

Dongsheng Biotech Co., Ltd. was established in 2005 in Guangzhou China.

We focus on R&D and produce fundamental molecular biology reagents. In the past 5 years, Dongsheng Biotech has successfully grown up from a start-up to a professional organization. Our products have worldwide sales and establish an excellent reputation.

We were ISO 9001:2008 certificated by BSI. We are also authorized import and export manufacture in China. With professional R&D group and strict control of the production process, we now have customers all over the world.

Our mission is to provide professional and high quality products for life science research. To make your research more professional and more accurate, we are always making our best efforts.



# **Make Your Research**

# **More Professional**

# **More Accurate**





Fax: +86-20-87791381

E-mail: dongshengbio@hotmail.com

Dongsheng Biotech provides wide range of molecular biology products for life science research.

### **Product Line:**

- 1. DNA Electrophoresis Products
- 2. PCR Products
- 3. DNA Purification Products
- 4, RNA Purification Products
- 5. Other Associative Reagents

.

### **Key Products:**

- Taq DNA Polymerases
- PCR Mix
- DNA Ladders
- PCR & Gel Extraction Kits
- TRIzol

 $\label{lem:all_product} Additionally, \textbf{all product} \ provides \ custom \ synthesis.$ 







### **Dongsheng Biotech Product Application area**

### Serve three major customer groups:

- (1) Laboratory reagent kits producers, a smaller group of customers (<50 major customers worldwide) who purchase substantial quantities of nucleotide monomers as raw material for specialized laboratory kits for use in research laboratories.
- (2) Diagnostic kits producers who use our nucleotide monomers as diagnostic agents in hospitals, blood collection agencies and pathology laboratories primarily for amplification.
- (3) Research laboratories in universities, industries, governments and hospitals & medical schools.



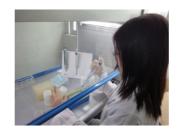


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### 1. DNA Electrophoresis Products

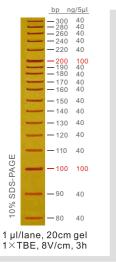
PCR and plasmid digestion by restriction enzymes are used for the production of DNA fragments for our ladders/markers. These products are stable during prolonged incubations at room temperature and multiple freeze-thaw cycles.

### Distinguishing features

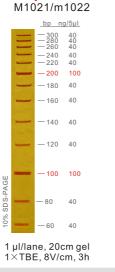
- Bright, sharp bands for highly accurate sizing
- Precise DNA concentration, quantitative
- · High resolution, low-background
- Stable and uniform with different batches



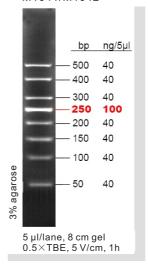
#### 10bp ladder M1011/M1012 bp ng/5µl -300 40



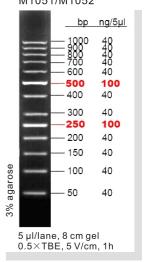
20bp ladder



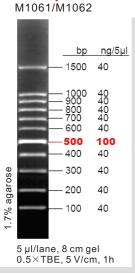
50bp ladder M1041/M1042



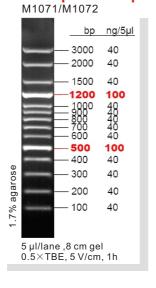
50bp ladder plus M1051/M1052



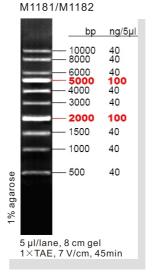
### 100bp ladder



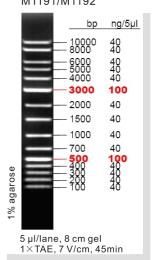
100bp ladder plus



1kb ladder



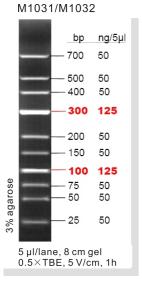
1kb ladder plus M1191/M1192



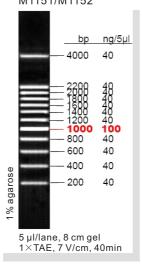
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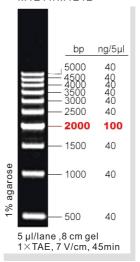
### Low ladder



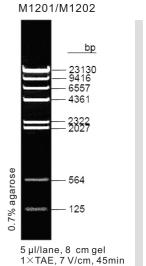
## **200bp ladder** M1151/M1152



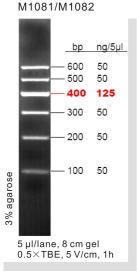
# **500bp ladder** M1211/M1212



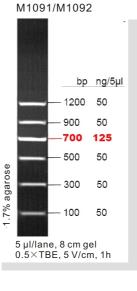
### Lambda DNA/Hind III



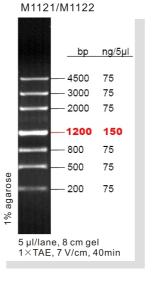
Marker 1



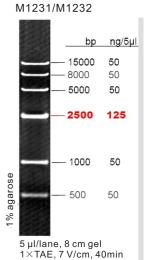
Marker 2



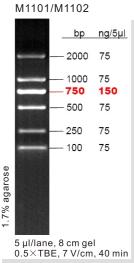
Marker 3



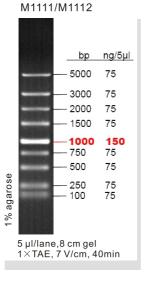
Marker 4



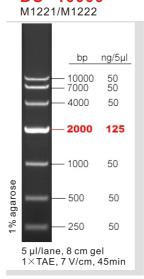
**DS**<sup>TM</sup>**2000** 



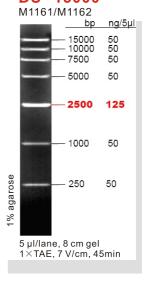
*DS*<sup>™</sup>5000



**DS**<sup>™</sup>10000



**DS**<sup>™</sup>15000



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# **Product List**

Product name	Cat #	Size	Range
10bp ladder	M1011	50µg	80-300bp
	M1012	5x50µg	·
20bp ladder	M1021	50µg	60-300bp
	M1022	5x50µg	
low ladder	M1031	50µg	25-700bp
	M1032	5x50µg	
50bp ladder	M1041	50µg	50-500bp
	M1042	5x50µg	
50bp ladder plus	M1051	50µg	50-1,000bp
	M1052	5x50µg	
100bp ladder	M1061	50µg	100-1,500bp
	M1062	5x50µg	
100 bp ladder plus	M1071	50µg	100-3,000bp
	M1072	5x50µg	
1kb ladder	M1181	50µg	500-10,000bp
	M1182	5x50µg	
1kb ladder plus	M1191	50µg	100-10,000bp
	M1192	5x50µg	·
200 bp ladder	M1151	50µg	200-4,000bp
	M1152	5x50µg	
500bp ladder	M1251	50µg	500-5,000bp
	M1252	5x50µg	·
$\lambda$ DNA/ $Hind$ $III$	M1201	50µg	125-23,130bp
	M1202	5x50µg	
Marker 1	M1081	50µg	100-600bp
	M1082	5x50µg	·
Marker 2	M1091	50µg	100-1,200bp
	M1092	5x50µg	
Marker 3	M1121	50µg	200-4,500bp
	M1122	5x50µg	·
Marker 4	M1231	50µg	500-15,000bp
	M1232	5x50µg	
<i>DS</i> <sup>™</sup> 2000	M1101	50µg	100-2,000bp
	M1102	5x50µg	
<i>DS</i> <sup>™</sup> 5000	M1111	50µg	100-5,000bp
	M1112	5x50µg	,
<i>DS</i> ™10000	M1221	50µg	250-10,000bp
	M1222	5x50µg	, 1
<i>DS</i> <sup>™</sup> 15000	M1161	50µg	250-15,000bp
	M1162	5x50µg	Γ

Website: www.dongshengbio.com

Product name	Cat #	Size
20x Swift DNA Electrophoresis Buffe	M9011	500ml
50xTAE Buffer	M9021	500ml
10xTBE	M9031	500ml
Gel Loading Dye(6x)	M9041	5x1ml
Gel Loading Dye, Blue(6x)	M9051	5x1ml
Gel Loading Dye, Three-colour (6x)	M9061	5x1ml
Gel Loading Dye, Orange(6x)	M9071	5x1ml
Gel Loading Dye, SDS (6x)	M9081	5x1ml

 $DS^{TM}$  is a collection of 30 highly-purified individual DNA fragments ranging from 10-15,000 bp.

DNA Francost Sina	Cat #	Amount
DNA Fragnent Size		Amount
10 bp	M8011	10µg
15 bp	M8021	10µg
20 bp	M8031	10µg
25 bp	M8041	10µg
35 bp	M8051	10µg
50 bp	M8061	10µg
75 bp	M8071	10µg
100 bp	M8081	10µg
150 bp	M8091	10µg
200 bp	M8101	10µg
250 bp	M8111	10µg
300 bp	M8121	10µg
400 bp	M8131	10µg
500 bp	M8141	10µg
600 bp	M8151	10µg
700 bp	M8161	10µg
800 bp	M8171	10µg
900 bp	M8181	10µg
1000 bp	M8191	10µg
1200 bp	M8201	10µg
1500 bp	M8211	10µg
2000 bp	M8221	10µg
2500 bp	M8231	10µg
3000 bp	M8241	10µg
4000 bp	M8251	10µg
5000 bp	M8261	10µg
6000 bp	M8271	10µg
8000 bp	M8281	10µg
10000 bp	M8291	10µg
15000 bp	M8301	10µg

We offer exceptionally pure DNA fragments that are free of any truncated or degraded molecules.

### Concentration

 $0.5 \,\mu g/\mu l$ 

### Storage Buffer (TE Buffer)

10 mM Tris-HCI (pH 7.6) with 1 mM EDTA

### **Quality Control**

Tested on an Agilent 2100 bioanalyzer and by appropriate gel electrophoresis. DNA concentration is determined by spectrophotometer. The absence of nucleases confirmed by appropriate tests..

### Storage

Store at -20°C

### **Custom DNA Ladders Service**

Bulk orders can be customized to fit your needs: Individual DNA fragments or mixtures.

Custom ladders of any combination and concentration.

Reference bands in a custom-made DNA ladder. Flexible formulations.

Ready-to-use versions premixed with any loading dye.

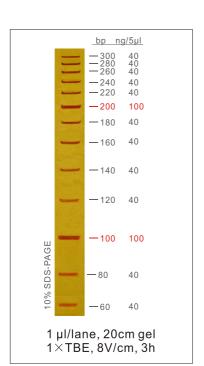
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From microgram to gram quantities.



# 20bp ladder

Cat.#SizeM102150μgM10225x50μg



Concentration: 128ng/µl

Recommended Loading: 2-5µl/lane

### Concentration

Typical Bands 100ng/5µl
Other Bands 40ng/5µl

### Recommended Electrophoresis Condition

5µI/lane, 8cm 5% Agarose Gel, 0.5xTBE, 5V/cm, 1h

### Contents(bp)

60, 80, <u>100</u>, 120, 140, 160, 180, <u>200</u>, 220, 240, 260, 280, 300

### Storage

1 year at room temperature, for long-time preservation, please store at -20℃.

### Description

20bp ladder is ideal for determining the size of double-stranded DNA from 60 to 300bp base pairs. The ladder consists of 13 linear double-stranded fragments. The 100bp and 200bp fragment are present at increased intensity to allow easy identification. All fragments are precisely quantified and mixed during the production. For 5 ul loading, all fragments except 100bp and 200bp are 40ng. The 100bp and 200bp fragment are 100ng. This ladder is pre-mixed with loading dye and is ready to use.

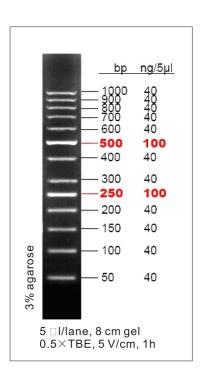
#### Recommendation

Do not heat before loading

For quantification, adjust the concentration of the sample to equalize. Visualize DNA by staining with ethidium bromide or with SYBY Green I.

# 50bp ladder plus

Cat.#SizeM105150μgM10525x50μg



Concentration: 128ng/µl

Recommended Loading: 2-5µl/lane

### Concentration

Typical Bands 100ng/5µl
Other Bands 40ng/5µl

### Recommended Electrophoresis Condition

5µl/lane, 8cm 3% Agarose Gel, 0.5xTBE, 5V/cm, 1h

### Contents(bp)

50, 100, 150, 200, **250**, 300, 400, **500**, 600, 700,800, 900, 1000

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### Storage

1 year at room temperature, for long-time preservation, please store at -20℃.

### Description

50bp ladder plus is ideal for determining the size of double-stranded DNA from 50 to 1000bp base pairs. The ladder consists of 13 linear double-stranded fragments. The 250bp and 500bp fragment are present at increased intensity to allow easy identification. All fragments are precisely quantified and mixed during the production. For 5 ul loading, all fragments except 250bp and 500bp are 40ng. The 250bp and 500bp fragment are 100ng. This ladder is pre-mixed with loading dye and is ready to use.

### Recommendation

Do not heat before loading

For quantification, adjust the concentration of the sample to equalize.

Visualize DNA by staining with ethidium bromide or with SYBY Green I.



# 100bp ladder plus

Cat.#SizeM107150μgM10725x50μg

Concentration: 136ng/µl

Recommended Loading: 2-5µl/lane

### Concentration

Typical Bands 100ng/5µl
Other Bands 40ng/5µl

### Recommended Electrophoresis Condition

5µl/lane, 8cm 1.7% Agarose Gel, 0.5xTBE, 5V/cm, 1h

### Contents(bp)

100, 200, 300, 400, <u>500</u>, 600, 700,800, 900, 1000, <u>1200</u>, 1500,2000,3000

### Storage

1 year at room temperature, for long-time preservation, please store at -20℃.

### Description

100bp ladder plus is ideal for determining the size of double-stranded DNA from 100 to 3000bp base pairs. The ladder consists of 14 linear double-stranded fragments. The 500bp and 1200bp fragment are present at increased intensity to allow easy identification. All fragments are precisely quantified and mixed during the production. For 5 ul loading, all fragments except 500bp and 1200bp are 40ng. The 500bp and 1200bp fragment are 100ng. This ladder is pre-mixed with loading dye and is ready to use.

#### Recommendation

Do not heat before loading

For quantification, adjust the concentration of the sample to equalize.

Visualize DNAby staining with ethidium bromide or with SYBY Green I.

# 1kb ladder plus

Cat.#SizeM119150μgM11925x50μg

ng/5µl bp 40 40 40 100 2000 40 1500 40 1000 40 40 700 agarose % 5 µl/lane, 8 cm gel 1×TAE, 7 V/cm, 45min

Concentration: 144ng/µl

Recommended Loading: 2-5µl/lane

### Concentration

Typical Bands 100ng/5µl
Other Bands 40ng/5µl

### Recommended Electrophoresis Condition

5µl/lane, 8cm 1% Agarose Gel, 1xTAE, 7V/cm, 45min

### Contents(bp)

100, 200, 300, 400, <u>500</u>, 700, 1000, 1500, 2000, <u>3000</u>, 4000, 5000, 6000, 8000, 10000

Fax: +86-20-87791381

### Storage

1 year at room temperature, for long-time preservation, please store at -20°C.

### Description

1kb ladder plus is ideal for determining the size of double-stranded DNA from 100 to10,000bp base pairs. The ladder consists of 15 linear double-stranded fragments. The <u>500bp</u> and <u>3,000bp</u> fragment are present at increased intensity to allow easy identification. All fragments are precisely quantified and mixed during the production. For 5 ul loading, all fragments except 500bp and 3,000bp are 40ng. The 500bp and 3,000bp fragment are 100ng. This ladder is pre-mixed with loading dye and is ready to use.

#### Recommendation

Do not heat before loading

For quantification, adjust the concentration of the sample to equalize.

Visualize DNA by staining with ethidium bromide or with SYBY Green I.



## 2. PCR Product

PCR (polymerase chain reaction) is a technique that is central to molecular biology research. It is used in applications such as cloning, gene expression analysis, genotyping, sequencing, and mutant genes.

Dongsheng Biotech offers different series of products to help you achieve PCR success. For enzymes, our Taq Polymerase, HS<sup>™</sup> Taq DNA Polymerase,  $FS^{TM}$  Tag DNA Polymerase, Pfu DNA Polymerase and Fusion Pfu DNA Polymerase provide high fidelity, efficient, sensitivity PCR performance. We also have Long Taq DNA Polymerase for long DNA amplification.

### We provide three series of PCR products for different uses:

Tag Polymerase Series **Taq Mix Series PCR Kit Series** 











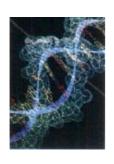
# **Properties of PCR enzymes**

Enzyme	Feature	Amplification size	Fidelity	Sensivity	PCR Product e	ends Application
Taq	• 5'→3' Activity	++	+	++	3'-dA	General PCR/RT-PCR DNA Mark/DNA Sequence TA Clone
<i>HS</i> ™ Taq	• 5'→3'Activity	+++	+	++++	3'-dA	General PCR/RT-PCR DNA Mark/DNA Sequence TA Clone
<i>FS</i> ™ Taq	• 5'→3' Activity • Fast-3 times of	+++ <sup>†</sup> Taq	+	++++	3'-dA	General PCR/RT-PCR DNA Mark/DNA Sequence TA Clone
Pfu	<ul><li>5'→3' Activity</li><li>5'-3' Exonclease</li></ul>	++ e activity	++++	+++	Blunt end	High fidelity PCR Blunt end vector cloning RT-PCR/cDNA clone and express/SNP
Taq Plus	<ul><li>5'→3' Activity</li><li>5'-3' Exoncleas</li><li>High fidelity-4 ti</li></ul>	-	+++	+++	3'-dA/ Blunt end	GC rich template Complicated template High repeat sequences
LongTaq	<ul><li>5'→3' Activity</li><li>5'-3' Exoncleas</li><li>High fidelity-4</li></ul>	•	+++	++	3'-dA/ Blunt end	GC rich template Long template Sequencing
Fusion pfu	<ul><li>5'→3' Activity</li><li>5'-3' Exoncleas</li></ul>	++++ e activity	++++	++++	Blunt end	High fidelity PCR Long fragment PCR













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# **Taq Polymerase Series**

# **Product List**

Product Name	Cat #	Size	Description
Taq DNA Polymerase	P1011 P1012* P1013 P1014* P1015	500U 500U 1,000U 1,000U 18,000U	Robust Taq DNA Polymerase for routine use incl.10x PCR Buffer, 6xloading buffer, Mg2+ plus or free. Supply with 5U/µl and 2.5U/µl packages.
HS <sup>™</sup> Taq DNA Polymerase	P1081 P1082 P1083 P1084	250U 500U 1,000U 18,000U	High specificity,incl. 10x HS <sup>™</sup> PCR buffer, 6x loading buffer, Mg2+ plus or free. Supply with 5U/µl and 2.5U/µl two packages.
Pfu DNA Polymerase	P1021 P1022* P1023 P1024*	250U 250U 1,000U 1,000U	High fidelity, incl.10x pfu buffer, 6xloading buffer, Mg2+ plus or free. Supply with 5U/µl and 2.5U/µl two packages.
Taq Plus DNA Polymerase	P1031 P1032* P1033 P1034*	250U 250U 1,000U 1,000U	Mix with Taq Polymerase and Pfu Polymerase, long amplification and high fidelity. Incl. 10xTaq Plus buffer, Mg2+ plus or free. Supply with 5U µI and 2.5U/µI two packages.
Long Taq DNA Polymerase	P1061 P1062* P1063 P1064*	250U 250U 1,000U 1,000U	For long PCR, incl. 10xLong Taq buffer, 6xloading buffer, Mg2+ plus or free. Supply with 5U/µl and 2.5U/µl two packages.
FS <sup>™</sup> Taq DNA Polymerase	P1071 P1072* P1073 P1074*	250U 250U 1,000U 1,000U	Fast, incl. 10xFS <sup>™</sup> PCR buffer, 6xloading buffer, Mg2+ plus orfree. Supply with 5U/µl and 2.5U/µl two packages.

# **Taq Mix Series**

Product Name	Cat #	Size	Description
Taq Mix	P2011 P2011	1ml 5x1ml	Pre mix with Taq DNA Polymerase, dNTPs, Mg2+ and Reaction buffer at optional conc- entrations
HS <sup>™</sup> Mix	P2081 P2082	1ml 5x1ml	Pre mix with HS <sup>™</sup> Taq DNA Polymerase, dNTPs, Mg2+ and Reaction buffer at optional concentrations
Pfu Mix	P2021 P2022	1ml 5x1ml	. Pre mix with Pfu DNA Polymerase, dNTPs, Mg2+ and Reaction buffer at optional conc- entrations.

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FS™ Taq Mix	P3011 P3012	1ml 5x1m	Pre mix with $FS^{TM}$ Taq DNA Polymerase, dNTPs, Mg2+ and Reaction buffer at optional concentrations.
Long Taq Mix	P2041 P2042	1ml 5x1ml	Pre mix with Long Taq DNA Polymerase, dNTPs, Mg2+ and Reaction buffer at optional concentrations.
FS <sup>™</sup> Taq Mix Direct for Blood	P3021 P3022	1ml 5x1ml	Pre mix with $FS^{TM}$ Taq DNA Polymerase, dNTPs, Mg2+ and Reaction buffer at optional concentrations.
FS <sup>™</sup> Taq Mix Direct for Tissue	P3031 P303	1ml 5x1ml	Contains $FS^{TM}$ Taq Mix T(2x) and sample preparation solution (Extraction solution and Neutralization solution
SYBR Green qPCR Mix	P2091 P2092	1ml 5x1ml	Mix reaction (2x) for Real Time PCR

### **PCR Kit and Relative Solution**

Product Name	Cat #	Size	Description
Taq Mix Kit	P2052	5x1ml	Contains 250U Taq DNA Polymerase and 5x1ml CR Reaction Mix (2x)
HS™ Kit	P2101	5x1ml	Contains 250U HS Taq DNA Polymerase and 5x1ml HS Reaction Mix (2x)
FS™ Kit	P3042	5x1ml	Contains 250U FS Taq DNA Polymerase and 5x1ml FS Reaction Mix (2x)
Long Taq Kit	P2072	5x1ml	Contains 250U Long Taq DNA Polymerase and 5x1ml Long Reaction Mix (2x)
Water (nuclease-free,	P9021	5x1ml	
molecular biology grade)	P9022	100ml	
	P9023	500ml	
MgCl <sub>2</sub> (25mM)	P9031	4x1.25ml	
PCR Enhancer	P9041	500ul	
10xPCR buffer	P9051	4x1.25ml	With 15mM Mg2+
	P9052	4x1.25ml	Without 15mM Mg2+
	P9053	6x1ml	With Mg2+ concentration set
PCR Sample Preparation	P9061	50 preps	To prepare blood or tissue sample for
Solution	P9062	250 preps	fast PCR
dNTP Mix (2.5mM)	P9011	1ml	
dNTP Mix (10mM)	P9012	0.5ml	
	P9013	1ml	
dATP(100mM)	P9071	0.5ml	
dTTP (100mM)	P9081	0.5ml	
dCTP (100mM)	P9091	0.5ml	
dGTP (100mM)	P9101	0.5ml	

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# Taq DNA Polymerase(recombinant)

# P1011

Taq DNA Polymerase (5U/µI)	100µl
10xPCR Buffer (Mg2+ plus)	1.4ml
6xloading buffer	1ml
P1012	
Taq DNA Polymerase (5U/µI)	100µl
10xPCR Buffer (Mg2+ plus)	1.4ml
dNTP Mix (2.5mM each)	1ml
6xloading buffer	1ml
P1013	
Taq DNA Polymerase (5U/µI)	200µl
10xPCR Buffer (Mg2+ plus)	2x1.4ml
6xloading buffer	1ml
P1014	
Taq DNA Polymerase (5U/µI)	200µl
10xPCR Buffer (Mg2+ plus)	2x1.4ml
dNTP Mix (2.5mM each)	2x1ml
6xloading buffer	1ml
P1015	
Taq DNA Polymerase (5U/µI)	36x100µl
10xPCR Buffer (Mg2+plus)	36x1.4ml

### Store at -20°C

#### Feature

Thermostable-half life is more than 40 min at 95°C

Generates PCR products with 3'-dA overhangs

Incorporates modified nucleotides (e.g. Tiotin-, digoxigenin-, fluorescently-labeled nucleotides



Lane M:DS<sup>™</sup> 5000 Lane 1 and 2:0.5 kb Lane 3 and 4:1 kb Lane 5 and 6:2 kb

Figure. Optimal results with different fragments using Taq DNA Polymerase

### Description

Tag DNA Polymerase is a themostable recombinant DNA Polymerase derived from Thermus aquaticus. Its molecular weight is 94 kDa. Taq DNA Polymerase can amplify DNA target up to 5kb (simple template). The elongation velocity is 0.9~1.2kb/min (70~75°C). It has 5' → 3' exonuclease activity but lacks 3' → 5' exonuclease activity that results in a 3'-dA overhangs PCR product.

### Storage Buffer

20mM Tris-HCI(pH 8.0), 100mM KCI, 3mM Mg<sup>2+</sup>, 1mM DTT, 0.1%NP-40, 0.1%Tween20, 0.2mg/ml BSA, 50% (v/v) glycerol.

#### 10xPCR Buffer

120mM Tris-HCI (pH 8.8), 500mM KCI, 1% Trition-X-100, 100mM Lycine, 25mM Mg<sup>2+</sup>

# Taq Mix (2x)

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### Composition of Taq Mix(2x)

Taq Mix(2x) is a mixture of Taq DNA Polymerase, 2x Taq buffer, dNTPs, 3mMMg<sup>2+</sup> and bromophenol blue. Taq mix buffer is a proprietary formulation optimized for robust performance in PCR.

### Description

Taq Mix (2x) is a premixed, ready-to-use solution containing Taq DNA Polymerase, dNTPs, Mg²+ and Reaction Buffer at optional concentrations for efficient amplification of DNA templates by PCR. To prepare the final PCR, only primers and template DNA are added. This pre-mixed formulation saves time and reduces contamination due to the fewer pipetting steps required for PCR set up. The mix retains all features of Taq DNAPolymerase.

Store at -20°C

#### Feature

Convenient-Taq DNA Polymerase in a ready-to-use mix
Fast- save time due to reduced number of pipetting steps
High yield of PCR products with minimal optimization
Reproducible- lower contamination and pipetting error risk





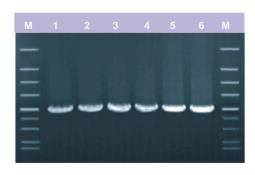


Figure. Stability of Taq Mix (2X).

Lane M: $DS^{TM}$  5000 Lane 1 and 2: PCR product generated with Taq Mix (at room temperature after 1 week) Lane 3 and 4: PCR product generated with Taq Mix (after 15 freeze-thaw cycles) Lane 5 and 6: PCR product generated with Tag Mix(stored at -20°C)

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# **HS<sup>™</sup> Taq DNA Polymerase**

P1011	
<i>HS</i> <sup>™</sup> Taq DNA Polymerase (5U/μl)	50µl
10x <i>HS</i> ™PCR Buffer(Mg²⁺ Plus)	1.4ml
6xloading buffer	1m
P1012	
HS <sup>™</sup> Taq DNA Polymerase (5U/μl)	100µl
10x <i>HS</i> ™PCR Buffer(Mg²⁺ Plus)	1.4ml
6xloading buffer	1ml
P1013	
<i>HS</i> <sup>™</sup> Taq DNA Polymerase (5U/μl)	200µl
10x <i>HS</i> ™PCR Buffer(Mg²⁺ Plus)	2x1.4ml
6xloading buffer	1ml
P1014	
Taq DNA Polymerase (5U/μΙ)	36x100µl
10x <i>HS</i> ™PCR Buffer(Mg²⁺ Plus)	36x1.4ml

### Store at -20°C

#### 10xPCR buffer

120mM Tris-HCI (PH 8.8), 500mM KcI, 1% Trition-X-100, 100mM Lycine, 25mM Mg<sup>2+</sup>

### Feature

Description

 $HS^{\text{TM}}$  Taq DNA Polymerase(High Specificity Taq DNA Polymerase) is a thermostable recombinant DNA Polymerase derived from Thermus aquaticus. It contants the different reaction buffer that is designed to provide highly spicific amplification of nucleic acid templates by PCR. It has 5' → 3' exonuclease activity but lacks of 3' → 5' exonuclease activity. The elongation velocity is 0.9~1.2kb/min (70~75°C). It is heat-stable and will synthesize DNA at elevated temperatures from single-stranded templates in the presence of a primer.

### Storage Buffer

20mM TrisCl(PH 8.0), 100mM KCl, 3mM Mg<sup>2+</sup>, 1mM DTT, 0.1%NP-40, 0.1%Tween20, 0.2mg/ml BSA, 50% (v/v) glycerol.

High yield amplification of complex templates

High PCR specificity-reduced effects of mispriming and primer-dimer formation

Enhanced PCR sensitivity

Convenient room temperature PCR set-up
Generates PCR products with 3-dA overhangs



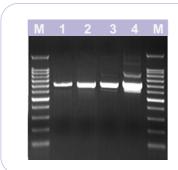


Figure . Comparison of PCR sensitivity and yield with different *HS*™ Taq DNA Polymerase concentrations.

# HS<sup>™</sup> Mix (2X)

P2011		
2xHS™ Mix	1ml	
Nuclease-free Water	1ml	
P2012		
<b>P2012</b> 2x <i>HS</i> <sup>™</sup> Mix	5x1ml	

### Composition of $HS^{TM}$ Mix (2x)

 $HS^{TM}$ Mix(2x) is a mixture of  $HS^{TM}$ Taq DNA Polymerase,  $2x HS^{TM}$  buffer, dNTPs,  $3mMMg^{2+}$  and bromophenol blue.  $HS^{TM}$  Taq mix buffer is a proprietary formulation optimized for robust performance in PCR.

### Description

 $HS^{\text{TM}}$  Mix (2x) is a premixed, ready-to-use solution containing  $HS^{\text{TM}}$  Taq DNA Polymerase, dNTPs,  $Mg^{2^+}$  and Reaction Buffer at optional concentrations for efficient amplification of DNA templates by PCR. To prepare the final PCR, only primers and template DNA are added.  $HS^{\text{TM}}$  Mix provides increased sensitivity, specificity, and yield, while allowing assembly of reactions at room temperature. This mix contributes to highly reproducible PCR by reducing the risk of pipetting errors, miscalculation and contamination. And also save time.

### Feature

Convenient-HS<sup>™</sup> Taq DNA Polymerase in a ready-to-use mix

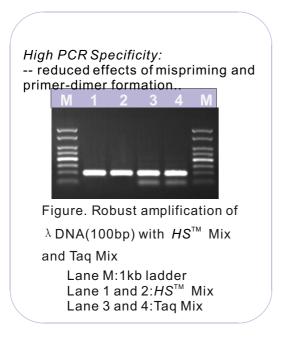
Fast- save time due to reduced number of pipetting steps

High yield of PCR products with minimal optimization

Reproducible- lower contamination and pipetting error risk

Higher sensitivity compared to conventional Taq DNA Polymerase

### Store at -20℃



### Long PCR products:

- -- up to 8 kb with viral DNA as template;
- -- up to 5 kb with genomic DNA as template.

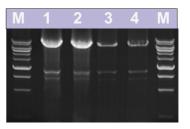


Figure. Robust amplification of human genomic DNA(8kb) with  $HS^{TM}$  Mix and Taq Mix

Lane M:1kb ladder Lane 1 and 2:*HS*™ Mix Lane 3 and 4:Taq Mix

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# 3. DNA Purification Kits

Dongsheng Biotech offers a broad range of kits for the isolation and purification of genomic and plasmid DNA from a wide variety of samples, including tissue, cells, blood, serum, bacterial, yeast, and plants samples.

We provide three series for different materials' purification:

**Plasmid DNA Extraction Kits DNA Product Purification Kits Genimic DNA Purification Kits** 













### **Guides of DNA Purification Kits**

Application	Material	Experimental requirement	Product Choose
		Sample size:1-5ml	High Purity Plasmid Miniprep Kit
		Sample size:1-5ml Safe, Healthy	Eco-Friendly Plasmid Miniprep Kit
		Sample size:100-200ml	Plasmid DNA Maxiprep Kit (High
Plasmid DNA	Bacteria	Simple, Convenient	Purifity, Silica-membrant Spin Column)
Extraction		Sample size:100-200ml	Plasmid DNAMaxiprep Kit (High Purifity, Silica-precipitation)
		Sample size:100-200ml Endotoxin-Free	Plasmid DNA Maxiprep Kit (Endotoxin-Free)
	Yeast	Sample size:1-5ml	Yeast Plasmid DNA Miniprep Kit
	Standard		Gel Extraction Kit
	Agarose Gel	Safe, Healthy	Eco-Friendly Gel Extraction Kit
DNA Product	PCR Product/	_	PCR and DNA Fragment Purification Kit
Purification	DNA Fragment Solution	Safe, Healthy	Eco-Friendly PCR and DNA Fragment Purification Kit
	Insect/ Animal Nematoda	Sample size: less than 10mg tissue	Quick Genomic DNA Extraction Kit
		Sample size: 0.I-1ml flesh blood/frozen blood	Blood Genomic DNA Extraction Kit
Genomic DNA Purification	Blood	Sample size: less than 200ul whole blood	Quick Blood Genomic DNA Extraction Kit
	Tissue/ Cell	Sample size: less than 10mg tissue/ 10 <sup>6</sup> -10 <sup>7</sup> culture cell	Quick Tissue/Culture Cells Genomic DNA Extraction Kit
	Bacteria	Sample size: 0.5-2ml/ less than 10 <sup>6</sup> -10 bacteria	Quick Bacteria Genomic DNA Extraction Kit
	Yeast	Sample size: less than 1-5x10 <sup>7</sup> Yeast	Quick Yeast Genomic DNA Extraction Kit

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# **Product List**

### **Plasmid Extraction Series**

Product Name	Cat#	Size
High Purity Plasmid Miniprep kit	N1011	50 preps
	N1012	100 preps
Eco-Friendly Plasmid Miniprep Kit	N1021	50 preps
	N1022	100 preps
Plasmid DNA Maxiprep Kit (High Purity, Silica-mimbrane	N1031	10 preps
Spin Column)		
Plasmid DNA Maxiprep Kit (High Purity, Silica preciptation)	N1041	10 preps
Plasmid DNA Maxiprep Kit (Endotoxin-free)	N1051	10 preps
Yeast Plasmid DNA Miniprep Kit	N1061	50 preps
	N1062	100 preps

### **PCP Product Purification Series**

Product Name	Cat #	Size
Gel Extraction Kit	N1071	50 preps
	N1072	100 preps
Eco-Feiendly Gel Extraction Kit	N1081	50 preps
	N1082	100 preps
PCR and DNA Fragment Purification Kit	N1091	50 preps
	N1092	100 preps

### **Genomic DNA Purification Serious**

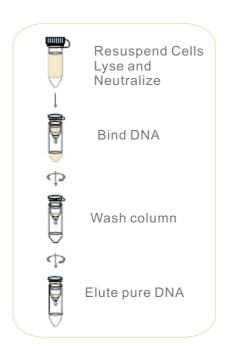
Product Name	Cat#	Size
Quick Genomic DNA Extraction Kit	N1111	50 preps
	N1112	100 preps
Blood Genomic DNA Extraction Kit	N1121	50 preps
	N1122	100 preps
Quick Blood Genomic DNA Extraction Kit	N1131	50 preps
	N1132	100 preps
Quick Tissue/Culture Cells Genomic DNA Extraction Kit	N1141	50 preps
	N1142	100 preps
Quick Bacteria Genomic DNA Extraction Kit	N1151	50 preps
	N1152	100 preps
Quick Yeast Genomic DNA Extraction Kit	N1161	50 preps
	N1162	100 preps
Marine Animal Tissue Genomic DNA Extraction Kit	N1171	50 preps
	N1172	100 preps

Website: www.dongshengbio.com

### **High Purity Plasmid Miniprep Kit**

Content	N1011	N1012
Size	50preps	100preps
RNase A	150ul	300ul
Resuspension Solution	15ml	30ml
Lysis Solution	15ml	30ml
Neutralization Solution	20ml	40ml
Wash Buffer PB	30ml	50ml
Wash Buffer W	30ml	2x30ml
Elution Buffer	5ml	10ml
(10mM Tris-HCI, pH 8.5)		
Spin Columns	50each	100each

For the initial use of the kit, dilute the Wash Buffer W with ethanol 1:1.5 (v/v).



### Description

High Purity Plasmid Miniprep Kit is designed for rapid and cost-effective small-scale preparation of high quality plasmid DNA from recombinant *E. Coli* cultures. The kit utilizes an exclusive silicabased membrane technology in the form of a convenient spin column. Each spin column can recover up to 40ug of plasmid DNA. The kit can be used for any size plasmid and cosmids purification efficiently.

### Feature

Fast-procedure takes only 30 min

Efficient-can get 10-40ug high copy plasmid DNA in one prep

High purity-purified DNA is ready for down-

### Application

stream applications

Fast isolation of high purity plamid DNA are ready for direct use in all applications, including: Restriction digestion
DNA Sequencing
In vitro transcription

### Storage

Store RNase at  $-20^{\circ}$ C. Other solutuions can be stored for up to 1 year at RT (15-25 $^{\circ}$ C)

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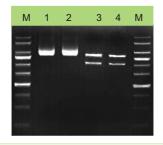


Figure .A5 kb PCR fragment was cloned into pCMV vector. Lane M: 1kb ladder

Lane 1 and 2: Plasmid DNA from overnight bacterial cultures of recombinant clones was purified using the Plasmid Miniprep Kit (N1011).

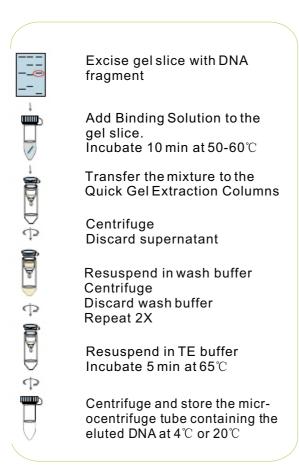
Lane 3 and 4: analyzed by double digestion with EcoRI and HindIII in 30 min at 37°C.



### **Gel Extraction Kit**

Content	N1711	N1712
Size	50preps	100preps
Binding Buffer BD	20ml	40ml
Wash Buffer PE	15ml	2x15ml
Elution Buffer	2.5ml	5ml
(10mM Tris-HCI,pH8.5)		
Spin Columns	50each	100each

For the initial use of the kit, dilute the Wash Buffer PE with ethanol 1:4 (v/v)



### Description

Gel Extraction Kit is deesigned to extract and purify DNA fragments of 50bp to 40kb from s tandard or low-melt agarose gels in either Tris acetate (TAE) or Tris borate (TBE). This membrane-based system, which can bind up to 40ug DNA, allows recovery of isolated DNA fragments in as little as 20 minutes.

#### Feature

**Fast**-procedure takes only 15 min.

**Efficient**-gather up to 85% recoveries in the range of 50bp-40kb.

Safe-not involve phenol extraction High Purity-purified DNA is ready for downstream applications.

### Application

Fast and efficient extraction of high purity DNA fragments are ready for direct use in all applications, including:

Restriction digestion

**PCR** 

DNA sequencing
In vitro transcription

### Storage

All solutions can be stored for up to 1 year RT (15-25 $^{\circ}$ C).

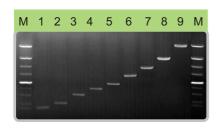


Figure . Recovery of  $DS^{\mathsf{TM}}5000$  individual 100-5000 bp DNA fragments from a 1% agarose gel using the DNA Gel Extraction Kit.

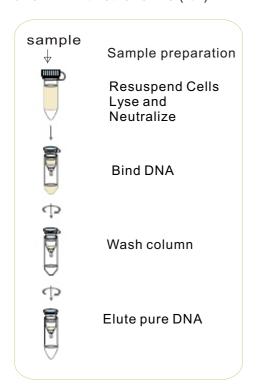
Lane M:  $DS^{TM}5000$ ; Lane1:100bp; Lane 2: 250bp; Lane 3: 500bp; Lane 4: 750bp; Lane 5: 1000bp; Lane 6: 1500bp; Lane 7: 2000bp; Lane 8: 3000bp

Lane 9: 5000bp.

### **Quick Genomic DNA Extraction Kit**

Content	N1111	N1112
Size	50preps	100preps
Solution DS	15ml	30ml
Solution MS	20ml	40ml
Proteinase K	1ml	2ml
Wash Buffer PS	30ml	60ml
Wash Buffer PE	15ml	30ml
Elution Buffer	5ml	10ml
(10mM Tris-HCI, pH 8.5)		
Spin Columns	50each	100each

For the initial use of the kit, dilute the Wash Buffer PE with ethanol 1:3 (v/v).



### Description

Quick Genomic DNA Extraction Kit provides a simple and rapid method for high quality genomic DNA purification from insect, nematode, mamalian tissue. The Genomic DNA system uses the silica-gel-menbrane technology for simple and fast isolation for genomic DNA without phenol/choroform.

#### Feature

Fast--procedure takes only 30 min

High Efficient--gather3-35ug of genomic

DNA from 10mg of tissue

**Universal**--purifies genomic DNA from various sources

**High Purity**--purified DNA is ready for downstream applications

### Application

Fast and efficient extraction of high purity DNA fragments are ready for direct use in all applications, including: Restriction digestion

PCR

labeling

In vitro transcription

### Storage

Store proteinase K at -20°C. Other solutions can be stored for up to 1 year at RT (15-25°C).

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Figure . Extract chicken blood DNA and human blood DNA by using the Quick Genomic DNA Extraction Kit.

Lane M: Lambda DNA/Hind III Lane1: chicken blood DNA

Lane 2: *EcoR* I restriction product for purified chicken

blood DNA

Lane 3: human blood DNA

Lane 4: *Eco*R I restriction product for purified human blood DNA



# 4. RNA Purification Kits

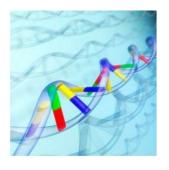


### RNA Summarization

Ribonucleic acid (RNA) is a biologically important type of molecule that consists of a long chain of nucleotide units. RNA is transcribed from DNA by enzyme called RNA Polymerase and is generally further processed by other enzymes. RNA is central to Protein synthesis.

Obtaining high quality, intact RNA is the first and often the most critical step in performing many fundamental molecular biology experiments, including Northern analysis, nuclease protection assays, RT-PCR, RNA mapping, in vitro translation and cDNA library construction. To be successful, however, the RNA isolation procedure should include some important steps both before and after and actual RNA purification.

Dongsheng Biotech offers an expansive portfolio of RNA Purification products that now includes purification and stabilization products. Our unique and trusted TRIzol, General RNA Extraction technologies provide high yields and quantity demanded by our RNA Research team. We offer products for RNA extraction and stabilization, as well as for purifying total RNA from animals, plants, bacterias and cells.







# **Product List**

### **RNA Purification Product**

Product Name	Cat #	Size
TRIzol	R1021	20ml
	R1022	100ml
General Eco-Friendly RNA Extraction Kit	R1031	50 preps
	R1032	100 preps
General RNA Extraction Kit	R1051	50 preps
	R1052	100 preps

# **Reverse Transcription Series**

Product Name	Cat #	Size
M-MLV (recombinant)	R1041	5,000U
RT-PCR Kit	R1011	20 preps
	R1012	100 preps

### **Other Products**

Product Name	Cat #	Size
RNasin (RNase inhibitor)	R2011	1,000U
Oligo d(T)₁₅ Primer	R2021	20µl
Random Primer (6mer)	R2031	20µl
DEPC-treated Water	R2041	100m
dNTPs (RNase-free)	R2051	20µl
RNA Stablization	R2071	100ml
Proteinase K	N9011	100mg
RNase A	N9041	100mg
Agarose	N9051	500g
Dnase I	N9061	1ml
Mini Spin Column	C1021	1000pcs
Midi Spin Column	C2011	100pcs
Maxi Spin Column	C3011	100pcs









E-mail: dongshengbio@hotmail.com



### **TRIzol**

	R1021	R1022
TRIzol	20ml1	100ml

Reagents required, but not supplied.

Chloroform
Isopropyl alcohol
75% ethanol (RNA-free)
DEPC-treated water

### **Application**

The isolated RNA is suitable for any downstream application, including: RT-PCR
Amplification for array analysis
Hybridization assays
In vitro translation

### Storage

Store at  $2-8^{\circ}$ C, propect from light. TRIzol is stable for up to 12 months when properly stored.

### RNA yield with different starting materials.

Starting Material	RNA Yield
Liver tissue	6-10µg/mg tissue
Spleen tissue	6-10 µg/mg tissue
Kidney tissue	3-4 µg/mg tissue
Skeletal muscle or	1-1.5 μg/mg tissue
brain tissue	
Placenta	1-4 µg/mg tissue
Cultured epithelial cells	8-15 μg/10 <sup>6</sup> cells
Fibroblasts	5-7 μg/10 <sup>6</sup> cells

### Description

TRIzol Reagent is a ready-to-use reagent for the isolation of total RNA from cells and tissues. The reagent, a mono-phasic solution of phenol and guanidine isothiocyanate, is an improvement the single-step RNA isolation method. During sample homogenization or lysis, TRIzol Reagent maintains the integrity of the RNA, while disrupting cells and dissolving cell components. TRIzol Reagent facilitates isolation of a variety of RNA species of large or small molecular size. Starting material can be either fresh or frozen tissue, or cell samples of almost any size.

### Feature

**Classical**-used for classical RNA isolation step

High Efficient-stable RNA yield
High Purity-purified RNA is ready for any
downstream application



### **General RNA Extraction Kit**

Content	R1051	R1052
Size	50preps	100preps
Solution RL	60ml	2x60ml
Wash Buffer RPI	18ml	36ml
Wash Buffer RW	12ml	24ml
DEPC-treated water	10ml	20ml
Rnase-free spin culumn	50each	100each
RNase-free	50each	100each
microcentrifuge tube		

For the initial use of the kit, dilute the Wash Buffer RPI with ethanol 3:2 (v/v), and dilute the Wash Buffer RW with ethanol 1:4(v/v).

Reagents required, but not supplied. Chloroform Ethanol (96-100%)

### **Application**

The isolated RNA is suitable for any downstream application, including:

RT-PCR

Amplification for microarray analysis Hybridization assays In vitro transation

### Description

General RNA Extraction Kit can isolate total RNA from a wide range of sample types and amounts. In general, samples are lysed and then homogenized in the presence of guanidinium isothiocyanate, a chaotropic salt capable of protecting the RNA from endogenous RNase. After homogenization, ethanol is added to the sample. The sample is then processed through a spin column containing a clean silica-based membrane to which the RNA binds. Any impurities are effectively removed by subsequent washing. The purified total RNA is then eluted in RNase-free water.

#### Feature

**High Efficient-**Stable RNA yield. Reliable performance of high-quality purified total RNA in downstream applications.

### Storage

Store at  $2-8^{\circ}$ C, propect from light. Kit contents are stable for up to 12 months when properly stored.

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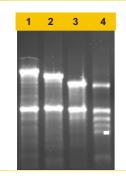


Figure . Extract different species of total RNA by using the Quick Genomic DNA Extraction Kit.

Lane1: Mouse tissue RNA Lane 2: Zebrafish tissue RNA

Lane3: Leguminosae leaf tissue RNA

Lane4: Nicotianatabacum leaf tissue RNA