

DEAE Persedex A-50

1. Basic product information

DEAE Persedex A-50 chromatography resin is a new type of highly cross-linked dextran chromatography resin. It has high flow rate, low back pressure, high dynamic binding capacity and good chemical stability. DEAE Persedex A-50 chromatography resin is convenient for scale-up, which can shorten production time, improve production efficiency and reduce the cost of large-scale production.

2. Chromatography resin parameters

Resin type	Weak anion exchange
Functional group	Diethylaminoethyl
Matrix	Dextran G50
Median particle size	80 µm
Total ionic capacity	3.0–4.0 mmol/g (dry)
Dynamic binding capacity	≥ 110 mg albumin/ml
Working temperature	4–30°C

3. Chemical resistance

pH stability*	2–9 (working range), 2–12 (CIP)
Chemical stability	All commonly used aqueous buffers, 8M urea, 6M guanidine hydrochloride
Temperature resistance	121°C, 0.1M NaCl solution for 30 minutes

* The physical and chemical properties and functions of the chromatographic resin have no obvious change after being placed in an environment of 40°C and pH 2–9 for 7 days.

4. Method of use

DEAE Persedex A-50 is in powder form and needs to be swollen when used. The swelling ratio depends on the buffer solution used. The swelling ratio in different solutions is different, and there are large differences. Do not use magnetic stirring during swelling, otherwise the gel will easily break.

4.1 Pre-processing

Weigh the required amount of DEAE Persedex A-50, put it in 50~100 times distilled water or loading equilibration solution, and allow to swell. It usually takes 1~2 days to swell at room temperature, and 2 hours in boiling water.

4.2 Column packing

Column packing is performed according to standard operating procedures. It is important to ensure that each material is at working temperature; the gel needs to be degassed before packing.

Note: If the column is packed at the maximum linear flow rate, the flow rate during the subsequent chromatographic separation should not exceed 75% of the column packing flow rate.

4.3 Equilibration

Equilibrate the column with 2–5 column bed volume (hereinafter CV), and make sure the conductance and pH of the eluent are exactly the same as that of the loading buffer.

4.4 Sample loading

Determine the loading amount according to the target product concentration and the loading capacity of the gel.

4.5 Cleaning

After loading the sample, equilibrate the column with loading buffer to wash away unbound proteins and impurities until the conductivity and pH of the eluent are exactly the same as those of the loading buffer.

4.6 Elution

For DEAE Persedex A-50, use the method of increasing the concentration of salt ions in the buffer, continuous elution or gradient elution.

4.7 Regeneration

First wash off the impurity protein on the column with 1~2M NaCl, then wash off the salt in the column with distilled water. Then process the DEAE Persedex A-50 again according to the pretreatment process.

5. Storage

Keep the unopened chromatography resin in the original container and store at 4~30°C in a well-ventilated, dry and clean place. Do not freeze. Wash the used column with 2–3 CV of 20% ethanol solution and store at 2~8°C.

6. Destruction and recycling

Since chromatography resin is difficult to degrade in nature, it is recommended that the waste chromatography resin is incinerated to protect the environment. For chromatography resin that has been in contact with biologically active samples such as viruses and blood, please follow the local biosafety requirements before destroying or disposing of it.

7. Packing method

Detailed information on resin packaging is available on request. Please contact your local distributor.

8. Ordering information

Product name: Product name DEAE Persedex A-50

Product Code in Catalogue	Product Volume
262-00025	25 g
262-00100	100 g
262-00500	500 g
262-01000	1 kg
262-05000	5 kg
262-10000	10 kg

Document version: v1.0. Copyright © 2026 LT Biotech, UAB. All rights reserved. Any redistribution or reproduction of part or all the content in any form is prohibited without the express written permission of LT Biotech, UAB.