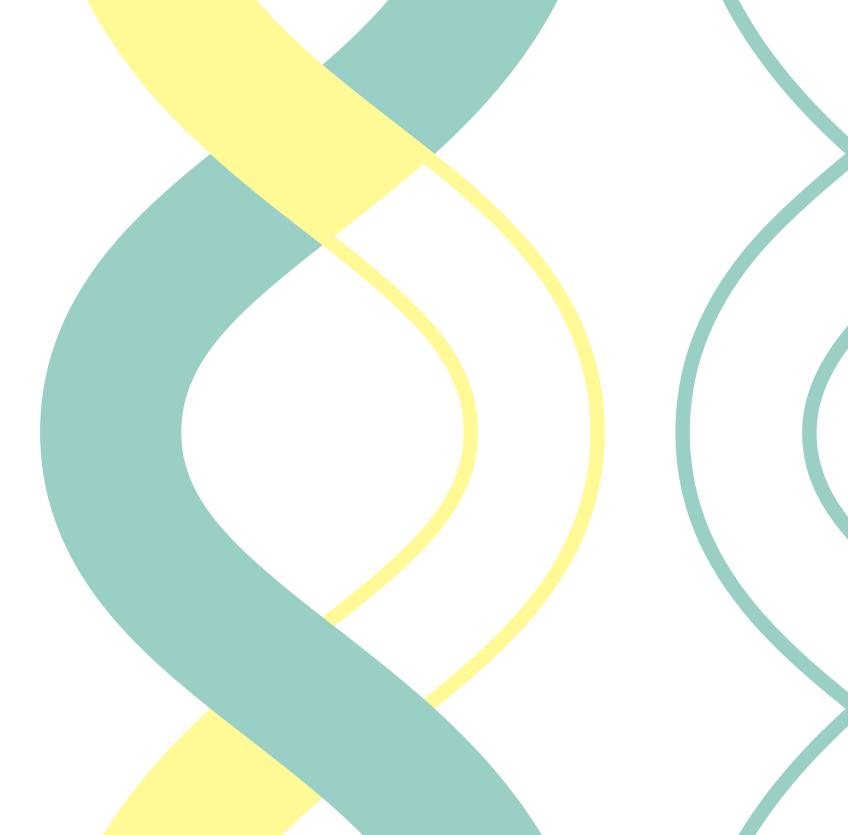
Selection guide

Nucleic acid purification

Choose from our wide range of solutions designed for rapid nucleic acid purification and amplification of DNA and RNA. From spin columns to amplification kits, we can support your purification workflows in multiple applications.





Isolation

Link	Product name	Product code	Format	Major subsequent applications	Capacity/scale	Typical yield	Total time	Advantages	Notes
\rightarrow	Sera-Mag™ Carboxyl	24152105050250	Magnetic bead (hydrophylic) 15 mL	DNA sample preparation and clean-up/size-	50 mg/mL			Wide application range. Convenient one-step or	Unkitted for kit development.
	Beads and SpeedBeads	24152105050350	Magnetic bead (hydrophylic) 100 mL	selection, proteomics and immunoassays.				two-step coupling. Fast magnetic separation. High performance.	Beads = single shell vs SpeedBeads = double shell
		24152105050450	Magnetic bead (hydrophylic) 1000 mL					portormanoc.	opocaboado acabio sitem
		44152105050250	Magnetic bead (hydrophobic) 15 mL						
		44152105050350	Magnetic bead (hydrophobic) 100 mL						
		44152105050450	Magnetic bead (hydrophobic) 1000 mL						
		45152105050250	Magnetic SpeedBead (hydrophylic) 15 mL						
		45152105050350	Magnetic SpeedBead (hydrophylic) 100 mL						
		65152105050250	Magnetic SpeedBead (hydrophobic) 15 mL						
		65152105050350	Magnetic SpeedBead (hydrophobic) 100 mL						
		65152105050450	Magnetic SpeedBead (hydrophobic) 1000 mL						
\rightarrow	Sera-Mag Oligo dT	38152103011150	Magnetic bead 1 mL	RT-PCR, cDNA library construction, cDNA	> 300 pmoL (dA)30	10 mg/mL	5–15 mins	Very high, specific poly A+ binding capacity ensures	
		38152103010150	Magnetic bead 5 mL	microarrays, affinity purification, primer extension and subtractive hybridization.	per mg			maximum extraction of mRNA. Fast reaction kinetics increase throughput and precision, also enabling	
		38152103010350	Magnetic bead 100 mL	extension and subtractive hybridization.				faster movement through viscous solutions. Uniform, nominal 1 µm diameter provides high surface area and excellent lot-to-lot reproducibility.	s n, nd
\rightarrow	SeraSil-Mag™	29357369	Magnetic bead 400 nm × 5 mL	DNA isolation from all sources for subsequent molecular or NGS applications.	8 μg of genomic DNA from a 200 μL alloquot	20 mg/mL in water (0.05% sodium azide)	120 minutes	Beads are monodispersed with narrow size distribution for consistent, reproducible results. Silanol hydroxyl groups on the bead surface give efficient, high purity isolation of nucleic acids. Tested for microbial contamination.	Unkitted for kit development
		29357371	Magnetic bead 400 nm × 60 mL						
		29357372	Magnetic bead 400 nm × 450 mL						
		29357373	Magnetic bead 700 nm × 5 mL						
		29357374	Magnetic bead 700 nm × 60 mL						
		29357375	Magnetic bead 700 nm × 450 mL						
\rightarrow	Sera-Xtracta™ Cell-Free DNA Kit	29437807	Magnetic bead 96 reactions	NGS, qPCR, ddPCR, other amplication and genotyping applications.	50–300 bp fragments	Up to 4 mL input volume of plasma or serum	< 120 mins	Scalability from 5–4 mL input plasma. Minimal copurification of high molecular weight gDNA.	
\rightarrow	Sera-Xtracta Virus/	29506009	Magnetic bead 96 reactions	PCR, Real Time PCR and qPCR.	1 copy per micorliter	Reproducible detection	30 minutes	High throughput; sensitive; scalable up to 400 μL	
•••••	Pathogen Kit	29514201	Magnetic bead 1000 reactions		minimum detection	of low viral titer from 10 pfu/mL	or less		
→	Sera-Xtracta Genomic DNA Kit	29429140	Magnetic bead 96 reactions	NGS, restriction enzyme digestion, PCR and other genotyping applications.	4–8 μg from 200 μL whole blood	Up to 200 μL	< 45 mins	Removes the need for RNAse treatment in most applications $A_{260}/A_{280} > 1.7$	Greater than 100 kb

Isolation (continued)

Link	Product name	Product code	Format	Major subsequent applications	Capacity/scale	Typical yield	Total time	Advantages	Notes	
\rightarrow	Nucleon BACC 1 Genomic DNA Extraction Kit	RPN8501	26 preparations	PCR, sequencing, restriction, BAC Constructs, Southerns.	50 μL–10 mL whole blood 1 × 10 ⁶ to 1 × 10 ⁷	whole blood whole blood.	30 mins	Purity, scalable, non-column format allows for high molecualr weight DNA, no phenol	Genreates high molecular weight DNA, Novel protein binding resin, protocol modifications for buccal	
	Nucleon BACC 3 Genomic DNA Extraction Kit	RPN8512	60 preparations		cultured cells	8–12 μg DNA/10 ⁶ cultured cells.			swabs, mouthwashes, soft tissue, sperm, gram negative bacteria	
→	Nucleon HT Genomic DNA Kit	RPN8509	50 preparations of up to 25 mg of hard tissue or 50 paraffin sections	PCR, sequencing, restriction, can be used with hard to process tissues e.g. FFPE, Southerns.	25 mg tissue sample 30 μm paraffin section	. •	_	Purity, non-column format gives high yields	Novel protein binding resin	
\rightarrow	Nucleon PhytoPure	RPN8510	50 × 0.1 g preps	PCR, RAPDs, restriction, for plant tissues	Up to 0.1 g	Variable depending on	1–1.5 hours	Purity, scalable, dedicated plant kit	Novel polysaccharide binding	
		RPN8511	50 × 1 g preps		Up to 1.0 g plan	species, age and part of plant used, approx. 10–60 µg per 0.1 g prep.			resin	
\rightarrow	RNASpin Mini Kit	25050070	20 preps	•	(qRT-PCR), Northern blot and microarray of 100 μg	Binding capacity	Up to 100 μg			
		25050071	50 preps			ot and microarray of 100 µg			total RNA since genomic DNA is removed via on-column DNase I treatment. Results can be obtained with even small amounts of sample. Lysis buffer is	
		25050072	250 preps	охропшонсь.				less susceptible to foaming to ensure valuable RNA sample is not wasted. Simple and convenient format suitable for all levels of expertise.		
\rightarrow	Blood genomicPrep	28904263	10 preparations	PCR, and restriction enzyme digests Suitable for use in molecular biology applications including standard, multiplex and quantitative PCR; cloning; haplotyping; restriction enzyme digestion; and genotyping	PCR, and restriction enzyme digests	Binding capacity	4–6 μg/200 μL	• •		
	Mini Spin Kit	28904264	50 preparations			60 µg	60 μg whole blood t	time < 20 minutes	s and less shearing. Demonstrated application to various types of whole blood including human, horse, rabbit, rat,	
		28904265	250 preparations					and mouse. Reduced pipetting volume changes, one centrifugation setting		
\rightarrow	Bacteria genomicPrep	28904258	50 preparations	Molecular biology applications including	Gram-negative:	4–8 μg of genomic DNA from 2 × 10 ⁹ bacteria cells	Total preparation time: 40 minutes	Rapid extraction and purification of high molecular		
	Mini Spin Kit	28904259	250 preparations		4–12 μg gDNA Gram-positive:			weight genomic DNA (gDNA) from Gram-negative (G-) and Gram-positive (G+) bacteria. Reduced lysis time.		
				and genetyping	5–10 μg gDNA		Bacteria), 55	Capable of handling input amounts ranging from 1×10^9 to 4×10^9 bacteria cells.		
\rightarrow	Tissue and cells	28904275	50 purifications	Restriction enzyme analysis, ligation,	Binding capacity	Up to 1.5 μg/mg of tissue	90 minutes	Rapid procedure produces high molecular weight		
	genomicPrep Mini Spin Kit	28904276	250 purifications	cloning, DNA sequencing and PCR	>35 µg	from an input tissue sample range of 5 to 50 mg, and up to 40 µg of genomic DNA per 5 × 10 ⁶ cells depending on cell type	45 minutes (Cultured cells)	gDNA with minimal shearing. Total DNA extraction time reduced significantly to 90 min Produces high-quality high molecular weight DNA and less degradation		
→	TriplePrep Kit	28942544	50 kit	Restriction enzyme digestion, PCR, sequencing, array CGH, RT-PCR, cDNA synthesis, expression array, SDS-PAGE, Western Blotting, 2-D DIGE, LCMS.	Binding capacity of 20–60 μg	80–160 μg	< 45 minutes	Fast: less than one hour. High yield: enough DNA and RNA, together with protein to allow gene, transcriptome and protein to be studied from one sample simultaneously. Easy to use: color-coded tubes.		

Amplification

Link	Product name	Product code	Format	Major subsequent applications	Capacity/scale	Typical yield	Total time	Advantages	Notes
→	GenomiPhi™ Single Cell DNA Amplification Kit	29108107 29108039	25 reactions (performed in tubes or microplates) 100 reactions (performed in tubes or microplates)	Whole-genome amplification from 1–1000 cells, used to amplify DNA samples for genotyping (single nucleotide polymorphism [SNP], short tandem repeats [STR], comparative genomic hybridization [CGH]), sequencing, or PCR assays, where starting materials are limited.	1 fg of input DNA	4–7 μg	< 3 hours	UV and enzymatic cleanup steps during manufacture keep kit free of detectable DNA. Propritary sample clean up step in protocol reduces unwanted background amplifictation.	Product Size > 10 kb. See Brochure: KA10877280120DF.
→	GenomiPhi V2 DNA Amplification Kit	25660030 25660031 25660032	25 reactions 100 reactions 500 reactions	Whole-genome amplification from nanogram amounts of DNA, used to amplify DNA samples for genotyping (single nucleotide polymorphism [SNP], short tandem repeats [STR], comparative genomic hybridization [CGH]), hybridization studies, cloning, sequencing, DNA archiving, transfection or PCR assays.	10 ng input DNA or cell lysate	4–7 μg	< 2 hours	Isothermal amplification (no thermocycler required). Phi29 DNA polymerase carries proofreading activity. Independent amplification products from random priming events. Uniform coverage of the genome.	Simple, robust, reliable setup for whole genome amplification (for all genomiphi kits) compared to other PCR based methods. Verified for use with FTA cards, blood or buccal swabs. Store at -70°C. Average product size > 10 Kb.
→	GenomiPhi V3 (RTG) DNA Amplification Kit	25660124 25600196 25660197	24 rxns (3 × 8-well strip) 96 rxns (1 × 96-well plate) 480 rxns (5 × 96-well plate)	Whole-genome amplification from nanogram amounts of DNA, used to amplify DNA samples for genotyping (single nucleotide polymorphism [SNP], short tandem repeats [STR], comparative genomic hybridization [CGH]), hybridization studies, cloning, sequencing, DNA archiving, transfection, or PCR assays.	10 ng input DNA or cell lysate	12-20 μg	< 2 hours	Preformatted, predispensed single dose lyophilized cake. Room temperature stable. Isothermal amplification (no thermocycler required). Uniform coverage of the genome. Phi29 DNA polymerase has proofreading activity. Independent amplification products from random priming events.	Verified for use with FTA cards, blood or buccal swabs. Average product size > 10 Kb. See App Note: 29-0245-46 AA 7.
→	TempliPhi™ Sequence Resolver Kit	28903529 28903530 28903531	20 reactions 50 reactions 200 reactions	Amplification of single or double stranded circular DNA from difficult to sequence viral isolates, glycerol stock, or bacterial culture, for the amplification of difficult to sequence templates (repeats, sequencing stops, and compressions) for successful DNA sequencing.	1–2 ng of input DNA	1 μg	18 hours	Simple isothermal workflow. Thermocycler not required. Consistent yield per reaction. Target-specific primer sequence/primers not needed as random priming is utilized.	800 bases. Phred20. Useful for both small (plasmid) and large (BAC) circular constructs.
→	TempliPhi 100/500 Amplification Kit	25640010 25640050	100 reactions (performed in tubes or microplates) 500 reactions (performed in tubes or microplates)	Amplification of single or double stranded circular DNA from viral isolates, glycerol stock, or bacterial culture prior to restriction anlaysis, labeling, sequencing, or transfection.	Small portion of a colony 0.2–0.5 µL liquid culture or glycerol stock or viral isolate 1–10 pg of purified plasmid	1–1.5 μg	on 4–6 hours	Simple isothermal workflow. Thermocycler not required. Consistent yield per reaction. Target-specific primer sequence/primers not needed as random priming is utilized.	Can be used to prepare templates for cell free protein expression, mRNA synthesis, circular viral genome enrichment, synthetic biology. Phi29 has proof reading activity. See Brochure: 28-9622-94 AD 10/2014.

Amplification (continued)

Link	Product name	Product code	Format	Major subsequent applications	Capacity/scale	Typical yield	Total time	Advantages	Notes
\rightarrow	TempliPhi 2000 Amplification Kit	28964286	2000 reactions (performed in tubes or microplates)	Amplification of single or double stranded circular DNA from viral isolates, glycerol stock, or bacterial culture prior to restriction anlaysis, labeling, sequencing, or transfection.	Small portion of a colony 0.2–0.5 µL liquid culture or glycerol stock or viral isolate 1–10 pg of purified plasmid	1–1.5 μg	< 20 mins hands on 4–6 hours amplification time	Simple isothermal workflow. Thermocycler not required. Consistent yield per reaction. Target-specific primer sequence/primers not needed as random priming is utilized.	Phi 29 has proof reading activity. See Brochure: 28-9622-94 AD 10/2014.
\rightarrow	Templiphi Large Construct Kit	25640080	1000 reactions	Prepare DNA for sequencing from very large single- or double-stranded circular constructs, for the amplification of very large sequence templates (BAC or Fosmid) for successful DNA sequencing.	1–10 ng BAC or Fosmid input DNA, glycerol stocks, bacterial culture	5 μg	18 hours	Simple isothermal workflow Thermocycer is not needed. Consistent yield per reaction. Target-specific primer sequence/primers not needed as random priming is utilized.	Phi 29 has proof reading activity.
→	PureTaq Ready-To-Go PCR Beads	27955701 27955702 27955801 27955901	Multiwell plate, 96 Reactions Multiwell plate, 5 × 96 Reactions 0.5 mL tubes, 100 Reactions 0.2 mL hinged tube with cap, 96 Reactions	Reliable and robust end point PCR, PCR amplification, compatible with TaqMan probes and MGB Eclipse Probes, intercalating dye for real time PCR systems, minimizes assay variability for large sample sets.	Pre-dispensed 25 μL reactions	NA	NA	Room temperature stable, sustainable, low non-specific background amplification, pre-dispensed reactions reduce training time.	See App Note: 07/2008 63-0054-46 AB.
\rightarrow	RAPD (RTG) Analysis Kit/Beads	27950001 27950201	100 reactions 100 reactions and 6 primers	Rapidly detect genomic polymorphisms, DNA profiling experiments using randomly amplified polymorphic DNA (RAPD) techniques, used for gene mapping, determining strain diversity, population analysis, taxonimic relationships, microbial QC testing.	ng amounts of DNA	NA	NA	Performatted, predispensed, single dose. Room temperature stable. Controls provided for development studies (kit).	Customer specific primer set is required to use RAPD beads. See App Note: 80-6334-41 Rev A / 5–96.
→	RTG RT-PCR Beads	27925901 27926601 27927601	0.2 mL hinged tube with cap 0.5 mL tubes 0.2 mL tubes	Reliable and robust end point Reverse Trascriptase-PCR, used for detection of RNA in a species, determining relative RNA levels, gene-specfic amplfication of known mRNAs.	Pre-dispensed 50 μL reactions	NA	NA	Long term stabiltiy at room temperature reduces shipping and freezer requirements, reagents are optimized for full-length cDNA synthesis to > 7.5 kb, rabbit globin mRNA and primer controls included.	M-MuLV reverse trascriptase and RNAguard. See Datafile: 11-0026-06 AC 04/2013.
→	Direct RT-qPCR Kit	29656615	AIC Mix, 100 preparations RT-qPCR Kit Mastermix and reconstitution buffer, 100 preparations	Real-time and endpoint RT-PCR	Reaction volume 20 μL	10 copies per reaction		Capable of detecting low viral loads even when the sample-transport media volume reaches 45% of the final reaction volume. Increased RNA stability in the sample. Detection of SARS-CoV-2 viral particles down to approximately 10 copies per reaction	

Clean-up

Link Product name	Product code	Format	Major subsequent applications	Capacity/scale	Typical yield	Total time	Advantages	Notes
→ Sera-Mag Select	29343045	5 mL liquid reagent containing Sera-Mag Carboxyl beads	Removal of unincorporated primers, linkers, probes in library construction.		> 80% recovery of 250 bp DNA fragments	Protocol dependant (e.g. SPRI, PCR)	Single reagent for PCR clean-up and size selection. Manufacturer of beads contained in reagent = security of supply. Direct replacement from AMPure. Options for customization.	Selection of fragment size is determined by adjusting ratio of reagent to fixed volume of sample.
	29343052	60 mL liquid reagent containing Sera-Mag Carboxyl beads	Fragment size selection for next-genreation sequencing (NGS).					
	29343057	450 mL liquid reagent containing Sera-Mag Carboxyl beads					Gustoffization.	
	29453302	20 × 5 mL liquid reagent containing Sera-Mag Carboxyl beads						
Sera-Mag Carboxyl	24152105050250	Magnetic bead (hydrophylic) 15 mL	Library prep, next-generation sequencing,	50 mg/mL		< 45 minutes	Wide application range. Convenient one-step or	Unkitted for kit development.
Beads and SpeedBeads	24152105050350	Magnetic bead (hydrophylic) 100 mL	SP3 Protocols				two-step coupling. Fast magnetic separation. High	Beads = single shell vs
эрееивеаиз	24152105050450	Magnetic bead (hydrophylic) 1000 mL					performance.	Speedbeads - double shell.
	44152105050250	Magnetic bead (hydrophobic) 15 mL						
	44152105050350	Magnetic bead (hydrophobic) 100 mL						
	44152105050450	Magnetic bead (hydrophobic) 1000 mL						
	45152105050250	Magnetic SpeedBead (hydrophylic) 15 mL						
	45152105050350	Magnetic SpeedBead (hydrophylic) 100 mL						
	65152105050250	Magnetic SpeedBead (hydrophobic) 15 mL						
	65152105050350	Magnetic SpeedBead (hydrophobic) 100 mL						
	65152105050450							
ExoProStar™ 1-Step	US77702	100 reactions, mixed enzymes Exo I and Alkaline Phosphatase	Sequencing, cloning, genotyping		100% recovery	30 minutes	Enzymatic PCR clean up of unicorporated primers and dNTPs. Scalable. No product loss, one pipetting step.	Temperature inactivation of the enzymes once reaction completes.
	US77705	500 reactions, mixed enzymes Exo I and Alkaline Phosphatase						
	US77720	2000 reactions, mixed enzymes Exo I and Alkaline Phosphatase						
	US77750	5000 reactions, mixed enzymes Exo I and Alkaline Phosphatase						
→ GFX™ PCR DNA and	28903466	10 reactions	Prior to next-generation sequencing,	100 µL liquid reaction.	90% from solution.	5 minutes	Purification and concentration of PCR products/DNA	No glass slurry, elution in low-
Gel Band Purification	28903470	100 reactions	cloning, PCR	Up to 900 mg agarose	60%–80% from	from solution,	(100 bp–10 kbp) from solution or from gel bands. Speed, purity, both gel band and PCR clean up in same kit.	Unkitted for kit development. Beads = single shell vs SpeedBeads = double shell. Temperature inactivation of the enzymes once reaction completes. No glass slurry, elution in low- salt buffer, for agarose gel slices buffered in Tris-borate-EDTA (TBE) or Tris-acetate-EDTA (TAE). Prepacked and pre-equilibrated with Sephadex G-50 resin.
KIT	28903471	250 reactions		slice	agarose	15 minutes from agarose		
AutoSeq G-50 Columns	27534001	50 columns, spin-column, microfuge, tabletop centrifuge	Molecular biology assays	10–100 μL	80% recovery	4 minutes	Rapid DNA purification by removal unincorporated fluorescent dye-terminators from cycle sequencing reactions prior to analysis on sequencing platforms. Excellent recovery.	determined by adjusting ratio of reagent to fixed volume of sample. Unkitted for kit development. Beads = single shell vs SpeedBeads = double shell. Temperature inactivation of the enzymes once reaction completes. No glass slurry, elution in low-salt buffer, for agarose gel slices buffered in Tris-borate-EDTA (TBE) or Tris-acetate-EDTA (TAE).
	27534002	250 columns, spin-column, microfuge, tabletopcentrifuge						
	27534003	1000 columns, spin-column, microfuge, tabletop centrifuge						

Clean-up (continued)

Link	Product name	Product code	Format	Major subsequent applications	Capacity/scale	Typical yield	Total time	Advantages	Notes
→	MicroSpin™ G-25 Columns	27532501	Spin-column, microfuge, tabletop centrifuge	Molecular biology assays	10–100 μL	90% recovery	4 minutes	Rapid purification of DNA > 10 bp for buffer exchange, desalting and removal of unincorporated nucleotides following synthesis and labeling reactions. Excellent recovery.	Prepacked and pre-equilibrated with Sephadex G-25 resin.
\rightarrow	MicroSpin G-50 Columns	27533001 27533002	50 columns, spin-column, microfuge, tabletop centrifuge 250 columns, spin-column, microfuge, tabletopcentrifuge	Molecular biology assays	10–100 μL	80% recovery	4 minutes	Rapid purification of DNA > 20 bp for buffer exchange, desalting and removal of unincorporated nucleotides following synthesis and labeling reactions. Excellent recovery.	Prepacked and pre-equilibrated with Sephadex G-50 resin.
\rightarrow	MicroSpin S-200 HR Columns	27512001	Spin-column, microfuge, tabletop centrifuge	Molecular biology assays	25–100 μL	70%–80% recovery	4 minutes	Rapid purification of labeled single-stranded or double- stranded DNA ≥ 50 bases. Excellent recovery.	Prepacked and pre-equilibrated with Sephacryl S-200 HR resin.
\rightarrow	MicroSpin S-300 HR Columns	27513001	Spin-column, microfuge, tabletop centrifuge	Molecular biology assays	25–125 μL	70%–80% recovery	4 minutes	Rapid purification of DNA > 20 mers (primers) and nucleotides; for example plasmid purification prior to sequencing templates. Excellent recovery.	Prepacked and pre-equilibrated with Sephacryl S-300 HR resin.
\rightarrow	MicroSpin S-400 HR Columns	27514001	Spin-column, microfuge, tabletop centrifuge	Molecular biology assays	25–50 μL prior to subsequent PCR or cloning; 51–100 μL prior to sequencing	50%–70% recovery	4 min/2 columns	For rapid purification of PCR products (> 200 bp) from unincorporated primers (< 32-mers) and nucleotides including desalting, buffer exchange, and primer removal using spin-column chromatography. Speed and excellent recovery.	Prepacked and pre-equilibrated with Sephacryl S-300 HR resin.
\rightarrow	NICK Columns	17085501 17085502	20 columns, gravity flow chromatography 50 columns, gravity flow chromatography	Molecular biology assays	Concentration ≤ 1 mg/mL; ≤ 100 μL	90% recovery	< 15 min	Rapid nick-translated > 20 DNA fragments and for separation of any labeled probe from unincorporated labeled nucleotides. Excellent recovery.	No centrifuge required.
→	ProbeQuant™ G-50 Micro Columns	28903408	Spin-column, microfuge, tabletop centrifuge	Molecular biology assays	25-50 μL	80% recovery	4 minutes	Rapid purification of labeled DNA > 20 bases from unincorporated labeled nucleotides. Excellent recovery	Prepacked and pre-equilibrated with Sephadex G-50 resin.
\rightarrow	Sephadex™ G-25 DNA Grade SF	17057202	Spin-column or gravity flow chromatography	Molecular biology assays	Varies with end user	90% recovery	Varies with end user	Size exclusion chromatography resin designed for desalting and buffer exchange, and for purification of DNA and oligonucleotides > 20 base pairs for use in spin columns and gravity flow chromatography. End user can pack their own column.	
→	Sephadex G-50 DNA Grade F	17057301 17057302	25 g, spin-column or gravity flow chromatography 100 g, spin-column or gravity flow chromatography	Molecular biology assays	Varies with end user	90% recovery	Varies with end user	Size exclusion chromatography resin designed for desalting and buffer exchange, and for purification of DNA and oligonucleotides > 20 bp. End user can pack their own column.	Can be spin columns or gravity flow chromatography.

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CY12531-06Jun21-SG

