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22nd - 28th Mar 2025

Early Detection of Rabies in Dogs and Other Animals: Recognizing Behavioral Changes and Clinical Progression for Effective Dog Bite Prevention - Part I

This is the first article of two in a series on "Early Detection of Rabies in Dogs and Other Animals: Recognizing Behavioral Changes and Clinical Progression for Effective Dog Bite Prevention"

Rabies is a deadly, zoonotic disease caused by a neurotropic virus of the Rhabdoviridae family that primarily affects the central nervous system of mammals. Dog bites and scratches account for 99% of human rabies cases, and the disease can be largely prevented through dog vaccination and bite prevention. Once clinical signs appear, rabies is almost always fatal, making early detection of rabies in dogs and prevention of dog bites crucial for prevention and control. Recognizing symptoms of rabies in dogs, such as aggression, excessive salivation, and paralysis, is important for preventing transmission to humans. However, early and subtle behavioural changes in animals are often overlooked, yet they can serve as crucial warning signs for timely intervention.

Rabies is primarily transmitted through the bite of an infected animal, with the virus present in saliva. It can also be transmitted through scratches or direct contact of infected saliva with mucosa (e.g. eyes, mouth) or open wounds. The incubation period in animals can range from a few weeks to several months. In dogs and cats, the incubation period is typically less than 6 months, with most cases occurring between 2 weeks and 3 months after exposure. This period can vary depending on the site of the bite and the amount of virus introduced. Animals such as dogs and cats typically develop symptoms and die within a short period; usually 10 to 14 days — once the virus reaches the brain, especially if the animal was already infectious (i.e., the virus was present in its saliva) at the time of the bite. This forms the basis for the standard 14-day observation period for animals following a bite.

In humans, the incubation period varies, with about 75% of cases falling between 3 weeks and

3 months. However, it can be as short as a few days or extend to several months. This relatively long incubation period in the majority offers a critical window for initiating post-exposure prophylaxis (PEP).

Early recognition of symptoms in animals, even when subtle — can trigger timely PEP for exposed individuals, and help prevent human exposure and reduce the spread of the virus within animal populations.

Clinical findings of rabies in animals

Clinical signs of rabies in animals can vary, but there are common neurological symptoms that suggest rabies infection. Regardless of species, the most reliable signs of rabies are:

• Acute behavioural changes: Rabies trigger sudden behavioural shifts, such as aggression in a normally docile animal or increased violence in an already aggressive one.

•**Progressive paralysis:** One of the key features of rabies is the gradual onset of paralysis. This usually begins with difficulty swallowing and progresses to complete paralysis, ultimately leading to death.

Early symptoms of rabies in dogs

The onset of rabies in dogs is marked by subtle behavioural changes. The prodromal phase, lasting approximately 1–3 days, is the first stage of the disease, during which animals exhibit non -specific symptoms that rapidly worsen. Early signs can include:

• Apprehension or nervousness: Dogs may appear uneasy, retreating into corners or seeking isolation.

• Irritability and hyper-excitability: A normally calm dog might suddenly become easily agitated, responding aggressively to minor stimuli.

• Loss of appetite (anorexia): Affected dogs may stop eating, a common indicator of illness.

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• **Behavioural changes:** Dogs may show uncharacteristic aggression, becoming vicious or excessively playful without provocation. Normally nocturnal animals may begin to display daytime activity, which is atypical for their species

These early signs of rabies can be subtle, and owners might mistake them for common behavioural issues or other illnesses. However, as the disease progresses, these symptoms intensify, making it critical to recognize them early. Identifying these signs helps individuals avoid potentially dangerous interactions, reducing the risk of bites.

How rabies progresses in dogs

Rabies progresses through three general phases: the prodromal, acute excitative (furious), and paralytic phases. The speed and severity of these phases can vary depending on the individual animal and the strain of the virus.

1. Prodromal phase: This initial phase lasts 1–3 days and is characterized by nonspecific behavioural changes, such as irritability, nervousness, and decreased appetite. These signs may intensify rapidly.

2. Acute excitative (furious) phase: The furious form of rabies is marked by severe aggression, with animals attacking readily, often without provocation. There is no paralysis during this phase. Infected animals appear anxious, with dilated pupils, and lose their natural fear of humans and other animals. They may roam extensively and attack other animals, humans, or moving objects. Some may ingest non-food objects such as sticks, and stones, or even attempt to chew through fences or cages. Infected dogs may lose their usual fear of humans and other animals, increasing the risk of unpredictable aggression and bites. This phase is often referred to as "mad dog syndrome."

2. Paralytic (dumb) phase: As the virus affects the nervous system, paralysis sets in. Dogs in this stage may exhibit muscle weakness, difficulty swallowing, and excessive salivation. The inability to swallow, coupled with foaming at the mouth, is a hallmark sign of rabies in dogs. The paralysis progresses rapidly, leading to coma and death.

As rabies advances, it can cause severe respiratory issues, making breathing difficult and causing choking sensations. Eventually, the nervous system deteriorates progressively, leading to death. It is essential for pet owners to be vigilant and recognize these signs promptly to seek immediate medical attention.

Behavior of rabid puppies: from subtle signs to severe aggression

In puppies, rabies may not initially present with the pronounced aggression typically observed in adult dogs, making early recognition more difficult. The early symptoms can be subtle. Initially, affected puppies may continue to seek human companionship, appearing overly affectionate or playful. However, within a few hours, this behaviour can escalate to unprovoked biting, even when being petted. As the infection progresses, they may become increasingly irritable and respond aggressively to minimal provocation, using their teeth, claws, or paws to attack. Dilated pupils, alert posture, and anxious expression are also characteristic features. They may lose their natural fear of humans and other animals and react to sound or movement with sudden aggression. In the furious form of rabies, these young animals may roam, attack other animals or people, and even swallow non-food objects such as facces, straws, or stones. Some may attempt to bite through cage wires or follow and snap at hands moving near them. These behavioural changes often go unrecognized in the early phase but rapidly worsen, making rabid puppies a significant public health concern due to their close contact with humans. Early recognition and caution are essential.

Compiled by: Dr Aruni Hathamuna Senior Registrar Epidemiology Unit Ministry of Health

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To be Continued...

Table 1: Sel		elec	ted	noti	otifiable diseases			s reported by Medical			cal	Officers of Health			lth	15 ^{th –} 21 st Mar 2025			(12 th Week)									
	C*	100	100	75	100	100	100	100	100	100	93	100	100	100	100	100	100	100	100	100	100	92	100	100	100	100	100	66
WRCD	*	100	100	100	100	100	92	95	100	100	100	100	100	100	100	14	29	œ	100	92	96	100	94	73	90	100	∞	84
	F	477	249	168	188	37	73	123	42	47	46	10	5	10	9	39	17	19	95	48	71	19	60	27	11	70	31	2095
Tuberculosis	ю			~		e	e	_	0	~	7	e		2	0	e	N	0	~	0	o O	5	4	~	_	80	4	
Tuber	A	47	25	18	12			7		·		,			U	.,		U	10	U	0,		7		1	~	7	190
ania-	В	~	14	~	22	68	0	~	75	27	0	0	0	7	0	~	б О	S	146	7	204	106	6	38	38	,	0	788
Leishmania-	A	0	0	0	0	9	0	0	~	2	0	0	0	~	0	0	2	0	19	0	9	10	2	0	~	0	0	20
		18	34	10	6	~	9	48	4	13	თ	0	တ	9	4	18	9	7	40	26	29	2	25	16	43	23	œ	414
Meningitis	8	~	e	0	2	0	0	5	0	0	0	0	0		0	С	2	0	ო	0		0		0	9	7		36
	A	146	221	179	113	39	67	211	123	113	91	-	12	£	7	20	41	34	236	49	82	50	124	42	135	239	51	2487
Chickenpox	В	20	17	26	12	9	5	20	7	16	7	0		~	0		9	с	20	4	ω	с	8	ო	ω	22	e	235 2
	A	0	0	0	0	0	0	0	0	0	~ ~	0	0	0	0	0	0	0	, -	0	0	0	0	0	0	0	0	5
Rabiies	ш	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	~	0	0	0	0	0	0	0	0	÷
р. Н.	A	5	2	7	S	4	0	4	. 	2	~	-	0	0	0	б		2	. 	~	9	б	÷	4	2	9		83
Viral Hep.	В	~	0	0	0	0	0	0	0	0		0	0	0	0	0	0	0	0	0		. 		0	0	0	0	CJ
	A	4	S	0	19	~	18	27	14	7	232	10	2	2	4	~		Q	16	21	÷	0	œ	19	42	2J		448
Typhus F.	В	0	c	0		0	0			0	18	0		0		0	0	0		ო	0	0	0	0	0		0	31
	A	112	191	174	84	52	40	230	110	136	108	43	12	33	35	28	50	49	223	105	152	64	96	166	426	177	37	2933
ospirosis	в										Ţ																	
Lept	A	5 5	t 12	3 12	с 8	33	с т	3 28	3	3 13	_	2 4	- 2	3	0 2	6	2 6	() T	9 14	4 5	7 3	2	6 0	3 21	3 45	12	7 0	222
F. Poisoning	В		44	13	~	14	44	26			11		~	20	Ŭ	70		16	19	7			Ŭ		13	15	1-	352
F. Poi	A	0	2	0	2	œ	0	0	0	0	0	~	0	0	0	10	0	0	0	0	0	~	0	0	~	~	0	26
ever	в	с	~	2	4	0	4	0	0	~	С	4	0	~	~	0	0	0	~	0	~	~	S	0	e	0	0	33
En. Fever	A	0	0	0	5	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	8
Encephalitis	В	ŝ	14	~	2	~	с С	2	2	2	-	0	0	0	0	00	2	~	7	0	5	2	2	ŝ	4	ŝ	0	68
Ence	A	0	~	0	0	0	0	0	0	0	0	0	0	0	0	0	~	0	က	0	0	0	~	0	0	0	0	9
itery	В	1	15	10	19	Ø	24	19	9	4	24	5	0	5	~	62	9	19	7	0	18	7	0	9	28	23	7	352
Dysentery	A	0	-	0	-	0	-	~	0	0	2	0	0	0	0	7	0	0	0	~	4	0	က	0	~	7	0	52
ever		2849	1820	457	775	427	56	529	248	432	442	39	77	24	24	692	48	322	305	259	249	68	218	237	624	348	144	11713
Dengue Fever	A B	242	125	35	59	20	9	35	18	30	25	0	2	~	0	28	ო	23	21	10	œ	9	9	24	56	32	9	821
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RDHS		Colombo	Gampaha	Kalutara	Kandy	Matale	Nuwara Eliya	Galle	Hambantota	Matara	Jaffna	Kilinochchi	Mannar	Vavuniya	Mullaitivu	Batticaloa	Ampara	Trincomalee	Kurunegala	Puttalam	Anuradhapura	Polonnaruwa	Badulla	Monaragala	Ratnapura	Kegalle	Kalmunai	SRILANKA

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Table 2: Vaccine-Preventable Diseases & AFP

22nd – 28th Mar 2025

15th - 21st Mar 2025 (12th Week)

Disease	No. o	f Case	s by F	Provinc	:e			Number of cases during current	Number of cases during same	Total number of cases to date in	Total num- ber of cases to date in	Difference between the number of cases to date			
	W	С	S	Ν	Е	NW	NC	U	Sab	week in 2025	week in 2024	2025	2024	in 2025 & 2024	
AFP*	00	00	00	00	00	00	00	00	00	01	02	15	18	-16.6%	
Diphtheria	00	00	00	00	00	00	00	00	00	00	00	00	00	0 %	
Mumps	01	01	00	00	00	00	01	00	04	07	05	63	69	-8.7 %	
Measles	00	00	00	00	00	00	00	00	00	00	10	01	168	-99.4%	
Rubella	00	00	00	00	00	00	00	00	00	00	00	00	01	-100%	
CRS**	00	00	00	00	00	00	00	00	00	00	00	01	00	0 %	
Tetanus	00	00	00	00	00	00	00	00	00	00	00	02	00	0 %	
Neonatal Tetanus	00	00	00	00	00	00	00	00	00	00	00	00	00	0 %	
Japanese Enceph- alitis	00	00	00	00	00	00	00	00	00	00	00	04	01	300 %	
Whooping Cough	00	00	00	00	00	00	00	00	00	00	00	08	01	700 %	

Key to Table 1 & 2

Provinces: W: Western, C: Central, S: Southern, N: North, E: East, NC: North Central, NW: North Western, U: Uva, Sab: Sabaragamuwa.

RDHS Divisions: CB: Colombo, GM: Gampaha, KL: Kalutara, KD: Kandy, ML: Matale, NE: Nuwara Eliya, GL: Galle, HB: Hambantota, NT: Matara, JF: Jaffna,

KN: Killinochchi, MN: Mannar, VA: Vavuniya, MU: Mullaitivu, BT: Batticaloa, AM: Ampara, TR: Trincomalee, KM: Kalmunai, KR: Kurunegala, PU: Puttalam, AP: Anuradhapura, PO: Polonnaruwa, BD: Badulla, MO: Moneragala, RP: Ratnapura, KG: Kegalle.

Data Sources:

Weekly Return of Communicable Diseases: Diphtheria, Measles, Tetanus, Neonatal Tetanus, Whooping Cough, Chickenpox, Meningitis, Mumps., Rubella, CRS, Special Surveillance: AFP* (Acute Flaccid Paralysis), Japanese Encephalitis

CRS** =Congenital Rubella Syndrome

NA = Not Available

Number of Malaria Cases Up to End of March 2025, 04 All are Imported!!!

Comments and contributions for publication in the WER Sri Lanka are welcome. However, the editor reserves the right to accept or reject items for publication. All correspondence should be mailed to The Editor, WER Sri Lanka, Epidemiological Unit, P.O. Box 1567, Colombo or sent by E-mail to chepid@sltnet.lk. Prior approval should be obtained from the Epidemiology Unit before publishing data in this publication

ON STATE SERVICE

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