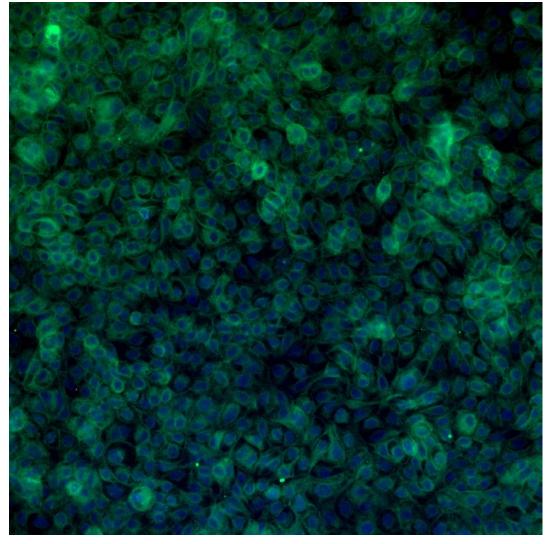


IPEC cells (Pig)

The reference antibody only works on mouse ileum, whereas it has been tested on pig jejunum and ileum, on a piglet cell line (IPEC) and on bovine mammary gland.

Recombinant monoclonal anti-Occludin Abcam Ab216327 1/400e



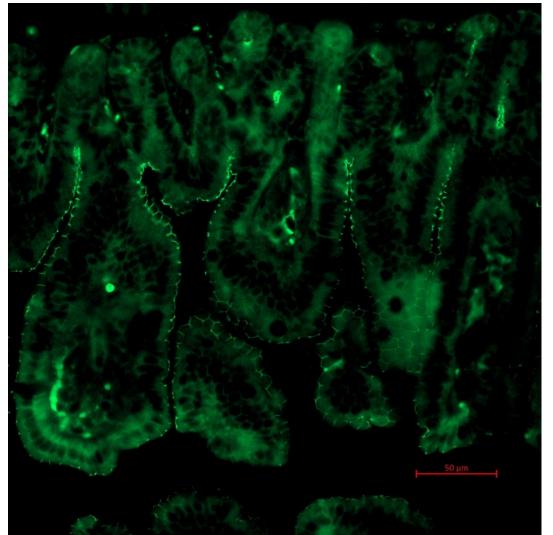


There is signal (green).

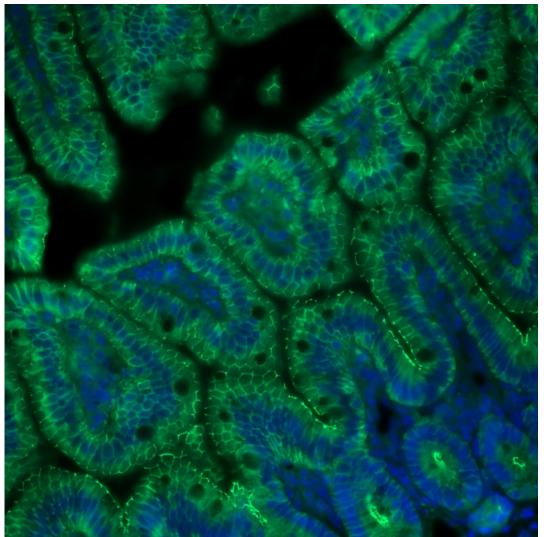


Mouse intestine

Reference polyclonal anti-Occludin 1/200e



Recombinant monoclonal anti-Occludin Abcam Ab216327 1/400e





The recombinant antibody detects Occludin (in green) on mouse ileum with a higher dilution than the control.

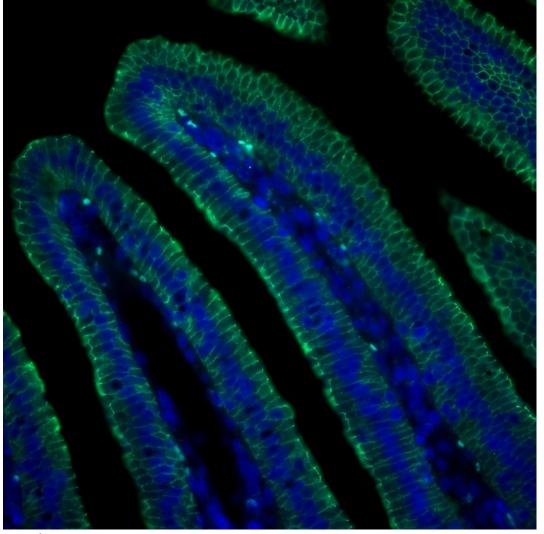






The reference antibody only works on mouse ileum, whereas it has been tested on pig jejunum and ileum, on a piglet cell line (IPEC) and on bovine mammary gland.

Recombinant monoclonal anti-Occludin Abcam Ab216327 1/400e





There is signal (green).

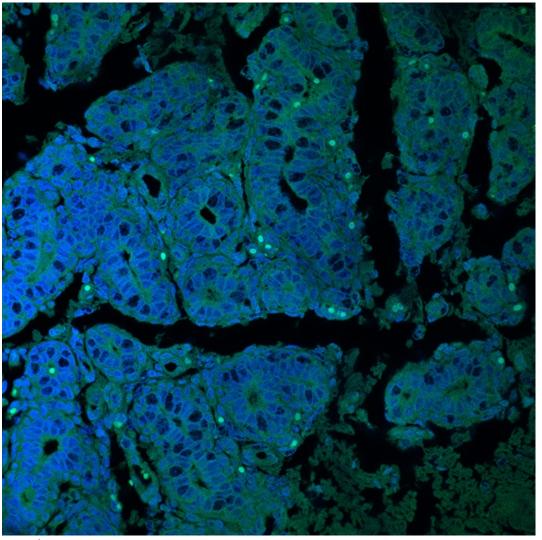


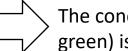
Bovine mammary gland



The reference antibody only works on mouse ileum, whereas it has been tested on pig jejunum and ileum, on a piglet cell line (IPEC) and on bovine mammary gland.

Recombinant monoclonal anti-Occludin Abcam Ab216327 1/400e



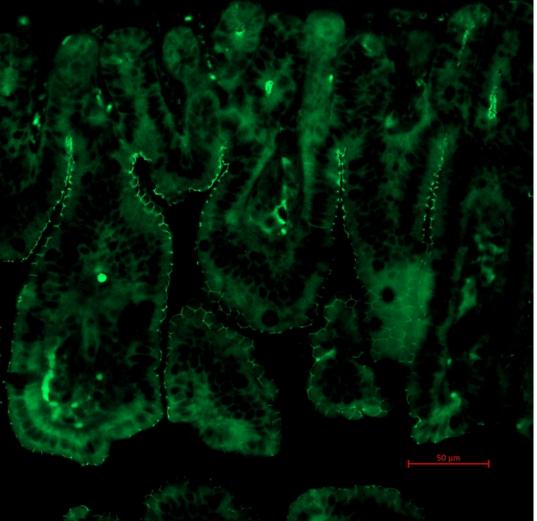


The conditions need to be optimised, the labelling (in green) is not clear, there is background noise and 'dots'.

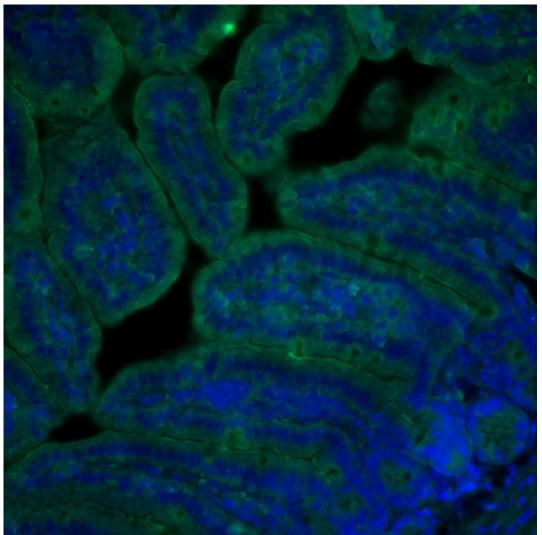


Mouse intestine

Reference polyclonal anti-Occludin 1/200e



Recombinant monoclonal anti-Occludin Absolute Antibody AbAb 37-5



No signal



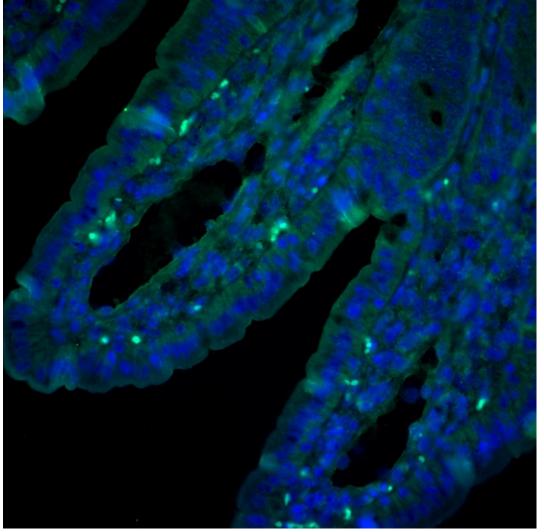


Pig intestine



The reference antibody only works on mouse ileum, whereas it has been tested on pig jejunum and ileum, on a piglet cell line (IPEC) and on bovine mammary gland.

Recombinant monoclonal anti-Occludin Absolute Antibody AbAb 37-5





There is a signal (green), but conditions need to be optimised: background noise, 'small dots' labelling.

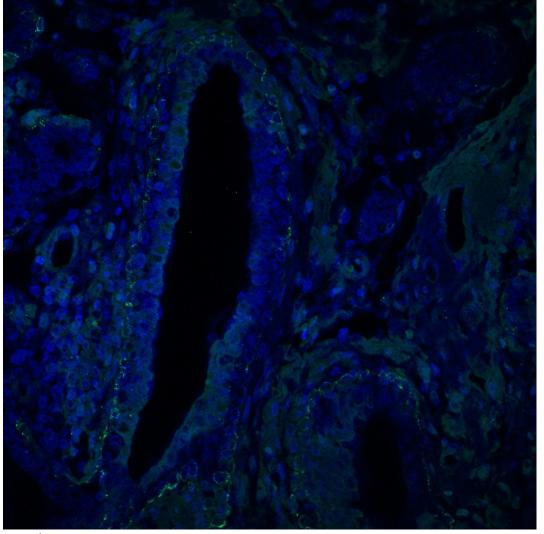


Bovine mammary gland



The reference antibody only works on mouse ileum, whereas it has been tested on pig jejunum and ileum, on a piglet cell line (IPEC) and on bovine mammary gland.

Recombinant monoclonal anti-Occludin Absolute Antibody AbAb 37-5





Conditions need to be optimised; labelling (in green) is very weak.

Materials and Methods

Experimental Strategy

The histology platform from Inrae St Gilles evaluated four recombinant Fc Rabbit anti-Occludin antibodies on formalin-fixed cells and paraffin-embedded (FFPE) tissue sections from pig and cow.

As no other antibody worked for these species (no positive control), all antibodies were tested on mouse tissue (internal & biological positive control).

Antibody performance was assessed based on staining intensity, specificity, and background signal. Validation criteria in tissues included a clear intense staining localized at the tight junctions.

Biological Material

All formalin-fixed paraffin-embedded (FFPE) samples were already available on the platform. No animal has been sacrificed for these tests.

- Intestinal porcine epithelial cells-jejunum 2 (IPEC-J2)
- Tissues:
 - o Paraffin-embedded sections (6 µm thick) of:
 - Mouse jejunum (from experiment "Candy S36 ileon").
 - Piglet jejunum (samples P22001, P117).
 - Heifer mammary gland (sample G4297).
- Sample preparation:
 - o Fixed in neutral buffered formalin, dehydrated, and embedded in paraffin.
 - o Sections mounted on slides and stored at 4°C until use.

Antibody Testing

- Reference antibody: Invitrogen 33-1500 anti-occludin monoclonal (mouse, clone OC-3F10)
- Abcam Ab216327 anti-Occludin (rabbit recombinant monoclonal Occludin antibody)
- Absolute Antibody AB 04036 AbAb 1-3 anti-Occludin (rabbit) recombinant
- Absolute Antibody AB 04037 AbAb 37-5 anti-Occludin (rabbit) recombinant
- Cell Signaling Technology #91131 E6B4R anti-Occludin (Rabbit) recombinant

Secondary Antibody

Goat anti-rabbit IgG Alexa Fluor 488 (Life Technologies, #A11008), diluted 1:500 in blocking buffer.

• Staining protocol (FFPE samples)

The recombinant anti-Occludin antibodies were tested under these conditions:

1. **Deparaffinization**:

o Automated process using the Gemini staining system:

- Xylene (10 min × 2) \rightarrow Ethanol series (100%, 90%, 80%, 70% 5 min each) \rightarrow Distilled H₂O (5 min) \rightarrow PBS 1X (5 min × 2).
- o Slides stored overnight in PBS 1X at 4°C.

2. Epitope Retrieval:

- Pressure cooker (Diagomics "cocotte") with preheated EDTA buffer (commercial, Diagomics).
- o Program: 110°C, 15 min (low-pressure mode).

3. **Blocking**:

1 h incubation with blocking buffer: PBS 1X + 10% horse serum + 3% Triton X-100.

4. Primary Antibody Incubation:

- o 50 μl/section, 1.5 h at 37°C.
- o Two dilutions tested for each antibody: 1:200 and 1:400 (prepared in blocking buffer).

5. Secondary Antibody Incubation:

o 50 μl/section, 30 min at room temperature.

6. Nuclear Counterstaining:

Mounting medium with DAPI (Gold Antifade Mountant with DAPI, 13 μl/section).

7. Reagent Details

Blocking buffer: PBS 1X (20 mM Tris-base, 150 mM NaCl, pH 7.6) + 10% horse serum + 3% Triton X-100.

Antibody diluent: Blocking buffer for primaries; PBS 1X + 0.2% BSA for secondaries.

Mounting medium: ProLong Gold Antifade with DAPI (Thermo Fisher, #P36935).

• Staining protocol (cells)

Pre-Fixation of plates with 4% PAF for 30 minutes. Then rinse them twice for 10 minutes with PBS before adding PBS until the day of immunostaining.

The day of immunostaining

1. Fixation:

10 min with Methanol (1ml in the insert + 1 ml in the well)

2. Permeabilization

5 min with permeabilization solution : PBS 1X + 0.5% Triton X100 (1ml in the insert + 1 ml in the well)

3. Primary Antibody Incubation:

- a. 100 ul/insert, 1.5 h at 37°C.
- b. Two dilutions tested for each antibody: 1:200 and 1:400

4. Secondary Antibody Incubation:

- a. 100 µl/insert, 1 h at 37°C.
- 5. Cutting the insert

6. Nuclear Counterstaining:

a. Mounting medium with DAPI (Gold Antifade Mountant with DAPI, 13 µl/insert).

7. Reagent Details

Permeabilization buffer: PBS 1X (20 mM Tris-base, 150 mM NaCl, pH 7.6) + 0.5 % Triton X-100.

Antibody diluent: PBS 1X + 0.2% BSA for primaries and secondaries.

Mounting medium: ProLong Gold Antifade with DAPI (Thermo Fisher, #P36935)

Image Acquisition

- **Microscope**: Zeiss ApoTome Imager 2 with Zen acquisition software.
- Settings:
 - o Objective: 40x oil immersion.
 - o Fluorescence filters: Alexa Fluor 488 (ex/em 495/519 nm) and DAPI (ex/em 358/461 nm).
 - o Z-stack imaging with Apotome optical sectioning to reduce background noise.
- Analysis:
 - o Three slides per tissue type (mouse, piglet, heifer) imaged.
 - o Staining intensity and localization assessed for Occludin at tight junctions.