

Table 2. Primer sets used to produce T7 RNA polymerase templates

Gene	F/R [†]	Primer sequence [‡]	Size, nt
<i>pgACBP</i>	F	<u>CCGGATCCTAATACGACTCACTATAGGGCGATGTCTCTCCAAGAAAAAT</u>	324
	R	<u>CCGGATCCTAATACGACTCACTATAGGGCGATGGAAGCTATGAGACCTC</u>	
<i>mgACBP</i>	F	<u>CCGGATCCTAATACGACTCACTATAGGGCGATGTCTCTCGACGAGCAAT</u>	297
	R	<u>CCGGATCCTAATACGACTCACTATAGGGCGGCGTATTTGGAGTGGAGTTT</u>	
<i>pgFAR</i>	F	<u>CCGGATCCTAATACGACTCACTATAGGGCGGGCACGAGGCGACGAAAC</u>	1,960
	R	<u>CCGGATCCTAATACGACTCACTATAGGGCGGGTAACTATAGAAATGGTTTAAC</u>	
<i>Bmpgdesat1</i>	F	<u>CCGGATCCTAATACGACTCACTATAGGGCGGATGCCTCCTAATTCAGTG</u>	990
	R	<u>CCGGATCCTAATACGACTCACTATAGGGCGGATTCATTCCATTGGTCG</u>	
<i>PBANR</i> [*]	F	<u>TAATACGACTCACTATAGGGAGATAGTGACTATTACGGCATT</u>	417
	R	<u>TAATACGACTCACTATAGGGAGATGACCCTCCTCTGTGAGC</u>	
<i>EGFP</i> [*]	F	<u>TAATACGACTCACTATAGGGAGAATGGTGAGCAAGGGCG</u>	717
	R	<u>TAATACGACTCACTATAGGGAGACTTGTACAGCTCGTCC</u>	

[‡] Nucleotide sequences corresponding to the T7 promoter region are underlined.

^{*} Modified T7 promoter sequences used for PCR amplification.

[†] F indicates forward primer; R indicates reverse primer.