



Oct 25, 2019

ChroPlate - ProteinA V.2

DOI

dx.doi.org/10.17504/protocols.io.8qphvvn

Alexandra Ehl¹, David Frommholz¹, Nadine Stefanczyk¹

¹DALEX Biotech



Anonymous

DALEX Biotech

Create & collaborate more with a free account

Edit and publish protocols, collaborate in communities, share insights through comments, and track progress with run records.

Create free account

OPEN  ACCESS



DOI: <https://dx.doi.org/10.17504/protocols.io.8qphvvn>

External link: <https://dalex-biotech.com/>

Protocol Citation: Alexandra Ehl, David Frommholz, Nadine Stefanczyk 2019. ChroPlate - ProteinA. **protocols.io**
<https://dx.doi.org/10.17504/protocols.io.8qphvvn>

License: This is an open access protocol distributed under the terms of the **Creative Commons Attribution License**, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited

Protocol status: Working

Official product protocol by DALEX Biotech



Created: October 25, 2019

Last Modified: October 25, 2019

Protocol Integer ID: 29167

Keywords: DALEX Biotech, Antibody, Antibodies, Protein Purification, Affinity Purification, Kit, Immunoglobulin, fast, simple, convenient, antibodies with chroplate filtration plate, quick high throughput antibody purification from various source, quick high throughput antibody purification, proteina purification guide for the isolation, proteina purification guide, chroplate filtration plate, antibody, purification, chroplate, proprietary resin, filtration, proteina, dalex biotech, well of the chroplate

Abstract

Purification Guide for the Isolation of Antibodies with ChroPlate Filtration Plates by DALEX Biotech.

Easy and quick high throughput antibody purification from various sources and species.

Each well of the ChroPlate has a binding capacity of > 1 mg (tested with human polyclonal Ig, binding varies between species and clones).

The proprietary resin does not shrink or swell in aqueous buffers.

High pressure stability.

pH stability short term 2 - 8, long term 3 - 8.

Excellent thermal stability up to 15 minutes at 80 °C in aqueous buffers at neutral pH.

Can be dried for long term storage (80 °C for > 2 h).

Guidelines

For optimal binding and purity, the pH of the sample should be 7.5-8.5 and should contain 150-300 mM NaCl. An easy way to achieve this is by adding 1/11 volume 0.5 M Tris, 2 M NaCl (pH 8.0) to your sample. For screening of binding condition for e.g. a monoclonal antibody this parameters might be varied.

Purification works best with an antibody concentration of 1-2 mg/ml in your sample.

It is advisable that all fractions are collected (Sample, flow through, wash, and eluate) in separate plates for analysis, e.g. SDS-PAGE.



Materials

Materials provided in the kit:

ChroPlate

Dummy plate

Wash buffer

Elution buffer

Neutralization buffer


Materials not provided in the kit:

Tween-20

0.5 M Tris, 2 M NaCl (pH 8.0)

Troubleshooting

Safety warnings

 The buffers in the kit include sodium azide (CAS No. 26628-22-8) as a preservative.
For safety information on this chemical(s) check <http://www.dguv.de/ifa/gestis-database>

Before start

Make sure your sample is free of particulate matter. You can remove particles by centrifugation or filtration (0.45 μm).



- 1 It is advisable to purify by centrifugation and not by vacuum filtration because of excessive frothing.

STEP CASE

Centrifugation 6 steps

Equilibration

- 2 Add 500 µl wash buffer to each well, place the ChroPlate on top of a deep-well plate, and centrifuge 2 minutes at 400 g in a swing-out rotor. For counterbalance of the centrifuge a dummy filter plate is included in the kit.

Load and Wash

- 3 Place the ChroPlate on a clean deep-well plate. Add up to 1.5 ml sample to every well. Centrifuge 5 minutes at 400 g in a swing-out rotor.

Note

It is advisable to keep the flow-through and wash fractions for later analysis, e.g. SDS-PAGE.

The centrifugation time depends on the sample's volume and viscosity. Volumes larger than 1 ml and viscous samples e.g. serum usually require more than 5 minutes centrifugation time.

For optimal binding and purity, the pH of the sample should be 7.5 - 8.5 and should contain 150 - 300 mM NaCl. An easy way to achieve this is by adding 1/11 volume 0.5 M Tris, 2 M NaCl (pH 8.0) to your sample.

- 4 Empty the deep-well plate or place the ChroPlate on a clean one. Add 500 µl wash buffer to each well and centrifuge 2 minutes at 400 g in a swing-out rotor.

Note

In case of unspecific hydrophobic and/or ionic interactions include up to 1 % Tween-20 and/or up to 0.5 M NaCl in the wash buffer.



- 5 Repeat the previous step.

Note

For increased purity, repeat the washing step a third time.

Elution

- 6 Place the ChroPlate on a clean deep-well plate. Add 300 μ l elution buffer to each well and centrifuge 2 minutes at 400 g in a swing-out rotor.
Repeat two more times.
- 7 Add one drop of neutralization solution to every well of the deep-well plate and mix gently.