Maximizing Light Utilization Efficiency and Hydrogen Production in Microalgal Cultures

Tasios Melis University of California - Berkeley Thursday, 10 June 2010 Project ID # PD036

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Overview

Timeline

- Start: 01-Dec-2004
- End: 30-Nov-2010
- Completion: 70%

Budget

- Total Project Funding
- DOE: \$1.2 M, UCB: \$450 k
- Funding for FY08

DOE: \$258 k, UCB: \$75 k

- Funding for FY10
- DOE: 0, UCB: \$75 k

Barriers addressed

 Low Light Utilization Efficiency in Photobiological Hydrogen Production due to a Large Photosystem Chlorophyll Antenna Size (Barrier X).



None: Sole
Source Effort



Objectives and Approach

Objective: Minimize the chlorophyll antenna size of photosynthesis to maximize solar conversion efficiency in green algae.

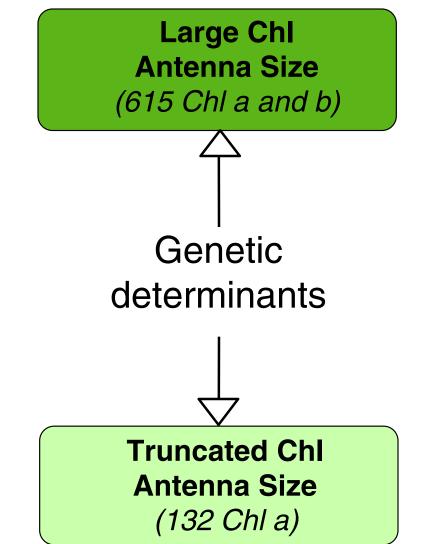
(Identify and characterize genes that regulate the ChI antenna size in the model green alga *ChIamydomonas reinhardtii.* Apply these genes to other green algae, as needed.)

<u>Approach</u>: Interfere with the molecular mechanism for the regulation of the chlorophyll antenna size.

(Employ DNA insertional mutagenesis and highthroughput screening to isolate tagged green algae with a smaller Chl antenna size.)



Regulation of the Chl antenna size

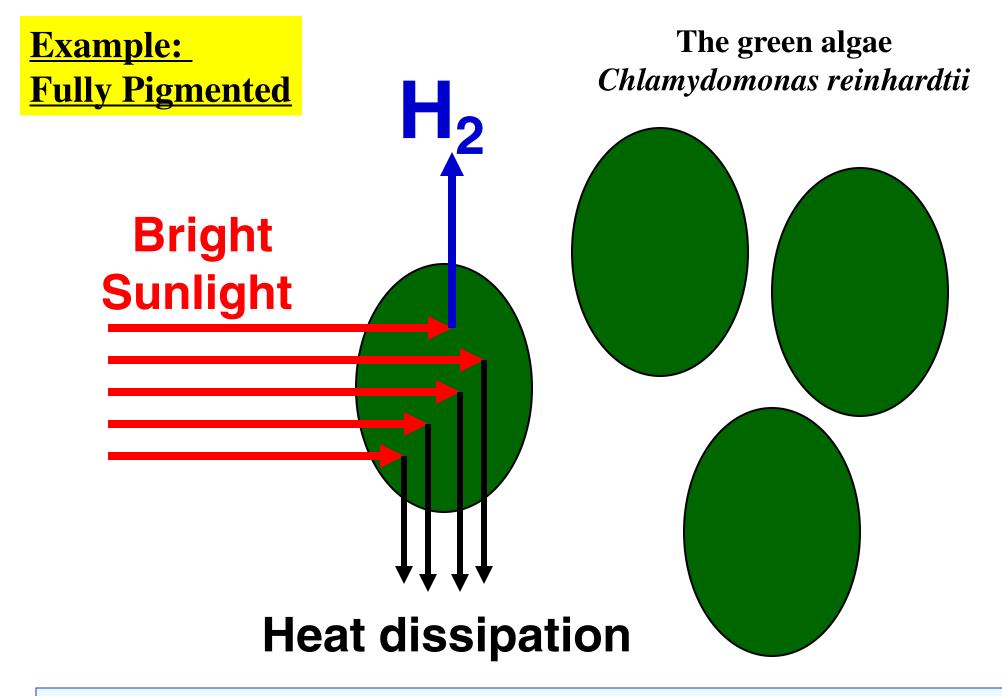


Interference with the genetic mechanism for the regulation of the ChI antenna size, to derive a permanently truncated ChI antenna size, is the goal of this R&D.



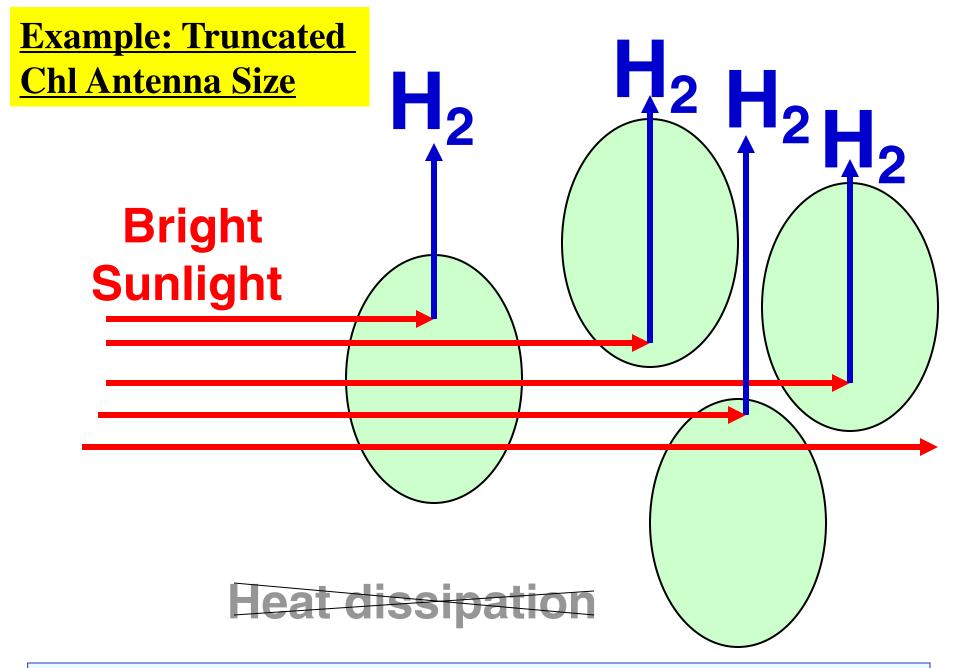
Hydrogen production in a backyard

Chlamydomonas reinhardtii mass culture



Fully pigmented cells over-absorb and wastefully dissipate bright sunlight.





Truncated Chl antenna cells permit greater transmittance of light and overall better solar utilization by the culture.



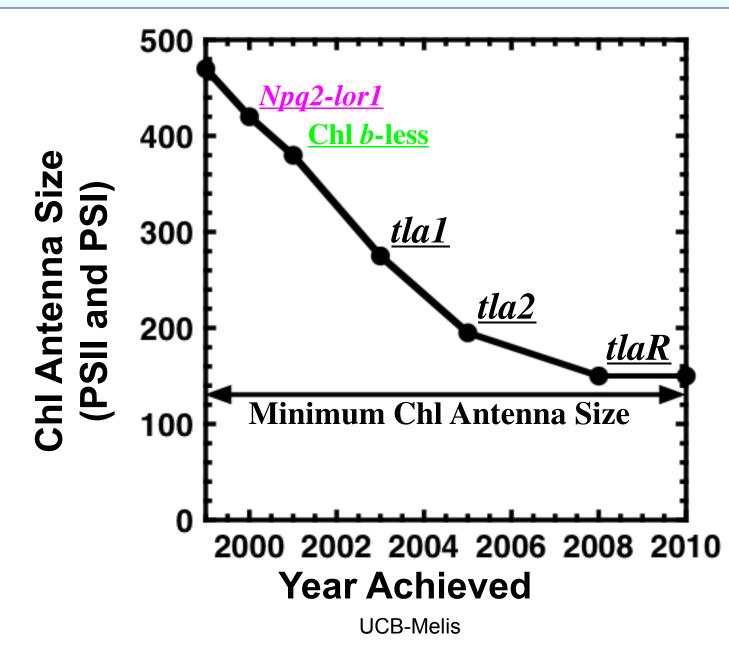
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Technical Barriers and Targets

- <u>Barrier X</u>: Low Light Utilization Efficiency in Photobiological Hydrogen Production due to a Large Photosystem Chlorophyll Antenna Size.
- Light Utilization Efficiency of wild type green microalgae (solar-to-chemical): ~3%
- <u>Theoretical maximum solar-to-chemical efficiency</u>: ~30%
- <u>Target for 2010</u>: Reach a 15% Utilization Efficiency of Absorbed Light Energy.
- <u>Ancillary Objective</u>: Identify and characterize genes that confer the "*tla*" property to microalgae.



Project Timeline Chlorophyll Antenna Size in Chlamydomonas



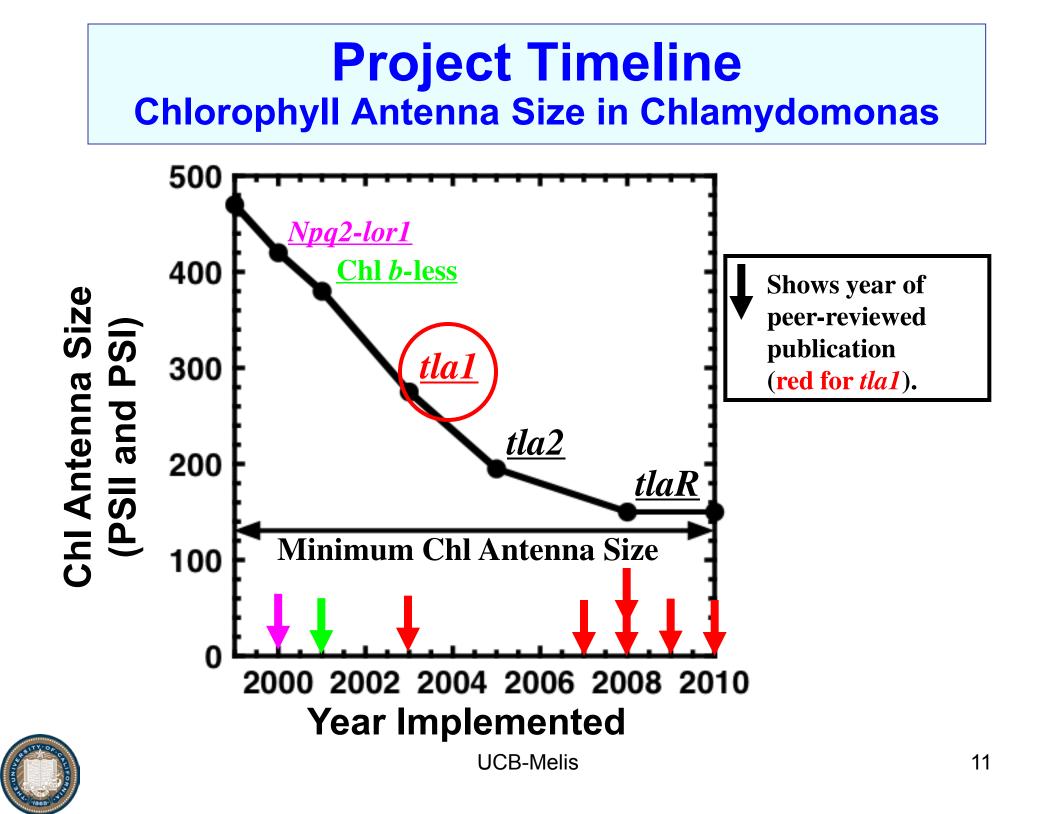


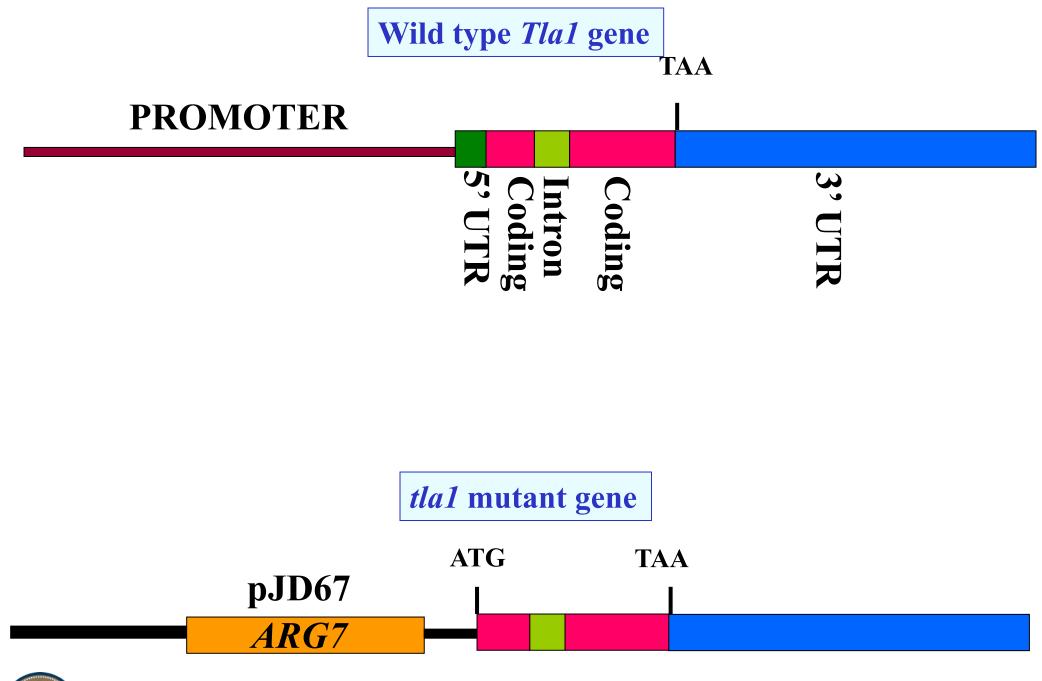
Progress achieved vs the DOE targets

Utilization Efficiency of Incident Solar Light Energy, E₀xE₁, %

	2000	2003	2005	2008	2010	2015
Program Targets	3%	10%			15%	20%
Progress	3%	10% <i>tla1</i>	15% <i>tla2</i>	25% tlaR		



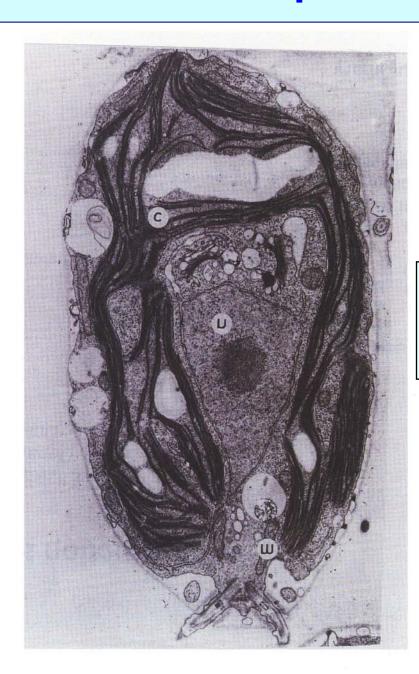






Contraction of the second seco

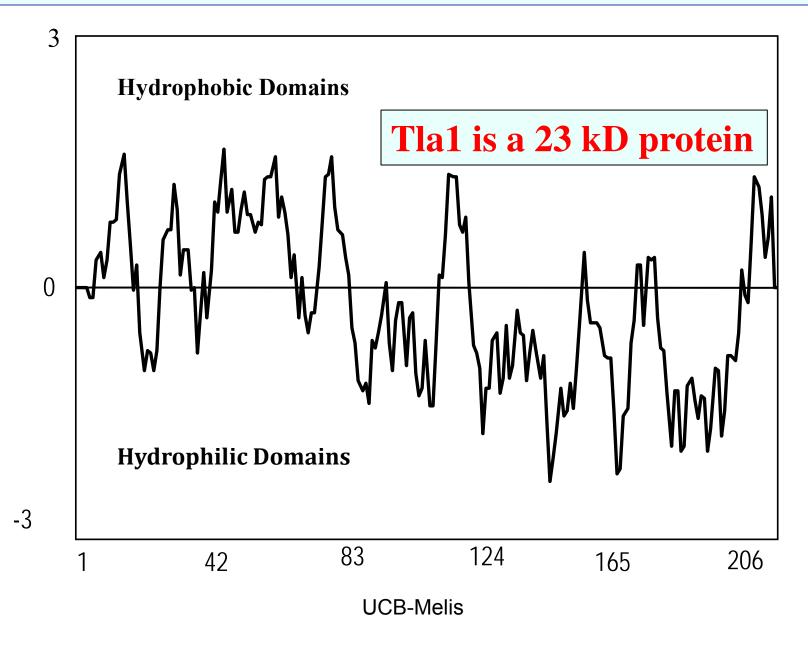
The unicellular green alga Chlamydomonas reinhardti



Environmentally friendly self-repairing and replicating microstructure

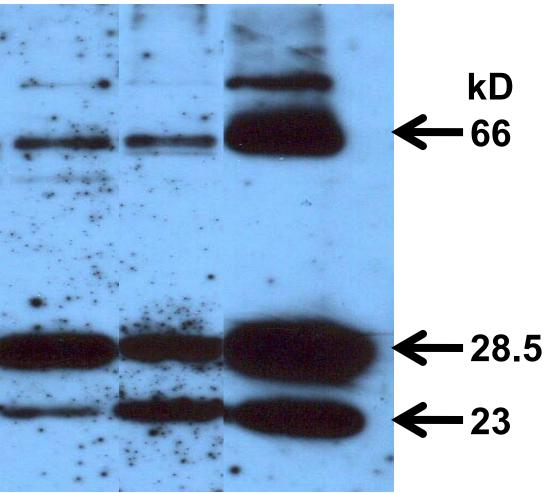
Localization of the Tla1 protein in *C. reinhardtii*

Current Technical Accomplishments Hydropathy plot of the Tla1 protein



• Tla1 polyclonal antibodies recognizing the 23 kD Tla1 protein, also cross reacted with a 28.5 kD protein.

tla1 ΔpetA WT





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Immuno-precipitation and isolation of the cross-reacting 28.5 kD protein, followed by mass spec analysis yielded the peptide sequence <u>R.TWFDDADDWLR.Q</u>, which is specific for the *psbD*/D2 photosystem-II reaction center protein.

1 mtiaigtyqe krtwfddadd wlrqdrfvfv gwsglllfpc ayfalggwlt gttfvtswyt 61 hglatsyleg cnfltaavst pansmahsll fvwgpeaqgd ftrwcqlggl wafvalhgaf 121 gligfmlrqf eiarsvnlrp ynaiafsapi avfvsvfliy plgqsgwffa psfgvaaifr 181 filffqgfhn wtlnpfhmmg vagvlgaall caihgatven tlfedgdgan tfrafnptqa 241 eetysmvtan rfwsqifgva fsnkrwlhff mllvpvtglw msaigvvgla lnlraydfvs 301 qeiraaedpe fetfytknil lnegirawma aqdqpherlv fpeevlprgn al



Current Technical Accomplishments A CLUSTAL 2.0.10 multiple sequence alignment of D2 and Tla1

revealed no similarity between the two proteins

- D2 MTIAIGTYQEKRTWFDDADDWLRQDRFVFVGWSGLLLFPCAYFALGGWLTGTTFVTSWYT 60 Tla1 -----MTFSCSADQT-ALLKILAHAAKYPS 24
 - : *.*: . *. :..:::

D2 HGLATSYLEGCNFLTAAVSTPANSMAHSLLFVWGPEAQGDFTRWCQLGGLWAFVALHGAF 120 Tla1 NSVNGVLVGTAKEGGSVEILDAIPLCHTTLTLAPALEIG----LAQVESYTHITGSV 77

D2 GLIGFMLRQFEIARSVNLRPYNAIAFSAPIAVFVSVFLIYPLGQSGWFFAPSFGVAAIFR 180 Tla1 AIVGYYQSDARFGPGD------ 99 .::*: : .:. . : ***:

D2 FILFFQGFHNWTLNPFHMMGVAGVLGAALLCAIHGATVENTLFEDGDGANTFRAFNPTQA 240

Tla1 -----KRLEQFCKAQA 130 * : * * . :::: : : * * : * *

D2 EETYSMVTANRFWSQIFGVAFSNKRWLHFFMLLVPVTGLWMSAIGVVGLALNLRAYDFVS 300

- Tla1 DNPFELFSKD-----GSKGWKR----ASADGGELALKNADWKKLRE 167
- D2 QEIRAAEDPEFETFYTKNILLNEGIRAWMAAQDQPHERLVFPEEVLPRGNAL 352
- Tla1 EFFVMFKQLKHRTLHDFEEHLDDAGKDWLNKGFASSVKFLLPGNAL----- 213



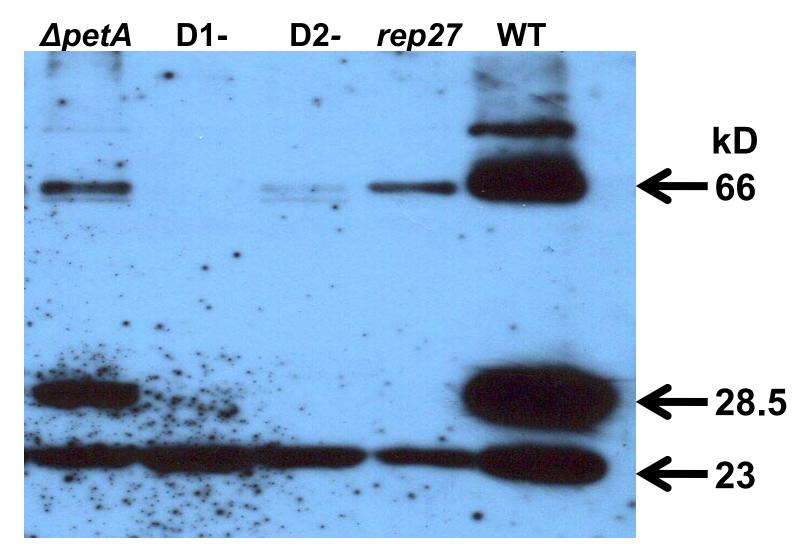
A CLUSTAL 2.0.10 partial sequence alignment of the C-termini of D2 and Tla1 revealed essential identity among 9 consecutive amino acids

D2	V-FPEEVLPRGNAL	13
Tlal	VKFLLP-GNAL	10
	* * • ** ***	

These results suggested that 9 amino acids from the C-terminus form a common epitope and serve as common antigenic determinants.



 The 28.5 kD protein cross reaction is absent in the D1-less, D2-less, and *rep27* mutants of *Chlamydomonas*.





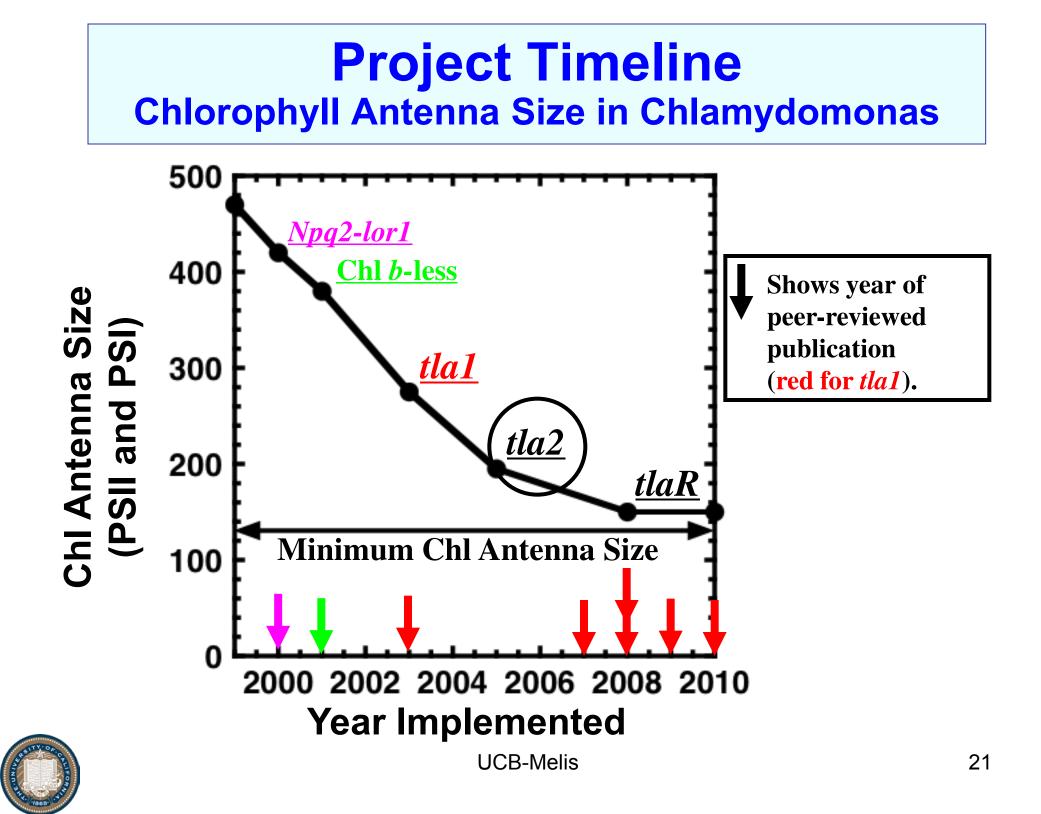


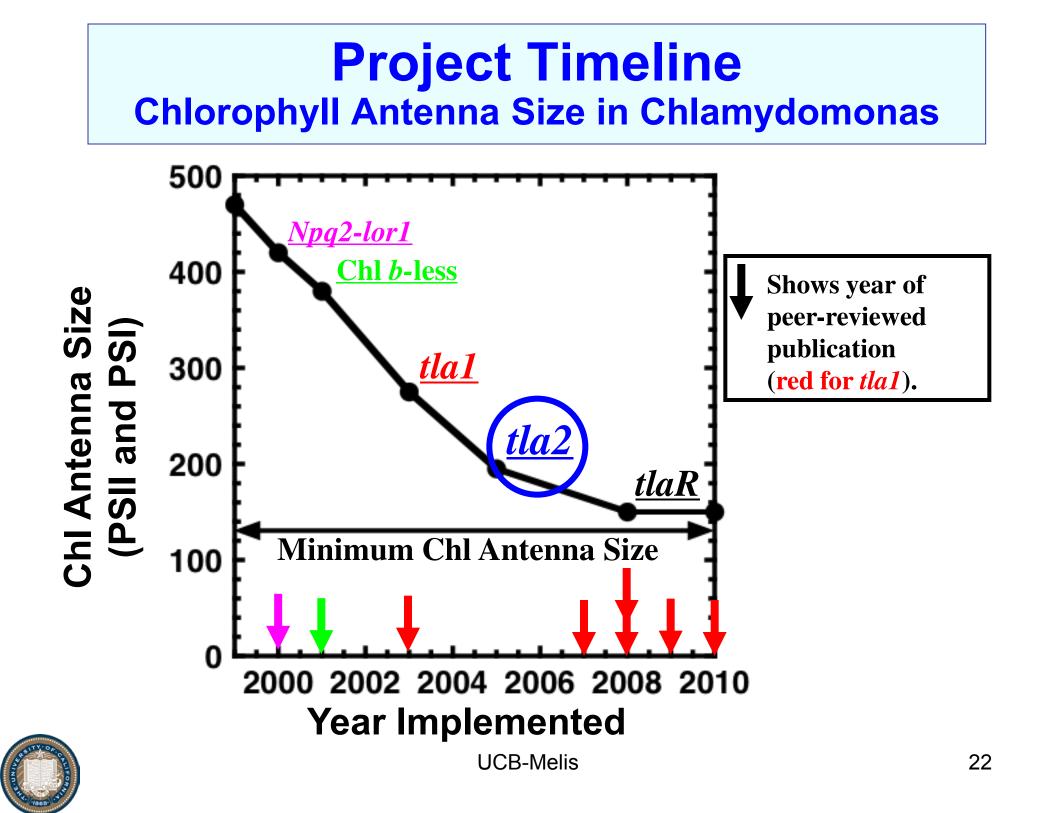
• Resolution:

The Tla1 and D2 proteins have a common 9 amino acid epitope in their C-terminus, that is antigenic enough to generate a strong antibody response against either protein.

This unexpected property has complicated the analysis of the Tla1 function, but it is now solved.

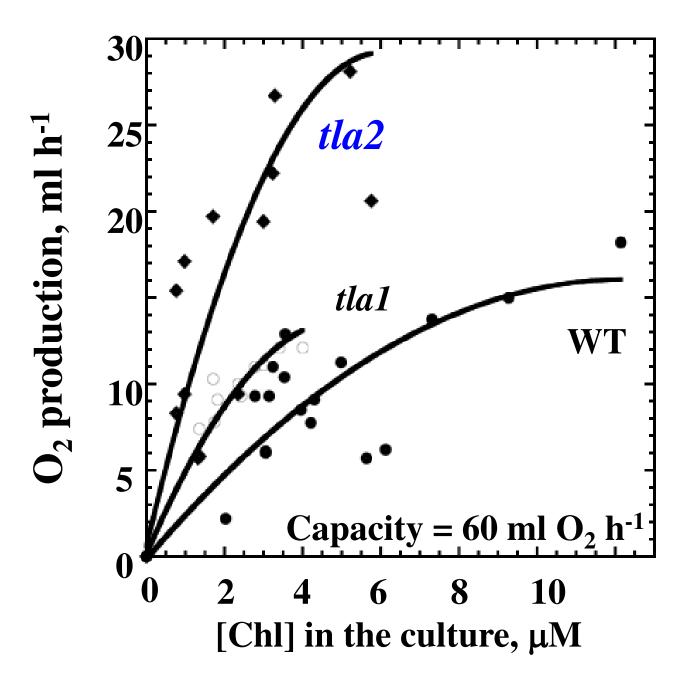






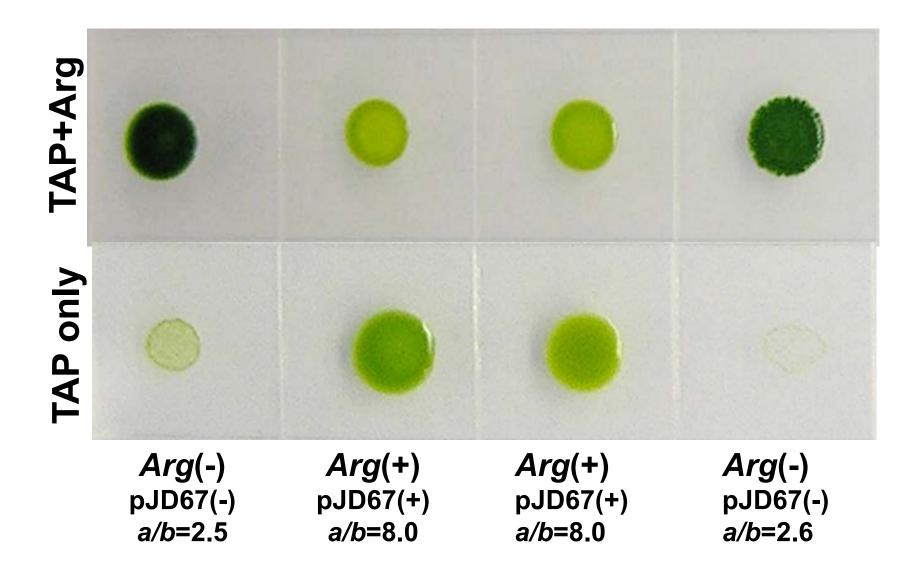
Productivity in Scale-up of Cultures

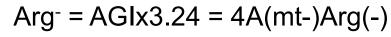
(tla2 outperforms both wild type and tla1 strains)





"Tetrad Analysis" of Progeny from a Single *tla2xArg(-)* Zygospore

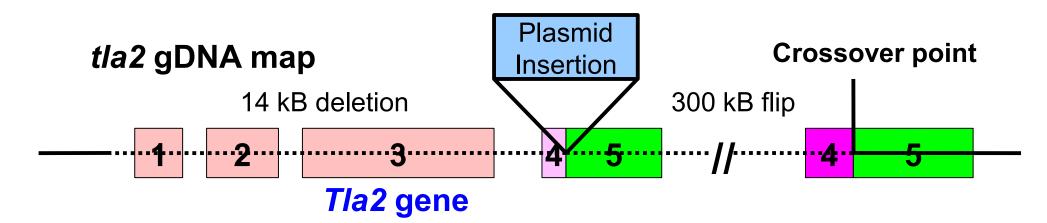


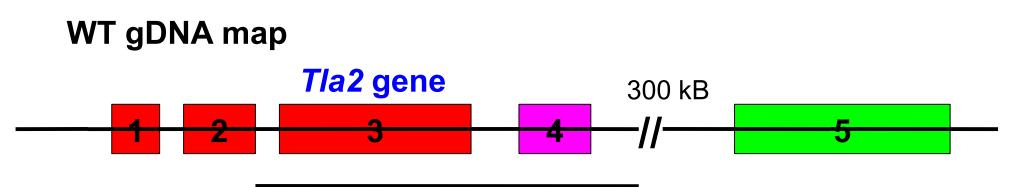




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tla2 gDNA Map and Complementation





BAC clone



Summary of tla2 Mutant Properties

- *Tla2 gene* knock-out.
- pJD67 plasmid Ori and Amp regions deleted.
- Putative *Tla2 gene* is ~6 kb in size.
- *Tla2* encodes a putative protein of ~36 kD.
- *Tla2* function is currently under investigation.



Summary of Accomplishments Analysis of the Tla1 protein and *tla2* mutant strain

- Sorted-put a nagging but unexpected antigenic complication affecting the analysis of the Tla1 protein.
- Cloned the gene conferring a truncated Chl antenna size to the *tla2* mutant.



Significance of Work

- First-time identification and documentation of two different genes (*Tla1 and Tla2*) that regulate the chlorophyll antenna size in photosynthesis.
- Findings could be applied in mass culture to increase solar conversion efficiencies and photobiological hydrogen production.



Current Work

Complete the cellular localization of the Tla1 protein.

Complete the genetic and molecular analysis of the *Tla2* gene; publish results.

Elucidate biochemical function of the *Tla2* gene.



Future Work

Continue work with the cloning of gene(s) conferring the "truncated Chl antenna" phenotype in the *tlaR* strain. (Entails molecular, genetic, biochemical, physiological and scale-up studies.)



Chl Antenna Size vs Light Utilization Efficiency Utilization Efficiency of Absorbed Light Energy

- Wild type antenna size = <u>470 Chl molecules</u> (100%) (PSII=230; PSI=240) Photon use efficiency of WT photosynthesis = ~6-10% <u>Utilization Efficiency of Absorbed Light Energy by WT: ~3-5%</u>
- *tla1* antenna size = <u>275 Chl molecules</u> (59% of control) (PSII=115; PSI=160) Photon use efficiency of *tla1* photosynthesis = ~20% <u>Utilization Efficiency of Absorbed Light Energy by *tla1*: ~10%</u>
- tlaX antenna size = <u>195 Chl molecules</u> (42% of control) (PSII=80; PSI=115) Photon use efficiency of *tlaX* photosynthesis = ~30% <u>Utilization Efficiency of Absorbed Light Energy by *tlaX*: ~15%</u>
- Long-term goal: 132 Chl molecules (28% of control) (PSII=37; PSI=95)
 Photon use efficiency of photosynthesis goal = ~60%
 <u>Utilization Efficiency of Absorbed Light Energy goal: ~30%</u>

