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List of abbreviations

| | |
|------------------------------------|--------------------|
| Binary ethylene amine | BEI |
| Bovine viral diarrhea | BVD |
| Bovine viral diarrhea virus | BVDV |
| Border disease virus | BDV |
| Cytopathic effect | CPE |
| Cell RNA | CRNA |
| Cod of federal regulation | CFR |
| Enzyme linked immunosorbant assay | ELISA |
| Hog cholera virus | HCV |
| Infectious bovine rhinotracheitis | IBR |
| Intra venous | I/V |
| Internal ribosome entry site | IRES |
| Madine derby bovine kidney | MDBK |
| Minimum essential medium | MEM |
| Mucosal disease | MD |
| National animal disease laboratory | NADAL |
| Non cytopathic | NCP |
| 3'nontranslated region | 3'UTR |
| Open reading frame | ORF |
| Parainflunza-3 | PI-3 |
| Persistent infected | PI |
| Polymerase chain reaction | PCR |
| Post vaccination | PV |
| Replicative intermediate | RI |
| Replicative form | RF |
| Revolution per minute | R.P.M. |
| Serum neutralization test | SNT |
| United state of America | USA |
| 5untranslated region | 5'UTR |
| Viral RNA | VRNA |
| 50% tissue culture infected dose | TCID ₅₀ |

summary

bovine viral diarrhoea virus is economically important pathogen of cattle through the world. The present study was planned for preparation, evaluation inactivated bovine viral diarrhoea virus vaccine contained genotype I&II and comparative with inactivated pneumo-3 vaccine contained BVDV,IBRV and PI-3V.

Vaccine prepared through propagation of the BVDV genotype I&II in Madin-Darby bovine kidney cell culture (MDBK) and used BAI for inactivation at 0.02% concentration for 6 hours at 37°C. add 20% alhydrogel as adjuvant.

The evaluation of prepared inactivated BVDV vaccine containing genotype I&II was conducted in laboratory as follows:

1. Sterility and purity tests:

these results were regulated through the American protocol, US code of Federal Regulations (CFR, 1997). The obtained results proved that the prepared vaccine were free from any bacterial, fungal and mycoplasma contamination as well it was proved that the prepared vaccine were free from other viral agent.

2. Safety test:

Safety test was performed on mice and guinea pigs and revealed that no clinical abnormalities were observed through 10 days of observation in mice and guinea pigs.

3. potency and challenge exposure tests:

The test were performed on fourteen cross breed calves (Friesian and local) about 6 month of age divided into two groups the result revealed that:

a. Vaccinated calves subgroup A :

No clinical finding were recorded in vaccinated calves up to one month post vaccination. the resulted of immune responses of calves post vaccination and the titer of neutralizing antibodies reached its peak at 2 months then begin to decline but remained protective till end of experiment.

b. Vaccinated challenged calves (challenge exposure test) subgroup B:

No clinical finding through out the challenged period and the immune response capable for protecting calves against experimental infection using virulent strain of genotype 1 and 2 of BVDv. the resulted of immune responses of calves post challenged, the mean of serum neutralizing antibody titers post challenge decreased at 3 day post challenged in which become 1.55 to genotype I and 1.50 to genotype II then titer where raised again till reached its peak at 21 days post challenge 2.25 genotype I, 2.15 genotype II.

c. The non vaccinated control infected calves subgroup c:

Showed high significant increase in the mean body temperature during the first eight days of challenge. The recorded body temperature

was 40.3°C on the 3rd day post infection and raised to 40.4 at the fourth day and remained high till eight days. The temperature started to subside gradually from the 9th day post infection and return to the normal on the nine day post infection. The observed clinical signs were dullness, off food, serous nasal discharge begin to appear with two days, faces become moisten and ocular discharge appeared. These previous symptoms lasted up to the 10th-12th days after which complete recovery occurred. Serum neutralize antibody titers reach its peak at 28 day 1.95.

d. The non vaccinated, non infected control calves subgroup D:

Neither clinical signs nor humeral immune response could be detected in these calves.

3. Evaluation of prepared vaccine under field condition:

The test was done on pregnant cows at last third of gestation result revealed that cow dams gave off springs immunized with BVDV genotype I&II antibodies within the protective level up to six months after delivery.

4. Comparative between inactivated bivalent vaccine containing genotype I&II and combined inactivated vaccine pneumo-3 containing (BVDV, BRV, PI-3V) revealed that bivalent vaccine give higher antibodies to BVDV than produced due to vaccination by combined inactivated vaccine pneumo-3 in cow dams and their off springs.