

Supplementary Information for:

**Assessment on the oil accumulation by knockdown of triacylglycerol lipase in the
oleaginous diatom *Fistulifera solaris***

Yoshiaki Maeda^a, Kahori Watanabe^a, Marshila Kaha^a, Yusuke Yabu^a, Tomoko Yoshino^a, Mitsufumi
Matsumoto^b, Tsuyoshi Tanaka^{a*}

^aDivision of Biotechnology and Life Science, Institute of Engineering, Tokyo University of
Agriculture and Technology, 2-24-16 Naka-cho, Koganei, Tokyo 184-8588, Japan,

^bBiotechnology Laboratory, Electric Power Development CO., Ltd., 1, Yanagisaki-machi,
Wakamatsu-ku, Kitakyushu 808-0111, Japan

*Corresponding author

E-mail address: tsuyo@cc.tuat.ac.jp

Tel: +81-42-388-7401

Fax: +81-42-385-7713

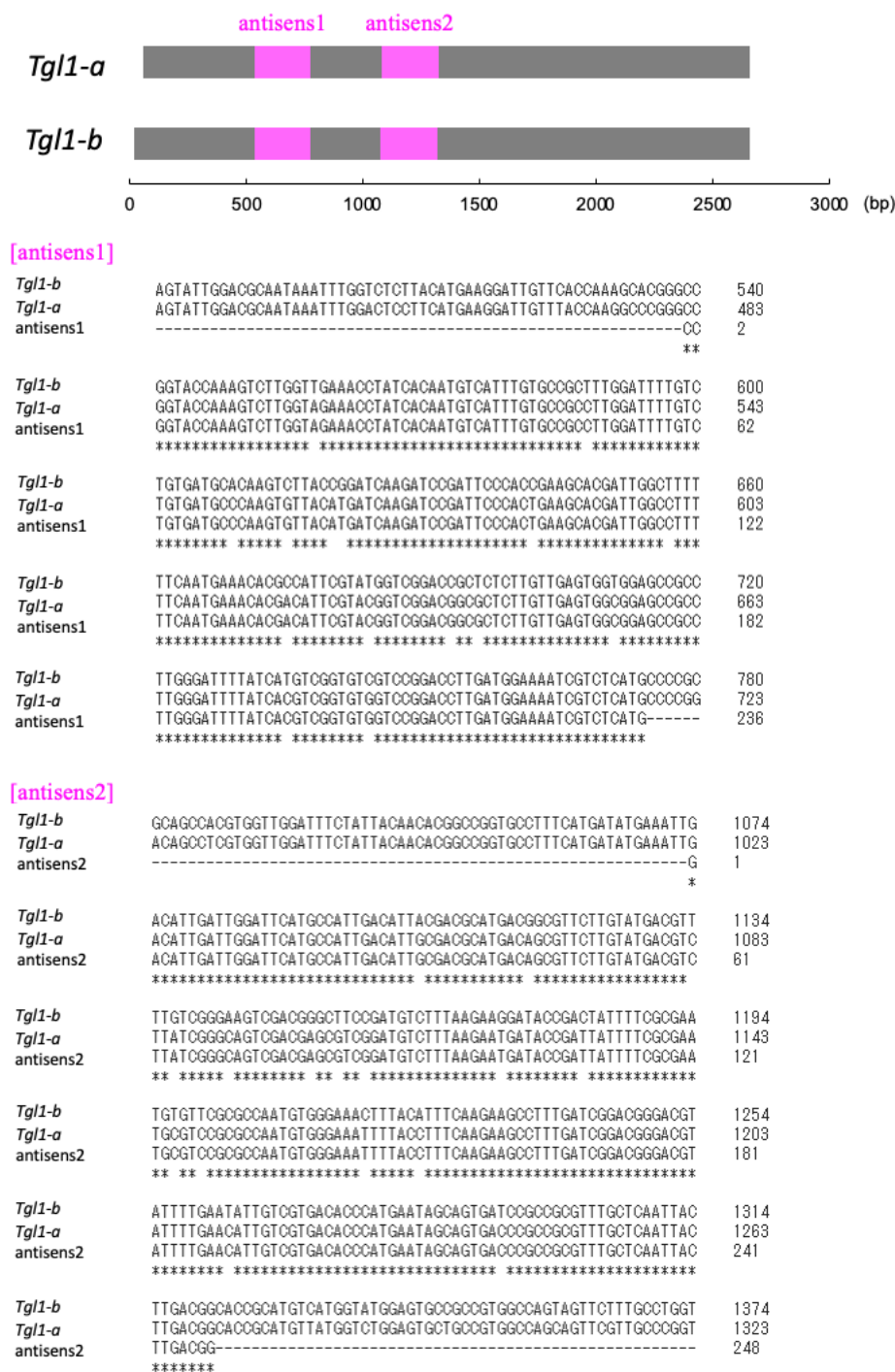


Fig. S1 Sequence alignments of *Tgl1* gene (*Tgl1-a* and *Tgl1-b*) and antisense fragments.

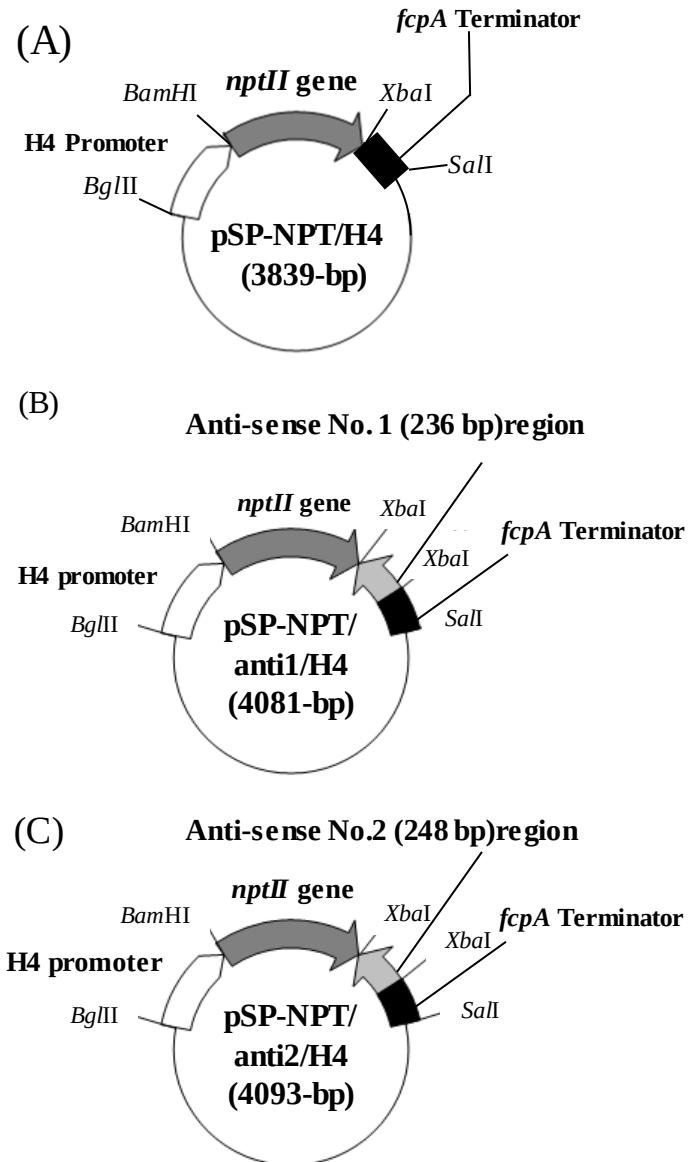


Fig.S2 Constructions of vectors used in this study. Restriction maps of the expression cassettes in plasmids pSP-NPT/H4 (A), pSP- ANT1 (B), and pSP-ANT2 (C).

Abbreviations: *H4* promoter, histone H4 gene promoter from *F. solaris*; *fcp A* terminator, fucoxanthin chlorophyll *a/c*-binding protein A gene terminator from *P. tricornutum*.

Table S1 Primers used in this study

Primer name	Sequence (5'-3')	Use
Tgl1_anti_2F	AAATTTTCTAGACATGAGACGATTTTCCATCA	Cloning by restriction enzyme
Tgl1_anti_2R	AAATTTTCTAGACCGGTACCAAAGTCTTGG	
Tgl1_anti_3F	AAATTTTCTAGACCGTCAAGTAATTGAGCAAA	
Tgl1_anti_3R	AAATTTTCTAGAGACATTGATTGGATTCATGC	
anti2_fwd	TGATCTAGACATGAGACGATTTTCCATC	Cloning by Gibson assembly
anti2_rev	GTTTCTAGACCGGTACCAAAGTCTTGG	
anti2-1_fwd	GGTACCGGTCTAGAAACAACTACCTCGACTTT	
anti2-1_rev	CGGAGGCAGATCTGGTTCCCGCATAG	
anti2-2_fwd	ACCAGATCTGCCTCCGGTACTCTTAC	
anti2-2_rev	GTCTCATGTCTAGATCAGAAGAACTCGTC	
anti3_fwd	TGATCTAGACCGTCAAGTAATTGAGCAAAC	
anti3_rev	GTTTCTAGAGACATTGATTGGATTCATGC	
anti3-1_fwd	ATCAATGTCTCTAGAAACAACTACCTCGACTTT	
anti3-1_rev	CGGAGGCAGATCTGGTTCCCGCATAG	
anti3-2_fwd	ACCAGATCTGCCTCCGGTACTCTTAC	
anti3-2_rev	CTTGACGGTCTAGATCAGAAGAACTCGTC	
anti_seq_F	GAATATCATGGTGGAAAATGG	Confirmation of introduction of antisense sequence
anti_seq_R	CATGGTGAAGACGAGCTAGT	
RNAi_nptII_F	TGAACAAGATGGATTGCAC	Sequence analysis of Neomycin cassette
RNAi_nptII_R	CAGAAGAACTCGTCAAGAAGG	
qPCR_GAPDH_F	ATTGGAGTCAATGGCTTTG	Quantitative RT-PCR to determine quantity of GAPDH gene expression level
qPCR_GAPDH_R	CCATGAACGGAATCGTACT	
qPCR_g13_anti2F	TTGGACTCCTTCATGAAGG	
qPCR_g13_anti2R	GATCATGTAACACTTGGGCA	
qPCR_g13_anti3F	CATTCCAGTACCGTCACG	Quantitative RT-PCR to determine quantity of Tgl1 gene expression level
qPCR_g13_anti3R	GACTGCCCGATAAGACGT	
qPCR_g10_anti2F	ATTGTTACCAAAGCACG	
qPCR_g10_anti2R	GATCCGGTAAGACTTGTGC	
qPCR_g10_anti3F	CCATAGTACAGTCACGACCTC	
qPCR_g10_anti3R	GACTTCCCGACAAAACGT	

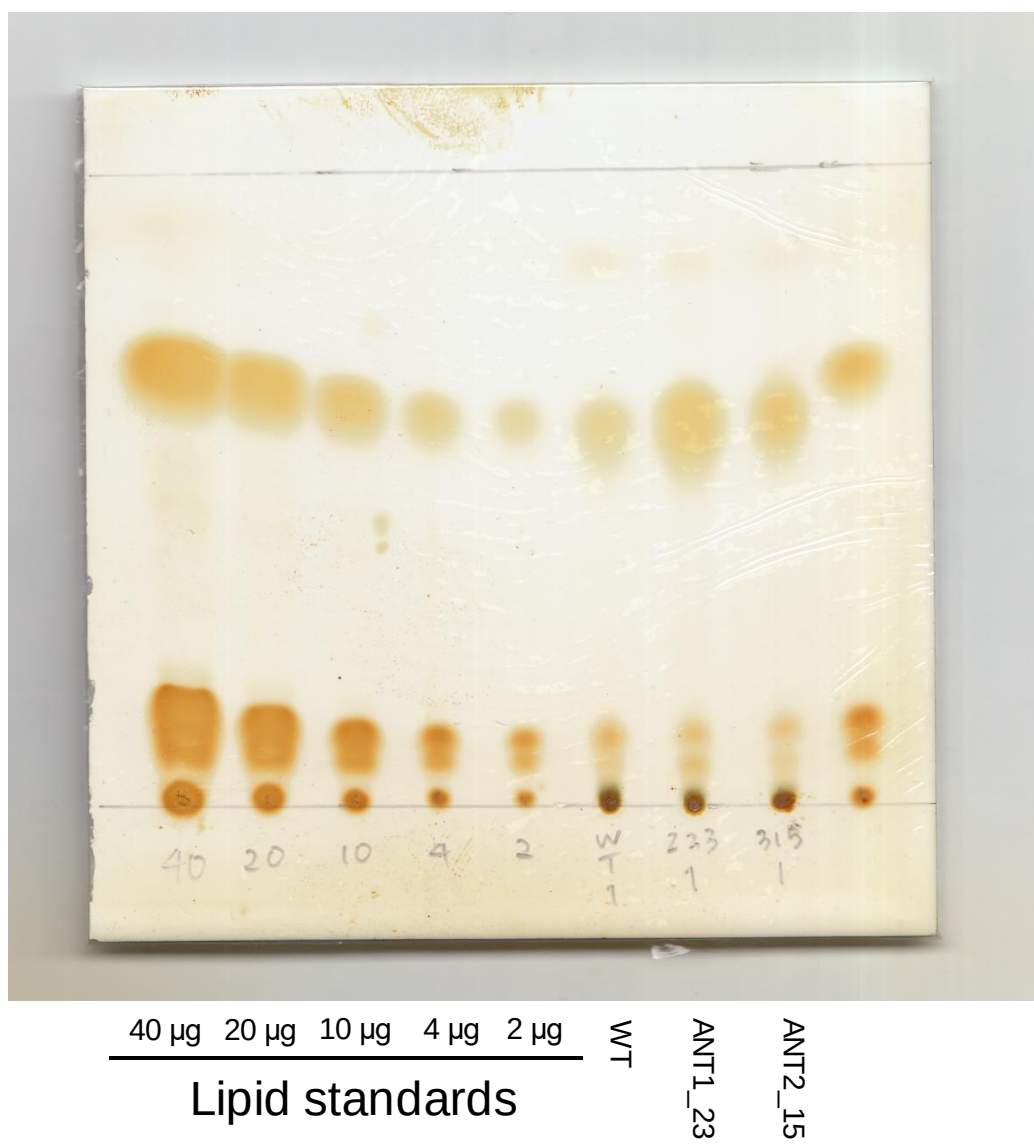


Fig. S3 Uncropped and non-inverted full-length image of the TLC plate shown in Figure 3 in the main text.