Guidelines for Bovine Viral Diarrhea Virus (BVDV) Testing

The NDSU-VDL offers polymerase chain reaction (PCR) and immunohistochemical (IHC) based assays to identify animals infected with BVDV. Which method will be most cost effective depends on the prevalence of persistently infected (PI) animals in the herd and number of animals being tested. In most instances, unless testing 6 or fewer animals or there is a high herd prevalence of PI animals, PCR testing will be the most cost effective method.

- Refer to [http://www.vdl.ndsu.edu/](http://www.vdl.ndsu.edu/) or current fee schedule for price and test day information.

Collection procedure for PCR testing: Care must be taken to avoid cross-contamination of samples. Use an ear notching tool that yields a 1-2 cm² notch. Disinfect the notching tool with 10% bleach solution between each animal and rinse the notching tool in clean water. Residual disinfectant on the notching tool may give false negative results, therefore rinsing is required. Place each ear notch in its own dry (NO FORMALIN), individually labeled red top blood tube or similar container (sterile container, must not contain preservatives). The laboratory will pool up to 15 ear notches per pool; individuals in positive pools will be automatically retested and charged accordingly.

- Ear notches must be free of dirt, residual disinfectant, feces, tattoo ink, or BVDV vaccine. Do not vaccinate or tattoo at the same time the samples are taken.
- Label samples and place on ice as soon as they are collected. Keep refrigerated or frozen until shipped. Samples may be kept in the freezer up to one month and shipped together on ice.

Collection procedure for IHC Testing: Cross contamination is less of a concern in IHC testing, therefore disinfecting the ear notch tool and rinsing in water is not required but is recommended to stop the transmission of other diseases. Place samples in individually labeled tubes containing sufficient formalin to cover the ear notch. DO NOT FREEZE. Samples should be tested within a couple weeks of collection. Prolonged storage in formalin will yield false negative results. The laboratory will place up to 6 samples per slide/charge.

Sample Submission: Complete the Herd Serology/PCR submission form. Enclose a copy with samples.

- Enter animal identification on the Sample Identification Form. Age, sex and breed are not required.
- Enclose a copy of the Sample Identification Form with the samples and email the electronic form (do not scan or fax) to ndsu.vetlab@ndsu.edu using the owner’s last name as the subject.

Interpretation of Results:

<table>
<thead>
<tr>
<th>Result</th>
<th>Interpretation</th>
</tr>
</thead>
<tbody>
<tr>
<td>Not Detected/Negative</td>
<td>The animal is not persistently infected with BVDV.</td>
</tr>
<tr>
<td>Detected/Positive</td>
<td>The animal is either persistently infected or acutely infected with BVDV.</td>
</tr>
<tr>
<td>Suspect</td>
<td>Very low level of virus was detected. This can occur during convalescence from acute infection or represent carryover of virus from improperly disinfected and/or rinsed notching tool.</td>
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</tbody>
</table>

Retesting of positive animals: IHC testing typically only identifies persistently infected animals, but we recommend retesting positive animals at least 3 weeks later to document persistent infection. PCR testing requires retesting of positive animals as it readily identifies both persistently infected and acutely infected animals.

Please contact the laboratory with any questions at 701-231-8307 or visit [www.vdl.ndsu.edu](http://www.vdl.ndsu.edu).

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