

CERTIFICATE OF EYE EXAMINATION

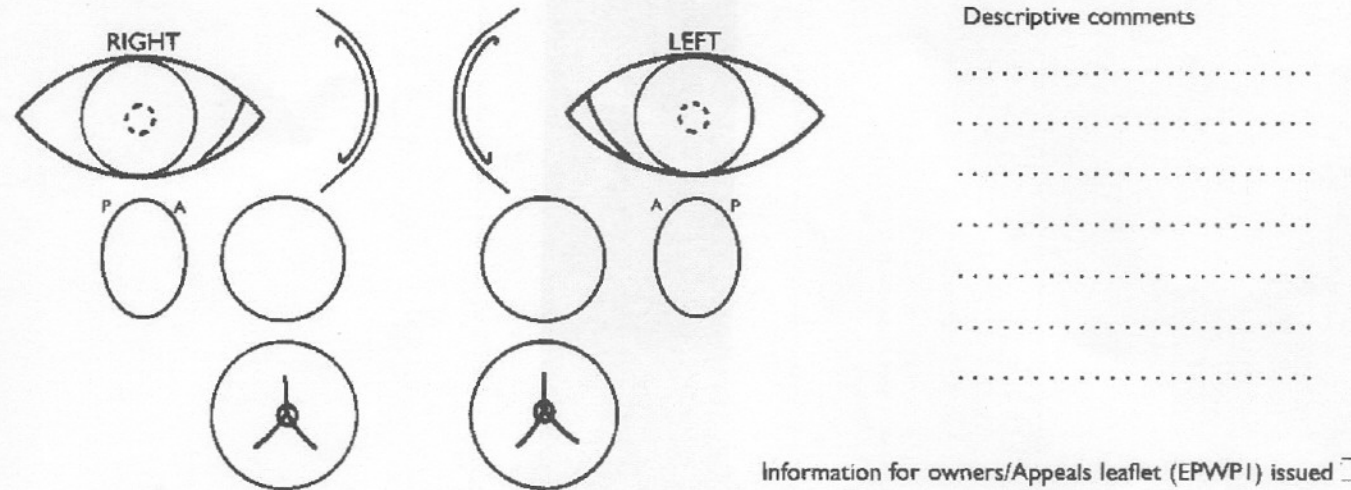
KC/ISDS registered name SANDBRAE BOLT from The BLUE Panellist's ref no MCJ10 969
 Breed CLCS M3 Glen Colour Ruby Registered no AS 03699701
 Owner's name M3 Glen Owner's veterinary surgeon CLYDE VET GP
 Owner's address 22 Lochaber Rd STRATHGON Date of birth 15/7/08 Sex M F
 Owner's telephone number 01352-522531 LANARK
 Previous examination: No Yes Date of last examination Microchip/tattoo No 956 000 001 985 017

I hereby declare that the dog submitted for examination under the BVA/KC/ISDS Eye Scheme is the one described above. I agree that the registration document should be stamped with the date of this examination and that the information obtained may be made available for research purposes and may be published (deletion of these statements invalidates the certificate). Any appeal against the results specified below must be made to the BVA (for details see leaflet EPWPI)

Date 16/10/10 Signed Alison Glen Owner/Agent

EXAMINATION OF EYE AND ADNEXA

Mydriatic: Ophthalmoscopy: Direct Indirect Biomicroscopy: Other
 Parts examined: Adnexa Cornea Iris Lens Vitreous Fundus
 Clinically unaffected
 Clinically affected



Descriptive comments

Information for owners/Appeals leaflet (EPWPI) issued
 I confirm that the scanned microchip/tattoo number matches the number on this certificate

INHERITED EYE DISEASE STATUS

This section applies only to those conditions in the breeds specified in Schedule A of the Procedure Notes current on the day of examination. These results will be sent to the Kennel Club and/or ISDS as appropriate.

CONGENITAL	CLINICALLY UNAFFECTED	CLINICALLY AFFECTED	NON-CONGENITAL	CLINICALLY UNAFFECTED	CLINICALLY AFFECTED
(CEA) Collie eye anomaly: - choroidal hypoplasia - coloboma	<input type="checkbox"/>	<input type="checkbox"/>	(GPRA) Generalised progressive retinal atrophy:	<input type="checkbox"/>	<input type="checkbox"/>
(MRD) Multifocal retinal dysplasia: <input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	(CPRA) Central progressive retinal atrophy:	<input type="checkbox"/>	<input type="checkbox"/>
(TRD) Total retinal dysplasia: <input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	(HC) Hereditary cataract: <input checked="" type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
(CHC) Congenital hereditary cataract: <input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>	(PLL) Primary lens luxation: <input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>
(PHPV) Persistent hyperplastic primary vitreous: <input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>			
(G) Goniodysgenesis: <input type="checkbox"/>	<input type="checkbox"/>	<input type="checkbox"/>			

The age of onset of non-congenital inherited eye disease varies in different breeds and between individual dogs. It is therefore important to follow any advice given at the time of this examination with regard to the necessity for and frequency of eye examination under the Scheme.

'Clinically affected' signifies that there is visible evidence of the inherited eye disease(s) specified, whereas 'clinically unaffected' signifies that there is no such evidence.

I have today examined the above animal under the BVA/KC/ISDS Eye Scheme with the results as shown
 Signed Alison Glen Name MCSAVIDSON Date 16/10/10