

## ENZYMES

# TTx DNA Polymerase

ORIGIN *E.coli (Recombinant)*

CAT# TTX Range

EC# 2.7.7.7

### ▶ ENZYME SPECIFICATION

Ribonuclease Activity: None detected

Endonuclease Activity: None detected

Nicking Activity: None detected

Glycerol Content: ≤ 0.1% (Glycerol free product specification only)

### ▶ HOT START TTx (DNA) KIT SPECIFICATION

Tested for functional performance by amplifying from dilutions of enterovirus RNA in combination with 5x Buffer for rTth/ TTx (DNA/ RNA)

The slope from analysis is as follows:  $-3.74 \leq \text{slope} \leq -3.01$

The correlation coefficients from both analyses are as follows:  $r \leq -0.980$

### ▶ HOT START TTx (RNA) KIT SPECIFICATION

Tested for functional performance by amplifying from dilutions of enterovirus RNA

The slope from analysis is as follows:  $-3.74 \leq \text{slope} \leq -3.01$

The correlation coefficients from both analyses are as follows:  $r \leq -0.980$

### ▶ PRODUCT FORMAT

PART #	DESCRIPTION	CONTENT/ FORMAT
TTX-119	TTx DNA Polymerase	TTx DNA Polymerase 1KU
TTX-129		TTx DNA Polymerase 10KU
TTX-159		TTx DNA Polymerase 100KU
TTX-219	TTx DNA Polymerase <Glycerol free>	TTx DNA Polymerase <Glycerol free> 1KU
TTX-229		TTx DNA Polymerase <Glycerol free> 10KU
TTX-259		TTx DNA Polymerase <Glycerol free> 100KU
HSTTX-101	Hot Start TTx (DNA) kit	Hot Start TTx DNA Polymerase (4U/uL) 62.5uL 2x Buffer for rTth/TTx (DNA) 1.25mL x2
HSTTX-111	Hot Start TTx (RNA) kit	Hot Start TTx DNA Polymerase (4U/uL) 62.5uL 5x Buffer for rTth/TTx (DNA/RNA) 1mL 50mM Mn (OAc) <sub>2</sub> 250uL 2mM dNTPs 1 mL

### ▶ DESCRIPTION AND APPLICATION

TTx DNA Polymerase has a high reverse transcription activity in the presence of Mn<sup>2+</sup> ions and allows for 1-Step 1-Enzyme RT-PCR, including reverse transcription and PCR steps. Effective for amplification from crude samples containing PCR inhibitors with high efficiency for both DNA and RNA. Suitable for amplification from low copies of template DNA/RNA and is efficient even in fast cycle condition. Glycerol free format available, enabling the making of a lyophilized mix.

In addition, TTx DNA Polymerase has a 5' → 3' exonuclease activity, so it can be used for real-time PCR using probe assays such as TaqMan<sup>®</sup> assay.

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**SEKISUI**  
DIAGNOSTICS

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## CHARACTERISTICS

### Enzyme features:

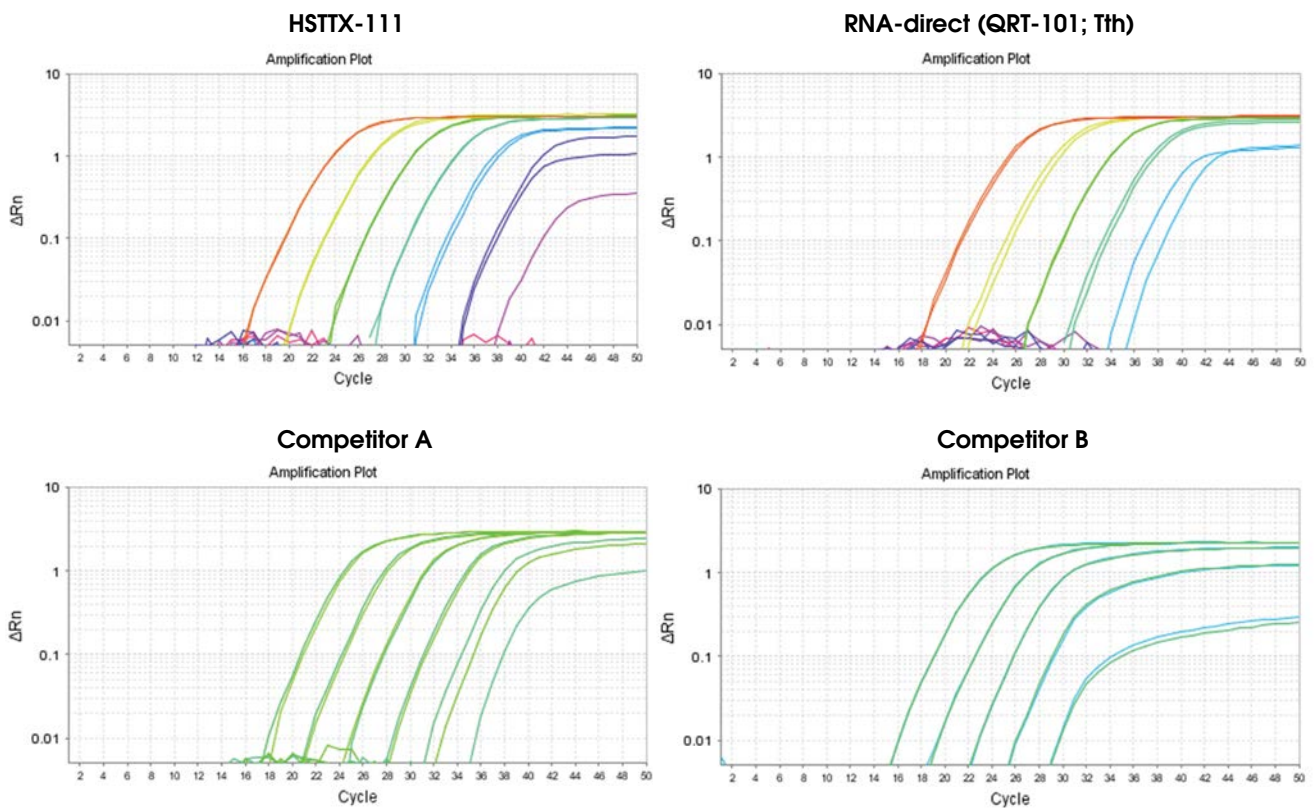
- Enables highly efficient 1-Step 1-Enzyme RT-PCR
- Effective amplification from crude samples
- Fast- High speed qPCR cycling detection
- Suitable for Multiplex PCR
- Lyo-ready glycerol free format available

### Highly efficient 1-Step 1-Enzyme RT-PCR

Influenza virus RNA was detected using TaqMan® probes. The Hot Start TTx (RNA) Kit obtained higher sensitivity than Tth DNA Polymerase based reagent (QRT-101), and 2 competitor PCR enzyme kits. By using TTx DNA Polymerase, efficient detection is possible even from low copies of template.

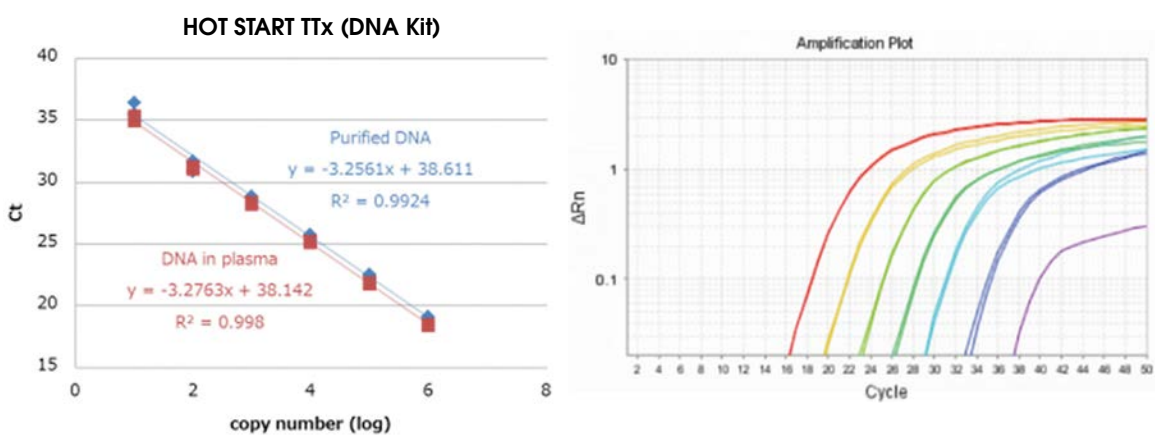
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TARGET GENE DILUTION	HSTX-111 Ct		RNA-DIRECT (QRT-101; TTH) Ct		COMPETITOR A Ct		COMPETITOR B Ct	
1/10 <sup>2</sup>	17.98	17.90	19.83	19.64	19.56	19.32	17.42	17.43
1/10 <sup>3</sup>	21.50	21.52	23.45	23.97	22.90	22.68	20.88	20.83
1/10 <sup>4</sup>	25.01	25.10	28.35	28.40	26.23	26.38	24.26	24.28
1/10 <sup>5</sup>	28.71	28.79	32.21	31.75	29.89	29.70	27.49	27.64
1/10 <sup>6</sup>	32.09	32.41	36.94	35.35	33.92	32.85	31.12	30.92
1/10 <sup>7</sup>	36.08	36.37	N/A	N/A	N/A	N/A	N/A	N/A
1/10 <sup>8</sup>	N/A	39.94	N/A	N/A	N/A	36.54	N/A	N/A
H <sub>2</sub> O	N/A	N/A	N/A	N/A	N/A	N/A	N/A	N/A



### Effective amplification from crude samples

African swine fever virus DNA was detected using TaqMan® probes. Performing the reaction in 20µL mixture with and without 2.5µL of plasma, Hot Start TTx (DNA) Kit was able to detect the African swine fever virus DNA.



### Fast- High speed qPCR cycling detection

African swine fever virus DNA was detected using TaqMan® probes. Only TTx polymerase could detect at high speed cycling conditions.

TARGET GENE SIZE	HSTTH-301 Ct		HSTTX-101 Ct		COMPETITOR A Ct	
2pg	19.57	21.94	18.88	19.09	12.45	13.62
0.2pg	28.46	N/A	21.91	22.09	17.25	18.25
0.02pg	32.72	29.93	24.63	24.65	20.78	19.99
2fg	N/A	N/A	27.44	27.47	24.02	22.44
0.2fg	N/A	N/A	30.99	30.89	27.04	26.82
0.02fg	N/A	N/A	34.34	34.38	30.63	29.08
0.002fg	N/A	N/A	35.21	36.85	34.00	35.56
NTC	N/A	N/A	N/A	N/A	N/A	N/A

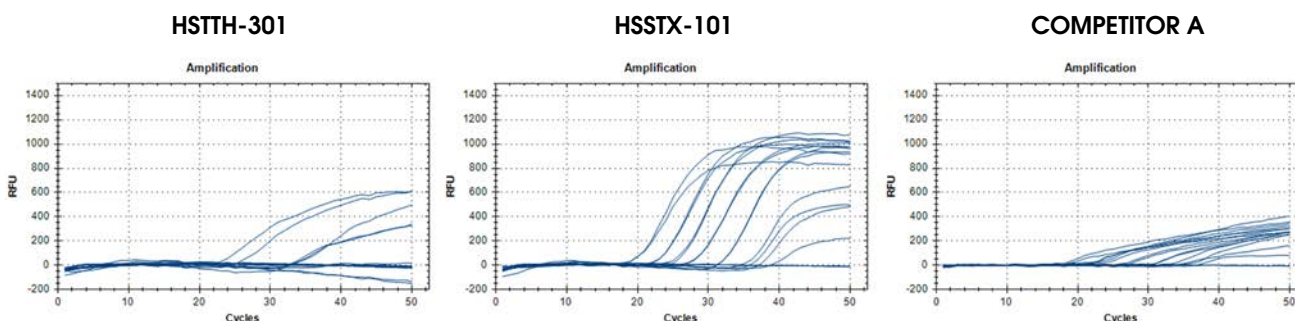
PCR cycling conditions

95°C, 2 min

95°C, 1 sec

60°C, 1 sec

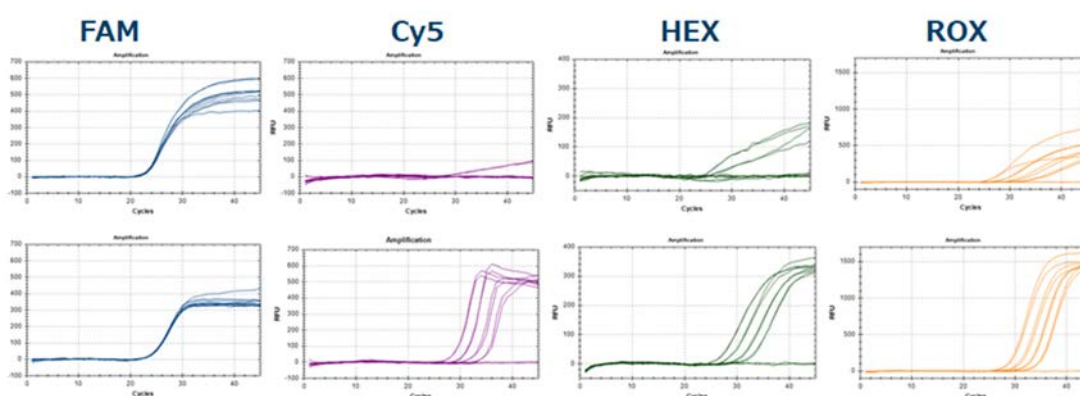
X50



**Suitable for Multiplex PCR**

The Hot Start TTx (DNA) Kit and Taq DNA Polymerase based real-time PCR reagents were mixed with a TaqMan Probe of primers to detect Salmonella, Shigella, and O157, and multiplex PCR was used to detect genes (1,000, 250, 63, and 15 copies).

**TAQ BASED REAGENT**



**HSTTX-101**

**THE AMERICAS**

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