

NEWCASTLE DISEASE

- **ETIOLOGY**: *Paramyxo virus.,* RNA Virus.
- ✓ First time recorded in New castle in 1926 in England and in Ranikhet in 1928 in Uttaranchal(India).
- ✓ It occurs throughout the year, but is most common in the summer.
 ✓ On the basis of virulence:
 - 1.LENTOGENIC VIRUS(MILD). 2. MESOGENIC VIRUS(MODERATE). 3.VELOGENIC VIRUS(SEVERE).

ND disease confused with Avian influenza :

- Avian influenza virus-HA-Rabbit but ND doesn't.
- Avian influenza virus doesn't produce disease in pigeons but ND can.
- ✓ ND Virus losses its HA activity at at5,90,180 minutes but pathogenicity lost at 180 minutes but AI Virus become noninfectious before HA property of the virus is lost.

- HOST : In poultry(virulent form), but in ducks, turkey, wild birds and pheasants have less severe form of disease.
 SPREAD :
- ✓ Inhalation and Ingestion.
- ✓ Feed, Water and Equipments.
- ✓ Movement of people.
- ✓ Air transmission is very important.
- ✓ In Humans, the virus causes conjunctivitis, headache and influenza like symptoms.

SYMPTOMS :

In velogenic form :

- ✓ Depression
- ✓ Closed eyes and facial swelling.
- ✓ Drooping wings and anorexia with greenish/yellowish diarrhea.
- ✓ Some times in neural form--- Torticollis, incordination or even paralysis of legs and arched back position of the body.



Paralysis of legs



Abnormal perching reflex



Greenish diarrhea



Coughing and gasping



Torticolli



Edema(Head),Conjunctivitis , Cornea edema(Eye)



Greenish diarrhea with white urates

* In mesogenic form :

- ✓ Severe respiratory distress.
- ✓ Marked drop in egg production.
- $\checkmark\,$ Some times soft shelled egg or shell less egg.
- In lentogenic form:
- ✓ Only mild respiratory distress.

DIAGNOSIS:

- ✓ Isolation of virus.
- ✓ P.M finding :

1. Pin point haemorrhage in proventriculus.

2.Enlarged and haemorrhagic caecal tonsil.

3. Intestinal haemorrhage in intestinal wall.

4. Necrotic spleen.

✓ Serological test :

1.H I Test 2. ELISA Test 3.CFT Test 4. V N Test.

✓ Biological test :

1. Inoculation --- Pigeon-----Produce disease in 6 day.

2. Immunized Poultry----not produce disease where as unimmunized poultry----- produce disease.



Fig 1:Proventriculus shows edematous glands with some areas of hemorrhages.





Fig 2: Caecal tonsils with necrosis and marked hemorrhagic lesions



Fig 3 :Severe hemorrhages on caecal tonsils and mucosa of the rectum.



Fig 4 : Entire length of intestinal mucosa shows hemorrhagic changes.

TREATMENT: No treatment PREVENTION AND CONTROL :

✓ Good hygiene, good management, and good biosecurity practices.

- ✓ Along with vaccination.
- There are 3 types of commercially available vaccines for RD disease:

1. Live lentogenic vaccines:

- ✓ They have F, Hitchner B1, LaSota and V4 vaccines.
- ✓ Least harmful vaccines(LaSota and B1 most widely used).
- ✓ Of these F has lowest disease producing power.

✓ LaSota is not used for first vaccination but often as a booster after one or more B1 strain/ F strain vaccines.

✓ Given by eye drop, drinking water or by machine producing spray.

2. Live mesogenic vaccines:

✓They are Mukteswar (R2b), Roakin, Komarow, and H (Hartford shire).

✓ They are capable of causing severe disease and must be given after earlier vaccination with least harmful vaccines (Live lentogenic vaccines).

✓ Capable of producing a high secondary immune response.

3. *Killed vaccines:*

- ✓ Prepared from both virulent and avirulent form.
- ✓ Given either by i.m or s.c route.

MATERNAL IMMUNITY :

✓ This may prevent the effectiveness of primary vaccination, so, the birds are either vaccinated at 3-4wks of age or vaccinated with live virus at one day old chicks by eye drops or coarse spray and revaccinate at 3-4 wks of age.

✓Killed vaccines successfully in one day old maternally immune chicks.