For Veterinary Use Only

READ ALL INSTRUCTIONS BEFORE BEGINNING THE TEST

KIDX™ CHW Ag Test Kit

[CAT No. CGM-CHG-11]

# Introduction

Canine Heartworm (CHW), *Dirofilaria immitis*, is a parasitic mosquitoborne nematode that causes canine dirofilariasis worldwide<sup>1</sup>. Adult *D. immitis* lives in the right side of the heart, the pulmonary artery, and associated blood vessels of infected dogs, causing pulmonary hypertension, congestive heart failure, and damage to other organs<sup>2</sup>. *D. immitis* harbors intracellular bacteria named *Wolbachia pipientis*, a Gram-negative bacteria. *Wolbachia* has the potential to play an important role in the pathogenesis and immune response to filarial infection<sup>1, 3</sup>. The immunopathology of filarial disease is extremely complex and the clinical manifestations of infection are strongly dependent on the type of immune response elicited by the parasite<sup>1</sup>. Dogs show a mild persistent cough, reluctance to exercise, fatigue after

moderate activity, decreased appetite, and weight loss at the early stage of the infection<sup>1, 2</sup>. If CHW infected dogs are not treated, large numbers of heartworms can develop a sudden blockage of blood flow within the heart leading to a life-threatening form of cardiovascular collapse.

CHW is a zoonotic parasite and is transmitted by mosquitoes. *D. immitis* is responsible for human pulmonary dirofilariasis throughout the world<sup>3, 4</sup>.

# • Principle

The RIDX<sup>™</sup> CHW Ag Test Kit is a lateral flow chromatographic immunoassay for the qualitative detection of canine heartworm antigens in canine blood. This kit shows two letters which are the test (T) line and the control (C) line on the surface of the device. If the CHW antigen exists in the sample, it binds to the gold-conjugated anti-canine CHW antibody. The antigen–antibody complex moves through the membrane by capillary force and responds to the CHW monoclonal antibody on the test line, resulting in a red line. The control line indicates that the test is performed correctly and should appear when the test is complete.

The highly selective and sensitive monoclonal antibody to *D. immitis* is used as capture in the kit. The RIDX<sup>™</sup> CHW Ag Test Kit can detect CHW antigens in canine blood with high accuracy.

### Performance

1. Clinical Sensitivity & Clinical Specificity

		I	Disease Status	3
		+	-	Total
RIDX™ CHW Ag Test	+	53	0	53
	-	2	30	32
	Total	55	30	85
Clinical Sensitivity: 96.36% (53/55, 95% CI: 97.68% ~ 99.00%)				
Clinical Specificity: 100% (30/30, 95% CI: 88.65% ~ 100%)				

Diagnostic Accuracy: 97.65% (83/85, 95% CI: 91.82% ~ 99.35%)

\* CI: Confidence Interval

2. Male and female *D. immitis* can be diagnosed.

### 3. Cross-Reactivity

Below potential cross-reactivity substances did not affect the performance of the RIDX^{\rm TM} CHW Ag Test Kit.

Pathogen	Titer	Result
Canine coronavirus	1 x 10 <sup>6</sup> TCID <sub>50</sub> /mL	Negative
Canine distemper virus	1 x 10 <sup>5</sup> TCID <sub>50</sub> /mL	Negative
Canine influenza virus	1 x 10 <sup>6</sup> EID <sub>50</sub> /mL	Negative
Canine parvovirus	1 x 10 <sup>6</sup> TCID <sub>50</sub> /mL	Negative
Escherichia coli	3.56 x 10 <sup>8</sup> CFU/mL	Negative
<i>Giardia</i> spp.	1.42 x 10 <sup>7</sup> <i>G</i> . cysts/100µL	Negative
<i>Salmonella</i> spp.	1 x 10 <sup>6</sup> CFU/mL	Negative

#### Kit Components

	Component	Number/Kit
1	CHW Ag Test device	10
2	Anticoagulant tube	10
3	Disposable dropper	10
4	Instructions for use	1

#### Storage & Stability

1. Store the test kit at 2~30°C (35.6~86°F). Do NOT freeze.

2. Do not store the test kit in direct sunlight.

3. The test kit is stable within the expiration date marked on the package label.

# Sample Preparation

#### [Whole blood]

1. Collect 1mL (0.5~1.5mL) of the whole blood sample and put it into an anticoagulant tube.

2. Close the cap on the anticoagulant tube and invert the tube 5 times to mix blood sample and EDTA.



3. The Anticoagulated whole blood samples should be used immediately after collection. If you cannot use the samples immediately, store them refrigerated (2~8°C/35.6~46°F) or keep them on ice. Do not freeze anti-coagulated whole blood samples. If you cannot use the samples within 24 hours, store them in a form of serum or plasma.

#### [Serum or plasma]

1. Prepare serum and plasma using a standard procedure of clinical laboratory.

2. Serum or plasma, either fresh or stored at  $2 \sim 8^{\circ}$ C ( $35.6 \sim 46^{\circ}$ F) for up to 72 hours, can be used. For longer storage, freeze at  $-20^{\circ}$ C ( $-4^{\circ}$ F).

# Test Procedure

1. All test components and samples must be at room temperature (15~30°C/59~86°F) before use.

2. Take canine blood sample (the anticoagulated whole blood, serum or plasma) using a dropper.

- 3. Add 2 drops of the sample solution into the sample hole (S).
- 4. Read test result at 10 minutes.



[Summary of Test Procedure]

# Interpretation of Results

#### 1. Positive result

Test (T) line and control (C) line within the result window indicate the presence of CHW antigens.



#### 2. Negative result

Only control (C) line appears in the result window.



# 3. Invalid results

If the control (C) line does not appear, the result might be considered invalid. The sample should be retested.



#### Precautions

1. This test kit is for veterinary *in vitro* diagnosis only especially canine. Do not use this test kit for other animals.

2. The test device is sensitive to humidity and heat. Use the test device within 10 minutes after removing the foil pouch.

3. Do not touch the membrane of the test device.

4. Do not use the test device if the foil pouch is damaged or the seal is open.

5. Do not use an expired test kit. The expiration date is marked on the package label.

6. Do not reuse the test components (device, dropper, anti-coagulant tube).

7. Do not mix components from different lot numbers because the components in this kit have been quality control tested as a standard batch unit.

8. Decontaminate and dispose of all samples, used kits, and potentially contaminated materials in the accordance with national and local regulations.

9. All samples should be handled as being potentially infectious. Wear protective gloves while handling samples. Wash hands thoroughly afterward.

# • References

1. McCall JW, Genchi C, Kramer LH, Guerrero J, Venco L. Heartworm Disease in Animals and Humans. *Adv Parasitol.* 2008; 66: 193–285.

2. Hoch H, Strickland K. Canine and feline dirofilariasis: prophylaxis, treatment, and complications of treatment. *Compend Contin Educ Pract Vet.* 2008; 30: 146–151.

 Simón F, Siles-Lucas M, Morchón R, González-Miguel J, Mellado I, Carretón E, Montoya-Alonso JA. Human and animal dirofilariasis: the emergence of a zoonotic mosaic. *Clin Microbiol Rev.* 2012; 25: 507–544.
Dantas-Torres F, Otranto D. Dirofilariosis in the Americas: a more virulent Dirofilaria immitis? *Parasit Vectors.* 2013; 6(1): 288.

# Symbol Descriptions

LIC	License number
CAT	Catalogue number
LOT	Batch code, Lot number
[]i	Consult instructions for use
$\sum_{n}$	Contains sufficient for $\langle n \rangle$ tests
2	Do not reuse
IVD	In vitro diagnostic medical device
	Temperature limitation
$\bigcirc$	Do not use, if the package is damaged
11	Upper side
	Manufacturer



SKYER, INC. #532, 416, Hwagok-ro, Gangseo-gu, Seoul, 07548, Republic of Korea TEL: +82-2-706-6801, FAX: +82-50-4096-6988 Technical Support: marketing@skyer.co.kr www.skyerdiagnostics.com

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