

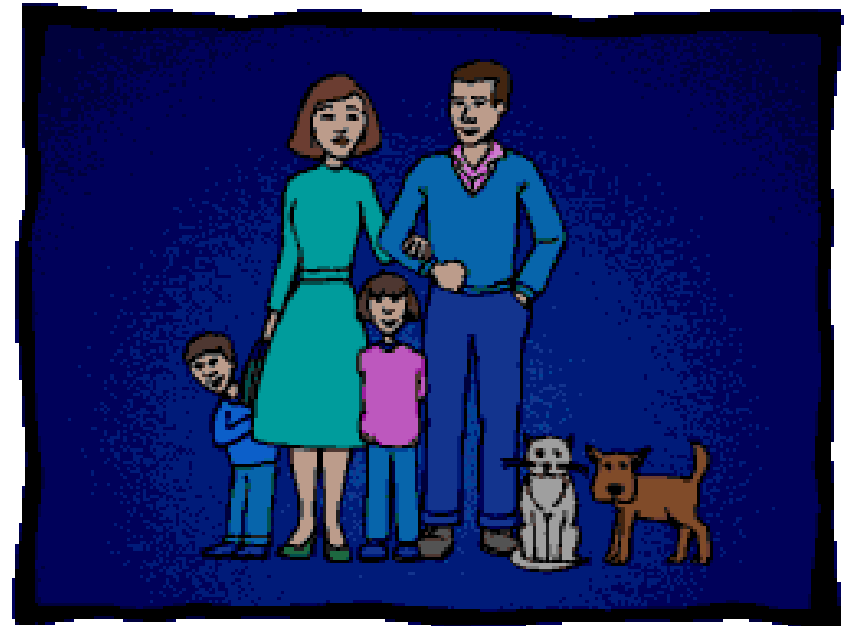
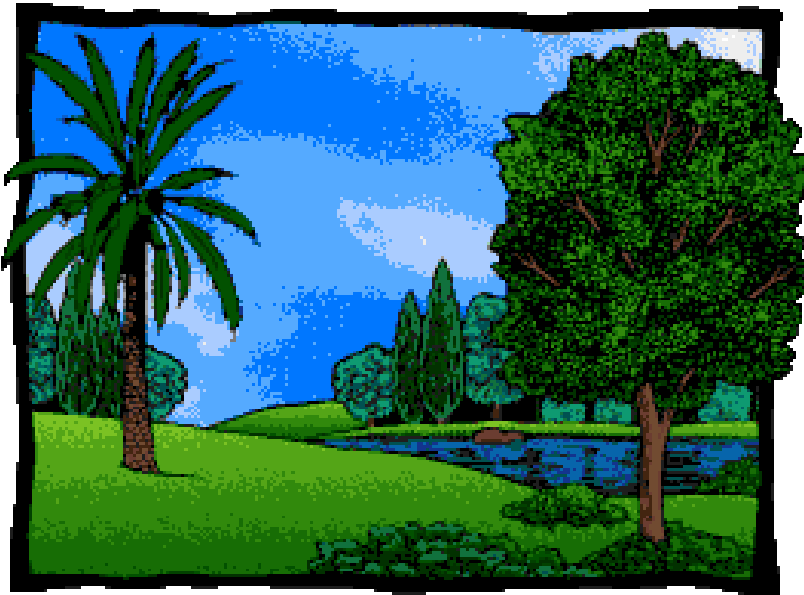
An Introduction to Viruses

Lecturer

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Virus infections are Universal



Introduction to Virology

- A virus is an obligate intracellular parasite containing genetic material surrounded by protein
- Virus particles can only be observed by an **electron microscope**



Introduction to Virology

- Recognizing the shape, size, and structure of different viruses is critical to the study of disease
 - Viruses have an inner core of nucleic acid surrounded by protein coat known as capsid
 - Most viruses range in sizes from 20 – 450 nanometers

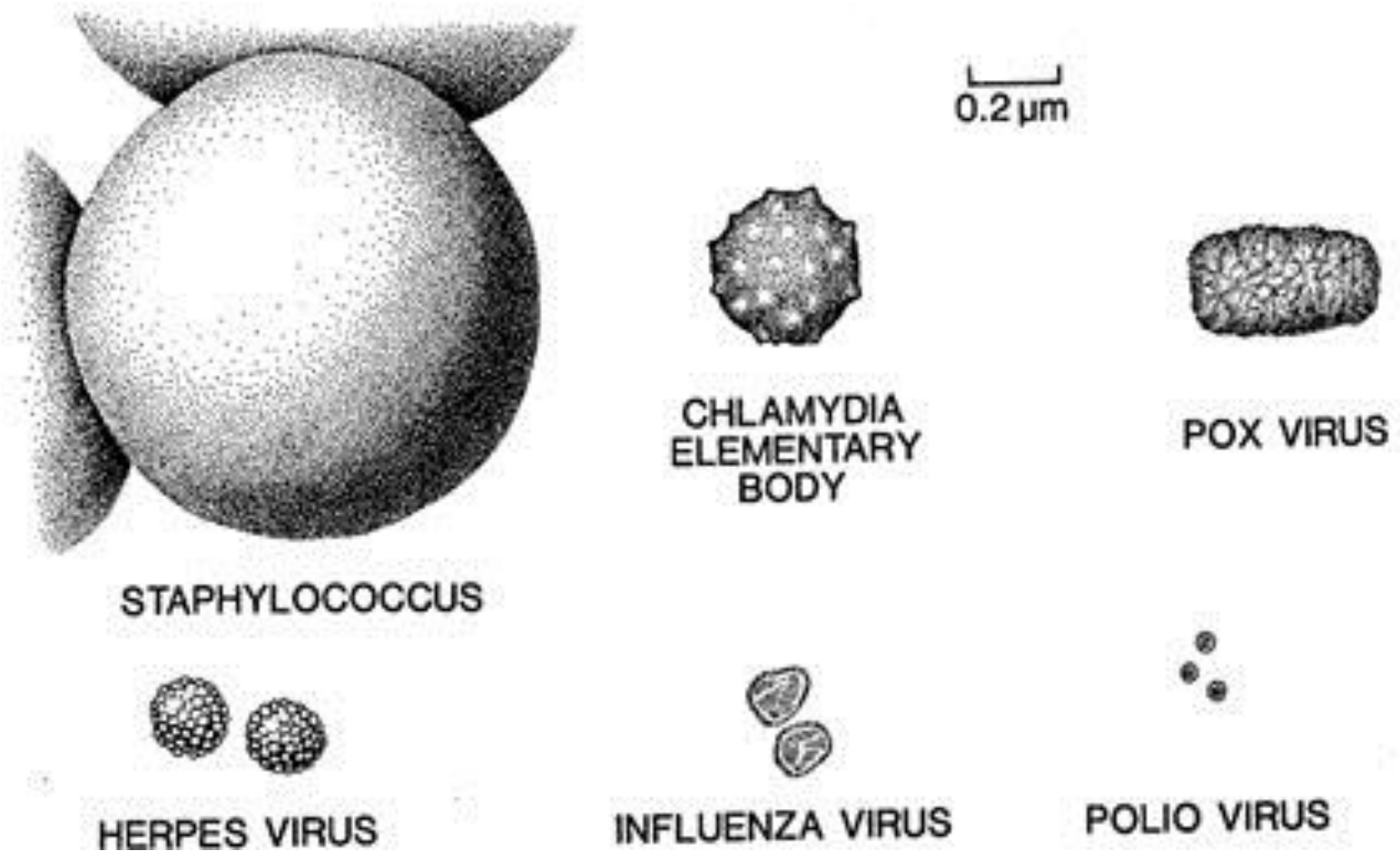
Viral Properties

- **Viruses are inert (nucleoprotein) filterable Agents**
- **Viruses are obligate intracellular parasites**
- **Viruses cannot make energy or proteins independent of a host cell**
- **Viral genome are RNA or DNA but not both.**
- **Viruses have a naked capsid or envelope with attached proteins**
- **Viruses do not have the genetic capability to multiply by division.**
- **Viruses are non-living entities**

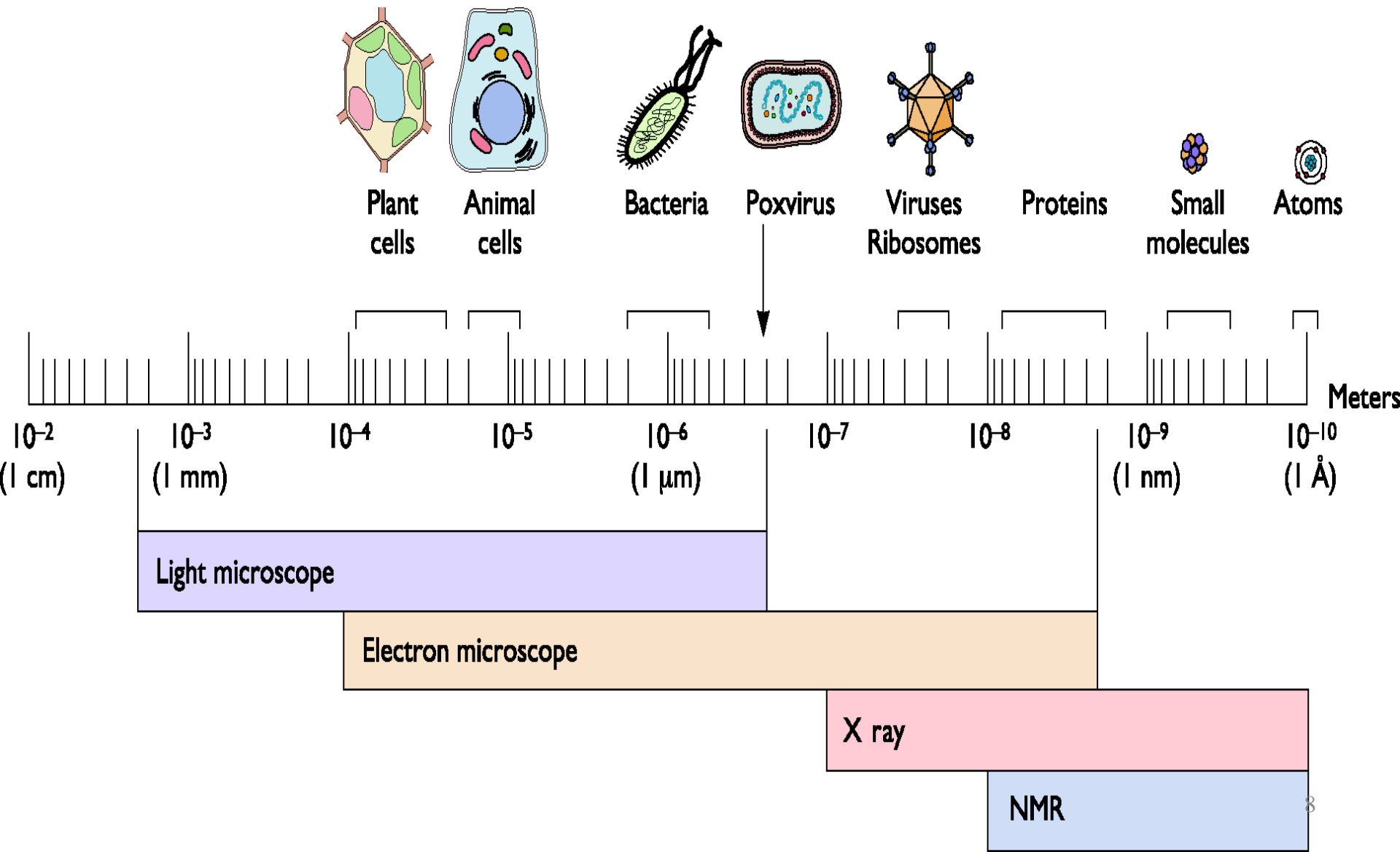
Virus vs. cells

Property	Viruses	Cells
Type of nucleic acid	DNA or RNA	DNA and RNA
Proteins	Few	Many
Lipoprotein membrane	Enveloped present in some viruses	Cell membrane present in all cells
Ribosomes	Absent	Present
Mitochondria	Absent	Present in eukaryotic cells
Enzymes	None or few	Many
Multiplication by binary fission	No	Yes (most cells)

Viruses are Ultramicroscopic



The size of viruses

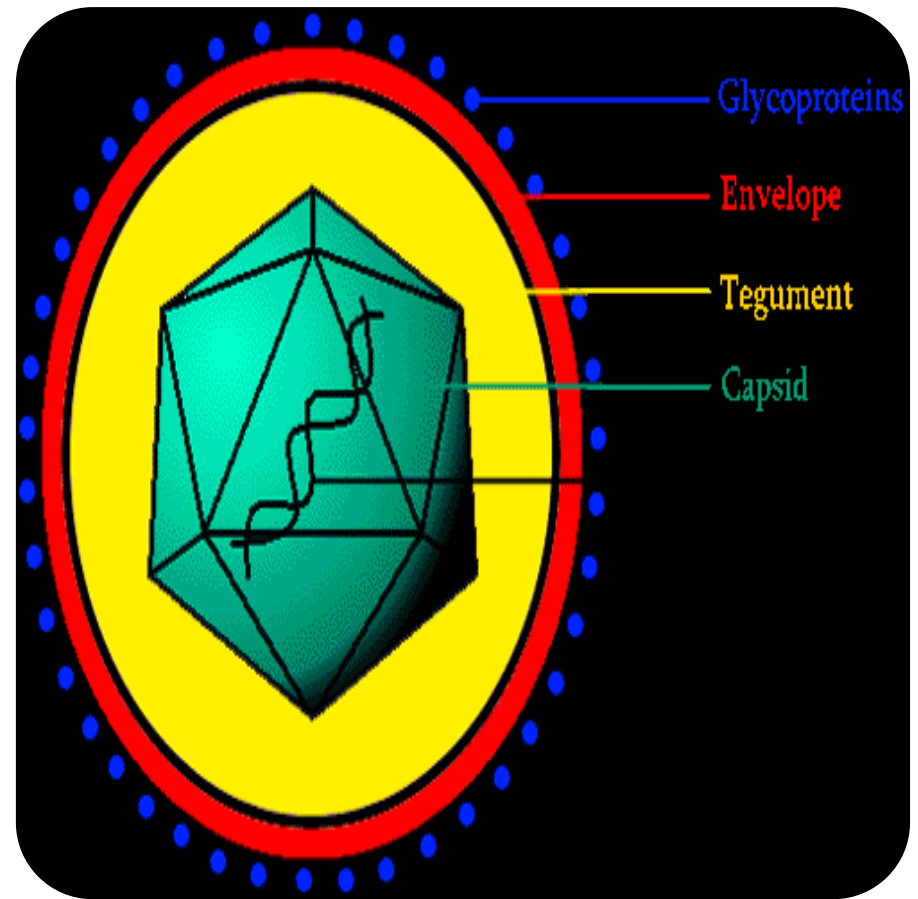


VIRAL STRUCTURE – SOME TERMINOLOGY

- virus particle = virion
- protein which coats the genome = capsid
- capsid usually symmetrical
- capsid + genome = nucleocapsid
- may have an envelope

Virion

- The complete infectious unit of virus particle
- Structurally mature, extracellular virus particles.



Viral Structure - Overview

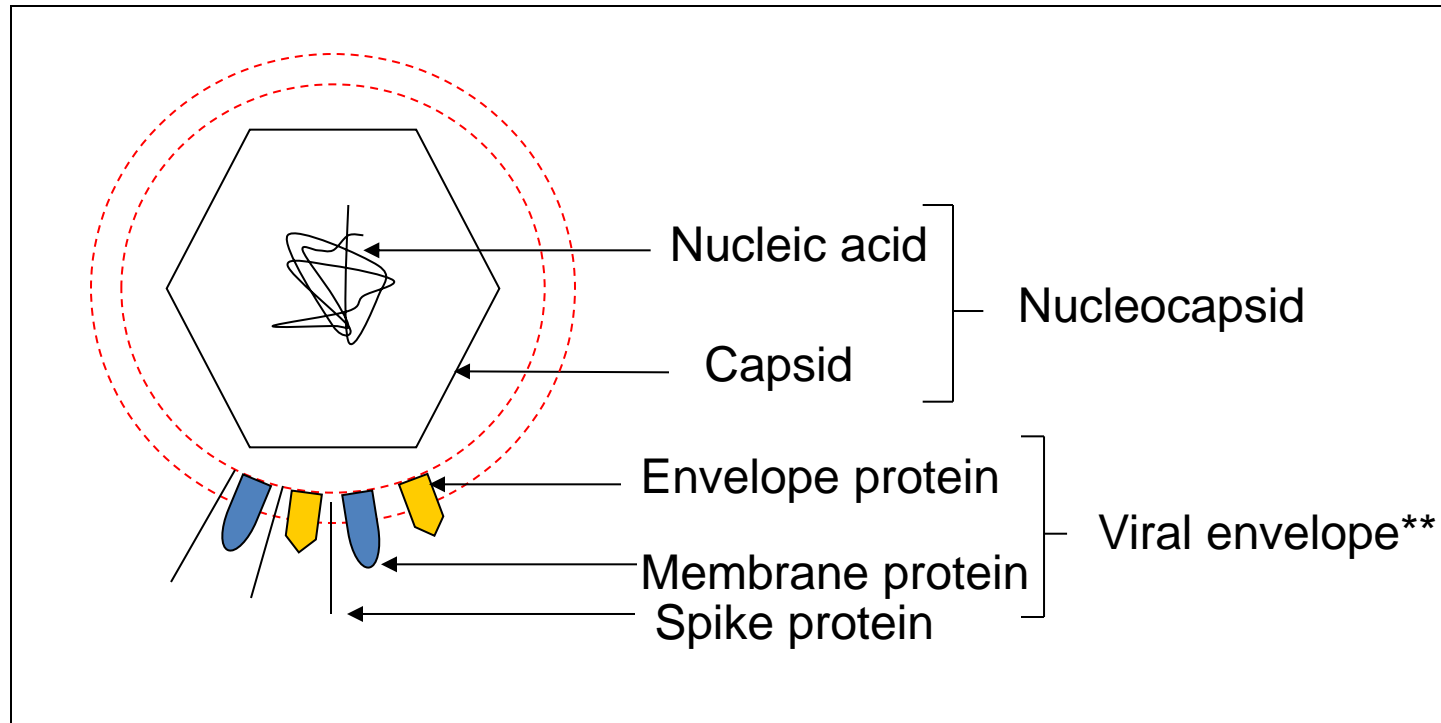


Fig 1. Schematic overview of the structure of animal viruses

** does not exist in all viruses

Distinguishing characteristics of viruses

- Obligate intracellular parasites
- Extreme genetic simplicity
- Contain DNA or RNA
- Replication involves disassembly and reassembly
- Replicate by "one-step growth"

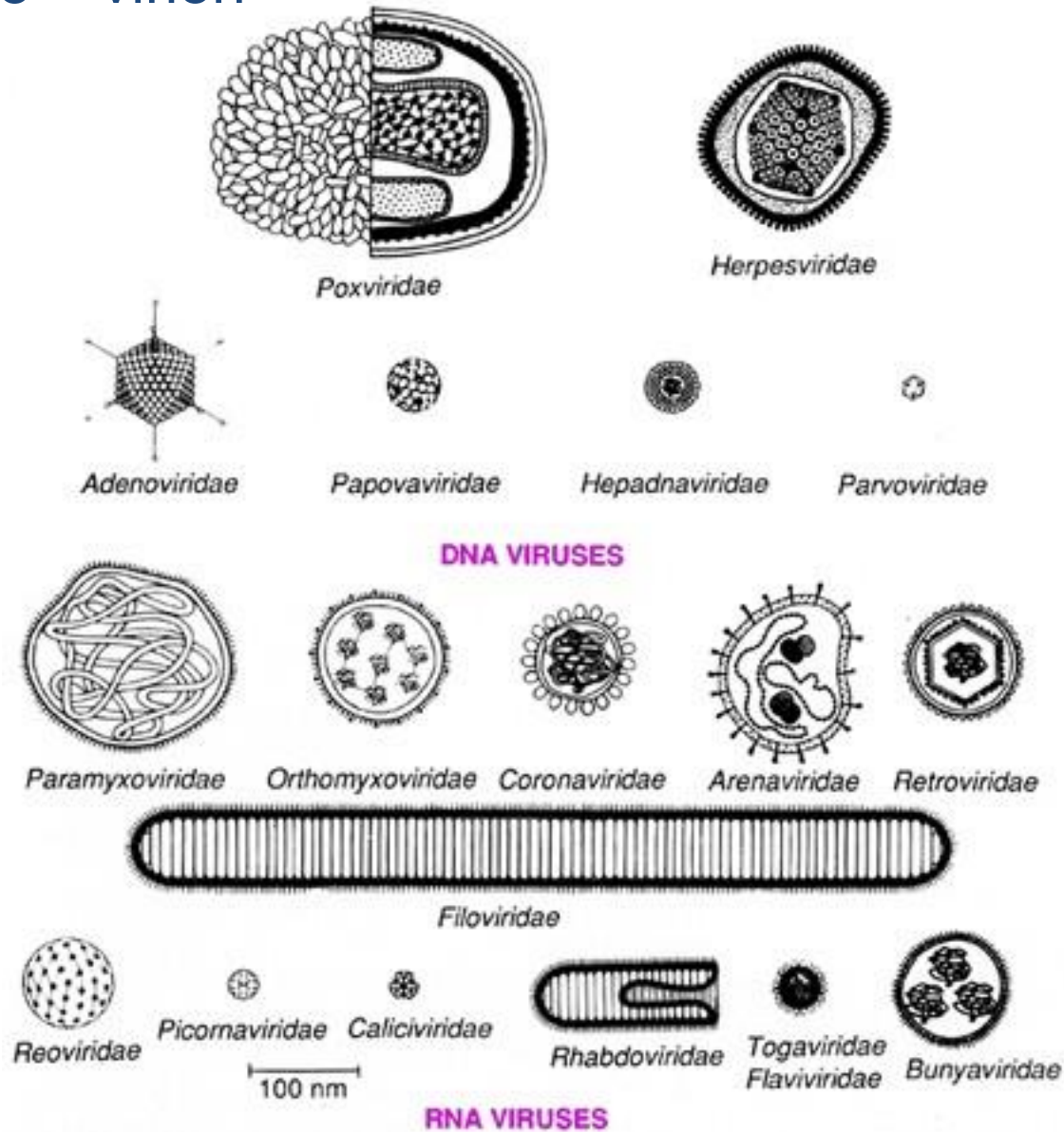
Naming viruses

- No taxa above Family (no kingdom, phylum, etc)
- Classified based on structures, size, nucleic acids, host species, target cells.
- 20 families of **animal** viruses (7 DNA, 13 RNA)
- Family name ends in – viridae
- Subfamily ends in — virinae
- Genus name ends in – virus
- Species
 - Example
 - Family – Herpesviridae
 - Subfamily - Herpesvirinae
 - Genus – Simplex virus
 - Common name – herpes virus (Herpes simplex virus I (HSV-I))
 - Disease – fever blisters, cold sores

How are viruses named?

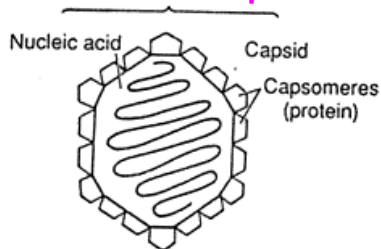
- Based on:
 - the disease they cause
poliovirus, rabies virus
 - the type of disease
murine leukemia virus
 - geographic locations
Sendai virus, Cocksackie virus
 - their discoverers
Epstein-Barr virus
 - how they were originally thought to be contracted
dengue virus (“evil spirit”), influenza virus (the “influence” of bad air)
 - combinations of the above
Rous Sarcoma virus

Virus particle = virion

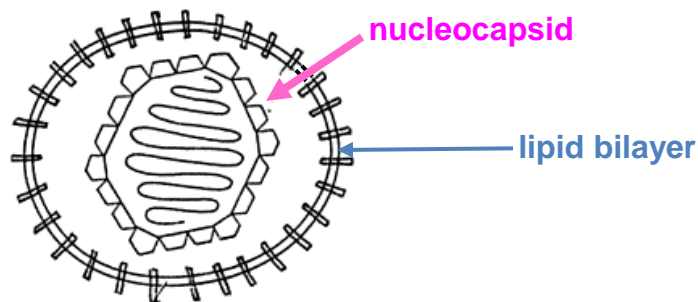


5 BASIC TYPES OF VIRAL STRUCTURE

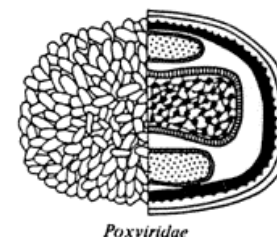
icosahedral nucleocapsid



ICOSAHEDRAL

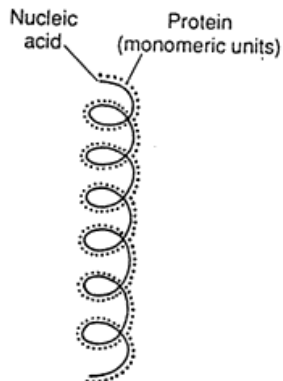


ENVELOPED ICOSAHEDRAL

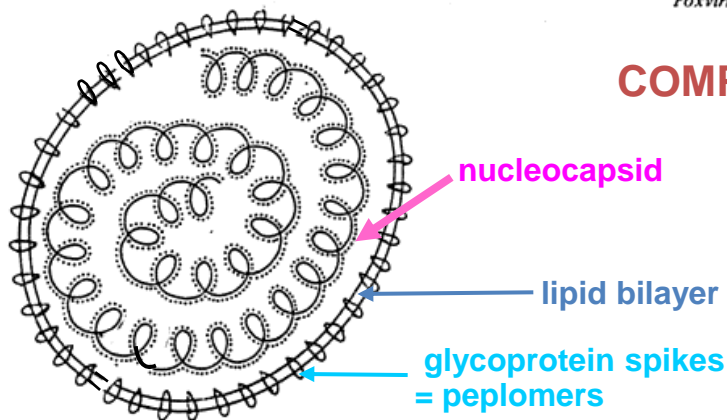


COMPLEX

helical nucleocapsid



HELICAL



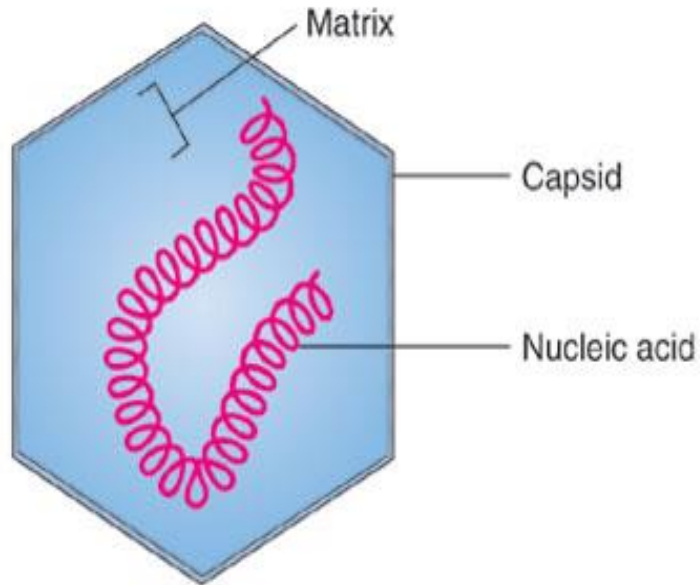
ENVELOPED HELICAL

Viral Structure

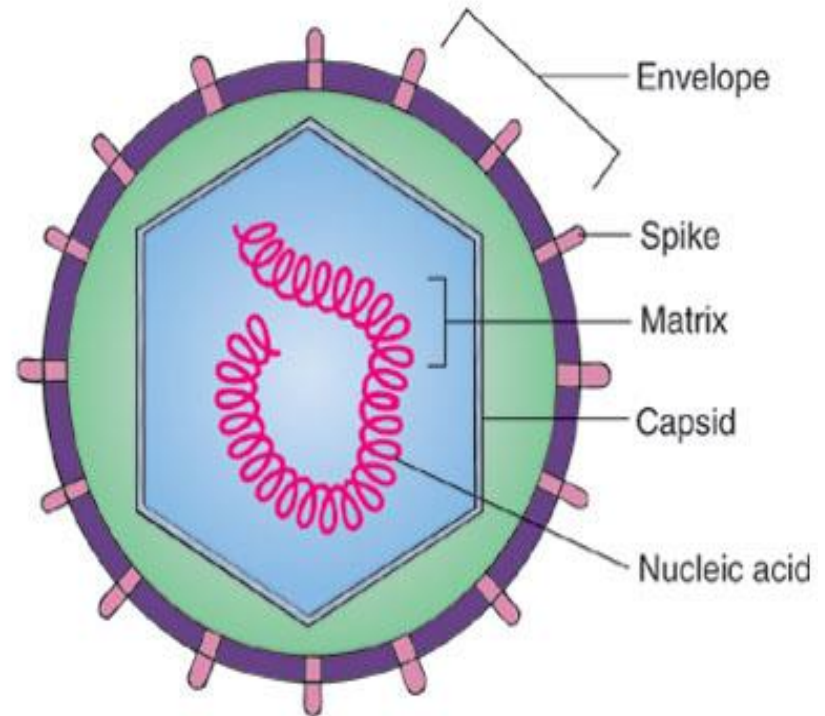
- Varies in size, shape and symmetry
- 3 types of capsid symmetry:
 - Cubic (icosahedral)
 - Has 20 faces, each an equilateral triangle. Eg. adenovirus
 - Helical
 - Protein binds around DNA/RNA in a helical fashion eg. Coronavirus
 - Complex
 - Is neither cubic nor helical eg. poxvirus

VIRAL STRUCTURE (virion)

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(a) Naked Nucleocapsid Virus



(b) Enveloped Virus

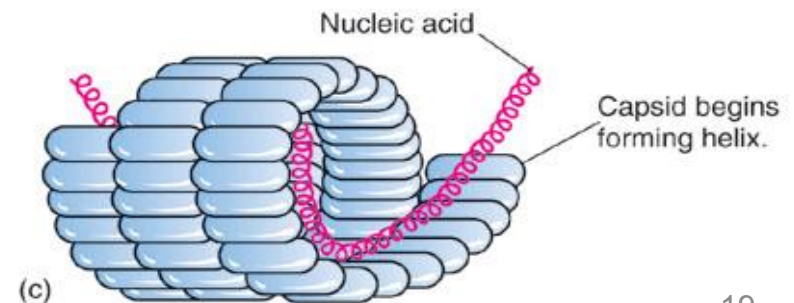
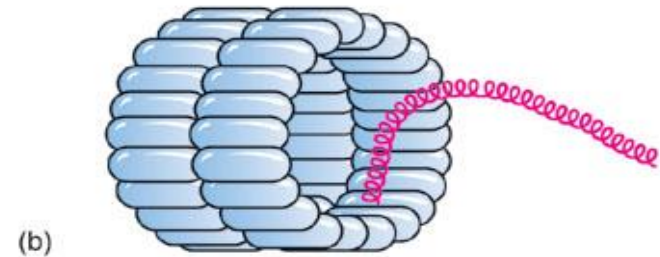
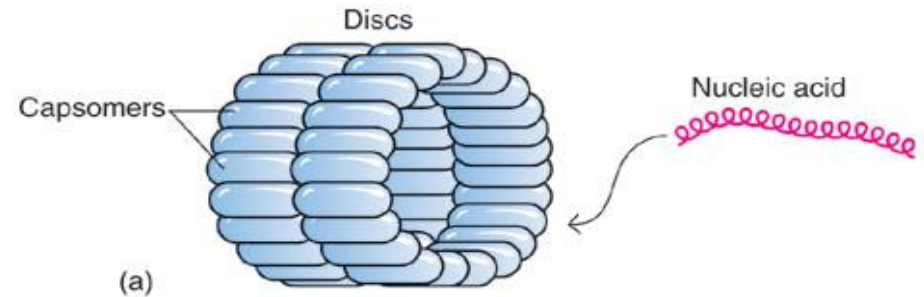
1. Protect genome during passage from one cell to another
2. Aid in entry process
3. Package enzymes for early steps of infection

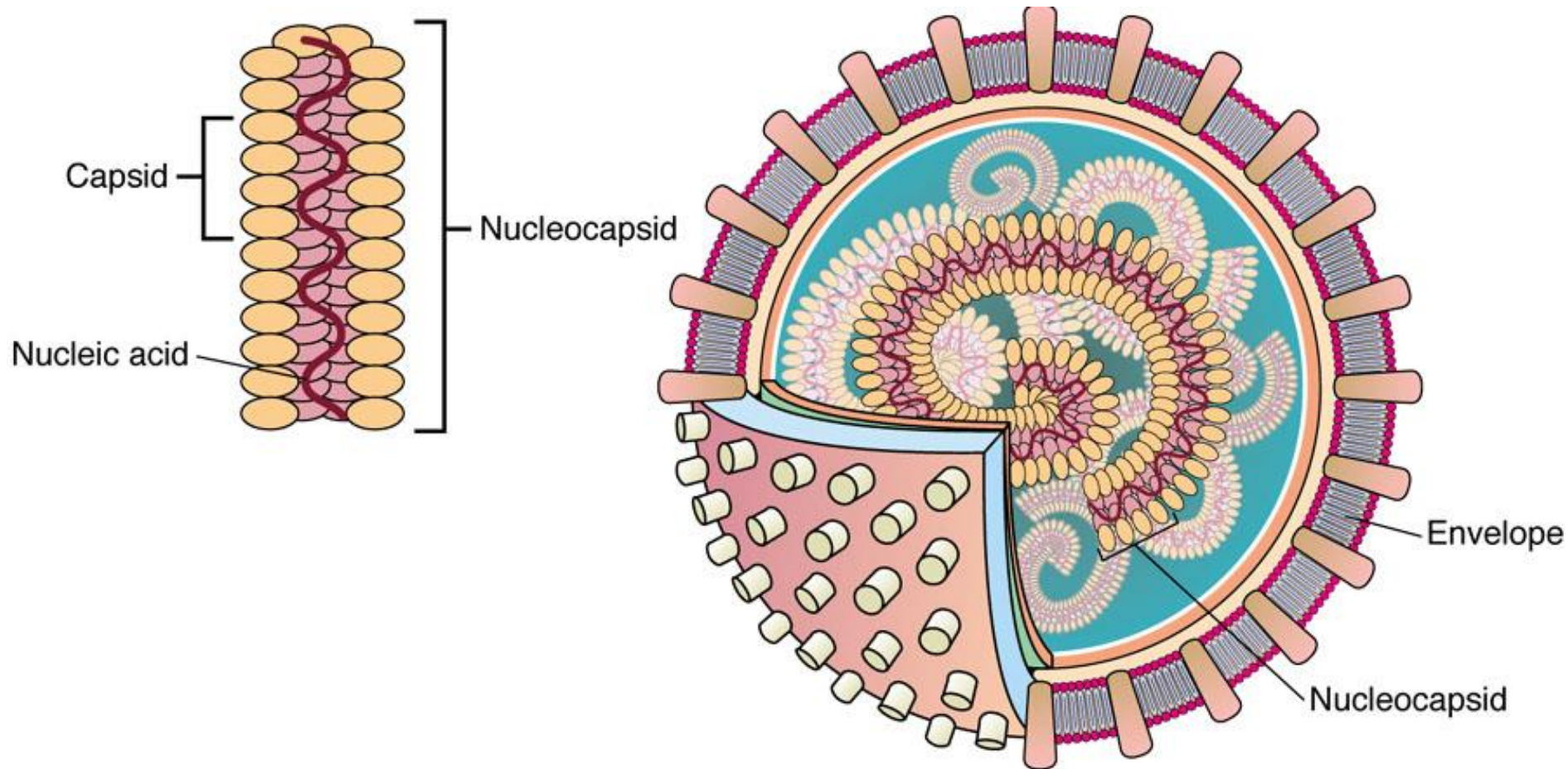
CAPSID STRUCTURE

1. Helical capsid

- Rod-shaped capsomers
- Coil around hollow center
- Nucleic acid is kept inside – wound-up within tube (**Helix**)

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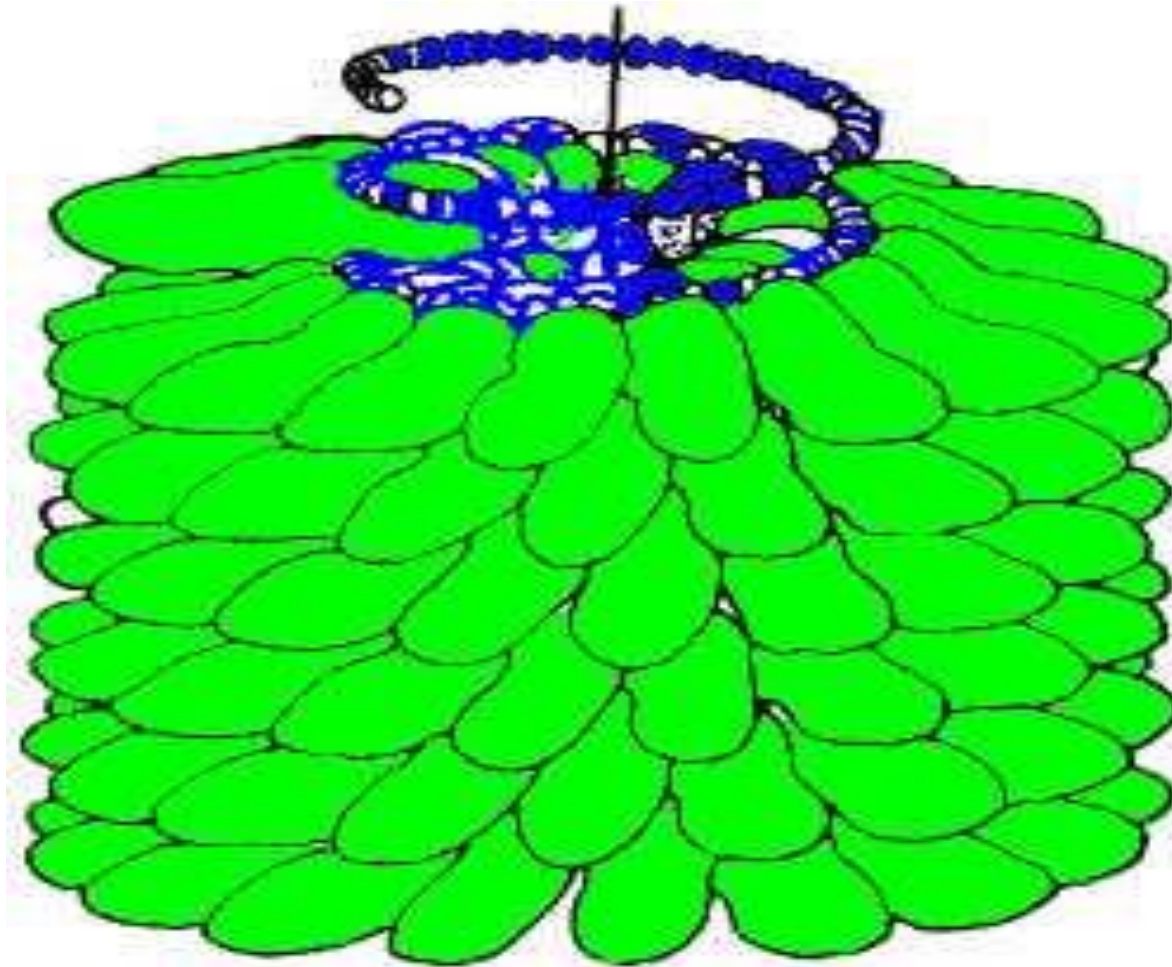




Helical – capsid surrounds DNA like hollow tube

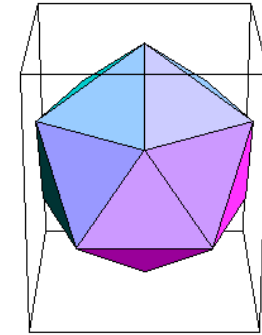
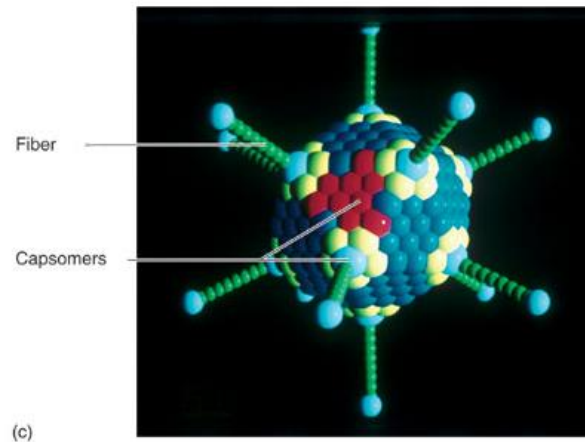
Ex: Influenza , measles, rabies (enveloped)

Helical symmetry

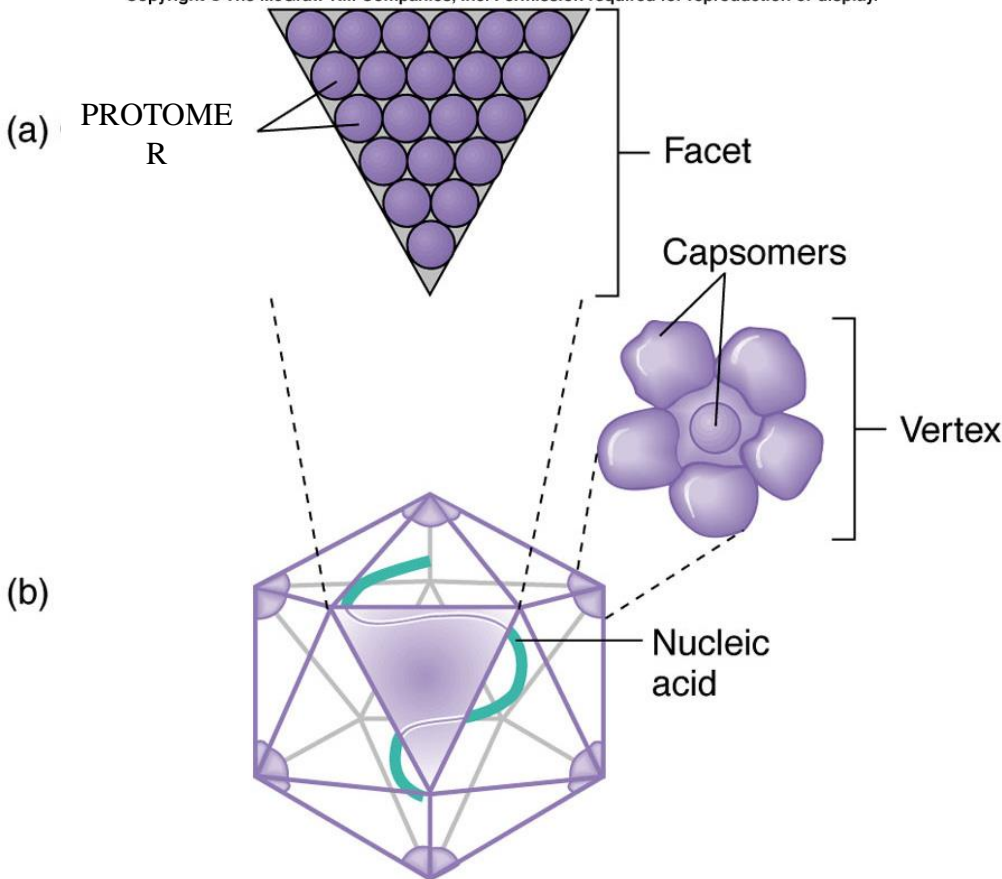


2. icosahedral

Morphological types

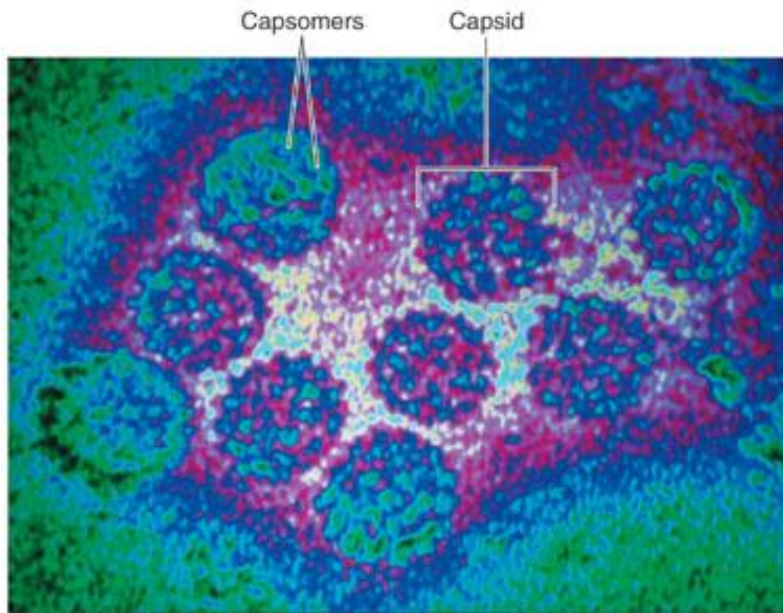


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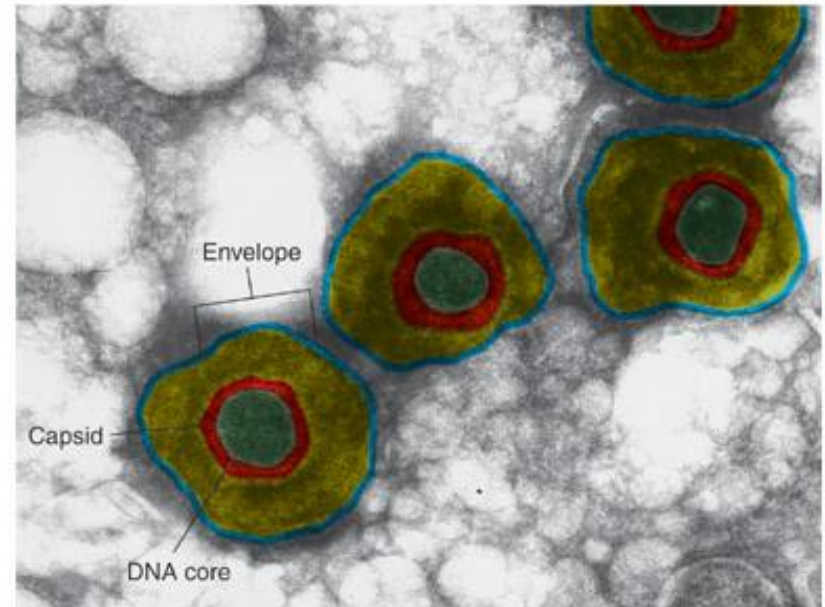


- 20-sided with 12 corners
- Vary in the number of capsomers
- Each capsomer may be made of 1 or several proteins
- Some are enveloped

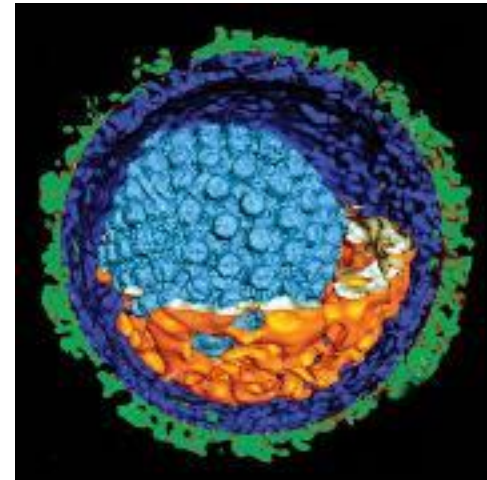
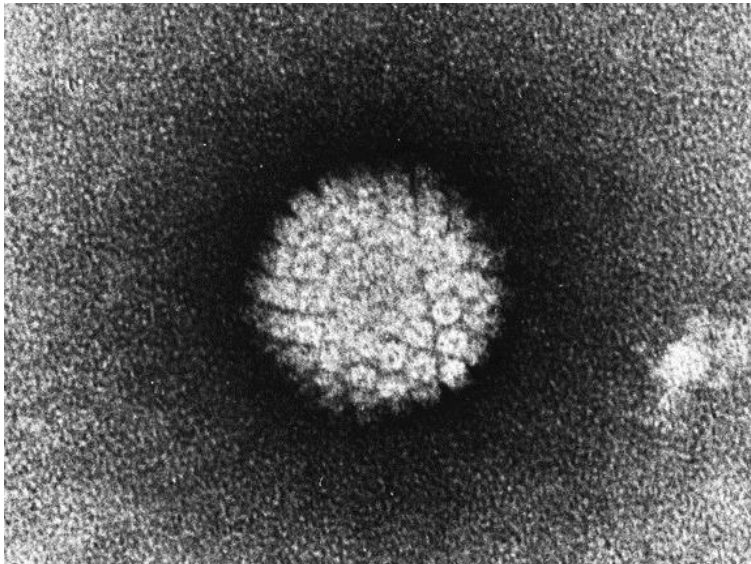




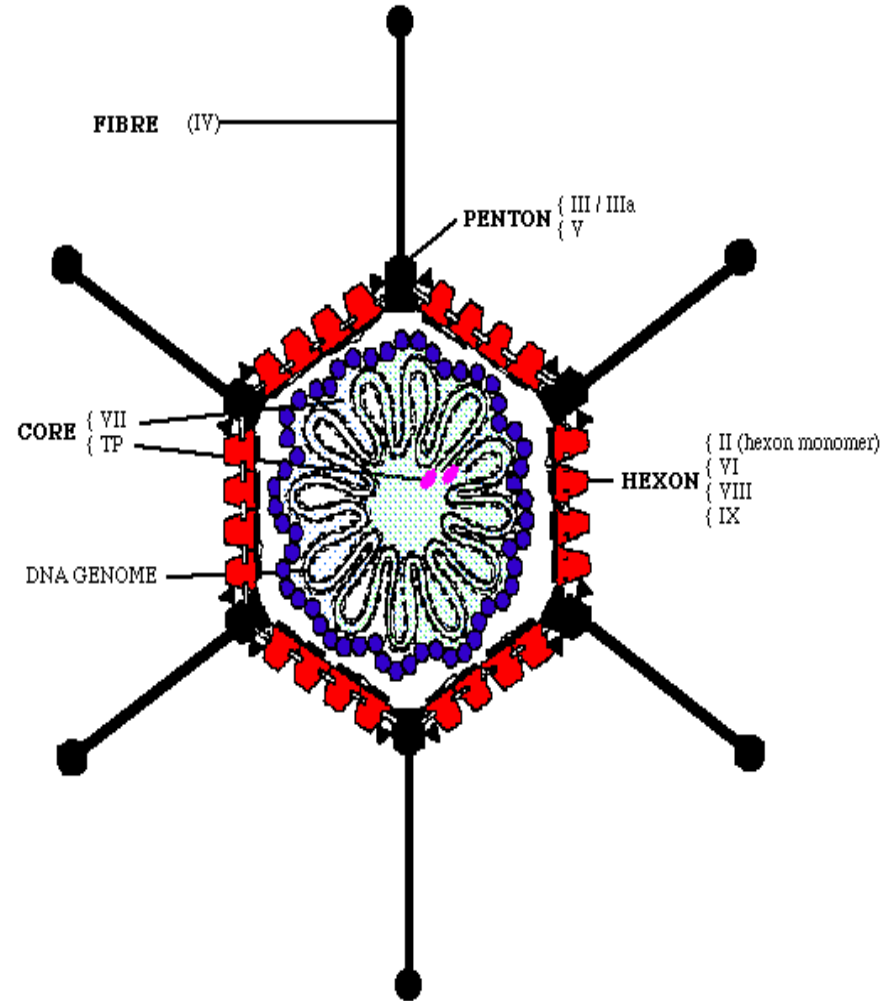
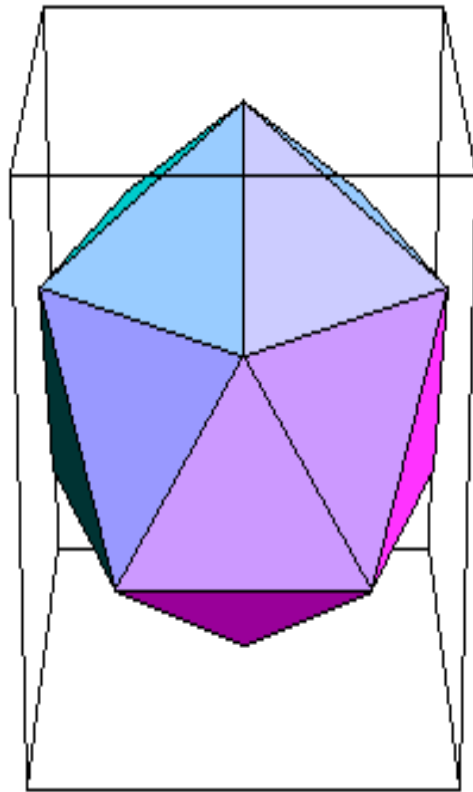
(a)



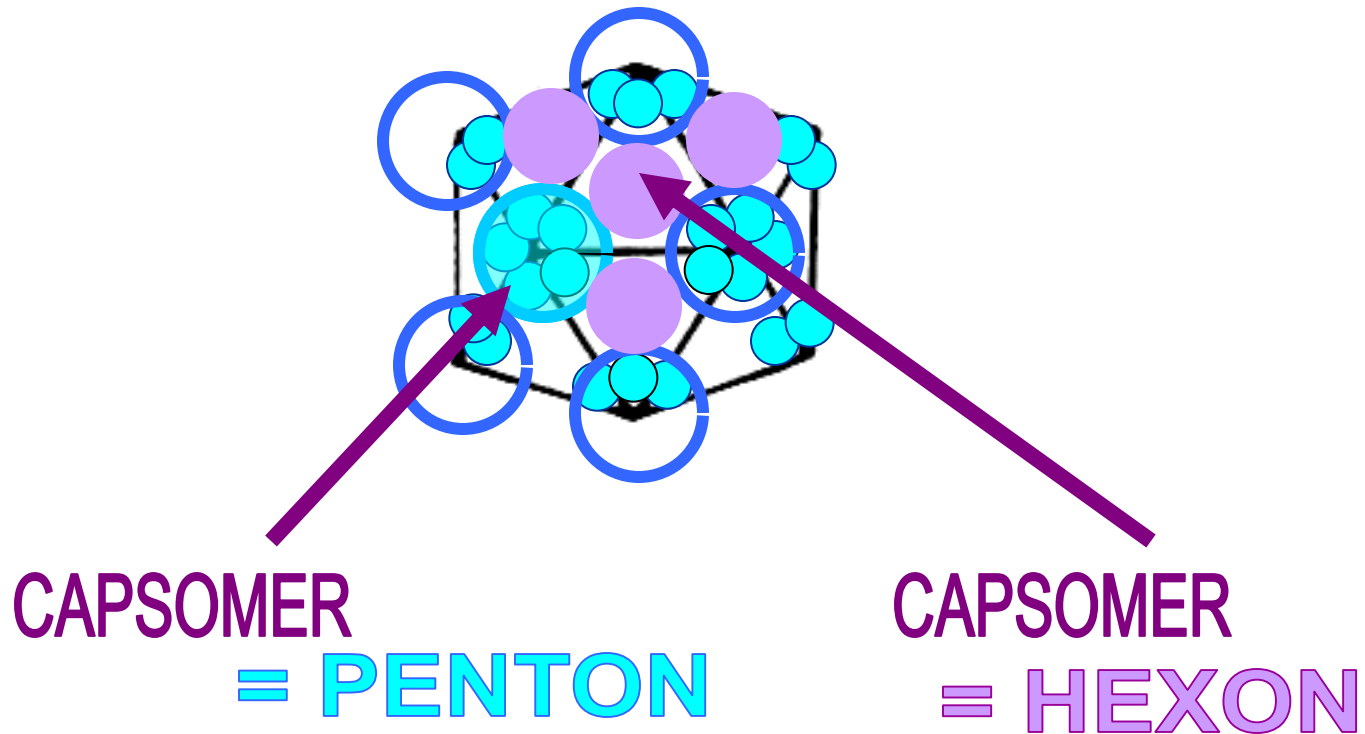
(b)

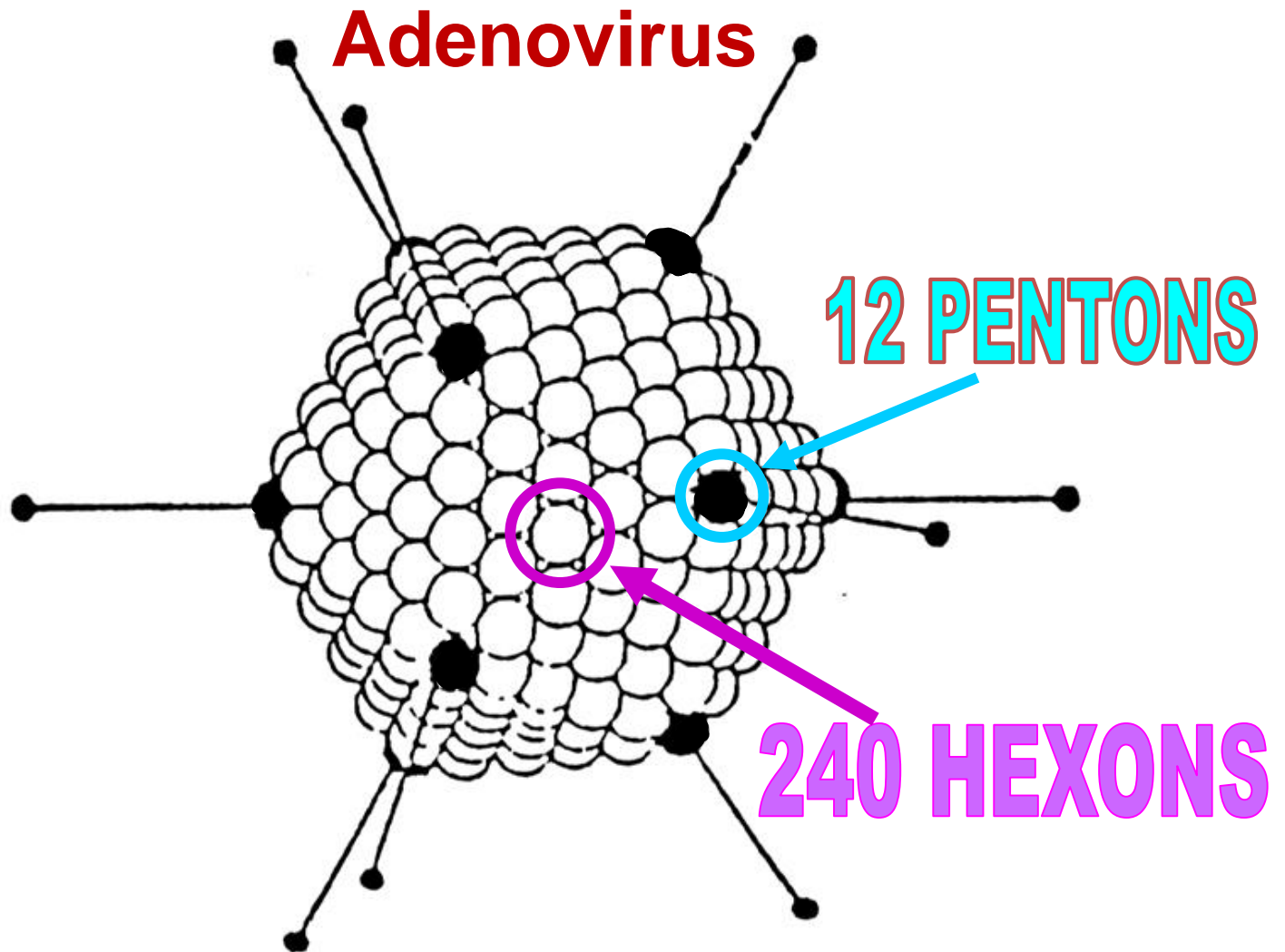


Cubic or icosahedral symmetry

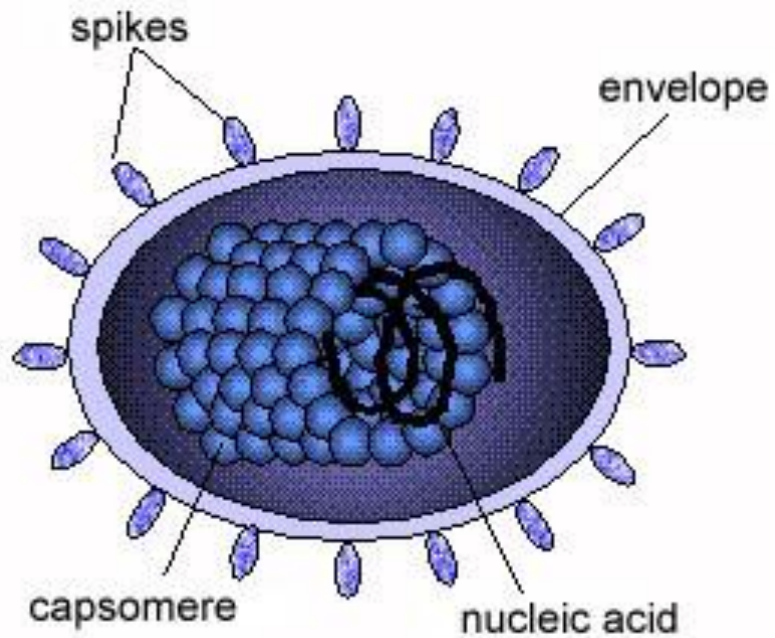


ICOSAHEDRAL SYMMETRY

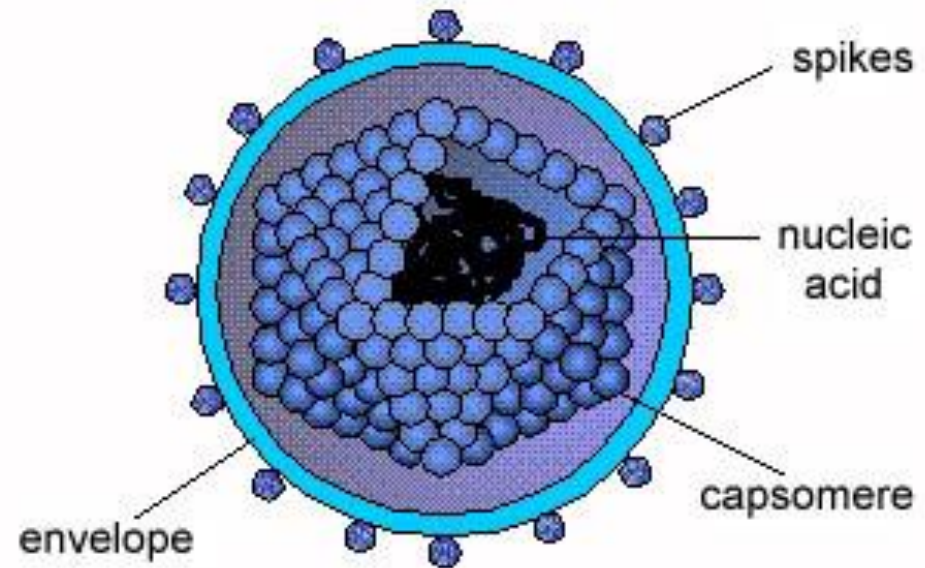




Enveloped helical virus



Enveloped icosahedral virus



Helical

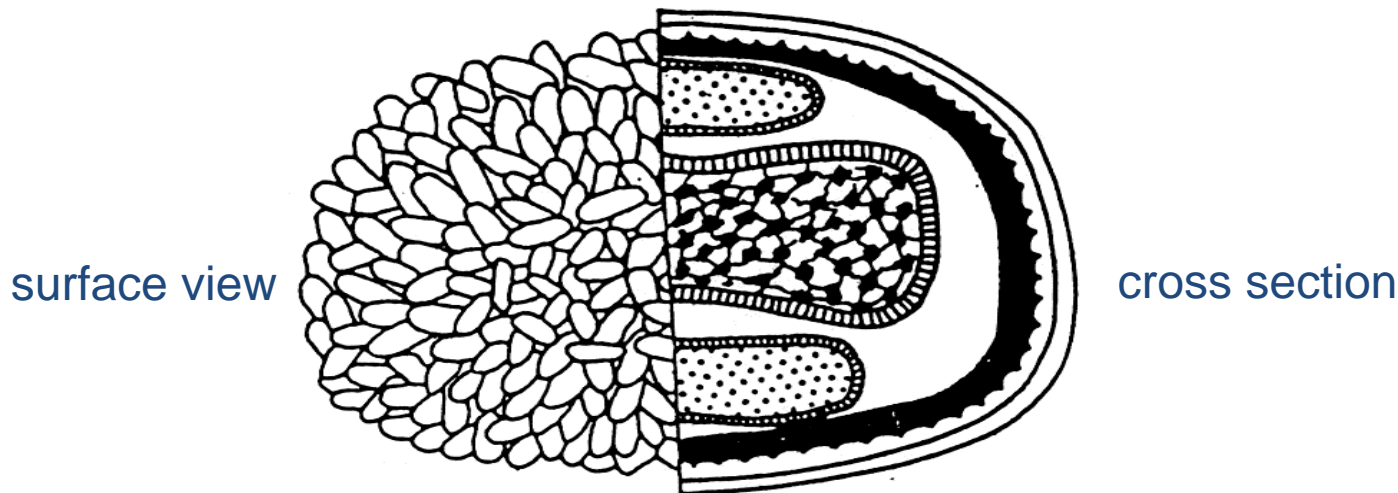
- **California Encephalitis Virus**
Coronavirus
Hantavirus
Influenza Virus (Flu Virus)
Measles Virus (Rubeola)
Mumps Virus
Para influenza Virus
Rabies Virus
Respiratory Syncytial Virus(RSV)

Icosahedral

- Adeno-associated Virus (AAV)
Adenovirus
B19
Coxsackievirus - A
Coxsackievirus - B
Cytomegalovirus (CMV)
Eastern Equine Encephalitis Virus (EEEV)
Echovirus
Epstein-Barr Virus (EBV)
Hepatitis A Virus (HAV)
Hepatitis B Virus (HBV)
Hepatitis C Virus (HCV)
Hepatitis Delta Virus (HDV)
Hepatitis E Virus (HEV)
- Herpes Simplex Virus 1 (HHV1)
Herpes Simplex Virus 2 (HHV2)
Human Immunodeficiency Virus (HIV)
Human T-lymphotrophic Virus (HTLV)
Norwalk Virus
Papilloma Virus (HPV)
Polio virus
Rhinovirus
Rubella Virus
Saint Louis Encephalitis Virus
Varicella-Zoster Virus (HHV3)
Western Equine Encephalitis Virus (WEEV)
Yellow Fever Virus

Complex viruses

- Have additional or special structures
- Examples:
- **Poxviruses** – lack normal capsid – instead, layers of lipoproteins and fibrils on surface

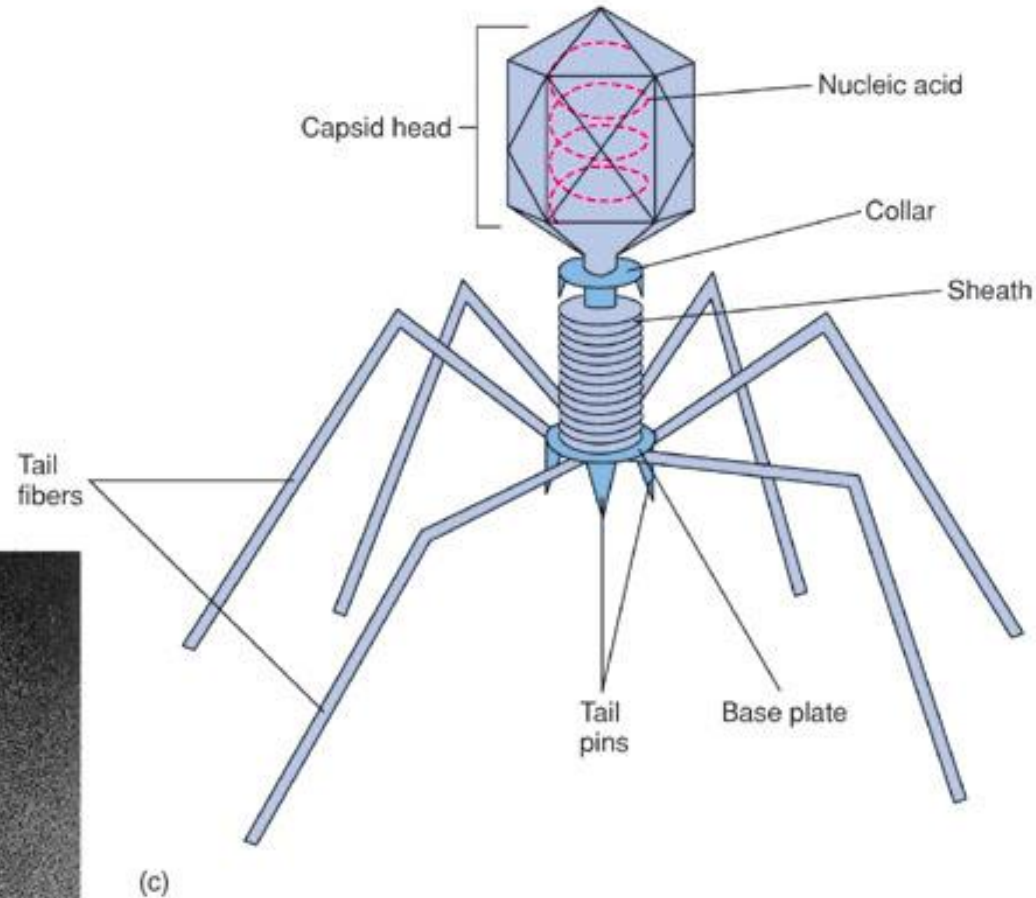
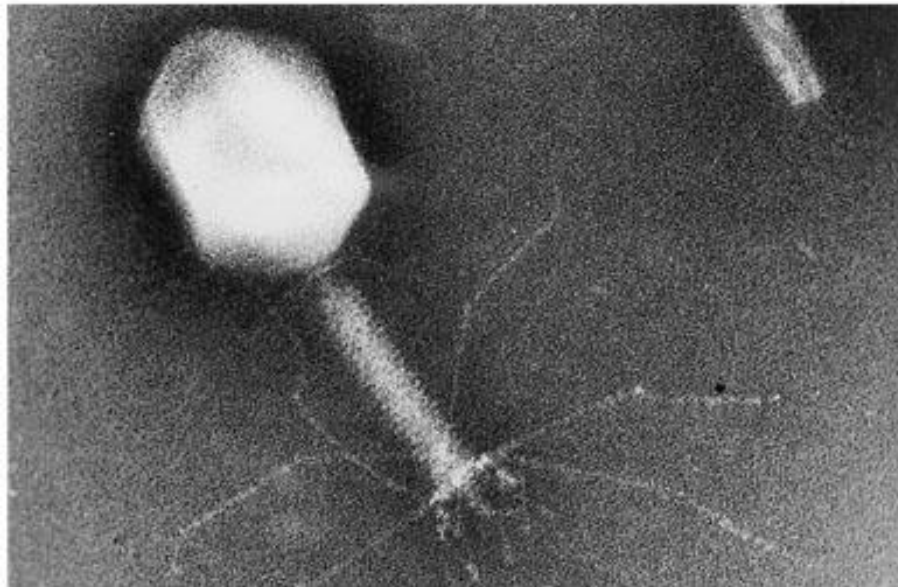
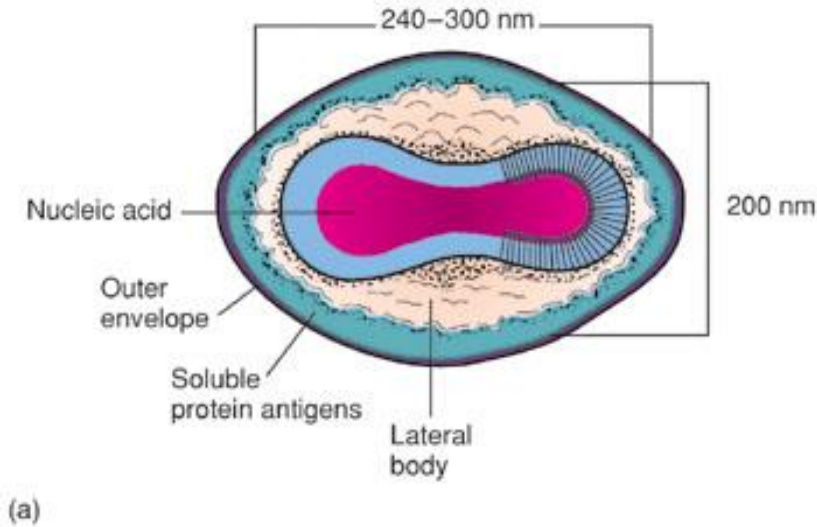


A bacteriophage

- A bacteriophage is any one of a number of viruses that infect bacteria. They do this by injecting genetic material, which they carry enclosed in an outer protein capsid. The genetic material can be ssRNA, dsRNA, ssDNA, or dsDNA ('ss-' or 'ds-' prefix denotes single-strand or double-strand) along with either circular or linear arrangement.

Phage - viruses have a polyhedral head, helical tail and fibers for attachment.

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Classification of viruses

- Nucleic acid
- Capsid
- Presence of envelope
- Replication strategy

CLASSIFICATION NUCLEIC ACID

- RNA or DNA
- segmented or non-segmented
- linear or circular
- single-stranded or double-stranded
- if single-stranded RNA
 - is genome mRNA (+) sense or complementary to mRNA (-) sense

ENVELOPE

- OBTAINED BY BUDDING THROUGH A CELLULAR MEMBRANE (except poxviruses)
- POSSIBILITY OF EXITING CELL WITHOUT KILLING IT
- CONTAINS AT LEAST ONE VIRALLY CODED PROTEIN
 - ATTACHMENT PROTEIN
- **LOSS OF ENVELOPE RESULTS IN LOSS OF INFECTIVITY**

Properties of naked viruses

- Stable in hostile environment
- Not damaged by drying, acid, detergent, and heat
- Released by lysis of host cells
- Can sustain in dry environment
- Can infect the GI tract and survive the acid and bile
- Can spread easily via hands, dust, fomites, etc
- Can stay dry and still retain infectivity
- Neutralizing mucosal and systemic antibodies are needed to control the establishment of infection

Naked viruses(Non Enveloped)

- Adeno-associated Virus (AAV)
Adenovirus
B19
Coxsackievirus - A
Coxsackievirus - B
Echovirus
Hepatitis A Virus (HAV)
Hepatitis E Virus (HEV)
Norwalk Virus

The Baltimore classification system

Based on genetic contents and replication strategies of viruses. According to the Baltimore classification, viruses are divided into the following seven classes:

1. dsDNA viruses
2. ssDNA viruses
3. dsRNA viruses
4. (+) sense ssRNA viruses (codes directly for protein)
5. (-) sense ssRNA viruses
6. RNA reverse transcribing viruses
7. DNA reverse transcribing viruses

where "ds" represents "double strand" and "ss" denotes "single strand".

Virus Classification

- the Baltimore classification

- All viruses must produce mRNA, or (+) sense RNA
- A complementary strand of nucleic acid is (–) sense
- The Baltimore classification has + RNA as its central point
- Its principles are fundamental to an understanding of virus classification and genome replication, but it is rarely used as a classification system in its own right

Viral genome strategies

- dsDNA (herpes, papova, adeno, pox)
- ssDNA (parvo)
- dsRNA (reo, rota)
- ssRNA (+) (picorna, toga, flavi, corona)
- ssRNA (-) (rhabdo, paramyxo, orthomyxo, bunya, filo)
- ssRNA (+/-) (arena, bunya)
- ssRNA (+RTase) (retro, lenti)

Sub-viral agents

- **Satellites**

- Contain nucleic acid
- Depend on co-infection with a **helper virus**
- May be encapsidated (satellite virus)
- Mostly in plants, can be human e.g. hepatitis delta virus
- If nucleic acid only = virusoid

- **Viroids**

- Unencapsidated, small circular ssRNA molecules that replicate autonomously
- Only in plants, e.g. potato spindle tuber viroid
- Depend on host cell polII for replication, no protein or mRNA

- **Prions**

- No nucleic acid
- Infectious protein e.g. BSE

Viroids & Prions

- **Viroids**

- ss RNA genome and the smallest known pathogens.
- Affects plants

- **Prions**

- Infectious particles that are entirely protein.
- No nucleic acid
- Highly heat resistant
- Animal disease that affects nervous tissue
- Affects nervous tissue and results in
 - Bovine spongiform encephalitis (BSE) “mad cow disease”,
 - scrapie in sheep
 - kuru & Creutzfeld-Jakob Disease (CJD) in humans

Viroids

- Viroids are small (200-400nt), circular RNA molecules with a rod-like secondary structure which possess no capsid or envelope which are associated with certain plant diseases. Their replication strategy like that of viruses - they are obligate intracellular parasites.
- Viroids do not encode any proteins and unlike satellites they are not dependent on the presence of another virus

Viroid replication

- Viroids utilize cellular RNA polymerases for their replication
- Replication is performed by “rolling circle mechanism”
- The resulting long RNA molecule is cut in pieces and ligated either autocatalytically or by cellular factors (depending on a viroid)
- So in a sense, at least some viroids are ribozymes...



Examples of plants, infected with various viroids

Hepatitis δ virus – a chimeric molecule, half viroid, half satellite

- Viroid like properties
 - Rod-like RNA molecule
 - Rolling circle replication
 - Self-cleaving activity
- Satellite like properties
 - Encodes a protein, which is necessary both for encapsidation and replication
 - Dependent on presence another virus – HBV
 - Genome larger than for viroids (1640 nt)

Prions

- Prions are rather ill-defined infectious agents believed to consist of a single type of protein molecule with no nucleic acid component. Confusion arises from the fact that the prion protein & the gene which encodes it are also found in normal 'uninfected' cells. These agents are associated with diseases such as Creutzfeldt-Jakob disease in humans, scrapie in sheep & bovine spongiform encephalopathy (BSE) in cattle.

Prions

Prions are proteinaceous transmissible pathogens responsible for a series of fatal neurodegenerative diseases (in humans, Creutzfeld-Jakob disease and kuru, in animals, bovine spongiform encephalopathy)

A prion (**proteinaceous infectious** particle, analogy for virion) is a type of infectious agent that does not carry the genetic information in nucleic acid!

Prions are proteins with the pathological conformation that are believed to infect and propagate the conformational changes of the native proteins into the abnormally structured form

Disease name	Natural host	Prion name	PrP isoform
Scrapie	Sheep, goat	Scrapie prion	OvPrP ^{Sc}
Transmissible mink encephalopathy (TME)	Mink	TME prion	MkPrP ^{Sc}
Chronic wasting disease (CWD)	Elk, mule deer	CWD prion	MDePrP ^{Sc}
Bovine spongiform encephalopathy (BSE)	Cattle	BSE prion	BovPrP ^{Sc}
Feline spongiform encephalopathy (FSE)	Cat	FSE prion	FePrP ^{Sc}
Exotic ungulate encephalopathy (EUE)	Greater kudu, nyala	EUE prion	NyaPrP ^{Sc}
Kuru	Human	Kuru prion	HuPrP ^{Sc}
Creutzfeldt-Jakob disease (CJD)	Human	CJD prion	HuPrP ^{Sc}
Gerstmann-Straussler-Scheinker syndrome (GSS)	Human	GSS prion	HuPrP ^{Sc}
Fatal familial insomnia (FFI)	Human	FFI prion	HuPrP ^{Sc}

Prion diseases: rare neurodegenerative disorders (one person per million)

1. Sporadic (85 %)

In the sixth or seventh decade, rapidly progressive (death in less than a year)

Creutzfeldt-Jakob disease (CJD)

2. Familial (inherited-15%)

Mutations in the PrP gene that favour the transition from the cellular form to the pathological form of PrP

Gerstmann-Straussler-Scheinker disease (GSS), fatal familial insomnia (FFI)

3. Transmissible (rare; a source of great concern)

Propagation of kuru disease in New Guinea natives (ritualistic cannibalism)

Recently, it has been discovered that BSE had been transmitted to humans in Europe after consumption of infected beef, producing a variant of the CJD called vCJD

Transmissible spongiform encephalopathy (TSE)=prion disease

A group of progressive conditions that affect the brain and nervous system of humans and animals and are transmitted by prions

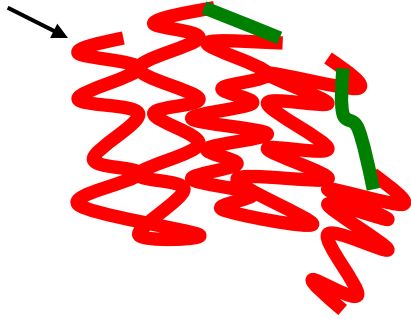
The pathology: vacuolar degeneration, neuronal loss, astrocytosis and amyloid plaque formation

The clinical signs: loss of motor functions (lack of coordination, ataxia, involuntary jerking movements), personality changes, depression, insomnia, confusion, memory problems, dementia, progressive tonic paralysis, death

Definitive diagnostic test: biopsy of brain tissue (histopathological examination and immunostaining for PrP^{Sc})

There is no cure

α -helix



Normal protein
(folded structure)

Conformational change



β -sheet



Disease-associated protein
(misfolded structure)

Aggregation

Gain of toxic
activity

Loss of biological
function

PrP^C

The normal protein
is called PrP^C (for cellular)

is a transmembrane glycoprotein
(neurons, lymphocytes); its function
is unknown; it binds Cu²⁺ (regulation
its homeostasis)

has dominant secondary structure α -
helix

is easily soluble

is monomeric and easily digested by
proteases

is encoded by a gene designated
PRNP located on the chromosome 20

PrP^{Sc}

The abnormal, disease-producing
protein

is called PrP^{Sc} (for scrapie)

has the same amino acid sequence
(primary structure)

has dominant secondary structure β -
sheets

is insoluble

is multimeric and resistant to
digestion by proteases

When PrP^{Sc} comes in contact with
PrP^C, it converts the PrP^C into more of
itself These molecules bind to each
other forming aggregates

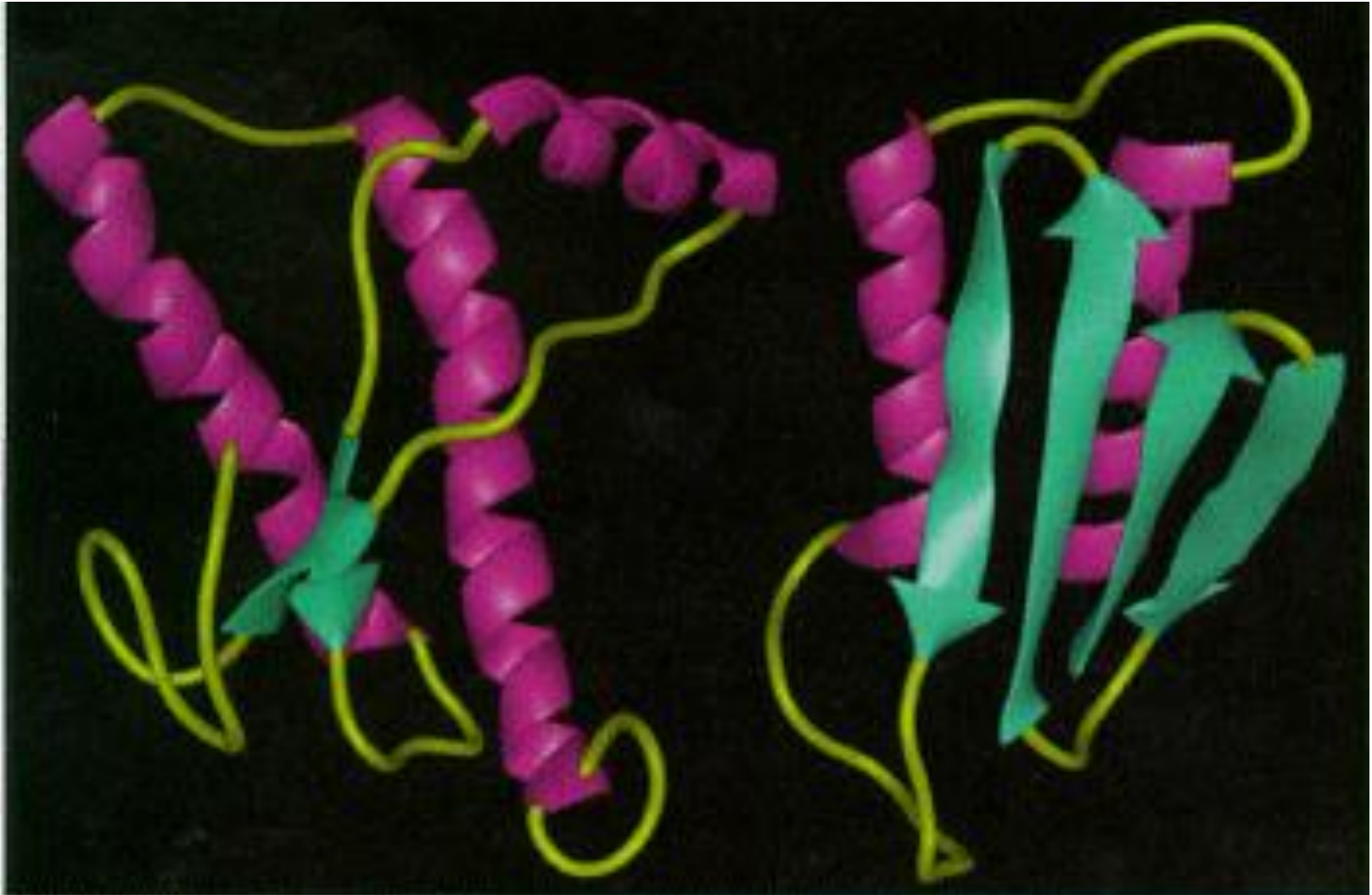
Molecular models of the structure of:

PrP^C

Predominantly α -helix (3)

PrP^{Sc}

β -sheets (40%), α -helix (30%)

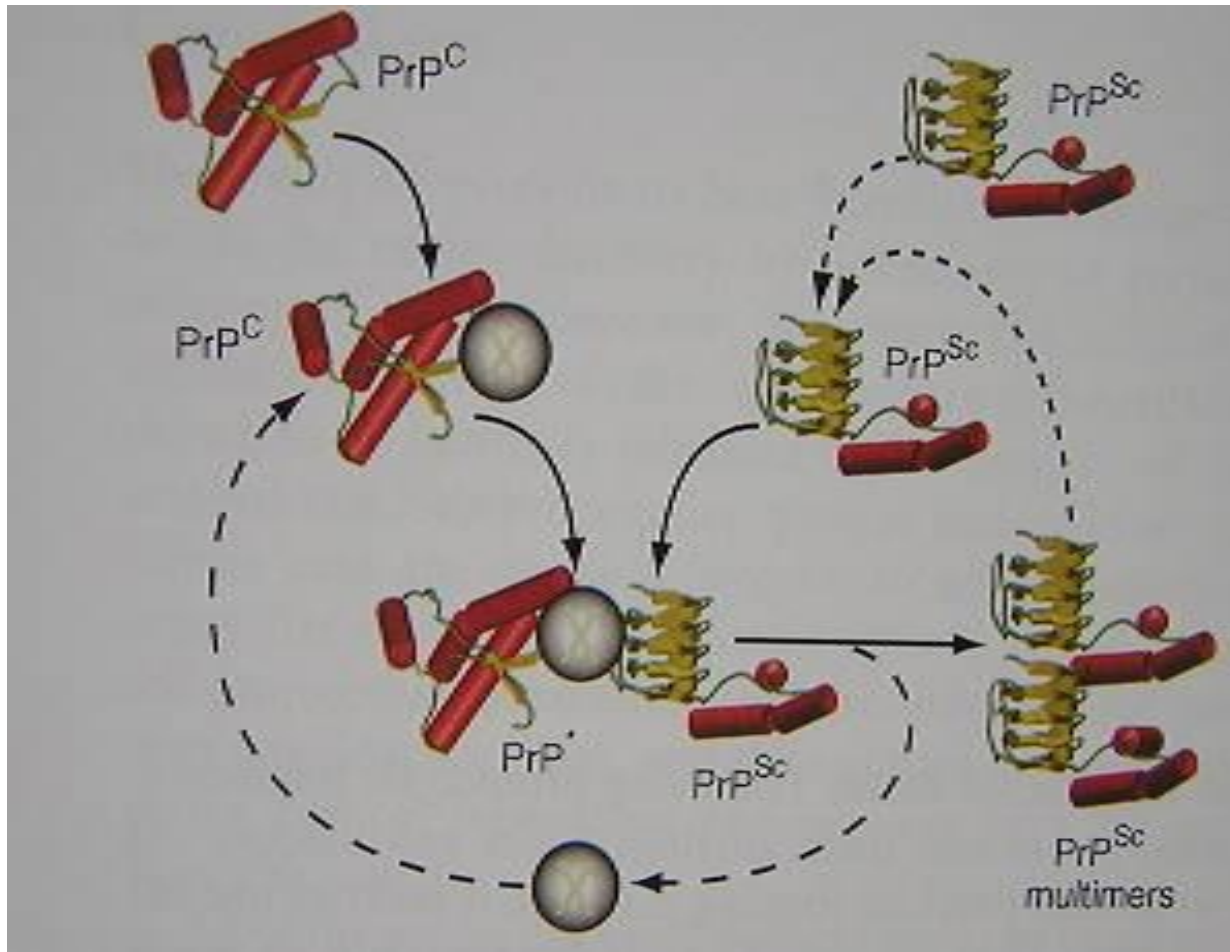


Replication cycle

The presence of an initial PrP^{Sc} : exogenous (infectious forms) or endogenous (inherited or sporadic forms)

This first prion will initiate PrP^{Sc} accumulation by sequentially converting PrP^{C} molecules into PrP^{Sc} in replication cycle

PrP^{Sc} molecules aggregate



Summary

The prions are proteins that carry information for self-reproduction (contradict the central dogma of modern biology)

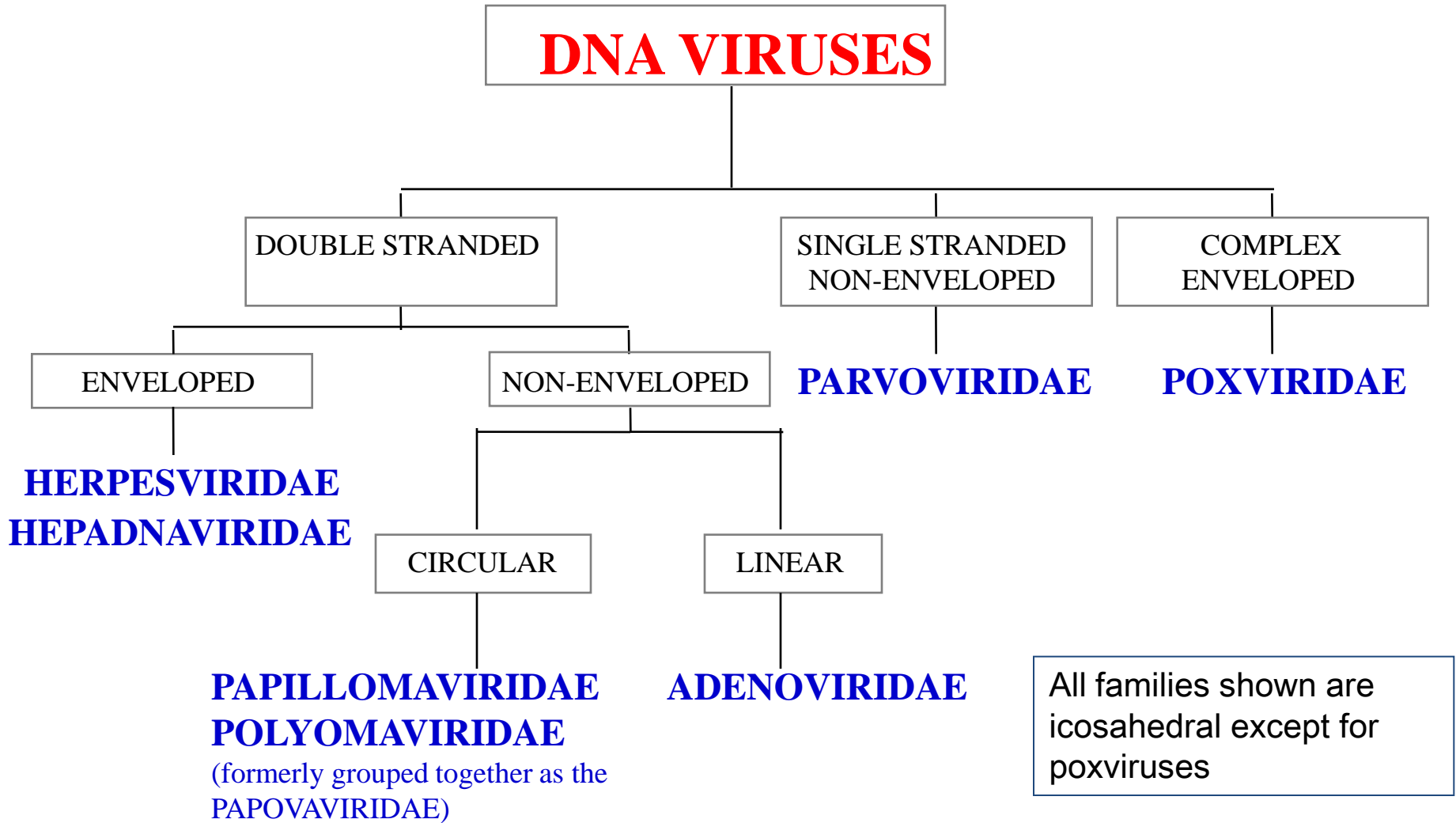
The prions are expressed in cells of healthy humans and animals; their abnormal conformations (PrP^{Sc}) are insoluble, resistant to digestion and aggregate

The PrP^{Sc} attacks the native prion PrP^{C} , changes its conformation into an abnormal form and causes an exponential production of insoluble proteins; they aggregate and form the fibrillar structure

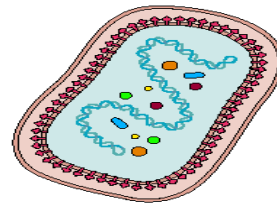
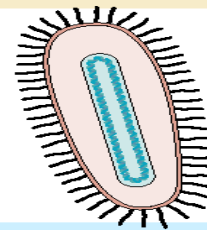
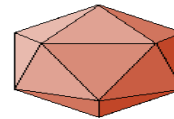
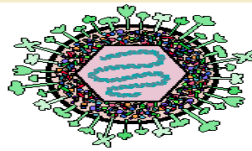
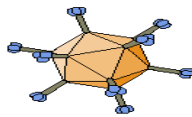
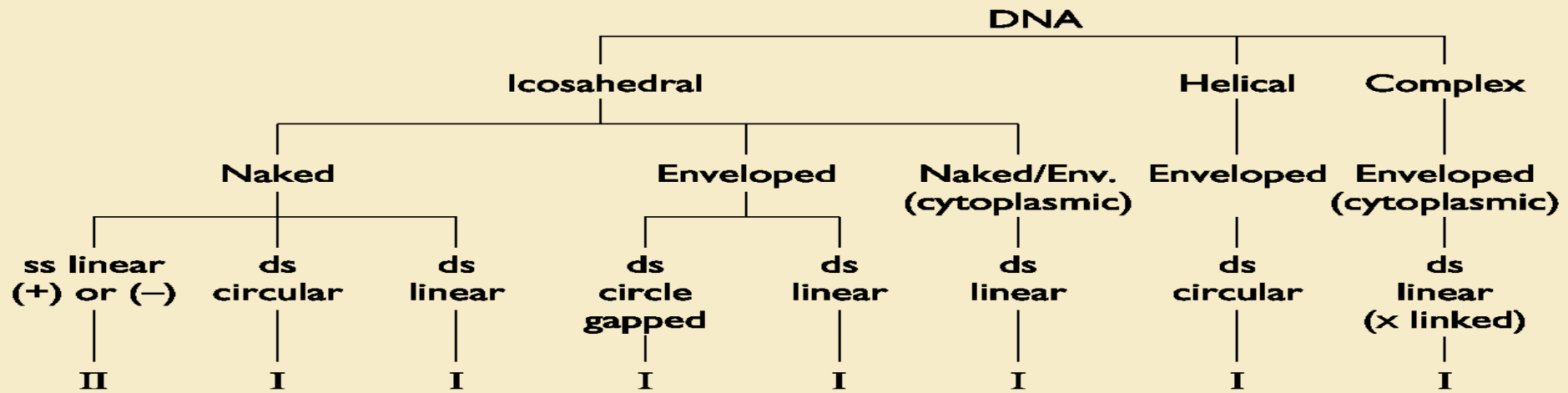
Prion disease are rare fatal degenerative disorders; a portion of them can be transmitted; this mechanism is not clear (e.g. transmission of BSE to human)

One part of the prion protein can cause apoptosis, or programmed cell death

Prions induce no immune reactions within the human



DNA viruses



Parvo

(-)

18-26

5

Papova

(-)

45-55

5-8

Adeno

(-)

70-90

36-38

Hepadna

(+)

42

3.2

Herpes

(-)

150-200

120-200

Irido

(-)

125-300

150-350

Baculo

(-)

60 X 300

100

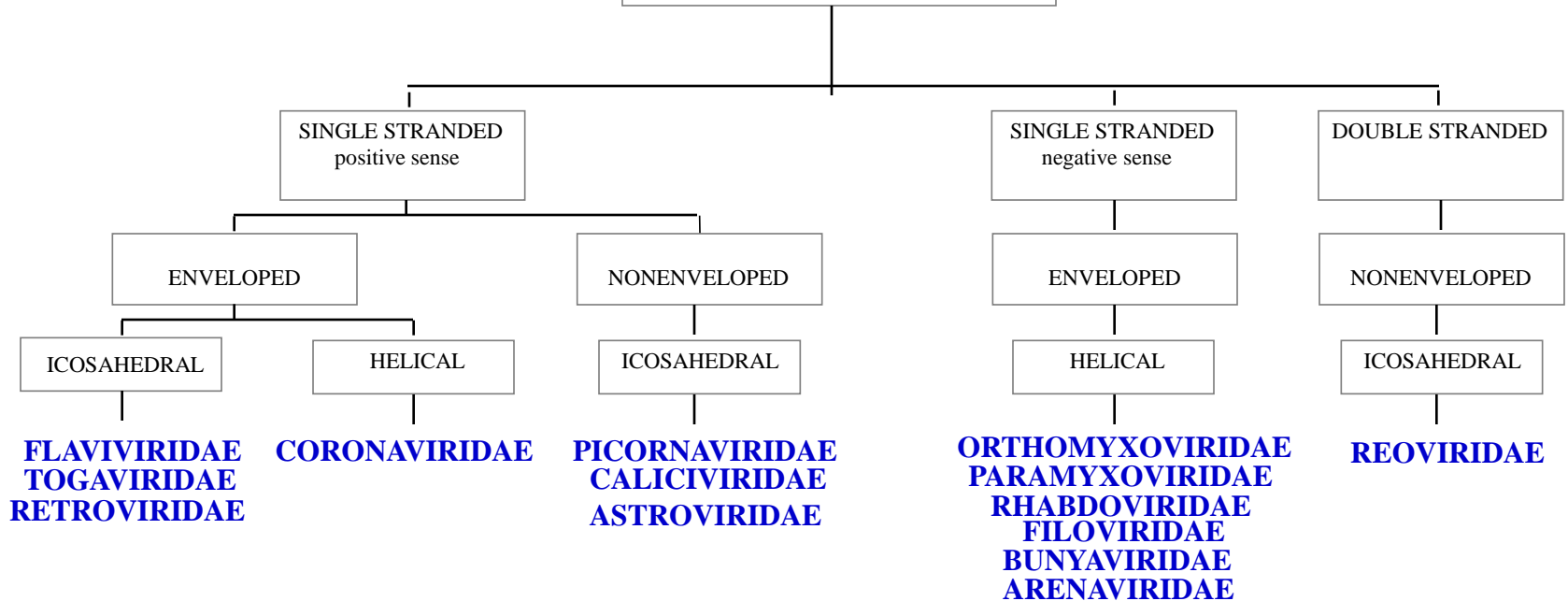
Pox

(+)

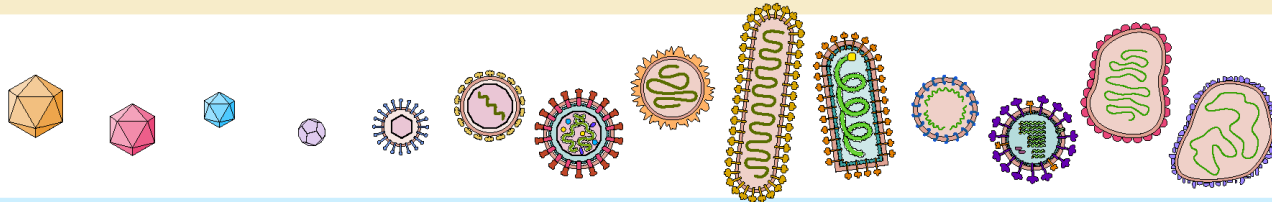
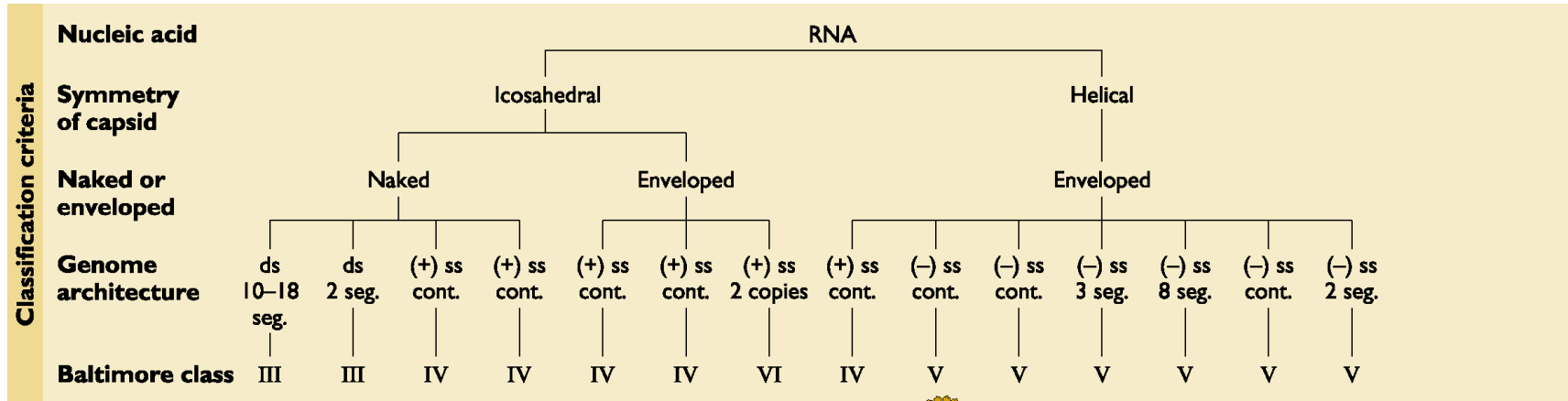
170-200
X 300-450

130-280

RNA VIRUSES



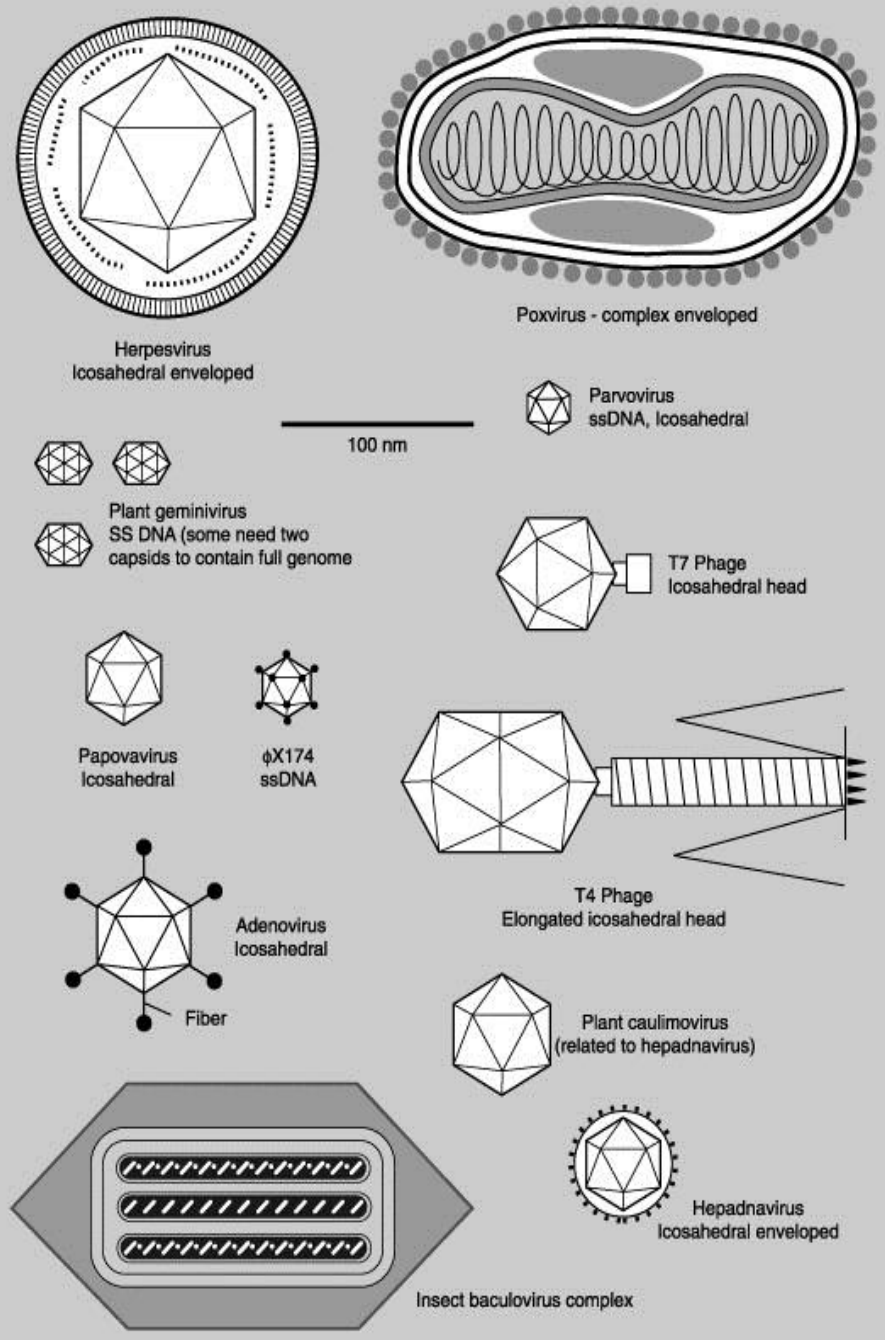
RNA viruses



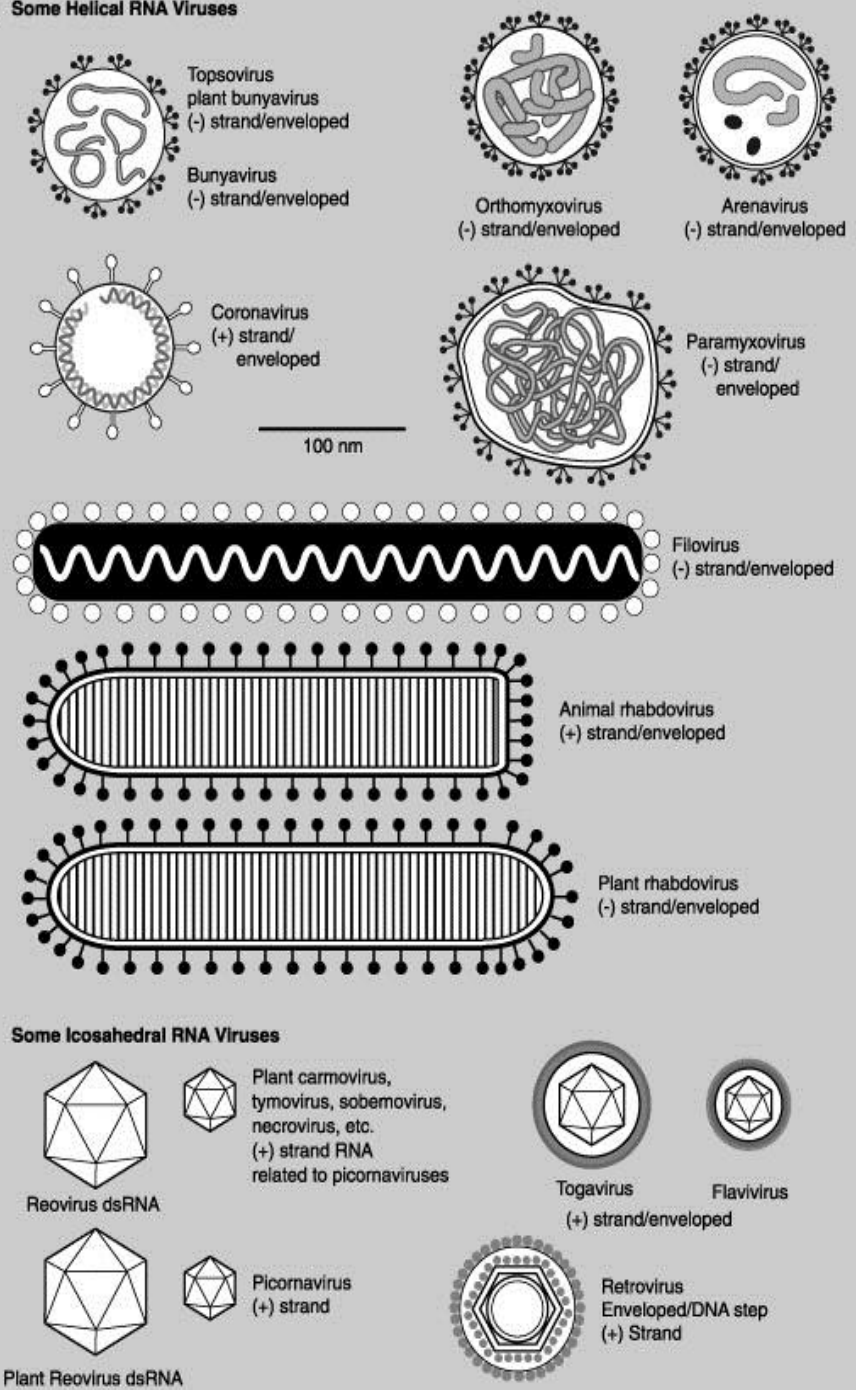
Properties	Family name	Reo	Birna	Calici	Picorna	Flavi	Toga	Retro	Corona	Filo	Rhabdo	Bunya	Orthomyxo	Paramyxo	Arena
	Virion polymerase	(+)	(+)	(-)	(-)	(-)	(-)	(+)	(-)	(+)	(+)	(+)	(+)	(+)	(+)
	Virion diameter (nm)	60–80	60	35–40	28–30	40–50	60–70	80–130	80–160	80 x 790–14,000	70– 85 x 130–380	90–120	90–120	150–300	50–300
	Genome size (total in kb)	22–27	7	8	7.2–8.4	10	12	3.5–9	16–21	12.7	13–16	13.5–21	13.6	16–20	10–14

From Principles of Virology Flint et al ASM Press

(a) Some DNA Viruses

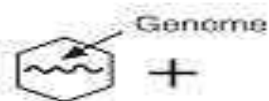


(b) Some Helical RNA Viruses



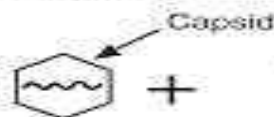
RNA Viruses

Picornavirus



C = 32
22-30 nm

Astrovirus



C = 32?
30-35 nm

Calicivirus



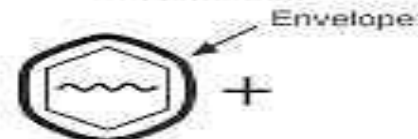
C = 32 (holes)
35-39 nm

Flavivirus



Icosahedral
45-50 nm

Togavirus



Icosahedral
70 nm

Coronavirus



Pleomorphic
120-160 nm

Retrovirus



Icosahedral
90-120 nm

Reovirus



C = 132
60-80 nm

Bunyavirus



90-120 nm

Orthomyxovirus



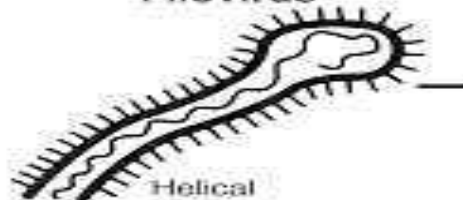
Helical, Pleomorphic
80-120 nm

Arenavirus



Pleomorphic
110-130 nm

Filovirus



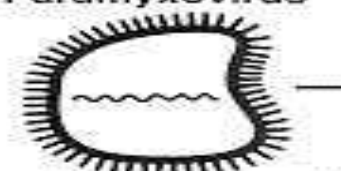
Helical
80x800-2500 nm

Rhabdovirus



Helical
60x180 nm

Paramyxovirus



Helical, Pleomorphic
150-300 nm

DNA Viruses

Circovirus



Icosahedral
17-22 nm

Parvovirus



C = 12
18-26 nm

Hepadnavirus



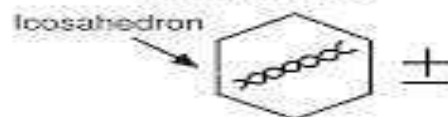
C = 180 Icosahedral
40-48 nm

Papovavirus



C = 72
45/55 nm

Adenovirus



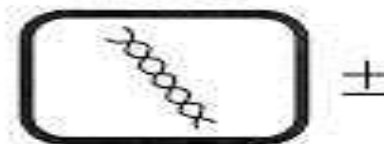
C = 252
75-80 nm

Herpesvirus



C = 162
150-200 nm

Poxvirus



Complex
240x300 nm

Dr. T.Y. Rao MD

BASIC STEPS IN VIRAL LIFE CYCLE

- **ADSORPTION**
- **PENETRATION**
- **UNCOATING AND ECLIPSE**
- **SYNTHESIS OF VIRAL NUCLEIC ACID AND PROTEIN**
- **ASSEMBLY**
- **RELEASE**