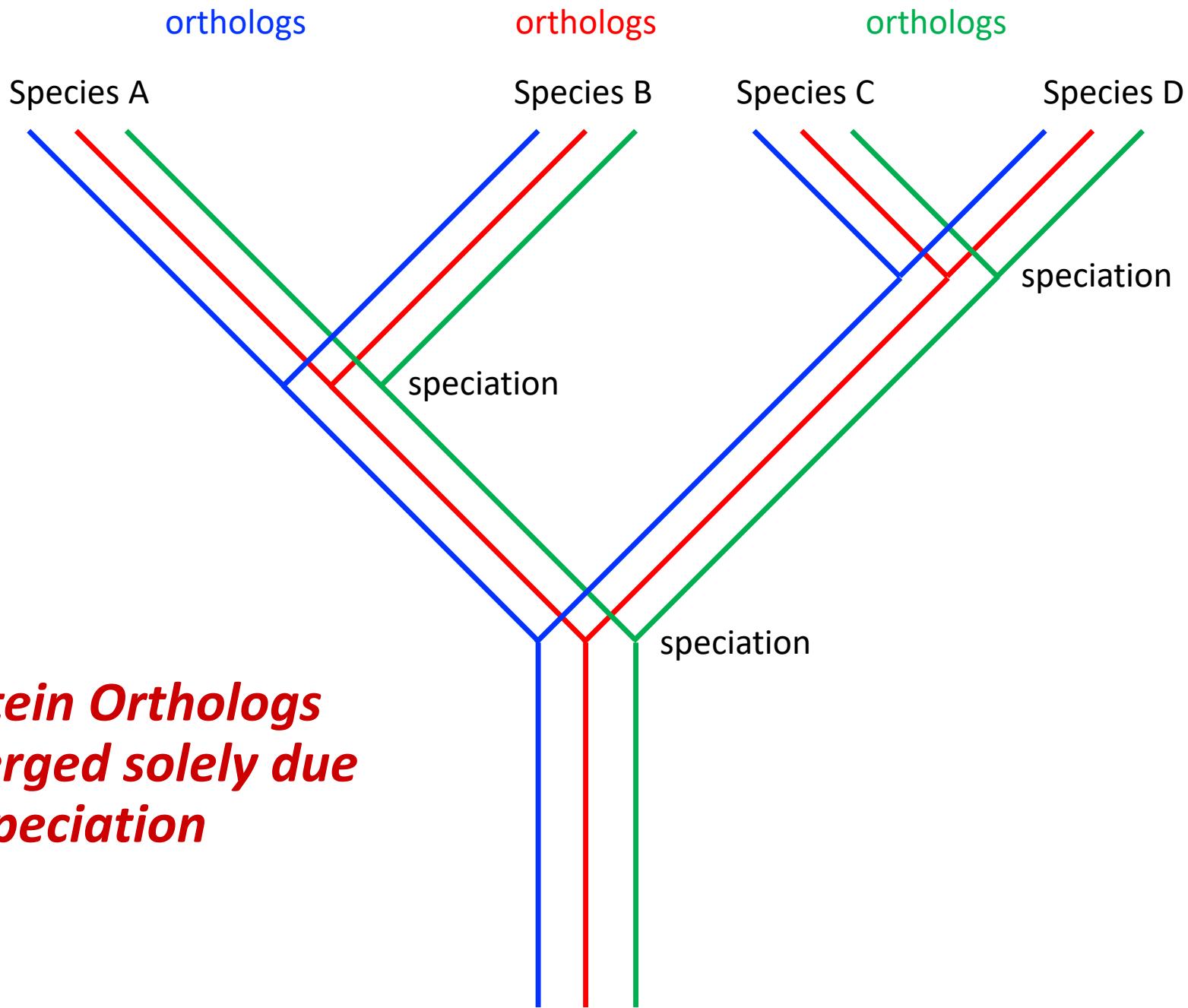
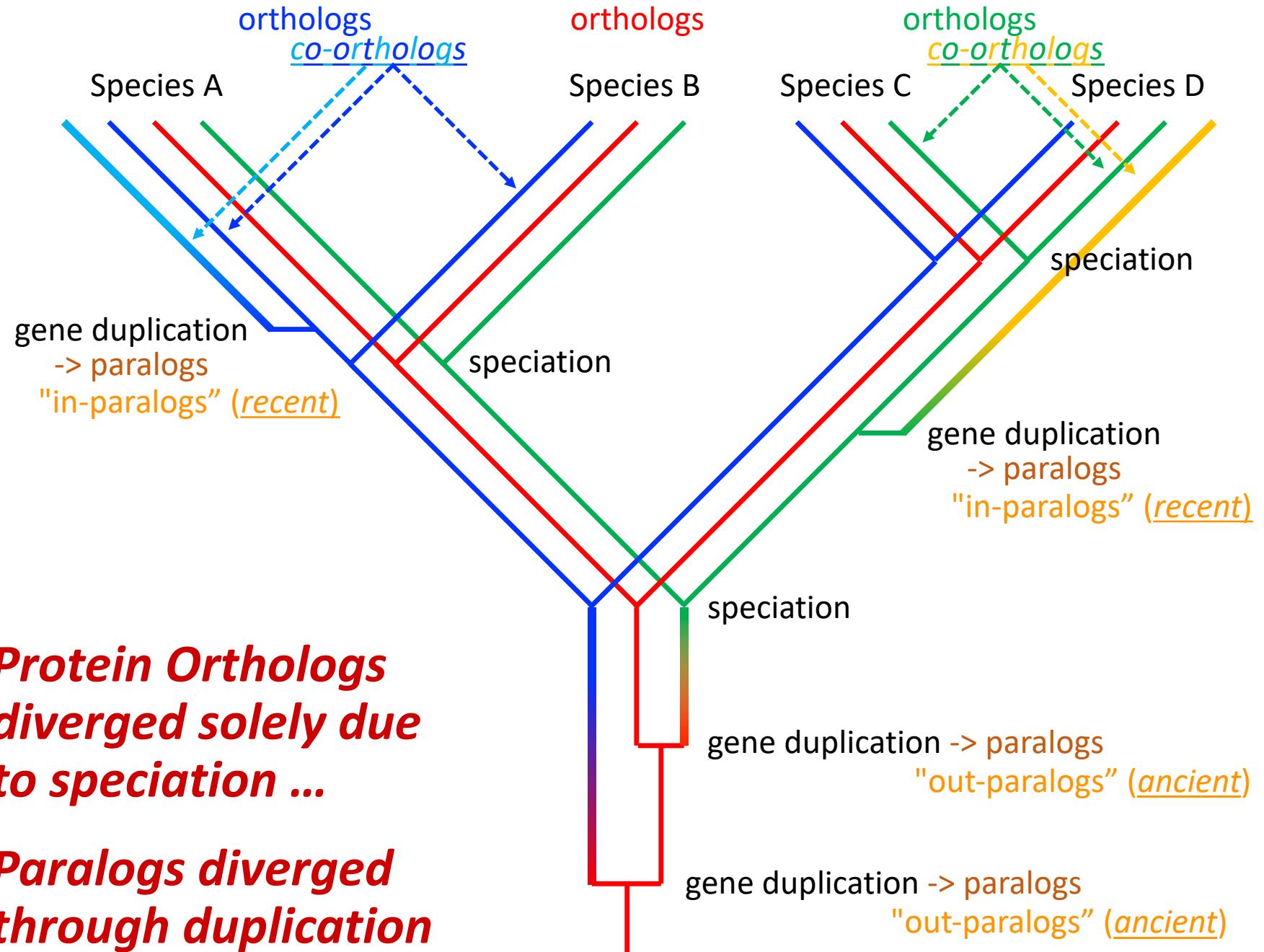


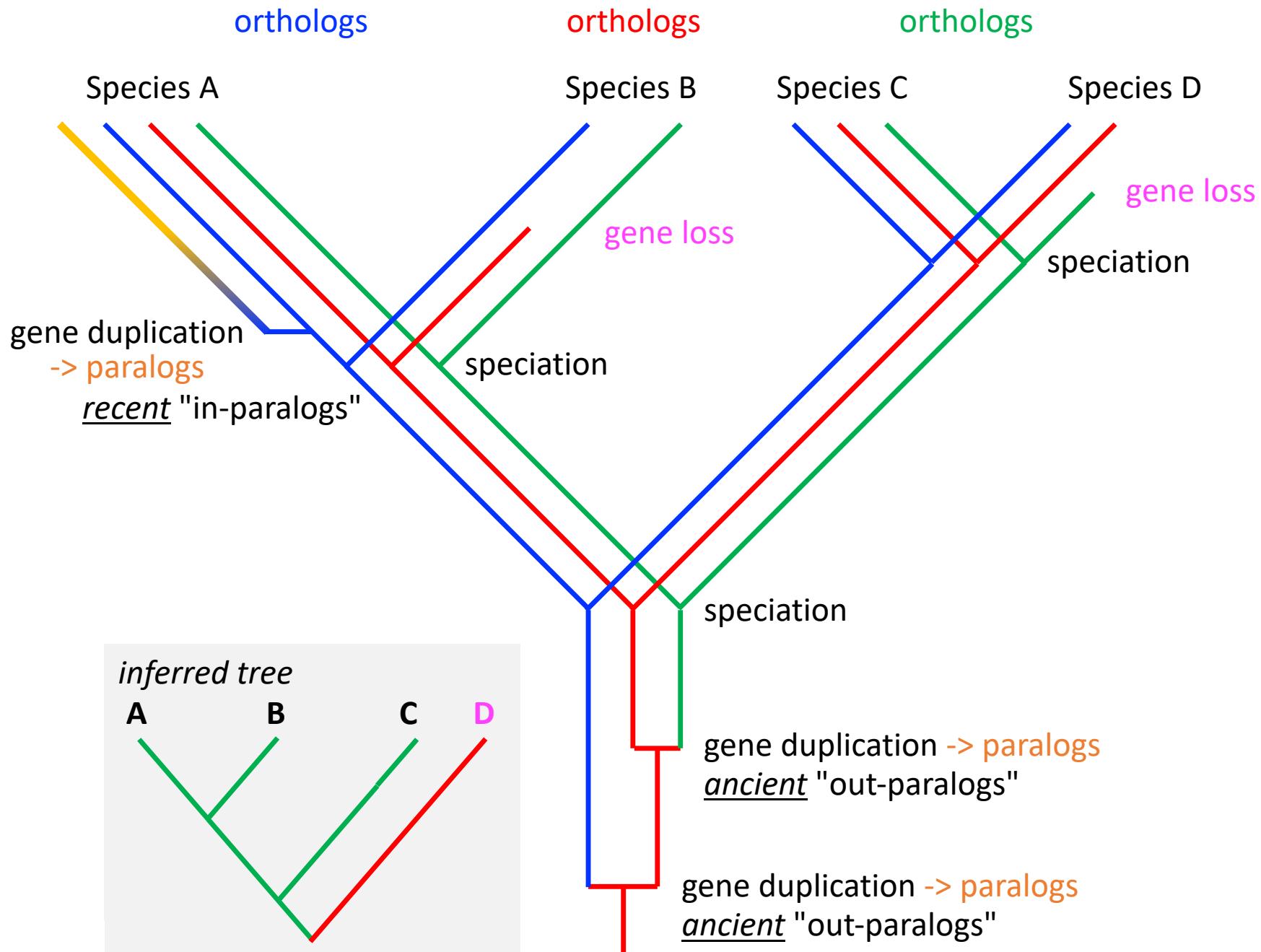
Protein Orthology for Comparative Genomics:

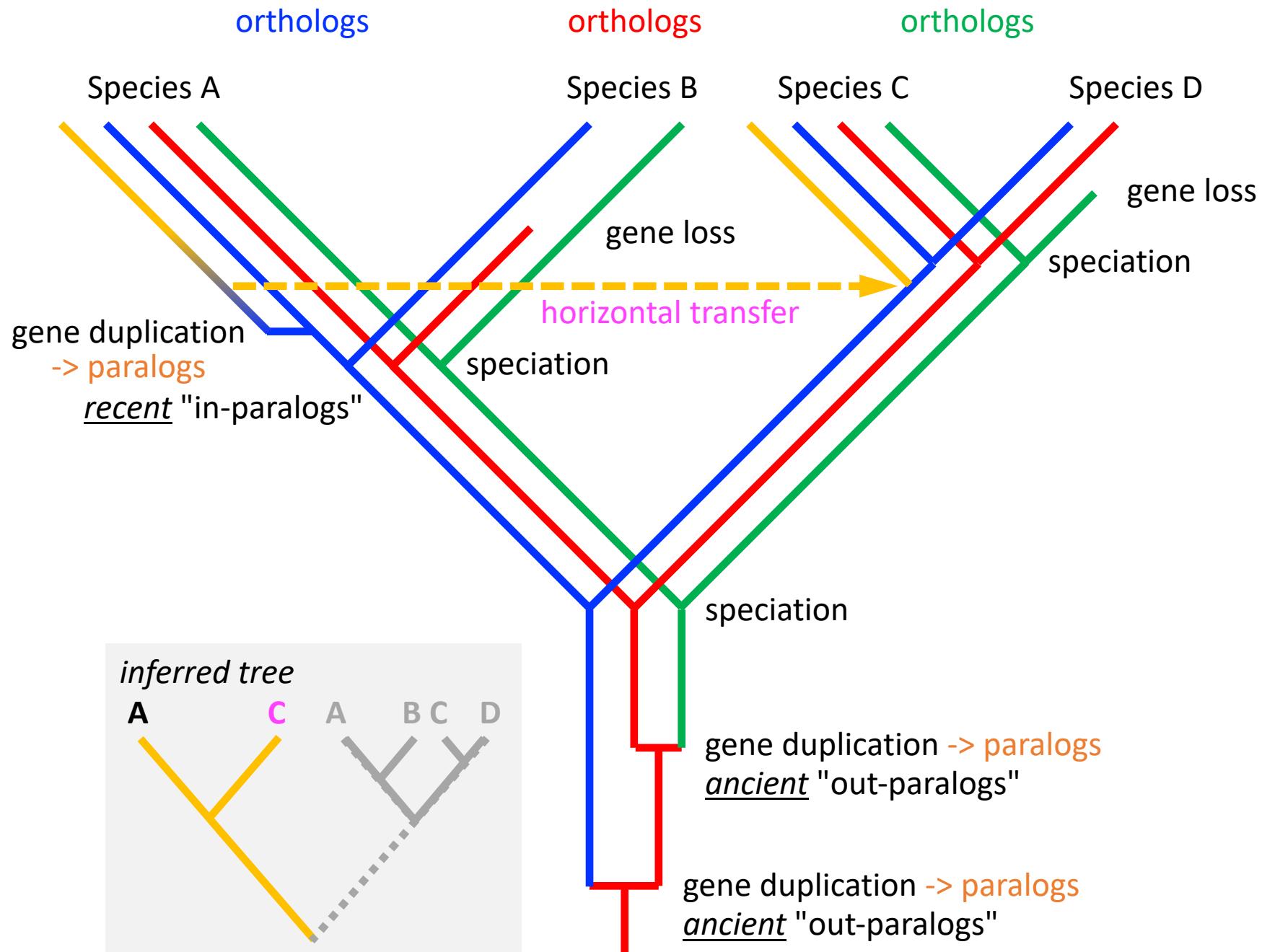
- ❖ ***Genome Comparisons*** (*synteny*)
- ❖ ***Functional Comparisons*** (*particularly for proteins*)
 - *how make reliable inferences ... based on functional data in other species?*
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- ❖ ***Methods based on:***
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 - ***structure*** (*ditto ... and not necessarily = function*)
 - ***expert curation*** (*labor-intensive, especially for eukaryotic taxa*)
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 - ***protein sequence similarity*** – e.g. OrthoMCL, OrthoFinder (*FungiDB, Mycocosm*)
 - ***domain structure*** – e.g. EggNOG, Compara (*Ensembl*)
 - ***phylogeny*** – e.g. OMA, OrthoFinder2
- Note: Benchmarking is difficult without a gold standard (Q4O)!***
- ❖ ***Orthology applications:***
 - ***Validation of genome assemblies & completeness*** (e.g. BUSCO)
 - ***Gene gains, losses, amplification, horizontal transfer*** (*phyletic pattern profiling*)
 - ***Functional inference for understudied taxa (GO terms, etc)***



***Protein Orthologs
diverged solely due
to speciation***







Protein Orthology for Comparative Genomics:

- ❖ ***Genome Comparisons*** (*synteny*)
- ❖ ***Functional Comparisons*** (*particularly for proteins*)
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 - ***Functional inference for understudied taxa (GO terms, etc)***

Genomic-scale identification of orthologous groups

Prokaryotic ortholog groups: COGs

Tatusov et al. *Science* 278:631-7, 1997

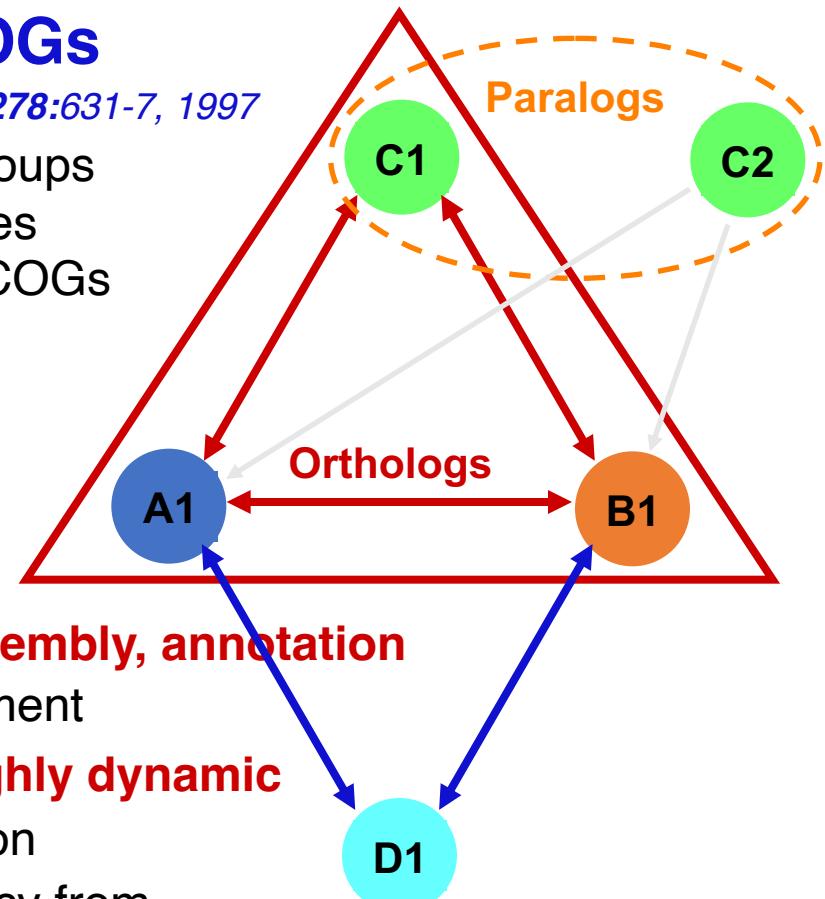
Major resource for prokaryotic orthologous groups

Triangles of reciprocal best hits from 3 lineages

Merge triangles with a common side to form COGs

One-way best hits added as paralogs

Extensive manual curation



Challenges in identifying eukaryotic ortholog groups:

Incomplete (meta)genome sequences, assembly, annotation

- Difficult to ensure correct ortholog assignment

Large genomes, extensive duplication, highly dynamic

- Places a premium on automated annotation
- Difficult to distinguish functional redundancy from functional divergence (in- vs out-paralogs)

Complicated protein domain structure

- Difficult to distinguish proteins with shared domains but distinct functions

Manual curation generally not feasible

Li et al, *Genome Res* 13:2178 (2003)
Chaudhary & Roos, *Nature BT* 23:1089 (2005)

Chen et al, *NAR* 34:363 (2006)

Chen et al, *PLoS1* 2:e383 (2007)

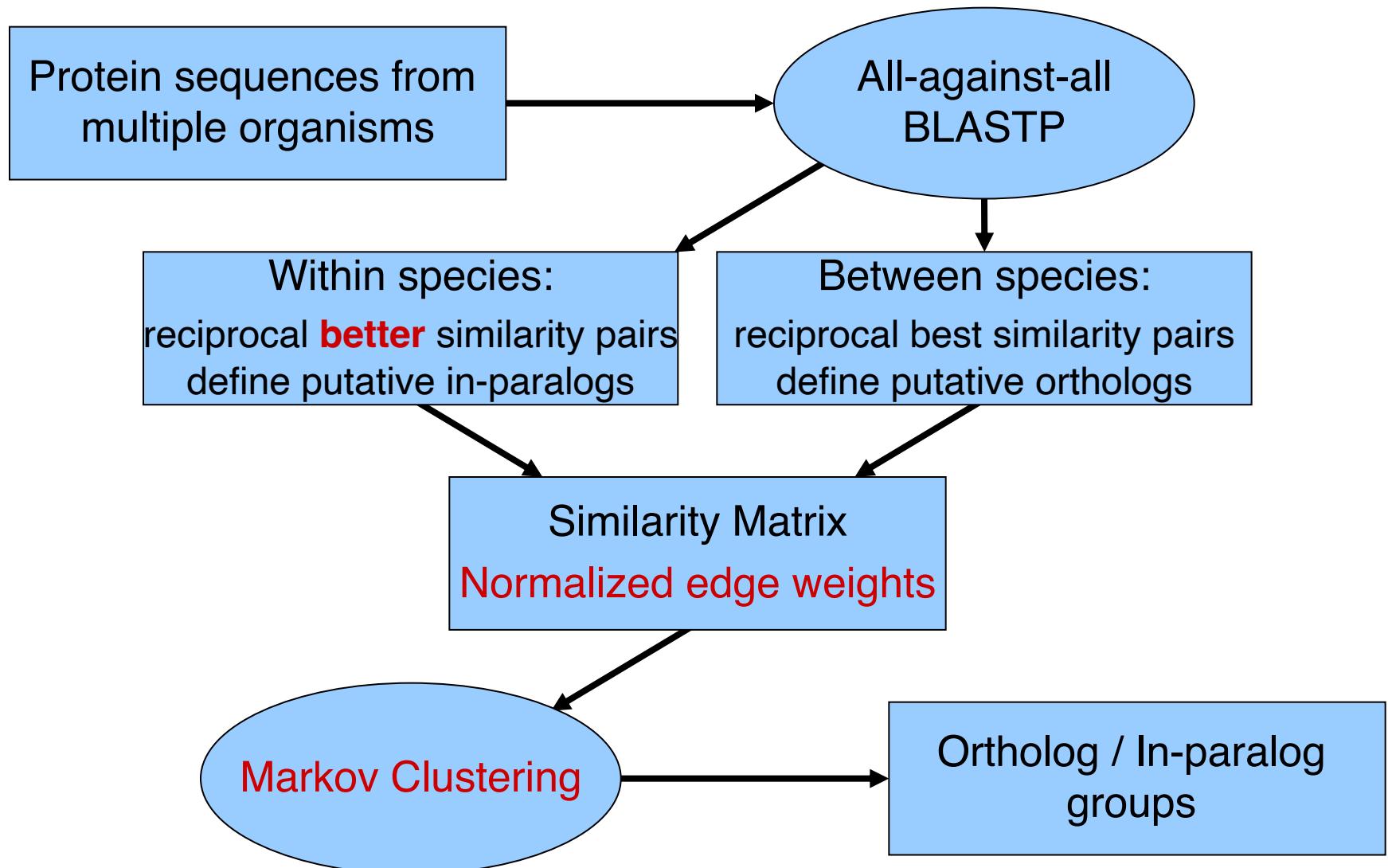
Morrison et al, *Science* 317:1921 (2007)

Peterson et al, *Prot Sci* 18:1306 (2009)

Coyne et al, *Genome Biol* 12:R100 (2011)

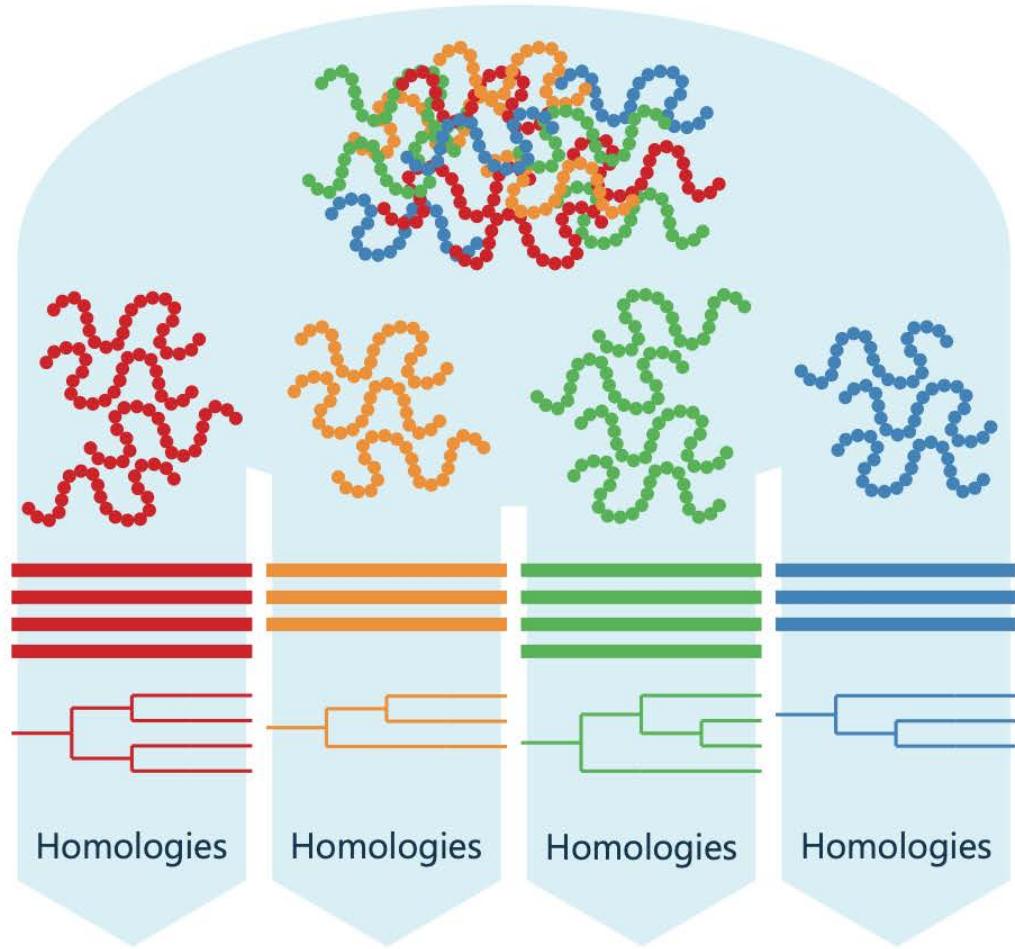
Fischer et all, *Curr Prot Bioinf* 6:1 (2011)

Identification of ortholog groups using OrthoMCL



Gene/protein trees

1. Representative translation of each gene from all species
2. All-vs-all HMM search to classify into families or clustering
3. Multiple protein alignment
4. Phylogenetic tree for each aligned cluster and reconciliation against NCBI taxonomy
5. Ortho-/parologue inference



https://fungi.ensembl.org/info/genome/compara/homology_method.html

Assessing Performance of Orthology Detection Strategies Applied to Eukaryotic Genomes

Feng Chen^{1,2}, Aaron J. Mackey^{2,3*}, Jeroen K. Vermunt⁴, David S. Roos^{2,3}

1 Department of Chemistry, University of Pennsylvania, Philadelphia, Pennsylvania, United States of America, 2 Department of Biology, University of Pennsylvania, Philadelphia, Pennsylvania, United States of America, 3 Genomics Institute, University of Pennsylvania, Philadelphia, Pennsylvania, United States of America, 4 Department of Methodology and Statistics, Tilburg University, The Netherlands

Orthology detection is critically important for accurate functional annotation, and has been widely used to facilitate studies on comparative and evolutionary genomics. Although various methods are now available, there has been no comprehensive analysis of performance, due to the lack of a genomic-scale 'gold standard' orthology dataset. Even in the absence of such datasets, the comparison of results from alternative methodologies contains useful information, as agreement enhances confidence and disagreement indicates possible errors. Latent Class Analysis (LCA) is a statistical technique that can exploit this information to reasonably infer sensitivities and specificities, and is applied here to evaluate the performance of various orthology detection methods on a eukaryotic dataset. Overall, we observe a trade-off between sensitivity and specificity in orthology detection, with BLAST-based methods characterized by high sensitivity, and tree-based methods by high specificity. Two algorithms exhibit the best overall balance, with both sensitivity and specificity >80%: INPARANOID identifies orthologs across two species while OrthoMCL clusters orthologs from multiple species. Among methods that permit clustering of ortholog groups spanning multiple genomes, the (automated) OrthoMCL algorithm exhibits better within-group consistency with respect to protein function and domain architecture than the (manually curated) KOG database, and the homolog clustering algorithm TribeMCL as well. By way of using LCA, we are also able to comprehensively assess similarities and statistical dependence between various strategies, and evaluate the effects of parameter settings on performance. In summary, we present a comprehensive evaluation of orthology detection on a divergent set of eukaryotic genomes, thus providing insights and guides for method selection, tuning and development for different applications. Many biological questions have been addressed by multiple tests yielding binary (yes/no) outcomes but no clear definition of truth, making LCA an attractive approach for computational biology.

Citation: Chen F, Mackey AJ, Vermunt JK, Roos DS (2007) Assessing Performance of Orthology Detection Strategies Applied to Eukaryotic Genomes. PLoS ONE 2(4): e383. doi:10.1371/journal.pone.0000383

INTRODUCTION

The rapid growth in the availability of genome sequence data, from an ever-increasing range of relatively obscure species, places a premium on the automated identification of orthologs to facilitate functional annotation, and studies on comparative and evolutionary genomics. Homologous proteins share a common ancestry, and may be characterized as either orthologs (which evolve by speciation only) or paralogs (which arise by gene duplication) [1,2]. Orthologs typically retain similar domain architecture and occupy the same functional niche following speciation, while (functionally redundant) paralogs are likely to diverge with new functions through point mutations and domain recombinations [3,4].

The concepts of orthology and paralogy are well-established in classical and molecular systematics [1], and have been extended to describe more complicated situations associated with extensive gene duplications commonly observed in eukaryotic species [4–6]. In- and out-paralogs are analogous to the phylogenetic concepts in- and out-groups, denoting genes duplicated subsequent or prior to speciation, respectively. Recent duplications yield in-paralogs that may exhibit a many-to-one or many-to-many ortholog relationship with genes in the other species (termed co-orthologs).

Several strategies have been employed to distinguish probable (co-)orthologs from paralog, as summarized in Table 1: phylogeny-based methods include RIO (Resampled Inference of Orthology) [7] and Orthotrappet/HOPS (Hierarchical grouping of Orthologous and Paralogous Sequences) [8,9]; methods based on evolutionary distance metrics include RSD (Reciprocal Smallest Distance) [10,11]; BLAST-based methods include Reciprocal Best Hit (RBH), COG (Cluster of Orthologous Groups)

[12–15]/KOG (euKaryotic Orthologous Groups) [15], and Inparanoid [5,16]. The problem of orthology detection is particularly acute for eukaryotic genomes, because of their large size, the difficulty of defining accurate gene models, the complexity of protein domain architectures, and rampant gene duplications [3,17]. To address these difficulties we previously developed the OrthoMCL algorithm [18], which improves on RBH by (i) recognizing co-ortholog relationships (Figure 1), (ii) using a normalization step to correct for systematic biases when comparing specific pairs of genomes, and (iii) using a Markov graph clustering (MCL) algorithm [19] to define ortholog groups. OrthoMCL and Inparanoid exhibit similar performance when comparing two species, but the former is extensible to cluster orthologs across

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Competing Interests: The authors have declared that no competing interests exist.

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= Current address: Informatics, GlaxoSmithKline, Collegeville, Pennsylvania, United States of America

ANALYSIS

OPEN

Standardized benchmarking in the quest for orthologs

Adrian M Altenhoff^{1,2}, Brigitte Boeckmann³, Salvador Capella-Gutierrez^{4–6}, Daniel A Dalquen⁷, Todd Deluca⁸, Kristoffer Forslund⁹, Jaime Huerta-Cepas⁹, Benjamin Linard¹⁰, Cécile Pereira^{11,12}, Leszek P Przybecki⁴, Fabian Schreiber¹³, Alan Sousa da Silva¹³, Damian Szklarczyk^{14,15}, Clément-Marie Train¹, Peer Bork^{9,16,17}, Odile Lecompte¹⁸, Christian von Mering^{14,15}, Ioannis Xenarios^{3,19,20}, Kimmien Sjölander²¹, Lars Juhl Jensen²², Maria J Martin¹³, Matthieu Muffato¹³, Quest for Orthologs consortium²³, Toni Gabaldón^{4,5,24}, Suzanna E Lewis²⁵, Paul D Thomas²⁶, Erik Sonnhammer²⁷ & Christophe Dessimoz^{7,20,28–30}

Achieving high accuracy in orthology inference is essential for many comparative, evolutionary and functional genomic analyses, yet the true evolutionary history of genes is generally unknown and orthologs are used for very different applications across phyla, requiring different precision-recall trade-offs. As a result, it is difficult to assess the performance of orthology inference methods. Here, we present a community effort to establish standards and an automated web-based service to facilitate orthology benchmarking. Using this service, we characterize 15 well-established inference methods and resources on a battery of 20 different benchmarks. Standardized benchmarking provides a way for users to identify the most effective methods for the problem at hand, sets a minimum requirement for new tools and resources, and guides the development of more accurate orthology inference methods.

Evolutionarily related genes (homologs) across different species are often divided into gene pairs that originated through speciation events (orthologs) and gene pairs that originated through duplication events (paralogs)¹. This distinction is useful in a broad range of contexts, including phylogenetic tree inference, genome annotation, comparative genomics and gene function prediction^{2–4}. Accordingly, dozens of methods⁵ and resources^{6–8} for orthology inference have been developed.

Because the true evolutionary history of genes is typically unknown, assessing the performance of these orthology inference methods is not straightforward. Several indirect approaches have been proposed. Based on the notion that orthologs tend to be functionally more similar than paralogs (a notion now referred to as the ortholog conjecture^{9–12}), Hulsen et al.¹³ used several measures of functional conservation (conservation levels, protein–protein interactions and protein domain conservation) to benchmark orthology inference methods. Chen et al.¹⁴ proposed an unsupervised learning approach based on consensus among different orthology methods. Altenhoff and Dessimoz¹⁵ introduced a phylogenetic benchmark measuring the concordance between gene trees reconstructed from putative orthologs and undisputed species trees. More recently, several 'gold standard' reference sets, either manually curated^{16,17} or derived from trusted resources¹⁸, have been used as benchmarks. Finally, Dalquen et al.¹⁹ used simulated genomes to assess orthology inference in the presence of varying amounts of duplication, lateral gene transfer and sequencing artifacts.

This wide array of benchmarking approaches poses considerable challenges to developers and users of orthology methods. Conceptually, the choice of an appropriate benchmark strongly depends on the application at hand. Practically, most methods are not available as stand-alone programs and thus cannot easily be

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Protein Orthology for Comparative Genomics:

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Transcriptional differences between Parasitic & Saprobic stage C. immitis & C. posadasii

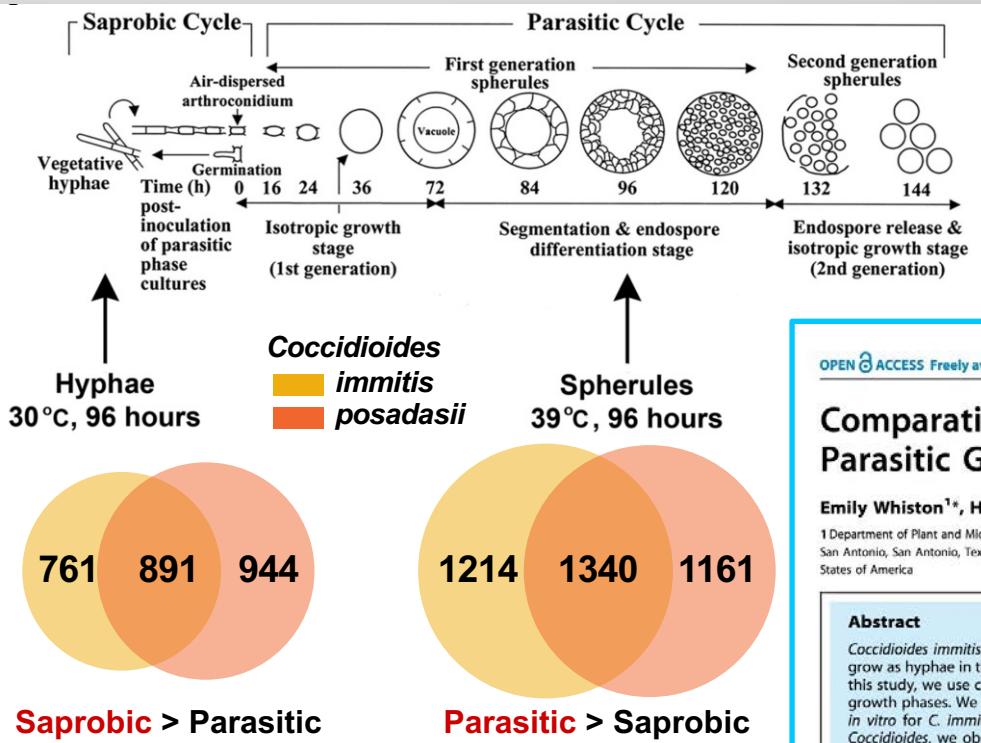


Table 1. Top 15 genes with significantly higher expression (up-regulated) in the saprobic phase.

Up in SAPROBIC stage *Coccidioides immitis* vs *C.posadasii*

Fold difference*	Annotation	Gene ID
185	Conserved protein (insect antifreeze protein repeat domain, predicted secreted)	CIMG_00925
166	Acetyltransferase	CIMG_07556
106	Acetamidase	CIMG_02374
101	Conserved hypothetical protein (predicted secreted)	CIMG_03870
94	Fungal hydrophobin (predicted secreted)	CIMG_06615
75	Conserved protein (PAN domain, predicted secreted)	CIMG_09824
53	Conserved protein (zinc-finger domain)	CIMG_00099
53	Conserved hypothetical protein	CIMG_06344
43	Putative serine proteinase	CIMG_09304
41	Cell wall synthesis protein (beta-glucosidase domain, SUN family, predicted secreted)	CIMG_05254
34	Hypothetical protein (predicted secreted)	CIMG_07839
31	Hypothetical protein	CIMG_13374
31	Helix-loop-helix transcription factor	CIMG_02390
29	Conserved hypothetical protein (pyridine nucleotide-disulphide oxidoreductase domain, predicted secreted)	CIMG_07557
24	Prp4 (C-terminal domain-containing protein, predicted secreted)	CIMG_07303

Table 2. Top 15 genes with significantly higher expression (up-regulated) in the parasitic phase.

Up in PARASITIC stage *Coccidioides immitis* & *C.posadasii*

Fold difference*	Annotation	Gene ID
125	Conserved hypothetical protein (DUF 536)	CIMG_09539
68	Heat shock protein 30 (Hsp20/alpha-crystallin domain)	CIMG_01749
31	Conserved hypothetical protein	CIMG_12822
30	Conserved hypothetical protein (YCI-related domain)	CIMG_07089
29	Conserved hypothetical protein	CIMG_13084
28	Hypothetical protein	CIMG_11522
28	Hypothetical protein	CIMG_05235
26	Polysaccharide deacetylase (Arp2/3 complex subunit Arc16)	CIMG_02628
24	Conserved hypothetical protein (predicted secreted)	CIMG_00509
23	Hypothetical protein	CIMG_11203
19	Spherule outer-wall glycoprotein (SOWgp, predicted secreted)	CIMG_04613
18	Conserved hypothetical protein	CIMG_10488
18	Hypothetical protein	CIMG_10670
17	Sphingosine hydroxylase	CIMG_01209
17	Conserved hypothetical protein	CIMG_04740

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A VEuPathDB Project

Site search, e.g. NCU06658 or *reductase or "binding protein"

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David

Download Data Files BETA makes downloading genome-scale files such as genome.fasta or GFF files easy and quick. Please check out the new Beta tool design and Contact Us with comments.

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Filter the searches below...

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VEuPathDB is packed with data, tools and visualizations that can help answer your research questions. We gather data from many sources, analyze according to standard workflows, and present the results for you to mine in a point and click interface. Here's how to get started:

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Explore the data and images on gene pages

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SNPs and CNV based on whole genome sequencing

Differential Expression Fold Change Percentile

Identify Genes based on *C. posadasii* C735 delta SOWgp Saprobic vs Parasitic Growth RNA-Seq (fold change)

For the Experiment: Saprobic vs Parasitic Growth unstranded

return protein coding Genes that are up-regulated with a Fold change >= 8 between each gene's minimum expression value (or a Floor of 10 reads) in the following Reference Samples: Saprobic Hyphae, Parasitic Spherules

and its maximum expression value (or the Floor selected above) in the following Comparison Samples: Saprobic Hyphae, Parasitic Spherules

Example showing one gene that would meet search criteria (Dots represent this gene's expression values for selected samples)

Up-regulated

Expression Value Comparison

Expression Value Reference

Reference Samples Comparison Samples

For each gene, the search calculates:

$$\text{fold change} = \frac{\text{comparison expression value}}{\text{reference expression value}}$$

and returns genes when fold change ≥ 8 .

You are searching for genes that are up-regulated between one reference sample and one comparison sample.

Get Answer

Build a Web Services URL from this Search >

Give this search a name (optional)

Give this search a weight (optional)

News »

FungiDB 63 Released WED MAY 03 2023

FungiDB 62 Released THU MAR 09 2023

FungiDB 61 Released THU DEC 15 2022

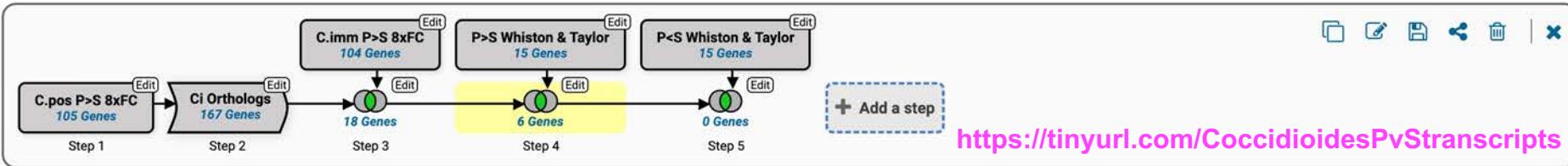
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Francis M. Ma... @fmarti... · May 6

The fungal grid: Fungal communication via electrical

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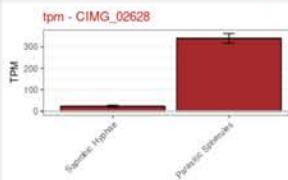
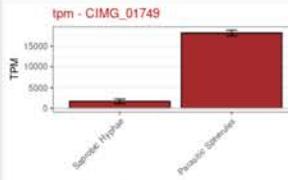
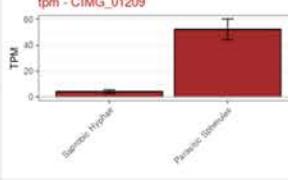
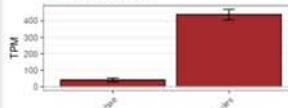
 Coccidioides Parasitic > Saprobic FC (Whiston & Taylor 2012) * 

<https://tinyurl.com/CoccidioidesPvStrancripts>

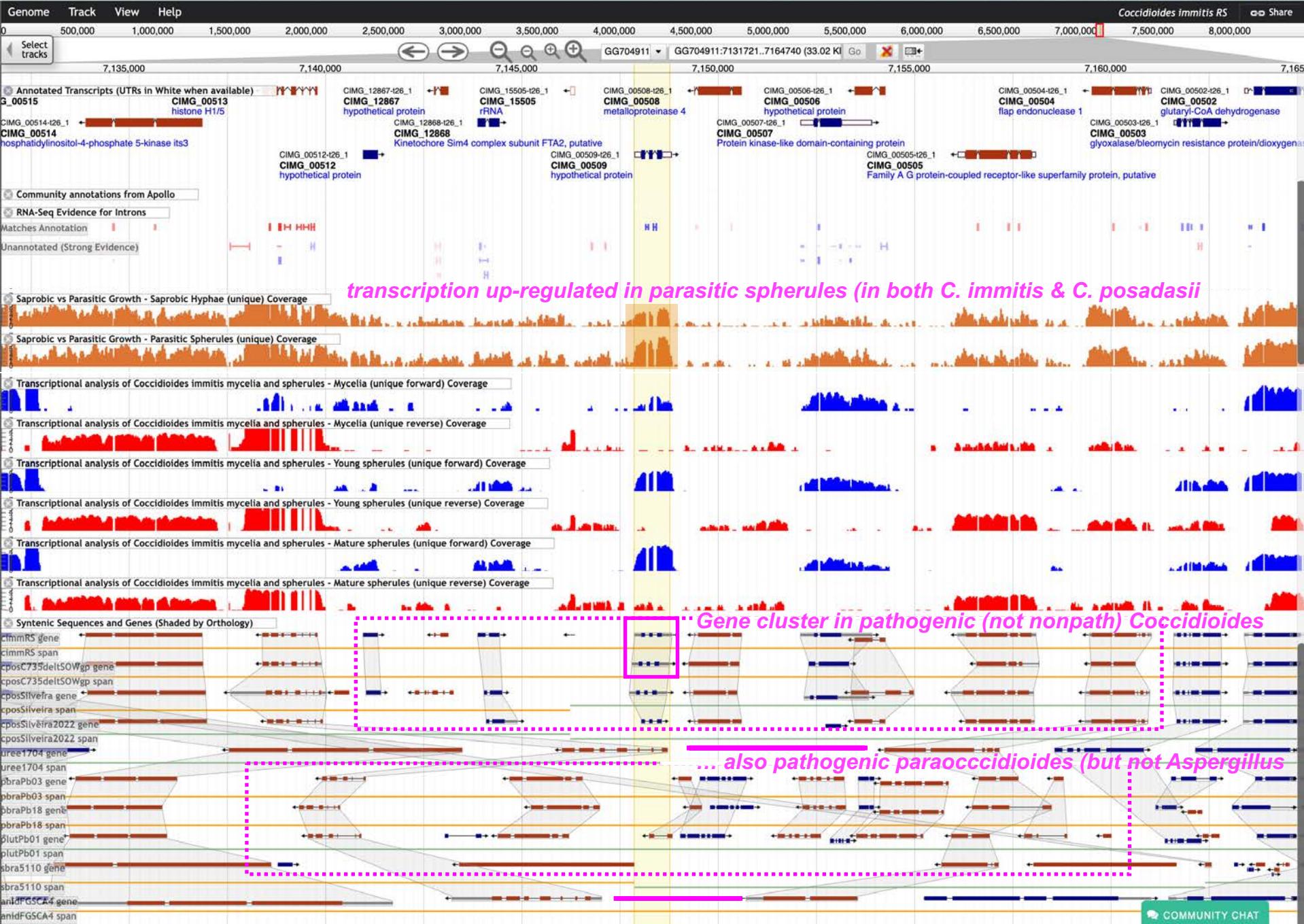
6 Genes (6 ortholog groups)

[Gene Results](#) [Genome View](#) [Analyze Results](#)

Rows per page: 100

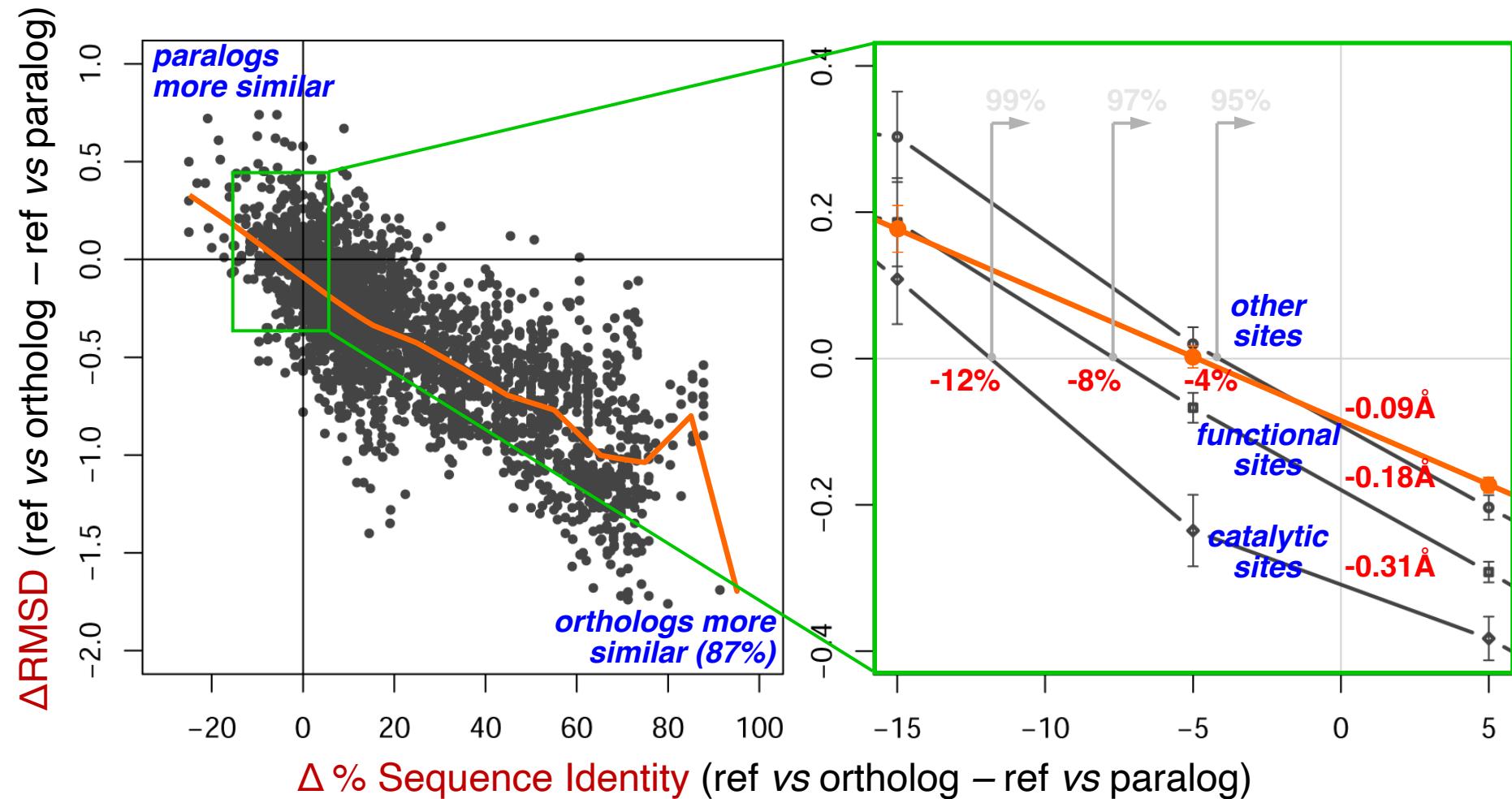
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	Gene ID	Transcript ID	Genomic Location (Gene)	Product Description	Computed GO Functions	Computed GO Processes	C.I. Saprobic/Parasitic RNA-Seq - tpm Graph
	CIMG_02628	CIMG_02628-t26_1	GG704911:1,361,297..1,362,421(−)	Arp2/3 complex subunit Arc16	catalytic activity;hydrolase activity, acting on carbon-nitrogen (but not peptide) bonds	carbohydrate metabolic process	
	CIMG_01749	CIMG_01749-t26_1	GG704911:3,854,425..3,855,421(+)	Heat shock protein 30	N/A	N/A	
	CIMG_01209	CIMG_01209-t26_1	GG704911:5,319,073..5,320,322(<)	C4-hydroxylase	iron ion binding;oxidoreductase activity	lipid biosynthetic process;obsolete oxidation-reduction process	
	CIMG_00509	CIMG_00509-t26_1	GG704911:7,147,867..7,148,818(+)	hypothetical protein	N/A	N/A	



Orthologs display greater structural conservation than paralogs ... particularly at functional sites

*Comparison of Reference / Ortholog / Paralog (AOP) triplets
499 catalytic sites (CSA), 4,041 functional sites (PDB)*



i Welcome to OrthoMCL where you will find the newest versions of our interface, features, tools and data. Since OrthoMCL went through a major algorithm update, we are keeping the old release 5 available in case needed. Here is a [form for sending your feedback](#) about the new site.

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- Avg % Homology
- Avg % Identity
- Avg % Match Length
- Avg E-Value
- EC Number
- Group ID(s)
- Group or Sequence ID
- Number of Sequences
- Number of Taxa
- PFam ID or Keyword
- Phyletic Pattern
- Text Terms

Proteins

- All Proteins
- BLAST
- EC Number

Overview of Resources and Tools



About OrthoMCL

OrthoMCL is a genome-scale algorithm for grouping orthologous protein sequences. Such orthologous sequences not only share evolutionary history, but also share function. Thus, ortholog prediction is important in predicting the function of newly identified proteins. Indeed, detection of orthologs has become more widespread with the rapid progress in genome sequencing and the discovery of protein sequences (Glover et al. 2019). Importantly, proteins in OrthoMCL groups have been shown to display a high degree of functional conservation (e.g., a group's proteins have consistent EC numbers) (Li et al. 2003), highlighting that OrthoMCL is useful for functional annotation of newly sequenced genomes.

OrthoMCL not only identifies groups shared by proteins from two or more species, but also groups representing species-specific gene expansion families. To achieve this, the OrthoMCL algorithm starts with reciprocal best BLAST hits within each proteome as potential in-


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[Orthology Phyletic Patterns](#)
[OrthoMCL tutorial](#)

Learn standard searches and ortholog group analysis within

[Search Strategies \(Basics\)](#)
[Search Strategies \(Advanced\)](#)


► Description

▼ Edge Options

Edge Type

- Ortholog
- Coortholog
- Inparalog
- Other Similarities

E-Value Cutoff

Max E-Value: 1E -35

A horizontal scroll bar with a grey track and a white slider.

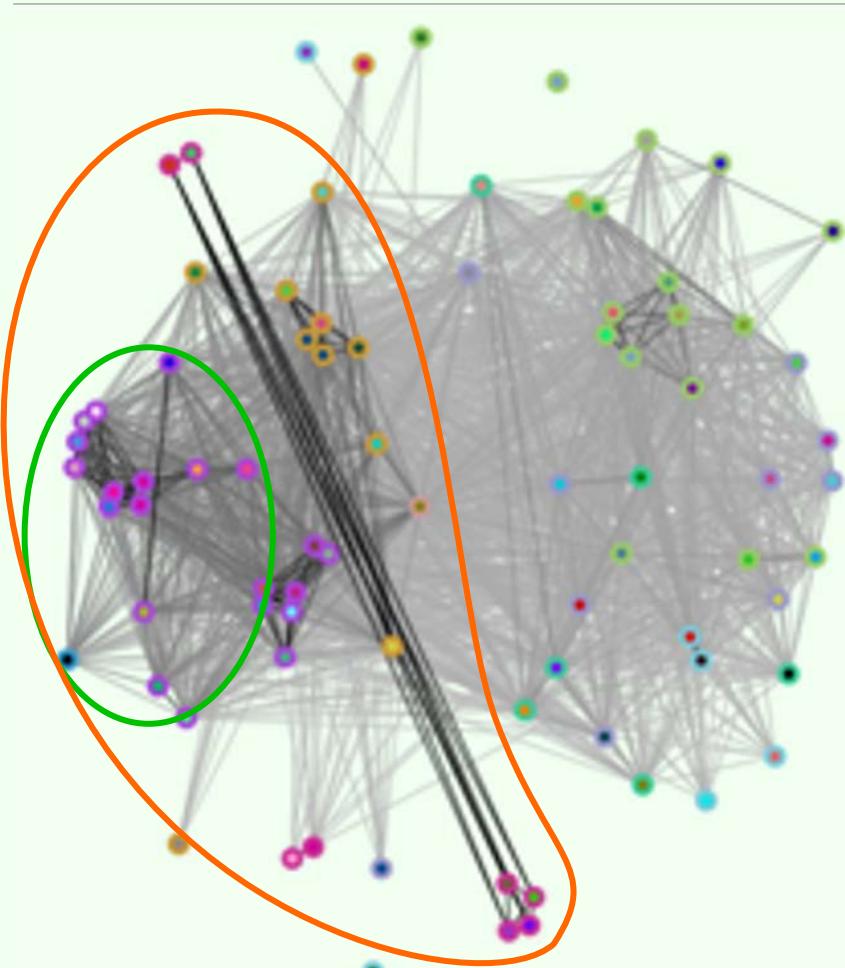
▼ Node Options

Show Nodes By

Taxa EC Numbers PFam Domains

Mouse over a taxon legend to highlight sequences of that taxon.

bant (1)	cbot (1)	cper (1)
lmon (1)	saur (1)	spne (1)
atum (1)	bsui (1)	bmal (1)
bpse (1)	cjej (1)	cbur (1)
ecol (1)	ftul (1)	gsul (1)
rsol (1)	sent (2)	sfle (1)
vcho (1)	wsuc (1)	yent (1)
ypes (1)	aaeo (1)	cpne (1)
ctep (1)	deth (1)	drad (1)
mtub (1)	mlep (1)	rbal (1)
syne (1)	tmar (1)	cmaq (1)
ckor (1)	halo (1)	hwal (1)
nmar (1)	tvol (1)	cmur (1)
ncan (1)	pber (1)	pcha (1)
npel (1)	ekoe (1)	puiv (1)



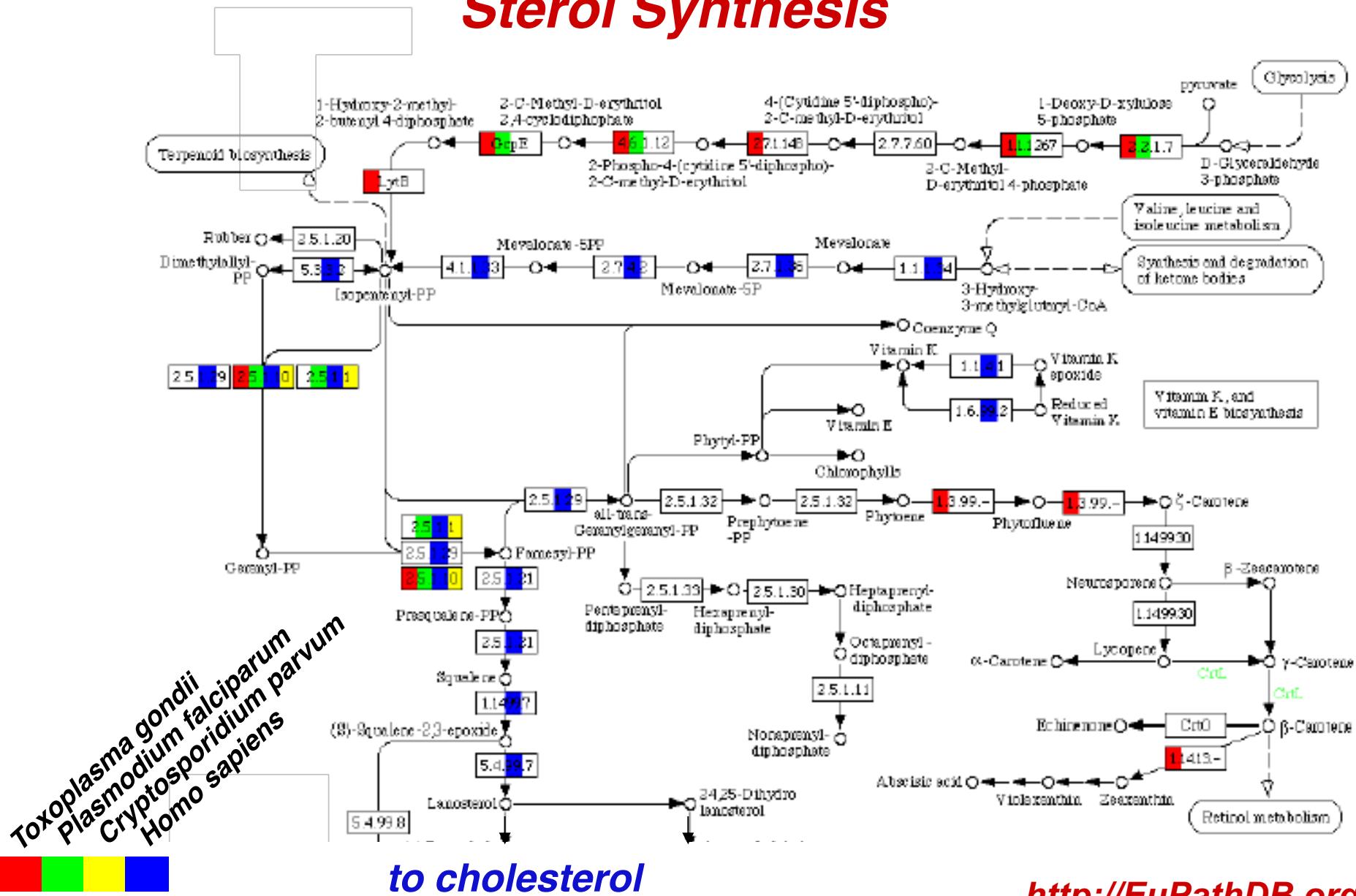
Sequence List

Node Detail

Search:

Accession	Taxon	
aaeo NP_214013	aaeo	399
afum Afu2g09840	afum	463
anid AN6032	anid	451
aory AO090011000651	aory	464
atha NP_194729	atha	554
atha NP_564953	atha	484
atum NP_354359	atum	261
bant YP_016674	bant	280
bmal YP_102541	bmal	296
bpse YP_333020.1	bpse	296
bsui NP_698035	bsui	279
calb calb_sc5314__orf19.579	calb	789
calb CAWT_04470	calb	789
cbot YP_001786179.1	cbot	395
cbur NP_820340	cbur	297
cgle XP_448052	cgle	807
cimm CIMG_07826T0	cimm	463
cjej YP_002344015	cjej	380
ckor YP_001737263	ckor	403
cmaq YP_001540023	cmaq	303
cmer CMQ278C	cmer	500
cmur CMU_032400	cmur	217
cneg CNAG_02786T0	cneg	736
cneo CNBG_3441T0	cneo	651
cper YP_695722	cper	269

Painting Orthologs onto KEGG Pathways: *Sterol Synthesis*



Orthology-based inference

- *Eukaryota (EUKA):

- *Alveolates (ALVE):

- Ciliates (CILI):

- *Apicomplexa (APIC):

- Coccidia (COCC):

- *Aconoidasida (ACON):

- Haemosporida (HAEM):

- Piroplasmida (PIRO):

- Amoebozoa (AMOE):

- Euglenozoa (EUGL):

- Viridiplantae (VIRI):

- Streptophyta (STRE):

- Chlorophyta (CHLO):

- Rhodophyta (RHOD):

- Cryptophyta (CRYP):

- Bacillariophyta (BACI):

- Fungi (FUNG):

- Microsporidia (MICR):

- Basidiomycota (BASI):

- Ascomycota (ASCO):

- Metazoa (META):

- ✗ Platyhelminthes (PLAT):

- ✗ Nematodes (NEMA):

*Apicoplast genes should be ...
present in:*

- *Toxoplasma*

- *Plasmodium*

- *Theileria*

- *Plants*

... but not in:
- *Cryptosporidium*
- *Mammals*

tthe

chom

cmu

xpar

ncan

tgon

pber

pcha

pfal

pkno

pviv

pyoe

bbov

tanr

xtpar

ddis

ehis

edis

einv

lbra

linf

lmaj

lmex

tbru

tbrg

tcon

tcru

tviv

atha

osat

ppat

rcom

crei

otau

cmer

gthe

tpse

ecun

cneo

lbic

pchr

afum

aory

cgle

cimm

cpos

dhan

egos

gzea

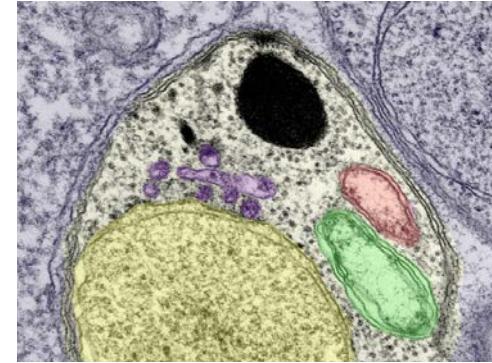
klac

xsmn

xbmaa

xcbri

xcele



Horizontal gene transfer facilitated the evolution of plant parasitic mechanisms in the oomycetes

Thomas A. Richards^{a,b,1}, Darren M. Soanes^a, Meredith D. M. Jones^{a,b}, Olga Vasieva^c, Guy Leonard^{a,b}, Konrad Paszkiewicz^a, Peter G. Foster^b, Neil Hall^c, and Nicholas J. Talbot^a

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Edited by W. Ford Doolittle, Dalhousie University, Halifax, Canada, and approved July 27, 2011 (received for review March 31, 2011)

Horizontal gene transfer (HGT) can radically alter the genomes of microorganisms, providing the capacity to adapt to new lifestyles, environments, and hosts. However, the extent of HGT between eukaryotes is unclear. Using whole-genome, gene-by-gene phylogenetic analysis we demonstrate an extensive pattern of cross-kingdom HGT between fungi and oomycetes. Comparative genomics, including the de novo genome sequence of *Hyphochytrium catenoides*, a free-living sister of the oomycetes, shows that these transfers largely converge within the radiation of oomycetes that colonize plant tissues. The repertoire of HGTs includes a large number of putatively secreted proteins; for example, 7.6% of the secreted proteome of the sudden oak death parasite *Phytophthora ramorum* has been acquired from fungi by HGT. Transfers include gene products with the capacity to break down plant cell walls and acquire sugars, nucleic acids, nitrogen, and phosphate sources from the environment. Predicted HGTs also include proteins implicated in resisting plant defense mechanisms and effector proteins for attacking plant cells. These data are consistent with the hypothesis that some oomycetes became successful plant parasites by multiple acquisitions of genes from fungi.

osmotrophy | pseudofungi | lateral gene transfer

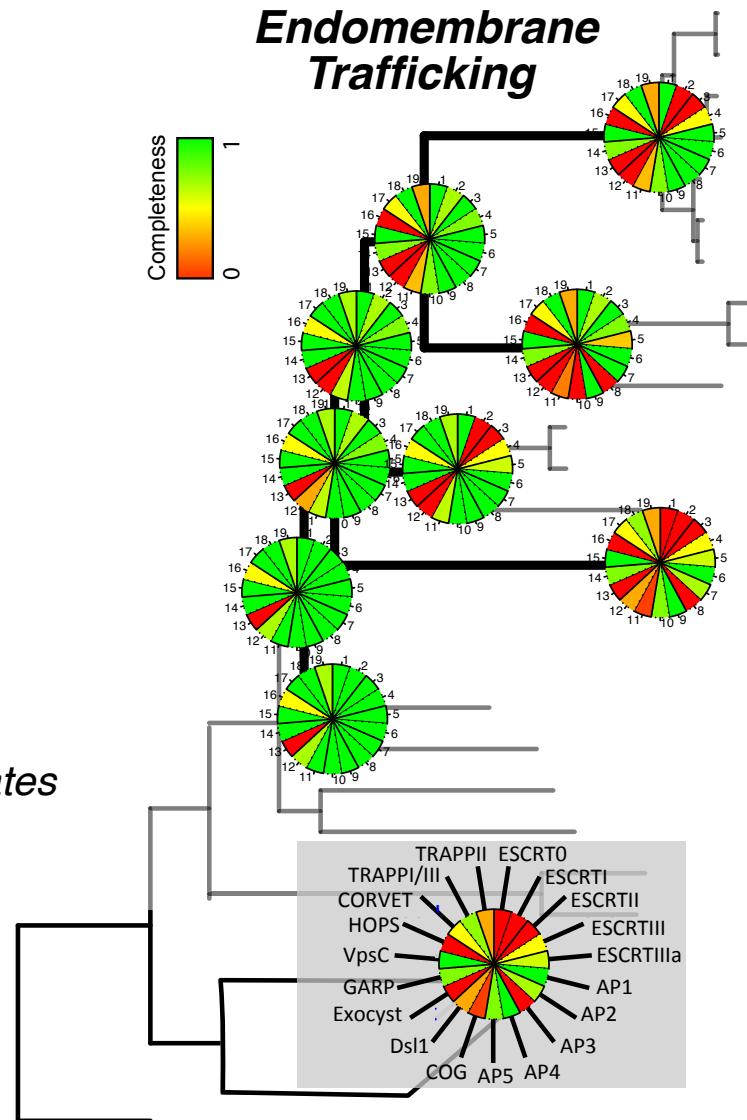
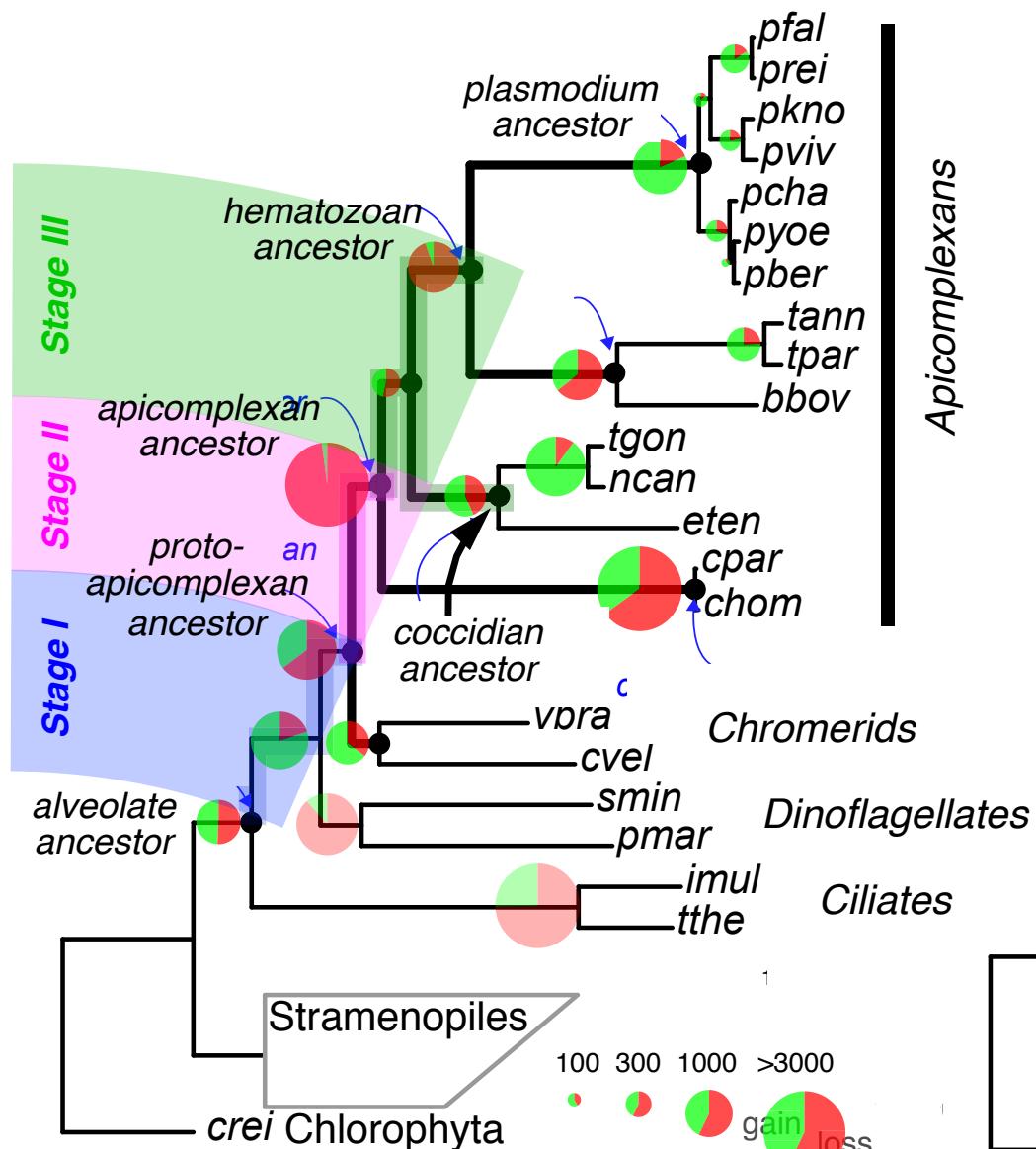
Horizontal gene transfer (HGT) involves the transfer of genetic material between reproductively isolated lineages (1, 2) and has been an important factor in prokaryotic evolution (3–5). By contrast, the role of HGT between eukaryotic genomes is

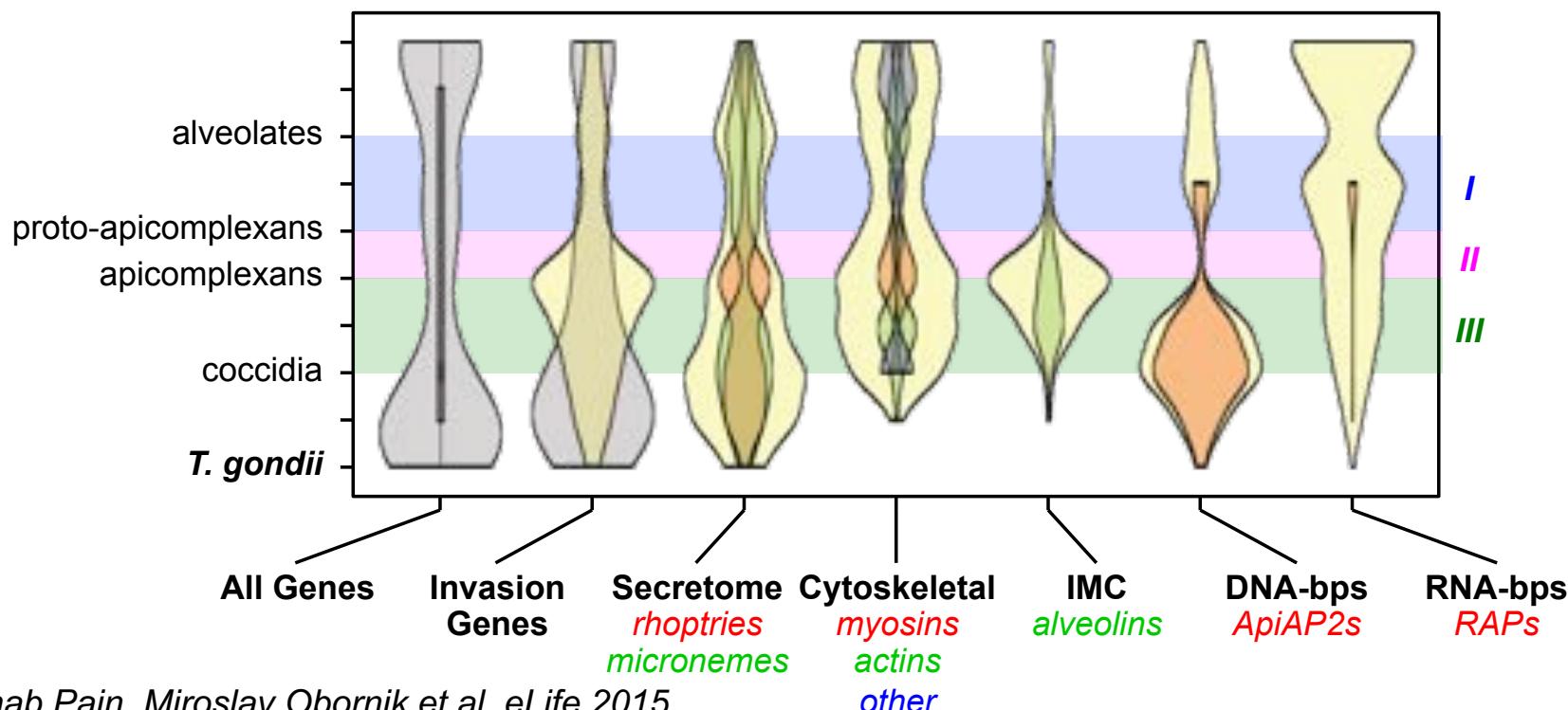
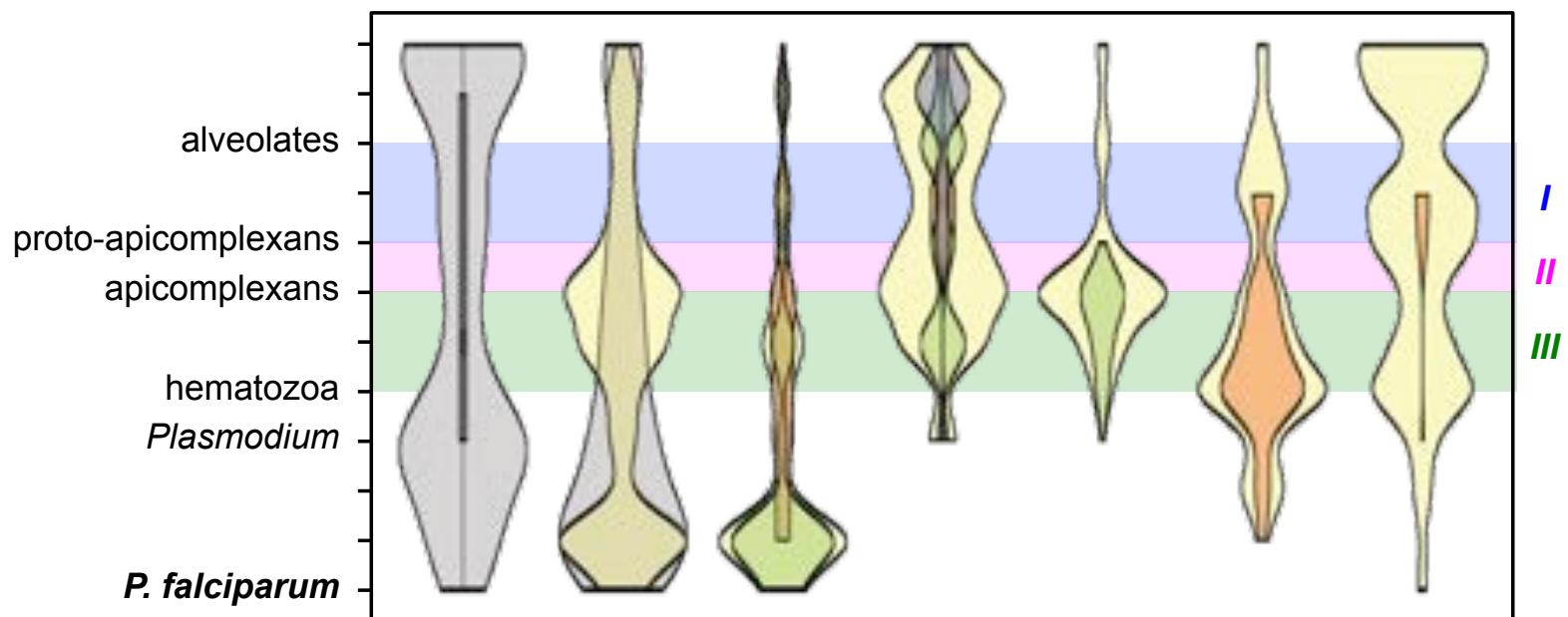
ramorum, for example, whereas the Irish potato famine of the 19th century was caused by the late blight parasite *Phytophthora infestans*. Important crop diseases caused by fungi include the devastating rice blast disease caused by *M. oryzae* and the rusts, smuts, and mildews that affect wheat, barley, and maize. In this study we report that HGT between fungi and oomycetes has occurred to a far greater degree than hitherto recognized (19). Our previous analysis suggested four strongly supported cases of HGT, but by using a whole-genome, gene-by-gene phylogenetic analysis we now reveal a pattern of 34 transfers and propose that these transfers have been fundamental to the evolution of plant parasitic traits within the oomycetes.

Results and Discussion

Identifying and Testing the Pattern of HGT Between Fungi and Oomycetes. Among the best methods for identifying HGT is to identify a gene phylogeny that places a taxonomic group (recipient) within the branches of a distantly related group (donor) in direct contradiction to the known phylogenetic relationships of the respective taxa (2, 6). To identify all potential gene transfers between fungi and oomycetes, we selected the predicted proteomes of the oomycete species *Phytophthora ramorum*, *Phytophthora sojae*, *Phytophthora infestans*, and *Hyaloperonospora parasitica* (also named *Hyaloperonospora arabidopsis*) (21–23). We processed each proteome separately, first excluding all putative transposable elements (*SI Materials and Methods*). Then

What happened when? Ortholog group gains and losses during apicomplexan evolution





Protein Orthology for Comparative Genomics:

- ❖ ***Genome Comparisons*** (*synteny*)
- ❖ ***Functional Comparisons*** (*particularly for proteins*)
 - *how make reliable inferences ... based on functional data in other species?*
- ❖ ***Protein Homology: Orthology & Paralogy***
- ❖ ***Methods based on:***
 - ***functional characterization*** (*typically unavailable at genomic scale*)
 - ***structure*** (*ditto ... and not necessarily = function*)
 - ***expert curation*** (*labor-intensive, especially for eukaryotic taxa*)
 - ***synteny*** (*common for closely-related taxa, but decays rapidly*)
 - ***protein sequence similarity*** – e.g. OrthoMCL, OrthoFinder (*FungiDB, Mycocosm*)
 - ***domain structure*** – e.g. EggNOG, Compara (*Ensembl*)
 - ***phylogeny*** – e.g. OMA, OrthoFinder2
- Note: Benchmarking is difficult without a gold standard (Q4O)!***
- ❖ ***Orthology applications:***
 - ***Validation of genome assemblies & completeness*** (e.g. BUSCO)
 - ***Gene gains, losses, amplification, horizontal transfer*** (*phyletic pattern profiling*)
 - ***Functional inference for understudied taxa (GO terms, etc)***