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PK DEFICIENCY AND IDENTITY MARKER REPORT

GUILLAUME GUY 2114 DEZERY MONTREAL HIW 2S2 CANADA	Case: CAT81024 Date Received: 19-Jan-2016 Print Date: 22-Jan-2016 Report ID: 5406-4908-7876-8071 Verify report at www.vgl.ucdavis.edu/myvgl/verify.html
Cat: ASIAFAUVE CAMEO OF BENGALLEOPARD DOB: 02/27/2015 Sex: Female Breed: Bengal Microchip: 967000009682876 Color: Brown	Reg: SBT022714 038
Sire: LEOPAWS DAKOTA Dam: ASAIFAUFE SASHA	Reg: SBT 030911 010 Reg: SBT 072411 028

PYRUVATE KINASE DEFICIENCY TEST RESULT

N/N

Result Codes:

- 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*

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: www.vgl.ucdavis.edu/services/pkdeficiency.php

*If your cat is diagnosed as homozygous for PK deficiency, we recommend that you contact your veterinarian for information on disease progression and management.

IDENTITY MARKERS

LOCUS	TYPE	LOCUS	TYPE
<i>FCA075</i>	R	<i>FCA220</i>	L
<i>FCA223</i>	G	<i>FCA678</i>	JM
<i>FCA698</i>	WC		