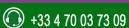
R **INTERFINE CHEMICALS** ANALYTICAL SCIENCES BIOCHROMATOGRAPHY **LIFE SCIENCES**



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Bio-Works RANGE

- Products
- Columns characteristics



Bio-Works Products

Technic	WorkBeads™	BabyBio™ Columns
Exclusion	WorkBeads™ 40 SEC WorkBeads™ 40/100 SEC WorkBeads™ 40/10000 SEC	BabyBio™ D-Salt
lon Exchange	WorkBeads™ 40 Q WorkBeads™ 40 S WorkBeads™ 40 DEAE	BabyBio™ Q BabyBio™ S Baby Bio™ DEAE
RecProtein Affinity	WorkBeads™ 40 Ni WorkBeads™ 40 IDA WorkBeads™ 40 TREN	BabyBio™ NiNTA
Antibody Affinity	WorkBeads™ Protein A	BabyBio™ Protein A
Affinity Antigen - Antibody	WorkBeads™ 40 ACT WorkBeads™ 40/10 000 ACT	

BIOCHROMATOGRAPHY ITM-20160929-P/E













Exclusion

- WorkBeads™ 40 SEC
- WorkBeads™ 40/10 000 SEC



High Performance Size Exclusion Chromatography Media for laboratory and process scale separation of proteins and molecules. Made from agarose, well established and well known in the biotech industry

- Excellent resolution
- Robust separation across a wide range of proteins and molecules
- Chemically stable media

Separation Ranges

Produc	Separation Range kDalton	Exclusion Limit kDalton
WorkBeads™ 40/100 SEC	10 - 150	150
WorkBeads™ 40 SEC	50-1200	1200
WorkBeads™ 40/10000 SEC	Very large molecules and particles	> 10 000

Media description

	WorkBeads™ 40 SEC	WorkBeads™ 40/100 SEC	WorkBeads™ 40/10 000 SEC
Agarose content %	7	9	5
Exclusion limit	1200kDa	150kDa	10 000kDa
Flow rate (cm/h)	600	600	600
Average particle size (µm)	45	45	45
pH stability	2-13	2-13	2-13

The media are preserved in 20% ethanol.



















BabyBio™ Dsalt

BabyBio™ Dsalt ready to use mini column available in 1mL and 5mL which allows quick, easy and convenient group separation of high and low molecular weight substances





- Designed for swifter and efficient desalting and buffer exchange applications
- Stronger, reproducible and easy to use column
- Easy scaling-up, columns can be coupled in series

Column characteristics

Target substance	Proteins and other biomolecules of similar size	
Medium	Highly cross-linked dextran	
Column volumes	1mL 5mL	
Column dimensions	7 × 28mm (1mL) 13 × 38mm (5mL)	
Recommended flow rate	1 ml/min (BabyBio Dsalt 1mL) 5 ml/min (BabyBio Dsalt 5mL)	
Max flow rate (conditions see Data sheet)	5 ml/min (BabyBio Dsalt 1mL) 12 ml/min (BabyBio Dsalt 5mL)	
Maximum back pressure	0.3MPa, 3bar, 43psi	
Chemical stability	Compatible with all standard aqueous buffers used for Protein purification.	
Recommended working range pH Stability	2-12	
Storage	+2°C to +25°C in 20% ethanol	



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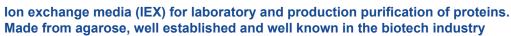






Ion Exchange

- WorkBeads[™] 40 Q
- WorkBeads™ 40 S
- WorkBeads™ 40 DEAE



- High throughput and resolution
- Reliable and reproducible
- High chemical stability for easy cleaning in place

Media description

	WorkBeads™ 40 Q	WorkBeads™ 40 S	WorkBeads™ 40 DEAE
Agarose content	7%	7%	7%
Protein capacity	BSA, 130mg/mL	IgG, 70mg/mL	BSA, 85mg/ml
lonic group	Quartenary amine	Sulphonic acid	Di-ethylaminoethyl
lonic capacity (mmol/mL)	0.18-0.25	0.18-0.25	0.11-0.16
Max flow rate (20cm bed height; 5 Bar) (cm/hr)	600	600	600



ITM-20160929-P/E

















BabyBio™ Q

BabyBio™ Q is a pre-packed, ready to use strong anion exchange column for easy and convenient purification of proteins.

- Rapid method screening and separations
- · High binding capacity and purity
- Easy ready to use columns

BabyBio™ S

BabyBio™ S is a pre-packed, ready to use strong cation exchange column for easy and convenient purification of proteins.

- · Rapid method screening and separations
- High binding capacity and purity
- Easy ready to use columns



BabyBio Q

Bio-Works

BabyBio™ DEAE

BabyBio™ DEAE is a pre-packed, ready to use strong cation exchange column for easy and convenient purification of proteins.

- · Rapid method screening and separations
- High binding capacity and purity
- Easy ready to use columns



Column characteristics

	BabyBio™ Q	BabyBio™ \$	BabyBio™ DEAE	
Medium	WorkBeads™ 40 Q	WorkBeads™ 40 S	WorkBeads™ 40 DEAE	
Matrix	F	Rigid, highly cross-linked agaro	se	
Average particle size		45 µm		
Ligand	Quaternary Amine	Sulfonate	Diethylaminoethyl	
lonic capacity (mmol/mL)	0.18-0.25	0.18-0.25	0.11-0.16	
Dynamic binding capacity1 (mg BSA/mL medium)	50	130	40	
Column volumes		1mL 5mL		
Column dimensions	7×28mm (1mL) 13×38mm (5ml)			
Recommended flow rate				
BabyBio 1mL		1mL/min		
BabyBio 5mL	5mL/min			
Max flow rates				
BabyBio 1mL		5mL/min		
BabyBio 5mL	20mL/min			
Maximum back pressure	0.3 MPa, 3 bar, 43 psi			
Chemical stability	Compatible with all standard aqueous buffers used for protein purification. 70% ethanol. Should not be stored at low pH for prolonged time.			
pH Stability	3-13 working range 2-13 cleaning			
Storage	+2°C to +25°C in 20% ethanol			













RecProtein Affinity

WorkBeads™ 40 Ni

High throughput agarose media for capture of His-tagged proteins

- High dynamic capacity
- High purity
- Excellent flow properties

Media description

	WorkBeads™ 40 Ni	
Agarose content	7%	
Metal Ion capacity (µmol Ni2+/mL)	50-60	
Average particle Size (µm)	45	
Protein capacity (mg/mL)	>60	
Max flow rate (20cm bed height; 5 Bar) (cm/hr)	600	
pH stability	2-13	

- WorkBeads[™] 40 IDA
- WorkBeads™ 40 TRFN

High Throughput Agarose Media for Immobilised Metal Affinity Chromatography (IMAC)

- Made from agarose, well established and well-known in the Biotechnology Industry
- Choice of IMAC chemistry to fit a large variety of proteins
- High flow characteristics

WorkBeads™ 40 IDA and WorkBeads™ 40 TREN immobilised metal affinity chromatography (IMAC) have optimal length spacer arm between the agarose backbone and the attached chelator.

The agarose beads are supplied in aqueous suspensions with 20 % ethanol as preservative and are immediate ready for use after washing.

WorkBeads 40™ IDA and WorkBeads™ 40 TREN media for selecting optimal IMAC conditions.

Media description

	WorkBeads 40 IDA	WorkBeads 40 TREN
Agarose content	7%	7%
Chelating group	Iminodiacetic acid (IDA)	Tris(2-ethylaminoethyl) amine (TREN)
Metal Ion capacity (µmol Cu2+/ml)	10-20 IDAlow 50-60 IDAhigh	10-20 TRENIow 50-60 TRENhigh
Average particle size (µm)	45	45
Protein capacity (mg/mL)	>60	>60
Max flow rate (20cm bed height; 5 Bar) (cm/hr)	600	600
pH stability	2-13	2-13



ITM-20160929-P/E







Bio-Works









BabyBio™ Ni-NTA

BabyBio™ Ni-NTA ready to use mini column available in 1mL and 5mL for quick, easy and convenient affinity purification of proteins carrying a polyhistidine tag (His)





- Swifter purification of His-tagged proteins
- Higher binding capacity and purity in one step
- · Simple and easy method giving reproducible results

Column characteristics

	BabyBio™ Ni-NTA	
Target substance	His-tagged proteins	
Medium	WorkBeads™ Ni-NTA	
Ligand	Nitrilotriacetic acid (NTA) charged with Nickel ions	
Static binding capacity	70 mg His-tagged protein/ml medium	
Dynamic binding capacity (conditions see Data sheet)	50 mg His-tagged protein/ml medium	
Column volumes	1mL 5mL	
Column dimensions	7×28mm (1mL) 13×38mm (5 ml)	
Recommended flow rate		
BabyBio™ 1mL	1mL/min	
BabyBio™ 5mL	5mL/min	
Max flow rates		
BabyBio™ 1mL	5mL/min	
BabyBio™ 5mL	20mL/min	
Maximum back pressure	0.3 MPa, 3 bar, 43 psi	
Chemical stability	Compatible with all standard aqueous buffers used for protein purification. 20% ethanol. Chelating substances (e.g, EDTA will strip off the Ni2+ ions) Stripped column: 10mM HCl (pH2), 10mM NaOH (pH12), 0.1M sodium citrate-HCl (pH3), 6M guanidine-HCl. Should not be stored at low pH for prolonged time.	
Recommended working range	7-9 short term	
pH Stability	2-12 cleaning (stripped column)	
Storage	+2°C to +25°C in 20% ethanol	



interchim[®]













Antibody Affinity

WorkBeads™ Protein A

WorkBeads™ Protein A is a new affinity resin designed for the purification of monoclonal and polyclonal antibodies.

- Higher dynamic binding capacity for monoclonal antibodies; excellent recovery & high purity
- Stronger coupling chemistry; high pH stability and low leakage
- Reliable, reproducible and efficient







ITM-20160929-P/E













BabyBio A

BabyBio A™ ready to use mini columns available in 1 ml and 5 ml. Products allow quick, easy and convenient purification of monoclonal and polyclonal antibodies.

- Swifter purification of polyclonal and monoclonal antibodies
- Higher binding capacity and purity in one step
- Simple and easy method giving reproducible results





Column characteristics

	BabyBio A	
Medium	WorkBeads Protein A	
Matrix	Rigid, highly cross-linked agarose	
Ligand	Recombinant protein A expressed in E. coli using animal-free medium	
Coupling chemistry	Bromohydrin	
Dynamic binding capacity	>40 mg human IgG/ml medium	
Column volumes	1mL 5mL	
Column dimensions	7×28mm (1mL) 13×38mm (5 ml)	
Recommended flow rate		
BabyBio 1mL	0.5-1mL/min	
BabyBio 5mL	1-4mL/min	
Max flow rates		
BabyBio 1mL	5mL/min	
BabyBio 5mL	15mL/min	
Maximum back pressure	0.3 MPa, 3 bar, 43 psi	
Chemical stability	Compatible with all standard aqueous buffers used for protein purification	
Recommended working range	7-9 short term	
pH Stability	3-10 short term 2-10 cleaning	
Storage	+4°C to +8°C in 20% ethanol	

















Affinity Antigen - Antibody

- WorkBeads™ 40 ACT
- WorkBeads™ 40/10 000 ACT

Activated Media for laboratory and process scale Affinity chromatography with User's choice of ligand.

- Faster to results; simple coupling procedure
- Stable at ambient temperature, aqueous coupling solution and neutral pH
- Suitable for coupling ligands containing Sulphydryl-, Amino- or Hydroxyl



- WorkBeads™ 40 ACT for coupling small molecules and peptides
- WorkBeads™ 40/10 000 ACT for coupling immunoglobulins and other large entities

	WorkBeads 40 ACT	WorkBeads 40/10 000 ACT
Agarose content	7%	5%
Flow rate (cm/h) (20cm bed height; 5 Bar) (cm/hr)	600	600
Average particle size (µm)	45	45
Degree of substitution approx (µmol/ml)	250	150
Activated groups	Bromide	Bromide

Buffer selection for WorkBeads™ ACT

Coupling conditions and selection of coupling buffers	Type of ligand Functional group of ligand	Coupling buffers
Organic molecules, peptides	Sulphydryl (-SH)	pH7 and higher.Sensitive ligands can be coupled at pH7.Better yield will be obtained at a higher pH.
Organic molecules, peptides	Amino (-NH2) R2-NH R3-N	 When the ligand is used in excess, dissolve the ligand in distilled water and let the basicity of the ligand determine the coupling pH.
Proteins polypeptides	Sulphydryl (-SH)	 pH7 and higher. Sensitive ligands can be coupled at pH7 but a better yield will be obtained at a higher pH.
Proteins polypeptides	Primary amino (-NH2)	 Coupling yield will increase at higher pH. A carbonate buffer of pH 8-8.5 gives often sufficient coupling without denaturation of sensitive polypeptides and proteins. Another possibility is to run the coupling reaction at lower temperature
All types	Hydroxyl (-OH)	 The low nucleophilicity of the hydroxyl group demands coupling condition at very high pH (pH>12). At a pH>12 cross-linking and hydrolysis will compete with the coupling procedure.











Bio-Works







BabyBio Columns characteristics

- Longer and thinner than competition : More efficient
- Slight funnel shape: sample is re-concentrate at the bottom of the column: thick peaks
- Plug in one peace : robustness
- Dispenser : design optimized. The sample distribution is perfect





BIOCHROMATOGRAPHY











