

PKD1 AND PERSIAN DERIVED PRA TEST REPORT

Provided Information: Case:

Name: RAYDUSOLEIL SARAH OF MINOOS Date Received: 12-Jan-2023 Report Issue Date: 17-Jan-2023

Registration: MC012XXX2-115423 Report ID: 0105-3154-0226-9188

Verify report at www.vgl.ucdavis.edu/verify

CAT143407

DOB: 01/16/2022 Sex: Female Breed: Maine Coon Microchip: 985141004407832 Color: BlackTortoiseshell

Sire: RIPPLE SPOD WAWELU Dam: MUSE COONS SAMANTHA OF RAYDUSOLEIL

Reg: SBT 031218 065 Reg: SBT 082219 074

Microchip: Microchip:

RESULT INTERPRETATION

PKD1	N/N	Normal - Does not possess the disease-causing PKD1 gene
PRA-pd		Not Requested



PKD1 AND PERSIAN DERIVED PRA TEST REPORT

Client/Owner/Agent Information:

GREG STAPLES 1014 SNIDER'S BAY ROAD GRAVENHURST ONTARIO P1P 1R2 CANADA
 Case:
 CAT143407

 Date Received:
 12-Jan-2023

Report Issue Date: 17-Jan-2023

Report ID: 0105-3154-0226-9188

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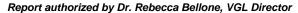
Name: RAYDUSOLEIL SARAH OF MINOOS

Additional Information

If testing for a disease or a disorder was performed and results indicate the animal is affected or at risk, we recommend contacting your veterinarian for further clinical evaluation and for additional information on disease and management.

For more detailed information on PKD1 and PRA-pd test results, please visit our website at: www.vgl.ucdavis.edu/services/pkd1.php www.vgl.ucdavis.edu/services/cat/PRApd.php

For terms and conditions of testing, please see www.vgl.ucdavis.edu/about/terms-and-conditions





DNA ANALYSIS CERTIFICATE

RAYDUSOLEIL SARAH OF MINOOS

Breed: Maine Coon

Sex: Female

Color: BlackTortoiseshell

DOB: 01/16/2022

Reg: MC012XXX2-115423

Alt. ID: 985141004407832

Case: CAT143407

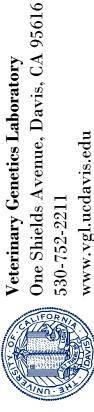
Report ID: 0105-3154-0226-9188

Print Date: January 17, 2023



Does not possess the disease-causing PKD1 gene.





Veterinary Genetics Laboratory

530-752-2211

www.vgl.ucdavis.edu

GRAVENHURST ONTARIO PIP 1R2 1014 SNIDER'S BAY ROAD GREG STAPLES CANADA



PK DEFICIENCY TEST REPORT

Provided Information:

Name: RAYDUSOLEIL SARAH OF MINOOS

Registration: MC012XXX2-115423

Case: CAT143407

Date Received: 12-Jan-2023
Report Issue Date: 17-Jan-2023

Report ID: 8499-2683-6492-2040

Verify report at www.vgl.ucdavis.edu/verify

DOB: 01/16/2022 Sex: Female Breed: Maine Coon Microchip: 985141004407832 Color: BlackTortoiseshell

Sire: RIPPLE SPOD WAWELU Dam: MUSE COONS SAMANTHA OF RAYDUSOLEIL

Reg: SBT 031218 065 Reg: SBT 082219 074

Microchip: Microchip:

PYRUVATE KINASE DEFICIENCY RESULT

N/K

Interpretation

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*



PK DEFICIENCY TEST REPORT

Client/Owner/Agent Information:

GREG STAPLES 1014 SNIDER'S BAY ROAD GRAVENHURST ONTARIO P1P 1R2 CANADA *Case:* CAT143407 *Date Received:* 12-Jan-2023

Date Received:12-Jan-2023Report Issue Date:17-Jan-2023

Report ID: 8499-2683-6492-2040

Verify report at www.vgl.ucdavis.edu/verify

Name: RAYDUSOLEIL SARAH OF MINOOS

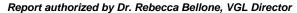
Additional Information

If testing for a disease or a disorder was performed and results indicate the animal is affected or at risk, we recommend contacting your veterinarian for further clinical evaluation and for additional information on disease and management.

For more detailed information on PK Deficiency test results, please visit our website at: www.vgl.ucdavis.edu/services/pkdeficiency.php

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.

For terms and conditions of testing, please see www.vgl.ucdavis.edu/about/terms-and-conditions







MAINE COON HCM (HYPERTROPHIC CARDIOMYOPATHY) TEST REPORT

Provided Information:

Name: RAYDUSOLEIL SARAH OF MINOOS

Registration: MC012XXX2-115423

Case: CAT143407

Date Received: 12-Jan-2023 Report Issue Date: 17-Jan-2023

Report ID: 2740-9028-3369-5160

Verify report at www.vgl.ucdavis.edu/verify

DOB: 01/16/2022 Sex: Female Breed: Maine Coon Microchip: 985141004407832 Color: BlackTortoiseshell

Sire: RIPPLE SPOD WAWELU Dam: MUSE COONS SAMANTHA OF RAYDUSOLEIL

Reg: SBT 031218 065 Reg: SBT 082219 074

Microchip: Microchip:

Maine Coon HCM Result

N/N

Interpretation

N/N Normal.

N/HCMmc One copy of the A31P mutation is present. Cat is 1.8 times more likely to develop HCM

than cats without the mutation.

HCMmc/HCMmc Two copies of the A31P mutation are present. Cat is 18 times more likely to develop

HCM than cats without the mutation.



MAINE COON HCM (HYPERTROPHIC CARDIOMYOPATHY) TEST REPORT

Client/Owner/Agent Information:

GREG STAPLES 1014 SNIDER'S BAY ROAD GRAVENHURST ONTARIO P1P 1R2 CANADA *Case:* CAT143407

Date Received: 12-Jan-2023
Report Issue Date: 17-Jan-2023

Report ID: 2740-9028-3369-5160

Verify report at www.vgl.ucdavis.edu/verify

Name: RAYDUSOLEIL SARAH OF MINOOS

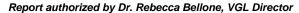
Additional Information

If testing for a disease or a disorder was performed and results indicate the animal is affected or at risk, we recommend contacting your veterinarian for further clinical evaluation and for additional information on disease and management.

For more detailed information on Maine Coon HCM test results, please visit our website at: www.vgl.ucdavis.edu/services/cat/MaineCoonHCM.php

The MHCM test only detects the A31P mutation associated with HCM in Maine Coon cats and outcrosses as described by Meurs et al. 2005. The A31P mutation is not the sole cause of HCM in Maine Coons. The other causes are not known at this time.

For terms and conditions of testing, please see www.vgl.ucdavis.edu/about/terms-and-conditions







MAINE COON SPINAL MUSCULAR ATROPHY TEST REPORT

Provided Information:

Name: RAYDUSOLEIL SARAH OF MINOOS

Registration: MC012XXX2-115423

Case: CAT143407

Date Received: 12-Jan-2023 Report Issue Date: 17-Jan-2023

Report ID: 2066-7306-7622-3080

Verify report at www.vgl.ucdavis.edu/verify

DOB: 01/16/2022 Sex: Female Breed: Maine Coon Microchip: 985141004407832 Color: BlackTortoiseshell

Sire: RIPPLE SPOD WAWELU Dam: MUSE COONS SAMANTHA OF RAYDUSOLEIL

Reg: SBT 031218 065 Reg: SBT 082219 074

Microchip: Microchip:

SMA Result

N/N

Interpretation

N/N No copies of SMA are present.

N/S 1 copy of SMA is present. Cat is normal but is a carrier. Breedings between carriers will be expected to produce

25% affected, 50% carriers and 25% normal kittens.

S/S 2 copies of SMA are present, cat is affected.



MAINE COON SPINAL MUSCULAR ATROPHY TEST REPORT

Client/Owner/Agent Information:

GREG STAPLES 1014 SNIDER'S BAY ROAD GRAVENHURST ONTARIO P1P 1R2 CANADA *Case:* CAT143407 *Date Received:* 12-Jan-2023

Date Received:12-Jan-2023Report Issue Date:17-Jan-2023

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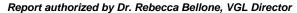
Additional Information

If testing for a disease or a disorder was performed and results indicate the animal is affected or at risk, we recommend contacting your veterinarian for further clinical evaluation and for additional information on disease and management.

For more detailed information on SMA test results, please visit our website at: www.vgl.ucdavis.edu/services/cat/SMA.php

The SMA test is specific for the mutation associated with SMA in Maine Coon cats and outcrosses.

For terms and conditions of testing, please see www.vgl.ucdavis.edu/about/terms-and-conditions







BLOOD GROUP TEST REPORT

Provided Information:

Name: RAYDUSOLEIL SARAH OF MINOOS

Registration: MC012XXX2-115423

Case: CAT143407

Date Received: 12-Jan-2023
Report Issue Date: 17-Jan-2023

Report ID: 9002-1845-4524-4199

Verify report at www.vgl.ucdavis.edu/verify

DOB: 01/16/2022 Sex: Female Breed: Maine Coon Microchip: 985141004407832 Color: BlackTortoiseshell

Sire: RIPPLE SPOD WAWELU Dam: MUSE COONS SAMANTHA OF RAYDUSOLEIL

Reg: SBT 031218 065 Reg: SBT 082219 074

Microchip: Microchip:

BLOOD GROUP RESULT

N/N

Interpretation

N/N Cat is Type A or Type AB

N/b Cat is a carrier of B factor; serotype could be Type A or Type AB

b/b Cat is Type B

N/c Cat is a carrier of AB factor; serotype could be Type A or Type AB

c/c Cat is type AB

c/b Cat is type AB; Carrier of B factor



BLOOD GROUP TEST REPORT

Client/Owner/Agent Information:

GREG STAPLES 1014 SNIDER'S BAY ROAD GRAVENHURST ONTARIO P1P 1R2 CANADA *Case:* CAT143407 *Date Received:* 12-Jan-2023

Report Issue Date: 17-Jan-2023

Report ID: 9002-1845-4524-4199

Verify report at www.vgl.ucdavis.edu/verify

Name: RAYDUSOLEIL SARAH OF MINOOS

Additional Information

If testing for a disease or a disorder was performed and results indicate the animal is affected or at risk, we recommend contacting your veterinarian for further clinical evaluation and for additional information on disease and management.

For more detailed information on Cat Blood Group test results, please visit our website at: www.vgl.ucdavis.edu/services/abblood.php

For terms and conditions of testing, please see www.vgl.ucdavis.edu/about/terms-and-conditions

