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PK DEFICIENCY AND IDENTITY MARKER REPORT

<p>SARA VEGA</p> <p>SANTIAGO, 8330269 CHILE</p>	<p>Case: CAT77070</p> <p>Date Received: 03-Aug-2015</p> <p>Print Date: 04-Aug-2015 Report ID: 0183-9401-8275-4120</p> <p>Verify report at www.vgl.ucdavis.edu/myvgl/verify.html</p>
<p>Cat: SABLE KLAN RHEA OF PERBAST Reg: SBV 112014 010</p> <p>DOB: 11/20/2014 Sex: Female Breed: Somali Microchip: Color: Blue</p>	
<p>Sire: UTRILLO VITTORIOSO DI GENS RUBRA Reg:</p> <p>Dam: DANAKIL'S BLUE BELLE Reg:</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

*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>	S	<i>FCA220</i>	L
<i>FCA223</i>	N	<i>FCA678</i>	J
<i>FCA698</i>	Na		