

Supporting information

Figure S1. Schematic illustration of the mutagenesis and screening strategy used to isolate antenna mutants.

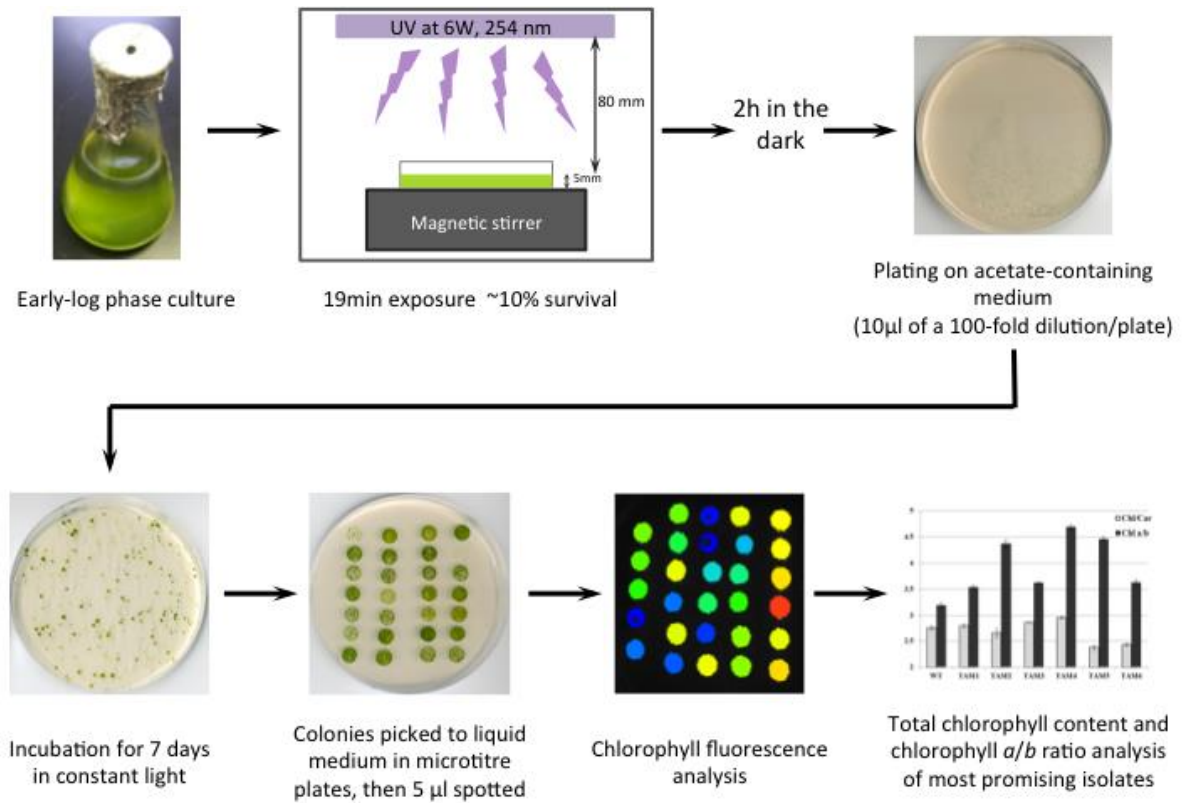


Figure S2. Functional antenna size of PSII and PSI measured in wild-type and TAM mutants

(A) Variable Chl fluorescence was induced with a green light of $15 \mu\text{mol photons m}^{-2} \text{s}^{-1}$, on dark-adapted cells ($\sim 1.0 \cdot 10^7$ cells/ml) in BG-11 medium supplemented with $50 \mu\text{M}$ DCMU. The reciprocal of time corresponding to two-thirds of the fluorescence rise ($T_{2/3}$) was taken as a measure of the PSII functional antenna size. (B) The kinetics of P700 oxidation (ΔAbs at 705 nm) were measured on thylakoids suspension ($75 \mu\text{g Chl/ml}$) treated with $50 \mu\text{M}$ DBMIB and 1 mM methylviologen, upon illumination with a 10 s pulse of red actinic light ($\lambda = 630 \text{ nm}$, $560 \mu\text{mol photons m}^{-2} \text{s}^{-1}$). Data are expressed as mean \pm SD, $n = 7$.

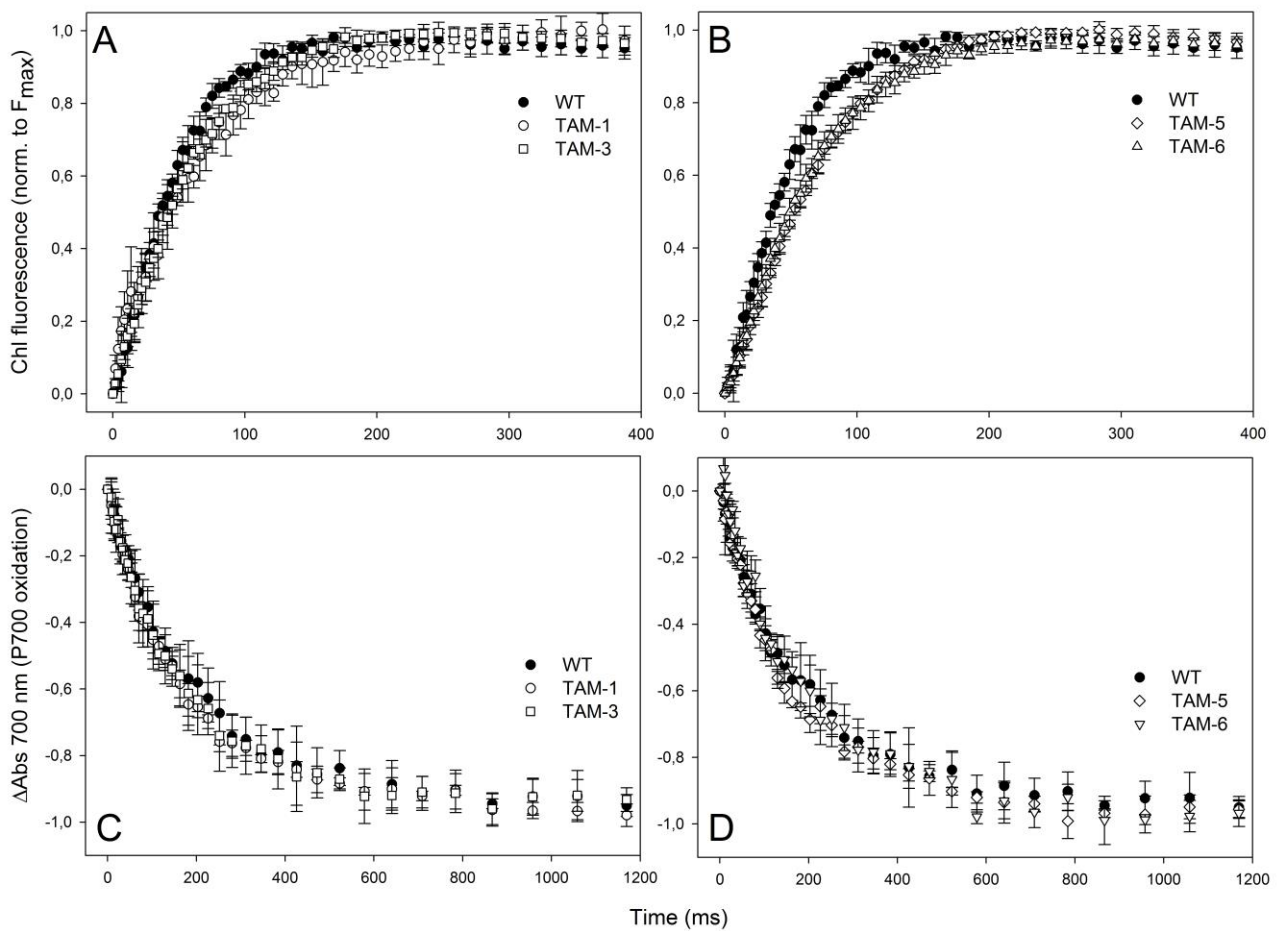


Figure S3. Light-saturation curves of photosynthesis. Curves were obtained with the *C. sorokiniana* wild-type and TAM mutants. Data are expressed as mean \pm SD, n = 4.

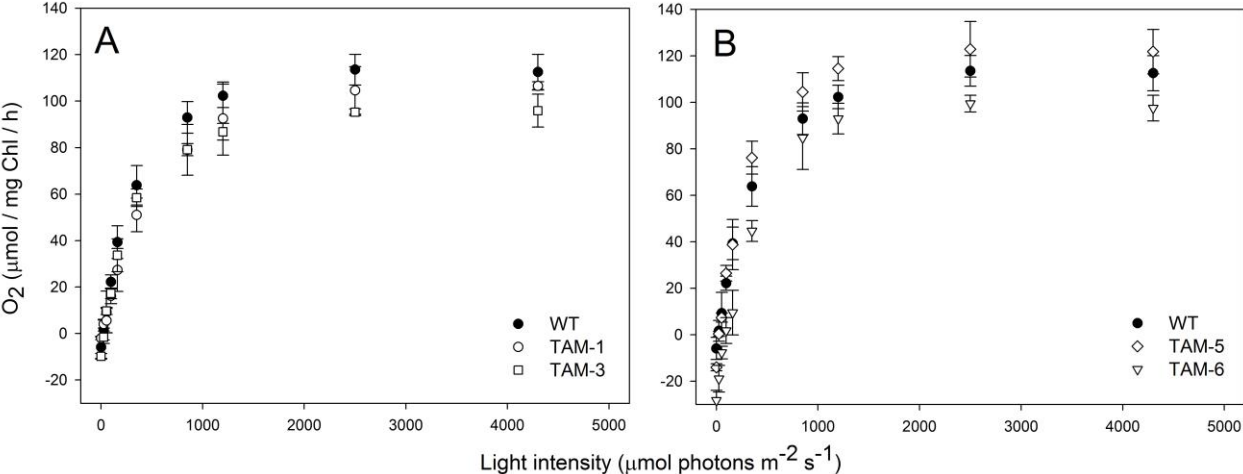


Figure S4. Kinetics of formation and relaxation of photoprotective energy dissipation in wild-type and TAM mutants. NPQ kinetics were measured on wild-type, TAM-2 and TAM-4 cells, grown photoautotrophically, upon illumination with $500 \mu\text{mol photons m}^{-2} \text{s}^{-1}$ of white actinic light. Symbols and error bars show means \pm standard deviation ($n = 3$).

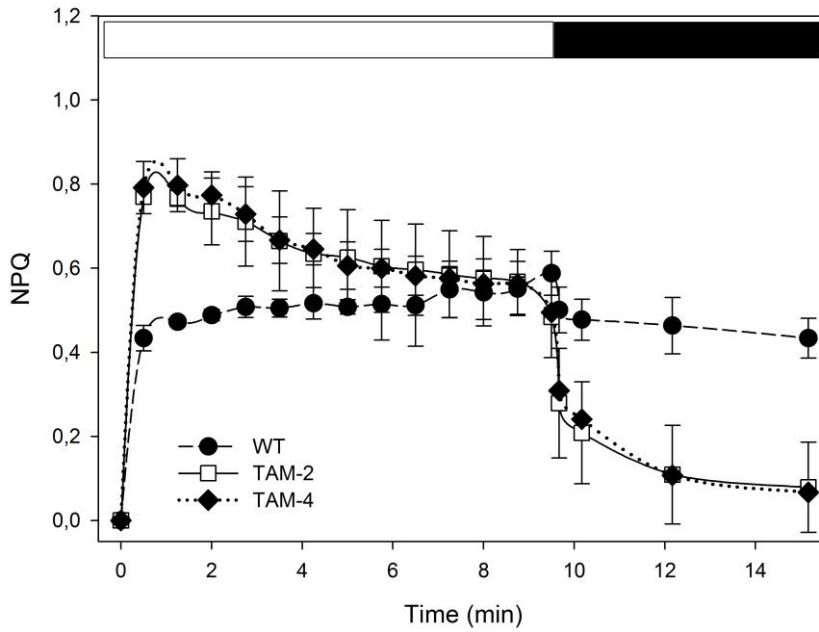


Figure S5

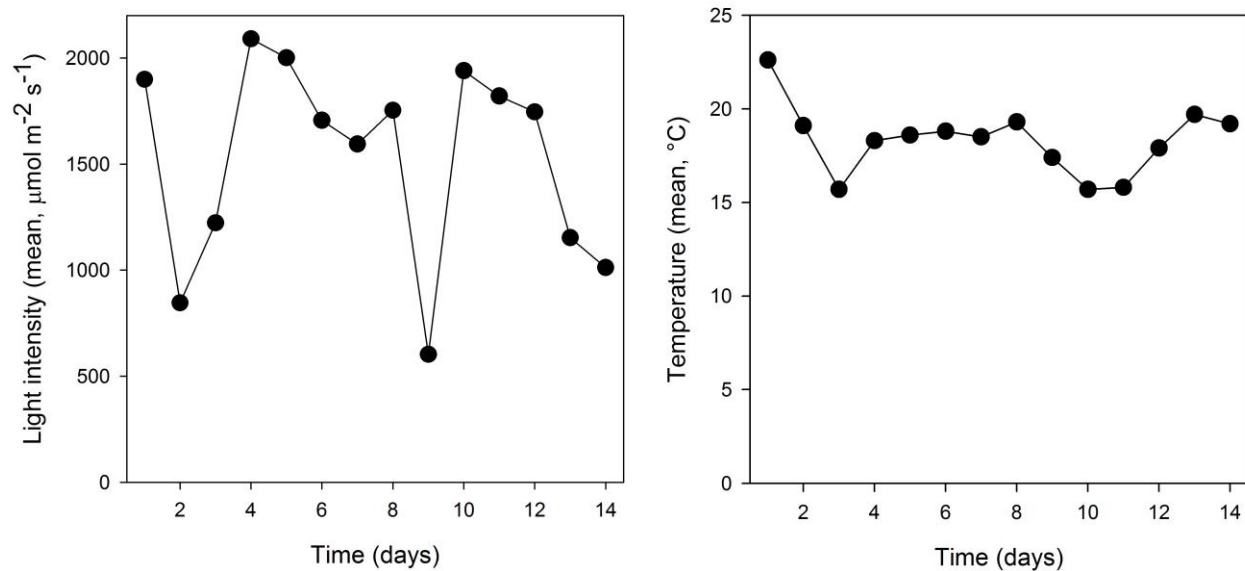


Figure S5. Daily mean irradiance on the reactors' surface (left panel) and mean atmospheric temperature (right panel) measured during the outdoor experiment.

Table S1. Pigment composition of wild-type and TAM mutants. Data are expressed as mean \pm SD. Significantly different values (ANOVA, $P < 0.05$) with respect to the wild-type, within the same column, are marked with different letters.

genotype	Chl / cell (pg)	Chl <i>a</i> / <i>b</i>	Chl / Car	PSII antenna size ($T_{2/3}^{-1} \cdot 10^3, ms^{-1}$)
WT	0.49 \pm 0.07 ^a	2.62 \pm 0.02 ^a	3.43 \pm 0.02 ^a	18.2 \pm 1.4 ^a
TAM-1	0.50 \pm 0.09 ^a	2.92 \pm 0.03 ^b	3.16 \pm 0.02 ^{b,c}	14.5 \pm 1.6 ^b
TAM-2	0.30 \pm 0.02 ^b	3.36 \pm 0.02 ^c	3.07 \pm 0.02 ^b	10.5 \pm 0.5 ^c
TAM-3	0.58 \pm 0.09 ^a	2.69 \pm 0.12 ^d	3.47 \pm 0.12 ^a	15.5 \pm 0.5 ^b
TAM-4	0.34 \pm 0.05 ^b	3.41 \pm 0.03 ^c	3.18 \pm 0.03 ^{b,c}	9.4 \pm 0.5 ^c
TAM-5	0.52 \pm 0.08 ^a	2.84 \pm 0.03 ^b	3.30 \pm 0.04 ^{a,c}	12.9 \pm 0.5 ^b
TAM-6	0.51 \pm 0.07 ^a	2.91 \pm 0.02 ^b	3.35 \pm 0.02 ^a	12.9 \pm 0.8 ^b

Table S2. HPLC analysis of carotenoid composition of wild-type and TAM mutants. Cells were dark-adapted before pigment extraction in DMFA. Data are expressed as mean \pm SD and normalized to 100 Chls. Significantly different values (ANOVA, $P < 0.05$) with respect to the wild-type, within the same column, are marked with different letters.

genotype	mol pigment / 100 mol Chls						
	neoxanthin	violaxanthin	antheraxanthin	lutein	zeaxanthin	α -carotene	β -carotene
WT	6.6 \pm 0.2 ^a	1.5 \pm 0.1 ^a	0.6 \pm 0.1 ^a	17.3 \pm 0.3 ^a	0.8 \pm 0.1 ^a	0.5 \pm 0.2 ^a	2.1 \pm 0.2 ^a
TAM-1	6.2 \pm 0.2 ^a	2.0 \pm 0.1 ^a	0.5 \pm 0.1 ^a	17.3 \pm 0.1 ^a	1.1 \pm 0.1 ^a	0.8 \pm 0.2 ^a	2.1 \pm 0.1 ^a
TAM-2	5.8 \pm 0.2 ^b	5.9 \pm 0.1 ^b	1.9 \pm 0.1 ^b	10.3 \pm 0.2 ^b	2.6 \pm 0.2 ^b	0.2 \pm 0.1 ^a	3.7 \pm 0.3 ^b
TAM-3	6.5 \pm 0.1 ^a	2.1 \pm 0.1 ^a	1.2 \pm 0.1 ^c	14.2 \pm 0.3 ^c	2.2 \pm 0.1 ^b	0.4 \pm 0.2 ^a	2.3 \pm 0.1 ^a
TAM-4	6.0 \pm 0.1 ^b	6.9 \pm 0.1 ^c	1.9 \pm 0.1 ^b	9.6 \pm 0.1 ^d	2.4 \pm 0.2 ^b	0.2 \pm 0.1 ^a	3.9 \pm 0.2 ^b
TAM-5	6.7 \pm 0.3 ^a	3.1 \pm 0.1 ^d	2.8 \pm 0.1 ^d	9.4 \pm 0.2 ^d	3.7 \pm 0.3 ^c	0.2 \pm 0.1 ^a	2.8 \pm 0.2 ^c
TAM-6	6.3 \pm 0.2 ^a	3.6 \pm 0.1 ^d	1.4 \pm 0.1 ^c	13.2 \pm 0.3 ^c	2.1 \pm 0.1 ^b	0.4 \pm 0.1 ^a	2.4 \pm 0.2 ^a

Table S3. Relative abundance of pigment-protein complex in the wild-type and TAM mutants.

Amount of pigment-protein complexes per cell were calculated by densitometric analysis of native PAGE and by Chls content per cell, and expressed as a percentage of the corresponding wild-type values. Data are expressed as means \pm standard deviation (n = 3). Significantly different values (ANOVA, P < 0.05) with respect to the wild-type, within the same column, are marked with different letters.

Genotype	Relative abundance of pigment-protein complexes per cell		
	PSI-LHCI	PSII core	Lhcb
WT	100 \pm 8 ^a	100 \pm 9 ^a	100 \pm 7 ^a
TAM-2	66 \pm 4 ^b	102 \pm 4 ^a	51 \pm 3 ^b
TAM-4	66 \pm 11 ^b	107 \pm 6 ^a	62 \pm 6 ^b