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Finalized Report

Accession Number: 029815-22

Sampled: 02/12/2022 Received: 02/16/2022 Finalized: 02/21/2022

Accession Related

607-253-3900

Not Tested

Item		Result	Reference Interval
1	Valentina - Bovine Guernsey Female Serum	Not Tested For: Johne's PCR	
		Reason: We received serum only when feces	s is required for
2	Elliana - Bovine Guernsey Female Serum	Johne's PCR testing. jlt9 2/16/22 Not Tested For: Johne's PCR	
_	Emand Boving Guerrico Francis Guerri	Reason: We received serum only when feces	s is required for
		Johne's PCR testing. jlt9 2/16/22	
3	Valencia - Bovine Guernsey Female Serum	Not Tested For: Johne's PCR	
		Reason: We received serum only when feces	s is required for
		Johne's PCR testing. jlt9 2/16/22	
4	Diamond - Bovine Guernsey Female Serum	Not Tested For: Johne's PCR	
		Reason: We received serum only when feces	s is required for
		Johne's PCR testing. jlt9 2/16/22	
5	Victoria - Bovine Guernsey Female Serum	Not Tested For: Johne's PCR	
		Reason: We received serum only when feces	s is required for
		Johne's PCR testing. jlt9 2/16/22	

Serology/Immunology

Director Dr. Bettina Wagner - 607-253-3900

Johne's (MAP) Commercial ELISA

Item		Result	Reference Interval
1	Valentina - Bovine Guernsey Female Serum	Negative; 0.209	
2	Elliana - Bovine Guernsey Female Serum	Negative; 0.232	
3	Valencia - Bovine Guernsey Female Serum	Negative; 0.206	
4	Diamond - Bovine Guernsey Female Serum	Negative; 0.096	
5	Victoria - Bovine Guernsey Female Serum	Negative; 0.414	

Test Interpretations

Finalized Report Report Generation Date: 2/21/2022 12:35:34PM

Reference Interval

Finalized Report

Accession Number: 029815-22

Sarah Greenleaf

Johne's (MAP) Commercial ELISA

A Positive result indicates the presence of antibodies, suggesting that the animal is likely to be infected with M. paratuberculosis. Confirmation of infection with M. paratuberculosis by fecal culture or PCR is highly recommended. A Negative result on a single animal can only be interpreted when the paratuberculosis history and test results of the entire herd are known. Development of antibodies is a late stage event in the course of infection. Infected animals can be antibody negative for months to years.

Important Additional Notes: As with any biological test, this test may occasionally give a false positive or false negative result due to local conditions within the herd or region. A test should be interpreted in the context of all available clinical, historical and epidemiological information relevant to the animal(s) or herd under test. As of 4/03/2017 the Cutoff is:

< 0.70 Negative >=0.70 Positive

Item

This test is validated and USDA licensed for the detection of MAP specific antibodies in bovine serum or plasma and caprine serum .

Virology

Director Dr. Diego Diel - 607-253-3900

Result

Bovine Leukosis Virus (BLV) ELISA

1	Valentina - Bovine Guernsey Female Serum	Negative 14%	
2	Elliana - Bovine Guernsey Female Serum	Negative 0%	
3	Valencia - Bovine Guernsey Female Serum	Negative 0%	
4	Diamond - Bovine Guernsey Female Serum	Negative 0%	
5	Victoria - Bovine Guernsey Female Serum	Negative 0%	
Bovine	Virus Diarrhea Ag Capt ELISA		
Item		Result	Reference Interval
Item 1	Valentina - Bovine Guernsey Female Serum	Result Negative	Reference Interval
Item 1 2	Valentina - Bovine Guernsey Female Serum Elliana - Bovine Guernsey Female Serum		Reference Interval
Item 1 2 3		Negative	Reference Interval
1 2	Elliana - Bovine Guernsey Female Serum	Negative Negative	Reference Interval

Test Interpretations

Bovine Leukosis Virus (BLV) ELISA

Currently the AHDC uses ELISA kits supplied by two different manufacturers for the detection of antibodies against BLV. If you receive a % inhibition value in addition to your result the blocking ELISA was used. This value is calculated for each test serum and the test kit was licensed using a cut- off value of 45%. Above 45% = positive. Animals with values between 35% and 45% are considered suspects and should be retested 3-4 weeks after the initial sampling. Testing of animals less than 6 months of age on either ELISA could lead to a false positive test result due to passively acquired maternal antibodies