OSTAIL podaci / Other information

Decretary Colors

All locales

All

Vlasnik / Owner

1 Wara Vlasic Treg 68 02021

Potpis uzgajivača / Breeder's Signature

2.

Potpis prijašnjeg vlasnika / Former Owner's Signature

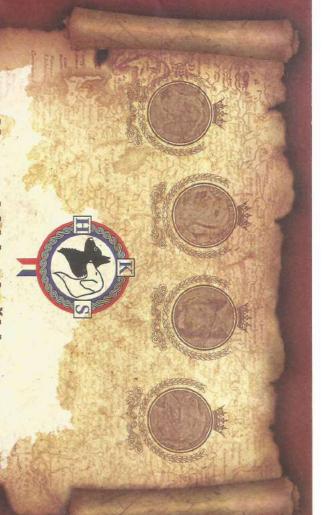
Potpis prijašnjeg vlasnika / Former Owner's Signature

Potpis prijašnjeg vlasnika / Former Owner's Signature



Hrvatski kinološki savez / Croatian Kennel Club Ilica 61, HR-10000 Zagreb, tel. + 385 (0)1 4846 124, www.hks.hr





Hrvatski kinološki savez Croatian Kennel Club

REPUBLIKA HRVATSKA / Republic of Croatia

Članica FCI / Member of FCI (Fédération Cynologique Internationale)

RODOWNICA PEDIGRAM



Pasmina / Breed

LABRADOR RETRIVER

LABRADOR RETRIEVER

Dlaka / Coat Oštenjen / Born On Ime psa / Name

Boja / Colour

Uzgajivač / Breeder

**ĐURINA 26, SELCI ĐAKOVAČKI 31415** 

Uzgajivačnica: BEST OF DISCOVERY, F.C.I.70/01

Grand Sires and Grand Dams

EPOCH'S MOCCASIN JOE

USAGCH USA-CH AKC SR68203801 Otac i majka

Otac / Sire

Ostenjen:

12.08.2015

COKOLADNA

CHM-HR HJCH BIHCHM BHFMCH CHHR-I

Broj rodovnice: HR 18888

GREENSTONE'S JOURNEY TO MY HEART

FILIC MARIO CHOCOLATE COKOLADNA 19.04.2018 NINA BEST OF DISCOVERY KRATKA / SHORT

1891.



HRVATSKI KINOLOŠKI SAVEZ Croatian Kennel Club

> Spot / Sex Pod brojem / Reg. Number Upisan u Hrvatsku rodovnu knjigu / Registered in Croatian Stud Book

HR 19465 LR **ŽENKA / FEMALE** 

Broj čipa / Microchip Number Tetovirani broj / Tattoo Number





100					2//5												
		CH STENVEYZ BOX OF CHOCOLATE  UKU 0011/3/04 HD:B/A	ED:0	CH GOWANVALE BALANCE KC AE02463802 HD:A	ED:0	MALLORN'S RING ANY BELLS FIN21713/07 HD:A	ED:free	CH CHABLAIS YOUR PLACE OR MINE CKC TA138694 HD:A		CH GREENSTONE'S SPECIAL BLEND AKC SR61364502		CH BANNER'S NO LINE ON THE HORIZON AKC SR67618601		CH EPOCH'S TREASURE AKC SR52301903		CH QUAIL CHASE BROADWAY JOE WINDFALL AKC SR47571508	Pradjedovi Great Grand Sires and Great Grand Dams
	CH STENVEYZ DESSERT CHOCOLATE RKF 1026750	CH STENVEYZ RED OCTOBER UKU 000575/07	OULSMI RUMAWAY TO GOWANVALE KC AC03118202	CH NIAKEN KRIS KCSB 3314 CI	CH MALLORN'S TERRACOTTA FIN17/17/002	MALLORN'S ROMEO FIN39321/04	CH CHABLAIS DELORME ZIGGY CKC PJ859909	BIG SKY'S STONE KUTTER CKC SR22980804	CH BOURNHALL'S MOCHA LATTE AT PUCKETTS AKC SR34191304	TEQUILAB'S AMADEUS AIR AKC SR59548601	EPOCH'S BAG OF TRICKS AT BANNER AKC SR28660602	CH ENDEAVOR PICCADILLY GOTTA WEAR SHADES AKC SR40600101	CH EPOCH'S BROWN BAGGIN' AKC SN91416901	CH WILCARE LEISURE SUIT LARRY JH AKG SR25847301	CH DUNE DANCER'S WINDFALL MARGARITA AKG SR11049901	CH LIBERTY'S BROAD RIDGE AKC SN91181201	Pra-pradjedovi Great Great Grand Sires and Great Great Grand Dams



ED:0

ED:0

IPO

HR 15923

CAMESWON AGREED ON EVERYTHING

IPO

Boja Ostenjen:

COKOLADNA

ED:0

7.07.2012

OCJENA: odličan

Broj rodovnice: HR 17311

IRISH ROSE BEST OF DISCOVERY

MALLORN'S RING MY BELL FI34818/10

Majka / Dam

ED:0

IPO

AKC SR75212702

GREENSTONE'S CUPPA DELIGHT



Matije Ivanića 19, 10000 Zagreb tel.: 01/3750-339 www.ljubimac.net ljubimac.ambulanta@gmail.com

# OFTALMOLOŠKI PREGLED - OPHTALMIC EXAMINATION

OFIS ZIVOTIME / AMINIAL DESCRIPTION		
IMEŽIVOTINJE / NAME: NINA BEST OF DISCON	very	enn.
DATUM ROĐENJA / DATE OF BIRTH: 19.04. 2018.		
VRSTA / SPECIES : PA_S SPOL / SEX :	£	
PASMINA/BREED: LARSCADOR RETRIVEL BOJA/COLOR:	COKO LADNA	
MIKROČIP / MICROCHIP No : 1911 00000990772 BROJ RODOVNIKA		-
VLASNIK / OWNER		
IME I PREZIME / FIRST NAME, SURNAME : VLA	zi c	
ADRESA / ADRESS: TRG 68, 47280	ozay Havatorus	
PREGLED OKA - EXAMINATION	N OF EYE	
MYDRIATIC: X OPHTALMOSCOPY: DIRECT X INDIRECT	BIOMICROSCOPY 🔂 OTHER	
CLINICALLY AFFECTED		CALLY
CLINICALLY UNAFFECTED	COLLIE EYE ANOMALY	X
		X
PPM PES		х
P A A P	PRIMARY VITREUS	Х
		X
		X
		X
		х
R	The state of the s	x
KOMENTAR / COMMENTS : After examination of both eyes of the dog I don't find any		X
patological or degenerative changes.		
	Vlatka-Antonija Csik DMV	
DATUM / DATE OF EXAMINATION : 12 9 2 2	VLATKA-ANTONIJA	CSIK





# 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

TEL: +385 (0)1 23 90 401



HEINZLOVA 55, ZAGPEB

# POTVRDA - CERTIFICATE

DISPLAZIJA LAKTOVA - ELBOW DYSPLASIA	DISPLAZIJA KUKOVA - HIP DYSPLASIA	×
	DISPLAZIJA LAKTOVA - ELBOW DYSPLASIA	×

**BROJ PROTOKOLA - PROTOCOL NO:** 5953/20

# PODACI O PSU - DOG DATA

PASMINA - BREED:	Labrador retriever	
IME PSA - DOG NAME:	Nina Best of Discovery	
SPOL - SEX:		
DATUM OŠTENJA - DATE OF BIRTH:	19.04.2018.	
BROJ RODOVNICE - PEDIGREE NUMBER:	HR 19465 LR	
BROJ MIKROČIPA - MICROCHIP NUMBER:	191100000990772	
VLASNIK - OWNER:	Ivana Vlašić	
ADRESA VLASNIKA - OWNER ADDRESS:	47280 OZALJ, Trg 68	

# OCJENA - CLASSIFICATION

KUKOVI - HIP	A	В	C	D	Ш
LAKTOVI - ELBOW	(e)		2	ప	

OCJENU IZDAO - THE EVALUATION WAS MADE BY:

Branimir Škrlin DVM

Postupak je izveden prema FCI pravilniku.

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

POTPIS - SIGNATOR

VETERINARSKI FAKULTET U ZAGREBU Zavod za rezdigenski jelyk ukrazvučnu dljagnostiku i fizikalnu terapiju





Sveučilište u Zagrebu Veterinarski Fakultet, Tel. 2390111 Heinzelova 55, Zagreb Odjel klinika Zavod za rendgenologiju, ultrazvuk i fizikalnu terapiju Laboratorij za RTG i UZV Vrijeme prijema: 15.9.2020. 10:18:09 Vrijeme unosa nalaza: 15.9.2020. 10:20:13

## Nalaz rendgenološke i/ili UZV pretrage

RTG-5953/2020

Broj uputnice: RTG-2423/20

Uputio/la: Branimir Škrlin, dr. med. vet.

Datum: 15.9.2020.

Vlasnik: Ivana Vlašić, OIB: -, 47280 OZALJ, Trg 68, tel. HR 19465 LR

Pacijent: kuja labrador retriver Nina Best of Discovery, datum rođenja: 19.4.2018, ID: 191100000990772

Što je rađeno

Rtg

Dijagnoza

HD-A; ED-0 (free);

Voditelj laboratorija

Prof. dr. sc. Damir Stanin

VETERINARSKI PredstojnikT U ZAGREPU Zavod za rendgenologiju, ukrazvučnu

Prof. dr. sc. Damir Stanin

Nalaz se odnosi na dostavljeni i pretraženi uzorak te se ne smije umražati bez pismenog odobrenja laboratorija.

Branimir Škrlin, dr. med. vet.

Vrijeme ispisa: 15.9.2020. 10:21:02



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www.eurovetgene.com

**REFERENCE NO.:** 2020 - 35345

**OWNER:** 

MATEJA TUŠČIĆ PALMOTIĆEVA 82 HR-10000 ZAGREB

**CROATIA** 

NAME/LABEL:

NINA BEST OF DISCOVERY

**SPECIES:** DOG

**BREED: LABRADOR RETRIEVER** 

**SEX:** FEMALE

MICROCHIP NO.: 191100000990772

TATOO NO.: NOT PROVIDED PEDIGREE NO.: HR 19465 LR

## **GENETIC REPORT**

**SAMPLE:** BLOOD

**SAMPLE TAKEN BY:** VLATKA-ANTONIJA CSIK, DVM, VETERINARSKA AMBULANTA LJUBIMAC, MATIJE

IVANIČA 19, 10000 ZAGREB, 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:** 





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**REFERENCE NO.:** 2020 - 35345

**OWNER:** 

MATEJA TUŠČIĆ PALMOTIĆEVA 82 HR-10000 ZAGREB

**CROATIA** 

NAME/LABEL:

NINA BEST OF DISCOVERY

**SPECIES:** DOG

**BREED: LABRADOR RETRIEVER** 

**SEX:** FEMALE

MICROCHIP NO.: 191100000990772

TATOO NO.: NOT PROVIDED PEDIGREE NO.: HR 19465 LR

# **GENETIC REPORT**

**SAMPLE:** BLOOD

**SAMPLE TAKEN BY:** VLATKA-ANTONIJA CSIK, DVM, VETERINARSKA AMBULANTA LJUBIMAC, MATIJE

IVANIČA 19, 10000 ZAGREB, 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.

**AUTHORIZED SIGNATURE:** 





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CROATIA

NAME/LABEL:

NINA BEST OF DISCOVERY

**SPECIES:** DOG

**BREED: LABRADOR RETRIEVER** 

**SEX:** FEMALE

MICROCHIP NO.: 191100000990772

TATOO NO.: NOT PROVIDED PEDIGREE NO.: HR 19465 LR

# **GENETIC REPORT**

SAMPLE: BLOOD

**SAMPLE TAKEN BY:** VLATKA-ANTONIJA CSIK, DVM, VETERINARSKA AMBULANTA LJUBIMAC, MATIJE

IVANIČA 19, 10000 ZAGREB, 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.

**AUTHORIZED SIGNATURE:** 





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**REFERENCE NO.:** 2020 - 35345

OWNER:

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**CROATIA** 

NAME/LABEL:

NINA BEST OF DISCOVERY

**SPECIES:** DOG

**BREED: LABRADOR RETRIEVER** 

**SEX:** FEMALE

MICROCHIP NO.: 191100000990772

TATOO NO.: NOT PROVIDED PEDIGREE NO.: HR 19465 LR

## **GENETIC REPORT**

SAMPLE: BLOOD

SAMPLE TAKEN BY: VLATKA-ANTONIJA CSIK, DVM, VETERINARSKA AMBULANTA LJUBIMAC, MATIJE IVANIČA 19,

10000 ZAGREB, 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 carriers and 25% of puppies will be clear.

**AUTHORIZED SIGNATURE:** 

MARIBOR, 29.09.2020



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 for potential false result is limited to the invoiced amount. With the test it is not possible to rule out the presence of other genetic changes which might affect the development of the disease. Testing is performed according to the latest scientific knowledge.



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**REFERENCE NO.:** 2020 - 35345

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CROATIA

NAME/LABEL:

NINA BEST OF DISCOVERY

**SPECIES:** DOG

**BREED: LABRADOR RETRIEVER** 

**SEX:** FEMALE

MICROCHIP NO.: 191100000990772

TATOO NO.: NOT PROVIDED PEDIGREE NO.: HR 19465 LR

# **GENETIC REPORT**

SAMPLE: BLOOD

**SAMPLE TAKEN BY:** VLATKA-ANTONIJA CSIK, DVM, VETERINARSKA AMBULANTA LJUBIMAC, MATIJE

IVANIČA 19, 10000 ZAGREB, 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:** 





**OWNER:**MATEJA TUŠČIĆ
PALMOTIĆEVA 82
HR-10000 ZAGREB
CROATIA

DATE:29.09.2020

# TEST REPORT NO. 204314

TEST: HEREDITARY NASAL PARAKERATOSIS (HNPK)

MUTATION: c.972 T>G in SUV39H2 gene

**RESULT:** CLEAR (NORMAL/NORMAL)

ANIMAL NAME: NINA BEST OF DISCOVERY

**SPECIES:** DOG **BREED:** LABRADOR RETRIEVER

MICROCHIP NO.: 191100000990772 PEDIGREE NO.: HR 19465 LR

**SAMPLE TYPE:** BLOOD

SAMPLE TAKEN BY: VLATKA-ANTONIJA CSIK, 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:** 

Results are valid for laboratory analysed samples only

GOOGTEST



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**REFERENCE NO.:** 2020 - 35345

**OWNER:** 

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**CROATIA** 

NAME/LABEL:

NINA BEST OF DISCOVERY

**SPECIES:** DOG

**BREED: LABRADOR RETRIEVER** 

**SEX:** FEMALE

MICROCHIP NO.: 191100000990772

TATOO NO.: NOT PROVIDED PEDIGREE NO.: HR 19465 LR

# **GENETIC REPORT**

**SAMPLE:** BLOOD

**SAMPLE TAKEN BY:** VLATKA-ANTONIJA CSIK, DVM, VETERINARSKA AMBULANTA LJUBIMAC, MATIJE

IVANIČA 19, 10000 ZAGREB, 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.

**AUTHORIZED SIGNATURE:** 





### HRVATSKI KINOLOŠKI SAVEZ CROATIAN KENNEL CLUB

ILICA 61 HR – 10 000 ZAGREB www.hks.hr hks@hks.hr



# **WORKING CLASS CERTIFICATE (WCC)**

Nom du chien Name des Hundes	NI	NA BEST	OF D	ISCOVE	RY		
Nombre del perro							
Breed & Variety					-		
ace & Variété	IAR	RADOR F	FTRIE	VFD			
asse & Varietät	באם	MADON	LINIL	VLK			
aza & Variedad							
ex					Date of bir	th	
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exo		Rüde/macho	Hü	ndin/hembra	Fecha de na		
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Numéro du tatoua	·	crochin					
Tätowier- oder Mi		7.1	1913	1000009	90772		
Número del tatua							
Owner	je o dei mich	Cilip					
Propriétaire		v. z					
Eigentümer		VLAŠIĆ I	VANA				
Propietario							
Country of legal re	esidence						
Pays de résidence		HR_47 0	OO KA	RIOVAC	Kralia	Zvonimi	a 10 Kroation
Land (Gesetzliche País de residencia This dog has pa	er Wohnsitz) a legal assed the Co	e chien a passé a	vec succès	Der genannt		at die	
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