

# **Short protocol**

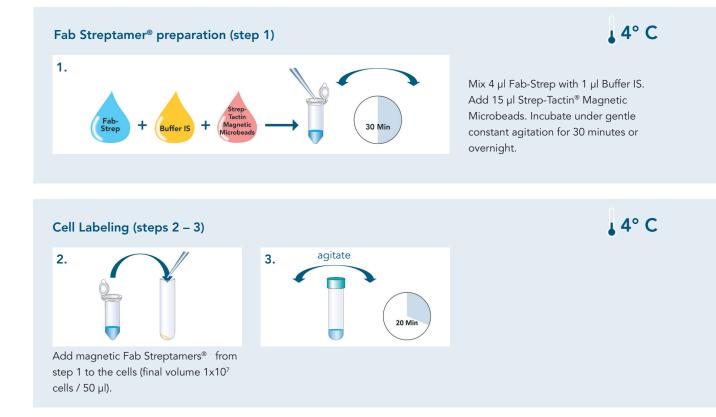


From up to 1 x 10<sup>7</sup> human PBMCs



Avoid foaming, which interferes with proper bead retention on the magnet!

Optional: Wash magnetic microbeads before use to remove sodium azide.

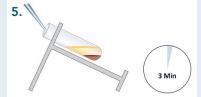


### Magnetic separation (positive selection; step 4 – 7)



Add 5 ml of buffer IS to the cell:bead preparation from step 3.





Incubate tube on StrepMan Magnet for 3 min, isolate target cells by removing unbound cells in 5 ml supernatant.

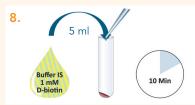


4° C

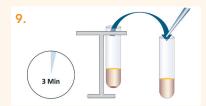
Remove the reaction tube from the StrepMan Magnet and carefully resuspend labeled cells with 5 ml Buffer IS.

Repeat steps 5 and 6 twice, pool target cell fractions, and centrifuge cells.

## Dissociation of Fab Streptamer® reagents (steps 8 – 10)



Resuspend final cell fraction in 5 ml 1 mM D-biotin working solution, mix by pipetting and incubate for 10 minutes.

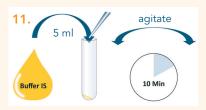


Place tube for 3 min onto StrepMan Magnet. Collect supernatant (target cell fraction).

# 1 x 8. and 9.

Repeat 8. and 9. once, pool target cell fractions, and centrifuge cells.

# Removal of Fab-Streps (step 11 - 13)



Remove Fab-Streps by washing target cell fraction in 5 ml Buffer IS and incubate for 10 min (under agitation).



Place tube for 3 min onto StrepMan Magnet. Collect supernatant (target cell fraction).



Repeat 11. and 12. once, and pool target cell fractions.

#### To perform further positive selections please repeat the protocol from step 1.

#### Please find relevant Fab Streptamer® products here:

http://www.iba-lifesciences.com/products\_fab\_streptamers\_cell\_isolation\_products.html

For detailed instructions please refer to the comprehensive Fab Streptamer® Microbeads manual at www.iba-lifesciences.com/technical-support.html.

Contact us for more information or technical assistance.

#### Fab Streptamer®