

# HaoKebio™ circRNA FISH Kit(Animal)

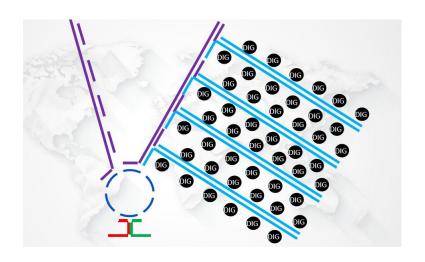
## Catalog Number: HKR13D-1

### **[Product Information]**

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Product Name	Catalog No	30T	Storage	Shelf Life
Wash Buffer (powder, dissolve in 10L ddH2O)	HKR13-1	10L		
Proteinase K (100x)	HKR13-2	30uL		
HRP-Mouse Anti-Digoxin (100x)	HKR13-3	30uL		
TSA Chromogenic Solution 570nm (Optional)	HKR13-4	3mL	Short-term:4°C  Long-term:-20°C	12 months
Blocking Buffer	HKR13-5	3mL		
Probe:	HKR13-6	3mL		
Pre-Amplification Probe	HKR13-7	3mL		
Amplification Probe	HKR13-8	3mL		

## **[Product Description]**

The circRNA hybridization kit includes a combination probe, a pre amplification structure, and an amplification probe. When two probes bind to adjacent positions of the target sequence, a small segment of the top sequence can bind to the pre amplification structure. The repeated sequence on the pre amplification structure can trigger a branched HCR to form a large nucleic acid aggregate. Each nucleic acid strand is labeled with digoxin, and the antibody recognizes digoxin for color development.



## **【Probe Information】**

## **Tissue Fixation**

Tissue Type	Animal	Plant	Frozen	Cell Climb	Cells
			Samples	Slides	
Treatment	Fix at RT	Vacuum fix	Dehydrate in	Fix at 4℃	scrape off
	for 12h,	for 1h, RT fix	15% sucrose	for 2h.	cells,fix in
	paraffin	for 12h,	at 4℃ for		4% PFA at
	embed	paraffin	8h,then in		4°C for 2h,
		embed.	30% sucrose		wash with
			at 4℃ for		PBS, agarose
			8h, OCT		embed.
			embed.		
Туре	mRNA	lncRNA	circRNA	miRNA	rRNA
Treatment	Fix at RT for	Fix at RT for	Fix at RT for	Fix at RT for	Fix at RT for
	12h	12h (<300bp:	12h	12h	12h (<300bp:
		24h)			24h)

## **[Storage and Shipping]**

Ship on wet ice; store at -20  $^{\circ}$ C for long-term or at 4  $^{\circ}$ C for short-term use. Shelf life: 6

months.

### [Protocol]

#### 1. Deparaffinization and Rehydration

Immerse slides sequentially in:Xylene I (15 min)  $\rightarrow$  Xylene II (15 min)  $\rightarrow$  Xylene III (15 min)  $\rightarrow$  100% ethanol (10 min)  $\rightarrow$  90% ethanol (10 min)  $\rightarrow$  80% ethanol (10 min)  $\rightarrow$  70% ethanol (10 min)  $\rightarrow$  Rinse with distilled water.

#### 2. Enzyme Repair

After slides are completely dry, draw a hydrophobic circle around the tissue using a histology pen (recommended: HKR14P In Situ Hybridization Pen). Place slides horizontally in a hybridization oven or humidified chamber. Add 100  $\mu$ L of Proteinase K repair solution (1X) onto the tissue and incubate at 37 °C for 30 min (15 min for cell samples). Rinse with distilled water to stop the reaction.

#### 3. Blocking

Remove excess liquid from slides. Add 100 μL of Blocking Buffer per slide and incubate at 37°C for 30 min. Wash once for 5 min. Wash steps: Place slide racks in a wash tank, add wash buffer (ensure samples are submerged), and shake at 60 rpm for 5 min.

#### 4. Probe Hybridization

Remove excess liquid from slides. Add 100 μL of probe per slide and incubate at 37 °C for 3 h or overnight (maintain humidity to prevent drying). Wash 5 times, 5 min each. Wash steps: As described above.

#### 5. Pre-Amplification Hybridization

Remove excess liquid from slides. Add 100  $\mu$ L of Pre-Amplification Probe to each slide and incubate at 37  $^{\circ}$ C for 3 h or overnight (ensure humidity). Wash 5 times, 5 min each. Wash steps: As described above.

#### 6. Probe Amplification

Remove excess liquid from slides. Add 100  $\mu$ L of Amplification Probe per slide and incubate at 37  $^{\circ}$ C for 1.5 h (maintain humidity). Wash 5 times, 5 min each. Wash steps: As described above.

#### 7. HRP-Mouse Anti-Digoxin

Remove excess liquid from slides. Add 100  $\mu$ L of HRP-Mouse Anti-Digoxin (1X) per slide and incubate at 37  $^{\circ}$ C in a humidified chamber for 40 min. Wash 5 times, 5 min each. Wash steps: As described above.

#### 8. Chromogenic Reaction

Remove excess liquid from slides. Add  $100~\mu L$  of TSA Chromogenic Solution per slide and incubate at RT for 10~min. Rinse with distilled water to stop the reaction.

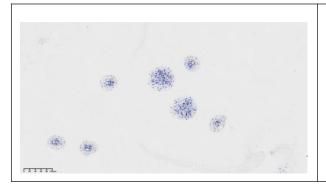
#### 9. DAPI Staining

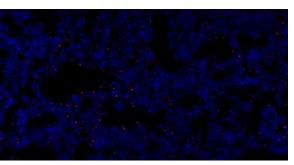
Add 50  $\mu$ L of DAPI staining solution per slide, incubate in the dark for 5 min, rinse with distilled water, and mount with anti-fade mounting medium.

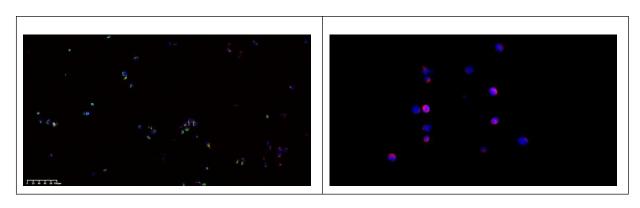
### **[Precautions]**

- 1. For research use only.
- **2.** Wear lab coats and disposable gloves for safety.

#### **Example Images**







## **【Source of Reagents】**

发表[中文论文]请标注:circRNA FISH Kit(Animal)(HKR13D-1)由杭州浩克生物技术有限公司提供; 发表[英文论文]请标注:circRNA FISH Kit(Animal)(HKR13D-1) were kindly provided by Hangzhou Hao ke Biotechnology Co., Ltd.