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PK DEFICIENCY AND IDENTITY MARKER REPORT

ANDREA A. S. VAN ELMPT VAALRIVIERSTRAAT 10A 1091 PD AMSTERDAM THE NETHERLANDS NETHERLANDS	Case: CAT64149 Date Received: 01-Apr-2014 Print Date: 04-Apr-2014 Report ID: 3524-3489-8036-7195 Verify report at www.vgl.ucdavis.edu/myvgl/verify.html
Cat: DETREVANDE'S KEEPING UP APPEARANCES DOB: 07/17/2013 Sex: Female Breed: Abyssinian Microchip: 528246002387971	Reg: (NL) MU.LO13.ABY.035.1
Sire: HELIOS DE SHAKIRI Dam: DETREVANDE'S CRAZY LIL' THING	Reg: LOOF 2012.12428 Reg: MU.L09.ABY.013.2

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>	QR	<i>FCA220</i>	L
<i>FCA223</i>	MN	<i>FCA678</i>	K
<i>FCA698</i>	T		