



## PCRonBlood™ for Mouse Blood

For 100 Reactions of 50µl pr Reactions

Cat. No.: 410100 (100 Reactions)

Cat. No.	Product Name	Size Reac	5x PCR on Blood-Hu	5x PCR on Blood Enhancer-Hu	Taq DNA Pol	dNTP (5 mM)
400100	PCRonBlood™ Human Blood	100	1 mL	1 mL	50 µl	200 µl
400200	PCRonBlood™ Human Blood	100	1 mL	1 mL	not included	not included

Cat. No.	Product Name	Size Reac	5x PCR on Blood-Mo	5x PCR on Blood Enhancer-Mo	Taq DNA Pol	dNTP (5 mM)
410100	PCRonBlood™ Mouse Blood	100	1 mL	1 mL	50 µl	200 µl
410200	PCRonBlood™ Mouse Blood	100	1 mL	1 mL	not included	not included

Store at -20°C. Reagent for in-vitro laboratory use only

### General Description

Mammalian body fluids including blood contain many substances that inhibit the activity of enzymes such as Taq DNA Polymerase. Therefore it is generally necessary to remove the inhibitory substances and purify DNA before performing DNA analysis on cells present in these samples. Traditional preparation of DNA samples from mammalian cells includes overnight treatment with proteinase K, followed by purification by phenol/chloroform extraction and EtOH precipitation. Recently, many kinds of DNA isolation kits have become commercially available, so DNA isolation has become simpler than before, but are still time-consuming. Finally, the DNA isolation procedure generally increases the likelihood of sample contamination with foreign DNA, including DNA from samples processed earlier.

A novel reagent PCRonBlood™ capable of effectively neutralizing the substances in mouse blood that inhibit DNA amplification. Use of PCRonBlood™ for Mouse Blood enables direct amplification of DNA from Mouse blood.

### Features of PCRonBlood™

- ◆ DNA present in blood can be amplified directly.
- ◆ No DNA extraction is required
- ◆ Only small volumes of blood are necessary for the procedure.
- ◆ Risk of cross contamination and/or procedural error is dramatically reduced.
- ◆ Three types of anticoagulants (sodium citrate, dipotassium EDTA and sodium heparinate) are acceptable.
- ◆ Blood (citrate or EDTA-treated blood is recommended) stored frozen for long periods can be analyzed.
- ◆ Easy PCR method permits mass screening and routine use in a broad field of applications.

### Protocol for PCRonBlood™ of Mouse Blood<sup>1</sup>

Prepare the PCR reaction mixture with  
5x PCRonBlood™-Mo  
(Addition of 5x PCRonBlood™ Enhancer-Mo  
may be useful)

↓  
Aliquot 50µl PCR reaction mixture to each PCR tube

↓  
Add 1.0µl<sup>2</sup> mouse blood to each tube<sup>3</sup>

↓  
Overlay mineral oil, if needed

↓  
PCR<sup>4-7</sup>

### Preparation of the PCR Reaction Mixture

5x PCRonBlood™-Mo	10 µl
5x PCRonBlood™ Enhancer-Mo	10 µl
dNTP (5 mM each)	2 µl
5'-Primer <sup>8</sup>	0.5 µM
3'-Primer <sup>8</sup>	0.5 µM
Taq DNA Pol (5 U/µl) <sup>8,9</sup>	0.5 µl
Distilled water	to 50 µl

1. We recommend that these procedures be carried out on ice (except when Hot Start PCR is used) to avoid any non-specific reactions and/or reduced sensitivity.

2. We recommend the use of 1.0µl Mouse blood in 50µl reaction volume or 0.5µl in 20µl reaction volume, as excess blood may in some cases reduce efficiency, depending on the individual blood samples and primer sequences used.

3. Any agitation should be strictly avoided.

4. Temperature profile for direct PCR of Mouse blood:

Pre-heating at 80°C for 15 min<sup>5</sup>  
94°C for 4.5 min

Followed by 40 cycles<sup>6</sup>

94°C for 30 sec  
Annealing Temperature<sup>7</sup> for 1 min

72°C for 1 min

Finally 72°C for 7 min

5. Pre-heating at 80°C for 15 min is useful when fresh blood is used.

6. Five more PCR cycles are required than for purified DNA.

7. We recommend decreasing annealing temperature in 1°C increments, when poor PCR product is detected.

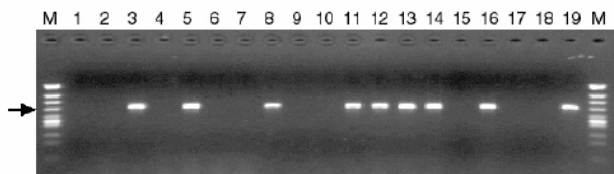
8. Included in this version of PCRonBlood™

9. PCRonBlood™ is effective for Taq DNA Polymerase with the exception of polymerase activated chemically by higher temperature (e.g. AmpliTaq Gold, Applied Biosystems, and HotStar Taq DNA Polymerase, Qiagen). However, Taq DNA Polymerase plus anti-Taq antibody permits hot start direct PCR.

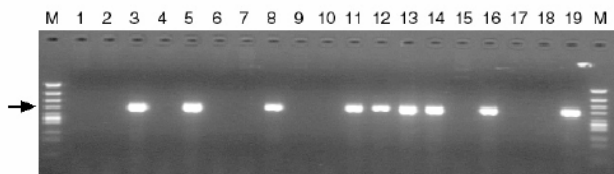
## Experimental Result

Detection of transgene in transgenic mice. PCRonBlood™ detection of the *lck* promoter-human D4-GDI transgene in 19 transgenic C57BL/6J mice using blood (a), was compared with the results of conventional PCR using purified DNA (b). The specific product is indicated at 512bp by an arrow.

### a. Direct PCR



### b. Conventional PCR



LaneM:  $\phi$ X174 RF DNA digested with *HincII*.

PCRonBlood™ using 1  $\mu$ l of heparinized blood was carried out in 50  $\mu$ l of PCRonBlood™ mixture. Conventional PCR using purified DNA obtained from tissue samples of mouse tails was carried out in 50  $\mu$ l of standard mixture PCR was performed at 80°C for 15 min, then 94°C for 4.5 min, followed by 40 cycles of amplification (94°C for 30 s, 55°C for 1 min, 72°C for 1 min) followed by final extension.

## Related Products

Description	Cat. No.
Taq DNA Polymerase (500 Units) with 10X Ammonium Reaction Buffer with 10X Standard Reaction Buffer	110303
Taq DNA Polymerase (500 Units) with 10X Combination Buffer	110403
Taq DNA Polymerase (500 Units) with 10X Mg <sup>++</sup> Free Ammonium Buffer	110503
Taq DNA Polymerase 2.0X Master Mix (100 Reac) with 2.0 mM MgCl <sub>2</sub>	150301
Taq DNA Polymerase 2.0X MaMi RED (100 Reac) with 1.5 mM MgCl <sub>2</sub> ,	180301
Taq DNA Polymerase 2.0X MaMi RED (100 Reac) with 2.0 mM MgCl <sub>2</sub>	190301
AccuPOL DNA Polymerase (500 Units)	210303
TEMPase Hot Start DNA Polymerase (500Units) with 10X TEMPase Buffer I with 10X TEMPase Buffer II	220303
UniPOL –Long Range PCR (100 Reac)	270701
Rapid Ligation Kit (50 React)	750300
RT-PCR One Tube (100 Reac)	740301
TEMPase Hot Start 2X Master Mix with TEMPase Buffer I (100 Reac)	230301
TEMPase Hot Start 2X Master Mix with TEMPase Buffer II (100 Reac)	230701
dNTP Mix (2 x 500 $\mu$ l) (12.5 mM of each dA, dC, dG and dT)	501004
dNTP Mix, (2 x 500 $\mu$ l) (10 mM of each dA, dC, dG and dT),	502004
GC5 Value Efficiency, 10 <sup>8</sup> Cfu/ $\mu$ g pUC19 Chemically Competent Cells, (10x 200 $\mu$ l)	812010
GC5 High Efficiency, 10 <sup>9</sup> Cfu/ $\mu$ g pUC19 Chemically Competent Cells, (10x 50 $\mu$ l)	805010
GC5 High Efficiency, 10 <sup>9</sup> Cfu/ $\mu$ g pUC19 Chemically Competent Cells, (5x 200 $\mu$ l)	802005
SuperPath GC10, 10 <sup>10</sup> Cfu/ $\mu$ g pUC19 ElectroCompetent Cells, (5x 80 $\mu$ l)	830805
SOC Medium, 10x 10mL	800000

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### NOTICE

In certain countries, patents cover the PCR process. This product is intended for researchers having a license to perform PCR or those not required to obtain a license.