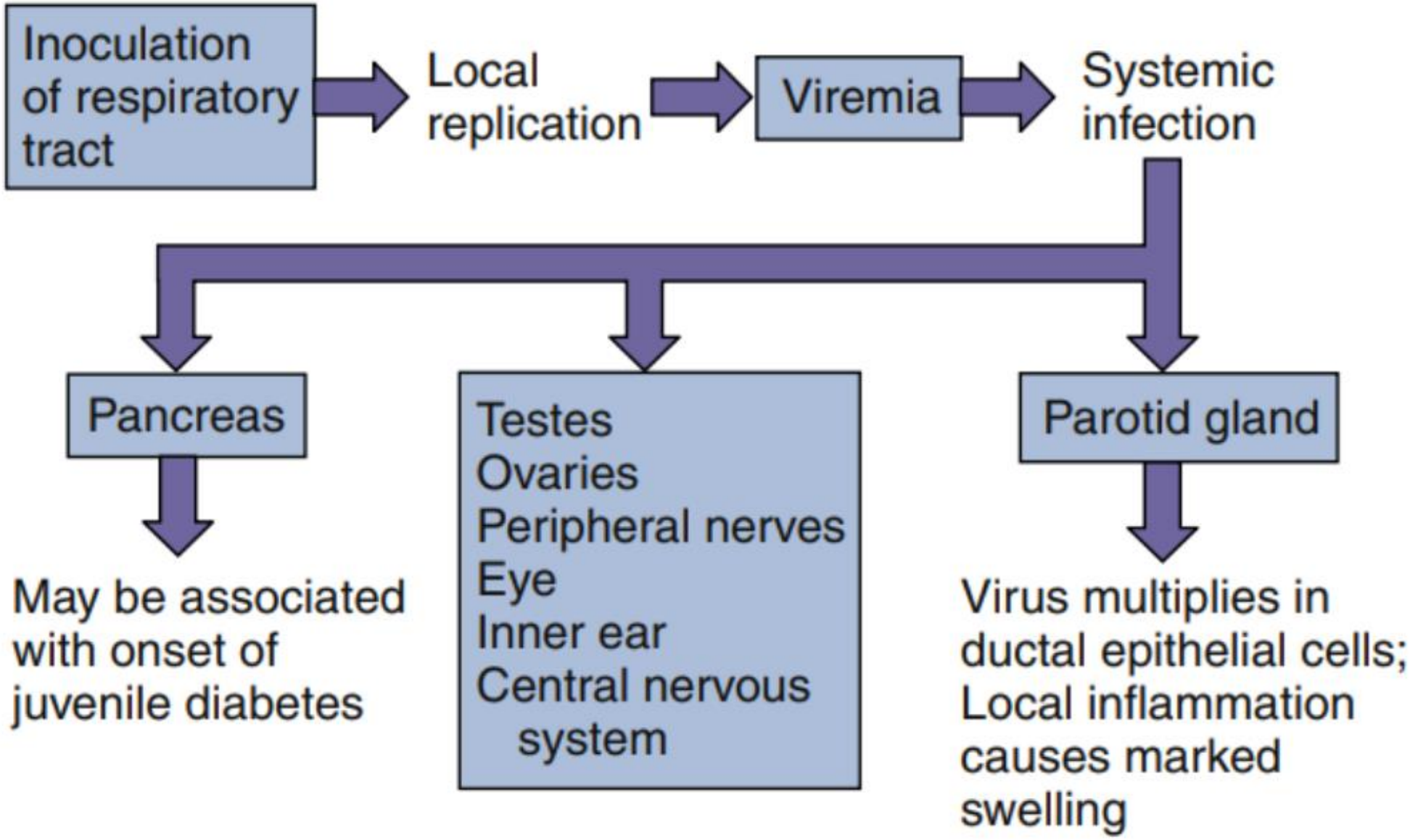
•Children aged 5-15 years are more vulnerable. The virus is excreted in the mouth until the 9th day of the disease, including the last three days of the incubation period. The disease is transmitted by **airborne droplets**, sometimes by **contact** with objects contaminated with saliva.

•Mumps is an **acute childhood infection** characterized by damage to the **parotid glands** and sometimes other organs. The virus multiplies in the epithelium of the mucous membranes of the upper respiratory tract, passes into the blood, spreads throughout the body, and enters the **salivary glands.**

•The virus infects testicles, ovaries, pancreas, thyroid gland, meninges and other organs and causes inflammation.



PATHOGENESİS



Mechanism of spread of mumps virus within the body.

Clinical manifestations of mumps

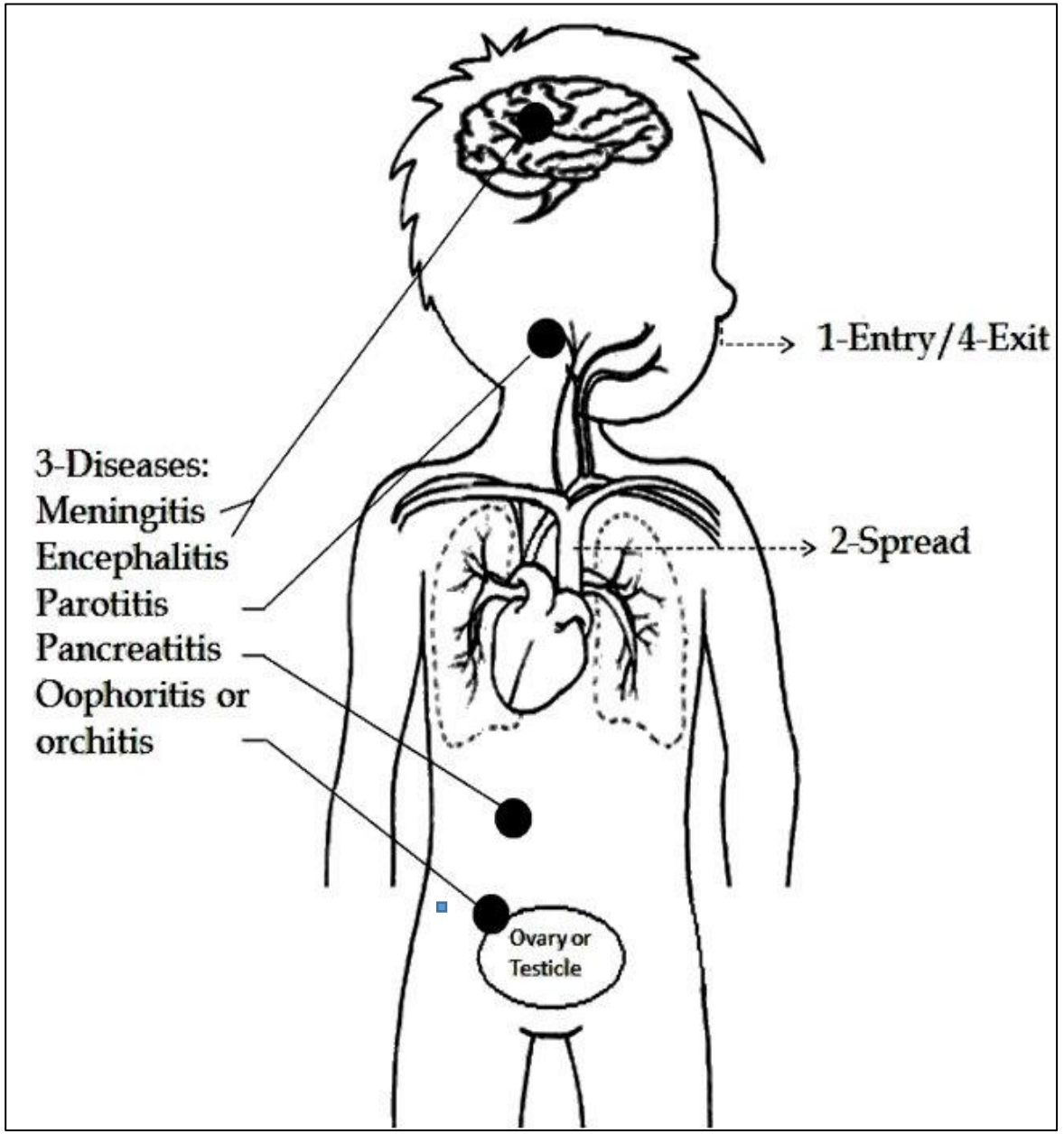
•The **latent period** lasts 2-4 weeks. The most characteristic symptom of the disease is the **swelling and pain of the parotid glands**. Swelling of one or both of the parotid glands gives the patient a characteristic appearance ***(Swine`s face).*** Other salivary glands can also be involved in the pathological process. In about one third of patients, parotitis has an asymptomatic course

•After puberty, mumps can be complicated by inflammation of the testicles **(orchitis)** in boys, and inflammation of the ovaries **(oophoritis)** in girls.

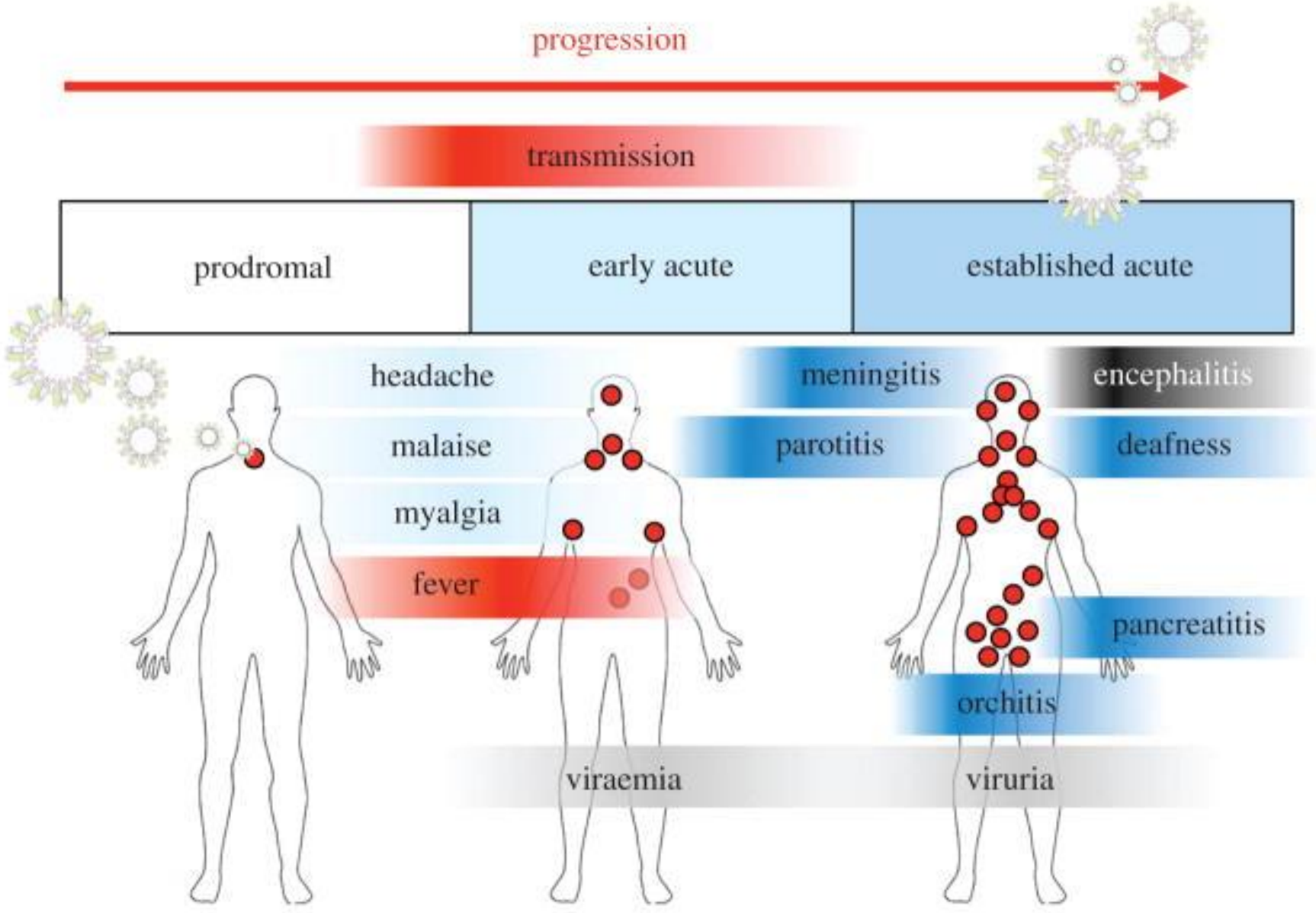
•**Aseptic meningitis and meningoencephalitis** are the most common complications of mumps.

•Mumps can be complicated by pancreatitis in about 4% of cases.

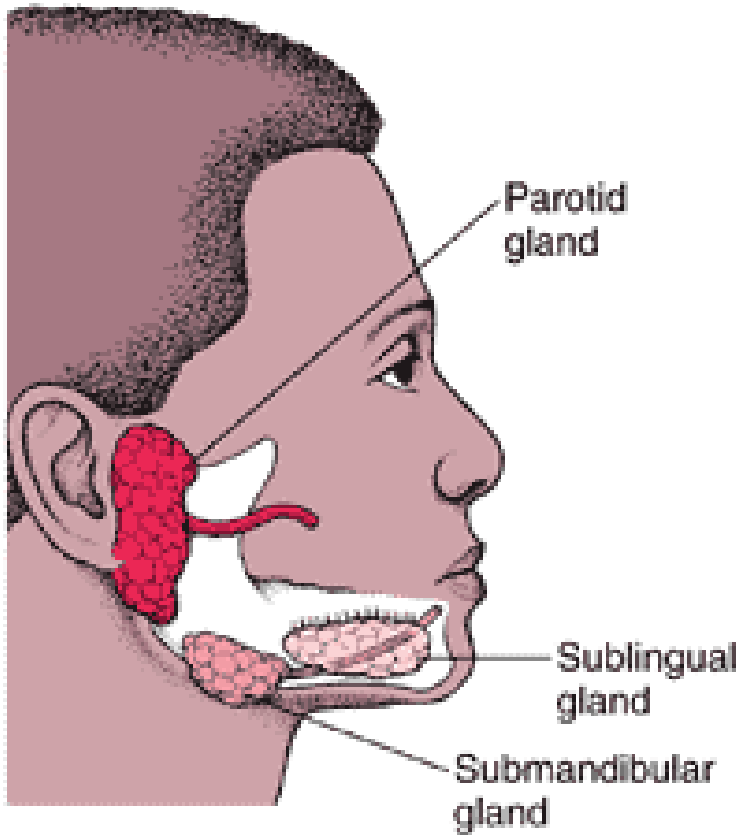
•**Lifelong immunity** is formed after the disease.

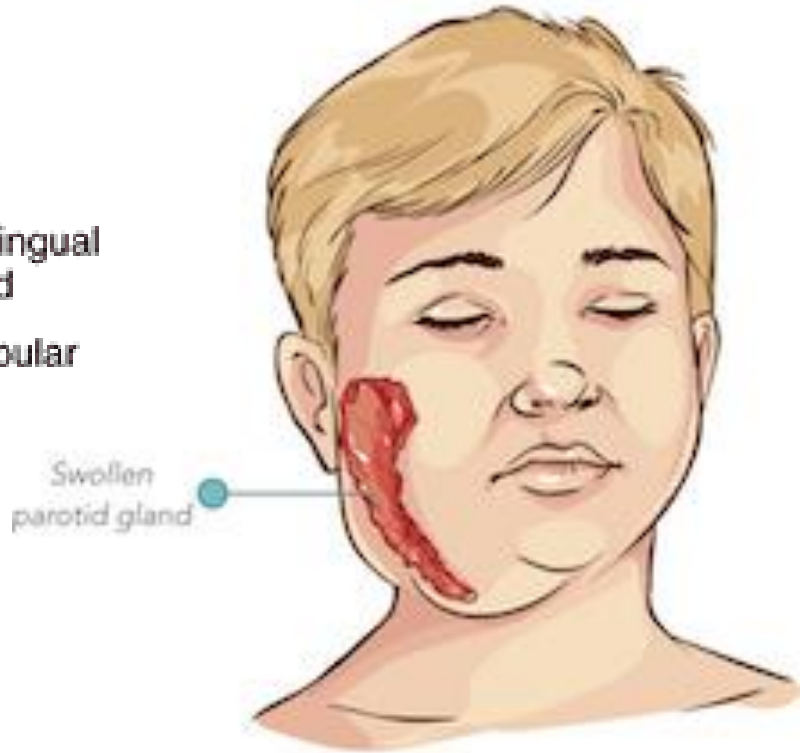


Clinical manifestations of mumps

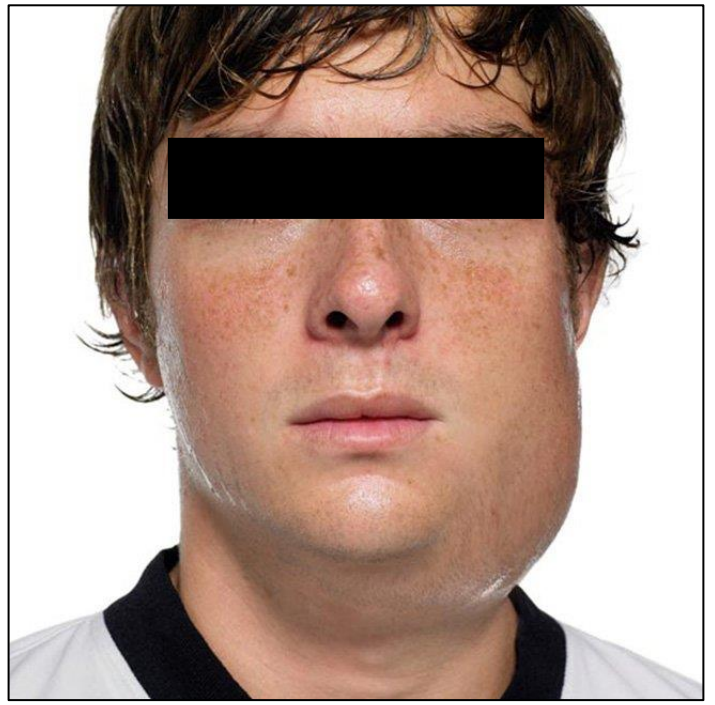


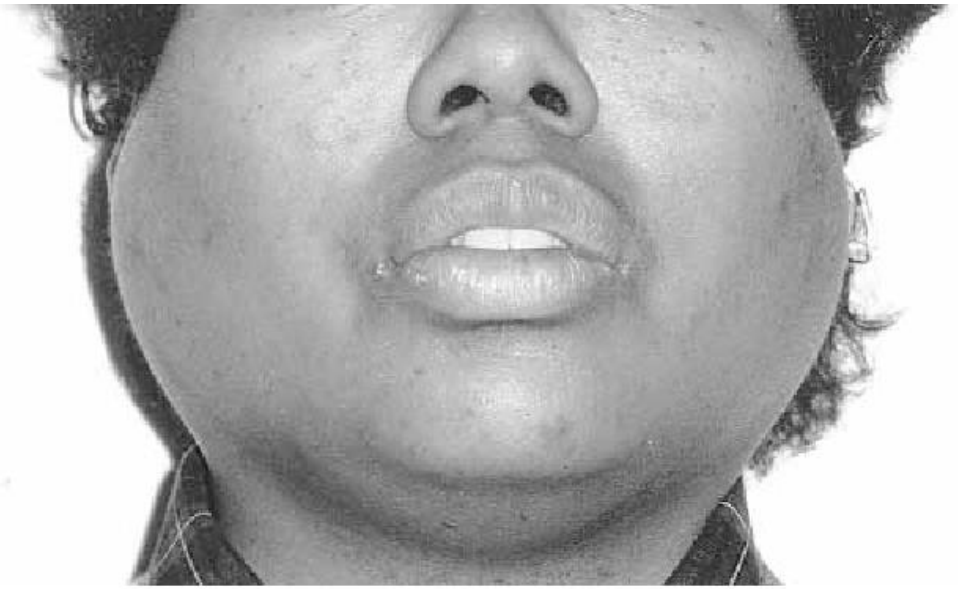
MUMPS





Characteristic mumps symptom





Swine`s face

Complications of mumps

- **Orchitis - 20-50 %**
- Meningitis and meningoencephalitis - 15 %
- Ovaritis - 5 %
- Pancreatitis - 2-5 %
- **Rare complications:** polyarthritis, diabetes, nephritis, thyroiditis, deafness, myocarditis.

Laboratory Diagnosis

- **No Laboratory confirmation needed.**
- Atypical infection needs laboratory Diagnosis.
- Virus isolated from

  Saliva

  Urine

  CSF.

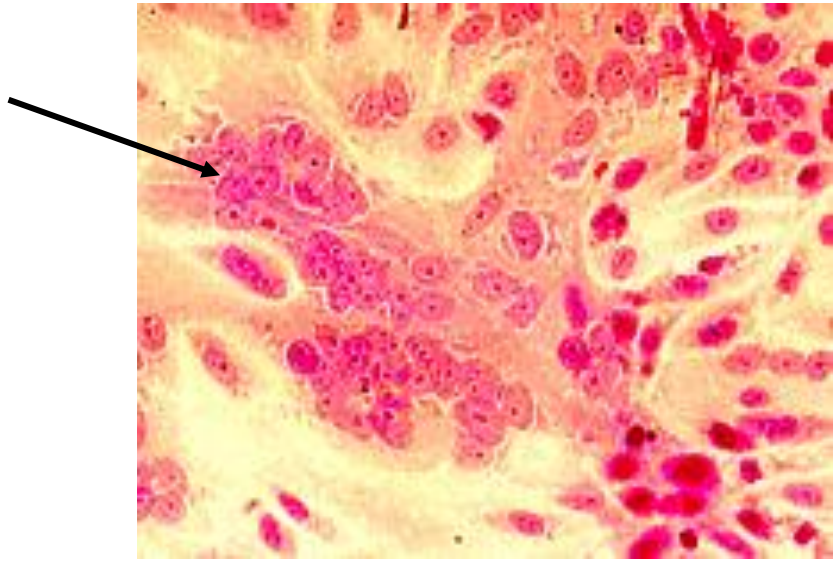
LABORATORY DIAGNOSIS OF MUMPS

1. **The virus isolation** from the saliva, liquor or urine in cell culture (or chicken eggs).

   CPE: giant multinucleated cells formation.

   Identification:

   HAI, NT, IF, CFT.



2. **Serology**: **HAI, NT, ELISA, CFT** (demonstrating IgM in the first serum and detecting IgG rise in paired sera).

3. **Molecular-genetic: PCR**

Treatment

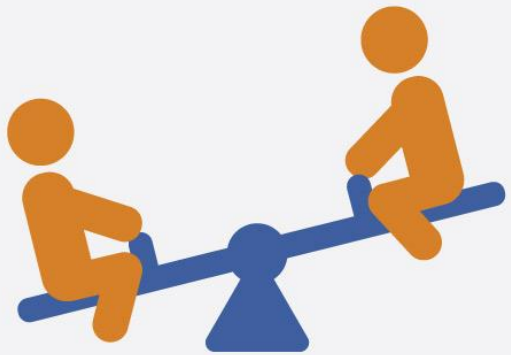
- There is no medication to treat mumps so self-care techniques are used e.g. bed rest, painkillers, plenty of water, a compress for swollen glands and soft food

- Good hygiene and staying away from others is important to prevent the spread of mumps

Specific immunoglobulin ***can be used to ensure a mild course of the disease.***

MMR VACCINE

Protecting children and adults from measles, mumps and rubella



FIRST DOSE

12-15

months old

SECOND DOSE

4-6

years old

Percentage of children protected after 2 doses:

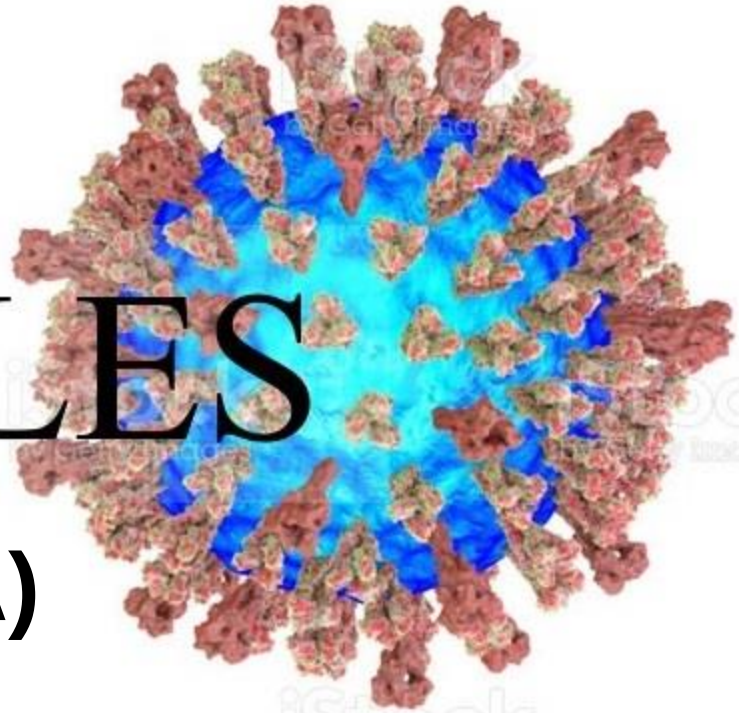
MEASLES **97%**

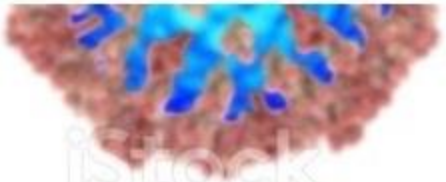
MUMPS **88%**

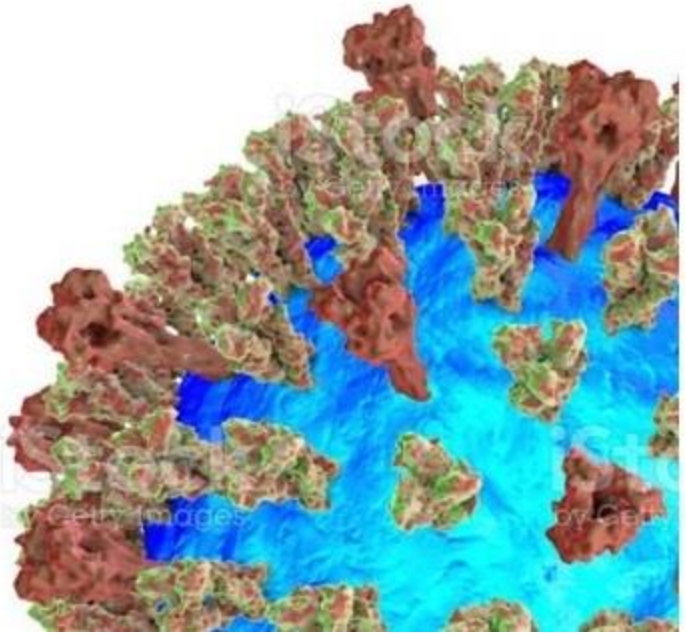
RUBELLA **97%**

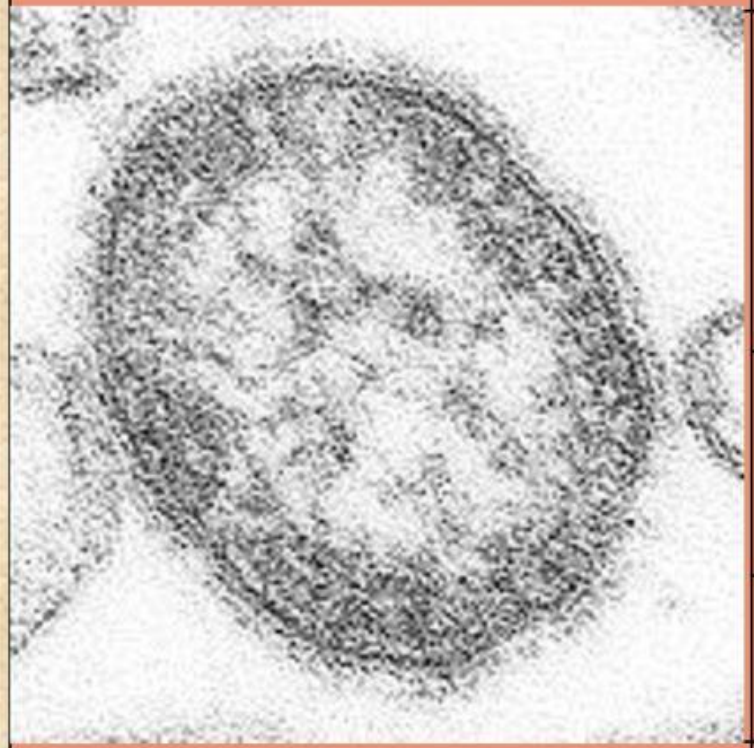
MEASLES

(RUBEOLA)



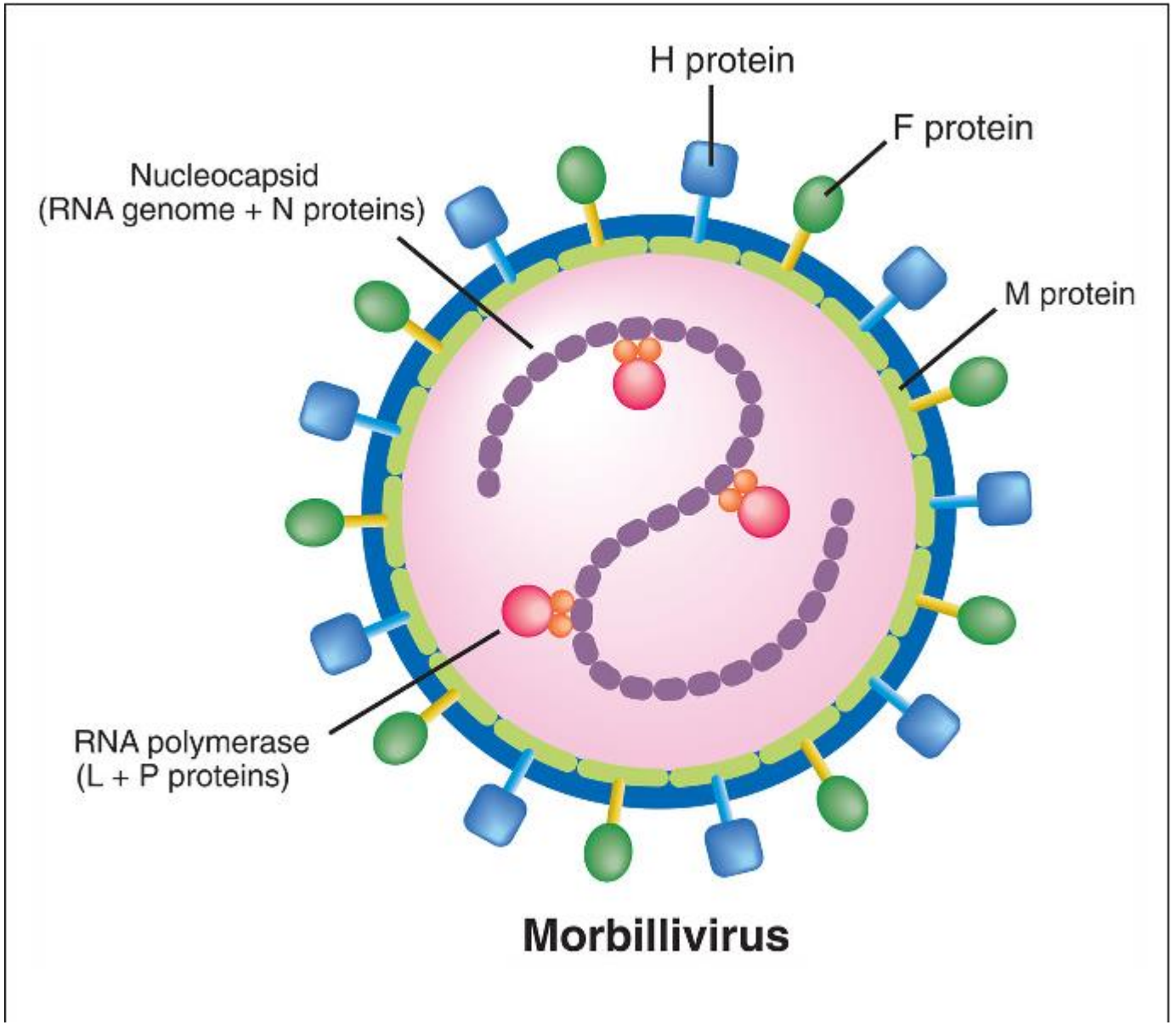




Measles	Virus classification	
	**Group:**	Group V ((-)ssRNA)
	Order:	*Mononegavirales*
	Family:	*Paramyxoviridae*
Measles virus	**Genus:**	*Morbillivirus*

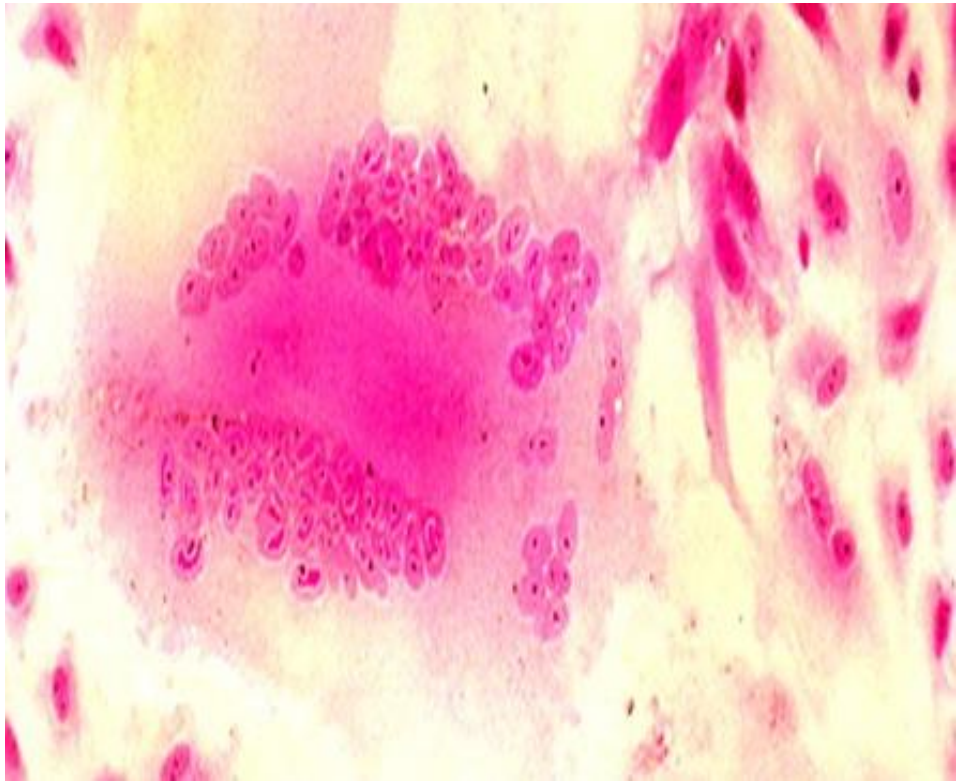
MEASLES (RUBEOLA) VİRUS

- **Enveloped**, pleomorphic spheres 100-300 nm diameter.
- Virions have an inner **helical** nucleocapsid that is a coiled helix of protein and RNA. Envelope has **hemagglutinin (H)** and **fusion (F)** glycoprotein spikes.
- Nucleic acid: **non-segmented, single-stranded, negative-sense RNA virus.**



MEASLES (RUBEOLA) VİRUS

- Measles virus is **cultured** in primary monkey and human kidney cell cultures. At this time, it causes a cytopathic effect with the formation of multinucleated giant cells - symplasts with intranuclear and cytoplasmic inclusions. Unlike other paramyxoviruses, it forms **intranuclear inclusions**.
- Measles virus is very **unstable** in the environment, it is inactivated after 3-4 hours at room temperature.



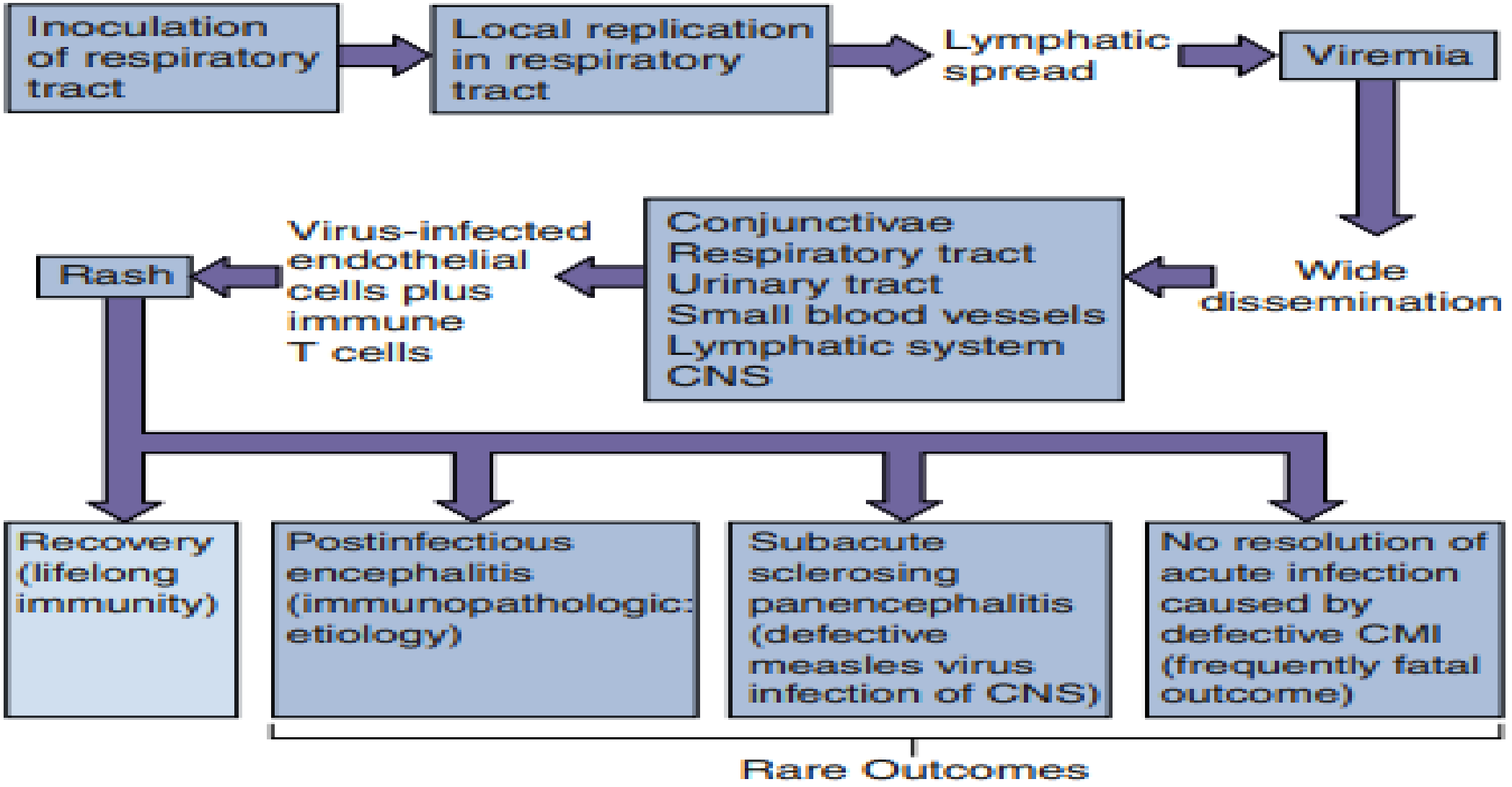
EPIDEMIOLOGY

- Infection sources
  - Patients of acute stage and viral carriers of atypical measles
- Transmission
  - Highly contagious, approximately 90% of susceptible contacts acquire the disease.
  - Respiratory secretions: maximal dissemination of virus occurs by droplet spray during the prodromal period (catarrhal stage).
  - Contagious from 5 days *before symptoms*, 5 days *after onset of rash*
  - Seasons: in the spring, peak in Feb-May

PATHOGENESIS AND PATHOLOGY

- Portal of entry
  - Respiratory tract and regional lymph nodes
  - Enters bloodstream (primary viraemia) → monocyte – phagocyte system → target organs (secondary viraemia)
- Target organs
  - The skin; the mucous membranes of the nasopharynx, bronchi, and intestinal tract; and in the conjunctivae, ect

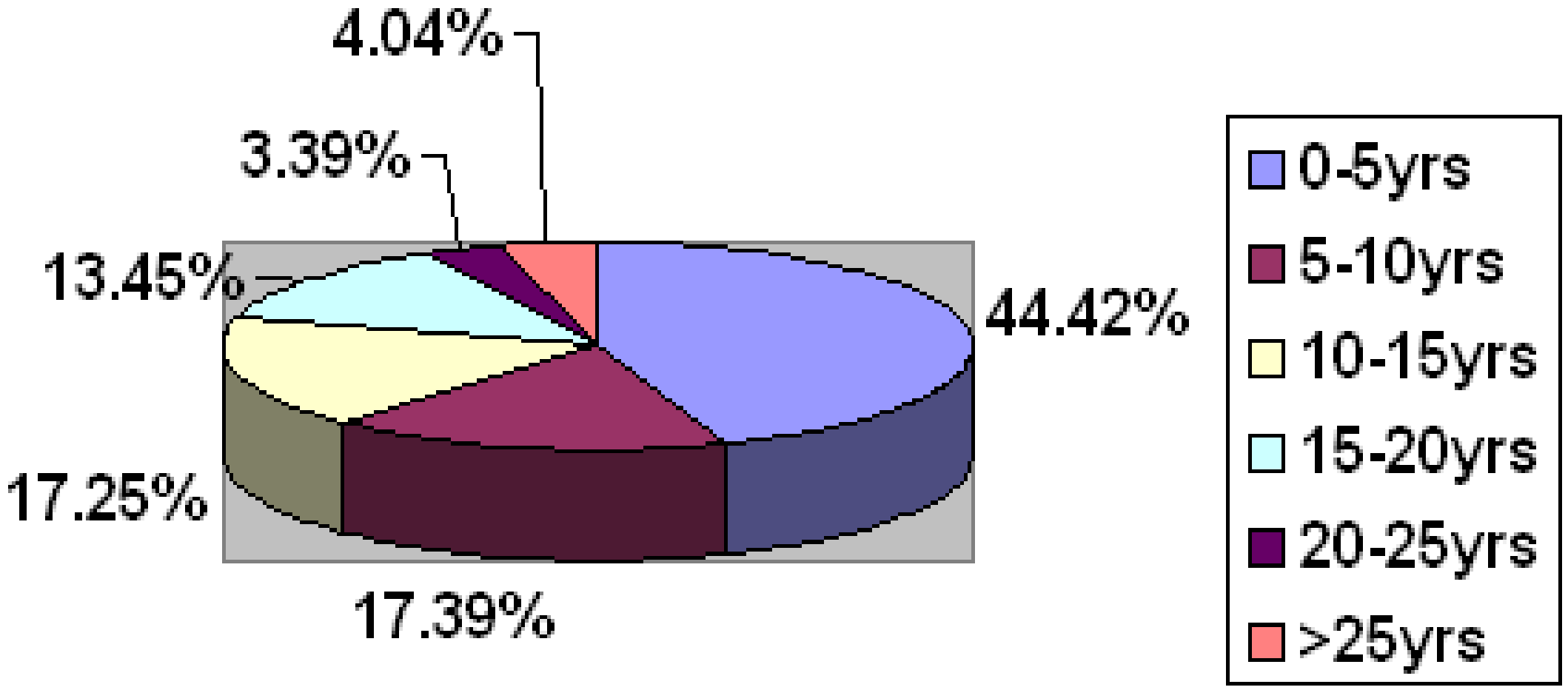
PATHOGENESIS



Mechanisms of spread of the measles virus within the body and the pathogenesis of measles. *CMI*, Cell-mediated immunity; *CNS*, central nervous system.

Age distribution of measles cases

- Measles is one of the most contagious of all human viruses, with about **40 million infections world wide each year, and one to two million deaths.**



CLINICAL MANIFESTATION

- **Measles** - a systemic infection, disseminated by viremia, with acute disease manifestations involving the lymphatic and respiratory systems, the skin, and sometimes the brain.

1. Incubation period (infection to symptoms) :

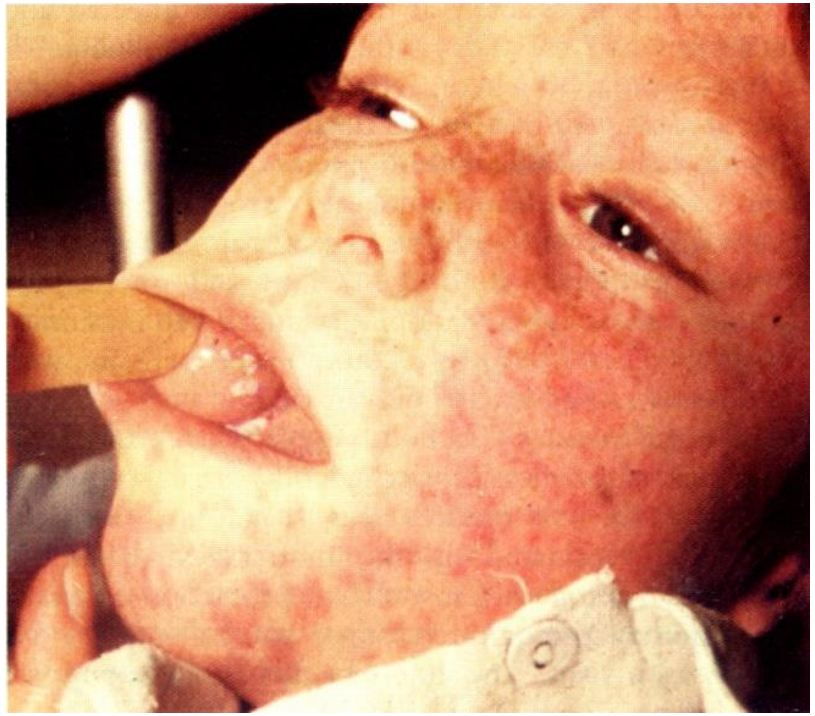
6-18days (average 10 days)

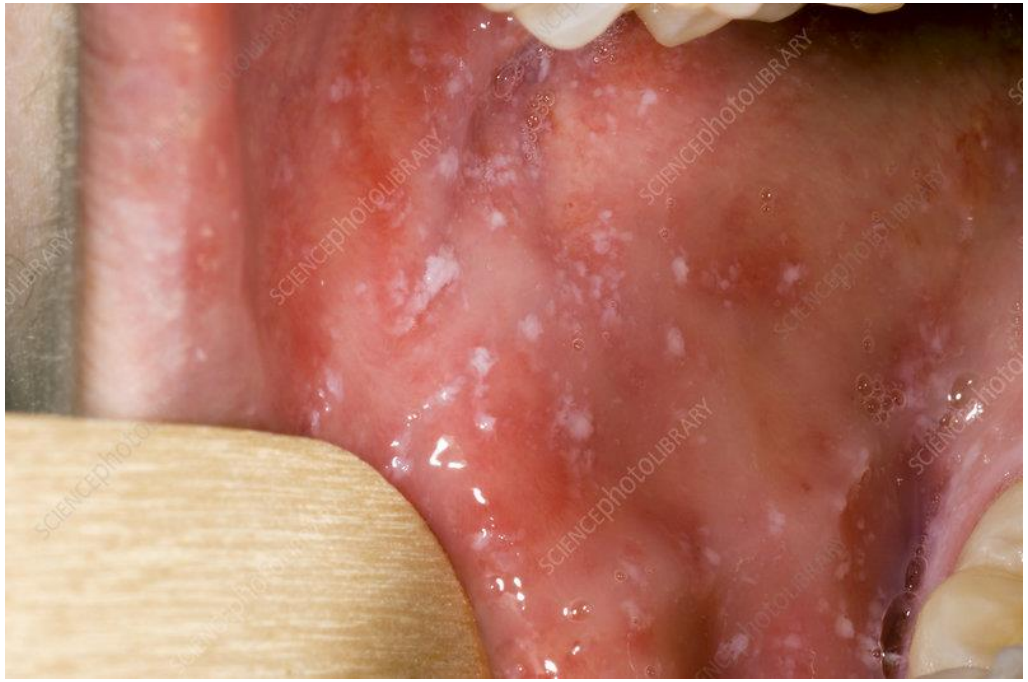
2. Prodromal period:

- 3-4 days
- Non-specific symptoms: fever, malaise, anorexia, headache
- Classical triad: cough, coryza, conjunctivitis (with photophobia, lacrimation)

Enanthem (Koplik spots):

- *Pathognomonic for measles*
- 24-48 hr before rash appears
- 1mm, grayish white dots with slight, reddish areolae
- Buccal mucosa, opposite the lower 2nd molars
- increase within 1day and spread
- fade soon after rash onset





3. Rash period

3-4days

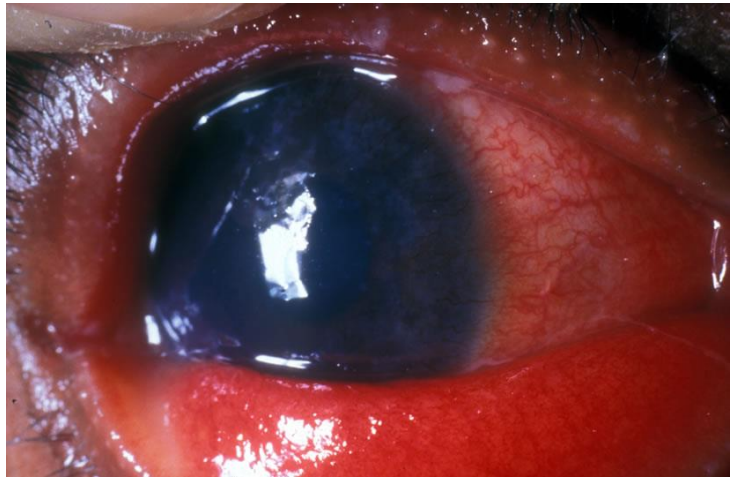
Exanthem:

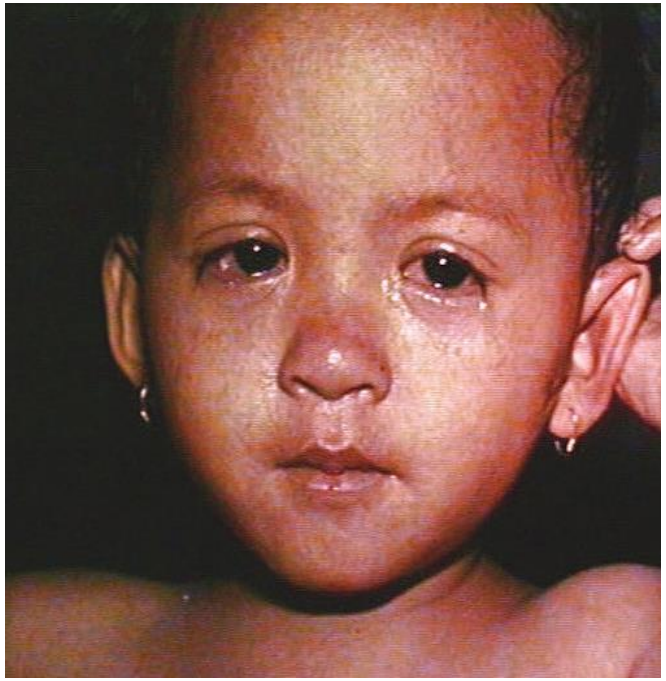
Erythematous, non-pruritic, maculopapular

- Upper lateral of the neck, behind ears, hairline, face → trunk → arms and legs →feet

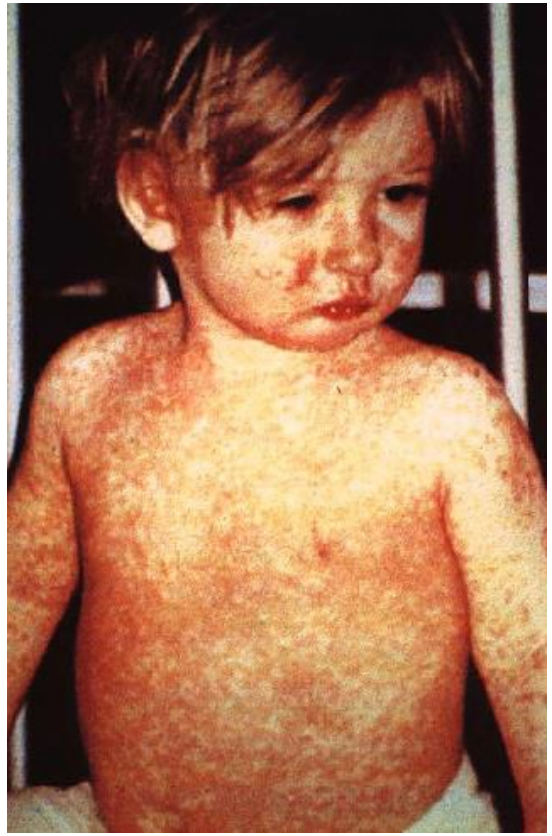
- The severity of the disease is directly related to the extent and confluence of the rash

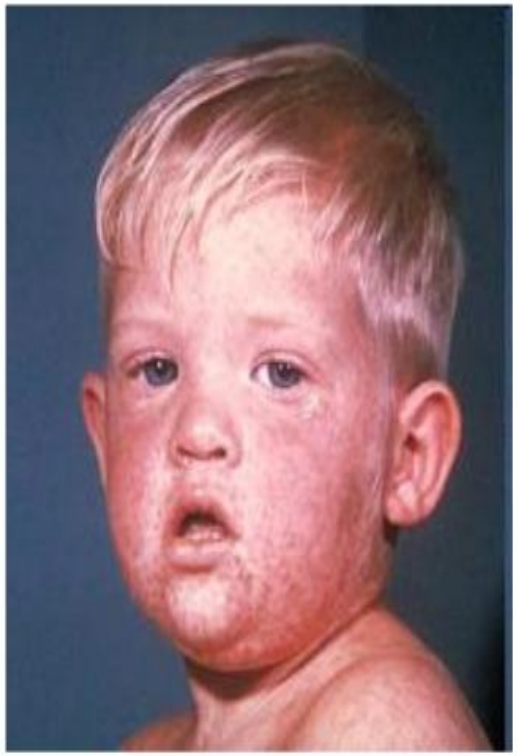
Conjunctivitis

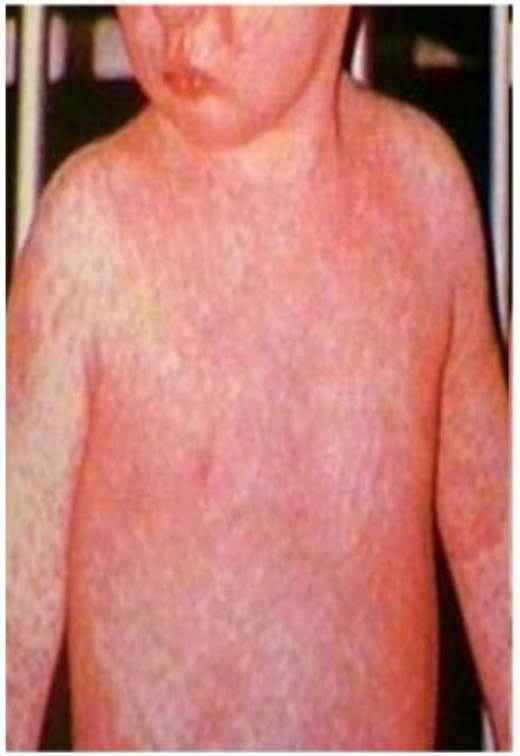




Maculopapular rash







CLINICAL MANIFESTATION

4. Recovery period

3-4days

Exanthem:

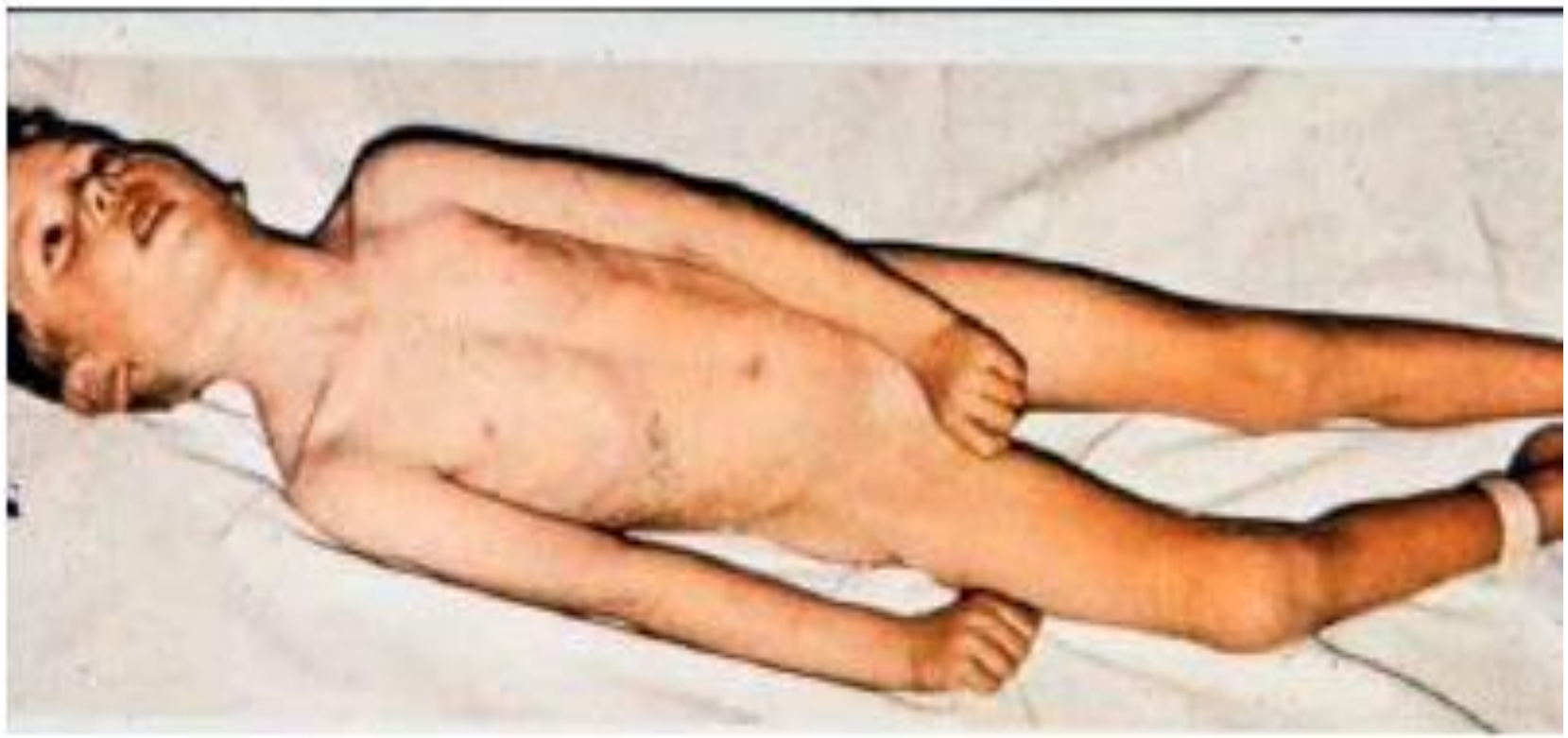
- Fades in order of appearance
- Branny desquamation and brownish discoloration

Entire illness – 10 days

Complications:

- **Pneumonia**, **otitis** media, optic neuritis
- Encephalitis, **subacute sclerosing panencephalitis (SSPE)**
- Hemorrhagic measles

Post - Measles Encephalitis



Immunity

- Infection confers **lifelong immunity** (after measles, permanent humoral immunity is formed). Recurrence is rarely observed.
- **Passive immunity** transmitted through the placenta in the form of IgG protects the child for 6 months after birth.
- Cell-mediated immunity causes rashes.
- Measles infection is accompanied by immunosuppression, which results in complications.

DIAGNOSIS

characteristic clinical picture:

- Measles contact
- Koplik spot
- Features of the skin rash
- The relation between the eruption and fever

Laboratory confirmation is rarely needed

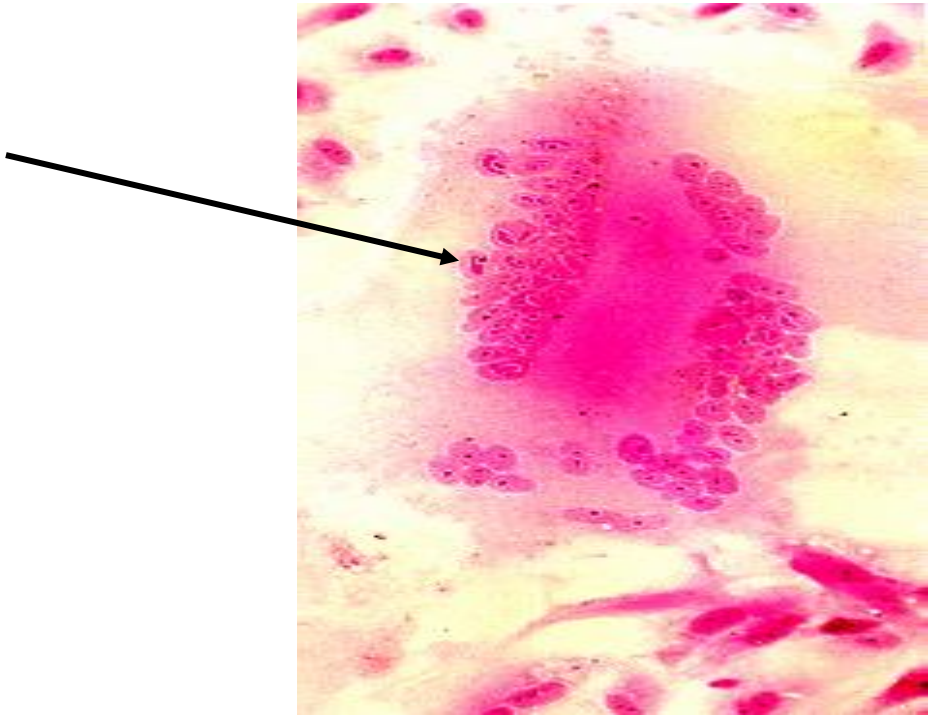
LABORATORY DIAGNOSIS OF MEASLES

Clinically Koplick's spots are pathognomonic.

1. **Detection of antigen** from nasopharyngeal aspirates and throat swab **by IF**.
2. **The virus isolation** in cell culture.

   CPE: giant multinucleated cells formation.

   Identification: **HAI, IF, NT.**
3. **Serology**: **HAI, NT, CFT.**



TREATMENT

- Supportive, symptom-directed
  - Antipyretics for fever
  - Bed rest
  - Adequate fluid intake
  - Be protected from exposure to strong light
- Antibiotics for otitis media, pneumonia
- High doses Vitamin A in severe/ potentially severe measles/ patients less than 2 years
  - 100,000IU—200,000IU

PREVENTION

- 1. Quarantine period

  5 days after rash appears, longer for complicated measles

- 2. Vaccine

  The initial measles immunization is recommended at 8mo of age

  A second immunization is recommended routinely at 7yr of age

- 3. Postexposure Prophylaxis

  Passive immunization with immune globulin (0.25mL/kg) is effective for prevention and attenuation of measles within 5 days of exposure.

Measles

IT ISN'T JUST A LITTLE RASH

Measles can be dangerous, especially for babies and young children.

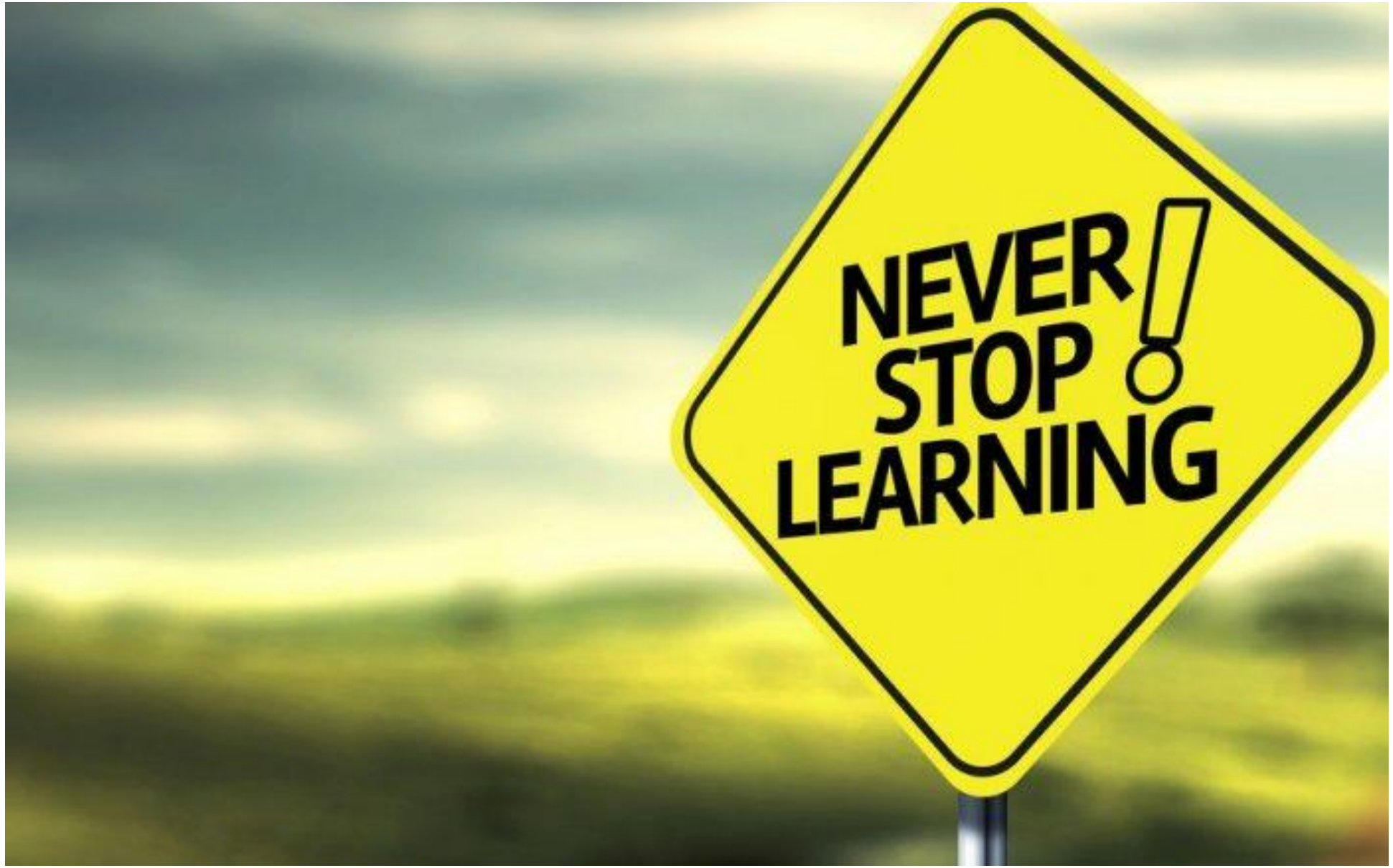
Protect your child from measles

Give your child the best protection against measles with **two** doses of measles-mumps-rubella (MMR) vaccine:

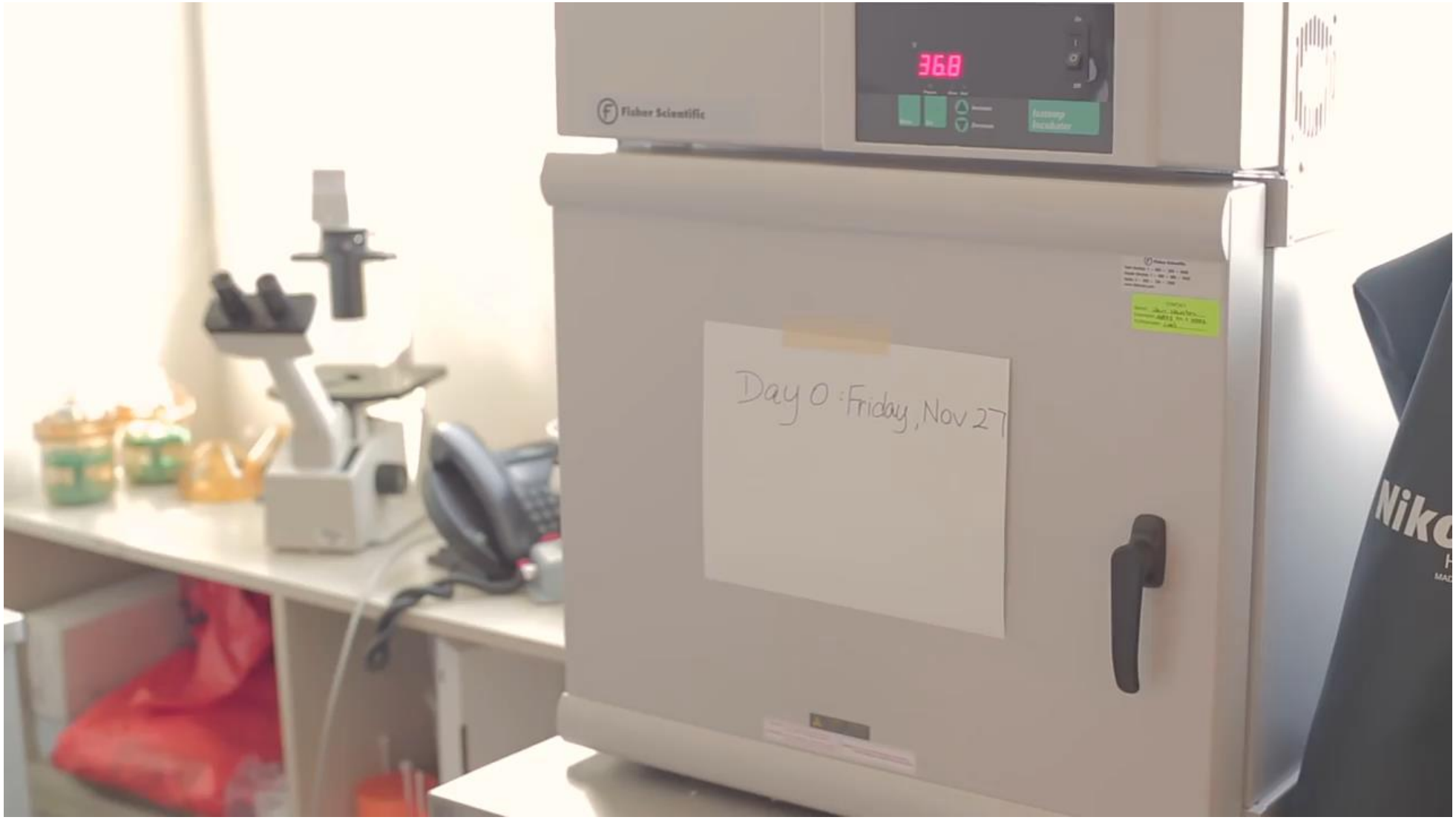
1st dose at **12-15 months**

2nd dose at **4-6 years**

Since vaccinations began in 1963, cases of **MEASLES** have dropped by **99%**


NEVER
STOP
LEARNING

İNOCULATION OF VİRUSES İN EMBRYONATED CHICKEN EGGS



DISSECTION AND EXAMINATION OF THE INFECTED EMBRYO