

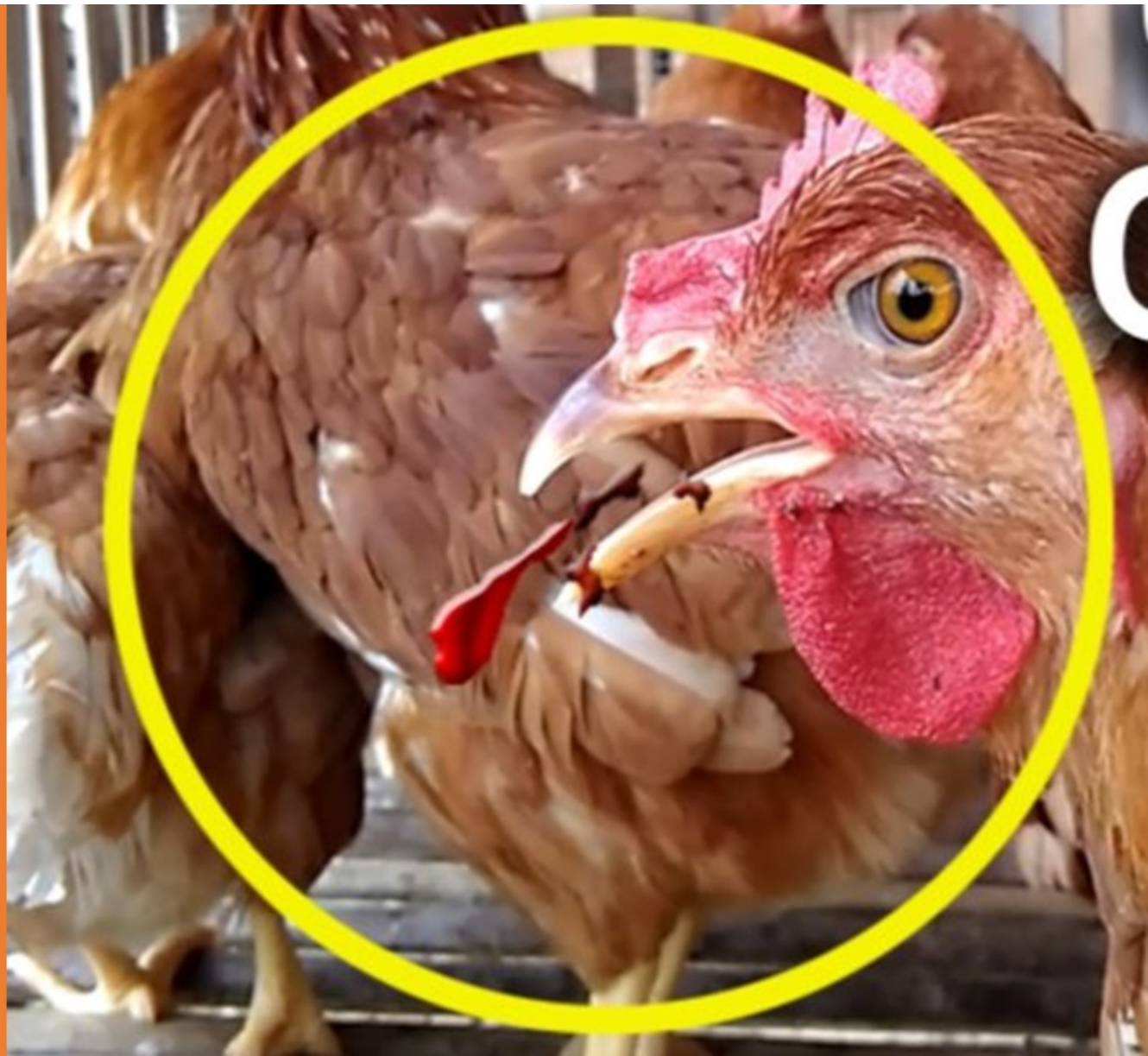
VMC 321: Systematic Veterinary Virology

Online lecture

Topic:
“Infectious laryngotracheitis virus”

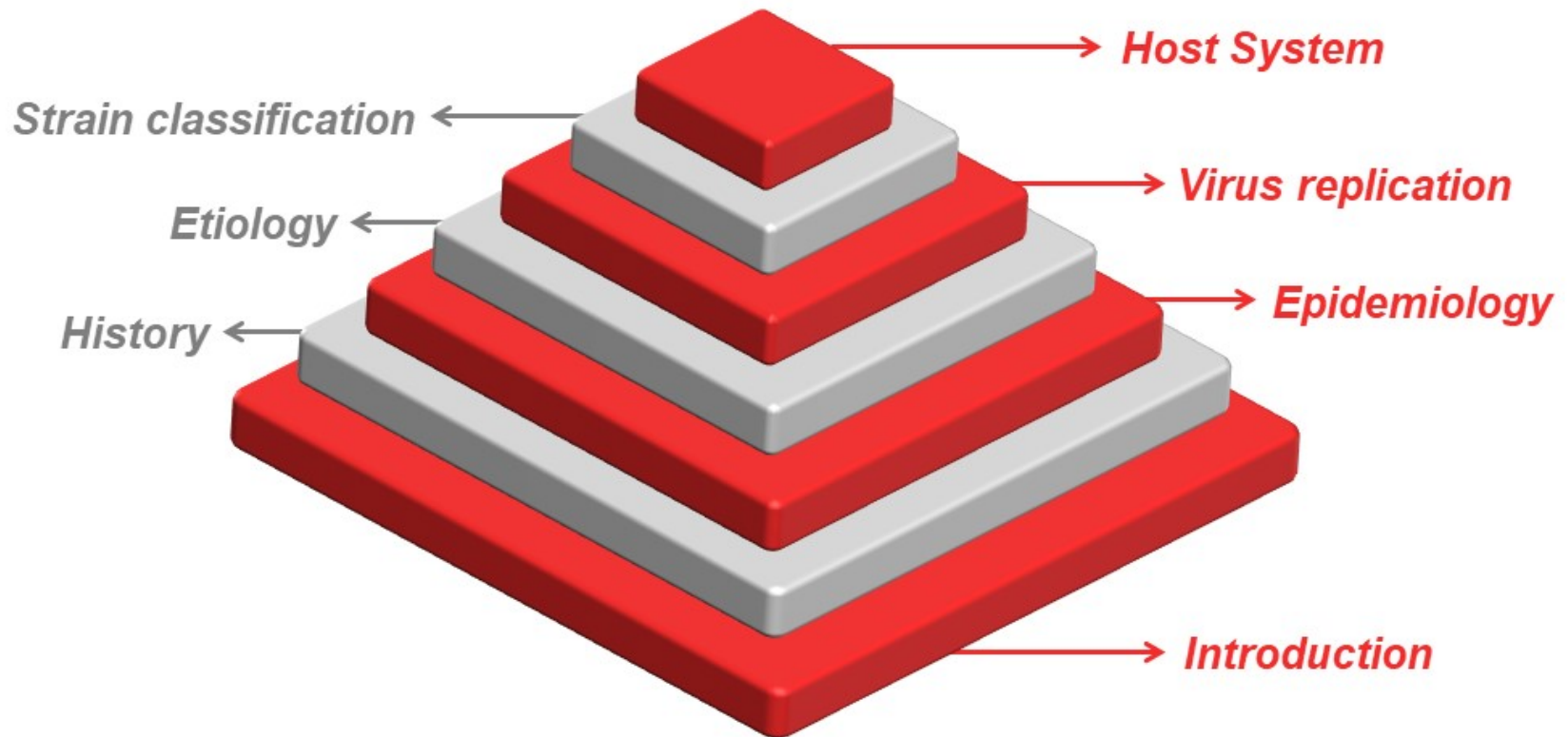
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Infectious laryngotracheitis virus

Contents





Introduction

- Infectious laryngotracheitis (ILT) is an acute, highly contagious infection of chickens and pheasants
 - Result in severe production losses due to mortality and/or decreased egg production
 - Severe epizootic forms of infection are characterized by signs of respiratory depression, gasping, expectoration of bloody mucus and high mortality
 - Mild enzootic forms of infection are encountered increasingly in developed poultry industries and manifest variously as mucoid tracheitis, sinusitis, conjunctivitis, general unthriftiness and low mortality
 - Included within List B of the **World Organization for Animal Health (OIE)**
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HISTORY

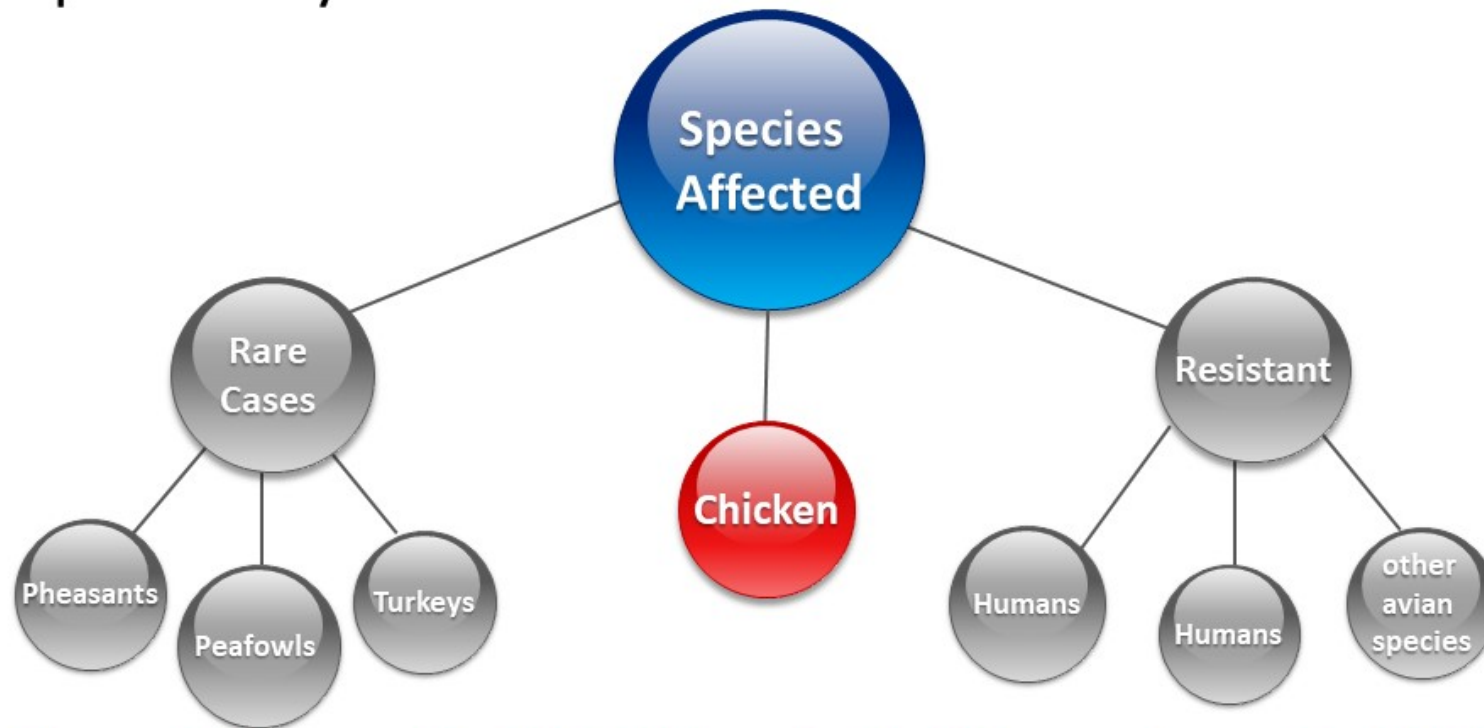
- The disease was first described in 1925 in Canada
- Followed by United States in 1926
- Given several different names including laryngotracheitis, infectious laryngotracheitis, and avian diphtheria
- The name infectious laryngotracheitis was adopted in 1931 by the **Special Committee on Poultry Diseases of the American Veterinary Medical Association**
- The cause of LT was first shown to be a filterable virus by Beaudette
- Laryngotracheitis was the first major avian viral disease for which an effective vaccine was developed



EPIDEMIOLOGY

- Distributed world-wide
 - May be present only in certain localities within a country or geographic region
 - The greatest incidence of disease is generally seen in areas of highly intensive
 - poultry production
 - Route of entry : upper respiratory and ocular routes
 - Transmission occurs : acutely infected bird , mechanical transmission (contaminated equipment and litter)
 - **No evidence** for vertical ILTV transmission to the egg or for shedding ILTV on shells of eggs laid by infected hens
 - Incubation period :
 - **6-12 days**, following natural exposure
 - **2-4 days**, following experimental inoculation
-

Susceptibility



- All ages of chickens are affected, but chickens older than 3 wk are most susceptible to ILTV
- ILTV can infect pheasants, pheasant-bantam crosses, and peafowl
- ILT can infect turkeys at about 100 d of age
- Humans are resistant

ETIOLOGY

Gallid herpesvirus 1

Genus *Iltovirus* ; Subfamily *Alphaherpesvirinae* ; Family *Herpesviridae*

Symmetry Icosahedral

Complete virus particle has a diameter of 195–250 nm and consists of an irregular envelope surrounding the nucleocapsid

DNA genome consists of a linear 155-kb ds molecule composed of **unique long (UL)** and **unique short (US)** regions flanked by **inverted repeats**

Nucleic acid of LTV is composed of DNA with a buoyant density of **1.704 g/mL**

HOST SYSTEM

I. Chicken Embryo

Embryonating chicken eggs are the most common method for propagating ILTVs

In chicken embryos, ILTV forms plaques on the chorioallantoic membrane (CAM).

The plaques can be observed 48 h after infection, and embryos can die in 2-12 d post infection (PI)

HOST SYSTEM

II. Cell Culture

ILTV can be isolated in primary cell cultures, such as chicken embryo liver (CEL), chicken embryo kidney (CEK), and chicken kidney (CK) cell cultures.

The sensitivity of ILTV isolation and propagation from field samples vary depending on the type of cell cultures. CEL was the most sensitive for isolation, followed by CK. The CEK and chicken embryo lung cells were less sensitive.

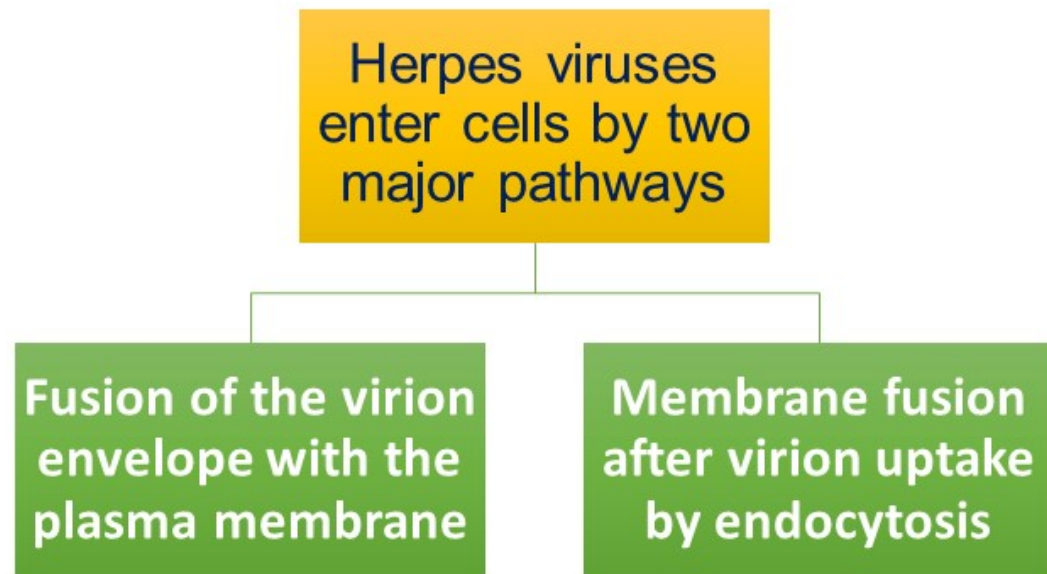
Chicken embryo fibroblasts can also be used

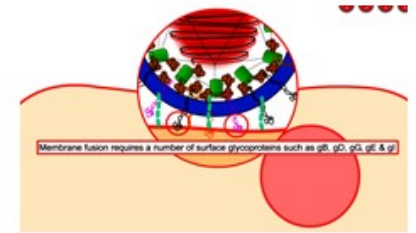
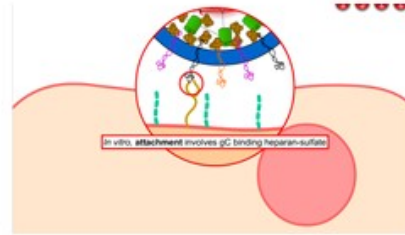
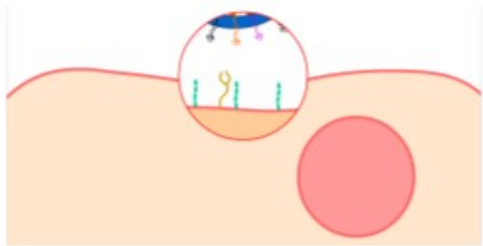
CPE consist of round refractile single cells or small syncytia in focal arrangement.

Altered cells were subsequently detached from the flask surface forming plaques and lyse

VIRAL REPLICATION

- Initiation of infection begins with receptor binding through the initial interactions of the virus genome with the host transcriptional machinery in the nucleus



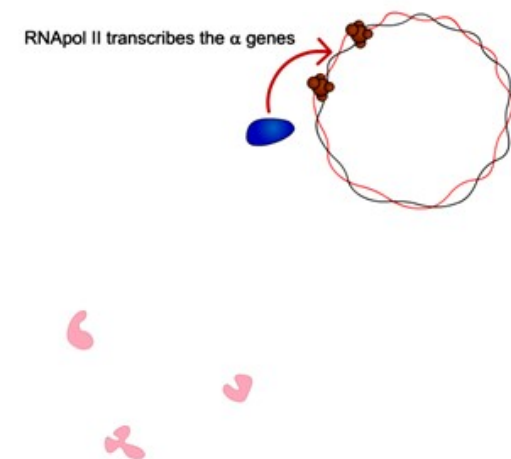
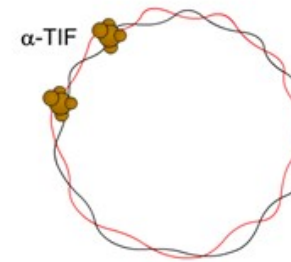


Fusion of virion and cellular membranes is mediated by gB and the gH /gL complex

Attachment of ILTV with host receptor

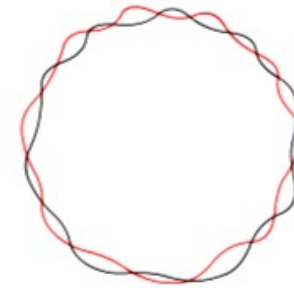
Transcription & replication

- Three classes of mRNA- α , β and γ are transcribed in sequence by **cellular RNA polymerase II**
- **α RNA (immediate early)** - processed appropriately to become mRNAs, are translated to form α proteins
- **β RNA (early RNA)** - the translation of which produces β (early) proteins
- **γ RNA (late) mRNAs**, which are transcribed from sequences situated throughout the genome are translated into γ proteins



VIRAL REPLICATION

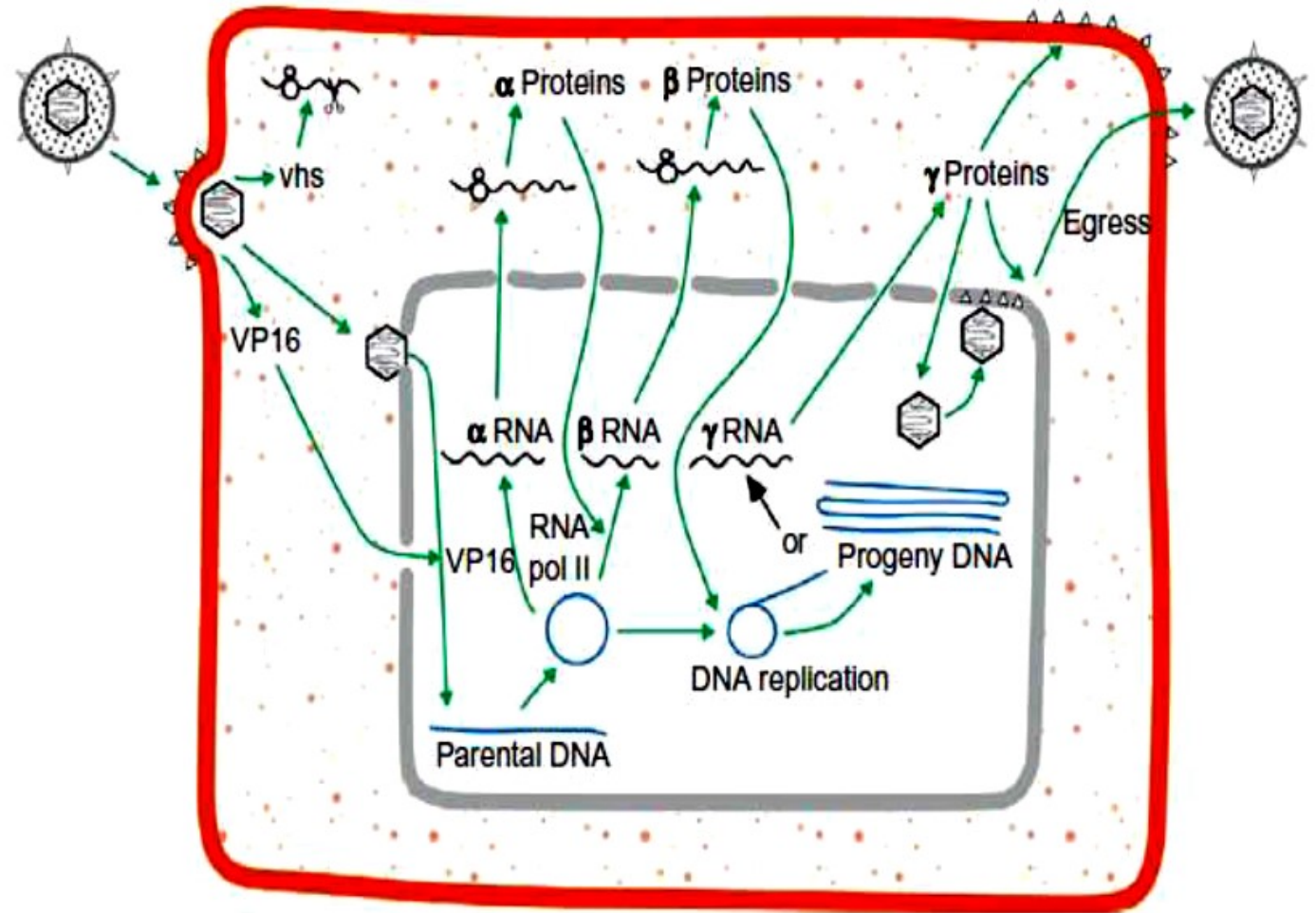
- Viral DNA replication then commences, utilizing some of the viral α and β proteins, in addition to host-cell proteins.



Using many viral enzymes such as helicases, primases and polymerases, the genome undergoes rolling circle type replication



Replication cycle



VIRAL REPLICATION

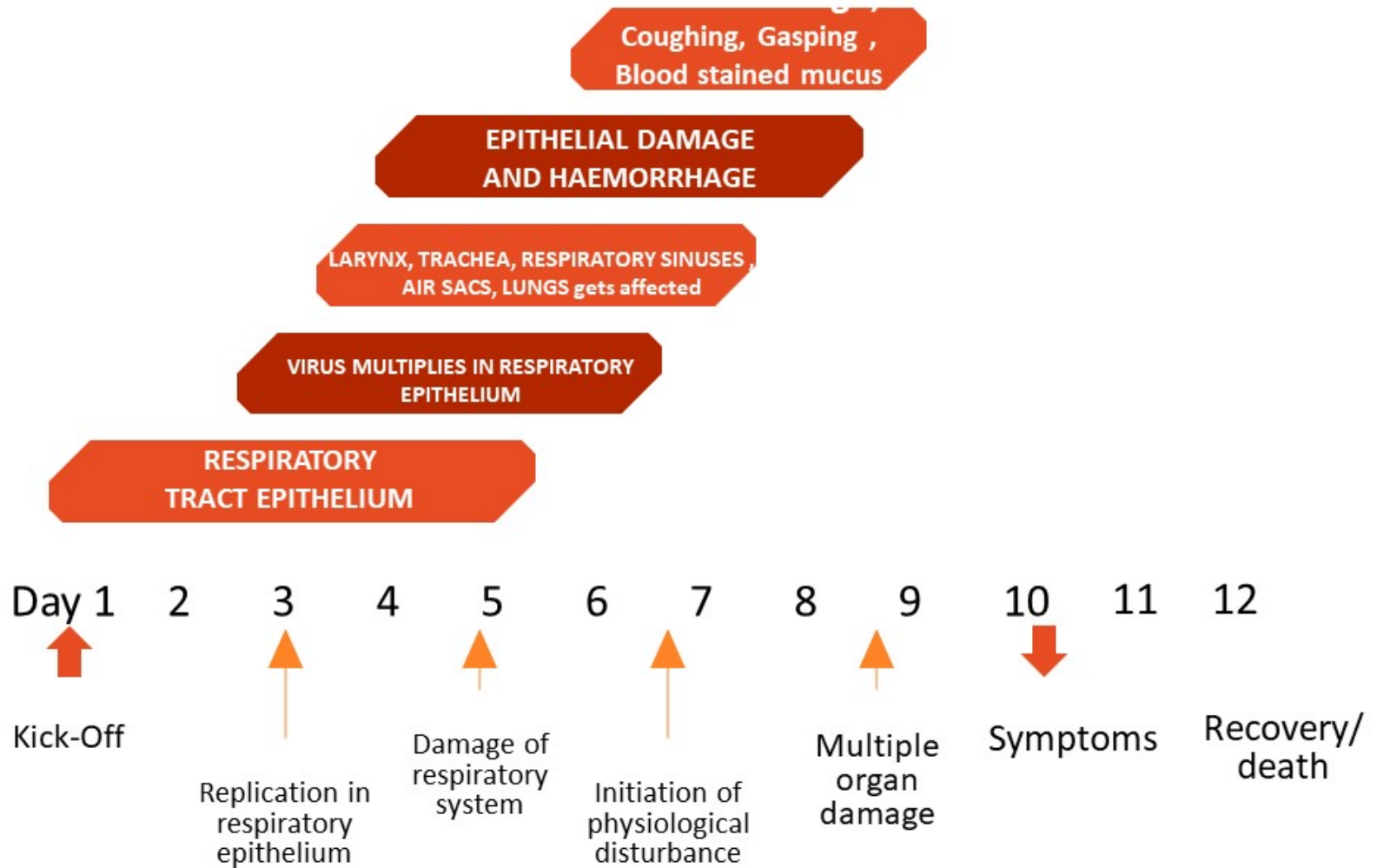
Maturation involves the completion of encapsidation of virion DNA into nucleocapsids and the association of nucleocapsids with altered patches of the inner layer of the nuclear envelope

Complete envelopment occurs by budding through the nuclear membrane

Mature virions accumulate within vacuoles in the cytoplasm and are released by exocytosis or cytolysis

Virus-specific proteins found in the plasma membrane, where they are involved in cell fusion

Pathogenesis Timeline



PATHOGENESIS cont...

Infectious virus usually is present in tracheal tissues and tracheal secretions for **6–8 days PI**

Extratracheal spread of LTV to trigeminal ganglia after 4-7 days of tracheal exposure

Trigeminal ganglion is the principal site of LTV latency

Reactivation of latent LTV from the trigeminal ganglia 15 months after vaccination of a flock has been reported from Germany

IMPORTANCE OF LATENT INFECTION

Rates of shedding of ILTV into the trachea increase by the stresses of either the onset of lay or mixing with unfamiliar birds

In this case, the latently infected chicken can act as an unsuspected reservoir host and enable ILTV to infect further susceptible chickens

CLINICAL SIGNS

Sub-acute form

- Nasal and ocular discharge
- Tracheitis
- Conjunctivitis
- Mild rales

Acute form (Severe form)

- Nasal discharge
- Moist rales
- Gasping (**Pump handle respiration**)
- Signs of respiratory distress (Dysnoea) including gasping and expectoration of bloody mucus
- High morbidity and moderate-to-high mortality
- Expectoration of blood stained mucus

Mild form

- Decreased egg production
- Watery eyes, conjunctivitis, swelling of infra-orbital sinuses,
- Mild tracheitis, persistent nasal discharge and hemorrhagic conjunctivitis mucoid tracheitis, sinusitis, conjunctivitis, general unthriftiness, and low mortality

CLINICAL SIGNS

Course of the infection varies with the severity of lesions

Generally, most chickens recover in 10–14 days

Epizootic forms of the disease cause high morbidity (90–100%) and variable mortality

Mortality can vary from 5% to 70% but usually is in the range of 10–20%

Mild enzootic forms of the disease result in morbidity as low as 5% and very low mortality (0.1–2%)

CLINICAL SIGNS

-RECOVERY

Infected birds that recover often become latent carriers and may intermittently shed the virus into the environment and infect other birds.

These recovered carrier birds are often the source of infection after introduction to naïve birds

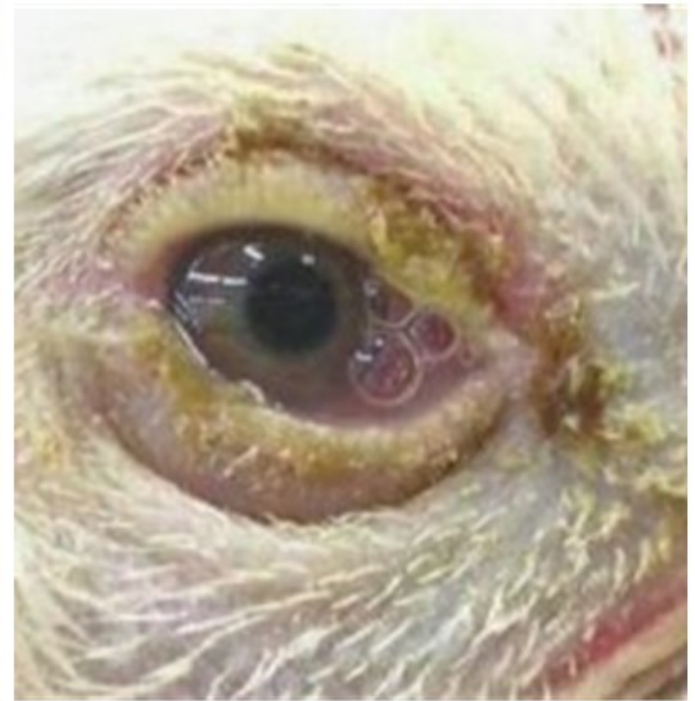


Open beak breathing

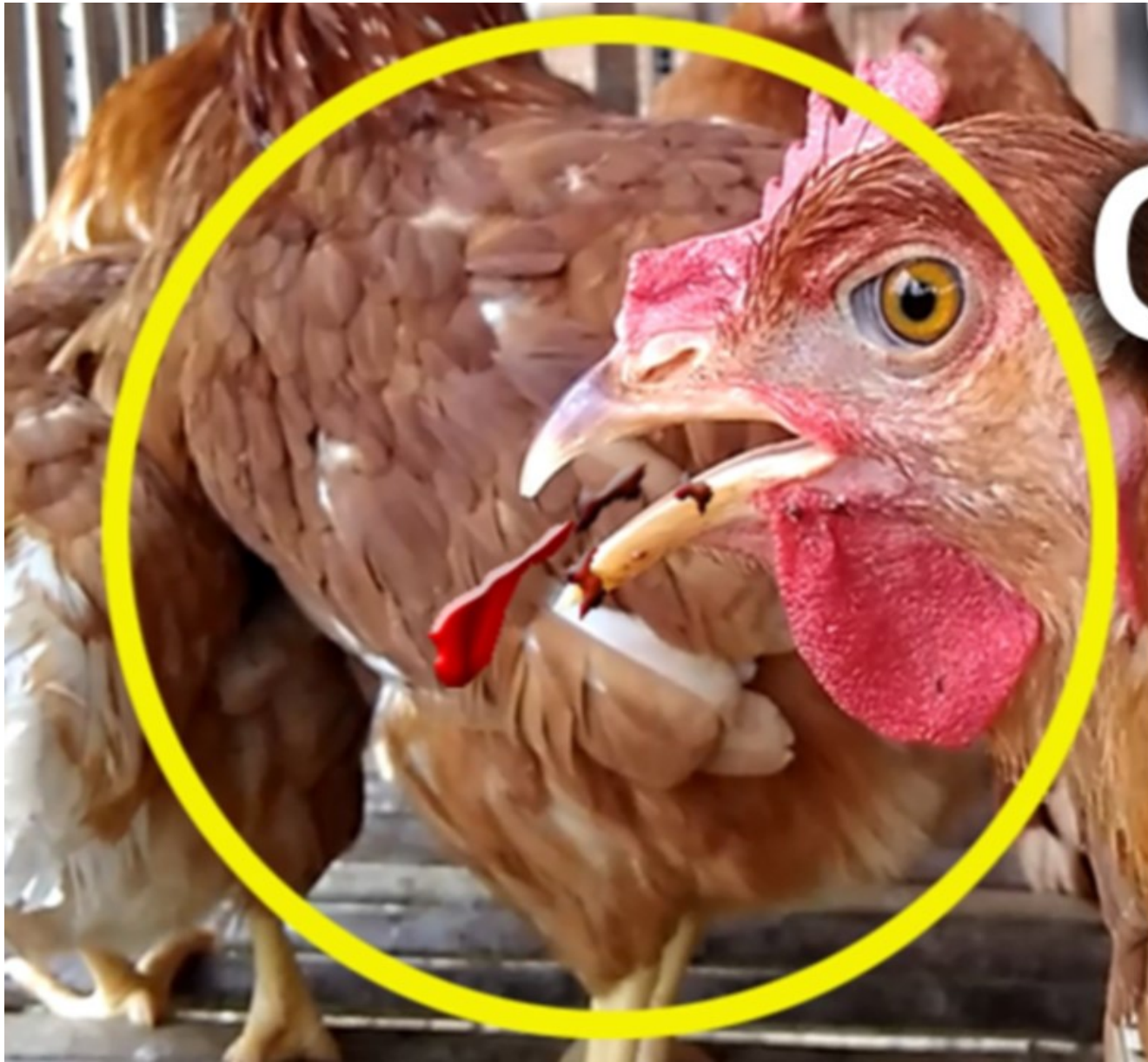
Chicken is stretching neck and opening beak while breathing



chicken suffer from depression and gasping



Swollen eyelid and conjunctivitis



Blood Coughing

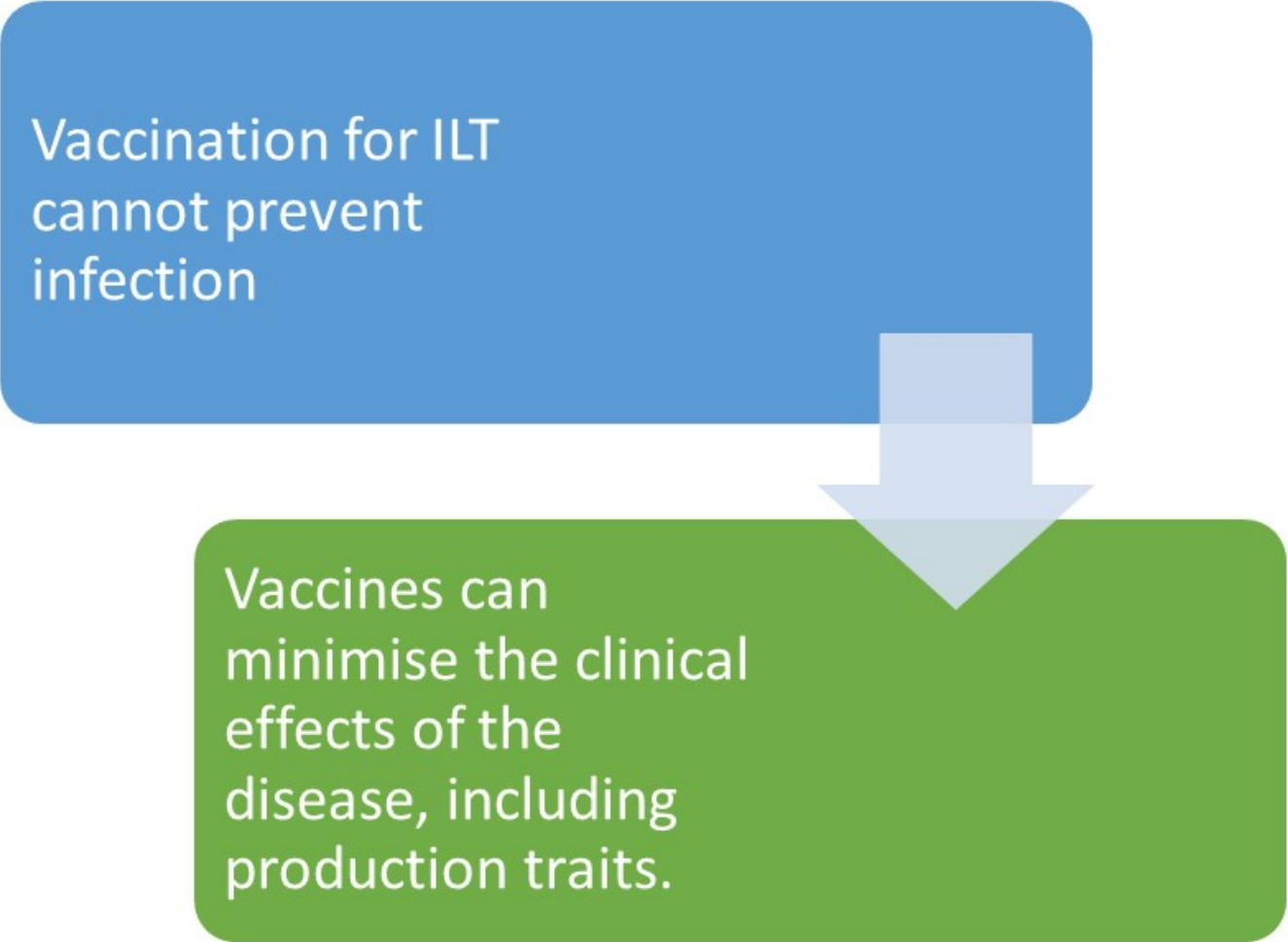
Chicken coughing blood



Lacrimation

INTERVENTION STRATEGIES

Vaccination for ILT cannot prevent infection



Vaccines can minimise the clinical effects of the disease, including production traits.

Vaccination

Type	Administration Route	Age
Modified Live Chick Embryo Origin (CEO)	✓ Eye Drop (preferred method) or Drinking Water or Coarse Spray	~ <ul style="list-style-type: none"> • Dose 1: 3–8 weeks of age. • Dose 2: 9–14 weeks of age, prior to moving to lay house/entering lay.
Modified Live Tissue Culture Origin (TCO)	✓ Eye Drop	~ <ul style="list-style-type: none"> • Dose 2: 10 weeks after dose 1, and prior to moving to layhouse/entering lay.
Vectored Pox-ILT	✓ Subcutaneous Injection	~ <ul style="list-style-type: none"> • Day of Hatch. May need to follow with CEO or TCO prior to entering lay if in high ILT challenge area.
Vectored Pox-ILT	✓ Wing-Web	~ 7–8 weeks of age.

Abbreviation

CEO: Chicken embryo origin

TCO: Tissue culture origin

FieldVet Channel : ILT



ACKNOWLEDGEMENT

The images and part of the content has been taken from www.google.com .The contributors are duly acknowledged

Thank you