

## VAS Taq

<b>Product:</b>	<b>VAS Taq</b>
<b>CATALOG#:</b>	<b>BN0150-0250</b>
<b>Units:</b>	<b>250 U</b>
<b>Concentration:</b>	<b>5 units/μl</b>
<b>Volume:</b>	<b>50 μl</b>
<b>Storage:</b>	<b>-20°C</b>

### Supplied form

20mM Tris-HCl pH8.0, 50%glycerol,  
100mM KCl, 0.1Mm EDTA, 1mM DTT,  
0.5%Tween20, 0.5%NP-40

### Description

VAS Taq is isolated and purified from an E.Coli. strain that carries the cloned DNA polymerase gene from *Thermus aquoticus* YT-1strain.

**Purity(SDS-PAGE) > 99%**

### 10X VAS Taq PCR Buffer

Composition: Tris-HCl, KCl, (NH<sub>4</sub>)<sub>2</sub>SO<sub>4</sub>,  
15mM MgCl<sub>2</sub>, pH8.3 .

### Unit definition

One unit is the amount of enzyme that will incorporate 10 nmoles of dNTPs into acid-insoluble products at 72°C in 30 min.

### PCR performance test

Good performance of DNA amplification by PCR is confirmed by using 0.2U Taq will amplified 2.5 kb DNA fragment in a 50μl reaction volume.

### General reaction mixture for PCR ( total 50μl )

VAS Taq (5 units/μl )	0.5μl
10X Taq Buffer	5.0μl
dNTPs(2.5mM)	4.0μl
Template	< 1μg
Primer 1	0.2~1.0μM
Primer 2	0.2~1.0μM
H <sub>2</sub> O	upto 50μl

### PCR products

As most PCR product amplified with VAS Taq have one A added at 3`- terminus, the obtained PCR product can be directly used for cloning into T-vector.