

PK DEFICIENCY TEST REPORT

Provided Information:	Case: CAT147621
Name: BENGALISSIMO MAURIZIO	Date Received: 23-Oct-2023
Registration: SBT 022723 085	Report Issue Date: 30-Oct-2023
	Report ID: 8985-5995-9979-0040
	Verify report at www.vgl.ucdavis.edu/verify
DOB: 02/27/2023 Sex: Male Breed: Bengal Microchip: 528210006976463 Color: SEAL SPOTTED LYNX TABBY POINT	
Sire: CHAMPAGNE ON ICE VOM WEINBERG	Dam: MOONLIGHT ASHQUINDI
Reg: SBT 063016 068	Reg: SBT 091720 115
Microchip:	Microchip:

PYRUVATE KINASE DEFICIENCY RESULT

N/N

Interpretation

- N/N No copies of PK deficiency, cat is normal
- N/K 1 copy of PK deficiency, cat is normal but is a carrier
- K/K 2 copies of PK deficiency, cat is or will be affected. Severity of symptoms cannot be predicted*

PK DEFICIENCY TEST REPORT

Client/Owner/Agent Information: JÁNOS FABIAN HAJNAL STREET 17. 7627 PECS BARANYA HUNGARY	Case: CAT147621 Date Received: 23-Oct-2023 Report Issue Date: 30-Oct-2023 Report ID: 8985-5995-9979-0040 Verify report at www.vgl.ucdavis.edu/verify
Name: BENGALISSIMO MAURIZIO	

Additional Information

If testing for a disease or a disorder was performed and results indicate the animal is affected or at risk, we recommend contacting your veterinarian for further clinical evaluation and for additional information on disease and management.

For more detailed information on PK Deficiency test results, please visit our website at:
www.vgl.ucdavis.edu/services/pkdeficiency.php

Erythrocyte Pyruvate Kinase Deficiency (PK deficiency) is an inherited, autosomal recessive, hemolytic anemia. Breedings between carriers will be expected to produce 25% affected kittens. Go to our website for a list of breeds at risk of PK deficiency due to a significant frequency of the mutation.

For terms and conditions of testing, please see www.vgl.ucdavis.edu/about/terms-and-conditions

Results are determined using PCR-based methods. The results relate only to the sample tested as identified by the submitter (for example, identity and/or breed).

Report authorized by Dr. Rebecca Bellone, VGL Director

Veterinary Genetics Laboratory · University of California Davis · One Shields Ave · Davis, CA 95616
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BENGAL PRA TEST REPORT

Provided Information:		Case:	CAT147621
Name:	BENGALISSIMO MAURIZIO	Date Received:	23-Oct-2023
Registration:	SBT 022723 085	Report Issue Date:	30-Oct-2023
		Report ID:	7950-7160-7968-3015
Verify report at www.vgl.ucdavis.edu/verify			
DOB: 02/27/2023 Sex: Male Breed: Bengal Microchip: 528210006976463 Color: SEAL SPOTTED LYNX TABBY POINT			
Sire:	CHAMPAGNE ON ICE VOM WEINBERG	Dam:	MOONLIGHT ASHQUINDI
Reg:	SBT 063016 068	Reg:	SBT 091720 115
Microchip:		Microchip:	

BENGAL PRA RESULT

N/N

Interpretation

- N/N Normal - no copies of the PRA-b mutation.
- N/PRA Carrier - 1 copy of the PRA-b mutation; vision will be normal.
- PRA/PRA Affected - 2 copies of the PRA-b mutation; cat will develop clinical signs of Bengal PRA.

BENGAL PRA TEST REPORT

Client/Owner/Agent Information: JÁNOS FABIAN HAJNAL STREET 17. 7627 PECS BARANYA HUNGARY	Case: CAT147621 Date Received: 23-Oct-2023 Report Issue Date: 30-Oct-2023 Report ID: 7950-7160-7968-3015 Verify report at www.vgl.ucdavis.edu/verify
Name: BENGALISSIMO MAURIZIO	

Additional Information

If testing for a disease or a disorder was performed and results indicate the animal is affected or at risk, we recommend contacting your veterinarian for further clinical evaluation and for additional information on disease and management.

For more detailed information on Bengal PRA test results, please visit our website at:
www.vgl.ucdavis.edu/services/cat/BengalPRA.php

For terms and conditions of testing, please see www.vgl.ucdavis.edu/about/terms-and-conditions

Results are determined using PCR-based methods. The results relate only to the sample tested as identified by the submitter (for example, identity and/or breed).

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