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**PK DEFICIENCY AND IDENTITY MARKER REPORT**

<p><b>ERIKA KARSEMEYER</b>                  3400 AVENUE G                  YUCAIPA, CA 92399</p>	<p><b>Case: CAT82526</b>  <b>Date Received: 09-Mar-2016</b>                  Print Date: 10-Mar-2016                  Report ID: 8397-6449-7565-9140                  Verify report at <a href="http://www.vgl.ucdavis.edu/myvgl/verify.html">www.vgl.ucdavis.edu/myvgl/verify.html</a></p>
<p><b>Cat: SHANGRIPAW SEBASTIAN OF WB</b> <b>Reg: SBT 042715 078</b>                  DOB: 04/27/2015 Sex: Male Breed: Bengal Color: BROWN (BLACK) SPOTTED TABBY</p>	
<p>Sire: PURRECIOUSPOTS JSTFYD OF SHANGRIPAW <b>Reg: SBT 031313 046</b>                  Dam: SHANGRIPAW LEGACEE <b>Reg: SBT 021413 056</b></p>	

**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](http://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
FCA075	PR	FCA220	L
FCA223	UV	FCA678	M
FCA698	Uc		