Preparation of samples for histological labelling

It is possible to make histological sections as well as perform Hematoxylin-Eosin-Saffron staining and/or immunohistochemical reactions on BIOMIMESYS®.

This protocol can be used with any cell type grown in BIOMIMESYS®. It uses standard reagents for making a histological section but also more specific reagents such as OTTIX and OTTIX shaper (DiaPath) and Entellan® (Millipore).

- **Sample Fixation**
  - Fix the samples using alcohol, formaldehyde and acetic acid (4%)
  - Fix these samples for 24h at an ambient temperature.

- **Paraffin cuts**
  - Dehydrate the samples in successive ethanol, acetone and xylenes washes
    1. Absolute ethanol for 1h.
    2. 4 washes in acetone for 1h each.
    3. 1 wash in acetone for 1h15min.
    4. 3 washes in xylene for 1h each.
  - Embed the samples in paraffin at 56 ° C for 1h and repeat to ensure all solvent is removed. hour.
  - Cut into 5 µm slices using a microtome.
  - Mount the sections onto untreated, degreased slides using glycerine albumen.

- **HES Staining (Haematoxylin-Eosin-Saffron)**
  - Deparaffinize the sections in OTTIX and OTTIX shaper (DiaPath).
  - Stain these sections successively with :
    1. Harris’ hematoxylin for 6 minutes.
    2. Eosin G for 1 minute.
    3. Saffron for 3 seconds.
- Dehydrate in the OTTIX/OTTIX shaper.
- Mount using a mounting medium (Entellan®, Millipore).

HES staining performed on a section of BIOMIMESYS® containing cells (matrix: purple; cytoplasm: pink; nuclei: purple)

- **Immunohistochemistry**
  - Deparaffinize the sections in OTTIX and OTTIX shaper (DiaPath).
  - Uncover the antigenic sites using 10 mM citrate buffer (pH 6) for 12 minutes (or 6 minutes if the cuts come off).
  - Heat the sections with the citrate buffer in a microwave oven.
  - Incubate the sections overnight with the primary antibody at 4 °C.
  - Inhibit endogenous peroxidase by using 5% hydrogen peroxide (30 vol), diluted in 3% PBS-BSA.
  - Incubate the sections for 45 minutes with secondary antibody coupled to peroxidase.
  - Add diaminobenzidine (DAB) [Dako, SK3468] to reveal the antigen-antibody complexes as brown markings.
  - Stain the sections using Mayer’s hematoxylin.
  - Mount the sections in the aqueous medium between slide and cover slip.
Staining
BIOMIMESYS® with Mayer's haematoxylin and marking fibroblasts with an anti-vimentin antibody (M0725, lot 00038159 [Dako])

Contact Information

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