

# 2020 Product Catalogue



## Reagents for Molecular Biology Research

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## **Overview of Vazyme**

#### Vazyme: InoVation in Enzyme Technology

With the faith of "InnoVation in Enzyme Technology", Vazyme Biotech Co., Ltd. has passionately focused on developing enzyme and antibody technologies and products for years. The Vazyme Biotech is now staffed by more than 1,000 employees. The headquarter is located in Nanjing of China with a R&D / manufacturing base that covers 25,000 m² and a GMP workshop of 4,000 m². Vazyme has developed a powerful sales network in China and is expanding into international markets.



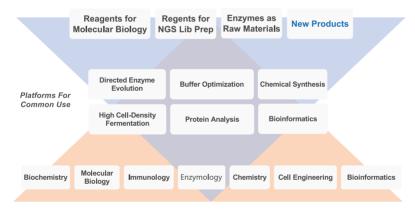




#### **Vazyme Technologies and Products**

With years of experience, Vazyme has developed six technology platforms that can be commonly used for R&D and manufacturing, including (1) directed enzyme evolution, (2) buffer optimization, (3) chemical synthesis, (4) high cell-density fermentation, (5) protein analysis, and (6) bioinformatics.

Based on the above platforms, Vazyme now provides a variety of products, solutions, and services, which fall into three major product lines, including (1) solutions for molecular biology research, (2) solutions for Next-Generation Sequencing (NGS) library preparation, and (3) enzymes as raw materials for industrial use.



### **Developing Technologies to Improve Human Health**

Fascinated by the enzyme and antibody technologies, we regard enzymes and antibodies as the key factor of the biotechnology industry. Vazyme's vision is to develop technology to improve human health.

Experts for Experts

## **2020 Vazyme Product Catalogue**

## **Reagents for Molecular Biology Research**

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### **PCR**

## High-Fidelity PCR

	Product Name	Size	Cat. No.#
HOT	Phanta Max Super-Fidelity DNA Polymerase	100 U / 500 U / 1,000 U	P505-d1/d2/d3
HOT	2 × Phanta Max Master Mix	1 ml / 5 ml / 15 ml	P515-01/02/03
New	2 ×Phanta Max Master Mix (Dye Plus)	1 ml / 5 ml / 15 ml	P525-01/02/03

### ■ Conventional PCR

Product Name	Size	Cat. No.#
Taq DNA Polymerase (Mg <sup>2+</sup> plus Buffer)	1,000 U / 5,000 U / 10,000 U	P101-01/02/03
Taq DNA Polymerase (Mg <sup>2+</sup> free Buffer)	1,000 U / 5,000 U / 10,000 U	P102-01/02/03
Taq DNA Polymerase (Mg <sup>2+</sup> plus Buffer, with dNTP)	1,000 U / 5,000 U / 10,000 U	P101-d1/d2/d3
Taq DNA Polymerase (Mg <sup>2+</sup> free Buffer, with dNTP)	1,000 U / 5,000 U / 10,000 U	P102-d1/d2/d3
2 × Taq Master Mix	5 ml / 15 ml / 50 ml	P111-01/02/03
2 × Taq Master Mix (Dye Plus)	5 ml / 15 ml / 50 ml	P112-01/02/03
Green Taq Mix	5 ml / 15 ml / 50 ml	P131-01/02/03

## High-Yield PCR

Product Name	Size	Cat. No.#
Taq Plus DNA Polymerase	250 U / 1,000 U / 3,000 U	P201-01/02/03
Taq Plus DNA Polymerase (with dNTP)	250 U / 1,000 U / 3,000 U	P201-d1/d2/d3
2 × Taq Plus Master Mix	5 ml / 15 ml / 50 ml	P211-01/02/03
2 × Taq Plus Master Mix II (Dye Plus)	5 ml / 15 ml / 50 ml	P213-01/02/03

## ■ Long-Fragment PCR

Product Name		Cat. No.#
Vazyme LAmp DNA Polymerase (Mg <sup>2+</sup> plus buffer)	125 U / 500 U	P301-01/02
Vazyme LAmp DNA Polymerase (Mg <sup>2+</sup> plus buffer, with dNTP)	125 U / 500 U	P301-d1/d2
Vazyme LAmp DNA Polymerase (Mg <sup>2+</sup> free buffer, with dNTP)	125 U / 500 U	P302-d1/d2
2 × Vazyme LAmp Master Mix	1 ml / 5 ml / 15 ml	P311-01/02/03
2 × Vazyme LAmp Master Mix (Dye Plus)	1 ml / 5 ml / 15 ml	P312-01/02/03

## ■ Direct PCR

	Product Name	Size	Cat. No.#
HOT	One Step Mouse Genotyping Kit	200 rxn	PD101-01
	One Step U* Probe Mouse Genotyping Kit	200 rxn	PD104-01
	Blood Direct PCR Kit V2	50 rxn / 200 rxn	PD103-01/02
	Plant Direct PCR Kit	50 rxn / 200 rxn	PD105-01/02

### Rapid PCR

	Froduct Name	3120	Cat. No.#
		5 ml / 15 ml	P222-01/02
HOT	2 × Rapid Taq Master Mix	50 ml (50 x 1 ml)	P222-03
		50 ml (10 x 5 ml)	P222-04



#### ■ Hot-Start PCR

	Product Name		Cat. No.#
HOT	AceTaq DNA Polymerase	250 U / 1,000 U / 3,000 U	P401-d1/d2/d3
	2 × AceTaq Master Mix	1 ml / 5 ml /15 ml	P411-01/02/03
	2 × AceTaq Master Mix (Dye Plus)	1 ml / 5 ml / 15 ml	P412-01/02/03
	Champagne Taq antibody	500 U	P121-01
HOT	Champagne Taq DNA Polymerase	500 U (2.5 / 5 / 10 U/μl)	P122-d1/d2/d3

## ■ Multiplex PCR

Product Name		Cat. No.#
Multiplex PCR Kit	50 rxn / 200 rxn / 1,000 rxn	PM101-01/02/03

## ■ Isothermal Amplification

Product Name		Cat. No.#
Bst DNA Polymerase Large Fragment	800 U / 8,000 U	P701-01/02

#### PCR-Related

Product Name	Size	Cat. No.#
PCR Enhancer	500 µl	P021-01
dNTP Mix (10 mM each)	1 ml / 5 ml	P031-01/02
dNTP Mix (2.5 mM each)	1 ml / 5 ml	P032-01/02
Heat-labile UDG	100 U / 500 U	P051-01/02
E.coli UDG	500 U / 5,000 U	P061-01/02

## Cloning / Mutagenesis

### ■ Fast Cloning

Product Name		
ClonExpress II One Step Cloning Kit	25 rxn / 50 rxn	C112-01/02
ClonExpress MultiS One Step Cloning Kit	10 rxn / 25 rxn	C113-01/02
ClonExpress Ultra One Step Cloning Kit	25 rxn / 50 rxn	C115-01/02

## ■ Fast Mutagenesis

Product Name		
Mut Express II Fast Mutagenesis Kit V2	10 rxn / 25 rxn	C214-01/02
Mut Express MultiS Fast Mutagenesis Kit V2	10 rxn / 25 rxn	C215-01/02

## ■ TA Cloning

Product Name		Cat. No.#
T4 DNA Ligase	40,000 U	C301-01
New 5min Universal Ligation Mix	50 rxn / 100 rxn	C311-01/02

## **■ TOPO Cloning**

Product Name		
5min TA/Blunt-Zero Cloning Kit	25 rxn / 50 rxn	C601-01/02



## Nucleic Acid Electrophoresis

#### GelRed Nucleic Acid Stain

	Product Name		Cat. No.#
HOT	Ultra GelRed Nucleic Acid Stain (10000 ×)	0.5 ml / 5 ml / 50 ml	GR501-01/02/03

#### DNA Marker

Product Name		Cat. No.#
DL2000 Plus DNA Marker	250 µl / 500 µl	MD101-01/02
DL5000 DNA Marker	250 μΙ / 500 μΙ	MD102-01/02
DL15000 DNA Marker	250 µl / 500 µl	MD103-01/02
100 bp DNA Ladder	250 μl / 500 μl	MD104-01/02

## Reverse Transcription

#### ■ Conventional RT-PCR

	Product Name	Size	Cat. No.#
НОТ	HiScript III Reverse Transcriptase	10,000 U	R302-01
	HiScript II 1st Strand cDNA Synthesis Kit	50 rxn / 100 rxn (20 μl / rxn)	R211-01/02
	HiScript III 1st Strand cDNA Synthesis Kit (+gDNA wiper)	50 rxn / 100 rxn (20 μl / rxn)	R312-01/02
	M-MLV(H-) Reverse Transcriptase	10,000 U	R021-01
	Murine RNase inhibitor	2,000 U / 10,000 U / 20,000 U	R301-01/02/03

## ■ RT-qPCR SuperMix

НОТ	Product Name		Cat. No.#
	HiScript II Q RT SuperMix for qPCR	100 rxn (20 μl / rxn)	R222-01
	HiScript III RT SuperMix for qPCR (+gDNA wiper)	100 rxn (20 μl / rxn)	R323-01
	HiScript II Q Select RT SuperMix for qPCR	100 rxn (20 μl / rxn)	R232-01
	HiScript II Q Select RT SuperMix for qPCR (+gDNA wiper)	100 rxn (20 μl / rxn)	R233-01

## ■ One-Step RT-PCR

Product Name		Cat. No.#
HiScript II One Step RT-PCR Kit	50 rxn (50 μl / rxn)	P611-01
HiScript II One Step RT-PCR Kit (Dye Plus)	50 rxn (50 μl / rxn)	P612-01

## ■ Single Cell Sequence Amplification

Product Name		Cat. No.#
Single Cell Sequence Specific Amplification Kit	200 rxn	P621-01

#### miRNA

## miRNA Reverse Transcription

Product Name	Size	Cat. No.#
miRNA 1st Strand cDNA Synthesis Kit (by stem-loop)	50 rxn / 100 rxn (20 μl / rxn)	MR101-01/02

### miRNA qPCR

Product Name	Size	Cat. No.#
miRNA Universal SYBR® qPCR Master Mix	125 rxn / 500 rxn (20 μl / rxn)	MQ101-01/02



## **qPCR**

## qPCR Master Mix (SYBR)

	Product Name	Size	Cat. No.#
HOT	ChamQ Universal SYBR® qPCR Master Mix	500 rxn / 2,500 rxn (20 μ <b>l</b> / rxn)	Q711-02/03
	AceQ Universal SYBR® qPCR Master Mix	500 rxn / 2,500 rxn (20 μl / rxn)	Q511-02/03
	AceQ qPCR SYBR® Green Master Mix	500 rxn / 2,500 rxn (20 μl / rxn)	Q111-02/03

## qPCR Master Mix (Probe)

	Product Name		Cat. No.#
	AceQ qPCR Probe Master Mix	500 rxn / 2,500 rxn (20 μl / rxn)	Q112-02/03
HOT	AceQ Universal U+ Probe Master Mix V2	500 rxn / 2,500 rxn (20 μl / rxn)	Q513-02/03
HOT	ChamQ Geno-SNP Probe Master Mix	500 rxn / 2,500 rxn (20 μl / rxn)	Q811-02/03

## One-Step qRT-PCR Mix

Product Name		Cat. No.#
HiScript II One Step qRT-PCR SYBR® Green Kit	250 rxn (20 μl / rxn)	Q221-01
HiScript II One Step qRT-PCR Probe Kit	250 rxn (20 μl / rxn)	Q222-01
HiScript II U+ One Step qRT-PCR Probe Kit	250 rxn (20 μl / rxn)	Q223-01

## Genome Editing

Product Name	Size	Cat. No.#
Cas9 Nuclease	50 pmol / 250 pmol	EN301-01/02
T7 Endonuclease I	50 pmol / 250 pmol	EN303-01/02

## In Vitro Transcription

	Product Name		
	T7 High Yield RNA Transcription Kit	50 rxn / 100 rxn	TR101-01/02
нот	T7 RNAi Transcription Kit	25 rxn / 50 rxn	TR102-01/02

## Nucleic Acid Isolation

## ■ Room Temperature RNA Isolation

	Product Name	Size	Cat. No.#
New	RNA-easy Isolation Reagent	100 ml / 200 ml	R701-01/02

## ■ Total RNA Isolation (Traditional)

Product Name	Size	Cat. No.#
RNA isolater Total RNA Extraction Reagent	100 ml	R401-01
Bacteria RNA Plus Reagent	20 ml	R402-01
Bacterial RNA Extraction Kit	100 rxn	R403-01



## Rapid Sample Treatment

	Product Name	Size	Cat. No.#
New	RoomTemp Sample Lysis Kit	250 rxn (5 ml) / 1000 rxn (20 ml) / 5000 rxn (100 ml)	P073-01/02/03

## ■ RNA Isolation (Column)

Product Name	Size	Cat. No.#
FastPure Cell / Tissue Total RNA Isolation Mini Kit	50 rxn	RC101
MiPure Cell / Tissue miRNA Kit	50 rxn	RC201
FastPure Plant Total RNA Isolation Kit (Polysaccharides / Polyphenolics-Rich)	50 rxn	RC401

## ■ DNA Isolation (Column)

,		
Product Name	Size	Cat. No.#
FastPure Blood DNA Isolation Mini Kit V2	50 rxn / 200 rxn	DC111-01/02
FastPure Cell/Tissue DNA Isolation Mini Kit	100 rxn	DC102
FastPure Bacteria DNA Isolation Mini Kit	100 rxn	DC103
FastPure Plant DNA Isolation Mini Kit	50 rxn	DC104
FastPure FFPE DNA Isolation Kit	50 rxn	DC105
Lysozyme	200 mg	DE103

## ■ Plasmid DNA Isolation

	Product Name	Size	Cat. No.#
	FastPure Plasmid Mini Kit	100 rxn	DC201
HOT	FastPure EndoFree Plasmid Maxi Kit	10 rxn	DC202

#### ■ Gel DNA Isolation

Product Name	Size	Cat. No.#
FastPure Gel DNA Extraction Mini Kit	100 rxn	DC301

### **■** Tissue Stabilizer

Product Name	Size	Cat. No.#
RNA Keeper Tissue Stabilizer	100 m <b>i</b>	R501-01
RNA Keeper -ICE Tissue Transition Buffer	100 ml	R502-01

#### **■** Exosome Isolation

Product Name	Size	Cat. No.#
VEX Exosome Isolation Reagent (from cell culture media)	50 m <b>l</b>	R601
VEX Exosome Isolation Reagent (from serum)	10 ml	R602
VEX Exosome Isolation Reagent (from plasma)	10 ml	R603



## Cell Biology / Protein Research

## Apoptosis

	Product Name	Size	Cat. No.#
HOT	TUNEL FITC Apoptosis Detection Kit	20 rxn / 50 rxn / 100 rxn	A111-01/02/03
	TUNEL BrightGreen Apoptosis Detection Kit	20 rxn / 50 rxn / 100 rxn	A112-01/02/03
	TUNEL BrightRed Apoptosis Detection Kit	20 rxn / 50 rxn / 100 rxn	A113-01/02/03
	Annexin V-FITC/PI Apoptosis Detection Kit	50 rxn / 100 rxn	A211-01/02
New	Annexin V-PE/7-AAD Apoptosis Detection Kit	50 rxn / 100 rxn	A213-01/02

## ■ C∈II Counting

	Product Name		Cat. No.#
HOT	CCK-8 Cell Counting Kit	500 rxn / 1,000 rxn	A311-01/02

### Cell Transfection

Product Name		Cat. No.#
ExFect 2000 Transfection Reagent	0.5 ml / 1 ml / 5 ml	T202-01/02/03

## ■ Dual Luciferase Reporter Assay

	Product Name	Size	Cat. No.#
HOT	Dual Luciferase Reporter Assay Kit	100 rxn	DL101-01

## Mycoplasma

	Product Name	Size	Cat. No.#
HOT	MycoBlue Mycoplasma Detector	20 rxn / 50 rxn	D101-01/02
	Myco-Off Mycoplasma Cleaner	100 µl / 500 µl / 1,000 µl	D103-01/02/03

## ■ ECL for Western Blotting

	Product Name	Size	Cat. No.#
HOT	High-Sensitive ECL Chemiluminescence Detection Kit (Ready-to-Use)	2 × 50 ml / 250 ml (A / B)	E412-01/02

#### ■ Protein Quantification

Product Name		Cat. No.#
Bradford Protein Quantification Kit	500 rxn / 1,000 rxn	E111-01/02
Detergent Compatible Bradford Protein Quantification Kit	500 rxn	E211-01
BCA Protein Quantification Kit	250 rxn / 500 rxn	E112-01/02



## **PCR**

## **Selection Guide**

Applications	Products (Cat.#)	Features	Applicable for
Conventional PCR	2× Taq Master Mix (#P111) 2× Taq Master Mix (Dye Plus) (#P112) Green Taq Mix (#P131)	No 3' -> 5' exonuclease activity. Excellent compatibility. Products contain A at 3'-end.	Colony PCR; Large-scale gene identification; TA Cloning for small fragments.
High-Yield PCR	2× Taq Plus Master Mix (#P211) 2× Taq Plus Master Mix II (Dye Plus) (#P213)	With fidelity 6-fold higher than Taq. Mixed products with 3"-end blunt or containing A.	PCR that requires some fidelity.
Rapid PCR	2× Rapid Taq Master Mix (#P222)	Amplification speed: up to 15 sec / kb.	Colony PCR.
Long-Fragment PCR	2× Vazyme LAmp Master Mix (#P311) 2× Vazyme LAmp Master Mix (Dye Plus) (#P312)	Efficiently amplify fragments > 20 kb.	Long-fragment amplification.
Hot-Start PCR	2× AceTaq Master Mix (#P411) 2× AceTaq Master Mix (Dye Plus) (#P412) Champagne Taq Antibody (#P121) Champagne Taq DNA Polymerase (#P122)	Excellent specificity. Excellent sensitivity.	Amplification that requires higher sensitivity and specificity; Amplification of genes with low copy or qPCR assay from complex templates (genomic DNA, cDNA).
Multiplex PCR	Multiplex PCR Kit (#PM101)	19-plex PCR in one single reaction.	Detection or typing of pathogens.
Direct PCR	One Step Mouse Genotyping Kit (#PD101) Blood Direct PCR Kit V2 (#PD103) Plant Direct PCR Kit (#PD105)	Easy and fast, without DNA purification.	One step mouse genotyping; Direct PCR from plant tissues; Direct PCR from blood.
High-Fidelity PCR	Phanta Max Super-Fidelity DNA Polymerase (#P505) 2× Phanta Max Master Mix (#P515) 2× Phanta Max Master Mix (Dye Plus) (#P525)	With super fidelity 53-fold higher than Taq; High resistance to PCR inhibitors.	High-fidelity PCR. Amplification of templates with high GC-content; Long-fragment (up to 40 kb) amplification.



## **High-Fidelity PCR**



2× Phanta Max Master Mix (#P515)

→ 2× Phanta Max Master Mix (Dye Plus) (#P525)



Super Fidelity: 53-fold higher than Tag DNA Polymerase.

Long Fragment: amplify fragments up to 40 kb.

Suitable for templates with high GC-content.

Suitable for Direct-PCR using crude materials as templates\*.

<sup>\*</sup> Validated crude materials: bacteria, fungi, whole blood, cultured cells, plant or animal tissue lysate, food lysates, etc.



#### **Selected Product Citations**

Zhao Q, et al. Metabolic coupling of two small-molecule thiols programs the biosynthesis of lincomycin A. *Nature*, 2015, 518(7537):115-9.

Tian Z, et al. An enzymatic [4+2] cyclization cascade creates the pentacyclic core of pyrroindomycins. *Nature Chemical Biology*, 2015, 11(4):259-65.

Han X, et al. Mapping the Mouse Cell Atlas by Microwell-Seq. Cell, 2018, 172(5):1091-107.

Cheng X, et al. Pacer Mediates the Function of Class III PI3K and HOPS Complexes in Autophagosome Maturation by Engaging Stx17. *Molecular Cell*, 2017, 65(6):1029-43.

Lv M, et al. Characterization of a C3 Deoxygenation Pathway Reveals a Key Branch Point in Aminoglycoside Biosynthesis. *Journal of the American Chemical Society*, 2016, 138(20):6427-35.



## **High-Yield PCR**

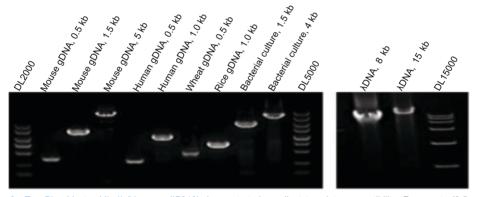


→ 2× Taq Plus Master Mix II (Dye Plus) (#P213)

#### **Features**

- \* Robust performance for high-yield PCR in most primer-template systems.
- \* Ready-to-use master mix with no need for operations on ice.
- \* PCR products can be directly loaded for electrophoresis with no need for loading buffer.

#### Validation Data



2× Taq Plus Master Mix II (Vazyme, #P213) demonstrated excellent template compatibility. Fragments (0.5 kb to 15 kb) were amplified from genomic DNA (mouse, human, wheat, rice), bacterial culture, and  $\lambda$ DNA, respectively. A specific corresponding band was observed in each PCR.

## ~W

#### Selected Product Citations

Zhang X, et al. (2014) Complementary sequence-mediated exon circularization. Cell, 159(1):134-47.

Yuan H, et al. Gyrl-like proteins catalyze cyclopropanoid hydrolysis to confer cellular protection. *Nature Communications*, 2017, 8(1).1485.



## Rapid PCR

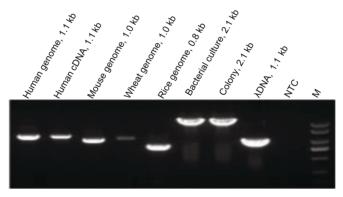


## 2× Rapid Taq Master Mix (#P222)

#### **Features**

- \* Rapid: amplification speed is 15 sec / kb, with an extreme speed of 1 sec / kb for fragments within 1 kb.
- \* Ready-to-use master mix with no need for operations on ice.
- \* PCR products can be directly loaded for electrophoresis with no need for loading buffer.
- \* Excellent stability: remains stable after 50 freeze-thaw cycles.

#### Validation Data



Fragments (1 kb - 2 kb) was amplified from genomic DNA (human, mouse, wheat, rice), cDNA (human), bacterial culture, colony, and λ DNA, respectively. The extension time was set as 1 sec / kb. Ten μl of PCR product was loaded for agarose gel electrophoresis. Specific bands were observed.

#### Selected Product Citations

Zhang B, et al. Enzyme-catalysed [6+4] cycloadditions in the biosynthesis of natural products. Nature, 2019, 568(7750):122-6.

Wang YS, et al. Molecular Basis for the Final Oxidative RearrangementSteps in Chartreusin Biosynthesis. J Am Chem Soc, 2018, 140(34):10909-14.



## **Multiplex PCR**

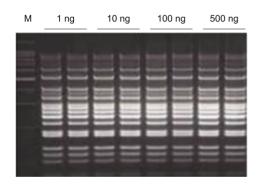


→ Multiplex PCR Kit (#PM101)

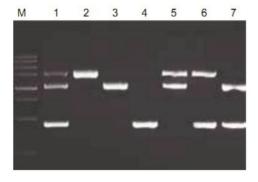
#### **Features**

- \* Multiplex: 19-plex PCR or even higher.
- \* Excellent target-to-target amplification uniformity and extremely low target preference.
- \* Highly sensitive amplification from trace amount of genomic DNA (>= 1 ng).

#### **Validation Data**



Uniform amplification coverage of different regions. Human genomic DNA was used as template for 19-plex PCR. The size of the amplicons ranged from 70 bp to 916 bp. The result indicated that Multiplex PCR Kit (Vazyme, #PM101) has a uniform amplification coverage of different regions for 1 ng-500 ng of template.



The Multiplex PCR Kit showed excellent compatibility with fragment length. Mouse genomic DNA was used as template for amplification of 1.55 kb, 1.07 kb, and 0.45 kb fragments, respectively. The result indicated that Multiplex PCR Kit (Vazyme, #PM101) is compatible with amplicons of various lengths in one single reaction system.

1: 3-plex PCR 2-4: 1-plex PCR 5-7: 2-plex PCR M: DL5000 DNA Marker



## **Cloning / Mutagenesis**

## Selection Guide

Applications			Applicable for
Fast Cloning	ClonExpress Ultra One Step Cloning Kit (#C115) ClonExpress II One Step Cloning Kit (#C112) ClonExpress MultiS One Step Cloning Kit (#C113)	Easy, fast, and efficient. No need to consider the restriction enzyme cutting sites on the inserts. Ligase-independent. Positive Clone Rate > 95%. Efficient doning of fragments of 50 bp - 10 kb.	Cloning or assembly of 1-5 fragments.
Fast Mutagenesis	Mut Express II Fast Mutagenesis Kit V2 (#C214) Mut Express MultiS Fast Mutagenesis Kit V2 (#C215)	Efficient amplification of any plasmids within 20 kb. Site-directed mutations of 1-5 discontinuous sites in one reaction.	1-5 separate site-directed mutagenesis on one plasmid.
TOPO Cloning	5min TA/Blunt-Zero Cloning Kit (#C601)	Cloning within 5 min. Positive Clone Rate > 95%	TA cloning. cloning with blunt ends.

## **TOPO Cloning**





#### **Features**

- \* Ready-to-use master mix.
- \* Suitable for both TA cloning and blunt-end cloning.
- \* Rapid cloning within 5 min.
- \* High cloning efficiency with Positive Clone Rate > 95%.
- \* Ampicillin and Kana dual resistance vector.

#### Workflow





## **Fast Cloning**



## → ClonExpress Ultra One Step Cloning Kit (#C115)

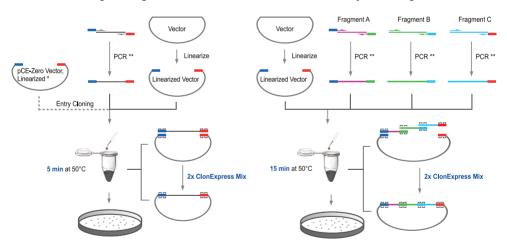
#### **Features**

- \* Cloning within 5 min.
- \* Ready-to-use super mix in one tube.
- \* Efficient cloning of fragments of 50 bp 10 kb with Positive Clone Rate > 95%.
- \* Suitable for cloning of 1 fragment, assembly of 2 5 fragments, and entry cloning.
- \* Independent of DNA ligase, significantly reducing the self-ligated colonies.

#### Mechanism

#### Cloning of 1 Fragment

#### Assembly of 2 - 5 fragments.



<sup>\*</sup> pCE-Zero Vector, Linearized, is supplied with ClonExpress Ultra One Step Cloning Kit (Vazyme, #C115).

## Selected Product Citations of ClonExpress

Wu N, et al. TBX6 null variants and a common hypomorphic allele in congenital scoliosis. New England Journal of Medicine, 2015, 372(4):341-50.

Ge J, et al. Architecture of the mammalian mechanosensitive Piezo1 channel. Nature, 2015, 527(7576):64-9.

<sup>\*\*</sup> It is highly recommended to use Vazyme's APP - "CE Design" - for easy primer design.



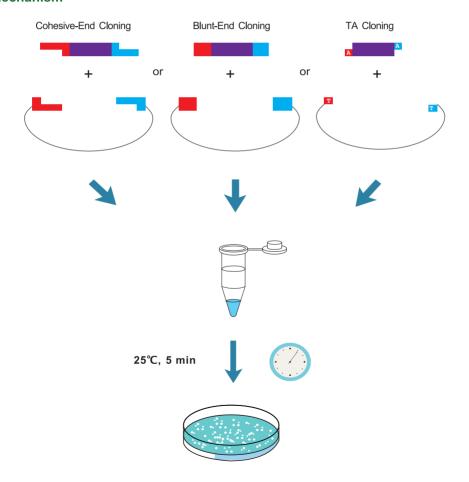


## → 5min Universal Ligation Mix (#C311)

#### **Features**

- \* Versatile: Suitable for TA cloning, blunt-end cloning, cohesive-end cloning, and ligation of linkers or adapters.
- \* Fast: Cloning within 5 min at 25°C.
- \* Efficient: Positive Clone Rate > 95%.

#### Mechanism





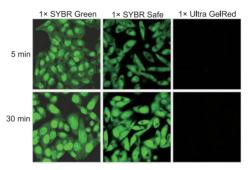
## GelRed



→ Ultra GelRed Nucleic Acid Stain (10000×) (#GR501)



No toxicity



Ultra GelRed is unable to cross cell membranes.

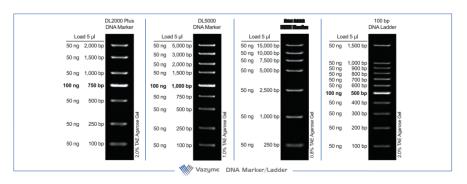
## DNA Marker/ Ladder



## → DNA Markers / Ladders

#### Stable

#### **Clear Bands**



## **Reverse Transcription**



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Applications	-	-						
RT-qPCR								
RT-PCR								
Features								
SuperMix								
Long-fragment cDNA								
Rapid removal of Genomic DNA								
Primers								
Oligo dT <sub>23</sub> VN / N6 Mix								
Optiona <b>l</b>								

	M-MLV (H-) (#R021)	HiScript II Reverse Transcriptase (#R201)	HiScript III Reverse Transcriptase (#R302)
Reaction temperature	37℃ - 42℃	42℃ - 55℃	37℃ - 50℃
Thermal stability	***	***	<b>☆☆☆</b>
RNase H activity	No	No	No
cDNA length	2 kb-3 kb	Up to 20 kb	Up to 20 kb
Template adaptability	***	አ <del>ለ</del> አለአ	<b>☆☆☆☆</b>

## RT-qPCR SuperMix



## $\rightarrow$ 1

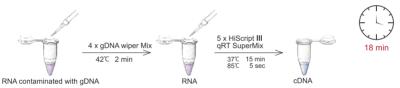
## HiScript III RT SuperMix for qPCR (+gDNA wiper) (#R323)

#### **Features**

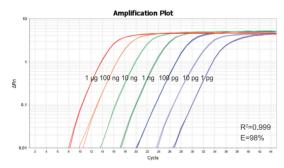
- \* Ready-to-use SuperMix: reverse transcription within 20 min by only adding template RNA.
- \* Excellent efficiency for low-input RNA or degraded RNA.
- \* Excellent tolerance for impurities (i.e. ethanol, isopropanol, phenol water, guanidine thiocyanate, humic acid).
- \* Lower C<sub>T</sub> value and higher efficiency than most other commercially available reverse transcription reagents.

#### Validation Data

#### 1. Easy & Fast



#### 2. Excellent Sensitivity



RNA of HeLa cells was serially diluted and reverse transcribed using HiScript III RT SuperMix for qPCR (+gDNA wiper) (Vazyme, #R323), followed by qPCR detection of gene ACTB. The results show an excellent linear relationship across a wide range of RNA concentrations. The target gene (ACTB) was detected in 1 pg of RNA.



## **qPCR**



Applications	Products (Cat.#)		
SYBR	ChamQ Universal SYBR QPCR Master Mix (#Q711)		
Probe	AceQ Universal U+ Probe Master Mix V2 (#Q513)		
SNP (TaqMan MGB Probe)	ChamQ Geno-SNP Probe Master Mix (#Q811)		

## qPCR Master Mix (SYBR)



→ ChamQ Universal SYBR® qPCR Master Mix (#Q711)

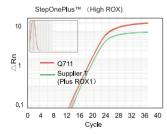
## Best Combination of Specificity + Sensitivity

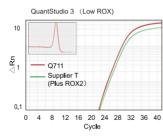
Unique Hot-Start Tag

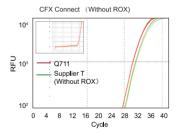
- Unique specificity-promoting Factors
- Optimal Concentrations of Mg<sup>2+</sup> and Dye
- Universal

#### Validation Data

#### Applicable for almost all qPCR instruments.







## *~*₩

#### Selected Product Citations

Xu L, et al. The transcription factor TCF-1 initiates the differentiation of TFH cells during acute viral infection. *Nature Immunology*, 2015, 47(3):538-51.

Guo C, et al. Cholesterol Homeostatic Regulator SCAP-SREBP2 Integrates NLRP3 Inflammasome Activation and Cholesterol Biosynthetic Signaling in Macrophages. *Immunity*, 2018, 49(5): 842-56.



## qPCR Master Mix (Probe)



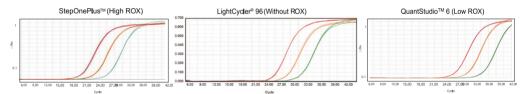
→ AceQ Universal U+ Probe Master Mix V2 (#Q513)

#### **Features**

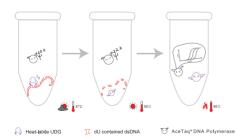
- \* Excellent sensitivity: Hot-start AceTaq and optimal buffer ensure high sensitivity and effectively inhibit non-specific amplification.
- \* Excellent linear relationship over a large range of input amount of template. Suitable for the detection of single-copy templates.
- \* Anti-contamination: the dUTP/UDG system eliminates possible contaminations and ensures reliable results.
- \* Universal: applicable for almost all qPCR instruments.

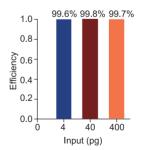
#### Validation Data

1. Applicable for almost all qPCR instruments.



#### 2. dUTP/UDG system.





For Vazyme #Q513, the removal rate of the contaminated template is as high as 99.6%, effectively ensuring the accuracy of experimental results. U-containing templates (4 pg, 40 pg) were added respectively to the reaction system to evaluate the removal efficiency of the contaminated template by Vazyme #Q513.



## qPCR Master Mix (Probe)



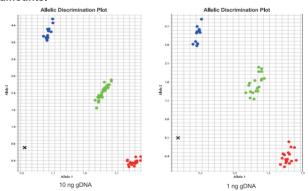
→ ChamQ Geno-SNP Probe Master Mix (#Q811)

#### **Advantages**

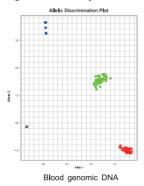
- \* Compatible with 1 ng 10 ng of input genomic DNA.
- \* Accurate genotyping of SNP sites with GC-content of 25% 73%.
- \* Excellent stability: stable signal and accurate genotyping results can be obtained both 72 hr pre-PCR and 72 hr post-PCR.
  - \* 72 hr pre-PCR: PCR reaction solutions were prepared and left in darkness (at room temperature) for 72 hr before PCR;
  - \* 72 hr post-PCR: after PCR, the samples were left in darkness (at room temperature) for 72 hr.
- \* Blood lysate can be directly used as a template for SNP genotyping, with no need for blood genomic DNA extraction.

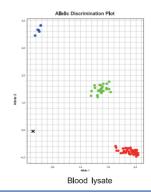
#### Validation Data

#### 1. Flexible input amounts.



#### 2. Direct genotyping with blood lysate.







## **Nucleic Acid Isolation**

## Selection Guide

	Rapid Sample Treatment	Blood	RoomTemp Sample Lysis Kit	P073
DNA Isolation & Purification	DNA Extraction (Column)	Blood	FastPure Blood DNA Isolation Mini Kit V2	DC111
		Cell/tissue	FastPure Cell/Tissue DNA Isolation Mini Kit	DC102
		Bacterial	FastPure Bacteria DNA Isolation Mini Kit	DC103
		Plant	FastPure Plant DNA Isolation Mini Kit	DC104
		FFPE	FastPure FFPE DNA Isolation Kit	DC105
		Lysozyme	Lysozyme	DE103
	Plasmid DNA extraction	Plasmid	FastPure Plasmid Mini Kit	DC201
		Endotoxin-free plasmid	FastPure EndoFree Plasmid Maxi Kit	DC202
	DNA Products recycle	Gel DNA Recycle / DNA purification	FastPure Gel DNA Extraction Mini Kit	DC301
RNA Isolation & Purification	RNA tissue Keeper	RNA Keeper for fresh tissue	RNA Keeper Tissue Stabilizer	R501
		RNA Keeper for thawing /frozen tissue	RNA Keeper-ICE Tissue Transition Buffer	R502
	RoomTemp RNA extraction	Total RNA	RNA-easy Isolation Reagent	R701
	Traditional total RNA extraction	Total RNA	RNA isolater Total RNA Extraction Reagent	R401
		Bacterial lysate	ysate Bacteria RNA Plus Reagent	
		Bacterial total RNA	Bacteria RNA Extraction Kit	R403
	Column RNA Extraction	Cell/tissue total RNA	FastPure Cell/Tissue Total RNA Isolation Mini Kit	RC101
		Cell/tissue miRNA	MiPure Cell/Tissue miRNA Kit	RC201
		Polysaccharide & Polyphenol-rich Plant total RNA	FastPure Plant Total RNA Isolation Kit (Polysaccharides & Polyphenolics-rich)	RC401
Exosome Isolation	Cell supernatant		VEX Exosome Isolation Reagent (from cell culture media)	R601
	Serum		VEX Exosome Isolation Reagent (from serum)	R602
	Plasma		VEX Exosome Isolation Reagent (from plasma)	R603

## **Room Temperature RNA Isolation**



 $\rightarrow$ 

RNA-easy Isolation Reagent (#R701)

#### **Features**

- \* Single-phase extraction with no need of multi-phase partitioning; all operations are at room temperature:
- \* Widely applicable to easy RNA isolation from all kinds of animal, plant, and microbial samples, as well as various cultured cells.
- \* Products with high purity and integrity, and with yield comparable to that of traditional Trizol extraction.
- \* No need to use toxic and harmful reagents such as chloroform.

#### Validated Samples

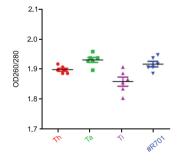
Animal tissue: liver, heart, muscle, brain tissue, kidney, zebrafish embryo, etc.

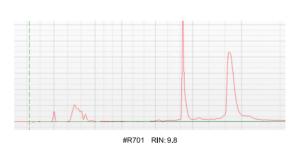
Plant tissue: rice, corn, Arabidopsis, wheat, buckwheat, tobacco, soybeans, potatoes, etc.

Cells: HEK293, Hela, hybridoma, B16-F10 melanoma cells, HepG2, H9c2 cardiomyocytes, MEF, etc.

Microorganisms: E. coli, mold, yeast, etc.

#### Validation Data





Total RNA from HEK293 cells ( $2 \times 10^6$ ) was purified using RNA-easy Isolation Reagent (Vazyme, #R701) and Trizol reagents from supplier Th, Ta, and Ti, respectively. The OD260/280 ratio of the yield RNA was determined using Onedrop, and RNA integrity was analysis using Agilent 2100 Bioanalyzer. The figure shows that the RNA extracted by Vazyme #R701 was excellent, and the purity was the same as that of Th and Ta, which was significantly higher than that of Ti (P = 0.0077).

## Plant RNA and DNA Isolation





FastPure Plant Total RNA Isolation Kit (Polysaccharides & Polyphenolics-rich) (#RC401)

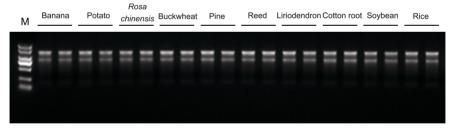
#### **Features**

- \* High purity.
- \* Rapid extraction of total RNA from plant tissues, especially from those rich in polysaccharide & polyphenol.
- \* Low genomic DNA residue.

#### Validated Samples

Pine needles, Eriobotrya japonica leaves, potato tubers, grape fruits, apples, pears, tobacco leaves, mature leaves and roots of wheat, peach fruit, lotus, chrysanthemum rhizome, bananas, Rosa chinensis, buckwheat leaves and seeds, poplar, Catharanthus roseus leaves, liriodendron, reed, rice plant, roots and leaves of cotton, strawberry leaf, Phoebe neurantha leaves, ginkgo (root, leaf, flower and fruits), Arabidopsis seeds, corn seeds, fungal hyphae, etc.

#### Validation Data



Total RNA was extracted using Vazyme #RC401 from 50 mg of banana fruit, potato tubers, rose petals, pine needles, reed leaves, Liriodendron leaves, cotton roots, soybean leaves, rice leaves, or 20 mg of buckwheat seed, respectively. The RNA products were loaded for agarose gel electrophoresis. Vazyme #RC401 showed great compatibility to above plants, especially to those that were rich in polysaccharide & polyphenol, and the RNA extracted using Vazyme #RC401 was with good integrity and high yield.

M: DL2000 Plus DNA Marker (Vazyme, #MD101). The elution volume was 100 µl and the loading amount was 4 µl-10 µl for agarose gel electrophoresis.

## **Rapid Sample Treatment**



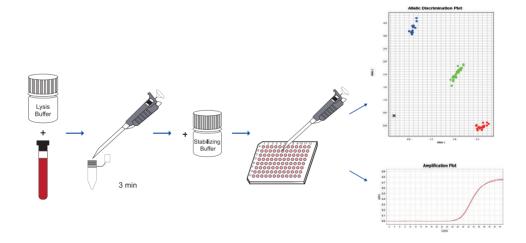


RoomTemp Sample Lysis Kit (#P073)

#### **Features**

- \* Replace cumbersome template extraction protocols with a simple one-step cell lysis procedure.
- \* Lyse samples in just **3 min** at room temperature, qPCR reagent is compatible with the fast program, total operation time is less than 1 hr.
- \* Lyse different anticoagulant blood, FTA card, buccal swab and other samples.
- \* The lysis reagent is consistent with the traditional kit for extracting the genome.

#### Workflow



Workflow of RoomTemp Sample Lysis Kit

## **Endotoxin-Free Plasmid**





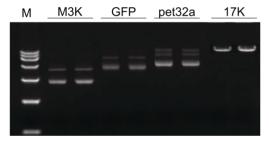
## FastPure EndoFree Plasmid Maxi Kit (#DC202)

#### **Features**

- \* Endotoxin residue < 0.1 EU / µg.
- \* Yield 0.2 mg-1.5 mg of high-copy plasmid DNA after rapid extraction.

#### Validation Data

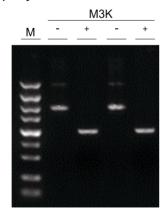
#### 1. High yield.



High extraction efficiency for plasmids with different sizes. Plasmids with different sizes (3.0 kb, 5.0 kb, 5.9 kb, and 17 kb) were extracted using FastPure EndoFree Plasmid Maxi Kit (Vazyme, #DC202), and detected by 1% agarose gel electrophoresis, respectively.

M: DL15000 DNA Marker (Vazyme, #MD103).

#### 2. High purity.



Extracted plasmid with FastPure EndoFree Plasmid Maxi Kit (Vazyme, #DC202) showed high purity and was suitable for restriction enzyme digestion. The extracted M3K plasmid was digested by restriction enzyme and detected by 1% agarose gel electrophoresis.

M: DL5000 DNA Marker (Vazyme, #MD102).

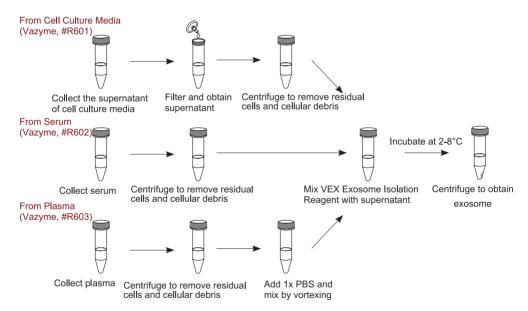
## **Exosome Isolation**



#### **Features**

- \* Easy isolation of exosomes by one-step precipitation, avoiding time-consuming ultra-centrifugation.
- \* Intact exosomes with high yield obtained by low-speed centrifugation.

### Workflow



## **Apoptosis Detection**

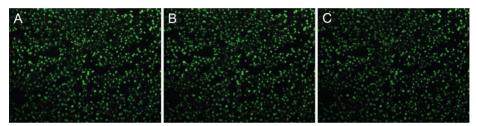


→ TUNEL BrightGreen Apoptosis Detection Kit (#A112)

#### **Features**

- \* Patented Bright factor makes the fluorescence brighter and more resistant to quenching.
- \* Robust activity of recombinant TdT enzyme ensures the high efficiency of fluorescence incorporation.
- \* Applicable for cells and FFPE or frozen tissue sections.

#### Validation Data



Mouse liver tissue sections were treated with DNase I and then stained with Vazyme #A112. Fluorescence signals were read after exposure to ultraviolet light (460 nm) for 1 min (A), 3 min (B), and 6 min (C), respectively,

### Selected Product Citations

Wang Y, et al. Genome-wide screening of NEAT1 regulators reveals cross-regulation between paraspeckles and mitochondria. *Nature Cell Biology*, 2018, 20(10):1145-58.

Hsu P, et al. Ythdc2 is an N(6)-methyladenosine binding protein that regulates mammalian spermatogenesis. Cell Research, 2017, 27(9):1115.

## **Apoptosis Detection**





→ Annexin V-PE/7-AAD Apoptosis Detection Kit (#A213)

#### **Features**

Red Fluorescent Label

Can be used to dual-label cells with green fluorescence.

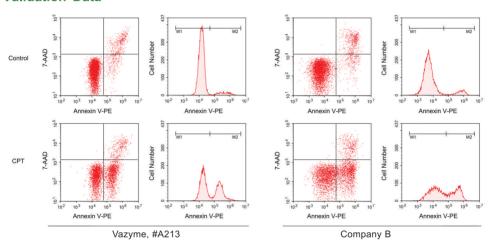
High Specificity

The apoptotic cells can be clearly clustered, and the population apoptosis trend is accurately determined.

Wide Range of Applications

Applicable for suspension cells, adherent cells

#### Validation Data



Apoptosis Detection Using Annexin V-PE/7-AAD (Vazyme, #A213)

Control: Jurkat cells (human T lymphoma cells) untreated with Camptothecin (CPT) were double stained with both Annexin V-PE and 7-AAD, and then detected by Flow Cytometer.

CPT: Jurkat cells untreated with Camptothecin (4 µM) for 4 hr were double stained with both Annexin V-PE and 7-AAD, and then detected by Flow Cytometer.

## **Cell Counting**

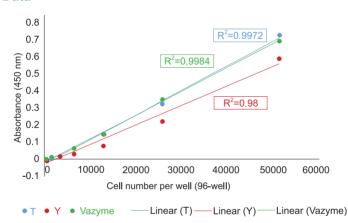


CCK-8 Cell Counting Kit (#A311)

#### **Features**

- \* Ready-to-use solution.
- \* High sensitivity, with excellent linear correlation and repeatability.
- \* Low cytotoxicity.

#### **Validation Data**



HEK293 suspension cells were serially diluted and inoculated to a 96-well plate. The cell density in each group (n = 3) is: 0, 400, 800, 1600, 3200, 6400, 12800, 25600, 51200 cells per well. CCK-8 reagents from Vazyme (#A311, green), Supplier T (blue), and Supplier Y (red) were used for cell counting, respectively. The R<sup>2</sup> value of Vazyme #A311 is > 0.99.

## Selected Product Citations

Zheng Q, et al. Thiopeptide antibiotics exhibit a dual mode of action against intracellular pathogens by affecting both host and microbe. *Chemistry & Biology*, 2015, 22(8):1002-7.

Liu Z, et al. Adiponectin reduces ER stress-induced apoptosis through PPAR $\alpha$  transcriptional regulation of ATF2 in mouse adipose. Cell Death & Disease, 2016, 7(11):e2487.

## **Cell Transfection**

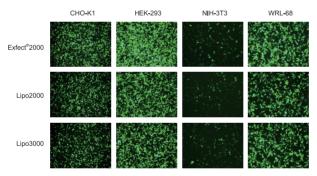


ExFect 2000 Transfection Reagent (#T202)

#### **Features**

- \* High transfection efficiency in a variety of cell lines.
- \* Low cytotoxicity to avoid damaging the normal physiological state of cells.
- \* Add directly to cells in culture medium, in the presence or absence of serum.
- \* Applicable for co-transfection with multiple plasmids.

#### Validation Data



ExFect 2000 exhibits higher transfection efficiency than that of Lipo2000 and Lipo3000. CHOK1, HEK293, NIH3T3 and WRL68 cells were transfected using various reagents in a 24-well plate, respectively. The expression of GFP was analyzed after transfection for 24 h.

## Selected Product Citations

Sun X, et al. Usp7 regulates Hippo pathway through deubiquitinating the transcriptional coactivator Yorkie. *Nature Communications*, 2019, 10(1):411.

Liu X, et al. 1-L-MT, an IDO inhibitor, prevented colitis-associated cancer by inducing CDC20 inhibition-mediated mitotic death of colon cancer cells. *International Journal of Cancer*, 2018, 143(6):1516-29.

## **Luciferase Assay**

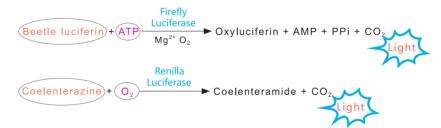


→ Dual Luciferase Reporter Assay Kit (#DL101)

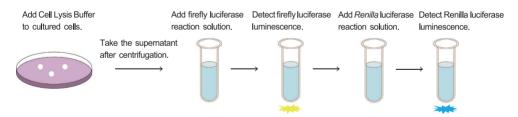
#### **Features**

- \* Robust luminescent signals: applicable for analysis of weak promoters and other genetic regulatory elements.
- \* Detection linear range covers up to 8 orders of magnitude ( $R^2 > 0.99$ ).
- \* Detection sensitivity of 10<sup>-18</sup> mole.

#### Mechanism



#### Workflow



## Selected Product Citations

Liu Z, et al. Circular RNA hsa\_circ\_001783 regulates breast cancer progression via sponging miR-200c-3p. *Cell Death & Disease*, 2019, 10:55

Wu H, et al. Ubiquitination is essential for avibirnavirus replication by supporting VP1 Polymerase activity. **Journal of Virology**, 2019, 93(3): e01899-18.

Wu H, et al. SUMO1 Modification Facilitates Avibirnavirus Replication by Stabilizing Polymerase VP1. *Journal of Virology*, 2019, JVI. 02227-18.

## **Mycoplasma Detection**



→ Myco-Blue Mycoplasma Detector (#D101)

#### **Features**

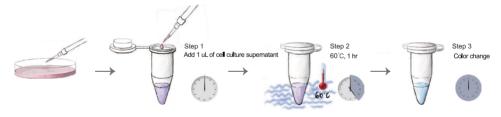
- \* Cell culture supernatant can be used directly for detection.
- \* Results are obtained after incubation at 60°C for 1 hr and can be determined by visual observation.
- \* Accuracy is higher than PCR method, and comparable to gPCR method.
- \* Suitable for detection of all kinds of mycoplasma that are commonly found in cell culture.

#### Validated Cell Lines

Validated cells and media serum include (but are not limited to):

- \* Suspension cells: CHO, NS0, 293F, mouse hybridoma, Sf9, BHK21, etc.;
- \* Adherent cells: Vero, MDCK, SP2/0, 293T, HepG2, HeLa, A549, MB-MDA231, L929, MEF, etc.;
- \* Medium: CD FortiCHO, CDM4, Expi 293 Medium, CD Hybridoma, Grace, DMEM, 1640, F12, etc.;
- \* Serum: fetal calf / calf serum; horse serum; Gibco KSR serum replacement, etc.

#### Workflow



#### Validated Data



Randomly selected 16 cell cultures, and mycoplasma were detected by three methods.



## In Vitro Transcription

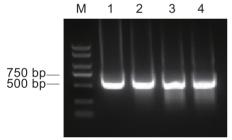


#### **Features**

- \* High yield: yields up to 80 µg of dsRNA in a single reaction.
- \* Magnetic bead purification: recovery efficiency up to 80%.
- \* Able to transcribe both siRNA (21 bp) and dsRNA (long fragment).

#### Validation Data

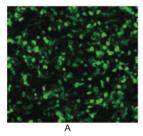
1. Excellent transcription efficiency.

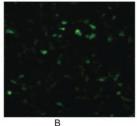


Agarose gel electrophoresis (2%) of 500 bp dsRNA.

- M: DL2000 Plus DNA Maker.
- 1 and 3: products before and after enzymatic hydrolysis of dsRNA, respectively;
- 2 and 4: products before and after enzymatic hydrolysis of dsRNA, respectively.

#### 2. Knock-down of GFP expression by transcribed siRNA.





293T cells were co-transfected for 24 hrs with both GFP plasmid and negative control GFP siRNA (A) or positive GFP siRNA (B).





#### **High-Fidelity PCR**

Zhao Q, Wang M, Xu D, et al. Metabolic coupling of two small-molecule thiols programs the biosynthesis of lincomycin A[J]. Nature. 2015 Feb5;518(7537):115-9. IF: 42.351

Zhang B, Wang K B, Wang W, et al. Enzyme-catalysed [6+4] cycloadditions in the biosynthesis of natural products[J]. *Nature*. 2019 Apr;568(7750):122-126. **IF:** 41.577

Ma Z, Zhu L, Song T, et al. A paralogous decoy protects Phytophthora sojae apoplastic effector PsXEG1 from a host inhibitor[J]. Science. 2017 Feb 17;355(6326):710-714. IF: 34.661

Han X, Wang R, Zhou Y, et al. Mapping the MouseCell Atlas by Microwell-Seq[J]. Cell. 2018 May17;173(5):1307. IF: 31.398

Zhang B, Li J, Yang X, et al. Crystal Structures of Membrane Transporter MmpL3, an Anti-TB Drug Target[J]. Cell. 2019 Jan 24:176(3):636-648.e13. IF: 31.398

Wang YS, Zhang B, Zhu J, et al. Molecular Basis for the Final Oxidative Rearrangement Steps in Chartreusin Biosynthesis[J]. J Am Chem Soc. 2018 Aug 29;140(34):10909-10914. IF: 14.357

Cheng X, Ma X, Ding X, et al. Pacer Mediates the Function of Class III PI3K and HOPS Complexes in Autophagosome Maturation by Engaging Stx17[J]. *Mol Cell*. 2017 Mar 16:65(6):1029-1043.e5. **IF:** 14,248

Wu H, Yin QF, Luo Z, et al. Unusual Processing Generates SPA LncRNAs that Sequester Multiple RNA Binding Proteins[J]. *Mol Cell*. 2016 Nov 3;64(3):534-548. **IF:** 13.958

Tian Z, Sun P, Yan Y, et al. An enzymatic [4+2] cyclization cascade creates the pentacyclic core of pyrroindomycins[J]. *Nat Chem Biol.* 2015 Apr;11(4):259-65. **IF:** 13.217

Zhang M, Zhou C, Wei Y, et al. Human cleaving embryos enable robust homozygotic nucleotide substitutions by base editors[J]. Genome Biol. 2019 May 22;20(1):101. IF: 13.214

Wang M, Zhao Q, Zhang Q, et al. Differences in PLP-Dependent Cysteinyl Processing Lead to Diverse S-Functionalization of Lincosamide Antibiotics[J]. *J Am Chem Soc.* 2016 May 25;138(20):6348-51. **IF:** 13.038

Lv M, Ji X, Zhao J, et al. Characterization of a C3 Deoxygenation Pathway Reveals a Key Branch Point in Aminoglycoside Biosynthesis[J]. *J Am Chem Soc.* 2016 May 25:138(20):6427-35. IF: 13

Sun X, Ding Y, Zhan M, et al. Usp7 regulates Hippo pathway through deubiquitinating the transcriptional coactivator Yorkie[J]. *Nature Communications*. 2019 Jan 24;10(1):411. IF: 12.353

Duan GF, Ye Y, Xu S, et al. Signal peptide represses GluK1 surface and synaptic trafficking through binding to amino-terminal domain[J]. *Nature Communications*. 2018 Nov 19:9(1):4879. **IF**: 12.353

Chen T, Xiang JF, Zhu S, et al. ADAR1 is required for differentiation and neural induction by regulating microRNA processing in a catalytically independent manner[J]. Cell Res. 2015 Apr;25(4):459-76. IF: 12.413

Ding W, Ji W, Wu Y, et al. Biosynthesis of the nosiheptide indole side ring centers on a cryptic carrier protein NosJ[J]. Nature Communications. 2017 Sep 5;8(1):437. IF: 12.124

Yuan H, Zhang J, Cai Y, et al. Gyrl-like proteins catalyze cyclopropanoid hydrolysis to confer Cellular protection[J]. Nature Communications. 2017 Nov 14;8(1):1485. IF: 12.124

Zhang X, Wang TT, Xu QL, et al. Genome Mining and Comparative Biosynthesis of Meroterpenoids from Two Phylogenetically Distinct Fungi[J]. *Angew Chem Int Ed Engl.* 2018 Jul 2;57(27):8184-8188. IF: 12.102

Ji X, Li Y, Xie L, et al. Expanding Radical SAM Chemistry by Using Radical Addition Reactions and SAM Analogues[J]. Angew Chem Int Ed Engl. 2016 Sep 19;55(39):11845-8. IF: 11.71

Ji X, Li Y, Ding W, et al. Substrate-Tuned Catalysis of the Radical S-Adenosyl-L-Methionine Enzyme NosL Involved in Nosiheptide Biosynthesis[J]. Angew Chem Int Ed Engl. 2015 Jul 27;54(31):9021-4. IF: 11.261

Yin Z, Chen C, Yang J, et al. Histone acetyltransferase MoHat1 acetylates autophagy-related proteins MoAtg3 and MoAtg9 to orchestrate functional appressorium formation and pathogenicity in Magnaporthe oryzae[J].

Autophagy. 2019 Jul;15(7):1234-1257. IF: 11.1

Xu D, Shan B, Sun H, et al. USP14 regulates autophagy by suppressing K63 ubiquitination of Beclin1[J]. Genes Dev. 2016 Aug 1;30(15):1718-30. IF: 10.042



#### **Conventional PCR**

Zhang B, Wang K B, Wang W, et al. Enzyme-catalysed [6+4] cycloadditions in the biosynthesis of natural products[J]. *Nature*. 2019 Apr:568(7750):122-126. **IF:** 41,577

Zhang XO, Wang HB, Zhang Y, et al. Complementary sequence-mediated exon circularization[J]. Cell. 2014 Sep 25;159(1):134-147. IF: 33,116

Wang Y S, Zhang B, Zhu J, et al. Molecular Basis for the Final Oxidative Rearrangement Steps in Chartreusin Biosynthesis[J]. J Am Chem Soc. 2018 Aug 29;140(34):10909-10914. IF: 14,357

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