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## PK DEFICIENCY AND IDENTITY MARKER REPORT

<p><b>OLGA DREIDT</b> [REDACTED] GERMANY</p>	<p><b>Case: CAT71350</b> <b>Date Received: 15-Dec-2014</b> <b>Print Date: 16-Dec-2014</b> <b>Report ID: 1115-1870-3053-1155</b> 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: RUSLANE RASPUTIN</b> <b>DOB: 06/25/2013</b> <b>Sex: Male</b> <b>Breed: Bengal</b> <b>Microchip: 276093400471745</b> <b>Color: Brown Spotted Tabby</b></p>	<p><b>Reg: SBT 062513 022</b></p>
<p><b>Sire: GOGEEES OLE</b> <b>Dam: RUSLANE RASCHEL</b></p>	<p><b>Reg: SBT 120909 002</b> <b>Reg: SBT 030712 070</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	P	FCA220	KL
FCA223	GM	FCA678	KN
FCA698	U		