

## AMV Reverse Transcriptase (Recombinant)

Product No : ME2301

Quantity : 500u



Lot :  
 Expiry Date :  
 Concentration : 20u/μl  
 Supplied with : 10X Buffer AMV-RT  
 (New improved buffer system)

Store at -70°C for long term periods  
 Store at -20°C for short term (1 month)



info@vivantechnologies.com

### Description:

Avian Myeloblastosis Virus (AMV) Reverse Transcriptase is an RNA dependent DNA polymerase ( $\alpha\beta$  holoenzyme) with molecular weight of 157,000 daltons. This enzyme can synthesize a complementary DNA strand initiating from a primer using either RNA (cDNA synthesis) or single-stranded DNA as a template.

### Features:

- Ultra pure recombinant protein.
- Maintains the RNA- and DNA-dependent DNA polymerase and RNase H activities.
- RNase H activities can be regulated over a wide range of temperatures.
- Capable of synthesizing cDNA over a wide range of temperatures.

### Unit Definition:

1u is defined as the amount of enzyme that is required to incorporate 1nmol of dTMP into acid-insoluble material in 10 minutes at 37°C using Poly(rA)•(dT)<sub>12-18</sub> as a template primer.

### Assay Conditions:

50mM Tris-HCl (pH 8.3), 40mM KCl, 6mM MgCl<sub>2</sub>, 4mM DTT, 0.4mM poly(rA)•(dT)<sub>12-18</sub> and 0.5mM [<sup>3</sup>H]-TTP (10-20c/m/pmol) in a reaction volume of 25μl.

### Reaction Buffer:

#### 10X Buffer AMV-RT

250mM Tris-HCl (pH 8.3), 500mM KCl, 50mM MgCl<sub>2</sub> and 20mM DTT.

### Storage Buffer:

200mM K-phosphate (pH 7.2), 0.2% Triton™ X-100, 2mM DTT and 50% glycerol.

**Thermal Inactivation:** 80°C for 10 minutes

### Application:

- First strand synthesis of cDNA.
- Synthesis of cDNA for cloning.
- cDNA labeling.
- Primer extensions and RNA sequencing.
- RT-PCR.
- Dideoxy sequencing of DNA and RNA.

### Quality Control:

All preparations are assayed for contaminating endonuclease, exonuclease and non-specific RNase, as well as single- and double-stranded DNase activities.

\*Due to viscosity of the enzyme, please briefly centrifuge the vial for full recovery of the enzyme to the bottom upon receipt. Please use positive displacement pipets or low-binding pipet tips for maximum transferring efficiency.

Product Use Limitation

This product is for research purposes and *in vitro* use only.

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DSME2301\_rev0\_010311

## AMV Reverse Transcriptase (Recombinant)

Product No : ME2302

Quantity : 2500u



Lot :  
 Expiry Date :  
 Concentration : 20u/μl  
 Supplied with : 10X Buffer AMV-RT  
 (New improved buffer system)

Store at -70°C for long term periods  
 Store at -20°C for short term (1 month)



info@vivantechnologies.com

### Description:

Avian Myeloblastosis Virus (AMV) Reverse Transcriptase is an RNA dependent DNA polymerase ( $\alpha\beta$  holoenzyme) with molecular weight of 157,000 daltons. This enzyme can synthesize a complementary DNA strand initiating from a primer using either RNA (cDNA synthesis) or single-stranded DNA as a template.

### Features:

- Ultra pure recombinant protein.
- Maintains the RNA- and DNA-dependent DNA polymerase and RNase H activities.
- RNase H activities can be regulated over a wide range of temperatures.
- Capable of synthesizing cDNA over a wide range of temperatures.

### Unit Definition:

1u is defined as the amount of enzyme that is required to incorporate 1nmol of dTMP into acid-insoluble material in 10 minutes at 37°C using Poly(rA)•(dT)<sub>12-18</sub> as a template primer.

### Assay Conditions:

50mM Tris-HCl (pH 8.3), 40mM KCl, 6mM MgCl<sub>2</sub>, 4mM DTT, 0.4mM poly(rA)•(dT)<sub>12-18</sub> and 0.5mM [<sup>3</sup>H]-TTP (10-20c/m/pmol) in a reaction volume of 25μl.

### Reaction Buffer:

#### 10X Buffer AMV-RT

250mM Tris-HCl (pH 8.3), 500mM KCl, 50mM MgCl<sub>2</sub> and 20mM DTT.

### Storage Buffer:

200mM K-phosphate (pH 7.2), 0.2% Triton™ X-100, 2mM DTT and 50% glycerol.

**Thermal Inactivation:** 80°C for 10 minutes

### Application:

- First strand synthesis of cDNA.
- Synthesis of cDNA for cloning.
- cDNA labeling.
- Primer extensions and RNA sequencing.
- RT-PCR.
- Dideoxy sequencing of DNA and RNA.

### Quality Control:

All preparations are assayed for contaminating endonuclease, exonuclease and non-specific RNase, as well as single- and double-stranded DNase activities.

\*Due to viscosity of the enzyme, please briefly centrifuge the vial for full recovery of the enzyme to the bottom upon receipt. Please use positive displacement pipets or low-binding pipet tips for maximum transferring efficiency.

Product Use Limitation

This product is for research purposes and *in vitro* use only.

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DSME2302\_rev0\_010311