

Intro. to Medical Virology



LECTURE FIVE

Objectives:

- General characteristics of viruses.
- Structure & symmetry of viruses.
- Classification of viruses.
- Steps of virus replication.
- Laboratory diagnosis of viral infections.

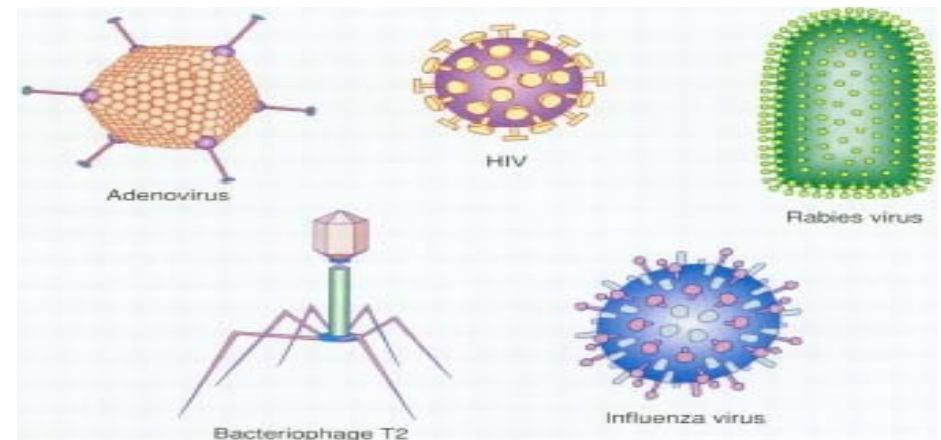
PROPERTIES OF MICROORGANISMS:

<i>characteristic</i>	<i>Parasites</i>	<i>Fungi</i>	<i>Bacteria</i>	<i>Viruses</i>
<i>Cell</i>	Yes	Yes	Yes	No
<i>Type of nucleus</i>	Eukaryotic	Eukaryotic	Prokaryotic	-----
<i>Nucleic acid</i>	Both DNA & RNA	Both DNA & RNA	Both DNA & RNA	DNA or RNA
<i>Ribosomes</i>	Present	Present	Present	Absent
<i>Mitochondria</i>	Present	Present	Absent	Absent
<i>Replication</i>	Mitosis	Budding or mitosis	Binary fission	<i>special</i>

CHARACTERISTICS OF VIRUSES

- A cellular organisms
- Tiny particles
- Obligate intracellular organisms
- Replicate in a manner diff from cells

Size ; 20-300 nm



VIRAL STRUCTURE

1-Viral genome

2-Capsid

3-Envelope

1-Viral genome:

- **DNA**

(Deoxyribonucleic acid)

All DNA Vs have ds

except

Parvoviruses

Single molecule

Or

- **RNA**

(Ribonucleic acid)

All RNA Vs have ss

except

Reoviruses

single / multiple

(+) polarity

(-) pol

2-Capsid

a protein coat

Subunits (capsomere)

Genome (NA) +

capsid

= nucleocapsid

Function:

Protects NA

Facilitates its entry into

cell

3-Envelope

Lipoprotein mb

(host lipid ,virus

specific protein)

- Budding

Envelope is derived

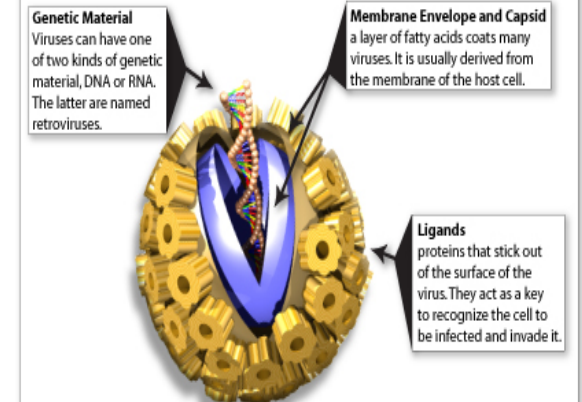
from cell mb

except herpesviruses

from nuclear mb

All Vs are haploid ,except retroviruses are diploid

VIRUS STRUCTURE



SYMMETRY (shape of the virus):

- based on arrangement of capsomere

1-Cubic symmetry (Icosahedral):

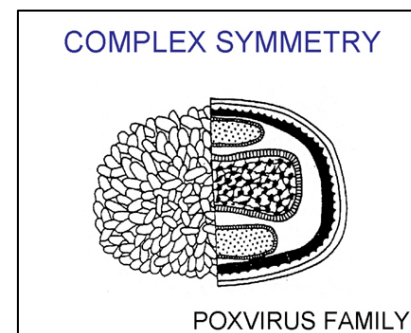
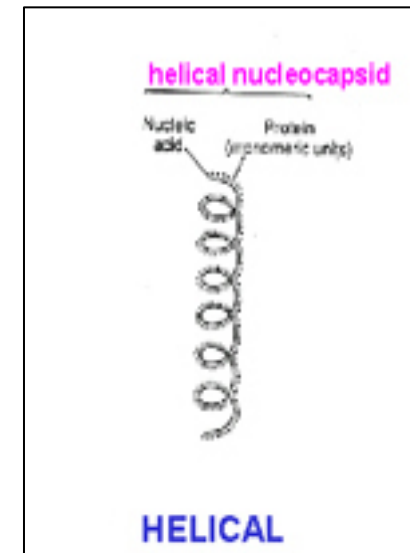
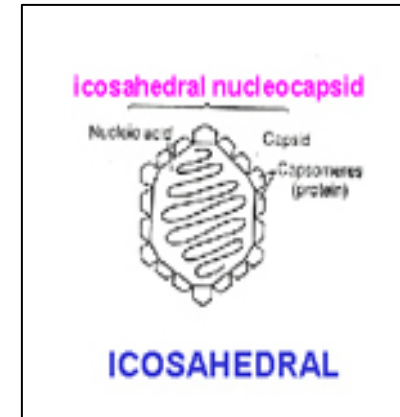
- Herpes virus
- Adenovirus

2-Helical symmetry:

- Pleomorphic(influenza v)
- Elongated (filoviruses)

3- Complex symmetry:

- poxviruses



VIRAL PROTEINS:

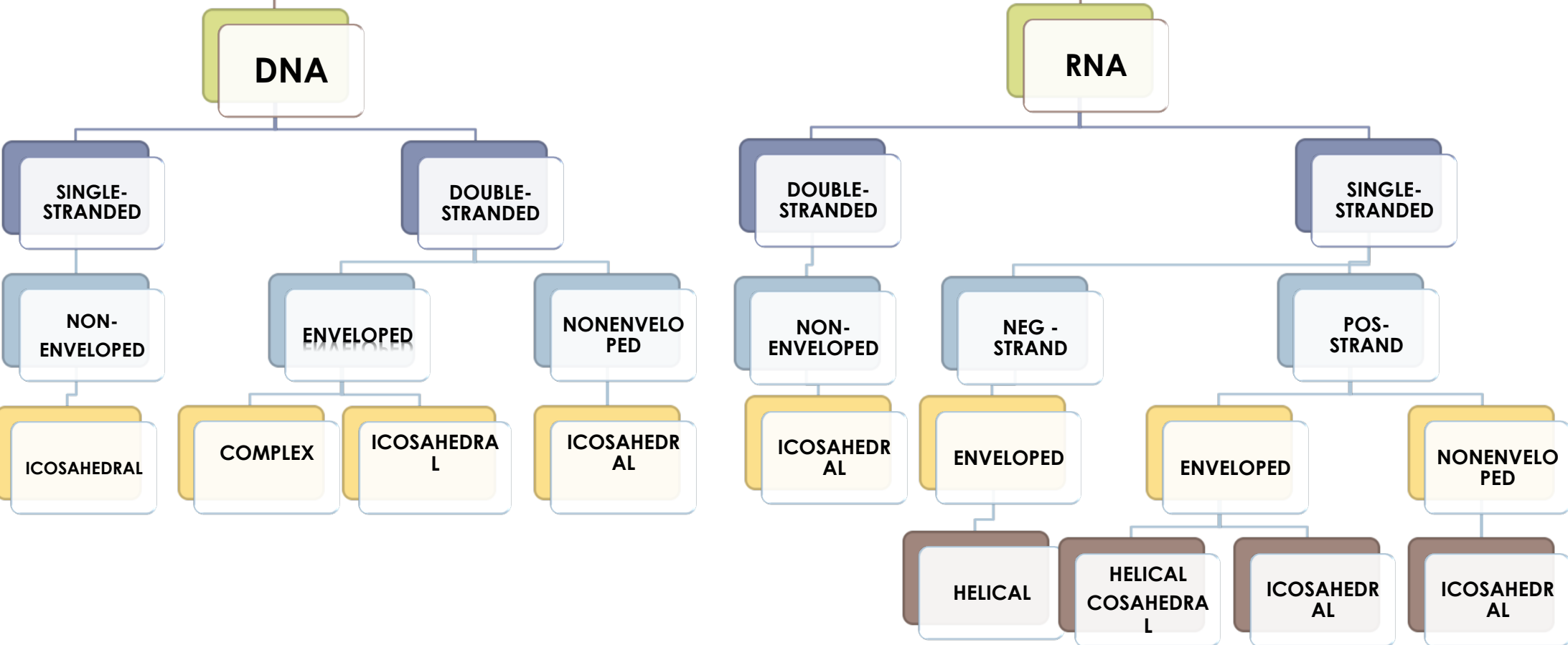
The outer viral proteins:

- Mediate attachment to specific Rs.
- Induce neutralizing Abs.
- Target of Abs.

The internal viral proteins:

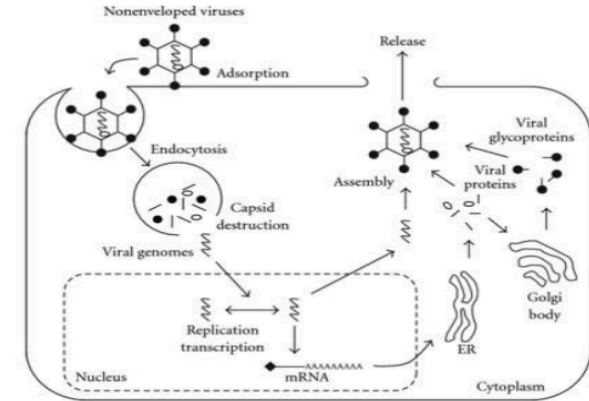
- Structural proteins (capsid proteins of enveloped Vs).
- Nonstructural proteins (enzymes):
 - All ssRNA Vs (-) polarity have **transcriptase** (RNA dependent RNA polymerase) inside virions
 - RetroVs & HBV **contain reverse transcriptase**

CLASSIFICATION OF VIRUSES



REPLICATION:

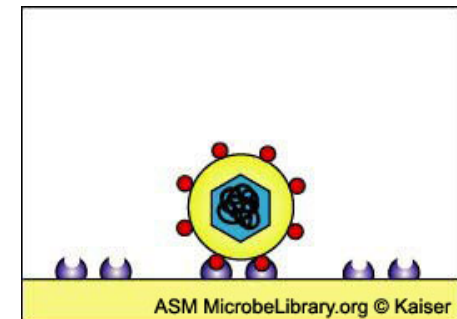
1. Adsorption (attachment)
2. Penetration
3. Uncoating
4. Synthesis of viral components
 - mRNA
 - Viral proteins
 - NA
5. Assembly
6. release



ADSORPTION (ATTACHMENT SITE) :

- 1 - some cells will have GLAYCOPROTIEN
- 2 - some will have SPECIAL FOLDING IN THE CAPSID PROTEINS.

WILL ATTACH TO A RESPTOIRER IN THE HOST CELL MEMBRAN .



PENETRATION :

ENDOCYTOSIS :

The host cells membrane will engulf the virus inside forming a ENDOCYTOTIC VESICULE :

FUSION :

The Envelope of the virus will fuse with the host cells membrane then the virus will enter the cytoplasm.

ENVOLPED VIRUSES :

Will break the vesicle throw fusion with membrane

NON-ENVOLPED VIRUSES :

Will break through the vesicle By lysis which will create pores

Either stay in the CYTOPLASM to continue to replicate

UNCOATING :

Release of viral genome

Or enter the NUCLUES

SYNTHESIS OF VIRAL COMPONENTS :

3/ replication of viral Genome:

The mechanism will continue for the replication, as we side, one virus will produce in one cell millions of viruses

1/ mRNA :

Viral Genome (+) → mRNA
Direct transcription of + single strand RNA

2/ Viral Protein :

mRNA → Enzymes/ structural protein (Capsomere)
Translated by the cells ribosomes

ASSEMBLY:

$NA + V. proteins = Virions$

1-Budding

(enveloped Vs)
-all Vs require their envelope from the cell mb*
-expect (herpes Vs.)
Require their envelope from the nuclear mb

RELEASE:

2- Cell lysis or rupture for (no enveloped) Vs

LABORATORY DIAGNOSIS OF VIRAL INFECTIONS:

- Microscopic examination.
- Cell culture.
- Serological tests .
- Detection of viral Ag.
- Molecular method .

MICROSCOPIC EXAMINATION

Light microscopy:

to see the effect of the virus in the host cell “**Histological appearance**”

Ex. Inclusion bodies

Owl's eye (CMV)

Electron microscopy:

To detect and identify the virus based on Morphology & size of virions

Ex.

- Diagnosis of viral GE such as rota, adenoviruses.
- Diagnosis of skin lesion caused by herpes, poxviruses.

This micro scope has many disadvantages

Ex:

- very expensive It is replaced by Ag detection & molecular tests

VIRUS CULTIVATION

Laboratory animal

Embryonated egg

Cell culture
Cells obtained from animals or humans

Very expensive
Very hard to handle

Cell culture	No of sub passages
Primary C/C	1 or 2
Diploid C/C (semi continuous)	20 or 50
Continuous cell line	indefinite

It is very IMP That we use a combination methods of cell culture

Detection of viral growth:

- Cytopathic effects
 - Others
- Any possible change In the cell culture

Ex: cell rounding
Syncytium .
“fusion of The inflected cells together → One Giant cell

PROBLEMS WITH CELL CULTURE:

1. Long incubation
2. Sensitivity is variable
3. Susceptible to bacterial contamination
4. Some Vs do not grow in c/c ex. HCV

RAPID CULTURE TECHNIQUE :

Since the normal cell culture had to take a month to grow they came up with a new method :

- **Shell Vial Assay**
Detect viral antigens
1-3 days

SEROLOGICAL TEST:

*Very common

*Very fast 1- 2 hours

Antigen detection:

sample	virus	test
Nasopharyngeal aspirate	Influenza V	IF
Skin scrapings	HSV	IF
Feces	Rotavirus	ELISA
Blood	HBV (HBsAg)	ELISA

Antibody detection:

Ex of techniques

Immunofluorescence (IF)

A- Direct
Antigen detection using an antigen Sample

B- Indirect
Antibody detection by using an antibody Sample

Enzyme-linked Immunosorbent test (ELISA)

A- Direct
antigen Detection Or specific IgM, IgG

B- Indirect
antibody detection
It uses an enzyme to react with an enzymatic substrate to show color

◆ colored wells indicate reactivity

MOLECULAR TEST:

- **Polymerase chain reaction (PCR)**
 - Amplification tech.
 - Viral genome
- **Uses of molecular test:**
 - Diagnosis of viral disease
 - Monitoring response to treatment

Thank you

قال تعالى: (وَالَّذِينَ تَبَوَّءُوا الدَّارَ وَالْإِيمَانَ مِنْ قَبْلِهِمْ يُحِبُّونَ مَنْ
هَاجَرَ إِلَيْهِمْ وَلَا يَجِدُونَ فِي صُدُورِهِمْ حَاجَةً مِمَّا أُوتُوا وَيُؤْثِرُونَ عَلَى
أَنْفُسِهِمْ وَلَوْ كَانَ بِهِمْ خَصَاصَةٌ وَمَنْ يوق شح نفسه فَأُولَئِكَ هُمُ
الْمُفْلِحُونَ)

DONE BY:



- Alhanouf AlMohanna



- AlJouhara AlDahsh



- Abudalaziz AlMani



- Amal Afrah



- Aya AlDayel



- Deema AlRajhi



- Dhaherah AlJohani



- Hanan Khoshaim



- Jawaher AlOmran



- Manal AlHamdan



- Nouf AlMasood



- Rawa AlOhali



- Reema Hazazi



- Reema AlHammad



- Wajda AlHathali

MCQs:

• **The genetic material in viruses is:**

a) DNA only b) **Either DNA or RNA** c) Both DNA & RNA d) RNA only

• **Which one of the following is type of virus symmetry:**

a) **Complex symmetry** b) Columnar symmetry c) Simple symmetry

• **Molecular test is used for:**

a) Prognosis b) Pathogenesis c) **Diagnosis**

• is an example of a virus that doesn't grow in cell culture :

a) **HCV** b) HIV c) CMV

• **Viruses have organelles such as mitochondria.**

a) T b) **F**

Related Videos:

- <http://youtu.be/uwiPidO7K2s>
- <http://youtu.be/s8jhJXgC-bk>
- http://youtu.be/L8oHs7G_syl
- <http://youtu.be/eS1GODinO8w>

