Picornaviridae

VMC 321

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ETYMOLOGY

- Picornaviridae
- 'Pico' very small
- · 'rna' RNA
- viridae suffix used for family of virus
- The *Picornaviridae* family includes small, icosahedral viruses with single-stranded, highly diverse positive-sense RNA genomes.

Taxonomy

| Group IV: (+) sense, ss RNA viruses | | | |
|-------------------------------------|----------------|------------------------------|----------------------|
| Order: Picornavirales | | | |
| Family | Genus | Species | Host |
| Picornaviridae | Aphthovirus | Foot-and-mouth disease virus | cloven footed animal |
| | Avihepatovirus | Duck hepatitis A virus | Duck |
| | Erbovirus | Equine rhinitis B virus | Equine |
| | Hepatovirus | Hepatitis A virus | Vertebrates |
| | Sapelovirus | Porcine sapelovirus | Pig |
| | Teschovirus | Porcine teschovirus | Pig |
| | | | |

Characteristics of members of the family *Picornaviridae*

| Characteristic | Description |
|----------------|---|
| Prototype | poliovirus 1, species Enterovirus C, genus Enterovirus |
| Virion | Non-enveloped, 30–32 nm virions comprising 60 protomers |
| Genome | 6.7–10.1 kb of positive-sense, non-segmented RNA with a poly(A) tail |
| Replication | RNA synthesis occurs in reorganized cytoplasmic replication organelles containing non-structural proteins derived from the 2BC-P3 region of the encoded polyprotein; RNA structures at the 5' and 3' ends of the genome direct initiation of RNA synthesis and uridylated 3B serves as primer for synthesis of both RNA strands |
| Translation | Directly from genomic RNA containing an internal ribosomal entry site (IRES) |
| Host range | Vertebrates (at least five of the seven classes) |

Virion Morphology

- Virions consist of a capsid, with no envelope, surrounding a core of ssRNA.
- Virus particles size: 30–32 nm in diameter
- Shape— Isometric; non-enveloped
- Symmetry Icosahedral
- Genome— linear; 7.2-8.4 kb; single segment; positive sense; single stranded RNA
- Molecular weight— 2.5 × 10⁶ k Da
- Virus has four major protein (VP₁, VP₂, VP₃, & VP₄) & one minor protein (VP₀)

Replication

- i. virion attaches to receptor, penetrates the host cell
- ii. uncoating occurs & VP_0 is removed from RNA segment by cellular enzyme
- iii. FMDV use entire RNA as mRNA, whole genome is translated & cleaves into different functional protein.

The Disease

Foot and Mouth Disease (FMD)

Susceptibile host:

- Cattle
- Buffalo
- Sheep
- Goat
- Deer
- Pig
- Camel
- Mithun
- Yak

Reservoir/Carrier:

- Wild ruminants
- Cattle- indicator host
- Sheep- maintenance host
- Pig- amplifying host

Transmission:

- spread at rapid rate through:
 - contact with infected animal
 - contaminated feed, Utensil
 - aerosol
 - ingestion

Pathogenicity:

- RNA itself is infectious
- Virus invades epithelium of upper respiratory tract or alimentary tract by via inhalation or ingestion and multiplies at the site of infection
- Formation of vesicle within 1-4 days
- Primary vesicle initiates viraemia and associated with fever
- Dissemination through viraemia and lodgment in distant epithelia & formation of secondary vesicle
- Stratified squamous epithelium are the predilection site of FMDV and vesicles are found in these tissues or mucous membrane.

Clinical sign:

- Appearance of vesicles on mucous membrane of tongue, lips, gums, cheeks, dental pad & skin of interdigital space, teats and udder
- Profuse salivation, drooling of foamy& ropy saliva and protrusion of tongue
- Pregnant animals abort
- Suckling calves die as a result of myocardial degeneration (tigeroid heart)

Diagnosis:

- 1. Clinical signs and symptoms
- 2. Isolation and identification of virus
 - Specimen for virus isolation: vesicular fluid, epithelial tissues from ruptured vesicle & transported in glycerol buffer
- 3. Demonstration of viral antigen in clinical specimen
 - Virus neutralization test (VNT)
 - Immunofluorescence test(IFT)
 - Complement fixation test (CFT)and
 - Enzyme linked immunosorbent assay(ELISA)

- 4. Demonstration of antibody in convalescent sera
 - > Liquid phase block Enzyme linked immunosorbent assay(LPB-ELISA)
 - Complement fixation test (CFT)
- 5. Detection of viral nucleic acid
 - In situ hybridization(with labeled gene probes)
 - Reverse trancriptase –polymerase chain reaction(RT-PCR)

Precaution:

| Vaccination | Age |
|---------------|------------------|
| Initial | 4 months & above |
| Booster | 1months of first |
| | vaccination |
| Revaccination | Every six months |

- 1. Vaccination
 - Trivalent FMD vaccine containing O, A, , Asia-1

 Dose- 2ml (Cattle & buffalo) ; 1 ml (Sheep & Goat)
- 2. Disease awareness, early detection, proper disposal of affected carcass, timely notification,
- 3. Strict biosecurity, isolation, zoning, quarantine, control of animal movement are prerequisites for effective control programmes aimed at eradication

Duck hepatitis virus (DHV)

- Three types of viruses are recognized:
- Type 1
- Type 2
- Type 3
- All the above three types causes hepatitis in duck, however, type1 is prevalent in India

The Disease

- DHV causes duck virus depatitis
- **Duck virus hepatitis**: Duck hepatitis is a highly fatal, rapidly spreading viral infection of young ducklings characterized primarily by hepatitis.
- Susceptible host: Ducklings
- **Reservoir**: Wild birds, recovered ducks and rats
- Transmission: Contact with infected bird
- Incubation period: 18-24hrs.

Pathogenicity:

- I. Duckling of 3-4 weeks old are infected; disease takes rapid cours
- II. Virus invades epithelium of upper respiratory tract by inhalation multiplies at the site of infection
- Mortality is as high as 100%; morbidity may be up to 95%

Clinical sign:

- I. The duckling reluctance to move & squat down
- II. Birds fall down on their sides; starts kicking their legs spasmodically and death ensues with head bent backward
- III. Death occurs within an hour

Diagnosis:

- 1. Clinical signs and symptoms
- 2. Post mortem lesions :-
 - ✓ Hepatomegaly with punctuate & ecchymotic haemorrhages;
 - ✓ Mottling & discolouration of liver;
 - ✓ Spleenomegaly & its mottling;
 - ✓ In few cases swelling of kidney & congestion of blood vessel is observed.

• 2. Isolation and identification of virus

- Specimen for virus isolation: Liver of infected bird
- Suspected sample is inoculated into SPF/antibody free embryonated duck egg (10-14day old) or chicken egg (9-11 day old) via allantoic sac; infected duck embryo dies within 24-72 hrs.; Chicken embryo dies within 5-8 days. The gross lesion of embryo shows stunting & sub cutaneous haemorrhages of body, oedema of abdomen & hind limb.Swollen, reddened & yellowish discoloration of liver
- Primary cell culture of duck embryo liver cells.
 - CPE includes rounding & necrosis of cells,

• 3. Demonstration of viral antigen in clinical specimen

- Virus neutralization test (VNT)
- > Immunofluorescence test(IFT) rapid & accurate diagnostic method
- Enzyme linked immunosorbent assay(ELISA)

4. Demonstration of antibody in convalescent sera

- Virus neutralization test (VNT)
- Enzyme linked immunosorbent assay(ELISA)

Thanks