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## PK DEFICIENCY AND IDENTITY MARKER REPORT

<p>GREG STAPLES 1014 SNIDER'S BAY ROAD GRAVENHURST ONTARIO P1P 1R2 CANADA</p>	<p><b>Case:</b> <b>CAT87523</b> <b>Date Received:</b> 08-Aug-2016 <b>Print Date:</b> 09-Aug-2016 <b>Report ID:</b> 1589-6810-4464-6015 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:</b> RAYDUSOLEIL STACKER PENTECOST OF MINOOS <b>DOB:</b> 01/29/2016 <b>Sex:</b> Male <b>Breed:</b> Maine Coon <b>Color:</b> Brown Classic Tabby and White</p>	<p><b>Reg:</b> SBT 012916 014</p>
<p><b>Sire:</b> HONOR VILLA PARK PL OF RAYDUSOLEIL <b>Dam:</b> TRISKEL AMY OF RAYDUSOLEIL</p>	<p><b>Reg:</b> SBT 062214 056 <b>Reg:</b> SBT 110412 033</p>

## PYRUVATE KINASE DEFICIENCY TEST RESULT

N/K

### 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	S	FCA220	L
FCA223	GW	FCA678	J
FCA698	S		