

RNAi pathway components and function in *Paramecium bursaria*

Finlay Maguire

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Overview

Motivation

RNAi in ciliates

Experimental RNAi induction in *P. bursaria*

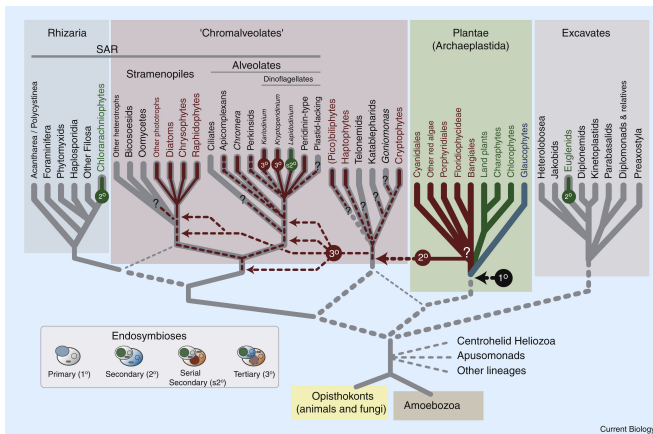
RNAi pathway components in active *P. bursaria* transcriptome(s)

In-silico analysis of potential endosymbiont 'cross-talk'

Conclusions

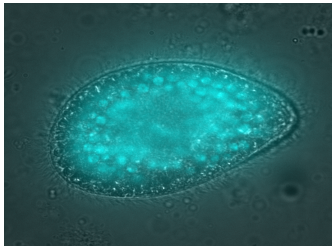
Why is *Paramecium bursaria*
potentially a good model for
(secondary photosynthetic)
endosymbiosis?

Broad diversity of plastid endosymbioses



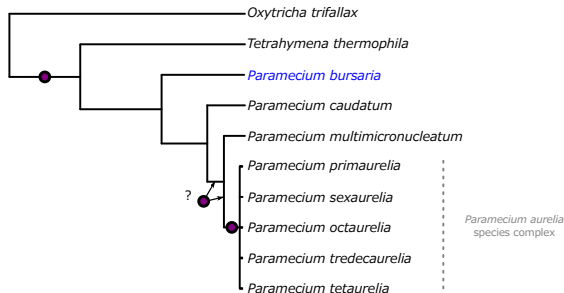
Reproduced from [Arc09].

Paramecium bursaria and its green algal endosymbionts



- ▶ 100 μm to 160 μm serial phagotrophic ciliate (nuclear dimorphism).
- ▶ ~ 300 endosymbiotic algae in stable heritable facultative(?) endosymbiosis.
- ▶ Multiple independent origins of these endosymbioses.
- ▶ Single cell transcriptome and genome of *P. bursaria*-*Micractinium reisseri* SW1-ZK.
- ▶ *P. bursaria* bulk transcriptome Yad1g1N [KSD⁺14].

RNAi pathways in the ciliates



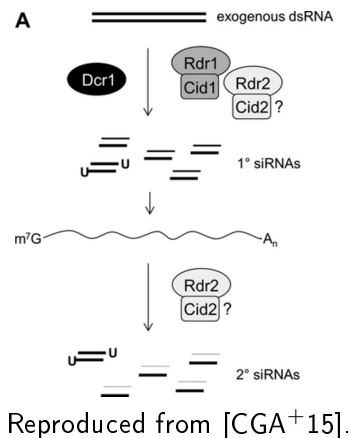
A good model needs a means to test hypotheses:

- ▶ Ciliate specific scnRNA system [MG04, CMM13].
- ▶ siRNA pathways present in *Paramecium tetaurelia* [GS01, GS02] (and *Tetrahymena thermophila* [CL06, YC05]):
 1. Transgene inducible pathway [GS01].
 2. Exogenous dsRNA inducible pathway (feeding or injection) [GS02].

Transgene pathway

- ▶ Microinjection and transformation of MAC with high-copy transgenes lacking 3' UTR [GS01]:
 1. 23nt siRNA generated from transgene transcripts (Dcr1, Rdr2, Rdr3 and Cid2) [LNS⁺09, MCT⁺14].
 2. mRNA cleavage (Ptiwi13 and Ptiwi14).

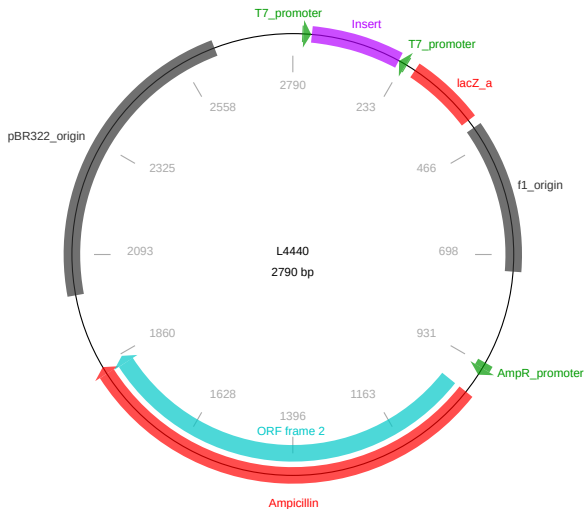
Exogenous dsRNA pathway



- ▶ Exogenous dsRNA via feeding (or microinjection) [GS02].
- ▶ 1° siRNA targeted cleavage (Ptiwi13) [BGK⁺11].
- ▶ Undefined role in MAC for 2° siRNA (Ptiwi12, Ptiwi15) [MCT⁺14, CGA⁺15, BGK⁺11].
- ▶ Pds1 involved in uptake of dsRNA from vacuole? [CGA⁺15].
- ▶ Activated at low levels by ssRNA from normal food bacteria [CGA⁺15].

So, can we experimentally induce
RNAi in *P. bursaria*?

Experimental feeding vector



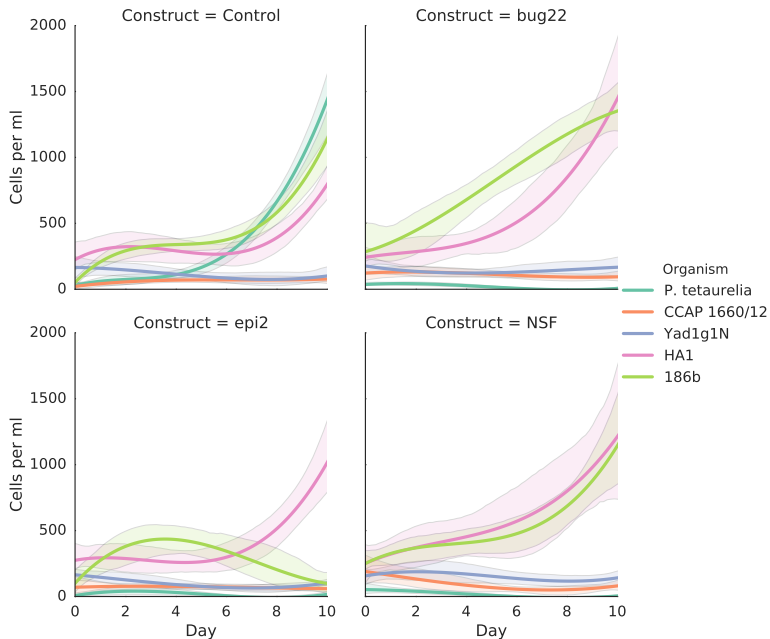
Transformed into *E. coli* with IPTG-inducible T7 polymerase and RNase III deficiency.

Construct inserts

Gene	Function	RNAi phenotype in <i>P. tetraurelia</i>
<i>epi2</i>	Epiplasmin	"Monstrous" cells
NSF	Membrane fusion factor	Lethal
<i>bug22</i>	Basal body/ciliary protein	Slow swimming and death



RNAi feeding had mixed results



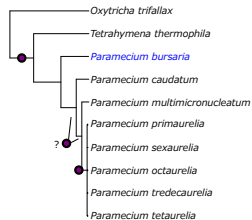
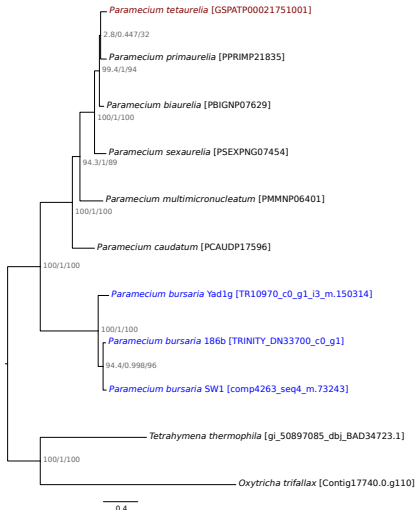
Are all the known RNAi pathway components present in the active transcriptome(s)?

Summary of known RNAi components

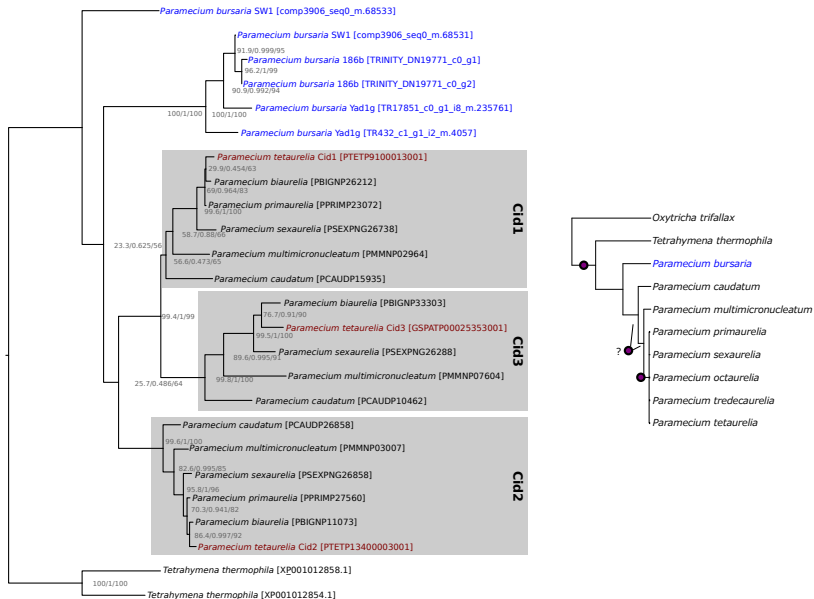
Pathway	Component	Function
transgene-induced siRNA	Rdr3 Ptiwi14	RdRP Piwi
both pathways	Rdr2 Dcr1 Ptiwi13 Cid2	RdRP Dicer Piwi Nucleotidyl transferase
exogenous dsRNA-induced siRNA	Rdr1 Cid1 Ptiwi12 Ptiwi15 Pds1	RdRP Nucleotidyl transferase Piwi Piwi Import of dsRNA?

Component phylogenies are as expected

Dcr1

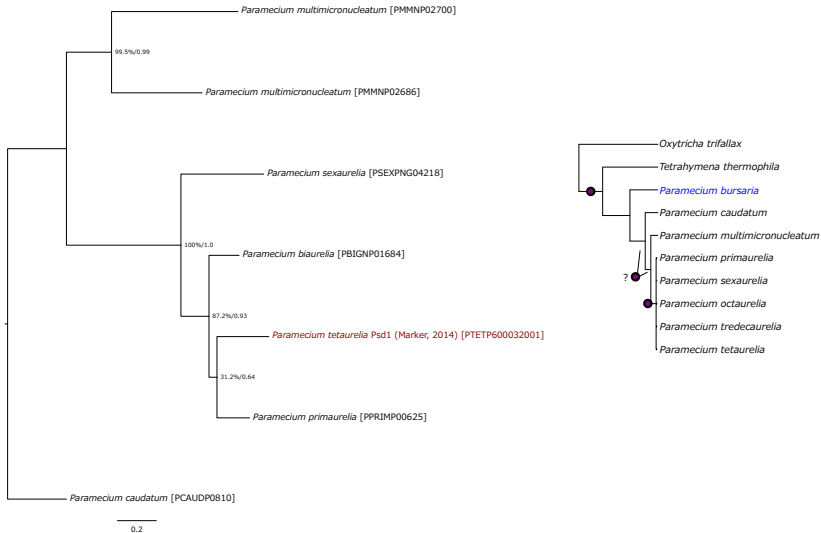


Cid ancestor

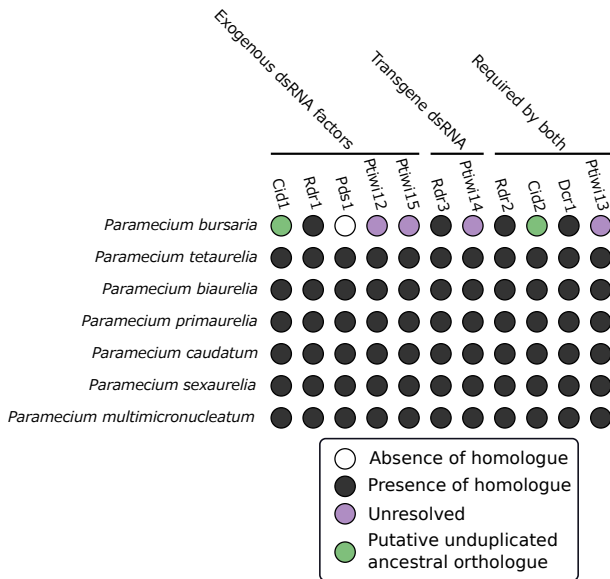


Pds1 absent

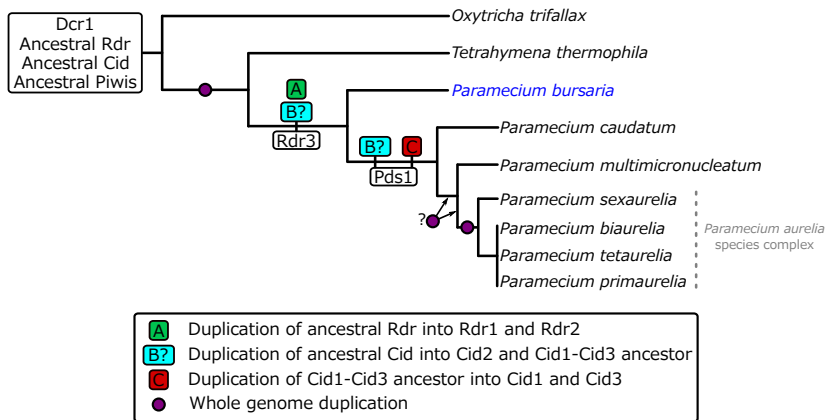
Pds1



Presence/absence of known pathway components

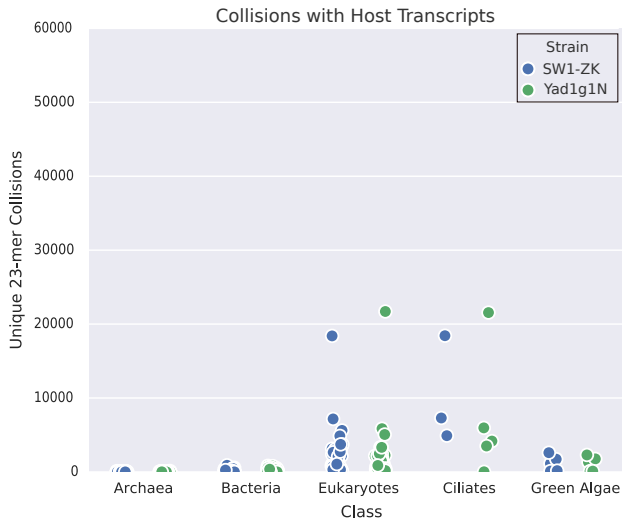


Putative RNAi component evolution scenario

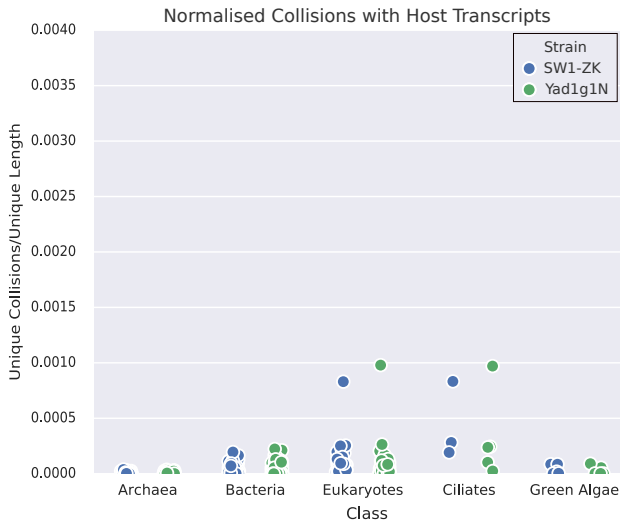


Could having a eukaryotic endosymbiont and RNAi activated by dsRNA in vacuoles be deleterious?

Higher level of collisions with eukaryotes



Collisions are a function of transcriptome size



Conclusions

- ▶ RNAi phenotypes not inducible in most *P. bursaria* strains via feeding.
- ▶ *P. bursaria* lacks Pds1 (in active transcriptome) thus may be unable to take up RNA from digestive vacuoles.
- ▶ High levels of 23-mer collisions between *P. bursaria* and eukaryotic endosymbiont transcriptomes may lead to deactivation of dsRNA uptake from vacuoles.
- ▶ Presence of other factors in active transcriptomes of *P. bursaria* indicate transgene and microinjected exogenous dsRNA pathways may function.

Acknowledgements

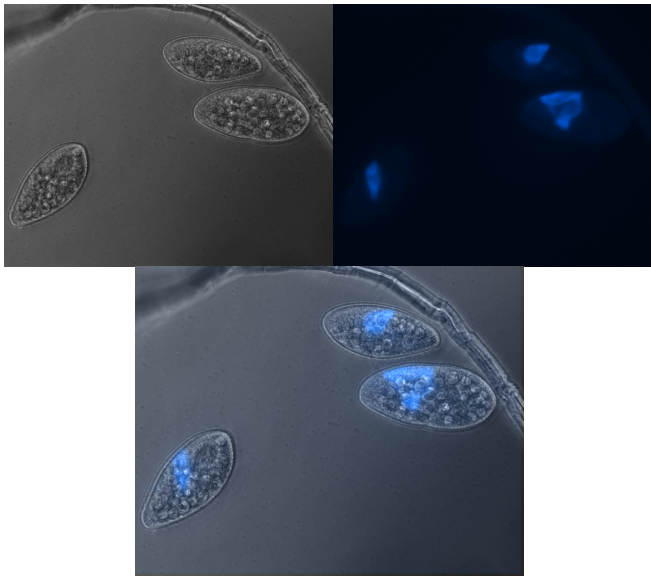
- ▶ Ben Jenkins (feeding experiments)
- ▶ David Milner (labwork)
- ▶ Tom Richards (PI)
- ▶ NHM-UCL PhD Studentship (main funding)



References

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Microinjection proved difficult



Psd1 Structure

