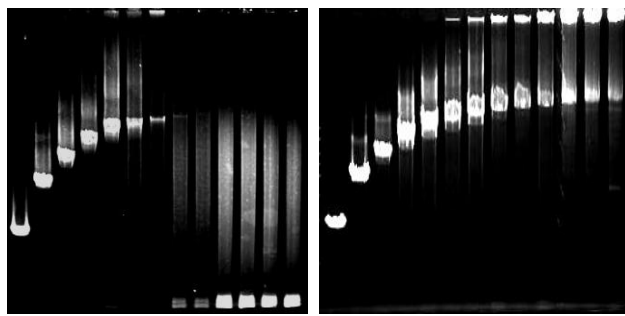


## Taq Blend with Pfu (+dNTPs)

<b>Product code</b>	02-120	02-120-5
<b>Size</b>	200 U	5 x 200 U
<b>Storage</b>	-20°C	
<b>Product Description</b>	Taq Blend with Pfu is optimized blend of Taq and Pfu DNA polymerases. The proof-reading 3'→5' exonuclease activity of Pfu increases the fidelity and robust amplification of Taq DNA polymerase. The reaction buffer has been formulated for robust yields and long PCR.	
<b>Concentration</b>	5 U/μl	
<b>Buffer</b>	35 mM Tris-HCl (pH 8.0), 50 mM KCl, 1 mM DTT, 0.1 mM EDTA, 50 % glycerol, 0.75 % Tween-20, 0.75 % Igepal CA-630	
<b>Component</b>	<b>Taq Blend with Pfu:</b> 5 U/μl Taq and Pfu DNA polymerase (02-Tbp 40 μl) <b>5 x Reaction Buffer for Taq Blend with Pfu</b> (02-B1b 2 ml) <b>dNTPs (2.5mM each)</b> (02-Dnt 640 μl)	
<b>Quality Assurance</b>	Greater than 95% purity as determined by SDS-PAGE (CBB staining). The absence of endonucleases 3→5 amplification was attained with λ DNA template was confirmed.	
<b>PCR product</b>	Good amplification result was obtained in PCR reaction using λ phage DNA as a template (Fig.2).	
<b>Data</b>	<p><b>Experimental Example</b></p> <p>Robustness of Taq Blend with Taq as compared Taq Economy.</p> <p><u>PCR conditions</u></p> <p>98°C 5 sec            94°C 1 min →            68°C 4-20 min (30 cycles)</p> <p>(extention time at 68°C)</p> <p>2-8kbp:4min 10-14kbp:7min 16-18kbp:10min 20-35kbp:20min</p> <p><u>Result</u></p> <p>Taq Blend with Pfu could amplify up to 35 kb template while Taq could amplify up</p> <p>2 4 6 8 10 12 14 16 18 20 25 30 35      2 4 6 8 10 12 14 16 18 20 25 30 35</p>  <p>Fig.1                      Fig.2</p>	
Please note: All products are FOR RESEARCH USE ONLY. NOT FOR USE IN DIAGNOSTIC PROCEDURES. NOT FOR MILITARY USE.		