

Infectious Bursal Disease

Vaccination

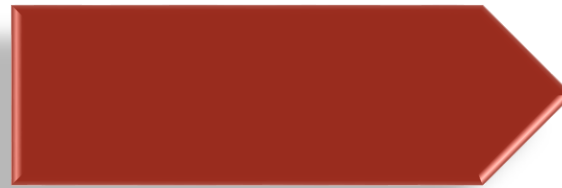
Plan of Talk

- › Hyperimmunisation of breeders
- › Vaccination and maternal immunity
- › Types of vaccines
 - Live vaccines
 - Choosing live vaccine
 - Timing of vaccination
 - Characteristics of good live vaccine

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Parent breeder chickens are hyperimmunised against IBDV.



To protect young chick during the first one to two critical weeks post hatch.



Live IBD Vaccine (2)

4 - 10 weeks of age

Inactivated IBD vaccine (1)

16 - 18 weeks of age or before onset of production

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IBD Vaccine and MDA (Maternal Derived Antibodies)

- › Since IBDV enters through the oral route, and should find its way to the bursa via the blood stream, any MDA in chick blood would neutralize the vaccine.

MDA Decrease

- › MDA is protective when present at a sufficiently high titre.
- › Due to metabolism and growth, the antibody titre declines to half ($t_{1/2}$) at a rate of:
 - › Broilers 3.5 days
 - › Broiler breeders 4.5 days
 - › Layers 5.5 days
- › 2 to 3 weeks post hatch, the susceptibility of a chicken flock to an IBDV infection increases.

MDA & Vaccination

- › Breeder flocks are immunized against IBD, so they would confer protective antibodies to their progenies.

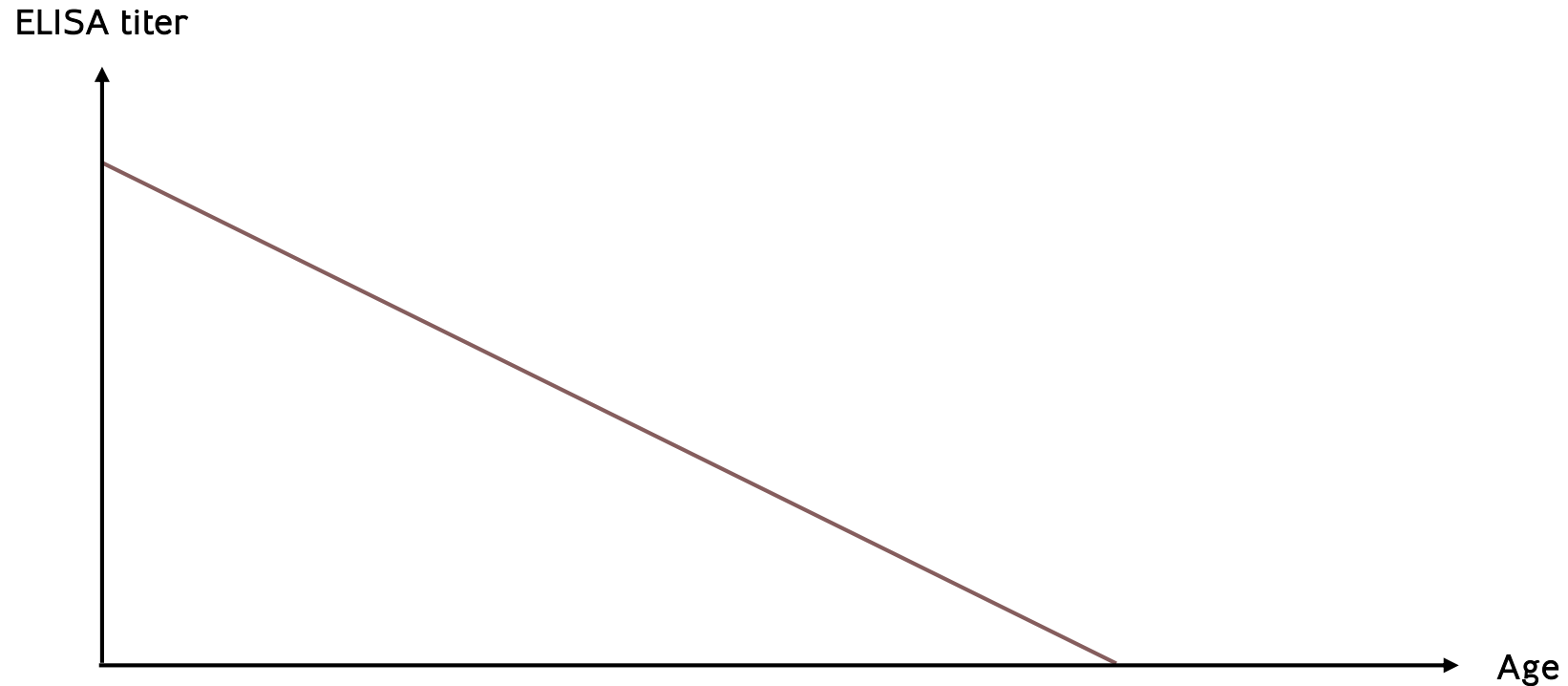
Advantage

- Protects chickens from early infection.

Disadvantage:

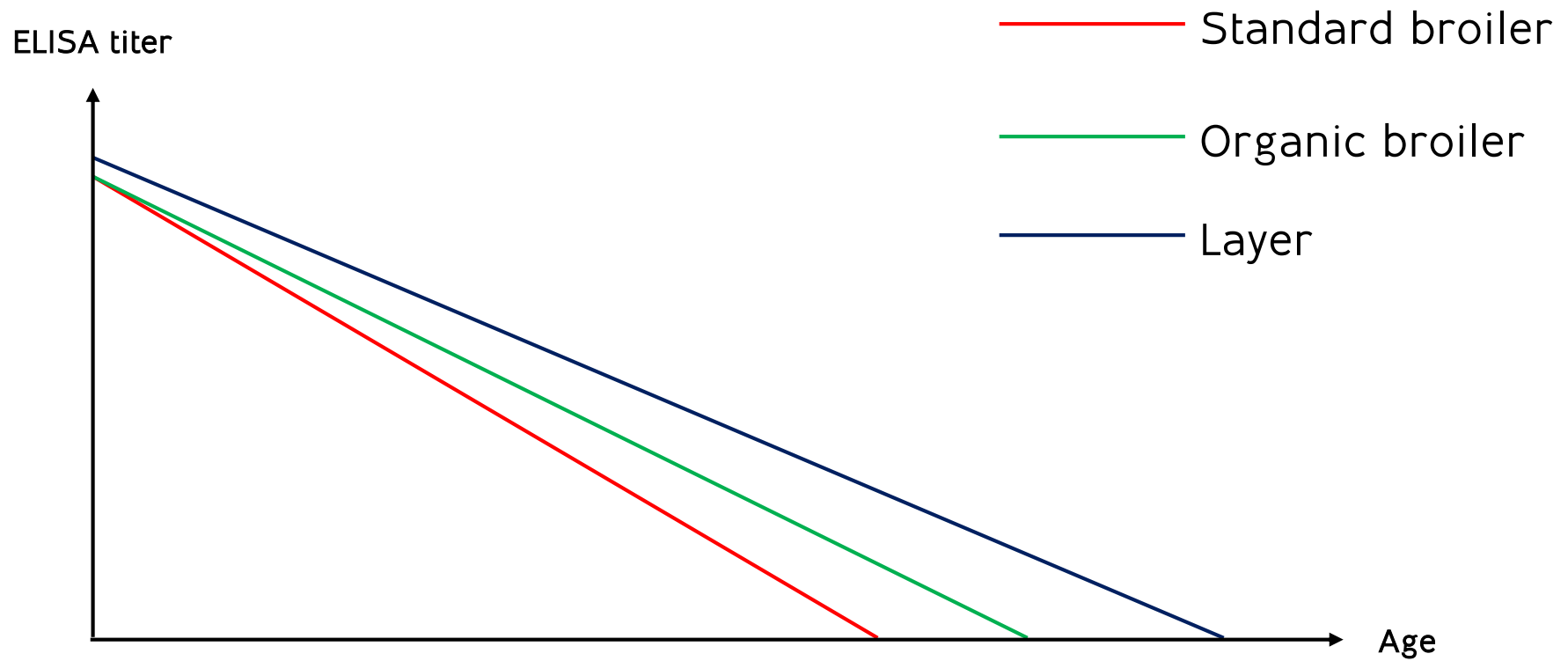
- Neutralizes live vaccines.

MDA Decrease

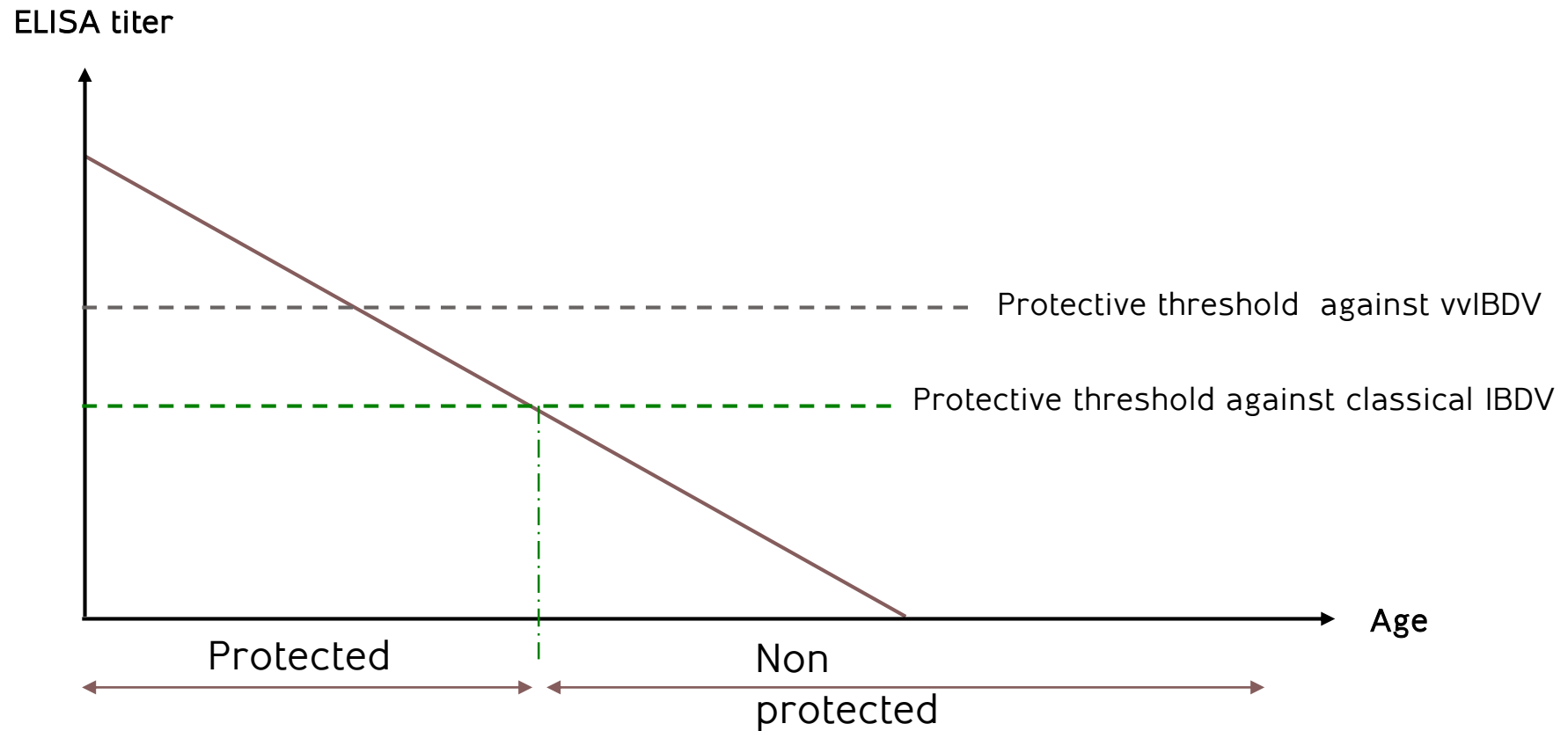


- Maternal antibody will normally protect chicks for 1-3 weeks
- By boosting the immunity in breeder flocks with oil adjuvanted vaccines, passive immunity may be extended to 4 or 5 weeks

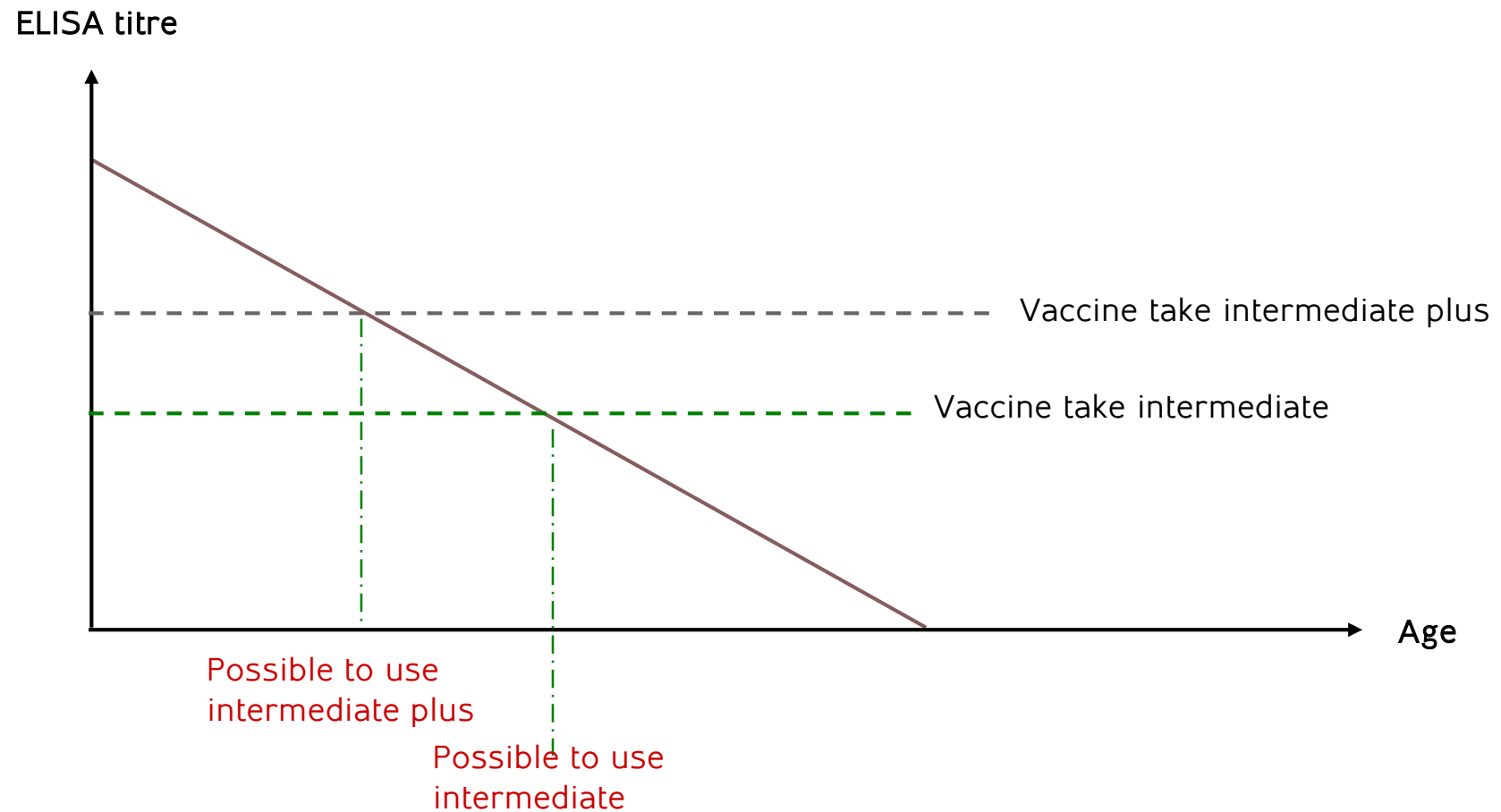
MDA Variation



MDA Protective Threshold



Cont. ...



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Types of IBD vaccines

1. Live attenuated vaccines
2. HVT Recombinant Vaccine
3. Virus-Antibody Complex Vaccines
4. Inactivated vaccines

Live Attenuated Vaccines

Live attenuated vaccines

- › They are administered as the primary defense in the young susceptible chicken.
- › Commercial live vaccines are classified into 3 groups:
 - a) Intermediate strains
 - b) Intermediate plus strains
 - c) Hot strains

HVT Recombinant Vaccine

Composition

- › The concept of recombinant vaccines is to insert genes of critical immunizing epitopes (VP2) of IBD virus into a vector virus HVT (herpesvirus of turkeys).
- › While the vector virus replicates, the VP2 also replicates at the same rate resulting immunization against both.

Precautions

- › The HVT is usually a slow replicating virus, it induces immune response after 14 days.
- › Adding a gene to the HVT may slows its replication and extends the 14 days.

Immune Complex Vaccine

Composition

Antibody specific for the virus

+

Vaccine virus

- › Both are mixed in an appropriate ratio.
- › The antibodies surround the vaccine virus.

Cont. ...

Significance of adding antibodies

- › The presence of antibodies protect vaccine virus from being neutralized by MDA.
- › Antibodies delay, by several days, the normal course of vaccine virus replication.
- › This process controls the time of releasing vaccine virus.

Precaution

- › The level of MDA should be adjusted to the vaccine virus strain.

Inactivated vaccines

- › They are administered to boost the immunity of parent birds.
- › Not used for layers

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Live Vaccines

Objective of using live vaccines:

- › Live vaccines are administered to achieve active immunity.

Classification of Live Vaccines

Live vaccines are classified into 3 groups according to their ability to break through levels of MDA.

1. Mild

- These vaccine strains are highly attenuated.
- They can break through very low levels of MDA.
- They are no longer applicable in the commercial environment.

2. Intermediate

- They are attenuated IBDV strains.
- They can break through MDA titres $\leq 6 \log_2 \text{VN}$

3. Intermediate Plus/Hot

- They are less attenuated IBDV strains
- They can break through MDA titres $\leq 8 \log_2 \text{VN}$

When to Use More Invasive Vaccine?

- › The main feature of the presence of very virulent IBDV field challenge is increased mortality.
- › vvIBDV can break through higher levels of MDA than intermediate IBDV vaccines infections can occur before it is possible to immunise the chicks with intermediate vaccines.
- › To be able to compete with such vvIBDV viruses and induce immunity in the face of still high maternal immunity, more invasive IBDV vaccines strains are required.
- › The main objective is to reduce mortality and the prevalence of vvIBDV in the flocks
- › Once this aim has been reached return to intermediate vaccines

General Recommendations

- › In farms with IBD problems or in "hot areas"
 1. Use a hot or intermediate plus vaccine 3 - 4 cycles;
 2. Followed by an intermediate vaccine
- › In "endemic" areas, multiple age farms and farms relying solely on hot vaccines
 1. Use continuously a hot vaccine.

Whenever possible try to switch back from a hot vaccine to an intermediate

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Choosing Live Vaccine

Choice of live vaccine depends on:

1. Virulence of field infection

- Mild and Intermediate vaccine strains can not protect against vvIBDV.
- Mild and Intermediate vaccine strains cannot be administered at an early age due to MDA interference.

2. Age of chickens to be vaccinated

- The earlier the vaccination the higher the level of MDA, requiring a stronger vaccine.

3. Age at which Gumboro outbreak occurs.

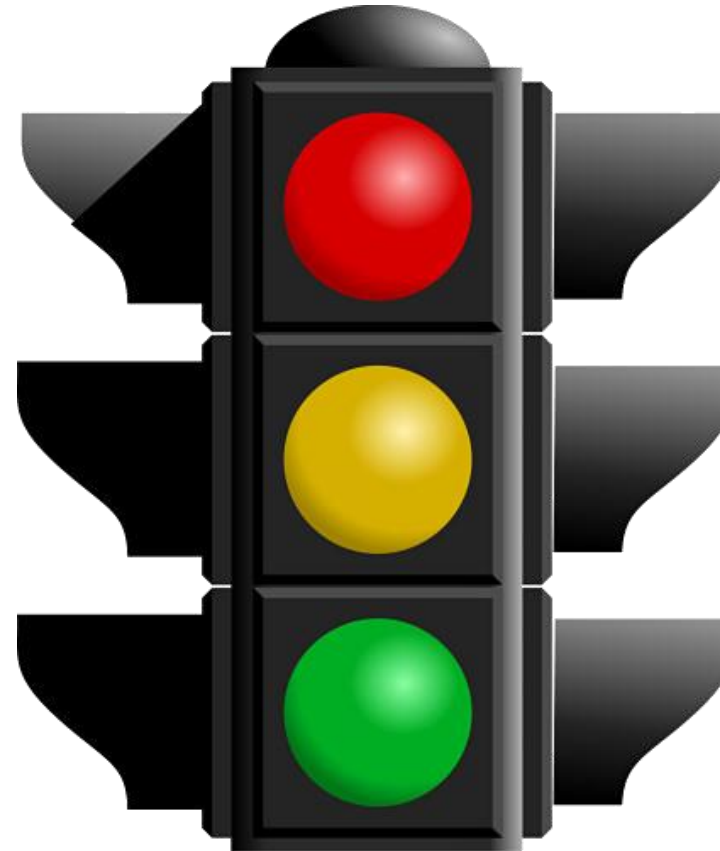
- Early outbreaks require earlier vaccination.
- Infections before 2 weeks require using vaccine that cover variant strains.
- Late infections requires intermediate plus or hot strains.

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Timing of Vaccination

- › **Too soon**
 - MDA neutralises vaccine
- › **Late**
 - Late protection
- › **Optimal**
 - Sooner the better



Cont. ...

There is no IBD vaccination schedule that can be routinely recommended.



Cont. ...

Factors influencing a vaccination schedule include:

1. Type of chicken to be vaccinated (broiler or commercial layer).
2. Level of MDA
 - The higher the start level of MDA the later the age of vaccination.
3. Uniformity of MDA
 - If the variation in MDA levels is too high ($CV > 30\%$) a second IBD live vaccination is required to effectively immunize the flock.
4. Field pressure.

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Characteristics of Good Life Vaccine

1. Early protection.
2. Minimum immunosuppression.
3. Highest antibody titer.
4. Cross protection between different strains.
5. Easy application.

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