IOWA STATE UNIVERSITY

Veterinary Diagnostic Laboratory

Pathology Submission Guide

BOVINE ABORTION

Specimens to submit: Tissues are received in best condition if removed at necropsy in the field.

Samples should include:

Brain Formalin-fixed (1/2 of entire tissue)

Dam's serum 3 - 5 ml from affected cows. Optional, see notes on abortion serology.

Heart Formalin-fixed (1/2 cm slice)

Ileum Formalin-fixed

Kidney/Liver/Spleen Fresh, formalin-fixed (1/2 cm slice)

Lung Fresh (cranioventral), formalin-fixed (1/2 cm slice)

Placenta (very important) 3 cotyledons, fresh; 2 cotyledons, formalin-fixed (please submit placenta

when possible - this increases the diagnostic success rate)

Skeletal Muscle Tongue and diaphragm formalin-fixed (1/2 cm slice)

Skin (lesions/ear notch) Formalin-fixed (1/2 cm slice)

Stomach content 1-3 ml in sterile syringe or tube, fresh

Thoracic fluid 1-3 ml in sterile syringe, fresh

Thymus/Adrenal gland Fresh, formalin-fixed (1/2 cm slice) *Alternatively, the entire fetus and placenta can be submitted.

SAMPLING SUGGESTIONS

1. Do NOT freeze fresh tissues; keep them chilled.

- 2. Always submit placenta if possible! Failure to submit placenta severely diminishes the diagnostic success rate of bovine abortion cases.
- 3. It may be useful to submit serum from affected and unaffected dams.

COMMON INFECTIOUS AGENTS DETECTED BY ROUTINE EXAMINATION, CULTURE, AND PCR OF FETAL AND PLACENTAL TISSUES

Bacteria Trueperella (Arcanobacterium) pyogenes, Bacillus spp., Brucella spp., Campylobacter spp.,

Histophilus somnus, Salmonella, Listeria monocytogenes, Leptospira, Ureaplasma,

Mycoplasma (PCR or culture), etc.

Fungi Aspergillus, Phycomycetes

Protozoa Toxoplasma gondii, Neospora caninum, Tritrichomonas foetus

Viruses IBR, BVD

AGENTS REQUIRING SPECIAL TESTS (BY REQUEST)

Eyeball (aqueous) Nitrate/nitrite

- If leptospirosis is suspected, extra effort should be made to deliver freshly aborted, chilled fetuses directly to the lab. PCR testing can be conducted on fetal tissues (kidney). Serology on affected dam sera is very helpful.
- Diagnosis of Neospora caninum abortion is based on histopathologic examination of brain, heart, skeletal muscle, liver, lung, and placenta for characteristic lesions. Presence of the organism can be confirmed by immunohistochemistry. Absence of serum antibody in the cow would rule out neosporosis.
- *Tritrichomonas foetus* infection is best diagnosed by placing preputial wash or fetal fluids/stomach contents directly into TF pouch for PCR.

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BOVINE CENTRAL NERVOUS SYSTEM DISORDERS

Specimens to submit: Tissues from euthanized or dead animals including:

Blood sample EDTA tube for lead analysis or cholinesterase inhibition

Eyeball (aqueous) Cations (calcium); nitrite

Brain (including brain DO NOT FIX BRAIN IF RABIES TESTING IS DESIRED

stem) Non-Rabies: split longitudinally, 1/2 brain fresh; 1/2 brain, formalin-

fixed

Colon Optional, nervous coccidiosis. Several partial loops with contents, fresh.

1 cm pieces, formalin-fixed

Liver Optional, lead toxicosis. Fresh Kidney Optional, lead toxicosis. Fresh

Peripheral Nerves Depending on clinical signs. Fresh and fixed Skeletal Muscle Depending on clinical signs. Fresh and fixed

Spinal cord Optional, locomotor involvement

Entire carcass or vertebral column, fresh OR

Dissected cord, fresh w/cross-sections (1/2 cm) of cord from 4-5 levels,

formalin-fixed

Rumen content Fresh/chilled

SAMPLING SUGGESTIONS

- 1. Entire head can be submitted. Chill all samples before shipment if possible.
- 2. Do NOT freeze fresh brain or head.
- 3. Fresh half of brain should be packed carefully to avoid crushing.
- 4. Fixed half of brain should be incised, at least once, transversely (not longitudinally) into the lateral ventricle to aid fixation if the brain is large.

COMMON AGENTS DETECTED BY ROUTINE EXAM AND/OR CULTURE

Bacteria Histophilus somnus, Listeria monocytogenes, Trueperella (Arcanobacterium)

pyogenes, etc.

Non-infectious Polioencephalomalacia

AGENTS REQUIRING SPECIAL TESTS (BY REQUEST)

Deficiencies Magnesium (serum, entire eyeball, fresh/chilled), calcium (serum,

fresh/chilled)

Parasites Coccidia (flotation; feces, fresh/chilled) - NO lesions in brain

Toxicoses Lead (whole blood in EDTA, liver, stomach contents, fresh/chilled),

organophosphate (whole blood in EDTA, brain, rumen, fresh/chilled)

Sodium (whole blood in EDTA, brain, rumen, fresh/chilled)

Viruses Rabies (FA - requires entire brain to be submitted fresh/chilled),

herpesviruses: IBR, pseudorabies (PCR - brain, fresh/chilled)

COMMENTS

• Cerebellum and brain stem are affected by most infectious causes of CNS disease and should always be included in submitted samples.

 Many toxic, nutritional, and metabolic causes of CNS disease do not induce lesions in the brain and must be diagnosed by analysis of other tissues. For most toxicoses, submission of rumen contents, complete feed, water and feed components, liver, kidney, serum, and whole blood (in EDTA) as well as brain would include the tissues necessary for diagnosis.

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BOVINE ENTERITIS - CALVES < 2 MONTHS OF AGE

Specimens to submit: Antemortem fecal samples are of value if collected on the first day of diarrhea.

Tissues collected from a euthanized or dead calf should include:

Abomasum Fresh and formalin-fixed
Cecal contents 10 ml fluid contents, fresh
Colon Several partial loops, fresh

2-3 1cm pieces, formalin-fixed

Ear notch Formalin-fixed

Ileum Two or three 10-15 cm segments, fresh

Three 1 cm pieces, formalin-fixed

Jejunum Two or three 10-15 cm segments, fresh

Three 1 cm pieces, formalin-fixed

Kidney/Liver/Lung Fresh and formalin-fixed Mesenteric lymph node Fresh and formalin-fixed

Rumen Fresh content and formalin-fixed tissue

Spleen/Thymus Fresh and formalin-fixed

Because autolysis occurs very quickly in bovine intestines, samples removed at necropsy in the field and properly preserved soon after death are usually better than a whole dead calf submitted to the lab.

SAMPLING SUGGESTIONS

- 1. Samples must be taken as soon after death as possible.
- 2. Fresh samples should be chilled quickly. DO NOT FREEZE.
- 3. Intestines do not need to be tied off at the ends.
- 4. Flush intestinal segments for histopathologic examination with formalin and drop in fixative. Or, gently open ends of 1 cm segments with scissors or forceps to expose mucosa as immersed. Do not split open.
- Pool all formalin-fixed tissues from each calf in one bag; individual calves can be pooled or kept separate as desired. Package fresh intestines separately from other tissues and each calf in a separate bag.

COMMON AGENTS DETECTED BY ROUTINE EXAM, CULTURE, PCR, AND/OR FECAL FLOTATION

Bacteria E. coli, Salmonella spp., Clostridium spp.

Parasites Cryptosporidia, Coccidia

Viruses Rotavirus, Bovine coronavirus, BVD virus: IHC on lymphoid tissue and

skin, PCR on fresh lymphoid tissue/lung

- In cases of necrotic enteritis, submit both necrotic and adjacent non-necrotic segments fresh and fixed
- In-house quick tests (acid-fast stained impression smears) may be of value for detection of cryptosporidia. The preferred site for impression smears/mucosal scrapings for cryptosporidia is ileum. As such, it is helpful if fresh ileum is submitted in a separate container.
- Colon is the preferred tissue in which to identify lesions of coronavirus enteritis, for laboratory confirmation with BCV IHC, and for observation of coccidia. Colon should be submitted with all calf diarrhea cases.



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BOVINE ENTERITIS - CALVES > 2 MONTHS OF AGE, FEEDLOT CATTLE, ADULTS

Specimens to submit: Fecal samples may be of value if collected on the first day of diarrhea. Tissues collected from a euthanized or dead calf should include:

Abomasum Fresh/chilled and formalin-fixed
Any other gross lesions Fresh/chilled and formalin-fixed
Colon Several partial loops, fresh/chilled

Three 1 cm pieces, formalin-fixed

Colon contents 10 ml fluid contents, fresh/chilled

Ileum Two or three 10-15 cm segments, fresh/chilled

Three 1 cm pieces, formalin-fixed

Jejunum Two or three 10-15 cm segments, fresh/chilled

Three 1 cm pieces, formalin-fixed Fresh/chilled and formalin-fixed Fresh/chilled and formalin-fixed Fresh/chilled and formalin-fixed

Rumen contents Fresh/chilled for pH

Spleen Fresh/chilled and formalin-fixed

Samples removed in the field are better than a whole dead animal submitted to the lab.

SAMPLING TECHNIQUES

Mesenteric lymph node

Liver

Rumen

- 1. Samples must be taken as soon after death as possible (within minutes).
- 2. Intestines do not need to be tied off at the ends.
- 3. Flush intestinal segments for histopathologic examination with formalin and drop in fixative. Or, gently open ends of 1 cm segments with scissors or forceps to expose mucosa as immersed. Do not split open.
- 4. Pool all formalin-fixed tissues from each calf in one bag; individual calves can be pooled or kept separate as desired. Package fresh intestines separately from other tissues and each calf represented in a separate bag. Chill fresh tissues before mailing. Do NOT freeze.

COMMON AGENTS DETECTED BY ROUTINE EXAM, CULTURE, PCR, AND/OR FECAL FLOTATION

Bacteria Salmonella spp., Clostridium perfringens; Mycobacterium avium subsp. paratuberculosis (Johne's disease) - PCR on fresh chilled feces;

histopathology and acid fast-stains on intestines and mesenteric lymph

nodes

Parasites Coccidia, GI nematodes

Viruses BVD virus, Bovine coronavirus: IHC on fixed ileum/colon, PCR on feces

- BVD mucosal disease diagnosis: Fixed ileum, spleen, mesenteric lymph nodes, skin, heart, lung, and ANY GROSS LESIONS for immunohistochemistry. Fresh/chilled spleen, lung, thymus, and mesenteric lymph node for PCR.
- Coccidiosis is a common cause of diarrhea in this age group. It is necessary to submit feces and/or colon to diagnose coccidiosis.



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BOVINE PNEUMONIA

Specimens to submit: Ante mortem samples from acutely affected calves should include:

Nasal swabs, Deep nasopharyngeal swabs, Use a long, Dacron-tipped swab that reaches deep into the nasal cavity Swabs to be used for virus PCR should penetrate the mucous layer to

Tracheal wash/lavage retrieve epithelial cells.

Submit separate swabs for bacterial culture and virus isolation in saline

or transport media. Do not freeze.

Swabs and/or lavage material can be submitted for PCR respiratory

panel (bacteria and viruses) as antemortem samples.

See <u>video</u> on sampling techniques.

Serum samples Acute and convalescent serum from 5-10 affected and 5-10 normal

calves. Hold acute samples and submit with convalescent.

Tissues collected from a euthanized or dead calf should include:

Ear notch Formalin-fixed

Heart/Liver Fresh and formalin-fixed

Lung Sample 3-4 areas of lung, generous portions of lesions and adjacent

unaffected lung, fresh

Four or more thin slices (1 cm) through affected and adjacent unaffected

lung, formalin-fixed

Lymph node/Thymus Fresh and formalin-fixed

Trachea Optional, if lesions are observed. Affected portion (10 cm) with larynx,

fresh. Several rings at edge of lesion, formalin-fixed.

SAMPLING SUGGESTIONS

- 1. Fresh tissues should be chilled before shipping. DO NOT FREEZE.
- 2. Samples for virus detection need to be taken from ACUTE animals at the onset of respiratory signs.
- 3. Swabs must be kept moist and cold before and during shipment.

COMMON AGENTS DETECTED BY ROUTINE EXAMINATION, CULTURE, AND/OR PCR

Bacteria Histophilus somni, Mannheimia haemolytica, Pasteurella multocida,

Mycoplasma bovis, Trueperella (Arcanobacterium) pyogenes

Viruses IBR, BRSV, BVD, BRCV, PI-3

- Acute lesions are most likely to hold active causative agents and are usually most prevalent at the
 interface of diseased/normal tissue. Chronic lesions in dependent tips or lobes may no longer
 contain primary pathogens.
- Nasal swabs may be of value to identify viruses if sampled in the early stages (exhibiting serous
 nasal discharge). Nasal swabs may also pick up resident bacterial flora but may be of value in
 certain acute cases. Ante mortem swabs from several affected calves can be pooled for PCR
 testing; and results can be compared with testing of swabs from unaffected calves.
- Tracheal washes submitted on ice may be used for both virus and bacteria identification.