

TEST REPORT

NO.: TY-230725X

Date: 2023.12.19

1. **Product name** : FeLV Ag Test (feline leukemia virus Ag Test)
2. **Company** : Bioguard
3. **Test period** : 2023.07.25-2023.12.18
4. **Date issued** : 2023.12.19
5. **Test result**

Sample:

51 FeLV infected-positive specimens.

77 FeLV non-infected-negative specimens.

Procedure:

- a) Add 1 drop (20ul) sample into test cartridge by dropper.
- b) Add 4 drops assay buffer into test cartridge slowly by dropper.
- c) Results interpretation.
Positive: the appearance of control and test line.
Negative: the appearance of control line only.
Invalid: no appearance of control line.

	Real time RT-PCR (+)	Real time RT-PCR (-)	Total
FeLV Ag Test (+)	50	1	51
FeLV Ag Test (-)	1	76	77
Total	51	77	128

6. Statistical analysis

Sensitivity: 98.04%

Specificity: 98.70%

Accuracy: 98.44%

7. Condition

Temperature 25±10°C, Relative Humidity 65±15%

8. Reference

- a. Slov Vet Res, 2011, 48(2): 57-64.
- b. Virology. 2005 Feb 5;332(1):272-83.

The requirements of the specification are only for reference, veterinarians should consider other clinical information to make the definitive diagnosis in practice.

TEST REPORT

NO.: TG-231025X

Date: 2023.12.07

- 1. Product name** : FIV Ab Test (feline immunodeficiency virus Ab Test)
2. Company : Bioguard
3. Test period : 2023.10.25-2023.12.06
4. Date issued : 2023.12.07
5. Test result

Sample:

40 FIV infected-positive specimens.

120 FIV non-infected-negative specimens.

Procedure:

- a) Add 1 drop (20ul) sample into test cartridge by dropper.
b) Add 4 drops assay buffer into test cartridge slowly by dropper.
c) Results interpretation.
Positive: the appearance of control and test line.
Negative: the appearance of control line only.
Invalid: no appearance of control line.

	Western blot (+)	Western blot (-)	Total
FIV Ab Test (+)	39	4	43
FIV Ab Test (-)	1	116	117
Total	40	120	160

6. Statistical analysis

Sensitivity: 97.50%

Specificity: 96.67%

Accuracy: 96.88%

7. Condition

Temperature 25±10°C, Relative Humidity 65±15%

8. Reference

- a. Veterinary clinical pathology, 2010, 39(2): 210-214.
b. Pesq. Vet. Bras. vol.30 no.10 Rio de Janeiro Oct. 2010.

The requirements of the specification are only for reference, veterinarians should consider other clinical information to make the definitive diagnosis in practice.