Product List / 2022

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Enzymes As You Need

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## **DNA & RNA ISOLATION KITS**

Product Name	Pack Size	Cat. No.	Description
GENOMIC DNA Isolation	on Kits	'	
NA AKCIJI  EXTRACTME GENOMIC DNA KIT universal	50 preps	EM13-050	Purification of genomic, mitochondrial, bacterial, parasite or viral DNA from solid tissues, physio- logical fluids (urine, cerebrospinal fluid, peritoneal fluid, pleural fluid, sputum), fresh and
	250 preps	EM13-250	frozen blood, mucosa membrane swabs (including buccal, nasal, pharyngeal and vaginal swabs), semen, hair, rodent tails, insects, bacteria, yeast and cell cultures.
<b>EXTRACT</b> ME	50 preps	EM03-050	Purification of high quality DNA from solid tissues (fresh, frozen, formalin-preserved or paraffin-
DNA TISSUE KIT	250 preps	EM03-250	-embedded), physiological fluids, hair, rodent tails, insects and cell cultures.
<b>EXTRACT</b> ME	50 preps	EM05-050	Purification of high quality (genomic, mitochondrial and viral) DNA from whole blood (fresh or
DNA BLOOD KIT	250 preps	EM05-250	frozen, human or other mammalian), plasma, serum, buffy coats, lymphocytes and body fluids.
<b>EXTRACT</b> ME	50 preps	EM06-050	Purification of high quality DNA from human and animal mucosa membrane swabs (including
DNA SWAB & SEMEN KIT	250 preps	EM06-250	buccal, nasal, pharyngeal and vaginal swabs) as well as from semen.



Product Name	Pack Size	Cat. No.	Description
RNA Isolation Kits			
<b>EXTRACT</b> ME	50 preps	EM39-050	Rapid and efficient purification of high-qualit viral RNA from swabs. The kit is specificall designed to isolate viral nucleic acid fror a variety of RNA viruses. The isolation protocc and buffer formulation were optimized for hig
VIRAL RNA KIT	250 preps	EM39-250	isolation efficiency and RNA purity. RNA bindin, capacity: ~120 µg. Purified RNA is eluted with th use of low ionic strength buffer and may be used directly in all downstream applications, such a RT-PCR, RT-qPCR, cDNA synthesis.
NA AKCIJI EXTRACTME	50 preps	EM09.2-050	Improved kit for rapid, efficient purification o high quality total RNA from up to 30 mg of tissu (fresh or frozen), or up to 10 <sup>7</sup> cultured cells. RN.
TOTAL RNA KIT	250 preps	EM09.2-250	binding capacity: ~230 µg. Significantly improved RNA yields and shortened processing time.
EXTRACTME	50 preps	EM12-050	For rapid, phenol-free extraction of RNA highl enriched in short RNA strands (< 200 nt). Superio yields and purity. Suitable for wide range of cell: tissues (including blood). This kit also allows par allel extraction of high quality long RNA strand
miRNA KIT	250 preps	EM12-250	(>200 nt) from the same sample. The kit contain three different types of columns: first one for DN, removal, second one for purification of long RNA and third one for purification of short RNA.
EXTRACTME	50 preps	EM15-050	Rapid, simultaneous isolation of high qualitingenomic DNA and total RNA from a singly biological sample, from up to 30 mg of tissur
RNA & DNA KIT	250 preps	EM15-250	or up to 10 <sup>7</sup> cultured cells. This kit is ideal for researchers interested in studying the genome and the transcriptome of a single sample.
EXTRACTME	50 preps	EM31.1-050	Rapid and efficient purification and concentratio of high quality RNA from tissue or cultured cell
TOTAL RNA MICRO SPIN KIT	250 preps	EM31.1-050	in a micro-spin column format (elution volum from 5 μl).
NA AKCIJI EXTRAZOL	200 ml	EM30-200	Ready-to-use reagent for the isolation of separat fractions of RNA, DNA and proteins from cell and tissue samples of human, animal, plant, yeast, obacterial origin, within one hour.
Bead-beating Tubes with ceramic filling	100 pcs	HPLM100 / HPLM100A	2 ml bead-beating tubes with 1 g cerami filling (1.4 mm) for soft tissue homogenization
	500 pcs	HPLM500 / HPLM 500A	Lysing Matrix D equivalent. Two different tube shapes that will fit to any bead-beater.



Product Name	Pack Size	Cat. No.	Description
PLASMID DNA Isolatio	n Kits		
<b>EXTRACT</b> ME	50 preps	EM01.1-050	Mini-scale extraction of plasmid DNA from broth culture or frozen cell pellets of recombinant
PLASMID MINI KIT	250 preps	EM01.1-250	Escherichia coli strains. Higher yields – column binding capacity 60 µg pDNA; one protocol for high/low copy plasmids.
EXTRACTME PLASMID MIDI KIT	10 preps	EM16-010	Ultrapure, transfection-grade plasmid DI isolation in medium scale (50–300 ml bacterial culture); vield: 200–600 µg DNA fro
	25 preps	EM16-025	100 ml culture; isolation time: 120–130 minutes (with DNA precipitation); centrifugation steps: 6000 x g (no need to have ultracentrifuge).
<b>EXTRACT</b> ME	10 preps	EM18-010	Ultrapure, transfection-grade plasmid DNA isolation in large scale (200–1000 ml of bacterial culture); yield: 1–1.5 mg DNA from 400 ml culture;
PLASMID MAXI KIT	25 preps	EM18-025	isolation time: 140–150 minutes (with DNA precipitation); centrifugation steps: 6000 $\times$ g (no need to have ultracentrifuge).

Product Name	Pack Size	Cat. No.	Description
DNA Fragments Purific	ation Kits		
NA AKCIJI EXTRACTME	50 preps	EM07.1-050	New upgraded kit for DNA purification after enzymatic reactions; the kit enables the purification of DNA fragments from 50 bp to 20 kb, as well as plasmid and genomic DNA;
DNA CLEAN-UP KIT	Sig (de 250 preps EM07.1-250 cap	significal improved recovery: up to 99% (depending on DNA fragment length); binding capacity: approx. 40 µg DNA; time required 10 min for 6 PCR purifications.	
EXTRACTME DNA	50 preps	EM26.1-050	DNA purification after enzymatic reactions & DNA
CLEAN-UP & GEL-OUT KIT	250 preps	EM26.1-250	fragments isolation directly from agarose gels – two options in one kit.



Product Name	Pack Size	Cat. No.	Description
Mini Spin Columns		'	
DNA CLEAN-UP mini spin columns	50 pcs	EM07.1C-050	Mini spin columns with silica resin with 2 ml receiving tubes used in EM07.1 kit.
DNA GEL-OUT mini spin columns	50 pcs	EM08C-050	Mini spin columns with silica resin with 2 ml receiving tubes used in EM26.1 kits.
PLASMID DNA mini spin columns	50 pcs	EM01.1C-050	Mini spin columns with silica resin with 2 ml receiving tubes used in EM01.1 kit.
SWAB & SEMEN DNA mini spin columns	50 pcs	EM06C-050	Mini spin columns with silica resin with 2 ml receiving tubes used in EM06 kit.
GENOMIC DNA mini spin columns	50 pcs	EM13C-050	Mini spin columns with silica resin with 2 ml receiving tubes used in EM03, EM05, EM13 kits.
TOTAL RNA mini spin columns	50 pcs	EM09.1C-050	Mini spin columns with silica resin with 2 ml receiving tubes used in EM09.1 and EM15 kits.
miRNA mini spin columns	50 pcs	EM12C-050	Mini spin columns with silica resin with 2 ml receiving tubes used in EM12 kit.
MICRO SPIN columns	50 pcs	EM28C-050	Micro spin columns with silica resin with 2 ml receiving tubes used in used in EM31 kits.

# **REAL-TIME PCR MASTER MIXES**

Product Name	Pack Size	Cat. No.	Description
<b>AMPLIFY</b> ME	200 rxns	AM01-020	The AMPLIFYME SG No-ROX Mix is a convenient enzyme mixture for fast and reliable quantitative Real-Time PCR, using SG dsDNA-binding dye.
SG No-ROX Mix	2000 rxns	AM01-200	Compatible with qPCR instruments that don't need ROX dye.
NA AKCIJI AMPLIFYME	200 rxns	AM02-020	The AMPLIFYME SG Universal Mix is a convenient enzyme mixture for fast and reliable quantitative Real-Time PCR, using SG dsDNA-binding dye.
SG Universal Mix	2000 rxns	AM02-200	Compatible with all types of qPCR instruments. Additional tubes with low and high concentration of ROX are included.
<b>AMPLIFY</b> ME	200 rxns	AM04-020	Convenient enzyme mixture for fast and reliable qPCR using probes, including TaqMan®, Scorpions® and molecular beacon probes. It is the best choice for your probe based Real-Time PCR assays, including
Probe No-ROX Mix	2000 rxns	AM04-200	singleplex and multiplex gene expression studies, genotyping experiments or diagnostic assays. Compatible with qPCR instruments that don't need ROX dye.
NA AKCIJI  AMPLIFYME	200 rxns	AM05-020	The AMPLIFYME Probe Universal Mix is a convenient enzyme mixture for fast and reliable qPCR using probes, including TaqMan®, Scorpions® and molecular beacon probes. It is the best choice for your probe based Real-Time PCR assays, including singleplex
Probe Universal Mix	2000 rxns	AM05-200	and multiplex gene expression studies, genotyping experiments or diagnostic assays. Universal – compatible with all types of qPCR instruments. Additional tubes with low and high concentration of ROX are included.
One-Step			
AMPLIFYME Probe One-Step	100 rxns	AM08.1-100	Ready-to-use, 2x concentrated Mix contains all ingredients necessary for Real-Time PCR based on probe detection technology: hot-start <i>Taq</i> polymerase, dNTPs, specially developed buffer,
No-ROX RT-qPCR Mix	500 rxns	AM08.1-500	stabilizers and enhancers. Additionally, Mu-MLV Reverse Transcriptase and RNase Inhibitor are included in separate tubes.
AMPLIFYME Probe One-Step	100 rxns	AM09.1-100	Ready-to-use, 2x concentrated Mix contains all ingredients necessary for Real-Time PCR based on probe detection technology: hot-start <i>Taq</i> polymerase, dNTPs, specially developed buffer,
Probe One-Step Universal RT-qPCR Mix	500 rxns	AM09.1-500	stabilizers and enhancers. Additionally, Mu-MLV Reverse Transcriptase and RNase Inhibitor and ROX solution are included in separate tubes.



# **PCR REAGENTS**

Product Name Pack Size Cat. No. Descriptio	n
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#### Thermostable DNA polymerases from *Thermus aquaticus* (*Taq* Polymerases)

	200 U (5 U/μl)	RP702A	<ul> <li>Taq DNA Polymerase suited to a wide range of</li> </ul>	
TaqNova	500 U (5 U/μl)	RP705A	applications, fast and very efficient; universal and easy-to-use; half-life of the enzyme is 45 minutes	
DNA Polymerase	1000 U (5 U/μl)	RP710A	at 95°C; shows 5'→3' exonuclease activity; does not have 3'→5' exonuclease activity; adds A on	
	2500 U (5 U/μl)	RP725A	the 3' ends.	
	200 Ս (5 Ս/µl)	RP1002	TaqNova DNA-free Polymerase is a 94 kDa recombinant, thermostable Taq DNA polymerase isolated from Thermus aquaticus. It is recommended for a wide range of applications which require DNA synthesis at extremely high temperatures.	
<i>TaqNova</i> DNA-free Polymerase	1000 U (5 U/μl)	RP1010	TagNova DNA-free Polymerase is an universal and easy-to-use DNA polymerase that works rapidly and effectively in various PCR conditions. It is highly purified from DNA contaminants (\$ 1 E. coli genome in 1 U of enzyme), enabling amplification of very	
NA AKCIJI	100 U/μl	RP1000HC (upon request)	conserved sequences (e.g. bacterial 165 rRNA reging without risk of false positive PCR results. The enzyme catalyzes DNA synthesis in a 5'-direction, shows no 3'-5' exonuclease activity. It has a 5'-3' exonuclease activity.	
2x PCR	100 rxns (50 μl)	RP85T	2x concentrated, ready-to-use PCR master mix with	
TaqNova-RED	1000 rxns (50 μl)	RP85T-10	<ul> <li>TaqNova polymerase, that facilitates an easy rapid PCR reaction set-up.</li> </ul>	
	200 U (5 U/μl)	RP902A	<ul> <li>Mixture of thermostable Taq DNA polymerase</li> </ul>	
TaqNovaHS	500 U (5 U/μl)	RP905A	and a highly specific monoclonal antibody, that acts as an inhibitor of the polymerization activity	
DNA Polymerase	1000 U (5 U/µl)	RP910A	(for Hot-Start PCR technique); high PCR specificity with minimal optimization; fast 2-minutes enzyme	
	2500 U (5 U/μl)	RP925A	activation time; very efficient.	
TaqNova Stoffel DNA Polymerase	1000 U (2 U/μl)	RP810	Highly active <i>Taq</i> DNA polymerase without 5'-3' exonuclease activity. <i>TaqNova</i> Stoffel DNA Polymerase works optimally at a broader range of MgCl <sub>2</sub> concentration (2-10 mM) as compared to <i>Taq</i> DNA polymerase – easier and faster optimization. It is also useful for multiplex reactions. In special applications <i>TaqNova</i> Stoffel DNA Polymerase has proven better specificity than regular <i>Taq</i> DNA polymerase. It is especially recommended for amplifications of small fragments from gDNA. The absence of the 5'-3' exonuclease activity makes it very suitable for cycle sequencing. It gives higher sequence intensity and low background.	

Product Name	Pack Size	Cat. No.	Description
PCR Enhancers			
DCD Aven to billion	100 rxns	RP50	PCR additive used for elimination of PCR inhibitors coextracted with DNA;
PCR Anty-inhibitor	500 rxns	RP51	amplification of problematic templates, isolated from: urine, stool, saliva, sputum, blood, swabs, biopsy materials etc.
Deoxyribonucleotides (	INTPs)		
dNTPs MIX 10 mM Total	1 ml	RP63	Deoxyribonucleotides Mix (2.5 mM dATP, 2.5 mM dCTP, 2.5 mM dCTP, 2.5 mM dTP); ultra-pure; supplied as lithium salts (greater stability).
dNTPs MIX 40 mM Total	1 ml	RP64	Deoxyribonucleotides Mix (10 mM dATP, 10 mM dCTP, 10 mM dGTP, 10 mM dTTP); ultra-pure; supplied as lithium salts (greater stability).
dNTPs MIX 100 mM Total	1 ml	RP65	Deoxyribonucleotides Mix (25 mM dATP, 25 mM dCTP, 25 mM dGTP, 25 mM dTTP); ultra-pure; supplied as lithium salts (greater stability).



# **REVERSE TRANSCRIPTION**

Product Name	Pack Size	Cat. No.	Description	
NA AKCIJI				
TRANSCRIPTME	20 rxns	RT31-020	10 pg – 5 μg of total RNA; optimal reaction temp. 50°C contains Enzyme Mix (Reverse Transcriptase and RNas	
RNA KIT cDNA synthesis kit	100 rxns	RT31-100	Inhibitor); 2x Master Mix (oligo(dT) primers, rando hexamers, dNTPs, MgCl <sub>2</sub> ) and RNase H.	
TRANSCRIPTME	10 000 U (200 U/μl)	RT32-010	Modified M-MuLV Reverse Transcriptase; 10 pg – 5 pof total RNA; has increased thermal stability (opt mum activity at 50°C); has no 3'-5' exonuclea	
M-MuLV Reverse Transcriptase	50 000 U (200 U/μl)	RT32-050	and reduced RNase H activity, which improves the synthesis of a full-length cDNA, even from long mRN templates, using random priming; gives high yields first strand cDNA up to 10 kb long.	
TRANSCRIPTME LYO M-MuLV Reverse Transcriptase	100 000 U	RT32L-100	Lyophilized version of M-MuLV Reverse Transcriptas increased thermal stability, that allows the reactit to be carried out at a higher temperature (optimu activity at 50°C); has no 3'→5' exonuclease or RNase activity, which improves the synthesis of a full-leng cDNA, even from long mRNA templates, using rando priming; gives high yields of first strand cDNA up 7 kb long.	
	250 U (5 U/μl)	RT34-025	RNase H is a 18.9 kDa recombinant endoribonucl which hydrolyses specifically the phosphodie bonds of RNA hybridized to DNA.The enzymes not degrade single and double-stranded DNA or u	
RNase H	1250 U (5 U/μl)	RT34-125	bridized RNA. It is a key enzyme in the removal of ml after first-strand cDNA synthesis. Treating cDNA v RNase H prior to PCR can improve sensitivity as I bonded to the cDNA template may prevent bindin the amplification primers in a PCR reaction.	
<i>RIBOPROTECT</i> <i>Hu</i> RNase Inhibitor	2000 U RT35-020 nant human placental prote coli. It inhibits ribonuclease eukaryotic enzymes such as RIBOPROTECT Hu is intend		RIBOPROTECT Hu RNase Inhibitor is a 50 kDa recomb nant human placental protein expressed in Escherich coli. It inhibits ribonuclease (RNase) activity of comme eukaryotic enzymes such as RNase A, RNase B, RNase RIBOPROTECT Hu is intended for use in application	
IMPROVED STABILITY!	10 000 U (40 U/μl)	RT35-100	where the presence of RNases may cause a haze RNA quality and experiment results, e.g. in RN. lation, cDNA synthesis, RT-PCR, in vitro transcri and translation, or RNase-free monoclonal anti preparation. Stable up to 58°C and at min. 0.5 – DTT concentration ranges.	
RIBOPROTECT Hu RNase Inhibitor Lyo-ready	10 000 U	RT35L-010	Formulation of <i>RIBOPROTECT Hu</i> Lyo-ready RNa: Inhibitor (glycerol-free) enables its usage direct in the lyophilization process. <i>RIBOPROTECT Hu</i> Ly	
	(40 U/μl)	RT35L-B (bulk)	ready is recombinant human placental RNase inhibit expressed in E. colistrain that completely inhibits RNa A, B, and C activity. Stable at least 4 weeks at 37° up to 3 freeze/thaw cycles acceptable.	

# **ENZYMES & PROTEINS**

Product Name	Form	Pack Size	Cat. No.	Description
Proteinase K		'		
		100 mg	RP100B	Recombinant Proteinase K from
	Powder	250 mg	RP101B	Tritirachium album expressed in Pichia pastoris is a broad spectrum serine protease. Our recombinant Proteinase K is
	Powder	1000 mg	RP102B	extensively purified to give highly active preparation devoid of any detectable nuclease activities.
NA AKCIJI		bulk	RP103B	It is widely used for digestion of proteins, including DNases and RNases during nucleic acid preparations without
MBG	Cake	on request	RP103B-C	compromising the integrity of the isolated DNA or RNA.  Proteinase K is fully active under denaturing
	Solution	1 ml (20 mg/ml)	RP107B-1	conditions (e.g. in the presence of urea and/ or SDS), what makes it ideal for digesting proteins in variety of applications. Solubility in water ≥20 mg/ml;
		5 ml (20 mg/ml)	RP107B-5	Activity ≥ 30 U/mg lyophilizate; Specific activity ≥ 40 U/mg protein; ≥ 800 U/ml liquid;
		bulk	RP107B	DNA content ≤ 10 pg/mg.
		100 mg	RP100N	Proteinase K NGS Grade is developed for most demanding applications.
Nec	Powder 250 mg	250 mg	RP101N	Additional purification technology resultion its significantly increased solubilit (≥50 mg/ml), increased specific activity
NGS		RP102N	(≥35 U/mg lyophilizate; ≥ 45U/mg protein) and remarkable purity with DNA content ≤0.1 pg/mg.	
		bulk	RP103N	Free of exonucleases, endonucleasesand ribonucleases.



Product Name	Pack Size	Cat. No.	Description	
Nucleases				
<b>Saltonase</b> (HL-Nuclease)	5000 U (20 U/µl)	EN32-050	Saltonase is a cold-active, heat-labile recombinant endonuclease produced in <i>E.coli</i> . Saltonase originates from psychrophilic bacteria and effectively digests all types of DNA and RNA substrates in different buffer conditions and a broad range of temperatures. It is very active in	
	25 000 U (20 U/μl)	EN32-250	demanding conditions, including low temperatures a environment with high salt content. These features m Saltonase extremely useful for removing undesired nucl acids contamination during purification of proteins laboratory and manufacturing workflows.	
<b>Masterase</b> (HL-dsDNase)	500 U (2 U/μl)	EN31-005	Masterase is a 43.3 kDa heat-labile recombinant endonuclease, derived from a cold water eukaryotic organism, expressed in <i>Pichia pastoris</i> . The enzyme displays high specific activity towards double-strandec DNA leaving single-stranded DNA or RNA undamaged in	
	2500 U (2 U/μl)	EN31-025	standard conditions. Masterase can be easily inactivated by heat treatment in moderate temperatures. It is intended for applications where the presence of dsDNA influences experiments' results in thermo-sensitive applications. The enzyme hydrolyzes phosphodiester linkages yielding oligonucleotides with a 5'-phosphate and a 3'-hydroxyl groups.	
<b>DNaseMe</b> (dsDNase)	5000 U (20 U/µl)	EN33-050	DNaseMe is a 42.8 kDa recombinant endonuclease, derived from marine amphipods, expressed in <i>Pichia pastoris</i> . The enzyme displays high specific activity towards double-stranded DNA leaving single-stranded DNA or RNA undamaged in standard conditions. DNaseMe is highly active in a broad spectrum of temperatures, buffer conditions and pH. The specific activity is similar to bovine DNase I however, DNaseMe is characterized	
	25 000 U (20 U/μl)	EN33-250	by higher stability in demanding reaction and storage conditions (e.g. high salt and detergent containing buffers elevated temperature). These features make DNaseMe extremely useful for rapid and "RNA safe" degradation of genomic DNA, where absence of ribonucleases is critical to maintain the integrity of RNA. The enzyme hydrolyzes phosphodiester linkages yielding oligonucleotides with a 5'-phosphate and a 3'-hydroxyl groups.	
RNase A (DNase-free)	50 mg	RP145	The Ribonuclease A is a 13.7 kDa (monomer) endoribo- nuclease isolated from bovine pancreas, which selectively cleaves single-stranded RNA 3' next to pyrimidine residues (cytosine, uracil). The RNase A is used to remove RNA during the isolation procedures of plasmid and genomic DNA. The enzyme is very active under a wide range of reaction conditions and difficult to inactivate.	
RNase H	250 U (5 U/μl)	RT34-025	RNase H is a 18.9 kDa recombinant endoribonuclease, which hydrolyses specifically the phosphodiester bonds of RNA hybridized to DNA. The enzymes does not degrade single and double-stranded DNA or unhybridized RNA. It is a key enzyme	
	1250 U (5 U/μl)	RT34-125	in the removal of mRNA after first-strand cDNA synthesis. Treating cDNA with RNase H prior to PCR can improve sensitivity as RNA bonded to the cDNA template may prevent binding of the amplification primers in a PCR reaction.	

Product Name	Pack Size	Cat. No.	Description		
Other Enzymes & F	Proteins	'			
T4 DNA Ligase	500 U	EN11-050	ATP-dependent recombinant enzyme used fo molecular cloning, site-directed mutagenesis, nic repair in duplex DNA, RNA or DNA/RNA hybrid		
	2500 U	EN11-250	Ligation Mediated PCR; concentration 5 Weiss U.		
Quick Ligase	50 rxns	EN12-050	ATP-dependent recombinant T4 DNA ligase for efficient ligation of DNA fragments with compatible		
	150 rxns	EN12-150	cohesive or blunt ends in 5 and 15 minute respectively. PEG included.		
Tth DNA Ligase	250 U (3750 CEU) (5 U/μl)	EN13-025	NAD-dependent recombinant ligase for Thermus thermophilus. The ligation will or only if oligonucleotides are perfectly paired the complementary target DNA and have no between them. Therefore, a single-base substitution be detected. High thermostability allows ligated using high-stringency hybridization conditions that the properties of SNPs. Equivalent of Ampligation (Epicentre).		
	2500 U (37 500 CEU) (5 U/μl)	EN13-250			
UDGase	500 U	EN19-050	Uracil DNA Glycosylase (UDG) catalyzes the releas of uracil from uracil-containing single-strande or double-stranded DNA, but not from RNA		
	2500 U	EN19-250	oligonucleotides. Widely used to control carry-ov contamination in PCR; concentration 1 U/µl.		
phi29 DNA Polymerase	1000 U (10 U/μl)	EN20-010	Very processive polymerase (up to 70 kb) with stron strand displacement activity, which allows for high efficient isothermal DNA amplification; possess		
	5000 U (10 U/μl)	EN20-050	a 3'→5' exonuclease (proofreading) activity a preferentially on ssDNA or RNA, therefore 3'-1 ified primers are recommended.		
TRANSCRIPTME M-MuLV Reverse Transcriptase	10 000 U (200 U/μl)	RT32-010	Modified M-MuLV Reverse Transcriptase; $10 \text{ pg} - 5 \text{ pof total RNA; concentration } 200  U/µl; has increase thermal stability (optimum activity at 50°C); has 3' \rightarrow 5' exonuclease and reduced RNase H activity$		
	50 000 U (200 U/μl)	RT32-050	which improves the synthesis of a full-length cl even from long mRNA templates, using ran priming; gives high yields of first strand cDN to 10 kb long.		
TRANSCRIPTME LYO M-MuLV Reverse Transcriptase	100 000 U	RT32L-100	Lyophilized version of M-MuLV Rever Transcriptase; increased thermal stability, th allows the reaction to be carried out at a higher terperature (optimum activity at 50°C); has no 3'-exonuclease or RNase H activity, which improve the synthesis of a full-length cDNA, even from lom RNA templates, using random priming; gives highelds of first strand cDNA up to 7 kb long.		



Product Name	Pack Size	Cat. No.	Description
Other Enzymes &	Proteins		
RIBOPROTECT Hu RNase Inhibitor IMPROVED STABILITY!	2000 U (40 U/µl)	RT35-020	RIBOPROTECT Hu RNase Inhibitor is a 50 kDa recombinant human placental protein expressed in Escherichia coli. It inhibits ribonuclease (RNase) activity of common eukaryotic enzymes such as RNase A, RNase B, RNase C. RIBOPROTECT Hu is intended for use in applications where the
	10 000 U (40 U/µl)	RT35-100	presence of RNases may cause a hazard to RNA quality and experiment results, e.g. in RNA isolation, cDNA synthesis, RT-PCR, in vitro transcription and translation, or RNase-free monoclonal antibody preparation. Stable up to 58°C and at min. 0.5 – 1 mM DTT concentration ranges.
RIBOPROTECT Hu RNase Inhibitor Lyo-ready	10 000 U (40 U/μl)	RT35L-010	Formulation of <i>RIBOPROTECT Hu</i> Lyo-ready RNase Inhibitor (glycerol-free) enables its usage directly in the lyophilization process. <i>RIBOPROTECT Hu</i> Lyo-ready is recombinant human placental RNase
		RT35L-B (bulk)	inhibitor expressed in <i>E. coli</i> strain that completely inhibits RNase A, B, and C activity. Stable at least 4 weeks at 37°C; up to 3 freeze/thaw cycles acceptable.

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