

Reference Protocol for Anti-CD11c (HS-375 017) Immunohistochemistry using DAB as Chromogen

Tissue Fixation

- 3.7% formaldehyde (24 h), 3.5 µM paraffin sections

Materials and Reagents

- | | |
|--|---------------------|
| • Food Steamer | Braun, Multigourmet |
| • Staining Containers with slide holders (e.g. Tissue-Tek) | |
| • Protein Block, Serum-Free | Agilent X0909 |
| • Antibody diluent | Agilent S2022 |
| • Biotinylated anti-rat antibody | Jackson 712-065-153 |
| • ABC HRP Kit, Standard | Vectorlabs PK-4000 |
| • ImmPACT DAB | Vectorlabs SK-4105 |
| • Hydrogen peroxide 30% | Merck 1.07298.0250 |
| • PBS (pH 7.4) | |
| • TBST (TBS, 0.05% Tween 20, pH 7.6) | |
| • Antigen Retrieval buffer:
Citrate Buffer (10 mM Citrate, 0.05% Tween 20, pH 6.0) | |
| • Xylene, 100% ethanol, 90% ethanol, 80% ethanol and 70% ethanol, 2-propanol | |
| • Optional: Hematoxylin Solution (Mayer's, Modified) or other nuclear counterstain | |
| • Optional: Avidin/Biotin Blocking Kit | Vectorlabs SP-2001 |
| • Non-aqueous mounting medium | |

Method

1. Deparaffinize and hydrate tissue sections

- | | |
|--------------------|------------|
| a. Xylol | 2 x 5 min |
| b. 100% EtOH | 2 x 2 min |
| c. 90% EtOH | 1 x 2 min |
| d. 80% EtOH | 1 x 2 min |
| e. 70% EtOH | 2 x 2 min |
| f. Deionized Water | 1 x 20 sec |
| g. PBS | 1 x 2 min |

*Keep the slides in PBS until ready to perform the Antigen Retrieval.
Do not allow the slides to dry out*

2. Antigen Retrieval (AR) using a food steamer

- a. Heat the steamer with a suitable staining container filled with Antigen Retrieval buffer to ~97°C
- b. Transfer the sections into the staining box, wait until the temperature reaches 97°C
- c. Incubate the sections in the steamer for 30 min
- d. Remove the staining container from the steamer and allow the slides to cool down for 20 min (target end temperature ~60°C)

3. Wash slides in PBS, 3 x 1 min

4. Blocking endogenous peroxidase activity

- a. Incubate the sections with 3% hydrogen peroxide in PBS (freshly prepared!) for 5 min

5. Wash slides in PBS, 2 x 1 min

6. Wash slides in TBST, 1 x 2 min

7. Optional: Perform Avidin-Biotin-Block according to manufacturer's instructions.

Note: Certain tissues (e.g. liver, kidney) contain high levels of endogenous biotin. The Avidin-Biotin blocking step is recommended when using the ABC system for these tissues. If the background problem persists, consider trying a polymer-based detection system instead of biotinylated secondary antibody / ABC system.

8. Block in Protein Block, Serum-Free for 10 min

9. Drain slides (do not rinse)

10. Apply primary antibody diluted in Antibody Diluent and incubate in a humidified chamber for 1 h at room temperature

Suggested dilution: 1:100 in Antibody Diluent

11. Wash slides in TBST, 3 x 2 min

12. Apply secondary antibody diluted in Antibody Diluent for 30 min at room temperature.

Suggested concentration: 5 µg/ml

Perform step 13 in the interim

13. Prepare the ABC-reagent: 5 ml PBS + 1 drop A + 1 drop B, incubate for 30 min

14. Wash slides in TBST, 3 x 2 min

15. Apply the ABC reagent for 30 min at room temperature

16. Wash slides in TBST, 3 x 2 min

17. Apply the DAB substrate, 1-10 min

***Observe the staining with a microscope!**

Development times may differ depending upon the level of antigen*

18. Stop the DAB reaction with deionized water

19. Optional: Counterstain

- a. Follow the manufacturer's instructions for counterstaining and bluing

20. Wash slides in deionized water for 1 min

21. Dehydrate tissue sections:

- a. 70% EtOH 2 x 10 sec
- b. 80% EtOH 1 x 10 sec
- c. 90% EtOH 1 x 10 sec
- d. 2-Propanol 2 x 1 min
- e. Xylol 3 x 2 min

22. Mount slides in a suitable organic mounting medium and add coverslip

Note: The SYSY standard protocol generates good results in the SYSY labs and may be used as a reference. However, to achieve the highest

specific signal and lowest non-specific background signal, the best antigen retrieval condition, antibody concentration, incubation temperature, and incubation time must be determined individually. Please also refer to our general protocols.