# August 2024



Regency Reds Amos WALA 00044556



British Veterinary Association/Kennel Club/International Sheep Dog Society (BVA/KC/ISDS) CANINE HEALTH SCHEMES EYE EXAMINATION CERTIFICATE

| et name AM03   | KC no.  | M  | ticrochip no. 900   | 20000075  | 2666                  |
|--|---|--|---|---|-----------------------|
| C registered name  | MA  |  | Date of previous exar   | mination  | -                     |
| eed ANSTRACIAN   | LASRADODIE COLOUR   | 26.0   |   | Date of birth 6/4/2   | a                     |
| wner's name and address  |   |  | Sex M L F L   | Date of birth   |                       |
|  |   |  |   |   |                       |
| wner's telephone number  | Lang Vela a. A  | Owner's email address  | 1.12  |   |                       |
| t's name and address //  | man was sv. in  | nes ta CASIBO  | VENE  |   |                       |
| t's telephone number   |   | Vet's email address  |   |   |                       |
| ereby declare that the dog submitte<br>ade available for research purposes   | ed for examination under the BVA/<br>s and may be published. Any appe   | KC/ISDS Canine Health Scheme is the all against the results specified below  | he one described above<br>w must be made to the   | and that the information obtaine<br>BVA (for details see EPWP1).  | ed may be             |
| inderstand and agree that the use o  | of a mydriatic agent  | AVT INE is necessary to t  |   | amination of the eye and that a lo  | ocal anaesthetic      |
| inderstand that the personal information   | Il be used where gonioscopy is rec<br>on provided in this form will be used t   | quired.<br>to administer the eve examination servi   | ice and will be retained fo   | or 7 years for accounting purposes  |                       |
| stem. My personal information may be   | a used from time to time to provide m   | e with relevant information relating to 0  | CHS services or for other   | lawful reasons.   |                       |
| gnature of Owner/Agent   | na  |  | <u></u>   | Date \$ 18/24   |                       |
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| vdriatio Ophthalmoscopy Direct   |   | y Gonioscopy Tonometry   |   |   |                       |
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| is section applies to the known inherit<br>CONGENITAL/NEONATAL   | INF<br>ted ocular conditions specified in the<br>CLINICALLY CLINIC/<br>UNAFFECTED AFFEC   | HERITED EYE DISEASE STAT<br>Procedure Notes. These results will be<br>ALLY N<br>TED (HC) Hereditary ca<br>(PLL) Primary lens   | US<br>e sent to the KC and/or IS<br>ON-CONGENITAL<br>taract<br>luxation   | he scanned microchip number<br>umber on the certificate<br>owners/Appeals leaflet (EPWP1) i<br>DS as appropriate.<br>CLINICALLY   | clinically            |
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| is section applies to the known inherit<br>CONGENITAL/NEONATAL<br>EA) Collie eye anomaly<br>– Choroidal hypoplasia<br>– Coloboma<br>RD) Multifocal retinal dysplasia<br>RD) Total retinal dysplasia<br>RD) Total retinal dysplasia<br>RD) Total retinal dysplasia<br>RD) Total retinal dysplasia<br>RD) Persistent hyperplastic primary of<br>A) Pectinate ligament abnormality<br>rade 0 1 2 3 Besult<br>R<br>L<br>Goniosc<br>0 = norm<br>nically affected with ocular conditions<br>stichiasis<br>topico ilia<br>chiasis<br>tropion<br>tropion<br>mbined entropion/ectropion   | INF<br>ted ocular conditions specified in the<br>CLINICALLY CLINIC/<br>UNAFFECTED AFFEC<br>UNAFFECTED AFFEC<br>UNAFFECTED AFFEC<br>COPY Grading Result:<br>mail, 1 = mildly affected, 2 = moderate<br>enot currently specified in the Proceed<br>Persistent pupillary membrane<br>Ocular Melanosis<br>Pectinate ligament abnormality<br>Lens luxation<br>Anterior Capsular Cataract<br>Anterior Cortical Cataract | HERITED EYE DISEASE STAT<br>Procedure Notes. These results will be<br>ALLY N<br>(HC) Hereditary ca<br>(PLL) Primary lens I<br>(POAG) Primary lens I<br>(POAG) Primary lens I<br>(POAG) Primary lens I<br>(POAG) Primary lens I<br>(PA) Progressive m<br>(IOP) Intraocular<br>(PRA) Progressive m<br>(IOP) Retinal pigme<br>'Clinically affected's<br>the inherited disease(s<br>unaffected' signifies the<br>by affected, 3 = severely affected.<br>ure Notes.<br>Posterior Cortical Catarac<br>Posterior Polar Subcapsu<br>Posterior Polar Subcapsu<br>Posterior Capsular Catarac<br>PHPV<br>Optic nerve hypoplasia<br>Posterior segment colobo | Li confirm that t<br>matches the m<br>Information for<br>US<br>esent to the KC and/or IS<br>ION-CONGENITAL<br>taract<br>luxation<br>angle glaucoma<br>pressure R mmHg L<br>etinal atrophy<br>ent epithelial dystrophy<br>ignifies that there is evide<br>s) specified, whereas 'Cli<br>hat there is no such evide                               | he scanned microchip number<br>umber on the certificate<br>owners/Appeals leaflet (EPWP1) i<br>DS as appropriate.<br>CLINICALLY<br>UNAFFECTED<br>mmHg<br>mmHg<br>mce of<br>nically<br>ance.<br>GPRA-like appearance<br>RPED-like appearance | clinically            |
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| is section applies to the known inherit<br>CONGENITAL/NEONATAL<br>EA) Collie eye anomaly<br>– Choroidal hypoplasia<br>– Coloboma<br>RD) Multifocal retinal dysplasia<br>RD) Total retinal dysplasia<br>RD) T | INF<br>ted ocular conditions specified in the<br>CLINICALLY CLINIC/<br>UNAFFECTED AFFEC<br>introduction of the specified in the Proceed<br>Persistent pupillary membrane<br>Ocular Melanosis<br>Pectinate ligament abnormality<br>Lens luxation<br>Anterior Capsular Cataract<br>Anterior Cortical Cataract<br>Perinuclear Cataract<br>Nuclear Cataract   | HERITED EYE DISEASE STAT<br>Procedure Notes. These results will be<br>ALLY N<br>(HC) Hereditary ca<br>(PLL) Primary lens I<br>(POAG) Primary open<br>(IOP) Intraocular<br>(PRA) Progressive n<br>(RPED) Retinal pigme<br>'Clinically affected's<br>the inherited disease(s<br>unaffected' signifies th<br>Ity affected, 3 = severely affected.<br>Ure Notes.<br>Posterior Cortical Catarac<br>Posterior Capsular Catara<br>PHPV<br>Optic nerve hypoplasia<br>Posterior segment colobo<br>Choroidal hypoplasia<br>MRD-like appearance   | Li confirm that t<br>matches the m<br>Information for<br>US<br>esent to the KC and/or IS<br>ON-CONGENITAL<br>taract<br>luxation<br>angle glaucoma<br>pressure R mmHg L<br>etinal atrophy<br>ent epithelial dystrophy<br>ignifies that there is evide<br>s) specified, whereas "Cli<br>hat there is no such evide<br>that there is no such evide | he scanned microchip number<br>umber on the certificate<br>owners/Appeals leaflet (EPWP1) i<br>DS as appropriate.<br>CLINICALLY<br>UNAFFECTED<br>mmHg<br>mmHg<br>mce of<br>nically<br>ance.<br>GPRA-like appearance<br>RPED-like appearance | clinically            |
| is section applies to the known inherit<br>CONGENITAL/NEONATAL<br>EA) Collie eye anomaly<br>– Choroidal hypoplasia<br>– Coloboma<br>RD) Multifocal retinal dysplasia<br>RD) Total retinal dysplasia<br>HOY Dersistent hyperplastic primary of<br>A) Pectinate ligament abnormality<br>rade 0 1 2 3 Besult<br>R<br>D 1 2 3 Besult<br>Goniosc<br>0 = norm<br>nically affected with ocular conditions<br>tichiasis<br>topic cilia<br>chiasis<br>tropion<br>mbined entropion/ectropion<br>Iti-ocular defects<br>meal lipid deposition  | INF<br>ted ocular conditions specified in the<br>CLINICALLY CLINIC/<br>UNAFFECTED AFFEC<br>introduction of the specified in the Proceed<br>Persistent pupillary membrane<br>Ocular Melanosis<br>Pectinate ligament abnormality<br>Lens luxation<br>Anterior Capsular Cataract<br>Anterior Cortical Cataract<br>Perinuclear Cataract<br>Nuclear Cataract   | HERITED EYE DISEASE STAT<br>Procedure Notes. These results will be<br>ALLY N<br>(HC) Hereditary ca<br>(PLL) Primary lens I<br>(POAG) Primary open<br>(IOP) Intraocular<br>(PRA) Progressive m<br>(RPED) Retinal pigme<br>'Clinically affected's<br>the inherited disease(s<br>unaffected' signifies th<br>by affected, 3 = severely affected.<br>ure Notes.<br>Posterior Cortical Catarac<br>Posterior Polar Subcapsu<br>Posterior Capsular Catarac<br>PHPV<br>Optic nerve hypoplasia<br>Posterior segment coloboc<br>Choroidal hypoplasia   | Li confirm that t<br>matches the m<br>Information for<br>US<br>e sent to the KC and/or IS<br>ON-CONGENITAL<br>taract<br>luxation<br>angle glaucoma<br>pressure R mmHg L<br>etinal atrophy<br>int epithelial dystrophy<br>ignifies that there is evide<br>s) specified, whereas 'CH<br>hat there is no such evide                                | he scanned microchip number<br>umber on the certificate<br>owners/Appeals leaflet (EPWP1) i<br>DS as appropriate.<br>CLINICALLY<br>UNAFFECTED<br>mmHg<br>mmHg<br>mce of<br>nically<br>ance.<br>GPRA-like appearance<br>RPED-like appearance | clinically            |

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| BRITISH VETER  | INARY ASSO  | OCIATION  | KENNEL   | CLUB H   | IP DYSPLASIA SCHEME   |  |  |
|--|---|---|--|--|---|--|--|
| To: British Veterinary Associatior<br>7 Mansfield Street, London V<br>Telephone: 020 7908 6380   | V1G 9NQ   | 21 - 2  | 08890  | 0  | THE ORIGINAL OF THIS<br>CERTIFICATE IS GREEN  |  |  |
| Section A - TO BE COMPLETED  | BY OWNER/AGENT  |   |  |  | at Real trans   |  |  |
| KC Registered Name Rest<br>Breed Australia<br>Name of owner Dest   | Las<br>Cont   | a Amo   | Sex M<br>Address   |  | Date of hirth 10 10Lc 12620   |  |  |
| <ul> <li>(d) I give permission for the result</li> <li>(e) I give permission for the result</li> <li>(f) I understand that once the suprocess</li> <li>(g) I understand that the personal</li> <li>7 years for accounting purpose</li> </ul> | ION OF ANY OF<br>rect and relate to<br>year old and has<br>f the certificate to<br>ts of the examina<br>ts to be published<br>bmission has bee<br>information provi | THESE ITEM<br>the dog subm<br>s not previousl<br>b be sent to the<br>tion to be use<br>d and included<br>an received by<br>ided as part of<br>ic system. My | hitted for radiogn<br>by been scored is<br>e geneticist retained<br>at a future da<br>d on the relevan<br>the Canine He<br>the scheme is o | ES THIS C<br>raphic exa<br>under this<br>ined by the<br>te for the<br>the for the<br>the KC docu-<br>alth Scher<br>only used | ERTIFICATE)<br>imination<br>Scheme<br>e breed society or other representative body<br>purpose of statistical research   |  |  |
| Section B - TO BE COMPLETED B  |   |   |  |  |   |  |  |
| (Section A must be completed in full before  | completing Section B)   | ERINARY SURGE   | EON  |  |   |  |  |
| Microchip/Tattee no. 9 00  | 2000  | 200   | 7521   | GGC  | Microchip/T <del>attoo</del> confirmed  |  |  |
| I certify that the radiograph relating<br>and in conformity with the provisio<br>Veterinary surgeon submitting r<br>Address  | ns of the Hip Dys<br>radiograph (BLO<br>Rose D  | plasia Schem  | e Procedure No<br>S)<br>1 Road   | otes.  | S. Meacock MRCVS  |  |  |
| Veterinary Surgeon's Signature   | har   | S. 1  | land   | FRAD   | Post code / 0 1 MAY 2021  |  |  |
|  | A DAY OF THE REAL PROPERTY OF THE REAL PROPERTY OF  |   | The second se                            |  | ovo bate  |  |  |
| Please submit the correct fe   | e for the radiogr   | aph to be pro   | cessed (chequ  | ues payal  | ble to BVA.) For current fees contact BVA   |  |  |
| Section C - TO BE COMPLETED E  | CERT  |   | E OF SC  | ORIN   | IG .  |  |  |
| HIP JOINT  | Score Range   | Right   | Left   |  |   |  |  |
| Norberg angle<br>Subluxation   | 0-6   | 2   | 0  | Г  |   |  |  |
| Cranial acetabular edge  | 0-6   | 2   | 2  |  | B The scores represent the opinion of the   |  |  |
| Dorsal acetabular edge   | 0-6   | 2   | ~  |  | BVA appointed scrutineers for the radiograph<br>submitted. The lower the score, the less<br>evidence of hip dysplasia present. Please<br>consult the current procedure notes and<br>breed mean score sheet for relevant details |  |  |
| Cranial effective acetabular rim   | 0-6   |   |  | e  |   |  |  |
| Acetabular fossa   | 0-6   |   |  |  |   |  |  |
| Caudal acetabular edge   | 0-5   |   |  |  | available from BVA)   |  |  |
| Femoral head/neck exostosis  | 0-5   |   |  | Ľ  |   |  |  |
| Femoral head recontouring  | 0-6   |   |  |  |   |  |  |
|  | 53 per column)  | 5   | 2 0  | 2  | Total score (max possible 106)  |  |  |
| WE HEREBY CERTIFY that the so<br>was produced using the scoring or<br>Signed   | core of the radiog  | Kennel Club H   | d for the dog id<br>lip Dysplasia S<br>S Signed  | entified at  |   |  |  |
| 18   |   |   |  |  | 03/18   |  |  |

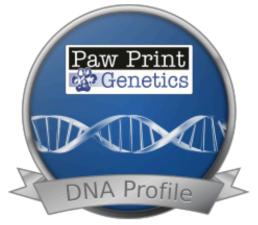
| BRITISH VETERINARY ASSOCI  | ATION/KENNEL CLUB E  | LBOW DYSPLASIA SCHEME   |
|--|--|---|
| To: British Veterinary Association<br>7 Mansfield Street, London W1G 9NQ<br>Telephone: 020 7908 6380   | 21-208800  | THE ORIGINAL OF THIS<br>CERTIFICATE IS GOLD   |
| Section A - TO BE COMPLETED BY OWNER/AGENT   | KC Registered Number   | not rein meral  |
| KC Registered Name<br>Breed Australian Labrad<br>Name of owner Debbic Comp   | sed Anos<br>codle sex M<br>Address   | Date of birth 10 104 1 3020   |
|  |  | Post code   |
| Sire:<br>Regency sid Rafe  | Dam:<br>Negenci  | ey red cherry   |
| <ul> <li>I hereby declare that (NB: DELETION OF ANY OF T</li> <li>(a) The particulars above are correct and relate to th</li> <li>(b) This dog is a minimum of one year old and has r</li> <li>(c) I give permission for a copy of the certificate to b</li> <li>(d) I give permission for the results of the examinati</li> <li>(e) I give permission for the results to be published</li> <li>(f) I understand that once the submission has been process</li> <li>(g) I understand that the personal information provid</li> <li>7 years for accounting purposes on an electronic</li> </ul> | he dog submitted for radiographic e<br>not previously been graded under t<br>be sent to the geneticist retained by<br>on to be used at a future date for th<br>and included on the relevant KC do<br>received by the Canine Health Sc<br>ed as part of the scheme is only us | examination<br>his Scheme<br>y the breed society or other representative body<br>he purpose of statistical research<br>ocuments<br>hemes office it cannot be withdrawn from the |
| Owner's/Agent's signature DACOCA   | bord   | Date / 01 MAY 2021  |
| Section B – TO BE COMPLETED BY SUBMITTING VETER<br>(Section A must be completed in full before completing Section B)   | RINARY SURGEON   |   |
| Microchip/Tettoo no. 9 0 0 2 0 0 0   | 00075266   | Microchip/T <del>atto</del> e confirmed   |
| I certify that the radiographs relating to the dog ident<br>and in conformity with the provisions of the Elbow Dy  |  | wing date/  |
| Address  | CK CAPITALS)<br>Pene, Shipton Road<br>Pr Wychwood OX7 6JT  | er S Meecock MRCVS NJD  |
| 1  | M. A   | Post code / 0 1 MAY, 2021   |
| Please submit the correct fee for the radiogram  |  |   |
|  | ons to be processed (cheques pa  | yable to BVA.) For current lees contact BVA   |
| Section C - TO BE COMPLETED BY SCRUTINEERS   |  |   |
| CERT   | FICATE OF GRAD   | ING   |
| RIGHT     LEFT       GRADE<br>(range 0-3)     O  | view of each elb   | re based on a flexed lateral and neutral lateral<br>bow and represent the opinion of the BVA<br>eers for the radiographs submitted. The lower                                   |
| OVERALL GRADE (max possible 3)   | the grade, the less<br>overall grade give<br>the highest grade   | ss evidence of elbow dysplasia present. The<br>n for both elbows is that given to the elbow with<br>e. Please consult the current procedure notes<br>s (available from BVA)     |
| WE HEREBY CERTIFY that the grade of the radiogram was produced using the grading protection of the BVA/k   | aphs submitted for the dog identifie   | ed above<br>me Date 2 3 JUN 2021  |
| Signed   | F/MRCVS Signed   | Diversion Date Date Date Date Date Date Date Date   |



### Paw Print DNA Profiling<sup>™</sup> Certificate

Call Name: Registered Name: Breed: Sex: DOB: Amos Regency Red Amos Australian Labradoodle Male April 2020 Laboratory #: Registration #: Certificate Date:

210450 -Jan. 21, 2021



This certificate displays a graphical representation of your dog's unique DNA profile



CKM

Christina J Ramirez, PhD, DVM, DACVP Medical Director

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Robert D. Westra, MS, DVM Assistant Medical Director

Paw Print Genetics<sup>®</sup> performed testing on the dog(s) listed on this certificate. Because this test is a DNA-based method, rare genomic variations may occur producing false results. If you think these results are in error, please contact the laboratory immediately for further evaluation. In the event of a valid dispute of results claim, Paw Print Genetics will do its best to resolve such a claim to the customer's satisfaction. If no resolution is possible after investigation by Paw Print Genetics with the cooperation of the customer, the extent of the customer's sole remedy is a refund of the fee paid. In no event shall Paw Print Genetics be liable for indirect, consequential or incidental damages of any kind. Any claim must be asserted within 60 days of the report of the test results. Genetic counseling is available at Paw Print Genetics.



## **Coat Color and Trait Certificate**

| Call Name:       | Amos                   | Laboratory #:     | 210450        |
|------------------|------------------------|-------------------|---------------|
| Registered Name: | Regency Red Amos       | Registration #:   | -             |
| Breed:           | Australian Labradoodle | Certificate Date: | Jan. 21, 2021 |
| Sex:             | Male                   |                   |               |
| DOB:             | April 2020             |                   |               |

#### This canine's DNA showed the following genotype(s):

| Coat Color/Trait Test                       | Gene   | Genotype                       | Interpretation  |
|---|--------|--------------------------------|---|
| coat color/ mait rest                       | Gene   | denotype                       | interpretation  |
| A Locus (Agouti)                            | ASIP   | a <sup>t</sup> /a <sup>t</sup> | Tricolor, black and tan                                       |
| B Locus (Brown)                             | TYRP1  | B/b                            | Black coat, nose and foot pads (carries one<br>copy of brown) |
| D Locus (Dilute)                            | MLPH   | D/D                            | Non-dilute (does not carry dilute)                            |
| E Locus (Yellow/Red)                        | MC1R   | e/e                            | Yellow/red  |
| IC Locus (Improper<br>Coat/Furnishings)     | RSPO2  | F/F                            | Furnishings   |
| K Locus (Dominant Black)                    | CBD103 | K <sup>B</sup> /K <sup>B</sup> | No agouti expression allowed                                  |
| S Locus (White Spotting, Parti, or Piebald) | MITF   | S/s <sup>p</sup>               | Limited white spotting, flash, parti, or<br>piebald (carrier) |

#### Interpretation:

This dog carries two copies of **a**<sup>t</sup> which results in tan points and can also present as a black and tan or tricolor coat color. However, this dog's coat color is also dependent on the E, K, and B genes. The tan point coat color is only expressed if the dog is also E/E or E/e at the E locus and k<sup>y</sup>/k<sup>y</sup> at the K locus. This dog will pass on **a**<sup>t</sup> to 100% of its offspring.

This dog carries one copy of one of the b mutations and has a B locus genotype of **B/b**. Thus, this dog typically will have a black coat, nose, and foot pads. However, this dog's coat color is dependent on the genotypes of many other genes. This dog will pass one copy of **B** to 50% of its offspring and one copy of **b** to 50% of its offspring. This dog can produce b/b offspring if bred to a dog that is also a carrier of a b mutation (B/b or b/b). Depending on the breed, b/b dogs may be referred to as brown, chocolate, liver or red.

This dog does not carry any copies of the  $d^1$  or  $d^2$  mutations and has a D locus genotype of **D/D** which does not result in the "dilution" or lightening of the pigments that produce the dog's coat color. This dog will pass one copy of **D** to 100% of its offspring and cannot produce d/d dogs.

This dog carries two copies of **e** which inhibits production of black pigment. The coat color of this dog will be yellow/red (including shades of white, cream, yellow, apricot or red). This dog will pass **e** on to 100% of its offspring.

This dog does not carry the mutation for weak furnishings or improper coat and will therefore have furnishings (proper coat). However, the overall coat type of this dog is dependent on the combination of this dog's genotypes at the L, Cu, and IC loci. This dog will pass **F** (furnishings, proper coat) to 100% of its offspring.

The K locus genotype for this dog is  $\mathbf{K}^{\mathbf{B}}/\mathbf{K}^{\mathbf{B}}$  which prevents expression of the agouti gene (A locus) and allows for solid eumelanin (black pigment) production in pigmented areas of the dog. However, this dog's coat color is also dependent on its genotypes at the E and B loci. This dog will pass on  $\mathbf{K}^{\mathbf{B}}$  to 100% of its offspring.

This dog carries one copy of **S** and one copy of  $s^{p}$  which results in limited white spotting, flash, parti, or piebald coat color due to the co-dominance of **S** and  $s^{p}$ . This dog will pass on one copy of **S** to 50% of its offspring and one copy of  $s^{p}$  to 50% of its offspring.

Paw Print Genetics<sup>®</sup> has genetic counseling available to you at no additional charge to answer any questions about these test results, their implications and potential outcomes in breeding this dog.

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Christina J Ramirez, PhD, DVM, DACVP Medical Director Robert D. Westra, MS, DVM Assistant Medical Director

Normal results do not exclude inherited mutations not tested in these or other genes that may cause medical problems or may be passed on to offspring. These tests were developed and their performance determined by Paw Print Genetics<sup>(b)</sup>. This laboratory has established and verified the tests' accuracy and precision. Because all tests performed are DNA-based, rare genomic variations may interfere with the performance of some tests producing false results. If you think these results are in error, please contact the laboratory immediately for further evaluation. In the event of a valid dispute of results claim, Paw Print Genetics will do its best to resolve such a claim to the customer's satisfaction. If no resolution is possible after investigation by Paw Print Genetics with the cooperation of the customer, the extent of the customer's sole remedy is a refund of the fee paid. In no event shall Paw Print Genetics be liable for indirect, consequential or incidental damages of any kind. Any claim must be asserted within 60 days of the report of the test results.

## About Amos



### Amos: The Perfect Family Dog

Looking for a handsome, affectionate, and adventurous companion? Look no further than Amos, our incredible Australian Labradoodle!
Full of personality and love, Amos enjoys playtime with endless games of fetch, walks in the park, and snuggles with his human family. An ideal addition to active households, he's always up for an exciting adventure, whether it's exploring new trails or conquering agility courses.
A social butterfly, Amos thrives on meeting new people and furry friends alike. Easy to train and well-behaved, he's a perfect choice for first-time dog owners.

But wait, there's more! Amos boasts a beautiful, soft, wavy coat that's hypoallergenic and low-shedding, making him a great fit for allergy sufferers. And did we mention he's a whiz in the agility ring? A quick learner, Amos is already excelling and sure to impress with his agility skills. Amos truly has it all! Contact South Downs Australian Labradoodles today to learn more about this amazing dog.

## About Your Breeder



Looking for a joyful, hypoallergenic companion?

South Downs Australian Labradoodles connects you with stunning, ethically bred pups in Sussex.

My name is Debbie, and I'm a passionate advocate for these incredible "designer dogs." Since 2020, I've dedicated myself to raising healthy, happy Australian Labradoodles in a loving home environment.



# Why Choose South Downs Australian Labradoodles?

- Small, nurturing home environment: As a small dedicated home breeder each litter of pups has our full attention, dedication and love from day one.
- Ethically bred, health-tested parents: We prioritise the well-being of our breeding dogs. All breeding dogs are fully health tested. These test include:- progressive Retinal Atrophy, Exercise induced collapse, Degenerative Myelopathy, Von Willebrand Disease and Hereditary Nasal Parakeratosis. To be affected by the above conditions puppies must receive a copy of the mutant gene from each parent, this is why health testing is so important. Hips and elbows are x-rayed and sent to the BVA for scoring. Only dogs with good hip and elbow scores are used for breeding. Eyes are tested every 18 months by the BVA eye scheme.
- Lifetime support: We're here to guide you on your puppy parenting journey, every step of the way.
- Gorgeous, authentic Australian Labradoodles: Prepare to be charmed by their intelligence, loyalty, and playful spirit.
- All our dogs family trees are available to be viewed on request showing many generations of genuine Australian Labradoodles and our membership to WALA.

Ready to welcome a furry friend into your life?

We'd love to connect! Contact us today to learn more about our upcoming litters and how you can become a South Downs Australian Labradoodle puppy parent.

> Tel Debbie: 07463228331 www.southdownsaustralianlabradoodles.co.uk