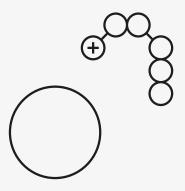
# Multimodal ion exchange chromatography

Multimodal ion exchange chromatography is also referred to as mixed-mode ion exchange chromatography. It utilizes ionic interaction in combination with hydrophobic and other types of interactions. The combined effect gives the resin unique selectivities that adds new possibilities in biomolecule separation.

## Target molecules

WorkBeads 40 TREN resin has a ligand that is positively charged below approx. pH 9. This resin can be used for several different applications, especially due to its higher salt tolerant properties, e.g., for alternative IEX selectivity, for sample cleanup in monoclonal antibody (mAb) purification processes to guard the protein A column from viruses and other host cell impurities, or as a polishing step in the mAb purification process.





## WorkBeads 40 TREN

- Differential selectivity due to higher salt tolerance and multimodal properties
- Reduced fouling of e.g. protein A resins by viruses and host cell impurity removal
- High binding capacity and purity
- · Available in several different GoBio prepacked columns

Structure of the ligand used in WorkBeads 40 TREN.

## **Applications**

## Comparison of prepacked GoBio Mini TREN and GoBio Mini DEAE

Columns: GoBio Mini TREN 1 mL

GoBio Mini DEAE 1mL
Binding buffer: 50 mM Tris-HCl, pH 7.4
Flution buffer: 50 mM Tris HCl 1 M Not

Elution buffer: 50 mM Tris-HCl, 1M NaCl, pH 7.4 Sample: 2.5 mL of 0.3 mg/mL apo-transferrin,

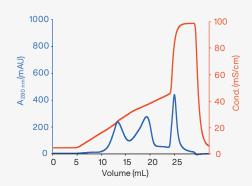
 $0.2\,mg/mL\,\alpha\text{-lactalbumin,}\,0.6\,mg/mL\,soybean$ 

trypsin inhibitor in binding buffer

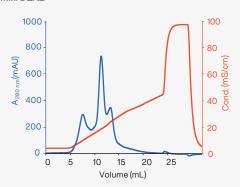
Flow rate: 1 mL/min (150 cm/h)

Gradient: 0 to 40% elution buffer in 20 CV

#### GoBio Mini TREN



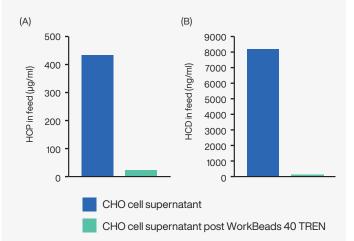
#### GoBio Mini DEAE

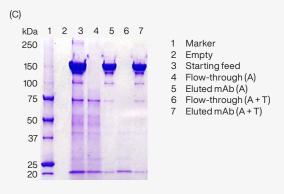


The peaks from left to right correspond to apo-transferrin, a-lactalbumin and soybean trypsin inhibitor. The blue line corresponds to the absorbance at 280 nm and the red line to the conductivity.

## Using WorkBeads 40 TREN as a guard column before protein A

Using WorkBeads 40 TREN upstream of protein A resins is an excellent option for eliminating the extensive bioburden on the protein A resin caused by the impurities from the host cells, and thus extending the lifetime of the protein A resin. The advantage of using WorkBeads 40 TREN upstream of WorkBeads affimAb is shown below. In this experiment up to 95% of HCP and 99% of HCD have been removed from the mAb feed loaded onto the protein A resin.

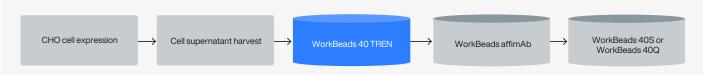




Levels of impurities in CHO cell supernatant before and after WorkBeads 40 TREN treatment. (A) HCP and (B) HCD in mAb sample loaded onto the protein A resin. (C) SDS-PAGE analyses of the feed, flow-through and eluted mAb, with or without WorkBeads 40 TREN (T) upstream of WorkBeads affimAb (A).

### Example of usage of WorkBeads 40 TREN in mAb purification processes

Flow-through mode, protection of Protein Aresin (guard column).



# Technical specifications

#### WorkBeads 40 TREN

Matrix	Rigid, highly cross-linked agarose	
Average particle size $(D_{v50})$	45 µm	
Ligand	Tris(2-aminoethyl)amine (TAEA)	
lonic capacity	130 – 200 µmol Cl <sup>-</sup> /mL resin	
Dynamic binding capacity <sup>2</sup>	50 mg BSA/mL resin	
Maximum flow rate	600 cm/h (20 cm bed height, 5 bar)	
Chemical stability	$Compatible\ with\ all\ standard\ aqueous\ buffers\ used\ for\ protein\ purification.\ Should\ not\ be\ stored\ at\ low\ pH\ for\ prolonged\ time.$	
Operational pH range <sup>3</sup>	2 to 13	
CIP and screening pH range <sup>3</sup>	2 to 14	
Storage	2 to 25°C in 20% ethanol	

The median particle size of the cumulative volume distribution.

# Ordering information

	Pack size	Article number
WorkBeads 40 TREN	25 mL	40 603 001
	150 mL	40 603 003
	1L	40 603 010
	5L	40 603 050
	10 L	40 603 060

Dynamic binding capacity determined at 4 minutes residence time (0.25 mL/min in 1 mL column) in 50 mM Tris-HCl, 50 mM NaCl, pH 8.0. Optimal flow rate during binding is depending on the sample. Within the operational pH range, the resin can be operated without significant change in function. Within the CIP (Cleaning-in-place) and screening pH range the resin can be subjected to the denoted pH range without significant change in function.

