

# Sample Preparation for Fixed Frozen Tissue using the RNAscope® Fluorescent Multiplex Assay

## Introduction

This Technical Note is for customers who wish to run the RNAscope® Fluorescent Multiplex Assay (Cat. No. 320850) with fixed frozen tissues. The required pretreatment reagents are RNAscope® Target Retrieval and RNAscope® Protease III (available in the RNAscope® Universal

Pretreatment Kit, Cat. No 322380). Refer to the user Safety Data Sheet (SDS) available on the ACD website. Materials required, but not provided by ACD, include sucrose, 1X PBS, OCT media, 100% EtOH, Superfrost® Plus slides (Fisher), and 4% paraformaldehyde (4% PFA).

## Workflow

### Part 1: Prepare the Tissue Sections

#### Fix Sample

1. If needed, perfuse the tissue with freshly prepared 4% paraformaldehyde (PFA) in 1X PBS or go directly to step 2.
2. Dissect tissue and immersion fix in freshly prepared 4% PFA for **24 HRS** at **4°C**.

#### Freeze Tissue

1. Immerse the tissue in 10% sucrose in 1X PBS at **4°C** until the tissue sinks to the bottom of the container (approximately **18 HRS** for brain tissue).

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**Note:** The time needed for the tissue to sink varies with tissue type and size.

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2. Immerse the tissue in 20% sucrose in 1X PBS at **4°C** until the tissue sinks to the bottom of the container.
3. Immerse the tissue in 30% sucrose in 1X PBS at **4°C** until the tissue sinks to the bottom of the container.
4. Freeze the tissue in OCT (Optimal Cutting Temperature) embedding media or TFM (Tissue Freezing Media) on crushed dry ice or iso-pentane or liquid nitrogen.
5. Store tissue blocks in an airtight container at **-80°C**.

#### Prepare Sections

1. Before sectioning, equilibrate the tissue blocks at **-20°C** for at least **1 HR** in a cryostat.
2. Section the blocks by cutting sections to a thickness of 7–15 µm. Mount the sections on SuperFrost® Plus slides (Fisher Scientific # 12-550-15).

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**IMPORTANT!** Use SuperFrost® Plus slides only. Other slide types may result in tissue loss.

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3. Air dry the slides for **2 HR** at **-20°C** and overnight at **-80°C**. If all the slides are not used immediately, store them at **-80°C** for up to **3 MONTHS**.
4. On the day of performing the RNAscope® assay, wash the slides with 200 mL 1X PBS in a Tissue-Tek® slide rack for **5 MIN** while moving the rack up and down to remove OCT.
5. Bake the slides for **30 MIN** at **60°C**.
6. Post-fix the slides by immersing them prechilled 10% NBF or 4% PFA in 1X PBS for **15 MIN** at **4°C**.

#### Dehydrate the tissue

1. Prepare 200mL 50% EtOH, 200 mL 70% EtOH, and 400 mL of 100% EtOH.
2. Remove the slides from the 10% NBF or 4% PFA, and immerse them in 50% EtOH for **5 MIN** at **RT**.
3. Remove the slides from 50% EtOH, and immerse them in 70% EtOH for **5 MIN** at **RT**.

4. Remove the slides from 70% EtOH, and immerse them in 100% EtOH for **5 MIN** at **RT**.
5. Remove the slides from 70% EtOH, and immerse them in 100% EtOH for **5 MIN** at **RT**.

### **Dry the slides**

1. Remove slides from 100% EtOH, and let them air dry for **5 MIN** at **RT**.

## **Part 2: Tissue Pretreatment**

### **Prepare Materials**

1. Bring the HybEZ™ Oven to **40°C**.
2. Place a wet humidifying paper in the Humidity Control Tray, leaving the EZ-Batch™ Slide Holder on the bench. Re-insert the covered tray into the oven and close the oven door. The tray should be pre-warmed for at least **20 MIN** before use.
3. Prepare 700 mL fresh 1X Target Retrieval in a beaker. Cover with foil, bring to a mild boil, and maintain uniform boiling at **99–100°C**. Do not boil for more than **30 MIN** before use.

### **Apply RNAscope® Target Retrieval**

1. With a pair of forceps *very slowly* submerge a slide rack containing the slides into boiling 1X Target Retrieval solution. Keep the slides in the solution for **5 MIN** only. Monitor the temperature closely and make sure it stays at 98–102°C

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**Note:** Depending on the tissue type, boiling time may need to be adjusted.

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2. *Immediately* transfer the hot slide rack to a staining dish containing distilled water at **RT**.
3. Wash the slides in distilled water by moving the rack up and down 3–5 times. Repeat with fresh distilled water.

4. Rinse the slides in fresh 100% EtOH by moving the slides up and down 3–5 times. Air dry.

### **Create Barrier**

1. Draw 2–4 times around the tissue using the Immedge™ hydrophobic barrier pen. Let the barrier dry completely **~1 MIN** or **OVERNIGHT** at **RT**.

### **Apply RNAscope® Protease III**

1. Place the slides in the EZ-Batch™ Slide Holder and add 2–4 drops of Protease III to each section. Use enough solution to completely cover the sections.
2. Place the EZ-Batch™ Slide Holder in the pre-warmed HybEZ™ Humidity Control Tray. Seal the tray and insert back into the HybEZ™ Oven. Incubate for **30 MIN** at **40°C**.
3. Wash the slides in the clear EZ-Batch™ Wash Tray by submerging the slide holder in the distilled water.
4. Wash the slides with slight agitation. Repeat with fresh distilled water.

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**IMPORTANT!** Proceed to the RNAscope® protocol using the *RNAscope® Fluorescent Multiplex Kit User Manual Part 2* (Catalog No. 320293) available at <http://www.acdbio.com/technical-support/user-manuals>.

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### **Obtaining Support**

For the latest services and support information, go to: <https://acdbio.com/technical-support/support-overview>.

At the website, you can:

- Access telephone and fax numbers to contact Technical Support and Sales
- Search through FAQs
- Submit a question directly to Technical Support

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