Serodiagnosis of Foot and Mouth Disease (FMD) Virus for Differentiation Between Naturally Infected and Vaccinated Cattle and Buffaloes

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ABSTRACT
FMD is a highly contagious viral disease of all cloven-footed animals and widely distributed all over the world. In this study, 465 serum samples were collected from 3 Nile delta governorates (Behaira, Mounofya and Kafer El-Sheikh) during 2009. The samples were used for detection of FMD antibodies to 3ABC non-structural proteins using commercial ELISA kit (Priocheck). The overall percentage of positive was 38.9%. The higher percentage of positive detected in Behaira (48%), then Mounofya (45.3%) while Kafer El-sheikh was the lowest (23.7%). The positive results of detection of antibodies against non-structured proteins of FMDV indicate that these samples come from natural infected animals.

Key words: Detection, FMD, FMD antibodies ELISA, non-structural proteins

INTRODUCTION
FMD is one of the most contagious epidemic diseases of livestock can spread very rapidly. It is caused by 7 immunologically distinct serotypes, O, A, C, Asia 1, South African Territories (SAT) 1, SAT 2 and SAT 3 which belong to the species Foot-and-mouth disease virus (genus Aphthovirus, family Picornaviridae). Several of these serotypes circulate currently or periodically in the Middle East and North Africa (Knowles and Samuel, 2003). It is characterized by the formation of vesicles in and around the mouth and on the feet, it reduces feeding and often causes lameness. Abortion, sterility, permanent decline in milk yield, decrease in meat production and reduction in breeding ability are common sequelae. Mortality can result and although low for adult animals, can be higher than 50% in the young. The virus needs to be eliminated to re-establish disease-free status; failure would have serious economic consequences (Commission of the European Communities, 1989; OIE, 1999).

Early warning is therefore essential to detect an incursion while it is still localized. Early and decisive reaction is required if the disease is to be contained and eventually eliminated without serious socio-economic consequences. To be effective, the control measures must be applied in the shortest possible time is crucial to success. Cattle are regarded as good indicator hosts, whereas, sheep tend to show few clinical signs and are often considered maintenance hosts for a relatively short period where movement and transport can be responsible for virus spread. Infected animals
may excrete the virus for up to a few days before exhibiting clinical signs. A proportion of recovered cattle, African buffaloes and sheep remain virus carriers for variable periods. Wild or feral ruminant or porcine animal populations may act as reservoirs for infection. Direct contact between animals is the most significant method of transmission, but the virus may persist for considerable periods in the environment (particularly in temperate climates) and mechanical transmission by fomites is also considerable. Windborne spread over considerable distances is possible in temperate climates.

Inactivated vaccines are widely used for FMD but vaccine strains must be carefully matched to prevailing field virus strains if a satisfactory level of protection is to be attained; vaccination cover must attain a level of at least 80% for effectiveness. Serological tests that allow discrimination between antibodies resulting from infection and vaccination (NSP ELISA tests) are now becoming available and should permit more accurate monitoring of control and eradication programmes based on mass vaccination. FMD vaccination is, unfortunately, still carried out in a haphazard manner in many countries, resulting in the disease remaining endemic for long periods.

A test which differentiates the vaccinated and infected animals antibodies would be of great value in FMD control. Several tests which are based on Non-Structural Proteins (NSP) have been described (Berger et al., 1990; Neitzert et al., 1991; Bergmann et al., 1993; Lubroth and Brown, 1995). For the screening of large numbers of samples an ELISA would be highly preferable. An indirect-trapping ELISA for the detection of antibodies against 3ABC has been reported (De Diego et al., 1997). The sensitivity of the assay on experimental sera post-infection was reported to be 100%. The specificity was reported to be more than 99%.

In Egypt, routine prophylactic vaccination has been conducted with a locally produced serotype O vaccine. The last outbreak of serotype O was in June 2000 and other serotypes have not been reported since 1972 when serotype A occurred (Ferris and Dawson, 1988). OIE (2006) and Knowles et al. (2007) described an FMD serotype-A virus responsible for recent outbreaks of disease in Egypt.

This study describes the using of commercial PrioCHECK, FMD NS ELISA kit, to cattle and buffaloes populations and the application of the developed kits to the serological surveillance system to monitor the progress of the FMD control program in three Nile Delta Governorates.

MATERIALS AND METHODS

Samples: Sera were randomly collected from vaccinated cattle and buffaloes in three Egyptian Governorates (Behaira, Mounofya and Kafer EL-Sheikh). Age of animals ranged from less than one year to more than 2 years. The samples data is showing in Table 1.

<table>
<thead>
<tr>
<th>Governorates</th>
<th>Total age</th>
<th>Age class-1**</th>
<th>Age class-2**</th>
<th>Age class-3***</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>B</td>
<td>C</td>
<td>class-1</td>
<td>B</td>
</tr>
<tr>
<td>Behaira</td>
<td>47</td>
<td>5</td>
<td>23</td>
<td>23</td>
</tr>
<tr>
<td>Kafer EL sheikh</td>
<td>55</td>
<td>11</td>
<td>28</td>
<td>30</td>
</tr>
<tr>
<td>Mounofya</td>
<td>95</td>
<td>12</td>
<td>33</td>
<td>45</td>
</tr>
<tr>
<td>Total</td>
<td>137</td>
<td>28</td>
<td>84</td>
<td>98</td>
</tr>
</tbody>
</table>

*Age class-1 (age less than 1 year), **Age class-2 (age less than 2 year), ***Age class-3 (age more than 2 year)
Table 2: Results of PrioCHECK, NSP of FMDV ELISA test

<table>
<thead>
<tr>
<th>Governorates</th>
<th>No. of cattle samples</th>
<th>Cattle No. of +ve (%)</th>
<th>No. of buffaloes samples</th>
<th>Buffaloes No. of +ve (%)</th>
<th>Total No. of samples</th>
<th>Over all +ve (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Behaira</td>
<td>106</td>
<td>45 (45)</td>
<td>50</td>
<td>27 (54)</td>
<td>150</td>
<td>72 (48)</td>
</tr>
<tr>
<td>孟fuseya</td>
<td>88</td>
<td>48 (54.54)</td>
<td>71</td>
<td>24 (33.8)</td>
<td>159</td>
<td>72 (45.3)</td>
</tr>
<tr>
<td>Kafer El sheikh</td>
<td>58</td>
<td>22 (22.44)</td>
<td>98</td>
<td>15 (25.9)</td>
<td>156</td>
<td>37 (23.7)</td>
</tr>
<tr>
<td>Total</td>
<td>246</td>
<td>115 (40.21)</td>
<td>219</td>
<td>66 (36.9)</td>
<td>465</td>
<td>181 (38.9)</td>
</tr>
</tbody>
</table>

Table 3: Seropositive percentage by Priocheck FMD NSP in relation to the age

<table>
<thead>
<tr>
<th>Age class</th>
<th>Results</th>
<th>1*</th>
<th>2**</th>
<th>3***</th>
<th>Total</th>
</tr>
</thead>
<tbody>
<tr>
<td>N</td>
<td>87</td>
<td>51</td>
<td></td>
<td>146</td>
<td>284</td>
</tr>
<tr>
<td>POS</td>
<td>50</td>
<td>33</td>
<td></td>
<td>98</td>
<td>181</td>
</tr>
<tr>
<td>Pos,%</td>
<td>36.5</td>
<td>39.29</td>
<td></td>
<td>40.2</td>
<td>38.9</td>
</tr>
<tr>
<td>Total</td>
<td>137</td>
<td>84</td>
<td></td>
<td>244</td>
<td>465</td>
</tr>
</tbody>
</table>

*Age class 1: Age less than 1 year; **Age class 2: Age less than 2 year; ***Age class 3: Age more than year

**ELISA kit:** The commercial PrioCHECK, FMD NS ELISA kit for in vitro detection of antibodies against FMD virus in serum of cattle, sheep, goat and pigs. The kit is used according to its instructions. Samples give percent of inhibition IP<50% considered negative and that give IP>50% considered positive.

**Statistical analysis:** This analysis was conducted using excel and Pivot table.

**RESULTS**

As shown in Table 2 the number of over all positive samples are 181 samples out of 465 samples (38.9%). The highest present of positive were found in Behaira Governorate (48%), Mounofya Governorate (45.3%) and the lowest one was Kafer El sheikh Governorate (23.7%). The positive percent in buffaloes (36.9%) was lower than cattle (40.21%) in the over all samples. Although, the percent of positive in buffaloes (54%) is higher than that of cattle (45%) in Behaira Governorate and buffaloes (25.9%) is higher than that of cattle (22.44%) in Kafer El sheikh Governorate.

As shown in Table 3 the percent of positive according to age at less than one year, less than 2 year, more than 2 years are respectively, 36.5, 39.29 and 40.2%, respectively. The highest percent of positive found at samples collected from animals more than 2 years. While the lowest found at samples collected from animals less than one year.

**DISCUSSION**

Foot and Mouth Disease (FMD) is the major disease constraint on international trade in livestock and their products. Effective vaccines and control measures have enabled the FMD unvaccinated, seronegative herds in compliance with strict international trade policies. However, the disease remains enzootic in many regions of the world; it is a serious problem for commercial trade with FMD-free countries (Bhattacharya et al., 2005).

Vaccination plays an important role in the control of FMD in Asia, Middle East, Africa and South America. In most FMD-free countries a non-vaccination policy is in place. Recent outbreaks
in Europe clearly demonstrated the risk of this policy. Using conventional diagnostic techniques, up to now it was not possible to distinguish FMD infected animals from purely vaccinated animals. In vaccinated areas disease control authorities had limited possibilities to monitor virus presence or circulation (Van Aarle, 2001). Art-vaccines are based upon highly purified antigens which are free from Non-Structural Proteins (NSP) of the FMD virus. Other vaccines may be partly purified and contain a reduced amount of NSP. Animals, antibodies against the Structural Proteins (SP) but not against NSP. Modern, state of the FMD virus infection induces antibodies against both SP as well as NSP. NSP-free or NSP-reduced vaccines in combination with a NSP-test lead to a so called marker-system (Van Aarle, 2001). An ELISA using baculovirus-expressed 3AB and 3ABC as the antigens has been demonstrated to successfully differentiate vaccinated from infected cattle and sheep (Sorensen et al., 1998).

In May 2006, the new bivalent vaccine was locally produced containing both O1 and A/Egypt/2006 local isolates and used for routine vaccination of Egyptian animals.

The results of FrioCHECK ELISA test in this study proved the presence of antibodies against NSP of FMDV in cattle and buffalo population in Behaira, Mounofya and Kafer El-Sheikh Governorates which may be attributed to natural infection of FMDV. The percent of positive was 38.9%. A proportion of vaccinated animals can become sub-clinically infected if they are subsequently exposed to the homologous virus and may be able to transmit infection for up to 14 days after vaccination, even when they become immune to the development of clinical disease (FAO), or due to the impurity of the inactivated vaccine (Chung et al., 2002).

As shown in Table 2, the number of total positive samples is 181 samples out of 465 samples (38.9%). The highest present of positive are found in Behaira Governorate (48%), Mounofya Governorate (45.3%) the lowest one is Kafer El Sheikh Governorate (23.7%). Recent outbreaks were reported in Behaira Governorates in September 2007 and January 2008 by serotype O (FAO, 2008).

The positive percent in buffaloes is lower than cattle in the over all samples (36.9%). This may be indicate the presence of high resistance of buffaloes to FMDV than cattle this is agreed with Ghoneim et al. (2010). Although, the percent of positive in buffaloes (25.9%) is higher than that of cattle (22.44%) in Kafer El sheikh Governorate.

As shown in Table 3, the percent of positive according to age at less than one year, less than 2 year, more than 2 years are respectively, 36.5, 39.29 and 40.2%. The highest percent of positive found at samples collected from animals above 3 years. While the lowest found at samples collected from animals less than one year. This indicates that the immunity afforded by vaccines does not last long (Geering and Lubroth, 2002).

To apply the differentiating diagnostic tool, the sera from vaccinated cattle and buffaloes would be preferred for evaluating the usefulness of these kits. The immune response to nonstructural proteins has been reported to develop later than that to structural proteins in the course of infection (De Diego et al., 1997; Sorensen et al., 1998). Following experimental infection, antibodies to the 3ABC antigen could not be detected earlier than 8 and 10, days in cattle, sheep (Sorensen et al., 1998), respectively. Since, all positive sera described in this report were collected from cattle and buffaloes vaccinated with FMD vaccine, the actual time of the first FMDV contact was unknown. Therefore, the earliest time after infection that the assay could detect an immune response to the 3AB antigen was undetermined.
CONCLUSION
As described above, the 3ABC PrioCHECK, FMD NS ELISA has promising sensitivity and specificity to distinguish FMDV-infected animals from vaccinated animals. This kit has also been demonstrated to be useful for monitoring the progress of the FMD eradication program in Egypt.

REFERENCES
Commission of the European Communities, 1989. Report from the Commission to the Council on a study carried out by the Commission on policies currently applied by Member States in the control of foot-and-mouth disease. SEC (89) 1731 final, Pages: 61.


