



VETERINARY GENETICS LABORATORY  
SCHOOL OF VETERINARY MEDICINE  
ONE SHIELDS AVENUE  
DAVIS, CALIFORNIA 95616-8744

TELEPHONE: (530) 752-2211  
FAX: (530) 752-3556

## PROGRESSIVE RETINAL ATROPHY (PRA) REPORT

HEIDE PETERSEN MAGLEMOSE 18 4960 HOLEBY DENMARK	<b>Case:</b> <b>CAT75321</b> <b>Date Received:</b> 04-Jun-2015 <b>Print Date:</b> 08-Jun-2015 <b>Report ID:</b> 2226-2663-8649-8022 Verify report at <a href="http://www.vgl.ucdavis.edu/myvgl/verify.html">www.vgl.ucdavis.edu/myvgl/verify.html</a>
<b>Cat:</b> <b>S*SPLENDID'S ISADORA</b> <b>DOB:</b> 12/06/2014 <b>Sex:</b> Female <b>Breed:</b> Siamese <b>Microchip:</b> 752098100728450 <b>Color:</b> seal-point	<b>Reg:</b>
<b>Sire:</b> *S SPLENDID'S CONCRETE <b>Dam:</b> S*YZINGS JAVIZZT	<b>Reg:</b> SVERAK LO 264112 <b>Reg:</b> SVERAK LO 306737

PRA-CEP290 Result	PRA-CRX Result
N/N	

### Result Codes for PRA-CEP290:

N/N	Normal, cat does not have rdAc mutation*
N/rdAc	Carrier, cat has one copy of rdAc mutation. Breeding between carriers is expected to produce 25% affected kittens
rdAc/rdAc	Affected

rdAc is a progressive retinal atrophy that causes late-onset blindness.\* Affected cats are born with normal vision, show retinal degeneration at about 7 months and are blind by age 3-5 years. The condition is inherited as an autosomal recessive, and there is no treatment. \* This test only detects the mutation in the CEP290 gene known to cause PRA-rdAc in Abyssinian, Somali, Ocicat, Siamese and related breeds, American Curl, American Wirehair, Bengal, Cornish Rex, Munchkin, Singapura and Tonkinese.

### Result Codes for PRA-CRX

N/N	Normal, cat does not have Rdy mutation**
N/Rdy	Affected, cat has one copy of the Rdy mutation. This cat will produce affected kittens 50% of the time when bred to a normal cat, or 75% of the time when bred to another cat with one copy of the Rdy mutation.
Rdy/Rdy	Affected, cat will always produce affected kittens.

Rdy is an early-onset retinopathy caused by a defective protein that is critical for eye development.\*\* Affected kittens display dilated pupils and sluggish pupillary reflexes around 2 weeks of age and often become blind by about 7 weeks of age. The condition is inherited as an autosomal dominant trait, and there is no treatment. \*\* This test only detects the mutation in the CRX gene known to cause PRA-Rdy in Abyssinian and Somali breeds.