



CAP DISCOVERY SET

Below is the IVT condition we used in our own lab to get high yield and high capping efficiency, if you have your own recipe, please feel free to use yours.

Components	Volume	
	Cap3011 Cap5011 Cap2811 Cap3311	Cap3411
Cap Analog	2 μ L	2 μ L
10xTranscription Buffer*	2 μ L	2 μ L
100mM ATP Solution	2 μ L	1 μ L
100mM CTP Solution	2 μ L	1 μ L
100mM GTP Solution	2 μ L	1 μ L
100mM NMPUTP Solution	2 μ L	1 μ L
T7 RNA Polymerase (250U/ μ L)	1 μ L	1 μ L
Inorganic Pyrophosphatase (1U/ μ L)	0.04 μ L	0.04 μ L
RNase Inhibitor (40U/ μ L)	1 μ L	1 μ L
RNase free ddH ₂ O	up to 20 μ L	up to 20 μ L
DNA Template	1 μ g	1-2 μ g

*Note on buffer compositions

CAP3011, CAP5011, CAP3311 and CAP2811 IVT buffer (10x Transcription Buffer): 400mM Tris-HCl, 450 mM Mg(CH₃COO)₂, 20mM spermidine, 100mM DTT, pH 7.9 at 25°C.

Cap3411 IVT buffer (10x Transcription Buffer): 400mM Tris-HCl, 250 mM Mg(CH₃COO)₂, 20mM spermidine, 100mM DTT, 0.1% Triton X-100, pH 7.9 at 25°C.

