

OLOSKO DRUSALO

DURDEVAC

HRVATSKI KINOLOŠKI SAVEZ

ISPIT PRIROĐENIH OSOBINA RETRIVERA

BROJ:

Organizator: K.D	Shedavac	
Mjesto: JUCDAC	Datum: 29,9.2022.	
Ime psa: LINA AUIM	1A MAGICA	
Pasmina: Le		
HR 20709	Oštenjen: 7.2.021	Spol: 2
Vlasnik: IVANA VLA	STE	
Adresa vlasnika: K-2v	ONIMIRA 10 KARD	LOVAC
Vodič:	1571	

	ISPITNE DISCIPLINE	OPIS RADA
4	PONAŠANJE NA PUCANJ	Koperan
2	VODLJIVOST	vorque
3	DONOŠENJE PERNATE DIVLJAČI	APPRATICA 12 Suloy
4	DONOŠENJE PATKE IZ DUBOKE VODE	1- 12 VODE

USPJEH:

POLOŽIO A 1 NIJE POŁOŻIO KINOLOŠKI SUDAC Petar Klarić HKS - FCI 4 1. (sudac) Hinološki save M (potpis suca) 2. (sudac) (potpis suca)

	EDO	AGH	Od		Boja: COKOLADNA OCJENA: odličan	Ostenjen 15.04.2018	NINA BEST OF DISCOVERY	Majka / Dam	600	HDA	Odi		Boja: COKOLADNA OCJENA: odličan	Oštenjen: 30.08.2019	GALLIVANT LIVE TO RIDE	Otac / Sire	Otac i majka Sire and Dam	LABRADOR RETRIVER LABRADOR RETRIVER LINA ANIMA MAGICA Diaka / Coat Boja / Celour Uzgajivač / Breeder Uzgajivač / Breeder Uzgajivačnica: ANIMA MAG
	EDO		3	IRISH ROSE BEST OF DISCOVERY	HD:A	Ē	CHM-HR HJCH BIHCHM BHFMCH CHHR-I	GREENSTONE'S JOURNEY TO MY HEART				GALLIVANT'S LIKE A PRAYER AKC SS03962306				GALLIVANT ADAMANTLY BROWN AKC SS07908803	Djedovi Grand Sires and Grand Dams	SICA, F.C. I. 101/20
やアクビン	ED:0	CAMESWON AGREED ON EVERYTHING HR 15923 HD A	EDO	FI34818/10 HD:A		GREENSTONE'S CUPPA DELIGHT AKC SR75212702		CH EPOCH'S MOCCASIN JOE AKC SR68203801		HEYBERN'S WAIT TILL YOUR FATHER GETS HOME AKC SR75573501		CH GALLIVANT BLACK RUSSIAN AKC SR95240602		GOFETCH MOLASSES COADT AKC SR86751601		CH SHEABOURNE'S GRIZZLY ADAMS AKC SR69734602	Pradjedovi Great Grand Sires and Great Grand Dams	Upisan u H Pod brojem Spol / Sex Tetovirani t Broj čipa / I Broj čipa / I
	CH STENVEYZ BOX OF CHOCOLATE	CH GOWANVALE BALANCE KC AE02463802	MALLORN'S RING ANY BELLS	CH CHABLAIS YOUR PLACE OR MINE	CH GREENSTONE'S SPECIAL BLEND	CH BANNER'S NO LINE ON THE HORIZON	CH EPOCH'S TREASURE	CH QUAL CHASE BROADWAY JOE WINDFALL AKC SR47571508	HEYBERN'S NEW BALANCE AKC SR3S093202	HEYBERN'S RHUMBA JUMPS AKC SR51090803	CH GALLIVANT BIOPINES KENYA AKC SR84819301	CH ASTER LIBERTI LABRO HAMBURG AKC SR82948401	BEECHCROFT JERSEY STAR AKC SR63677704	CH EPOCH'S MOCCASIN JOE AKC SR68203801	CH SHEABOURNE'S PRECIOUS ANGEL AKC SR22661101	CH PARADOCS OBSIDIAN AKC SR56731701	Pra-pradjedovi Great Great Grand Sires and Great Great Grand Dam	Irvatsku rodovnu knjigu / Registered in Croatian Stud n / Reg. Number broj / Tattoo Number Microchip Number 191100002131144 HRV STUREY EX-operation

U Zagrebu, 8.04.2021



REFERENCE NO.: 2023 - 057350/01 OWNER: IVANA VLAŠIĆ KRALJA ZVONIMIRA 10 HR-4700 KARLOVAC CROATIA NAME/LABEL:

LINA ANIMA MAGICA SPECIES: DOG BREED: LABRADOR RETRIEVER SEX: FEMALE MICROCHIP NO.: 191100002131053 HRV TATOO NO.: NOT PROVIDED PEDIGREE NO.: HR 20709 LR

GENETIC REPORT

SAMPLE:	BLOOD
SAMPLE TAKEN BY:	SINIŠA DRUGA, DVM VETERINARSKA STANICA OZALJ, ODVOJAK KARLOVAČKE CESTE 110, 47280 OZALJ, CROATIA
REQUESTED TEST:	NARCOLEPSY - LABRADOR RETRIEVER
RESULT:	CLEAR (WT/WT)

COMMENT :

The test examines presence or absence of HCRTR2 gene mutation (c.1103+5G>A) described as the cause of narcolepsy in Labrador Retrievers. The disease is characterized by daytime sleepiness, cataplexy, and striking transitions from wakefulness into rapid eye movement sleep. Narcolepsy is inherited as an autosomal recessive trait.

Regarding to the presence of tested mutation animals are classified in three groups:

- Clear (wt/wt) mutation is not present, normal genotype
- Carrier (mut/wt) one of two alleles carries tested mutation, disease is not clinically manifested
- Affected (mut/mut) both alleles carry tested mutation, disease is clinically manifested

For each group different breeding strategies should be followed. Breeding of affected and carrier animals should be avoided. If particularly valuable animal is classified as affected, it should be bred only with clear animal. In such case, all first generation siblings will be carriers. If a carrier is bred with clear animal, 50% of siblings are expected to be clear. In case two carriers are bred, 25% of siblings are expected to be clear and 50% are expected to be carriers. However, 25% of siblings are expected to be affected, therefore such breeding practice is discouraged.

AUTHORIZED SIGNATURE:



MARIBOR, 17.07.2023

Results are valid for laboratory analysed samples only. Accuracy of the data about animal identity is the sole responsibility of the customer/owner. Laboratory is not responsible for false results which arise due to inaccurate animal identity data, false sample labels etc. To the extent the law allows, the maximal compensation



REFERENCE NO.: 2023 - 057350/01 OWNER: IVANA VLAŠIĆ KRALJA ZVONIMIRA 10 HR-4700 KARLOVAC CROATIA NAME/LABEL: LINA ANIMA MAGICA SPECIES: DOG BREED: LABRADOR RETRIEVER SEX: FEMALE MICROCHIP NO.: 191100002131053 HRV TATOO NO.: NOT PROVIDED PEDIGREE NO.: HR 20709 LR

GENETIC REPORT

SAMPLE: BLOOD

SAMPLE TAKEN BY: SINIŠA DRUGA, DVM VETERINARSKA STANICA OZALJ, ODVOJAK KARLOVAČKE CESTE 110, 47280 OZALJ, CROATIA

REQUESTED TEST: OBESITY

RESULT: CARRIER (MUT/WT)

COMMENT :

The test examines the presence of POMC gene mutation (c.345_346insC), responsible for an increased tendency to obesity in Labrador retriever. Genetic change is associated with higher body weight, greater motivation for food and obesity. POMC gene mutation is inherited as an autosomal recessive trait with incomplete penetrance.

Regarding to the presence of tested mutation animals are classified in three groups:

- Clear (wt/wt) mutation is not present, normal genotype
- Carrier (mut/wt) one of two alleles carries tested mutation, trait is not clinically manifested
- Affected (mut/mut) both alleles carry tested mutation, trait is clinically manifested

The genetic change is inherited in an autosomal recessive mode of inheritance with incomplete penetrance. That means the trait is not developed in every homozygous animal carrying deleterious mutation. Other genetic or environmental factors cannot be excluded in development of this trait. The test cannot exclude other genetic defects, which may be involved in development of the disease.

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LINA ANIMA MAGICA SPECIES: DOG BREED: LABRADOR RETRIEVER SEX: FEMALE MICROCHIP NO.: 191100002131053 HRV TATOO NO.: NOT PROVIDED PEDIGREE NO.: HR 20709 LR

GENETIC REPORT

SAMPLE:	BLOOD
SAMPLE TAKEN BY:	SINIŠA DRUGA, DVM VETERINARSKA STANICA OZALJ, ODVOJAK KARLOVAČKE CESTE 110, 47280 OZALJ, CROATIA
REQUESTED TEST:	CENTRONUCLEAR MYOPATHY (CNM)
RESULT:	CLEAR (WT/WT)

COMMENT :

The test examines presence or absence of PTPLA gene mutation (c.191_192ins236bp) described as the cause of centronuclear myopathy (CNM) in Labrador Retriever. The disease is characterized by hypotonia, generalized muscle weakness, abnormal postures, stiff hopping gait, exercise intolerance and increased collapse when exposed to cold. CNM is inherited as an autosomal recessive trait.

Regarding to the presence of tested mutation animals are classified in three groups:

- Clear (wt/wt) mutation is not present, normal genotype
- Carrier (mut/wt) one of two alleles carries tested mutation, disease is not clinically manifested
- Affected (mut/mut) both alleles carry tested mutation, disease is clinically manifested

For each group different breeding strategies should be followed. Breeding of affected and carrier animals should be avoided. If particularly valuable animal is classified as affected, it should be bred only with clear animal. In such case, all first generation siblings will be carriers. If a carrier is bred with clear animal, 50% of siblings are expected to be clear. In case two carriers are bred, 25% of siblings are expected to be clear and 50% are expected to be carriers. However, 25% of siblings are expected to be affected, therefore such breeding practice is discouraged.

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GENETIC REPORT

SAMPLE:	BLOOD
SAMPLE TAKEN BY:	SINIŠA DRUGA, DVM VETERINARSKA STANICA OZALJ, ODVOJAK KARLOVAČKE CESTE 110, 47280 OZALJ, CROATIA
REQUESTED TEST:	EXCERSISE INDUCED COLLAPSE (EIC)
RESULT:	CLEAR (WT/WT)

COMMENT :

The test examines presence or absence of DNM1 gene mutation (c.767G>T) described as the cause of exercise induced collapse (EIC) in several dog breeds. EIC is a syndrome characterized by collapse episodes following strenuous exercise. DNM1 gene defect is inherited as an autosomal recessive trait.

Regarding to the presence of tested mutation animals are classified in three groups:

- Clear (wt/wt) mutation is not present, normal genotype
- Carrier (mut/wt) one of two alleles carries tested mutation, disease is not clinically manifested
- Affected (mut/mut) both alleles carry tested mutation, disease is clinically manifested

For each group different breeding strategies should be followed. Breeding of affected and carrier animals should be avoided. If particularly valuable animal is classified as affected, it should be bred only with clear animal. In such case, all first generation siblings will be carriers. If a carrier is bred with clear animal, 50% of siblings are expected to be clear. In case two carriers are bred, 25% of siblings are expected to be clear and 50% are expected to be carriers. However, 25% of siblings are expected to be affected, therefore such breeding practice is discouraged.

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LINA ANIMA MAGICA SPECIES: DOG BREED: LABRADOR RETRIEVER SEX: FEMALE MICROCHIP NO.: 191100002131053 HRV TATOO NO.: NOT PROVIDED PEDIGREE NO.: HR 20709 LR

GENETIC REPORT

SAMPLE: BLOOD

SAMPLE TAKEN BY: SINIŠA DRUGA, DVM VETERINARSKA STANICA OZALJ, ODVOJAK KARLOVAČKE CESTE 110, 47280 OZALJ, CROATIA

REQUESTED TEST: RETINAL DYSPLASIA/OCULOSKELETAL DYSPLASIA (RD/OSD)

RESULT: CLEAR (WT/WT)

COMMENT :

The test examines presence or absence of COL9A3 gene mutation (g.49,699,847insG) described as the cause of Retinal Dysplasia/Oculoskeletal Dysplasia (RD/OSD) in Labrador retriever. The disease is characterized by short-limbed dwarfism and ocular defects.

Regarding to the presence of tested mutation animals are classified in three groups:

- Clear (wt/wt) mutation is not present, normal genotype
- Carrier (mut/wt) one of two alleles carries tested mutations, partial expression of RD/OSD
- Affected (mut/mut) both alleles carry tested mutations, full expression of RD/OSD

Carrier dogs typically show symptoms of Retinal Dysplasia - partial expression of RD/OSD. Affected dogs show symptoms of dwarfism and ocular defects - full expression of RD/OSD. Heterozygote animals pass the mutation to their siblings therefore mating of two carriers should be avoided, as 25% of puppies will be affected, 50% of puppies will be clear.

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LINA ANIMA MAGICA SPECIES: DOG BREED: LABRADOR RETRIEVER SEX: FEMALE MICROCHIP NO.: 191100002131053 HRV TATOO NO.: NOT PROVIDED PEDIGREE NO.: HR 20709 LR

GENETIC REPORT

SAMPLE:	BLOOD
SAMPLE TAKEN BY:	SINIŠA DRUGA, DVM VETERINARSKA STANICA OZALJ, ODVOJAK KARLOVAČKE CESTE 110, 47280 OZALJ, CROATIA
REQUESTED TEST:	PROGRESSIVE RETINAL ATROPHY (PRA-PRCD)
RESULT:	CLEAR (WT/WT)

COMMENT :

The test examines presence or absence of PRCD gene mutation (c.5G>A) described as the cause of one form of progressive retinal atrophy (PRA) in several dog breeds. PRA-PRCD is a late onset disease characterized by progressive degeneration of retinal cells. PRCD gene defect is inherited as an autosomal recessive trait.

Regarding to the presence of tested mutation animals are classified in three groups:

- Clear (wt/wt) mutation is not present, normal genotype
- Carrier (mut/wt) one of two alleles carries tested mutation, disease is not clinically manifested
- Affected (mut/mut) both alleles carry tested mutation, disease is clinically manifested

For each group different breeding strategies should be followed. Breeding of affected and carrier animals should be avoided. If particularly valuable animal is classified as affected, it should be bred only with clear animal. In such case, all first generation siblings will be carriers. If a carrier is bred with clear animal, 50% of siblings are expected to be clear. In case two carriers are bred, 25% of siblings are expected to be clear and 50% are expected to be carriers. However, 25% of siblings are expected to be affected, therefore such breeding practice is discouraged.

AUTHORIZED SIGNATURE:



MARIBOR, 17.07.2023

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REFERENCE NO.: 2023 - 057350/01 OWNER: IVANA VLAŠIĆ KRALJA ZVONIMIRA 10 HR-4700 KARLOVAC CROATIA NAME/LABEL:

LINA ANIMA MAGICA SPECIES: DOG BREED: LABRADOR RETRIEVER SEX: FEMALE MICROCHIP NO.: 191100002131053 HRV TATOO NO.: NOT PROVIDED PEDIGREE NO.: HR 20709 LR

GENETIC REPORT

SAMPLE:	BLOOD
SAMPLE TAKEN BY:	SINIŠA DRUGA, DVM VETERINARSKA STANICA OZALJ, ODVOJAK KARLOVAČKE CESTE 110, 47280 OZALJ, CROATIA
REQUESTED TEST:	SKELETAL DYSPLASIA 2 (SD2) - DWARFISM
RESULT:	CLEAR (WT/WT)

COMMENT :

The test examines presence or absence of COL11A2 gene mutation (c.143G>C) described as the cause of skeletal dysplasia 2 (SD2) in Labrador Retriever. The disease is characterized by a very subtle phenotype where mild dwarfism with short-legged phenotype is observed. COL11A2 gene defect is inherited as an autosomal recessive trait.

Regarding to the presence of tested mutation animals are classified in three groups:

- Clear (wt/wt) mutation is not present, normal genotype
- Carrier (mut/wt) one of two alleles carries tested mutation, disease is not clinically manifested
- Affected (mut/mut) both alleles carry tested mutation, disease is clinically manifested

For each group different breeding strategies should be followed. Breeding of affected and carrier animals should be avoided. If particularly valuable animal is classified as affected, it should be bred only with clear animal. In such case, all first generation siblings will be carriers. If a carrier is bred with clear animal, 50% of siblings are expected to be clear. In case two carriers are bred, 25% of siblings are expected to be clear and 50% are expected to be carriers. However, 25% of siblings are expected to be affected, therefore such breeding practice is discouraged.

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OWNER: IVANA VLAŠIĆ KRALJA ZVONIMIRA 10 HR-4700 KARLOVAC CROATIA

DATE:17.07.2023

TEST REPORT NO. 211035

TEST: HEREDITARY NASAL PARAKERATOSIS (HNPK)

MUTATION: c.972 T>G in SUV39H2 gene

RESULT: CLEAR (NORMAL/NORMAL)

ANIMAL NAME: LINA ANIMA MAGICA

SPECIES: DOG

BREED: LABRADOR RETRIEVER

MICROCHIP NO.: 191100002131053 HRV

PEDIGREE NO.: HR 20709 LR

SAMPLE TYPE: BLOOD

SAMPLE TAKEN BY: SINIŠA DRUGA, DVM

RESULT COMMENT:

Clear (normal/normal): tested mutation is not present, normal genotype.

Carrier (normal/mutation): one allele carries tested mutation, disease is not clinically manifested.

Affected (mutation/mutation): both alleles carry tested mutation, disease is clinically manifested.

AUTHORIZED SIGNATURE:	DOGMITEST
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LOUBIMARSKA AMBULANTA

OFTALMOLOŠKI PREGLED - OPHTALMIC EXAMINATION

OPIS ŽIVOTINJE / ANIMAL DESCRIPTION

IME ŽIVOTINJE / NAME : Lina Anima Magica	
DATUM ROĐENJA / DATE OF BIRTH : 7.2.2021.	2
VRSTA / SPECIES : DOG SPOL / SEX :	F
PASMINA / BREED : LABRADOR RETRIVER BOJA / COLOR :	BROWN
MIKROČIP / MICROCHIP Nº : 191100002131053 BROJ RODOVNI	KA / PEDIGREE Nº : HR 20709LR
VLASNIK / OWNER	
IME I PREZIME / FIRST NAME, SURNAME : IVANA VLAŠIĆ	
ADRESA / ADRESS : KRALJA ZVONIMIRA 10 , KARLOVAC	
PREGLED OKA - EXAMINATIO	N OF EYE
MYDRIATIC : X OPHTALMOSCOPY : DIRECT X INDIRECT V	BIOMICROSCOPY
PARTS EXAMINED ADNEXA CORNEA IRIS LENS CLINICALLY AFFECTED	VITREOUS FUNDUS CLINICALLY CLINICALLY AFFECTED UNAFFECTED
PMP PA PA <td>COLLIE EYE ANOMALY X RD RETINAL DYSPLASIA X ESISTENT PUPILLARY MEMBRANE X PHPV PERSISTENT HYPERPLASTIC X PRIMARY VITREUS X CUNICA VASCULOSA LENTIS X CATARACT X CORNEAL DYSTROPHY X PRIMARY LENS LUXATION X RETINAL DEGENERATION (PRA) X ENTROPION X OPTIC NERVE HYPOPLASIA X Vlatka-Antonija Csik DMV</td>	COLLIE EYE ANOMALY X RD RETINAL DYSPLASIA X ESISTENT PUPILLARY MEMBRANE X PHPV PERSISTENT HYPERPLASTIC X PRIMARY VITREUS X CUNICA VASCULOSA LENTIS X CATARACT X CORNEAL DYSTROPHY X PRIMARY LENS LUXATION X RETINAL DEGENERATION (PRA) X ENTROPION X OPTIC NERVE HYPOPLASIA X Vlatka-Antonija Csik DMV
DATUM / DATE OF EXAMINATION : 5.1.2023.	VLATKA-ANFONIJA CSIK dr. med. vet. OVLAŠTENI VZUZRINAR br. 2042



HRVATSKI KINOLOŠKI SAVEZ CROATIAN KENNEL CLUB ILICA 61 HR – 10 000 ZAGREB <u>www.hks.hr</u> <u>hks@hks.hr</u>



WORKING CLASS CERTIFICATE (WCC)

Name of the dog Nom du chien Name des Hundes Nombre del perro	LINA ANIM	A MA	AGICA			
Breed & Variety Race & Variété Rasse & Varietät Raza & Variedad	LABRADOR R	ETRIE	VER			
Sex Sexe Geschlecht Sexo	male/mâle Rüde/macho	fe Hü	male/femelle indin/hembra	Date of birth Date de nais Wurfdatum Fecha de nac	sance	07. Februar 2021
Livre des Origines et nu Zuchtbuch- und Zuchtb Libro de orígenes y nún	ion number Iméro d'enregistrement uchnummer nero de registro	HR 2	0709 LR	6		
Tattoo or microchip nu Numéro du tatouage ou Tätowier- oder Microch Número del tatuaje o d	mber u de microchip nipnummer lel microchip	191:	10000213	31053		
Owner Propriétaire Eigentümer Propietario	VLAŠIĆ IV	ANA				
Country of legal resider Pays de résidence légal Land (Gesetzlicher Wol País de residencia legal	hrsitz) HR-47 00	0, KA	RLOVAC,	Kralja Z	vonimir	a 10, Kroatien
This dog has passed required working trial is is entitled to be enter in the working class at FCL shows (according the decision taken by General Assembly of FCL in Madrid, 1983).	the Ce chien a passé ave and l'épreuve exigée l'au red à s'inscrire dans la t all travail de toute to expositions FCI the décision de l'Ass the Générale de la FCI à 1983).	c succès itorisant a classe es les (selon semblée Madrid	Der genannte erforderliche bestanden und bei allen FCI-A Gebrauchshund werden (gemäß Generalversamr 1983).	Hund ha Arbeitsp ist damit bere usstellungen eklasse gemel Beschluss de mlung in t	t die rüfung chtigt, in die ldet zu er FCI- Madrid	perro ha aprobado la la exigida para poder birse en la clase trabajo de las exposiciones de la FCI n resolución de la blea General de la FCI en id en 1983).
Place (town, COUNTRY) Lieu (ville, PAYS) et dat Ort (Stadt, LAND) und D Lugar (ciudad, PAÍS) de) and date of the trial te de l'épreuve Datum der Prüfung la prueba		MOLVE	KRO	ATIEN	01.10.2023
Type of the trialType d'épreuvePrüfungstypTipo de prueba	EITSPRÜFUNG FÜR IEVER MIT BRINGEN		Qualificatio Qualificatio Qualifikatio Calificación	n / points n / points n / Punkte / puntos	Insgesamt Bewertung	104 Punkte g: (3) Gut
Name of the judge(s) Nom du (des) juges Name des (der) Richter(Nombre del juez (de los	s) jueces)	Petar KL	ARIĆ		/	
Date, date, Datum, fe 25.10.2023.	cha Stampt	ignature	cachet/sigr E	iature – Sten Jojan Matak Gene	npel/Unterso covic, Dr.Ve eralsekretär	chrift –sello/firma et.Med.

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ZAVOD ZA RENDGENOLOGIJU, ULTRAZVUČNU DIJAGNOSTIKU I FIZIKALNU TERAPIJU VETERINARSKI FAKULTET SVEUCILISTA U ZAGREBU

DEPARTMENT OF RADIOLOGY, ULTRASOUND DIAGNOSTICS AND PHYSICAL THERAPY FACULTY OF VETERINARY MEDICINE UNIVERSITY OF ZAGREB



HEINZLOVA 55, ZAGREB

TEL: +385 (0)1 23 90 401



POTVRDA - CERTIFICATE

13.7.2022.	DATUM SNIMANJA - X-RAYS MADE ON:
X	DISPLAZIJA LAKTOVA - ELBOW DYSPLASIA
X	DISPLAZIJA KUKOVA - HIP DYSPLASIA

BROJ PROTOKOLA - PROTOCOL NO: RTG-5967/22

\leq	
M SNIMANJA - X-RAYS MAD	
E ON:	
13.7.2	

PODACI O PSII - DOG DATA

PASMINA - BREED:	labrador retriver
IME PSA - DOG NAME:	Lina Anima Magica
SPOL - SEX:	female
DATUM OŠTENJENJA - DATE OF BIRTH:	7.2.2021.
BROJ RODOVNICE - PEDIGREE NUMBER:	HR 20709 LR
BROJ MIKROČIPA - MICROCHIP NUMBER:	191100002131053
VLASNIK - OWNER.	Ivana Vlašić
ADRESA VLASNIKA - OWNER ADDRESS:	47000 KARLOVAC, Kralja Zvonimira 10

13.7.

	•) -	KTOVI - EI ROW
_	A B	JKOVI - HIP

DCJENU IZDAO - THE EVALUATION
I WAS MADE BY: Ana Javor, dr. med. vet.

Postupak je izveden prema FCI pravilniku.

The procedure has been performed according to the rules of FCI.

POTPIS - SIGNATURE:

ZIG - STAMP:

VETERINARSKI FAKULTET U ZAGREBU Zavod za reedgezologiju, ufarazvašau angenetten i fizileninn sonspiju