

Supplementary Table S2. Primers used in this work.

Primer name	Sequence (5'-3') <sup>a, b</sup>	Experiment
leuAF	TGCTCCCTACCAACTCTATTC	Selective marker amplification, PCR validation
leuAR	GTCGAGTTGACCAGAATGTAC	Selective marker amplification
dmt2-F5'	GCCTGCACGGTATACTGACG	<i>dmt2</i> deletion construct
dmt2-R5'+ leuA	<u>GAATAGAGTTGGTAGGGAGCATCAAGAAGGTGGAGGAGAAACA</u>	<i>dmt2</i> deletion construct
dmt2-F3'+ leuA	<u>GTACATTCTGGTCAACTCGACAGCATAGTAAAAGATAGATGGAGCGT</u>	<i>dmt2</i> deletion construct
dmt2-R3'	CACGCGGAATCTGAATGACG	<i>dmt2</i> deletion construct
dmt3-F5	CGGTTGCCCTTTCTTTCTCG	<i>dmt3</i> deletion construct
dmt3-R5+leuA	<u>GAATAGAGTTGGTAGGGAGCATCTCTTTGTGATGAAGGTTGGA</u>	<i>dmt3</i> deletion construct
dmt3-F3+leuA	<u>GTACATTCTGGTCAACTCGACAGCCATTTTGTGATTTGAGGT</u>	<i>dmt3</i> deletion construct
dmt3-R3	GTCCCGGTAAGTACAACGCA	<i>dmt3</i> deletion construct
pkaR4-TR-F	AGTCTGGCCTTGCCGTGTTT	RT-qPCR
pkaR4-TR-R	GCAACGACCGTAGCTGCTCT	RT-qPCR
dmt1-F+ XhoI	TCCCTCGAGAGGTCCCCCTTGACACGTGGTTC	<i>dmt1</i> ORF
dmt1-R+ SacII	TCACCGCGGAGTGCACCATGGATTCTTGTC	<i>dmt1</i> ORF
P1	GAAAGCCTGCATTGGATTGGC	<i>dmt2</i> replacement confirmation
P2	CGAAAAGTGCCCTATCGG	<i>dmt2</i> replacement confirmation
P3	GTACACTGTTGCTCTCCATGTG	<i>dmt3</i> replacement confirmation
P4	AGCAAATCGTCGTGGATGGT	<i>dmt3</i> replacement confirmation
pyrGF	TGCCTCAGCATTTGGTACTTG	Selective marker amplification
pyrGR	GTACACTGGCCATGCTATCG	Selective marker amplification
85076 F	ACAATCTCGGCCTCGGAACAAGTGC	<i>dmt1</i> deletion construct
85076 RpyrG	<u>CAAGTACCAATGCTGAGGCATGCACCTTGCTGTCATGGCAACAAGC</u>	<i>dmt1</i> deletion construct
85076 FpyrG	<u>CGATAGCATGGCCAGTGTA</u> CGCTTGACCAATGGATACCTACTGAC	<i>dmt1</i> deletion construct
85076 R	ATCCCCCTTGACACGTGGTTCGTAGC	<i>dmt1</i> deletion construct
115786 F	CTCCATGCTGATATCTGTACGCTGT	<i>dmt3</i> deletion construct
115786 RpyrG	<u>CAAGTACCAATGCTGAGGCATGAGGTT</u> CAGCTGTGATTGGCTCCA	<i>dmt3</i> deletion construct
115786 FpyrG	<u>CGATAGCATGGCCAGTGTA</u> CTGCTGCTGGTGATTGACAGTTGCTG	<i>dmt3</i> deletion construct
115786 R	GATGGCTACTGACAAGATCGTTGCTG	<i>dmt3</i> deletion construct
106998 F	AGAGCACGCGGAATCTGAATGACG	<i>dmt2</i> deletion construct
106998 RpyrG	<u>CAAGTACCAATGCTGAGGCAGTCTATCCACGACATCAGAGCGAGGA</u>	<i>dmt2</i> deletion construct
106998 FpyrG	<u>CGATAGCATGGCCAGTGTA</u> CCGAAAATAGTGTGCGCCAGCAGGATG	<i>dmt2</i> deletion construct

106998 R	CTTGTGGACATGGAAGATGCTGCT	<i>dmt2</i> deletion construct
CarRP11	CATACAAAGCACGAGTTTCC	PCR validation
dmt1-TR-F	TGCCTCGTAAACCAACCACT	PCR validation
qPCR-EF1-F	GTCCGTGATATGCGTCAAACC	RT-qPCR
qPCR-EF1-R	AGCGGCCTTGGTGACCTTAC	RT-qPCR
Udmt1F	ACAATCTCGGCCTCGGAACAAGTGC	Southern blot validation
Ddmt1-pyrGR	CAAGTACCAATGCTGAGGCATGCACCTTGCTGTCATGGCAACAAGC	Southern blot validation
Udmt2F	AGAGCACGCGGAATCTGAATGACG	Southern blot validation
Udmt2-pyrGR	CAAGTACCAATGCTGAGGCAGTCTATCCACGACATCAGAGCGAGGA	Southern blot validation
Udmt3F	CTCCATGCTGATATCTGTACGCTGT	Southern blot validation
Ddmt3-pyrGR	CAAGTACCAATGCTGAGGCATGAGGTTGAGCTGTGATTGGCTCCA	Southern blot validation

<sup>a</sup> Underlined sequences are complementary to selection marker *pyrG* or *leuA*

<sup>b</sup> Bold bases corresponding to the cutting sites to restriction enzymes