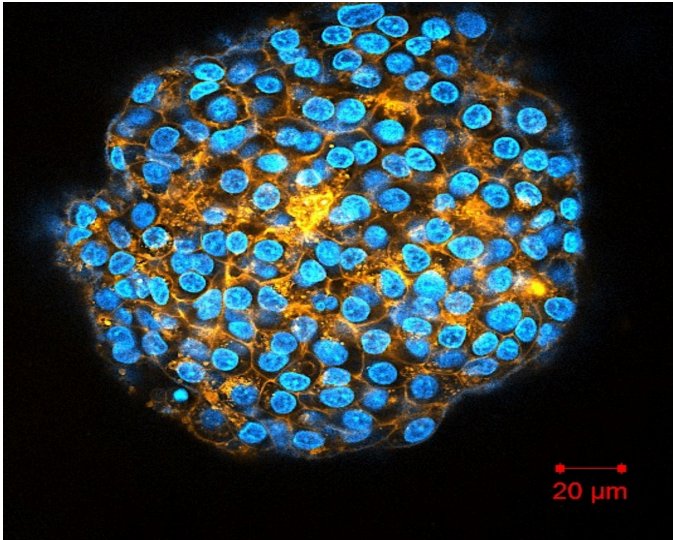


# Cello-IF: The Smarter Way to Label

All-in-one Reagent for Immunofluorescence Labelling of Organoids, Spheroids and Cells



Liver Cancer Spheroid labelled using Cello-IF technology - Tok et al (2022)

Simplify the steps of immunofluorescence labelling and maximize your results. The Cello-IF innovative technology allows you to stain organoids, spheroids or cells in just 8 hours, while still in hydrogel or extracellular matrix, preserving delicate structures and cellular integrity.

This all-in-one reagent removes the need for harvesting, clearing, transferring and centrifuging, preventing sample loss and ensuring consistent accurate results.

## Are you having issues with unsuccessful labelling of precious samples?

### Features

- High quality images in hours, not days.
- Reduced number of reagents and steps.
- Label organoids, spheroids, and cells while still in hydrogel or extracellular matrix.
- Reduces background noise for high-resolution images.
- Boosted antigenicity for clearer visualization of target.

### Benefits

- Faster Results.
- Streamlined Workflows.
- Effortless labelling.
- Enhanced Clarity.
- Optimize detection.

Request a free sample: [info@amsbio.com](mailto:info@amsbio.com)

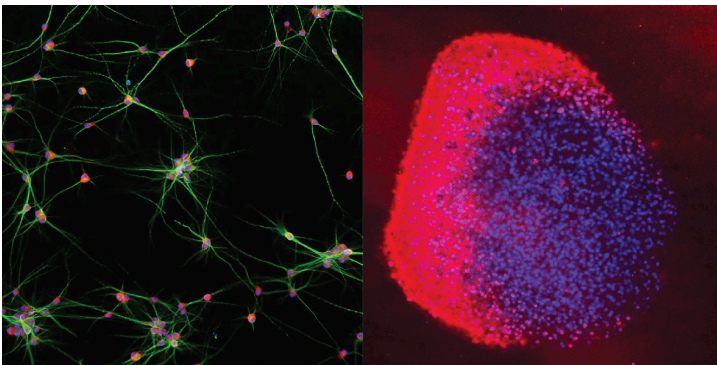
## PROTOCOL FOR USE

1. Fix in 4% paraformaldehyde (PFA) for 1 hr at 37°C.
2. Rinse with PBS X3, 10 min at 37°C.
3. Incubate for 1.5-2 hrs at 37°C with primary antibodies diluted in Cello-IF.
4. Wash with Cello-IF X3, 10 min at 37°C.
5. Incubate for 1.5-2 hrs with secondary antibodies diluted in Cello-IF.
6. Wash with PBS X3, 10 min at 37°C.
7. Mount & image under microscope.



|                          | Traditional Workflow | Workflow with CELLO-IF |
|--------------------------|----------------------|------------------------|
| Reagents                 | >3                   | 1                      |
| Steps                    | 24                   | 8                      |
| Transferring & Pipetting | >3                   | 0                      |
| Time                     | ~44 hours            | 8 hours                |

Available in two formats for labelling both 3D and 2D samples producing high quality images.



**"We really didn't think imaging the neuronal organoids would be as easy because they are so fragile and difficult to section with traditional methods. Using Cello™-IF simplified our workflow and it has great potential to make high-throughput imaging of organoids feasible. We got beautiful images using Cello-IF on both 2D and 3D cultures."**

*Didem Demirbas, PhD - Jenny Lai, MD-PhD Candidate, Boston Children's Hospital, Harvard Medical School*

*Left: Induced Pluripotent Stem Cell Derived Neurons (MAP2 - green, CREB - red, DAPI - blue).*

*Right: Brain Organoid (PAX2 - red, DAPI - blue).*

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