

# Genomics on the Half Shell: Making Science more Open

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**Steven B. Roberts**

Associate Professor

School of Aquatic and Fishery Sciences

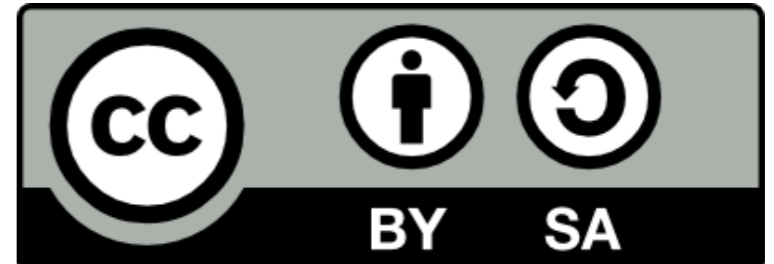
University of Washington

[robertslab.info](http://robertslab.info)

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---

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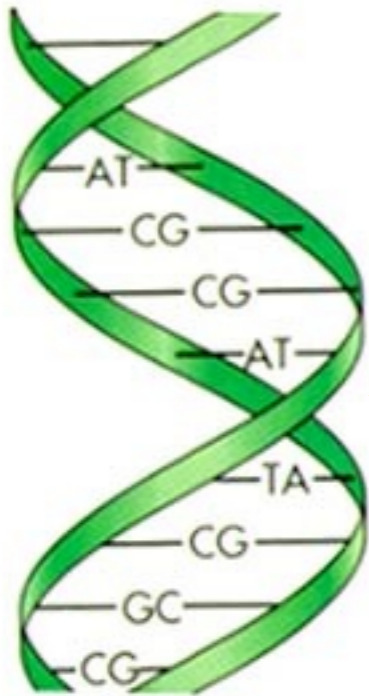
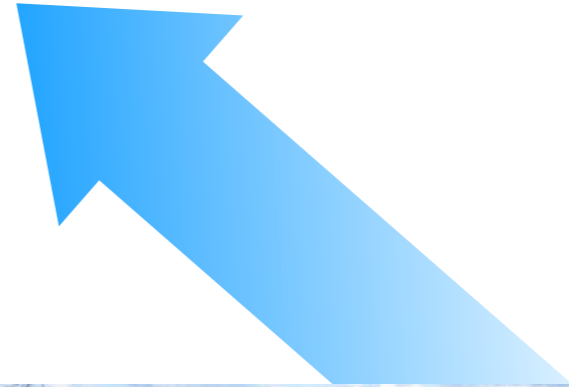
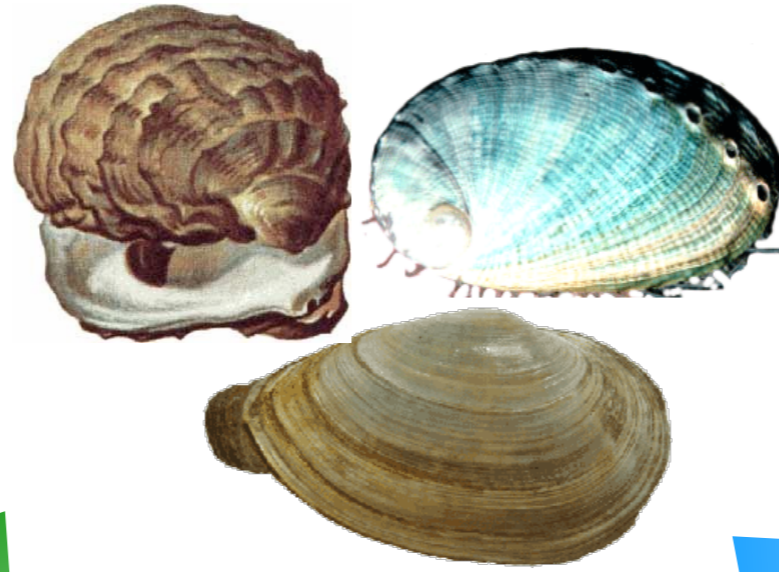
everything else...



BIV...VE BANDSTAND





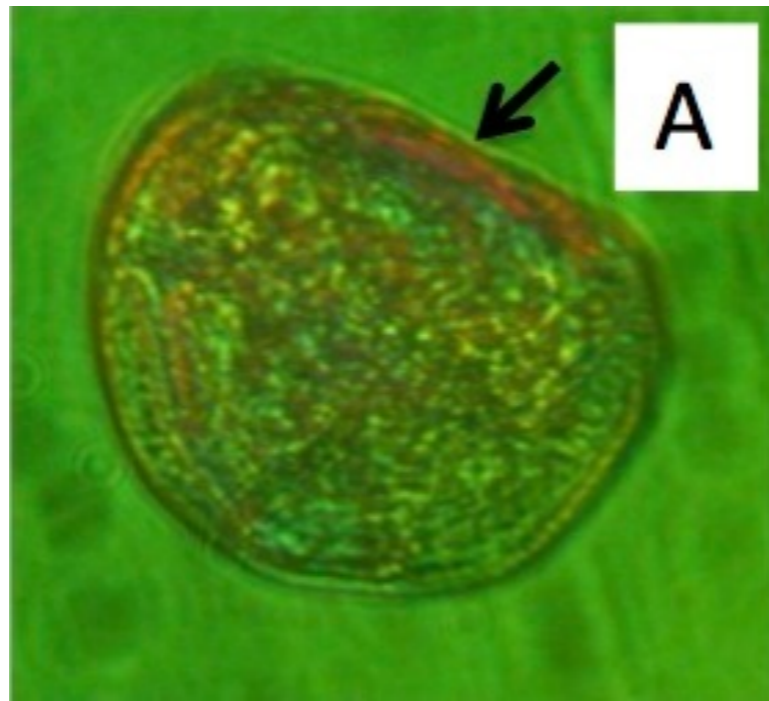






**Transcriptome**  
**Proteome**  
**DNA**  
**Methylation**





**Biology**

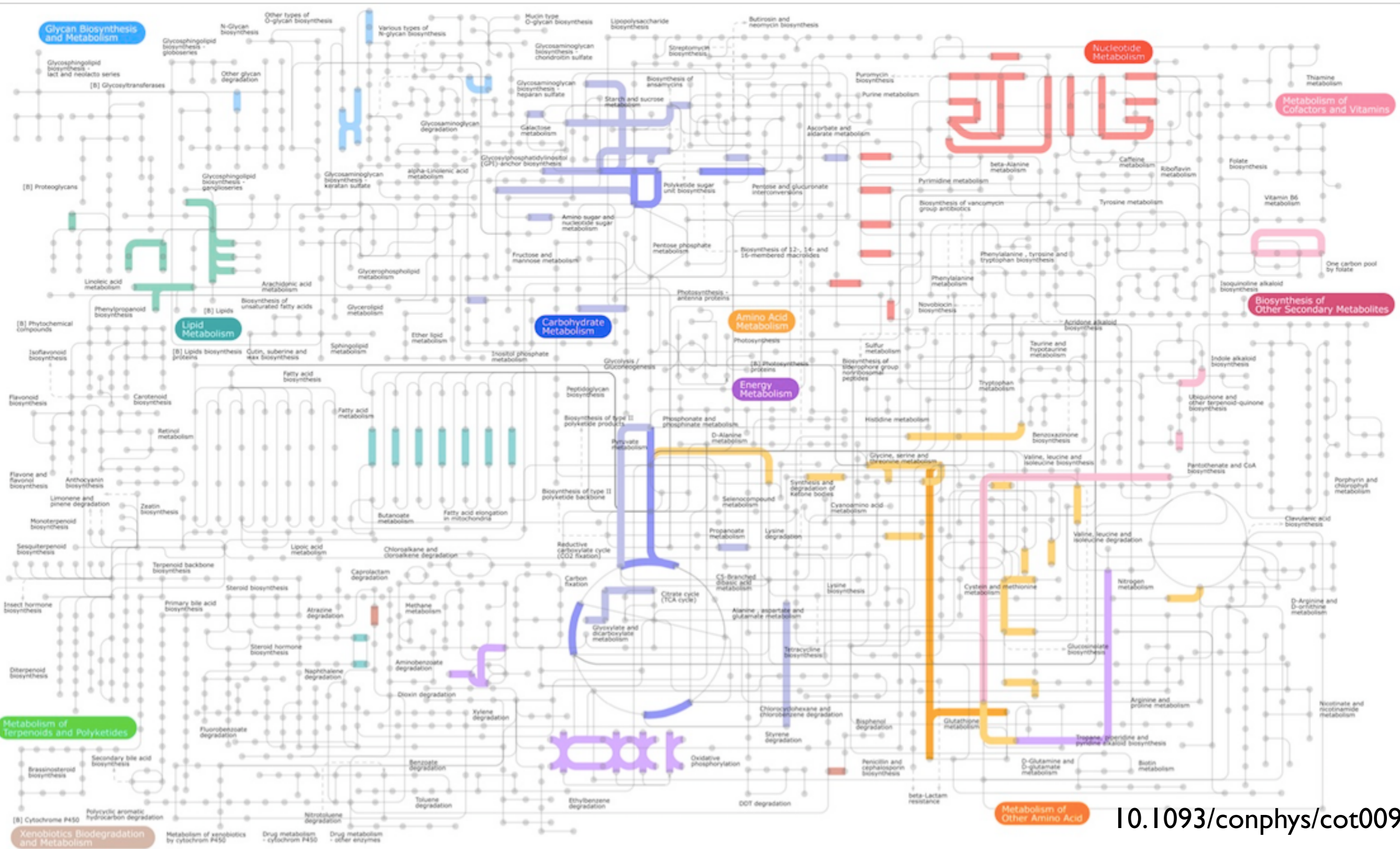
Environment

# Ocean Acidification

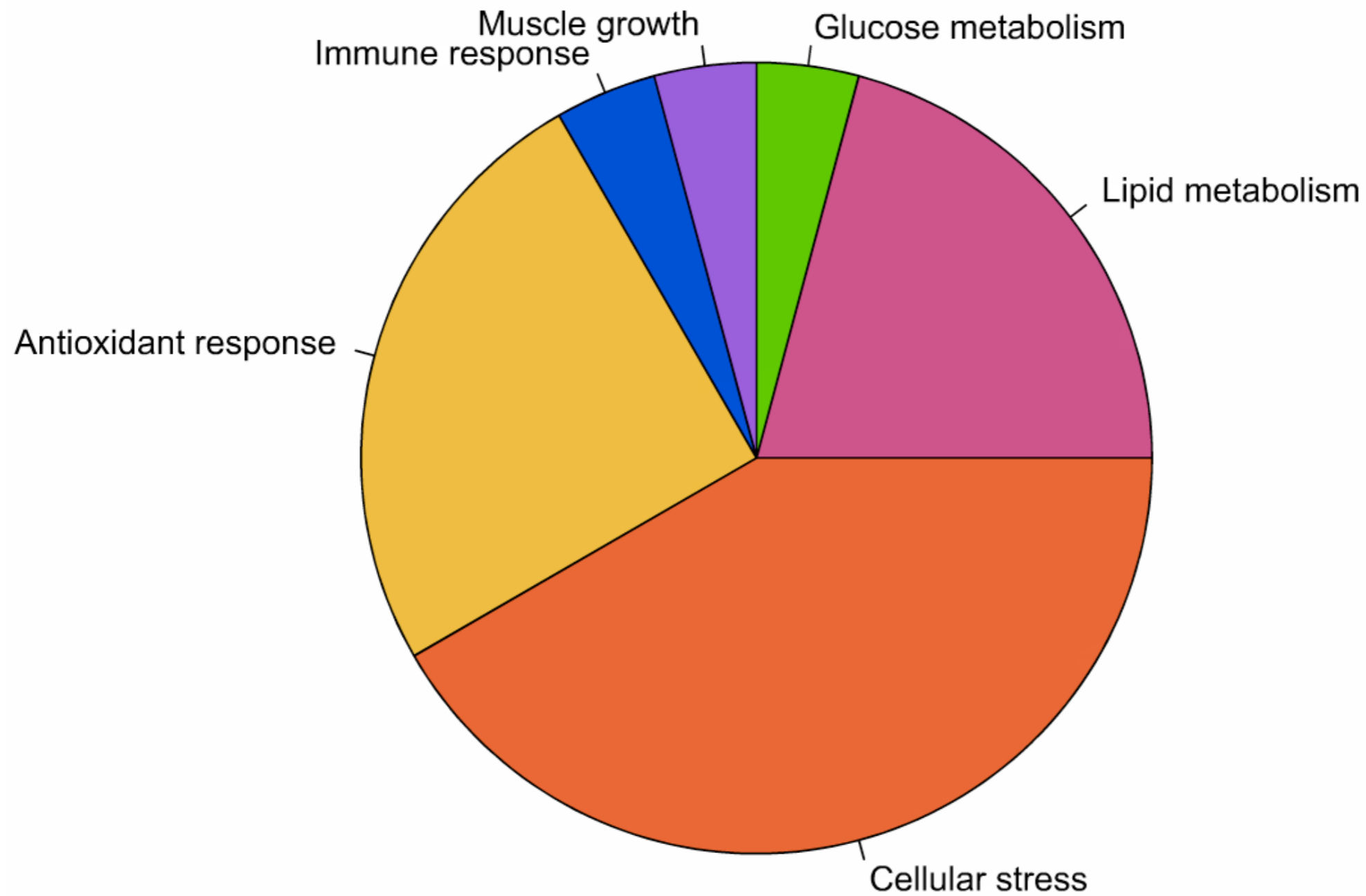


**Elevated pCO<sub>2</sub> causes developmental delay in early larval Pacific oysters, *Crassostrea gigas*.  
Timmins-Schiffman et al 2012**

# Shotgun Proteomics



# Shotgun Proteomics



# Ocean Acidification

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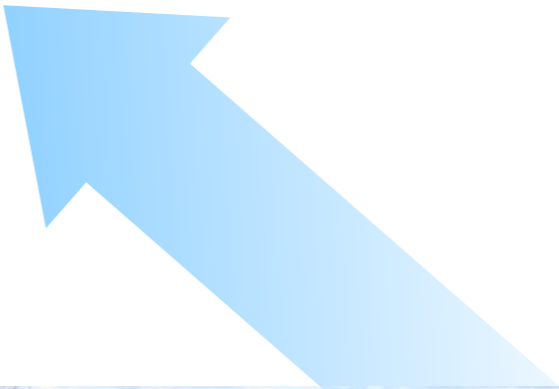
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**Transcriptome**

**Proteome**

**DNA  
Methylation**





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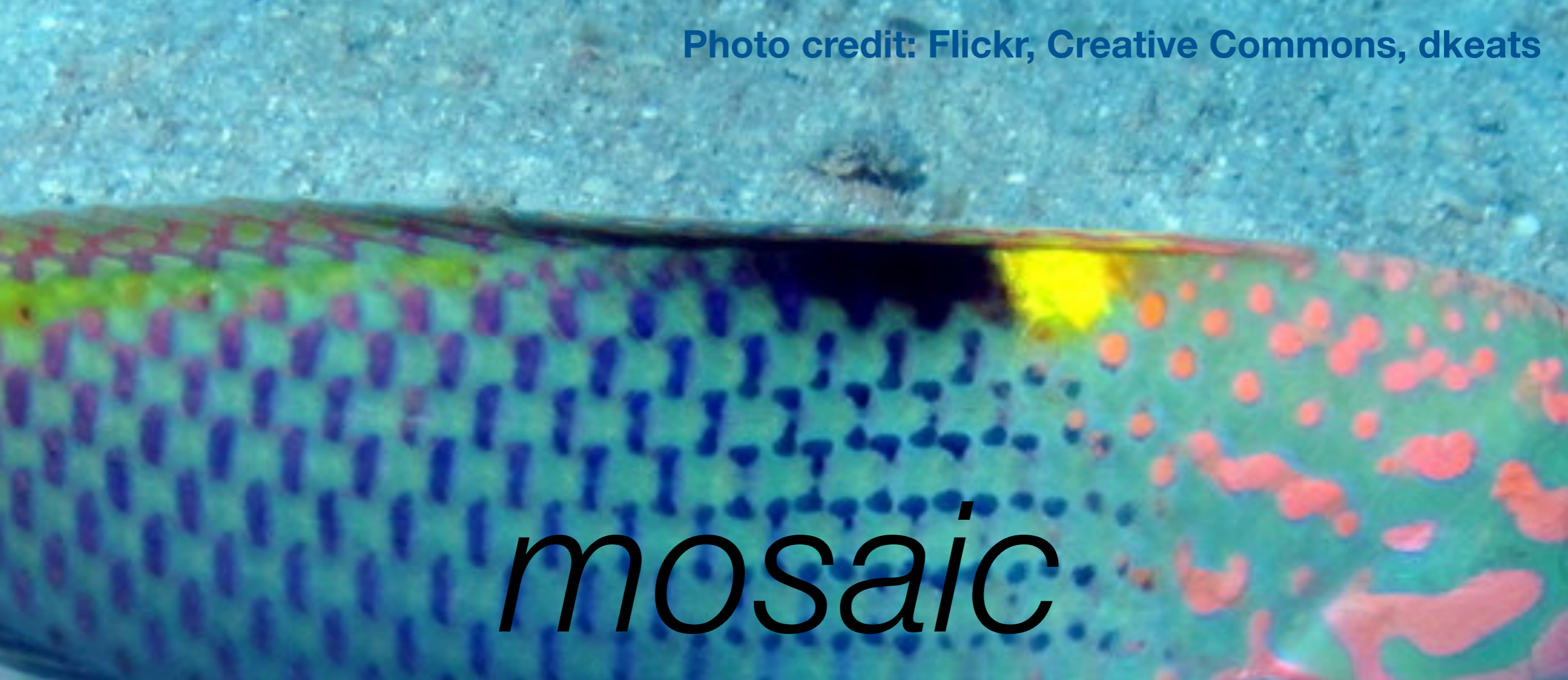
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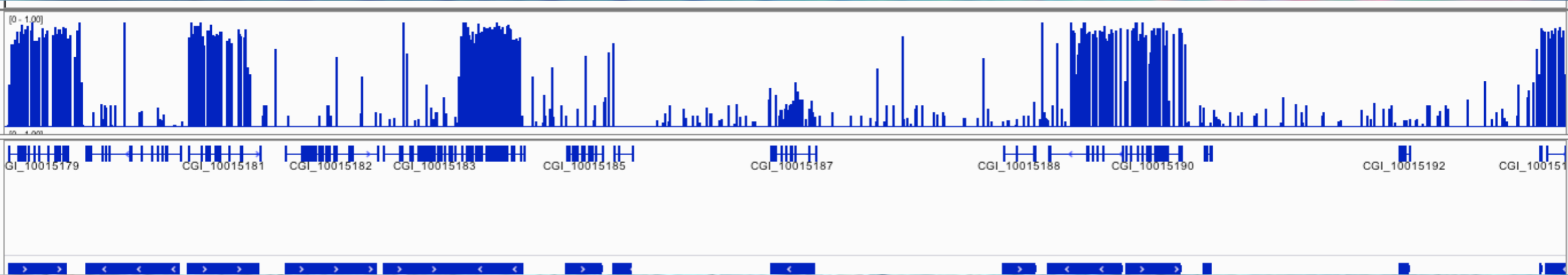
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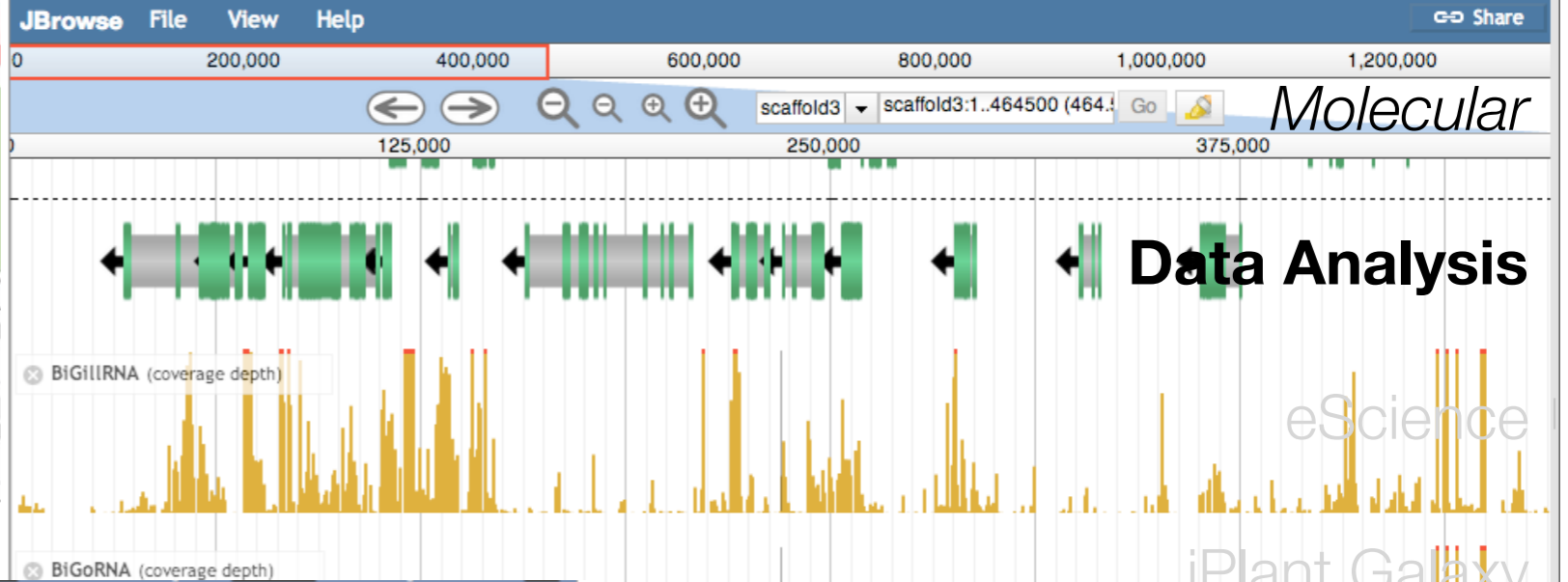
*mosaic*



associated with gene bodies

# GENOME VIEW

Crassostrea gigas (oyster\_v9, INSDC Assembly GCA\_000297895.1, Sep 2012): 7,658 scaffolds (vGCA\_000297895.1, id22811): softmasked by I



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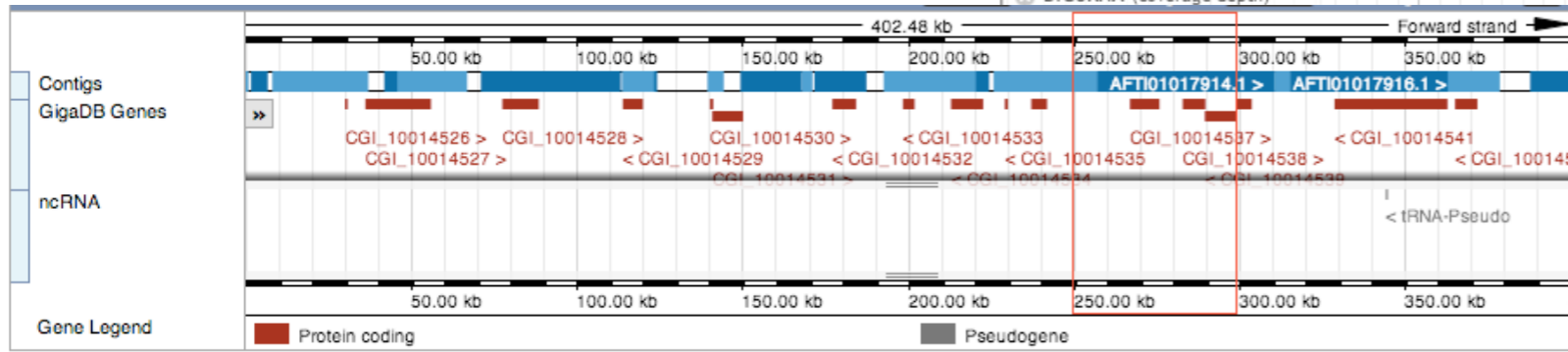
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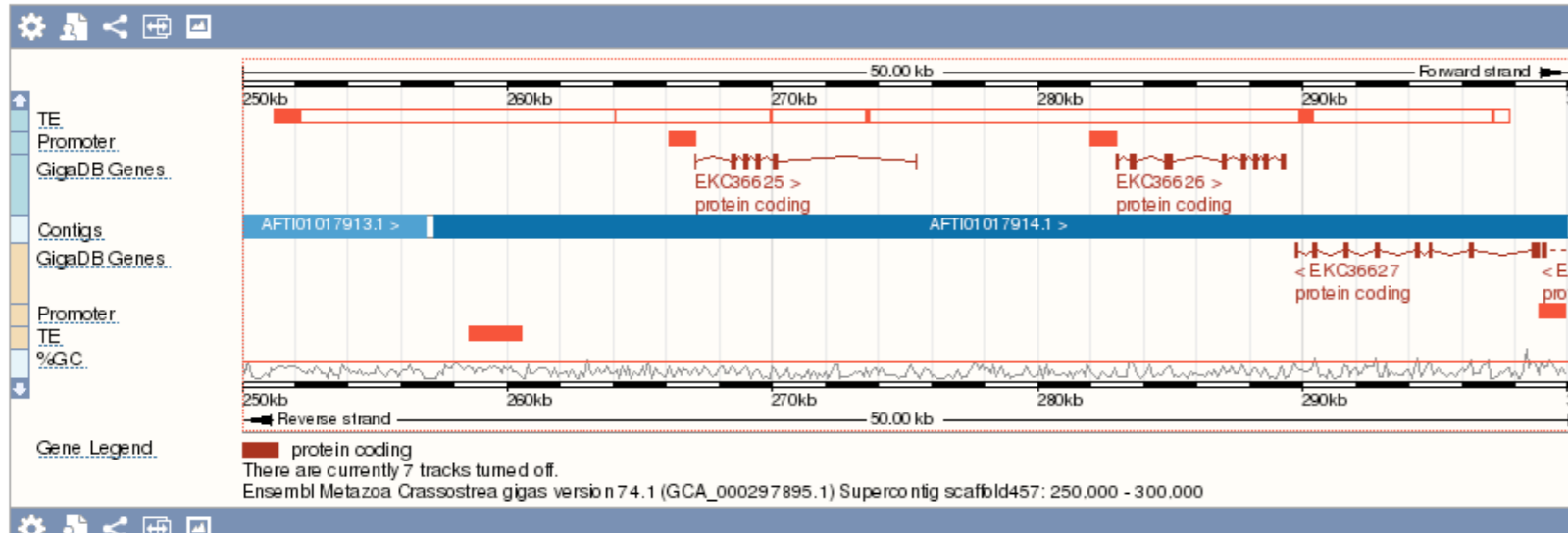
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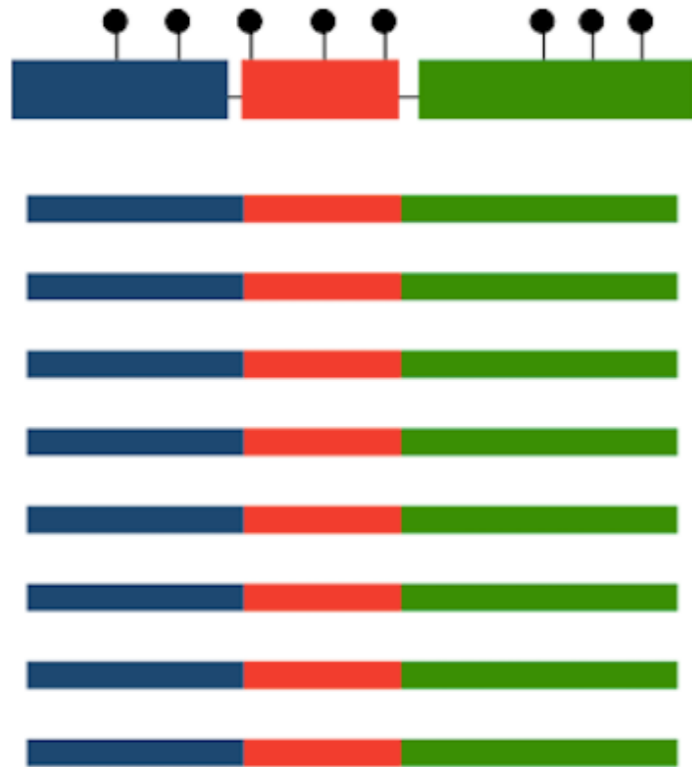
Location:

Gene:





# Stochastic Variation



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## **A context dependent role for DNA methylation in bivalves**

*Mackenzie R. Gavery and Steven B. Roberts*

[10.1093/bfgp/elt054](https://doi.org/10.1093/bfgp/elt054)

[10.6084/m9.figshare.880763](https://doi.org/10.6084/m9.figshare.880763)



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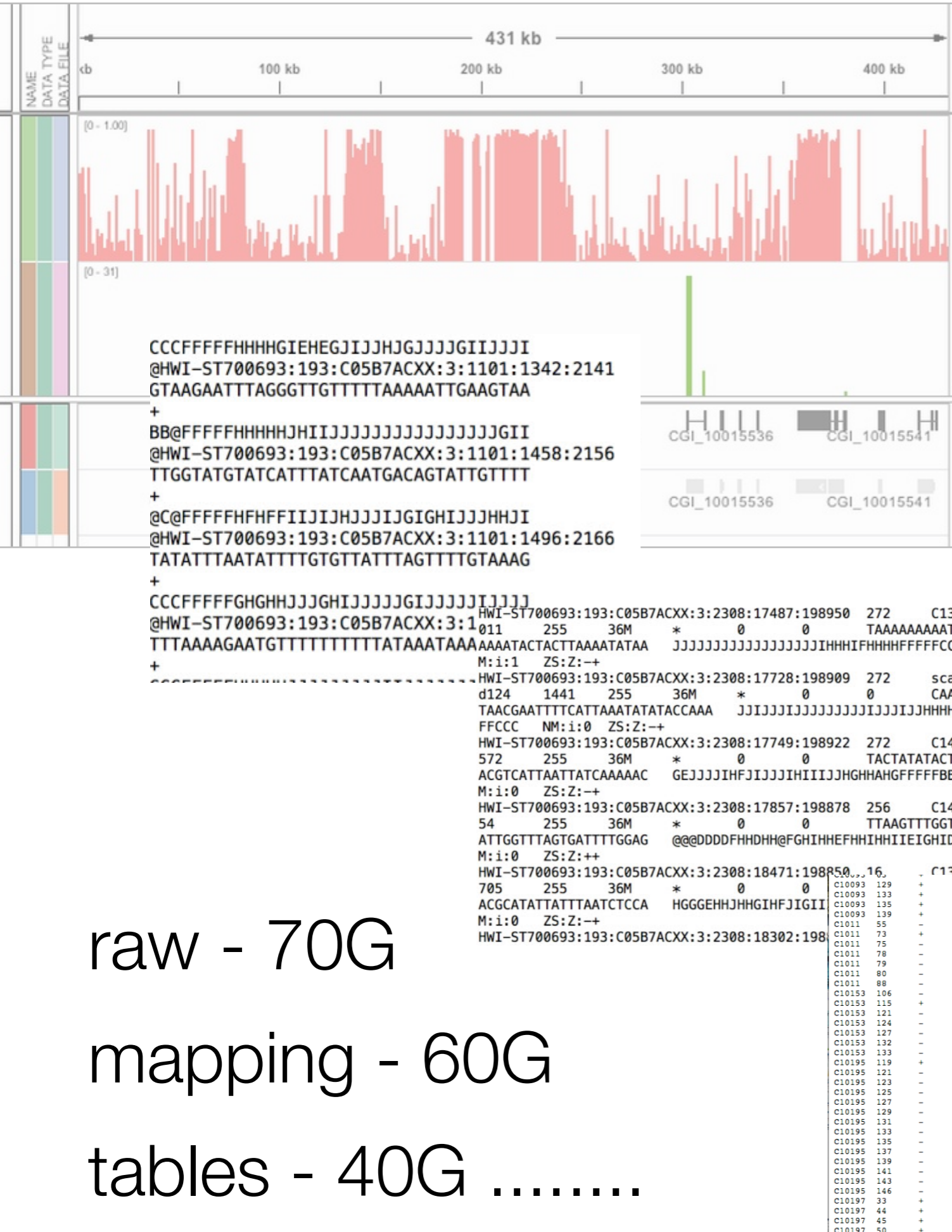
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raw - 70G  
 mapping - 60G  
 tables - 40G .....

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static

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Orthologs

CpG statistics

Structural Elements

Other species genomes

Transposable Elements

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Transcription Factors  
Binding Sites

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transcripts

dynamic

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Genetic Variation

Epigenetic Features

RNA-Sequencing

Single Nucleotide Polymorphisms

DNA Methylation

Expressed Sequence Tags

Amplified Fragment  
Length Polymorphisms

Histone Modification

Expression Microarrays

Simple Sequence Repeats

miRNA Expression

Genomic  
Data Types

Size  
Growth  
Location  
Environment  
Stage  
Treatment  
Tissue  
Trait  
Strain

Biology

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**Phenotype**

Yield

Disease Resistance

Increased Growth Rate

Tissue Quality

Fecundity

Appearance

**Gene Expression**

**Epigenetics**

**Genetics**

**Environment**

- Single Nucleotide Polymorphisms
- Simple Sequence Repeats
- Amplified Fragment Length Polymorphisms

- DNA Methylation Patterns
- miRNA Expression
- Histone Modifications

Temperature

Diet



Biology

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**Data Tables**

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*transcripts*

*dynamic*

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Amplified Fragment  
Length Polymorphisms

Histone Modification

Expression Microarrays

Simple Sequence Repeats

miRNA Expression

**Genomic  
Data Types**

Size  
Growth  
Location  
Environment  
Stage  
Treatment  
Tissue  
Trait  
Strain



# eScience Institute

## Querying Disparate Oyster Datasets | qDOD

The goal of this project is to produce a web-based interface for querying and visualizing *Crassostrea gigas* genomic datasets. This site serves as a portal for documenting our efforts, providing user access, as well as a means to gather feedback.

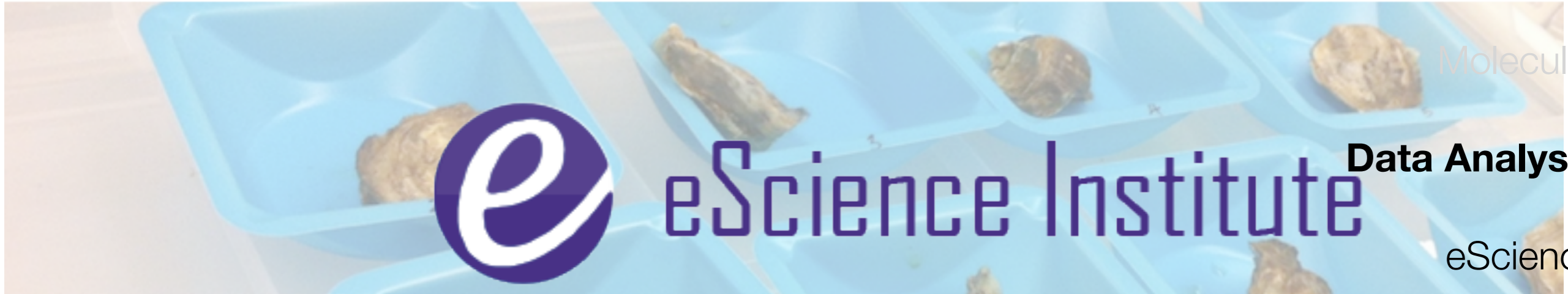
### Preliminary Phase: Aggregating Datasets

Using [SQLShare](#) as a platform we have already begun to aggregate and format data. Anyone can view (and contribute) using the tag "qDOD". Below is a table describing some of the relevant datasets. "Snapshot" provides you with a screenshot of the data in SQLShare and "Direct Link" brings you directly to the data in SQLShare. You can also [open the table in a new webpage](#).

qDOD online			
qDOD_Cgigas_gene_fasta	sequence fasta file. Exon only.	<a href="http://goo.gl/ogCxl">http://goo.gl/ogCxl</a>	<a href="https://sqlshare.esc">https://sqlshare.esc</a>
qDOD_Zhang_Gil_gene_RNA-seq	Gill RNA-seq data (gene based)	<a href="http://goo.gl/8oISR">http://goo.gl/8oISR</a>	<a href="https://sqlshare.esc">https://sqlshare.esc</a>
qDOD_Zhang_Mgo_gene_RNA-seq	Male Gonad RNA-seq data (gene based)	<a href="http://goo.gl/6buVz">http://goo.gl/6buVz</a>	<a href="https://sqlshare.esc">https://sqlshare.esc</a>

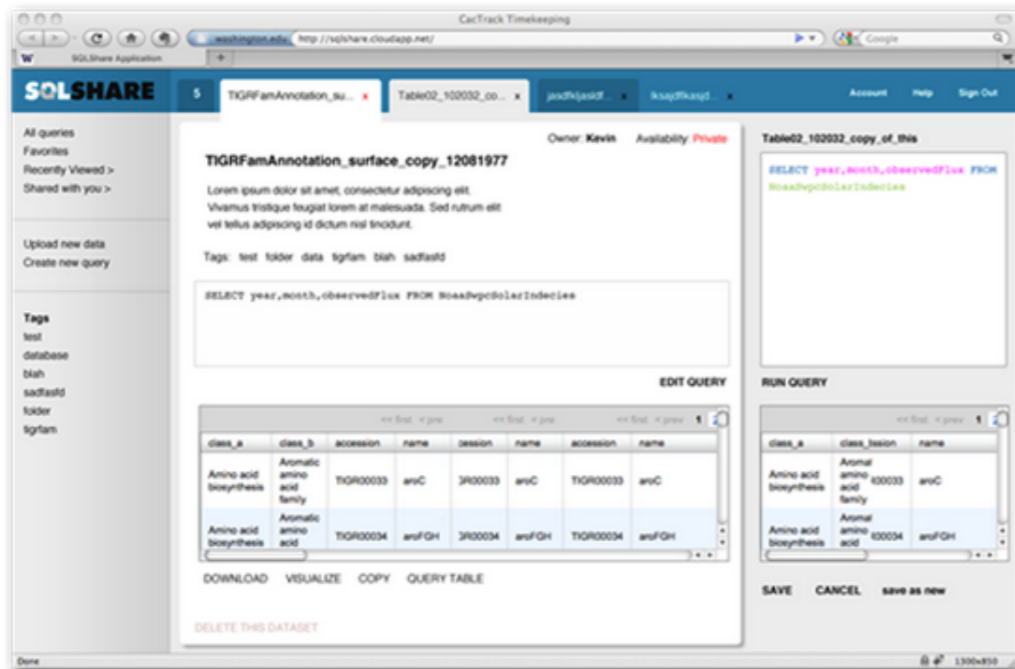
**DATA**





## SQLSHARE

SQLShare is an easier way to store and share your data. Get answers to your research questions right now.



Log in using your account:



Don't have an account?

Create a [Google Account](#) and start using SQLShare quickly.

### Upload

Upload any tabular data and start analyzing instantly. No need to install, configure, or design a database.

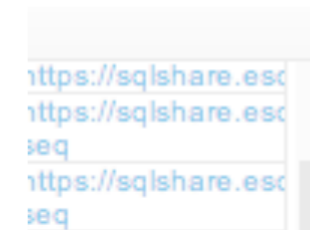
### Modify

Exercise the full power of SQL even with zero programming experience: joins, subqueries, set operations.

### Share

Analyze and compare your data collaboratively. Derive new datasets and share them with your colleagues.

Anyone can view (and contribute) using the links you generate. Share a screenshot of the data in the table in a new webpage.



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  - techtrip 34
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  - skyline 24
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  - ssgcid 18
  - qdod2 18
  - qdod 18
  - swissprot 17
  - sun 16
  - tsg 16

**Your Datasets**

Filter dataset by keyword:

Name	Sharing / Owner	Modif
qdod_proteome_blast_mouse	sr320@washington.edu	Jan 2
qDOD_v9_gene GFF format file of oyster genes ~28k <span style="background-color: #e0e0e0; padding: 2px;">gene</span>	sr320@washington.edu	Nov
_qdod_goslim_graphstest	sr320@washington.edu	Oct 2
SNP_RNAseqLibrary_SB_BiGill SNP table from RNA-seq library - SB gill tissue pool (BiGill complement) <span style="background-color: #e0e0e0; padding: 2px;">qdod2</span>	sr320@washington.edu	Oct 2
BiGill_meth_Zhang_exp Gene-centric data including length, CG, percent methylation (gill) and tissue specific RPKM data from Zhang et a <span style="background-color: #e0e0e0; padding: 2px;">qdod2</span>	sr320@washington.edu	Oct 2
qDOD_Cgigas_gene_fasta Tabular format of Cgigas gene sequence fasta file Derived using Dataset: Genomic data from the Pacific oyste <span style="background-color: #e0e0e0; padding: 2px;">qdod2</span>	sr320@washington.edu	Oct 2
qDOD Cgigas Gene Descriptions (Swiss-prot) Description and evalues associated with Cgigas 28k genes Derived using Dataset: Genomik <span style="background-color: #e0e0e0; padding: 2px;">blast</span>	sr320@washington.edu	Oct 2
file0	sr320@washington.edu	Aug 1
BiGill meth with SP	sr320@washington.edu	Aug 1
SPID and GO Numbers Swiss-Prot IDs and corresponding GO numbers <span style="background-color: #e0e0e0; padding: 2px;">qdod</span>	sr320@washington.edu	Aug 1
Cgigas_larvae_RNAseq_OsHV_GO	sr320@washington.edu	Jul 2
qDOD_Cgigas_GO_GOslim_DISTINCT	sr320@washington.edu	Jul 2
Cgigas Larvae RNA-Seq OsHV UR10 RNA-seq data with descriptions of larvae exposed to OsHV. (>= 10 UniqueReads) <span style="background-color: #e0e0e0; padding: 2px;">oyster</span>	sr320@washington.edu	Jul 1
Cgigas Larvae RNA-Seq OsHV RNA-seq data with descriptions of larvae exposed to OsHV <span style="background-color: #e0e0e0; padding: 2px;">oshv</span>	sr320@washington.edu	Jul 1
Zhang_Mgo_gene_RNA-seq_IGV <span style="background-color: #e0e0e0; padding: 2px;">sperm</span>	sr320@washington.edu	Jun 2
Zhang_Gil_gene_RNA-seq_IGV IGV format <span style="background-color: #e0e0e0; padding: 2px;">rna-seq</span>	sr320@washington.edu	Jun 2
BiGill_methratio_Gene_Genomic_GFF GFF formatted file indicated DNA methylation on oyster genes <span style="background-color: #e0e0e0; padding: 2px;">qdod</span>	sr320@washington.edu	May
TJGR_GeneBased_CDS_GFF GFF format file with exons indicated for genes in oyster genome <span style="background-color: #e0e0e0; padding: 2px;">qdod</span>	sr320@washington.edu	May
BiGill_Gene_Methratio_VD	sr320@washington.edu	May
oyster_v9_mRNA GFF GFF (gene) from Zhang et al. Column9 modified for Joining <span style="background-color: #e0e0e0; padding: 2px;">qdod</span>	sr320@washington.edu	May
Cgigas gene length CDS only	sr320@washington.edu	Ma

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skyline 24  
oceanography 23  
ssgcid 18  
qdod2 18  
qdod 18  
swissprot 17  
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Name	Sharing / Owner	Modi
qdod_proteome_blast_mouse	🔒 sr320@washington.edu	Jan 2
qDOD_v9_gene GFF format file of oyster genes ~28k gene		Nov
_qdod_goslim_graphstest		Oct 2
SNP_RNAseqLibrary_SB_BiGill SNP table from RNA-seq library - SB qdod2		Oct 2
BiGill_meth_Zhang_exp Gene-centric data including length, CG, per qdod2		Oct 2
qDOD_Cgigas_gene_fasta Tabular format of Cgigas gene sequence qdod2		Oct 2
qDOD Cgigas Gene Descriptions (Swiss-prot) Description and evalu blast		Oct 2
file0	🔒 sr320@washington.edu	Aug 1
BiGill meth with SP	🔒 sr320@washington.edu	Aug 1
SPID and GO Numbers Swiss-Prot IDs and corresponding GO numbers qdod	🔒 sr320@washington.edu	Aug 1
Cgigas_larvae_RNAseq_OsHV_GO	🔒 sr320@washington.edu	Jul 2
qDOD_Cgigas_GO_GOslim_DISTINCT	🔒 sr320@washington.edu	Jul 2
Cgigas Larvae RNA-Seq OsHV UR10 RNA-seq data with descriptions of larvae exposed to OsHV. (>= 10 UniqueReads) oyster	🔒 sr320@washington.edu	Jul 1
Cgigas Larvae RNA-Seq OsHV RNA-seq data with descriptions of larvae exposed to OsHV	🔒 sr320@washington.edu	Jul 1
Zhang_Mgo_gene_RNA-seq_IGV	🔒 sr320@washington.edu	Jun 2
Zhang_Gil_gene_RNA-seq_IGV IGV format rna-seq	🔒 sr320@washington.edu	Jun 2
BiGill_methratio_Gene_Genomic_GFF GFF formatted file indicated DNA methylation on oyster genes qdod	🔒 sr320@washington.edu	May
TJGR_GeneBased_CDS_GFF GFF format file with exons indicated for genes in oyster genome qdod	🔒 sr320@washington.edu	May
BiGill_Gene_Methratio_VD	🔒 sr320@washington.edu	May
oyster_v9_mRNA GFF GFF (gene) from Zhang et al. Column9 modified for Joining qdod	🔒 sr320@washington.edu	May
Cgigas gene length CDS only	🔒 sr320@washington.edu	Ma

## Use Cases

- Joining on Annotations
- File Conversion
- Querying Gene Tables

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Name	Sharing / Owner	Modi
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BiGill_meth_Zhang_exp Gene-centric data including length, CG, per qdod2		Oct 2
qDOD_Cgigas_gene_fasta Tabular format of Cgigas gene sequence qdod2		Oct 2
qDOD Cgigas Gene Descriptions (Swiss-prot) Description and evalu blast		Oct 2

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- Joining on Annotations
- File Conversion
- Querying Gene Tables

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sr320@washington.edu	May
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## Predominant intragenic methylation is associated with gene expression characteristics in a bivalve mollusc

Mackenzie R. Gavery, Steven B. Roberts

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Zhang_Gil_gene_RNA-seq_IGV IGV format	rna-seq	
BiGill_methratio_Gene_Genomic_GFF GFF formatted file indicated DNA methylation on oyster genes	qdod	
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BiGill_Gene_Methratio_VD		
oyster_v9_mRNA GFF GFF (gene) from Zhang et al. Column9 modified for Joining	qdod	
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TJGR\_CCD\_d... 11

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qdod_proteome_blast_mouse	sr320@washington.edu	Jan 2
qDOD_v9_gene GFF format file of oyster genes ~28k gene	sr320@washington.edu	Nov
_qdod_goslim_graphstest	sr320@washington.edu	Oct 2
SNP_RNAseqLibrary_SB_BiGill SNP table from RNA-seq library - SB gill tissue pool (BiGill complement) qdod2	sr320@washington.edu	Oct 2
BiGill_meth_Zhang_exp Gene-centric data including length, CG, percent methylation (gill) and tissue specific RPKM data from Zhang et al. qdod2	sr320@washington.edu	Oct 2
qDOD_Cgigas_gene_fasta Tabular format of Cgigas gene sequence fasta file Derived using Dataset: Genomic data from the Pacific oyster qdod2	sr320@washington.edu	Oct 2
qDOD Cgigas Gene Descriptions (Swiss-prot) Description and evaluaes associated with 28k genes Derived using Dataset: Genomic data from the Pacific oyster blast	sr320@washington.edu	Oct 2

## Use Cases

- Joining on Annotations
- File Conversion
- Querying Gene Tables



Predominant intragenic methylation is associated with gene expression charac

## DNA methylation landscape of genomic features

In order to examine relationships between DNA methylation and genomic features, data from BSMAP (i.e., *methratio*) was converted to genomic feature tracks (i.e., generic feature format [GFF] files). Conversion was done using SQLShare (Howe et al., 2011), with the files and corresponding query language published (Gavery & Roberts, 2013).

The distribution of methylated CpGs with respect to specific genomic features was determined using BEDtools (i.e., *intersectBED*) (Quinlan & Hall, 2010). For this analysis, a CpG locus was considered methylated if at least half of the reads remained unconverted after bisulfite treatment. Genomic features that were examined include: exons and introns (Fang et al., 2012), putative promoters (defined as 1 kb upstream of open reading frames), and transposable elements. Putative transposable elements were

Gavery M, Roberts S. 2013. Crassostrea gigas high-throughput bisulfite sequencing (gill tissue). [figshare](#). Retrieved 18:44, Oct 21, 2013 (GMT)

sr320@washington.edu	Aug 3
sr320@washington.edu	Aug 3
sr320@washington.edu	Aug 3
sr320@washington.edu	Jul 2
sr320@washington.edu	Jul 2
sr320@washington.edu	Jul 1
sr320@washington.edu	Jul 1
sr320@washington.edu	Jun 2
sr320@washington.edu	Jun 2
sr320@washington.edu	May
sr320@washington.edu	May
sr320@washington.edu	May
sr320@washington.edu	May
sr320@washington.edu	May

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Name

qdod\_proteome\_blast\_mouse

qDOD\_v9\_gene GFF format file of oyster genes ~28k  
gene

\_qdod\_goslim\_graphstest

SNP\_RNAseqLibrary\_SB\_BiGill SNP table from RNA-  
qdod2BiGill\_meth\_Zhang\_exp Gene-centric data including  
qdod2qDOD\_Cgigas\_gene\_fasta Tabular format of Cgigas  
qdod2qDOD Cgigas Gene Descriptions (Swiss-prot) Descrip  
blast

BiGill\_CpG\_methylation.igv

preview

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BiGill\_exon\_clc\_rpk.igv

preview

download

BiGill\_igv\_charlie.xml

preview

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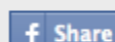
Query to derive\_CG\_AllData\_IGV.txt

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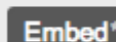
0



0



0



Cite this:

Gavery, Mackenzie; Roberts, Steven (2013): Crassostrea gigas high-throughput bisulfite sequencing (gill tissue). figshare.  
<http://dx.doi.org/10.6084/m9.figshare.749728>\*The embed functionality can only be used for non commercial purposes... [more](#)

## DNA methylation landscape of genomic features

In order to examine relationships between DNA methylation and genomic features, data from BSMAP (i.e., methratio) were converted to genomic feature tracks (i.e., generic feature format). Conversion was done using SQLShare (Howe et al., 2013) files and corresponding query language published (Howe et al., 2013).

The distribution of methylated CpGs with respect to genomic features was determined using BEDtools (Quinlan & Hall, 2010). For this analysis, a CpG was considered methylated if at least half of the reads were unmethylated after bisulfite treatment. Genomic features examined include: exons and introns (Fang et al., 2012), promoters (defined as 1 kb upstream of open reading frames), and transposable elements. Putative transposable elements

## Description

This fileset contains genomic feature tracks from methylation-enriched high-throughput bisulfite sequencing and RNA-seq analysis for Pacific oyster (*Crassostrea gigas*) gill tissue. Feature tracks were developed to be viewed with Integrative Genomics Viewer (<http://www.broadinstitute.org/igv/>) in conjunction with the *C. gigas* genome (Fang et al. 2012). All data and instructions are also available at <http://oystergen.es/bigill>.

File descriptions:

*BiGill\_CpG\_methylation.igv* - Location and proportion of methylation for all analyzed CpG dinucleotides with greater than 5x coverage.

*BiGill\_exon\_clc\_rpk.igv* - Exon-specific gene expression values (RPKM) from RNA-seq analysis.

*BiGill\_igv\_charlie.xml* - A session file, which loads methylation and RNA-seq feature tracks as well as the location of *C.gigas* genome features.

*Query to derive\_CG\_AllData\_IGV.txt* - Query (SQLShare) used to derive the methylation feature track from the original methratio output (<http://goo.gl/5LGq9Q>)

Reference:

Fang X, Li L, Luo R, Xu F, Wang X, Zhu Y, Yang L, Huang Z. 2012. Genomic data from the Pacific

207  
views0  
shares

Published on 20 Sep 2013 - 19:17 (GMT+8)  
Filesize in total is 92.16 MB

## Categories

- Physiology
- Marine Biology
- Bioinformatics
- Genetics
- Molecular Biology

## Authors

Mackenzie Gavery  
Steven Roberts

## Tags

- epigenetics
- Gene regulation
- invertebrate
- oyster
- dna methylation

## Export

- Export to Ref. Manager
- Export to Endnote
- Export to Mendeley



Your datasets	
All datasets	
Shared datasets	
Recent activity...	18
Recently viewed »	
Upload dataset	
New query	
YOUR TOP VIEWED	
qDOD Cgigas ...	18
BiGo_Larvae_j...	16
TJGR_CCD_d...	11
BiGill_RNAseq...	10

## Your Datasets

Filter dataset by keyword: 

Name	Sharing / Owner	Modi
qdod_proteome_blast_mouse	sr320@washington.edu	Jan 2
qDOD_v9_gene GFF format file of oyster genes ~28k gene	sr320@washington.edu	Nov
_qdod_goslim_graphstest	sr320@washington.edu	Oct 2
SNP_RNAseqLibrary_SB_BiGill SNP table from RNA-seq library - SB gill tissue pool (BiGill complement) qdod2	sr320@washington.edu	Oct 2
BiGill_meth_Zhang_exp Gene-centric data including length, CG, percent methylation (gill) and tissue specific RPKM data from Zhang et a qdod2	sr320@washington.edu	Oct 2
qDOD_Cgigas_gene_fasta Tabular format of Cgigas gene sequence fasta file Derived using Dataset: Genomic data from the Pacific oyst qdod2	sr320@washington.edu	Oct 2
qDOD Cgigas Gene Descriptions (Swiss-prot) Description and evaluaes associated with 28k genes. Derived using Dataset: Cgigas blast	sr320@washington.edu	Oct 2

## Use Cases

- Joining on Annotations
- File Conversion
- Querying Gene Tables

Secondary stress: proteomics

Original input file had some peptides of charge state >2, so had to redo everything with fixed input file.

SR discovered that for some proteins, a peptide was sequenced multiple times and so had multiple expression values. From the unique protein associations file in SQLshare, I summed the expression values for all identical peptides.

```
SELECT [peptide sequence], SUM([2_01 TotalArea]) AS CG2_01, SUM([2_02 TotalArea]) AS CG2_02, SUM([2_03
TotalArea]) AS CG2_03, SUM([5_01 TotalArea]) AS CG5_01, SUM([5_02 TotalArea]) AS CG5_02, SUM([5_03
TotalArea]) AS CG5_03, SUM([8_01 TotalArea]) AS CG8_01, SUM([8_02 TotalArea]) AS CG8_02, SUM([8_03
TotalArea]) AS CG8_03, SUM([11_01 TotalArea]) AS CG11_01, SUM([11_02 TotalArea]) AS CG11_02, SUM([11_03
TotalArea]) AS CG11_03, SUM([26_01 TotalArea]) AS CG26_01, SUM([26_02 TotalArea]) AS CG26_02, SUM([26_03
TotalArea]) AS CG26_03, SUM([29_01 TotalArea]) AS CG29_01, SUM([29_02 TotalArea]) AS CG29_02, SUM([29_03
TotalArea]) AS CG29_03, SUM([32_01 TotalArea]) AS CG32_01, SUM([32_02 TotalArea]) AS CG32_02, SUM([32_03
TotalArea]) AS CG32_03, SUM([35_01 TotalArea]) AS CG35_01, SUM([35_02 TotalArea]) AS CG35_02, SUM([35_03
TotalArea]) AS CG35_03, SUM([221_01 TotalArea]) AS CG221_01, SUM([221_02 TotalArea]) AS CG221_02,
SUM([221_03 TotalArea]) AS CG221_03, SUM([224_01 TotalArea]) AS CG224_01, SUM([224_02 TotalArea]) AS
CG224_02, SUM([224_03 TotalArea]) AS CG224_03, SUM([227_01 TotalArea]) AS CG227_01, SUM([227_02
TotalArea]) AS CG227_02, SUM([227_03 TotalArea]) AS CG227_03, SUM([230_01 TotalArea]) AS CG230_01,
SUM([230_02 TotalArea]) AS CG230_02, SUM([230_03 TotalArea]) AS CG230_03,
SUM([242_01 TotalArea]) AS CG242_01, SUM([242_02 TotalArea]) AS CG242_02, SUM([242_03 TotalArea]) AS
CG242_03, SUM([245_01 TotalArea]) AS CG245_01, SUM([245_02 TotalArea]) AS CG245_02, SUM([245_03
TotalArea]) AS CG245_03, SUM([248_01 TotalArea]) AS CG248_01, SUM([248_02 TotalArea]) AS CG248_02,
SUM([248_03 TotalArea]) AS CG248_03, SUM([251_01 TotalArea]) AS CG251_01, SUM([251_02 TotalArea]) AS
CG251_02, SUM([251_03 TotalArea]) AS CG251_03
```

i20@washington.edu	Aug 1
i20@washington.edu	Aug 1
i20@washington.edu	Aug 1
i20@washington.edu	Jul 2
i20@washington.edu	Jul 2
i20@washington.edu	Jul 1
i20@washington.edu	Jul 1
i20@washington.edu	Jun 2
i20@washington.edu	Jun 2
i20@washington.edu	May
i20@washington.edu	May
i20@washington.edu	May
i20@washington.edu	May
i20@washington.edu	May

# Visualize Dataset

**SQLSHARE** Graphs

SQL

[Share](#)

[Embed](#)

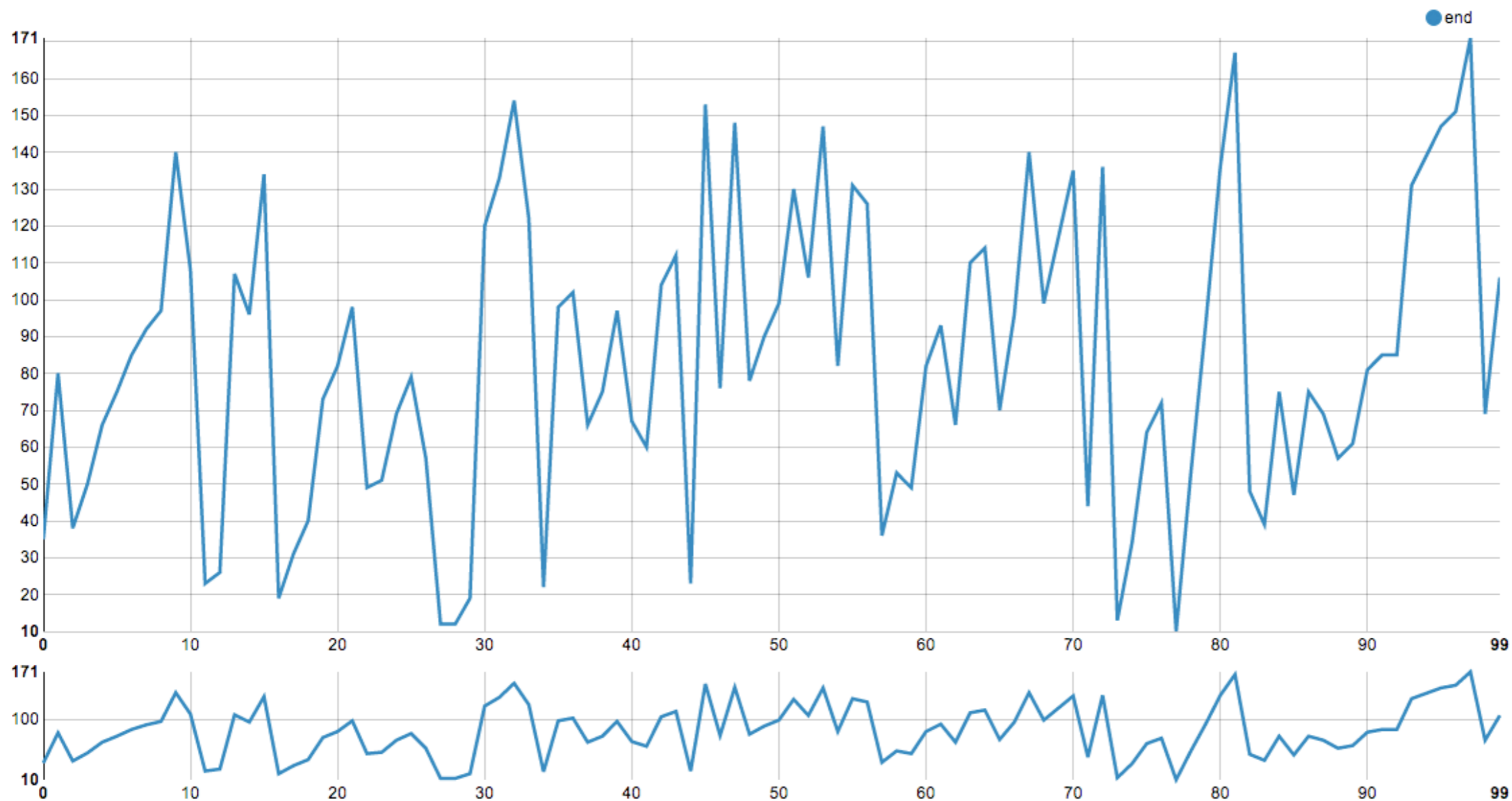
```
SELECT TOP 100 * FROM (SELECT * FROM [che625@washington.edu].[BiGO_MethylatedCGs_oysterv9]) x
```

[Click to edit](#)

## Data Analysis

Logged in: [sr320@washington.edu](#)

Graph [Share](#) [Embed](#)



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Notebooks

Rationale

Platforms

enScience

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anything else...

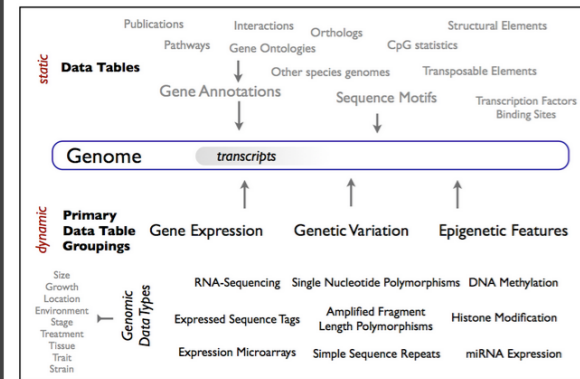




# qdod: Querying Disparate Oyster Datasets

This repository provides access to genomic data and workflows (IPython notebooks) that are being integrated as part of effort to increase efficiency of biological discovery. The [wiki](#) associated with this repository will serve as the *primary means for documentation*. Currently the documentation is focused on 1) **describing current datasets** and 2) **providing workflow tutorials**.

In brief, data in the form of delimited text files is aggregated into [SQLShare](#) where they can be easily queried. Below is schematic representation of the different types of datasets.



During the initial phases the focus is on the Pacific oyster and primary data from the [Roberts Lab](#).

## Select IPython Notebooks

- [Static Data Tables - Universal](#)
- [Static Data Tables - Annotations](#)

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Currently the documentation is focused on 1) **describing current datasets** and 2) **providing workflow tutorials**.

## A. Raw Data

- [Select NGS Data via Roberts Lab](#)

## B. Datasets in SQLShare

- [Universal](#)
- [Generic Oyster Datasets](#)
- [Tissue Specific Oyster Datasets](#)

## C. Tutorials

- [Simple Gene Search](#)
- [Standard SQLShare Queries](#)
- [Annotating Genes](#)
- [File Format Conversions](#)

## D. Genome Browser Feature Tracks

- [Canonical Tracks](#)
- [Bisulfite sequencing \(gill tissue\)](#)
- [Reference Genome Files](#)

Please use [GitHub's Issue feature](#) to ask question, report problems, or suggest features.

Last edited by sr320, 9 days ago

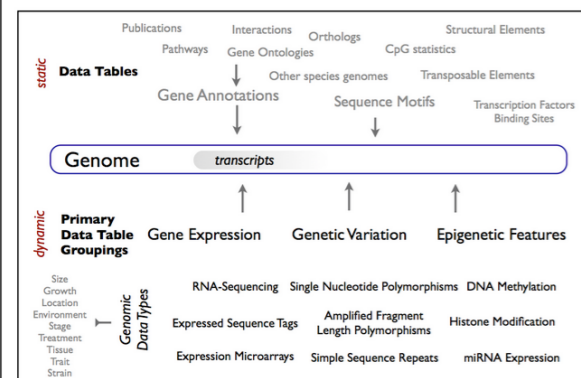


[github.com/sr320/qdod/wiki](https://github.com/sr320/qdod/wiki)

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## Select Genomic Data

ID	Platform	Molecule	Tissue	Length	Files
BB3	SOLiD	RNA	gill	25 x 1	<a href="#">csfasta; qual</a>
DH3	SOLiD	RNA	gill	25 x 1	<a href="#">csfasta; qual</a>
DH2	SOLiD	RNA	gill	25 x 1	<a href="#">csfasta; qual</a>
GE	SOLiD	RNA	larvae	50 x 1	<a href="#">csfasta; qual</a>
GC	SOLiD	RNA	larvae	50 x 1	<a href="#">csfasta; qual</a>
SBunmeth	SOLiD	DNA	gill	25 x 1	<a href="#">csfasta; qual</a>
SBmeth	SOLiD	DNA	gill	25 x 1	<a href="#">csfasta; qual</a>
BSseqGill	Illumina	DNA	gill	36 x 1	<a href="#">fastq</a>
ETStageseq	Illumina	RNA	gill		<a href="#">zip</a>
BSseqSperm	Illumina	DNA	sperm	72 x 2	<a href="#">fastq1; fastq2</a>
BiGillRNA	Illumina	RNA	gill	50 x 2	<a href="#">fastq1; fastq2</a>
BiGoRNA	Illumina	RNA	sperm	50 x 2	<a href="#">fastq1; fastq2</a>

Currently the documentation is focused on

### A. Raw Data

- [Select NGS Data via Roberts Lab](#)

### B. Datasets in SQLShare

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- [Tissue Specific Oyster Datasets](#)

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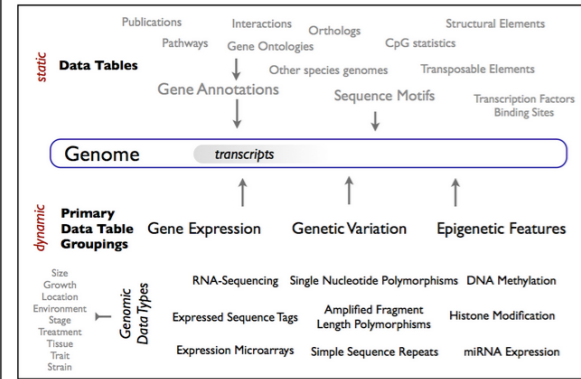
everything else...



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DH3	SOLiD	RNA	gill	25 x 1	<a href="#">csfasta; qual</a>

## Data Snapshots

Select datasets available from SQLShare. Tag: `qdod2`  
<https://sqlshare.escience.washington.edu/sqlshare/#s=tag/qdod2>

## Universal

Dataset	Screenshot	more
<a href="#">UniprotProtNamesReviewed_yes20130610</a>		etc
<a href="#">SPID and GO Numbers</a>		etc

Currently the documentation is focused on

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### B. Datasets in SQLShare

- [Universal](#)
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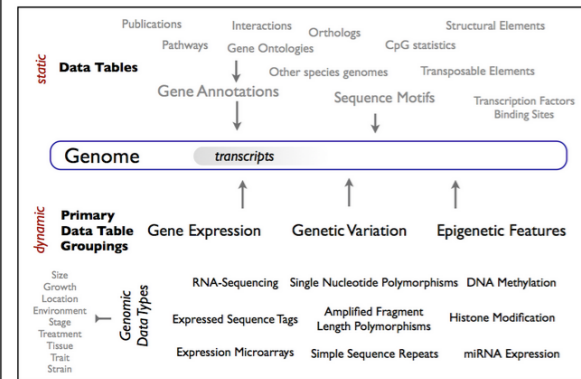
everything else...



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- [Reference Genome Files](#)

Please use [GitHub's Issue feature](#) to ask

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BB3	SOLID	RNA	gill	25 x 1	<a href="#">csfasta; qual</a>
DH3	SOLID	RNA	gill	25 x 1	<a href="#">csfasta; qual</a>

## Data Snapshots

Select datasets available from SQLShare. Tag: [qdod2](#)  
<https://sqlshare.escience.washington.edu/sqlshare/#s=tag/qdod2>

## Universal

## Workflow 1: Annotating Oyster Genes

This workflow will take focus on taking a simple SQLShare table that has gene IDs and associated expression data and will take you through the steps of figuring out the name, function, etc of each gene.

### Initial Data Table: Oyster larvae RNA-seq - OshV exposure

SCREENSHOT

[solid0078\\_20091105\\_RobertsLab\\_GE\\_F3 trimmed RNA-Seq.txt](#) Viewable by everyone

OshV RNA-seq on Version 9 transcriptome

[Click here to add a tag](#)

```
SELECT
["Feature ID"] as ID,
["Unique gene reads"] as UniqueReads,
["Total gene reads"] as TotalReads,
["RPKM"] as RPKM
FROM [sr320@washington.edu].[table_solid0078_20091105_RobertsLab_GE_F3 trim
```

DATASET PREVIEW Rows 1 - 100 of 28027 | Columns 4 of 4

ID	UniqueReads	TotalReads	RPKM
CGI_10000001	0	10	5.23
CGI_10000002	5	5	2.756
CGI_10000003	0	0	0
CGI_10000004	0	0	0
CGI_10000005	0	0	0

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## The iPlant Collaborative

The iPlant Collaborative develops cyberinfrastructure and computational tools to solve Grand Challenges in plant science

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## The iPlant Collaborative

The iPlant Collaborative develops cyberinfrastructure and computational tools to solve Grand Challenges in plant science

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Environment

Molecular

**Data Analysis**

### Discovery Environment



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# Data Analysis

The screenshot displays the Discovery Environment interface, which is used for data analysis. It features a teal header with the 'Discovery Environment' logo and a notification badge showing '39'. The interface is divided into three main panels:

- Data Panel:** Shows a file browser with a navigation pane on the left containing folders like 'labshare', 'qdod', 'BiGo\_larvae', and 'Cgigas\_v9'. The main pane lists files with columns for Name, Last Modified, and Details.
- Apps Panel:** Displays a list of applications categorized into 'Workspace (8)', 'Public Apps (434)', and 'General Utilities (73)'. The 'Workspace' section lists apps like 'Uncompress files wi...', 'RNAseq2bedgraph', and 'FastQC 0.10.1 (mul...)' with their respective integrators and ratings.
- Analyses Panel:** Shows a table of completed and failed analyses. The table has columns for Name, App, Start Date, End Date, and Status.

Name	App	Start Date	End Date	Status
genomeCoverageBed_Bi...	genomeCov...	2014 Feb 15 05:01:...	2014 Feb 15 05:11:...	Completed
genomeCoverageBed_Bi...	genomeCov...	2014 Feb 15 04:14:...	2014 Feb 15 04:18:...	Completed
RNAseq2bedgraph_anal...	RNAseq2be...	2014 Feb 15 04:07:...	2014 Feb 15 09:55:...	Completed
RNAseq2bedgraph SE_BB3	RNAseq2be... SE	2014 Feb 12 14:49:...	2014 Feb 12 14:49:...	Failed
RNAseq2bedgraph_BiGo...	RNAseq2be...	2014 Feb 12 14:36:...	2014 Feb 12 20:00:...	Completed
genomeCoverageBed_a...	genomeCov...	2014 Feb 12 11:22:...	2014 Feb 12 11:28:...	Completed
bamToBed_analysis1	bamToBed	2014 Feb 12 07:50:...	2014 Feb 12 07:54:...	Completed
VCF to GFF3_analysis1	VCF to GFF3	2014 Feb 11 06:58:...	2014 Feb 11 06:59:...	Completed
Find SNPS - mpileup_analysis1	Find SNPS - mpileup	2014 Feb 11 04:13:...	2014 Feb 11 06:37:...	Completed
TopHat2-PE_analysis1	TopHat2-PE	2014 Feb 10 17:23:...	2014 Feb 10 21:38:...	Completed

# Data Analysis



The screenshot displays the Discovery Environment interface, which is used for data analysis. It features several panels:

- Data Panel:** Shows a file browser with a navigation pane on the left and a main pane on the right. The navigation pane shows a folder structure under 'sr320', including 'labshare', 'qdod', 'BiGo\_larvae', and 'Cgigas\_v9'. The main pane shows a list of files with columns for Name, Last Modified, and other details.
- Apps Panel:** Shows a list of applications available in the workspace. The 'Workspace' section includes 'Apps under development (6)', 'Favorite Apps (2)', 'My public apps (0)', and 'Public Apps (434)'. The 'Public Apps' section lists applications like 'Uncompress files wi...', 'RNAseq2bedgraph', 'RNAseq2bedgraph SE', and 'FastQC 0.10.1 (mul...)'.
- RNAseq2bedgraph Panel:** A detailed view of the 'RNAseq2bedgraph' application. It shows the analysis name 'RNAseq2bedgraph\_analysis1' and provides instructions for running the application, including selecting a reference genome and providing a reference genome file in FASTA format.



Username:   
 Password:   
 Log In Enter As Guest  
 Forgot Password? Register

**D N A**

**S U B W A Y**

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Select an Image

Search by name, emi, tag, d

Featured Images

- iPlant Centos 6.5 NoGUI Base  
2436bf2f-13a7-4118-a349-d3529f79ae16  
centos centos6  
nodesktop  
nogul base
- Ubuntu 12.04 - iPlant Base  
75fdca4-d49d-4b2d-b919-a3297bc6d7ae
- NGS Viewers v3.2 (03-15-2013)  
ac8e0cfb-5f0e-4c63-bcf9-525356304c45  
NGS
- IBP Workbench v5

My Projected Resource Usage

Request More Resources

0% You are using 0 of 168 allotted AUs.

5% You will use 1 of 16 allotted CPUs.

3% You will use 4 of 128 allotted GBs.

Configuration

Launch Instance

Select Image

First, select an image on the left. Then you can re-name it, change its size, or tag it.

Name

Description

Instance Size

- tiny1 (1 CPUs, 4 GB memory, 30 GB disk)
- tiny2 (1 CPUs, 8 GB memory, 60 GB disk)
- small1 (2 CPUs, 8 GB memory, 60 GB disk)
- small2 (2 CPUs, 16 GB memory, 120 GB disk)
- medium1 (4 CPUs, 8 GB memory, 80 GB disk)
- medium2 (4 CPUs, 16 GB memory, 160 GB disk)
- medium3 (4 CPUs, 32 GB memory, 240 GB disk)
- large1 (8 CPUs, 16 GB memory, 160 GB disk)
- large2 (8 CPUs, 48 GB memory, 320 GB disk)
- large3 (8 CPUs, 64 GB memory, 480 GB disk)
- ultra1 (16 CPUs, 32 GB memory, 400 GB disk)

Tools Load Data

search tools

Get Data  
Send Data  
Lift-Over  
Text Manipulation  
Convert Formats  
FASTA manipulation  
Filter and Sort  
Join, Subtract and Group  
Extract Features  
Fetch Sequences  
Extract Genomic DNA using coordinates from assembled/unassembled genomes  
Fetch Alignments  
Get Genomic Scores  
Operate on Genomic Intervals  
Statistics  
Graph/Display Data  
Regional Variation  
Multiple regression  
Multivariate Analysis  
Evolution  
Motif Tools  
Multiple Alignments  
Metagenomic analyses  
Genome Diversity  
NGS TOOLBOX BETA  
Phenotype Association  
NGS: QC and manipulation  
NGS: Mapping  
Bowtie2 is a short-read aligner  
Lastz map short reads against reference sequence  
Map with Bowtie for SOLID

Map with Bowtie for SOLID (version 1.1.2)

Will you select a reference genome from your history or use a built-in index?:  
   
 Built-ins were indexed using default options

Select the reference genome:  
   
 If your genome of interest is not listed - contact Galaxy team

Is this library mate-paired?:

FASTQ file:  
   
 Must have Sanger-scaled quality values with ASCII offset 33

Bowtie settings to use:  
   
 For most mapping needs use Commonly used settings. If you want full control use Full parameter list

Suppress the header in the output SAM file:  
   
 Bowtie produces SAM with several lines of header information by default

**What it does**  
 Bowtie is a short read aligner designed to be ultrafast and memory-efficient. It is developed by Ben Langmead and Cole Trapnell. Please cite: Langmead B, Trapnell C, Pop M, Salzberg SL. Ultrafast and memory-efficient alignment of short DNA sequences to the human genome. Genome Biology 10:R25.

**Know what you are doing**  
 ⚠ There is no such thing (yet) as an automated gearshift in short read mapping. It is all like stick-shift driving in San Francisco. In other words = running this tool with default parameters will probably not give you meaningful results. A way to deal with this is to **understand** the parameters by carefully reading the [documentation](#) and experimenting. Fortunately, Galaxy makes experimenting easy.

**Input formats**  
 Bowtie accepts files in Sanger FASTQ format. Use the FASTQ Groomer to prepare your files.

History

Ensembl  
735.8 MB

3: Crassostrea gigas.GC  
A\_000297895.1.21.gff3

2: Crassostrea gigas.GC  
A\_000297895.1.21.gtf

~420,000 lines  
format: gtf, database: cgigas\_E\_sm

uploaded gtf file

1. Seqname	2. Source	3. Feature
scaffold39040	protein_coding	exon
scaffold39040	protein_coding	CDS
scaffold39040	protein_coding	start_cod
scaffold39040	protein_coding	exon
scaffold39040	protein_coding	CDS
scaffold39040	protein_coding	exon

1: Crassostrea gigas.GC  
A\_000297895.1.21.dna\_sm.genome.f  
a

7,658 sequences  
format: fasta, database: cgigas\_alpha\_v012

uploaded fasta file

```
>C12728 dna_sm:supercontig supercontig
TTTTTCTTATTTTATTTGTACCAGTTAATCCTGTGAg
ccggtttttatgtctctcttctgTTGACTTTATTATTCTC
TATGAATTATTATGTATCTTCATGGGATATTTTGC
ACACTTGTCTTTTCGGTGTAT
>C12722 dna_sm:supercontig supercontig
```

Biology  
 Environment  
 Molecular  
**Data Analysis**  
 eScience  
 iPlant Galaxy  
 Notebooks  
 Rationale  
 Platforms  
**Open Science**  
 Data  
 everything else...

**Galaxy** Analyze Data Workflow **Shared Data** Visualization Cloud Help User Using 13%

Published Workflows | [rna-seq-helin-group](#) | RNA-seq differential analysis (reads, 2 conditions, 2 replicates) +

**Galaxy Workflow 'RNA-seq differential analysis (reads, 2 conditions, 2 replicates)'**  
 Annotation: Workflow based on Tophat and cuffdiff. Inputs: 4 fastq files (example), 1 gtf file (annotations).  
 Outputs: bam, bigwig, xls, and html reports with quality controls.

**Step** **Annotation**

Step 1: Input dataset  
**Gene Annotations**  
*select at runtime*

Step 2: Input dataset  
**Group1 R1**  
*select at runtime*

Step 3: Input dataset  
**Group1 R2**  
*select at runtime*

Step 4: Input dataset  
**Group2 R1**  
*select at runtime*

Step 5: Input dataset  
**Group2 R2**  
*select at runtime*

Step 6: Input dataset  
**Pseudo reads BAM file**  
*select at runtime*

Step 7: Tophat for Illumina  
**RNA-Seq FASTQ file**  
 Output dataset 'output' from step 2  
**Use a built in reference genome or own from your history**  
 Use a built-in genome  
<https://usegalaxy.org/library/index>

**Access published resources**  
 Data Libraries Beta  
 Published Histories  
 Published Workflows  
 Published Visualizations  
 Published Pages

**About this Workflow**

**Author**  
 rna-seq-helin-group

**Related Workflows**  
[All published workflows](#)  
[Published workflows by rna-seq-helin-group](#)

**Rating**  
 Community (3 ratings, 5.0 average) ★★★★★  
 Yours ★★★★★

**Tags**  
 Community: illumina cuffdiff rnaseq tophat  
 Yours: +

Reads computationally generated from genome sequence to avoid zero coverage genes.

Biology

Environment

Molecular

**Data Analysis**

eScience

iPlant Galaxy

Notebooks

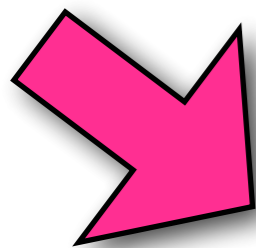
Rationale

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everything else...



**SQLSHARE**



**Hyak**



Biology

Environment

Molecular

**Data Analysis**

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iPlant Galaxy

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everything else...



SQLSHARE

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SQLSHARE

Galaxy

iPlant

BiGoRNA_genomecov.bedgraph	Yesterday, 4:41 AM	209.8 MB
BiGo_egg_redo4.sam	Feb 13, 2014, 5:47 PM	278.2 MB
BiGo_egg_redo3.sam	Feb 13, 2014, 5:26 PM	98.4 MB
solid0078_20091105_BB3 mapping.bam	Feb 13, 2014, 1:18 PM	1.46 GB
solid0078_20091105_BB3 mapping.sam	Feb 13, 2014, 1:15 PM	4.99 GB
solid0078_20091105_BB3 RNA-Seq.bam	Feb 13, 2014, 1:14 PM	706.3 MB
solid0078_20091105_BB3 RNA-Seq.sam	Feb 13, 2014, 1:13 PM	2.65 GB
BiGoRNA_bg1_sorted.bedgraph	Feb 12, 2014, 3:31 PM	269.8 MB
cgigasgenome3.txt	Feb 12, 2014, 3:26 PM	117 KB
solid0078_20091105_BB3.1.fastq	Feb 12, 2014, 11:49 AM	4.46 GB
BiGoRNA_GTGTCTAC_1.sorted.bam.bai	Feb 12, 2014, 9:43 AM	1.6 MB
BiGoRNA_bg1.bedgraph	Feb 12, 2014, 9:31 AM	269.8 MB
cgigasgenome3.tab	Feb 12, 2014, 8:33 AM	117 KB
cgigasgenome2.tab	Feb 12, 2014, 8:31 AM	558.4 MB
cgigasgenome1.tab	Feb 12, 2014, 8:31 AM	558.3 MB
BiGoRNA_GTGTCTAC_1.sorted.bam	Feb 12, 2014, 8:29 AM	1.42 GB
igv_session_cgigas_ensembl_1.xml	Feb 12, 2014, 8:28 AM	2 KB
BiGoRNA_GTGTCTAC_1.bam	Feb 12, 2014, 8:16 AM	1.42 GB
filtered_106A_Female_Mix_GATCAG_L004_R1.bam.fai	Feb 11, 2014, 8:52 AM	Zero bytes
filtered_106A_Female_Mix_GATCAG_L004_R1.bam.bai	Feb 11, 2014, 4:35 AM	9.1 MB
filtered_106A_Female_Mix_GATCAG_L004_R1.bam	Feb 11, 2014, 4:24 AM	8.58 GB
fish546_of_BowtieOut.sam	Feb 10, 2014, 1:32 PM	23.98 GB
fish546_of_BowtieOut.sorted.sam	Feb 10, 2014, 12:56 PM	3.7 MB
fish546_of_BowtieOut.sorted.sam.sai	Feb 10, 2014, 12:53 PM	4 bytes
fish546_14_Trinity.fasta.fai	Feb 10, 2014, 12:37 PM	4.3 MB
fishfe6_oCD-HITout.fa	Feb 10, 2014, 12:02 PM	88.1 MB
Emma_Defense_Talk.mp3	Feb 10, 2014, 6:19 AM	112.3 MB
fish546_14_Trinity_ls_length2.tab	Feb 9, 2014, 12:53 PM	163.3 MB
fish546_14_Trinity_ls_length.tab	Feb 9, 2014, 12:26 PM	162.9 MB
fish546_14_Trinity.tab	Feb 9, 2014, 12:16 PM	162.8 MB
data.csv	Feb 9, 2014, 10:25 AM	121 bytes
fish546_14_Trinity.fasta	Feb 9, 2014, 10:24 AM	165.1 MB
282014Emma_Defense_Talk.m4a	Feb 8, 2014, 2:00 PM	112.1 MB
f546_bubREVIGO.r	Feb 8, 2014, 10:46 AM	7 KB
f546REVIGO_treemap.r	Feb 8, 2014, 10:45 AM	6 KB
f546REVIGO.xqmm1	Feb 8, 2014, 10:35 AM	30 KB

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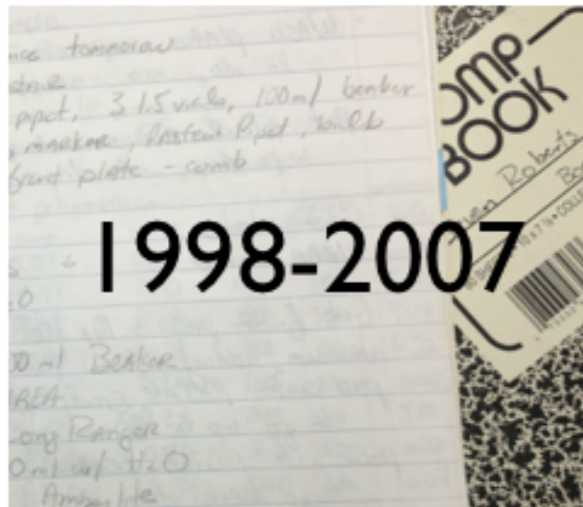
Platforms

Open Science

Data

everything else...

# The Evolution of My Lab Notebook



1998-2007



2008-2009



2010-2013



now

# Open Notebook Science

... there is a URL to a laboratory notebook that is freely available and indexed on common search engines. It does not necessarily have to look like a paper notebook but it is essential that all of the information available to the researchers to make their conclusions is equally available to the rest of the world.

—Jean-Claude Bradley

**Biology**

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# Open Notebook Science

**Biology**

Environment

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**Data Analysis**

eScience

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# Open Notebook Science

UsefulChem

☆ All Reactions Edit 2 2



**List of all experiments**

**NOTE: All solubility experiments by the Bradley group past this point are r challenge wiki.**

[EXP286](#) Synthesis of aldol condensation product of phenanthrene-9-carboxaldehyde and NaOH as the catalyst. -**Matthew McBride**

[EXP285](#) Synthesis of trans-dibenzalacetone in ethanol/water (1:1 vol) using NaOH as th

[EXP284](#) Synthesis of trans-dibenzalacetone in ethanol/water (1:1 vol) using NaOH as th

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## IheartAnthony's Research

Do you?

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## Experiments

### Active Experiments

#### Repeating Crumley

Experiment Introduction: Steve asked me to try and repeat an experiment by

# Open Notebook Science

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- on ChemSpider
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- ferences
- periment Format
- tra Credit
- ner03 Draft

## ☆ All Reactions

Edit 2 2



### List of all experiments

**NOTE: All solubility experiments by the Bradley group past this point are r challenge wiki.**

[EXP286](#) Synthesis of aldol condensation product of phenanthrene-9-carboxaldehyde and NaOH as the catalyst. -**Matthew McBride**

[EXP285](#) Synthesis of trans-dibenzalacetone in ethanol/water (1:1 vol) using NaOH as th

[EXP284](#) Synthesis of trans-dibenzalacetone in ethanol/water (1:1 vol) using NaOH as th

## IheartAnthony's Research

Do you?

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OpenWetWare  
Share your science.

### navigation

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- Add a Lab Notebook

### research

- Materials
- Protocols
- Resources

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## User:Andy Maloney/Notebook/Lab Notebook of Andy Maloney/Table of Contents/Table of Contents Category View

< User:Andy Maloney | Notebook | Lab Notebook of Andy Maloney | Table of Contents

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- 2 DNA unzipping
- 3 Kinesin & Microtubules
- 4 Wet lab install
- 5 Instruction Pages
- 6 Meetings & Questions
- 7 Classes
- 8 Sketchup pages

## Optical tweezers pages

### Active Experiments

### Repeating Crumley

Experiment Introduction: Steve asked me to try and repeat an experiment by

# Open Notebook Science

## Lab Notebook

(Introduction)

### 📡 Entries

#### Notes

31 Jan 2014

pageviews: (not calculated)

Scratch notes on uncertainty.

Function definition: [multiple\\_uncertainty.m](#)

Settings:

```
f=@(x, h) max( (x-h) * (1 - (x-h) ./ 100)
+ (x-h), 0);
x_grid = [0:5:100];
h_grid = x_grid; % Must be same dimension
s as x_grid, or L91 errors...
Tmax = 5;
sigma q = 0.1;
```

[Read more](#)

#### Notes

06 Jan 2014

pageviews: 3

- fishbase API discussions
- knitcitations (push previous updates to CRAN)
- data extraction from sardine collapse story (see EML in data/, [news article](#))

#### Notes

09 Jan 2014

pageviews: 8

- fishbase API discussions continue
- some discussion / work on rfigshare continues (As Karthik writes comprehensive test suite, yay).
- Possible ontological term for simulated data? modeling and simulation operation looking for example using this term now...
- [Request for term on OBI](#)
- Moore DDD investigators, full call

.@NOAA gives a glimpse of what the future of [Read more](#)

#### Open Science Literature Highlights

02 Jan 2014

pageviews: 40

A student recently asked me for some recommendations for an article on an open science topic for a journal club. Since I haven't jotted these down in one place before, though I might copy my reply here for

### Carl Boettiger

Theoretical Ecology and Evolution

I'm Carl Boettiger, a post-doctoral researcher with Marc Mangel and Steve Munch in the Center for Stock Assessment Research, at UC Santa Cruz, working on regime shifts in ecology and evolution. Such shifts mark the most dramatic events in complex systems. What causes these transitions to occur? Can we detect or forecast these changes? How do we robustly manage ecosystems that experience such tipping points? Understanding and forecasting these events challenges the capacity of both our statistical methods and the data available. My research aims to extend and quantify the limits of understanding through nonparametric Bayesian inference, high performance computing and big data



#### Notes

07 Jan 2014

pageviews: 6

#### Coding

- fishbase API discussions
- rfishbase updates pushed to CRAN
- Merges and minor bug fixes for rfigshare
- RNeXML: After some cooperative sleuth-work, we successfully resolved issue #23, uncovering a bug (missing feature really) in xmlint.

as usual, detailed activity on github. [Read more](#)

#### Notes

10 Dec 2013

pageviews: 64

Consider the model

$$dX_t = \alpha(\theta - X_t)dt + \sigma dB_t \quad (1)$$

carlboettiger.info/lab-notebook

# Open Notebook Science



genefish.wikispaces.com

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Search



## Lab Notebooks

Sam's Notebook  
Mac's Notebook  
Emma's Notebook  
Claire's Notebook  
Steven's Notebook  
Halley's Notebook  
Katie's Notebook  
Brent's Notebook  
Doug's Notebook  
Charles' Notebook  
Jessica's Notebook  
Hannah's Notebook  
Jake's Notebook  
Yanouk's Notebook

Featured Pages  
crassostreome

## home

Edit 0 220 ...

This wiki has been developed as a resource for lab personnel and students to access information and publish research activities using an [open notebook science](#) based system. All lab notebooks can be accessed via the side menu. The Roberts Lab is in the [School of Aquatic and Fishery Sciences](#) within the [College of Environment](#) at the [University of Washington](#). More information can be found concerning [research](#), [personnel](#), and [outreach](#) on the [Roberts Lab Official Webpage](#).

## Laboratory Reference Material

- [Data and Resource Sharing Plan](#)
- [Laboratory Protocols](#)
- [Emergency Contact Information](#)
- [UW Lab Safety Manual](#)
- [Code Repository](#)
- [UW Biosafety Manual](#)
- [Chemical Inventory](#)

## Lab Activity and Communication

- [Lab Meetings](#)
- [IPUS: Information for Prospective Undergraduate Students](#)
- [Lab Calendar](#)

## Data Repositories

- [The Eagle](#)
- [CLC Genomics Server](#) (password protected)
- [Primer Database](#)
- [NGS Library Info](#)
- [crassostreome](#)
- [UW HTSEQ](#) (gyccaiMS)

The screenshot shows the Roberts Lab Website with a navigation menu at the top: People, Research, Products, Outreach, Courses, Contact. Below the menu is a banner for Lab Notebooks with the text: "Why do you put your lab notebook and data online? The usual follow up to that question is 'what if someone steals your research?' I will not disagree this is a possibility. Others are concerned with intellectual property and I respect persons have their...". Below the banner are three columns of text: Research, Outreach, and News and Notes. The Research section describes the lab's focus on characterizing physiological responses of marine organisms to environmental change. The Outreach section describes the lab's efforts to engage with students, scientists, and the general public. The News and Notes section lists recent events, including a Student Spotlight Award at Aquaculture 2013 and a new website launch. At the bottom of the screenshot are social media links for Labcam, Connect, Access, and Funding.

# Open Notebook Science



- Wiki Home
- Projects
- Recent Changes
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- Members



## Lab Notebooks

- Sam's Notebook
- Mac's Notebook
- Emma's Notebook
- Claire's Notebook
- Steven's Notebook
- Halley's Notebook
- Katie's Notebook
- Brent's Notebook
- Doug's Notebook
- Charles' Notebook
- Jessica's Notebook
- Hannah's Notebook
- Jake's Notebook
- Yanouk's Notebook

## Katie's Notebook

Edit 0 75

**February 4, 2014**

This weekend, I downloaded Colony onto Thomas's computer (since it is Windows). I have been running the 2011 data on it since Saturday afternoon. As of this morning, it was making some pretty good progress. I will post the results on here when they are done! Tonight, I am going to download "HP-Rare" onto his computer so I can look at the rarefaction curve of this data. I am also working on organizing the south sound data from David Stick so that I can analyze that and compare it to the data I am working on now.

I made a Public Folder in my dropbox where I will keep all of the Microsat data so that there is a live file anyone can look at. Here is the link:

[https://www.dropbox.com/sh/5839mey4mqn2y28/cydXmRe\\_zd](https://www.dropbox.com/sh/5839mey4mqn2y28/cydXmRe_zd)

So far, there are the allele frequencies from David Stick's south sound wild Oly's and another file which will have the GenAIEx relatedness values.

I am working on Excel with this data and doing the same things that I have already done to the restoration grade data. (I will also put this through Colony when I get the other files through).

I also started running the 2010 restoration grade oly data through Colony downstairs in Jake's office.

**January 30, 2014**

In FSH 207 running Colony. I \*think\* I am now about 30% of the way through with the 2011 data.

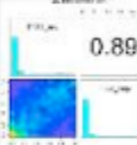
 **Oly.MidResult5**  
[Details](#) [Download](#) 1 KB

**January 29, 2014**

# Open Notebook Science


Search

**che625's notebook**


**Combining technical replicates for methylkit**  
Yesterday Female correlation file.list <-  
list('/Volumes/web/Mollusk/bs\_larvae\_exp/methylkit\_files/round2/CgF\_nov.txt', '/Volumes/web/Mollusk/bs\_...  


**methylkit troubleshooting**  
Yesterday Used Giles' code for formatting my files for methyl kit. Also used the R code listed below but kept getting errors. Will keep troubleshooting and try re-formatting my files. Used the files listed in

**Query for using methylkit**  
Yesterday SELECT chr as chr, pos as start, '+' as strand, cast (CT\_count as float) as CT\_count, cast (C\_count as float) as C\_count, cast (C\_count as float) / cast (CT\_count as float) as

**methyl kit with larvae data**  
Yesterday library(methylKit) library(data.table) library(GenomicRanges) file.list <- list('/Volumes/web/Mollusk/bs\_larvae\_exp/methylkit\_...  



**Filtering Larvae data**  
6 days ago Ran all files through BSMAP and methratio. Now need to filter so context= CG and a coverage cutoff of 5x or greater. Here is the query I am using to filter the data, and am changing the file

**Bioinformatics 2014- Module 6**  
6 days ago See Module 1 for De novo assembly with Trinity using oyster gonad RNA-Seq data: <https://www.evernote.com/shard/s232/sh/0596b21b-...>  


**Untitled Note**  
Last week For next figure look at this file: /Volumes/web/cnidarian/BiGo\_gene\_pmeth\_expression.csv  
Porzio<- read.csv('/Users/claireolson/Desktop/Porzio\ et\ al\ 2011\

**Beautiful graphics in R- experimenting with plotly**  
Last week # Required packages from CRAN .pkgs = c("RJSONIO", "RCurl", "devtools") # Install required packages from CRAN (if not) .inst <- .pkgs %in% installed.packages()

**Figure 1 for beautiful graphics in R**  
Last week With labels: Exon<- read.table('/Volumes/web/Mollusk/174gm\_analysis/Whole\_Genome\_Analyses/intersectBedfiles\_Steven

**Bioinformatics 2014- Module 4**  
Last week My file of p-values for all genes in gill v gonad: [http://eagle.fish.washington.edu/Mollusk/FIS\\_H546\\_2014/DESeq%20results%20for%20Tissue%2](http://eagle.fish.washington.edu/Mollusk/FIS_H546_2014/DESeq%20results%20for%20Tissue%2)  


che625's notebook

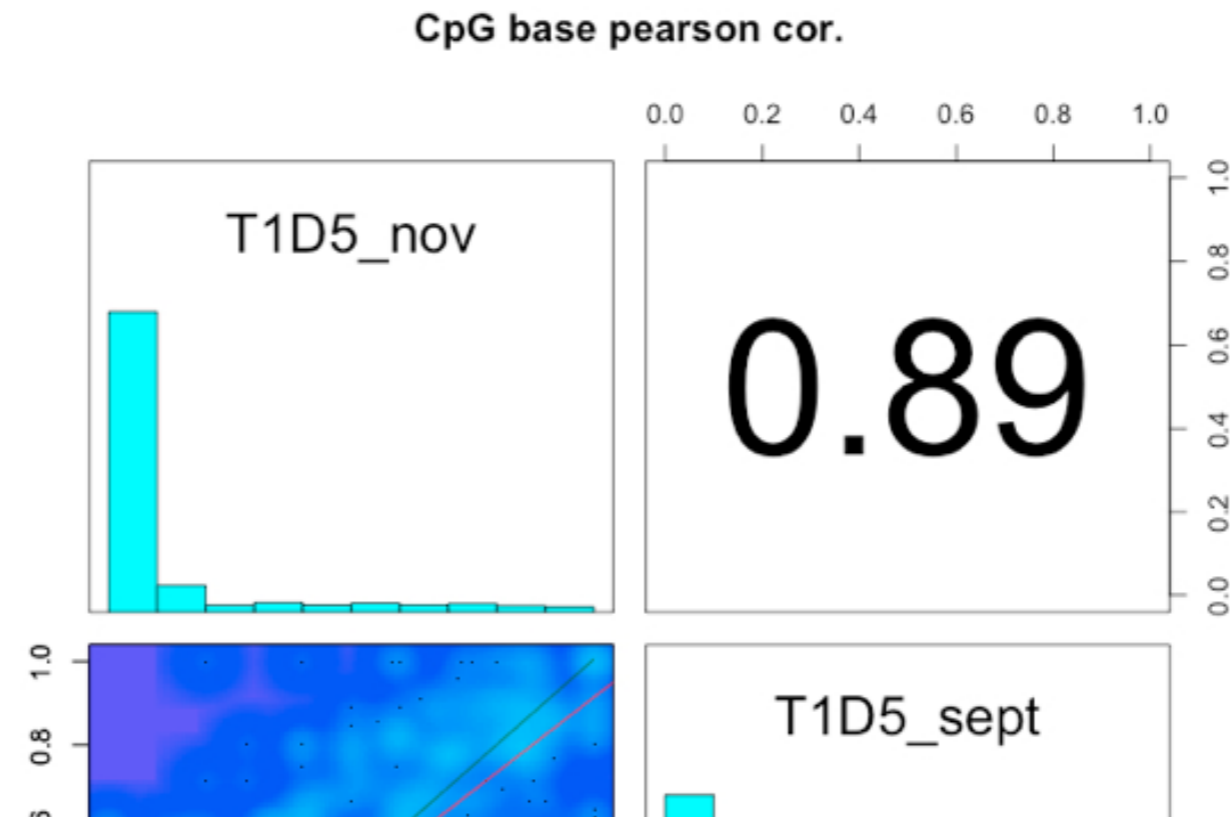
Created: Feb 18, 2014 Modified: Feb 18, 2014

### Larvae T1D5 correlation

```
file.list <- list('/Volumes/web/Mollusk/bs_larvae_exp/methylkit_files/round2/CgLarv_T1D5_nov.txt',  
'/Volumes/web/Mollusk/bs_larvae_exp/methylkit_files/round2/CgLarv_T1D5_sept.txt')  
  
myobj<-read(file.list, pipeline=list(fraction=TRUE, header=T, chr.col=1, start.col=2, end.col=3, strand.col=4, coverage.col=freqT.col=7),  
            sample.id=list("T1D5_nov", "T1D5_sept"), assembly="v9", treatment=c(1,0))  
  
meth<-unite(myobj, destrand=FALSE)  
  
head(meth)  
  
nrow(meth)  
  
getCorrelation(meth, plot=T)
```

**2373 loci make up this plot:**

**CpG base pearson cor.**



evernote.com/pub/che625/che625snotebook

# Open Notebook Science

IP[y]: Notebook BiGo\_methratio Last Checkpoint: Sep 13 09:53 (autosaved)

File Edit View Insert Cell Kernel Help

Code Cell Toolbar: None

## DNA methylation of Oyster Sperm based on Genomic Features

methratio file in SQLShare [https://sqlshare.escience.washington.edu/sqlshare#s=query/sr320%2540washington.edu/clean\\_BiGo\\_methratio\\_v1](https://sqlshare.escience.washington.edu/sqlshare#s=query/sr320%2540washington.edu/clean_BiGo_methratio_v1)

### BiGO methratio GFF

```
SELECT
chr as seqname,
'methratio' as source,
'CpG' as feature,
pos as start,
pos + 1 as [end],
ratio as score,
strand,
'.' as frame,
'.' as attribute
FROM [sr320@washington.edu].[clean_BiGo_methratio_v1]
where
context like '__CG_'
and
CT_Count >= 5
```

[https://sqlshare.escience.washington.edu/sqlshare#s=query/sr320%2540washington.edu/BiGo\\_methratio\\_GFF\\_boop&q=](https://sqlshare.escience.washington.edu/sqlshare#s=query/sr320%2540washington.edu/BiGo_methratio_GFF_boop&q=)

```
In [13]: # ie python fetchdata.py -d "[sr320@washington.edu].[BiGo_Methylation_oysterv9_GFF]" -f tsv -o /Volumes/web/cnidar
# running on commandline because cannot get to work in IPython
```

```
python fetchdata.py -d "[sr320@washington.edu].[BiGo_methratio_GFF_boop]" -f tsv -o
/Volumes/web/cnidarian/BiGo_methratio_boop.gff
```

```
In [15]: #fetchdata failed
```



# Open Notebook Science



## Automating a Workflow: Beyond Blast - to GO Slim

The concept is that you can take a fasta file in a working directory and end up with GO slim information all within a single notebook that is automated. Currently this work by writing (and overwriting) as scratch file to SQLShare. Assumptions are that you are working in a directory with fasta file named `query.fa`. And blast algorithms are in `PATH`.

```
In [13]: #allows plots to be shown inline
%pylab inline

Populating the interactive namespace from numpy and matplotlib

In [4]: #Setting Working Directory
wd="/Volumes/web/whale/fish546/qpx_go_val"
#Setting directory of Blast Databases
dbd="/Volumes/Bay3/Software/ncbi-blast-2.2.29\+/db/"
#Database name
dbn="uniprot_sprot_r2013_12"
#Blast algorithm
ba="blastx"
#Location of SQLShare python tools; you can empty (") if tools are in PATH
spd="/Users/sr320/sqlshare-pythonclient/tools/"

In [5]: cd {wd}
/Volumes/web/whale/fish546/qpx_go_val

In [5]: !{ba} -query query.fa -db {dbd}{dbn} -out {dbn}_{ba}_out.tab -evalue 1E-50 -num_threads 4 -max_hcps_per_subject 1 -
BLAST Database error: No alias or index file found for protein database [/Volumes/Bay3/Software/ncbi-blast-2.2.29+
/db/uniprot_sprot_r2013_12] in search path [/Volumes/web/whale/fish546/pipeline_test_dir4::]

In [6]: !head -1 {dbn}_{ba}_out.tab
QPX_transcriptome_v1_Contig_2 sp|P52712|CBPX_ORYSJ 43.75 416 213 12 2095 869 6 40
7 3e-98 326

In [17]: #Translate pipes to tab so SPID is in separate column for Joining
!tr '|' '\t' <{dbn}_{ba}_out.tab> {dbn}_{ba}_out2.tab

In [18]: !head -1 {dbn}_{ba}_out2.tab

In [8]: #Uploads formatted blast table to SQLshare; currently has generic name and meant to be temporary: Warning will over
!python {spd}singleupload.py -d scratchblast_out {dbn}_{ba}_out2.tab

...

In [9]: !python {spd}fetchdata.py -s "SELECT * FROM [sr320@washington.edu].[scratchblast_out]blast Left Join [sr320@washing

In [10]: !head -2 {dbn}_join2goslim.txt

...

In [11]: !python {spd}singleupload.py -d scratchjoin_slim {dbn}_join2goslim.txt
processing chunk line 0 to 18037 (0.0718240737915 s elapsed)
pushing uniprot_sprot_r2013_12_join2goslim.txt...
parsing 9A18D989...
finished scratchjoin_slim

In [12]: #Sets GO aspect
!python {spd}fetchdata.py -s "SELECT Distinct Column1 as query, Column3 as SPID, GOSlim_bin FROM [sr320@washington.

In [13]: !head justslim.txt

...

In [15]: from pandas import *
```

Set some variables

blast

convert file format

upload to SQLShare  
(python client)

join in SQLShare -  
download

read in pandas

matplotlib generates  
graph of GOslim

# Open Notebook Science

[https://github.com/sr320/ipython\\_nb](https://github.com/sr320/ipython_nb)

This repository Search or type a command Explore Gist Blog Help

sr320 / ipython\_nb Unwatch 1

My Lab Notebook — Edit

206 commits 2 branches 0 releases 1 contributor

branch: master ipython\_nb / +

File	Description	Time
sm		
sr320 authored 3 days ago	Latest commit 717a24e7f6	
.ipython_checkpoints	testing post	4 days ago
fish546	new tools	4 days ago
img	catching up with class	a month ago
tools	sm	3 days ago
.gitignore	en ignore	7 months ago
BiGill_Gene_Methylation.ipynb	sm	6 months ago
BiGill_RNAseq.ipynb	DMG annotations	2 months ago
BiGill_Trans_Elements.ipynb	Adding all ipynb files from cnidarian	7 months ago
BiGill_array.ipynb	DMG annotations	2 months ago
BiGo - methratio error.ipynb	Adding all ipynb files from cnidarian	7 months ago
BiGo_GFF_dev.ipynb	Adding all ipynb files from cnidarian	7 months ago
BiGo_RNAseq.ipynb	sm	a month ago
BiGo_larvae.ipynb	DMG annotations	2 months ago
BiGo_larvae_2.ipynb	bsmap on egg QC	4 days ago

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# Open Notebook Science

https://github.com/sr320/ipython\_nb

This repository Search or type a command Explore Gist Blog Help

sr320 / ipython\_nb Unwatch 1

My Lab Notebook — Edit

206 commits 2 branches 0 releases 1 contributor

branch: master ipython\_nb / +

File	Author	Commit	Time
sm	sr320	717a24e7f6	3 days ago
.ipynb_checkpoints	testing post		4 days ago
fish546	new tools		4 days ago
img	catching up with class		a month ago
tools	sm		3 days ago
.gitignore	en ignore		7 months ago
BiGill_Gene_Methylation.ipynb	sm		6 months ago
BiGill_RNAseq.ipynb	DMG annotations		
BiGill_Trans_Elements.ipynb	Adding all ipynb files from cnidarian		
BiGill_array.ipynb	DMG annotations		
BiGo - methratio error.ipynb	Adding all ipynb files from cnidarian		
BiGo_GFF_dev.ipynb	Adding all ipynb files from cnidarian		
BiGo_RNAseq.ipynb	sm		
BiGo_larvae.ipynb	DMG annotations		
BiGo_larvae_2.ipynb	bsmap on egg QC		

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## IPython Notebook Viewer

**Open Science**

A simple way to share your IPython Notebooks

Share your own notebook, and browse others'

Enter a gist number or url

Go!

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# Open Notebook Science

*a very new experiment*

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# Open Notebook Science

*a very new experiment*

The screenshot shows the sr320.info website interface. At the top, there is a search bar with a magnifying glass icon, a green 'SEARCH' button, and an orange 'RSS' button. The main content area is divided into two columns. The left column displays a list of notebook entries, each with a date banner (e.g., 'FEB 14'), a title in red (e.g., 'qDOD RNAseq[0214\_0646]'), and a light blue bar containing the text 'Posted 2 days ago' and 'nb, ipython,'. The right column contains an 'About' section with a description of the project, a list of included content (IPython Notebook snapshots, iPlant DE alerts, GitHub commits, and more), and a contact request. Below this is a navigation menu with links for 'Home', 'Ask me anything', and 'Archive'. At the bottom of the right column, there is a link for 'The Details of my Lab Notebook'. The bottom of the left column shows a date banner for 'FEB 13' and a 'Notes' entry with the text 'Worked on BSMAP pipeline'.

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February 2014 81 posts

The screenshot shows a grid of blog posts from February 2014. The posts are organized into a grid with varying numbers of columns per row. The posts include:

- ift\_05\_uniprot\_sprot\_r2013\_12\_join2goslim.txt** (eagle.fish.washington.edu)
- notes** (Running ift\_05 through fasta2slim pipeline)
- notes** (looking at new assembly of all libraries- and new trim)
- BlackAb\_Annot[0215\_1334]** (ift.tt)
- ift\_BlackAbalone\_sp\_go\_pat h** (eagle.fish.washington.edu)
- The Details of my Lab Notebook** (I use IPython Notebook as my lab notebook, hosted on GitHub, with the working directories in Dropbox. I use one primary repository which currently has subdirectories for fish546 (class) and 'tools' which represent reusable workflows. I maintain a second repository for a larger unit)
- notes** (Only 1.7% map of BSGill to Ensembl v)
- BiGill\_CpG\_Ensembl[0215\_0710]** (ift.tt)
- notes** (BSMAP needs redo on new genome)
- qDOD\_RNAseq[0215\_0557]** (ift.tt)
- sr320 pushed to master at sr320/qdod** (ift.tt)
- bb3 bam uploading, and bedgraph** (Feb 15, 2014)
- http://ift.tt/1f0caCH** (ift.tt)
- http://ift.tt/1f0c9Pf** (ift.tt)
- grep "[ATCG] [ATCG]CG[ATCG]" <file>** (When run on what you gave me it gives me C12722 104 + TACGT 1.000 1.00 1 1 1 1 0.207 1.000 C12722 134 + TGCGG 1.000 1.00 1 1 1 1 0.207 1.000)
- iPlant DE Alert** (The status of genomeCoverageBed\_BiGo has changed to Completed. Description: Start Date: Sat Feb 15 2014 05:14:28 GMT-0700 (MST) Results Folder: /iplant/home/sr320/analyses/genomeCoverageBed\_BiGo-2014-02-15-05-14-28.745)
- notes** (running genomecoveragebed sep to get bedgraph for BiGo)
- github: sr320 pushed to master at sr320/ipython\_nb** (sr320 pushed to master at sr320/ipython\_nb)
- notes** (Trying RNAseq2Bedgraph with BiGoRNA)
- notes** (Relook at Black Abalone transcriptome)
- github: sr320 pushed to master at sr320/qdod** (ift.tt)
- BlackAb\_Annot[0214\_1530]** (ift.tt)
- github: sr320 pushed to master at sr320/ipython\_nb** (ift.tt)
- notes** (Used some one-liners to help Hannah convert csv data- still mystery non-integer)
- w3\_RNAseq[0214\_1323]** (ift.tt)
- Notes** (Worked on BSMAP pipeline)
- notes** (Worked on automating IPython snapshots.)
- qDOD RNAseq[0214\_0646]** (ift.tt)
- w6\_denovo[0214\_0643]** (ift.tt)
- CC\_ampk.html** (ift.tt)
- Posting ip?** (-)
- iPlant DE Alert: RNAseq2bedgraph\_BiGoRNA**
- iPlant DE Alert: RNAseq2bedgraph SE\_BB3**
- iPlant DE Alert: genomeCoverageBed\_analys**
- BiGo\_larvae\_2.html** (ift.tt)
- iPlant DE Alert: bamToBed\_analysis1 status**

sr320.info

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Sharing



Photo credit: Flickr, Creative Commons, speechless

# Example



Based on Ensembl release 40 - Aug 2006

### Use Ensembl to...

- Search Contig Browser
- Data mining [BioMart]
- Export data
- Download data
- Digital Differential Display
- Venn Diagrams

### Docs and downloads

- About Contig Browser
- Contig Browser data

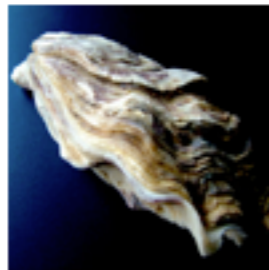
### Other links

- Home
- Sigenae
- Agenae



## Browse the *Crassostrea gigas* contigs database

### About the *Crassostrea gigas* contigs database



#### Assembly

The Oyster EST contig browser aims to produce and maintain an automatic annotation of Oyster EST libraries. This database [GigasDatabase](#) was initiated within the frame of the [AquaFirst](#) European project, it now gathers EST sequences produced by a [Marine Genomics Europe](#) project (GOCE-CT-2004-505403) and a [Genoscope](#) project. [GigasDatabase](#) is regularly updated in the context of the ANR project "Gametogenes" (ANR-08-GENM-041).

#### Annotation

Contigs were annotated searching sequence homologies against following databases:

- UniProtKB/Swiss-Prot Release 2011\_03 of 08-Mar-2011
- RefSeq Protein Index Blast of 09-Jan-2011
- Pfam Release 24.0 of Jul-2009
- Sigenae Oyster Contigs V8
- RefSeq RNA Index Blast of 09-Jan-2011
- TIGR Fugu FGI 3.0

### Statistics

Assembly:	pcg8, Mar 2011
Number of base pairs (Contigs):	67,875,621
Number of sequences (EST/mRNA):	1,013,570
Total number of contigs:	82,312
Number of singlets:	20,148
Polymorphism:	Mar 2011
Number of putative SNPs:	29,493
Number of contigs including SNP:	10,392

### Example Data Points

Jump directly to the example entry points:

Contig: [BQ426804.p.cg.8](#)

SNP: [snp\\_EW777925\\_683](#)

# Example



Based on Ensembl release 40 - Aug 2006

Use Ensembl to...

Browse the *Crassostrea gigas* contigs database

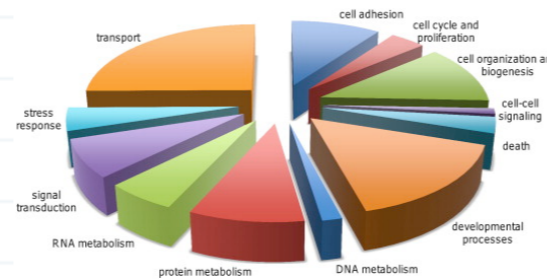
Search Contig Browser

About the *Crassostrea gigas* contigs

Statistics

Table 1. Summary of assembly and RNA-Seq statistics for de novo assembly and reference mapping (GigasDatabase v8).

		De novo assembly	Reference mapping
Assembly	Mapped reads	8,407,963	29,107,760
	Unmapped reads	36,944,698	16,244,901
	Contigs	18,510	77,433
	Average contig length	276	554
	Average contig coverage	62	16
RNA-Seq	Differentially expressed genes	2991	427
	Enriched GO biological process	15	3



Contigs were annotated searching sequence homologies against following databases:

- ▶ Sigenae Oyster Contigs V8
- ▶ RefSeq RNA Index Blast of 09-Jan-2011
- ▶ TIGR Fugu FGI 3.0

Assembly:	pcg8, Mar 2011
Number of base pairs (Contigs):	67,875,621
Number of sequences (EST/mRNA):	1,013,570
Total number of contigs:	82,312
Number of singlets:	20,148
Polymorphism:	Mar 2011
Number of putative SNPs:	29,493
Number of contigs including SNP:	10,392

Example Data Points

Jump directly to the example entry points:

Contig: [BQ426804.p.cg.8](#)

SNP: [snp\\_EW777925\\_683](#)

Characterizing short read sequencing for gene discovery and RNA-Seq analysis in *Crassostrea gigas*

Mackenzie R. Gavary, Steven B. Roberts\*



# Example



Based on Ensembl release 40 - Aug 2006

Use Ensembl to... Browse the *Crassostrea*

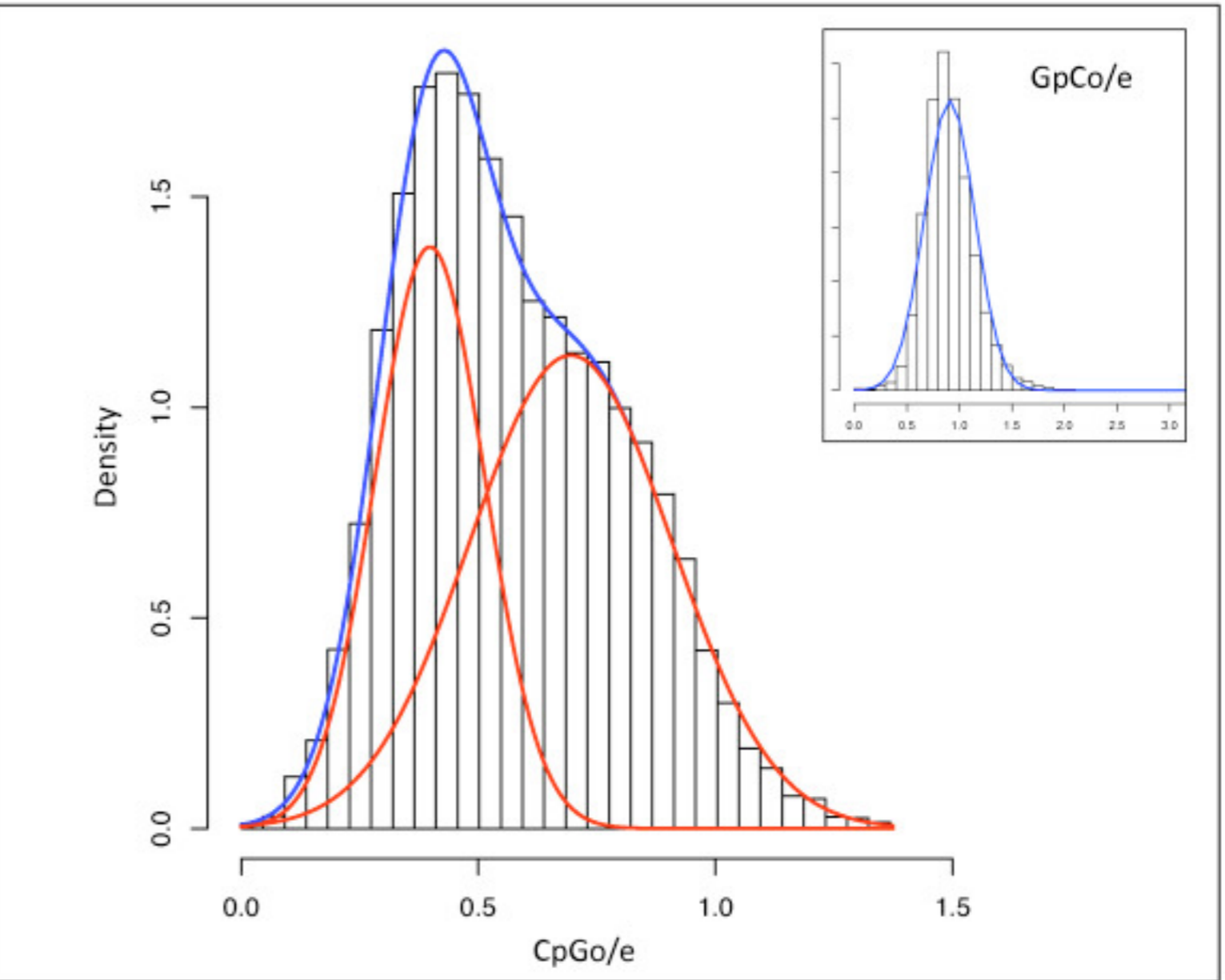
Search Contig Browser About the *Crassostrea*

Table 1. Summary of assembly and RNA-Seq statistics for de novo assembly (v8).

		De novo assembly	Reference
Assembly	Mapped reads	8,407,963	29,107,000
	Unmapped reads	36,944,698	16,244,000
	Contigs	18,510	77,433
	Average contig length	276	554
	Average contig coverage	62	16
	Contigs annotated to GO Slim	3931	7296
RNA-Seq	Differentially expressed genes	2991	427
	Enriched GO biological process	15	3

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- RefSeq Protein Index Blast
- Pfam Release 24.0 of Janel
- Sigenae Oyster Contigs
- RefSeq RNA Index Blast
- TIGR Fugu FGI 3.0



- Home
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Research article

Highly accessed

Open Access

**DNA methylation patterns provide insight into epigenetic regulation in the Pacific oyster (*Crassostrea gigas*)**

Mackenzie R Gavery and Steven B Roberts\*

DOI: [10.1186/1471-2163-10-103](https://doi.org/10.1186/1471-2163-10-103)

# Living in an Ivory Basement

Stochastic thoughts on science, testing, and programming.

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[teaching](#)

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## Loligo pealeii (squid) data dump

A few months back, I announced [the khmer protocols project](#), an effort to write down an explicit, open protocol for transcriptome and metagenome assembly. This project was started [during the summer of 2013 at the Woods Hole Marine Biological Lab](#), in collaboration with [Joshua Rosenthal](#). I'd met Josh at the [CephSeq meeting](#) the year before, and we reconnected at MBL in 2013; there, motivated in part by discussions at CephSeq, I started writing the eel-pond protocol.

During the summer, Josh and I found that we were both incredibly frustrated by the general failure of researchers to share data pre-pub, and so we started to chat seriously about some ideas we'd independently had about incentivizing the opening of transcriptome data. This eventually led to [the Open Marine Transcriptome project](#), which is still in its formative stages.

However, in the meantime I am happy to report that Josh assembled a bunch of squid transcriptomes and is willing to make them available to all. In addition, Josh did some skim Illumina sequencing (~40x coverage) of the squid genome, and my lab assembled it; we are making a really basic draft genome available as well. (Here, by squid, we mean *Loligo pealeii*, or the [Longfin inshore squid](#) -- "loligo" for short. It is also sometimes called "Doryteuthis pealeii".)

It should be noted that one of the transcriptomes was made from carefully dissected Giant Fiber Lobe neurons. These are the cell bodies for the well-studied squid giant axon. Thus this transcriptome should provide "molecular support" for all the cell biology and neurophysiology that has focused on this most famous of preps.

We are pleased to announce the availability of this data, in two formats.

First, you can download the assembled and annotated transcriptome data in FASTA format:

- [Loligo pealeii buccal ganglion transcriptome \(v1.0\)](#)
- [Loligo pealeii giant fiber lobe \(v1.0\)](#)
- [Loligo pealeii optic lobe \(v1.0\)](#)
- [Loligo pealeii stellate ganglion \(without giant fiber lobe\) \(v1.0\)](#)
- [Loligo pealeii vertical lobe \(v1.0\)](#)

Mon 17 February 2014

By [C. Titus Brown](#)

In [science](#).

tags: [open data f-yeah](#)

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# Living in an Ivory Basement

Stochastic thoughts on science, testing, and programming.

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## Loligo pealeii (squid) data dump

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Mon 17 February 2014

By [C. Titus Brown](#)

In [science](#).

tags: [open data f-yeah](#)

**Why aren't you trying to get a Science or Nature paper out of all this? Aren't your reputations going to suffer for doing all this work without trying to milk the data for all it's worth?**

We're pretty sure our reputations won't suffer from making a bunch of useful data available. Heck, we're pretty sure *your* reputation wouldn't suffer from making a bunch of useful data available (hint).

First, you can download the assembled and annotated transcriptome data in FASTA format:

- [Loligo pealeii buccal ganglion transcriptome \(v1.0\)](#)
- [Loligo pealeii giant fiber lobe \(v1.0\)](#)
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- [Loligo pealeii vertical lobe \(v1.0\)](#)

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The Olympia oyster (*Ostrea lurida*) is the only native oyster on the west coast of the US.

## Alleviating Regulatory Impediments To Native Shellfish Aquaculture

Our approach is to simultaneously address local adaptation in three genetically differentiated populations of Olympia oysters by evaluating genotype-by-environment interactions. We will reciprocally transplant seed produced from wild parents collected from contrasting environments into all environments. This very large reciprocal transplant experiment can test for a home field advantage in survival, maturation and growth in Olympia oysters. The overall goals of this project are to increase our knowledge of local adaptation in Olympia oysters to address concerns that interbreeding between potentially maladapted cultured and wild stocks could negatively impact wild populations. Accordingly, in order to attain these goals, the specific objectives of this proposal are to 1) Evaluate fitness components and performance of seed from different origins in a reciprocal transplant experiment and 2) Characterize genetic and epigenetic

Check out our [wiki](#) for up to date data and more details.

## Genomic Resources

### Transcriptomes

[O.lurida transcriptome version3 \[10/13\]](#)

[O.lurida transcriptome version2](#)

[O.lurida transcriptome version1](#)

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# Steven Roberts

Associate Professor (**Marine Biology**)  
University of Washington



## Research statistics

10100 views

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- Molecular Biology
- Ecology
- Bioinformatics
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## Related authors

Emma Timmins-Schiffman  
 Claire Ellis  
 Mackenzie Gavery  
 Samuel White  
 Michael O'Donnell

## Recently used tags

- thraustochytrid
- qpx
- quahog
- Gene regula
- oyster
- preprint
- aquaculture
- ma-seq

## Working with...

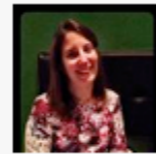
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elene dorfmeier



Caroline Storer  
Graduate Student



Colleen Burge  
Postdoctoral research ass...



Emma Timmins-Schiffman  
PhD candidate



Claire Ellis

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Herring Hepatic Transcriptome 34300 contigs.fa	Dataset	4872 views 16 shares
Herring Testicular Transcriptome 31545 contigs.fa	Dataset	747 views 18 shares
Bay scallop population structure on Cape Cod	Fileset	662 views 14 shares
Hard clam transcriptome contigs	Dataset	453 views 5 shares
DNA methylation as a mechanism to increase adaptive potential in invertebrates	Paper	412 views 1 shares
Development of genetic markers to assess disease resistance in the eastern oyster	Paper	320 views 9 shares

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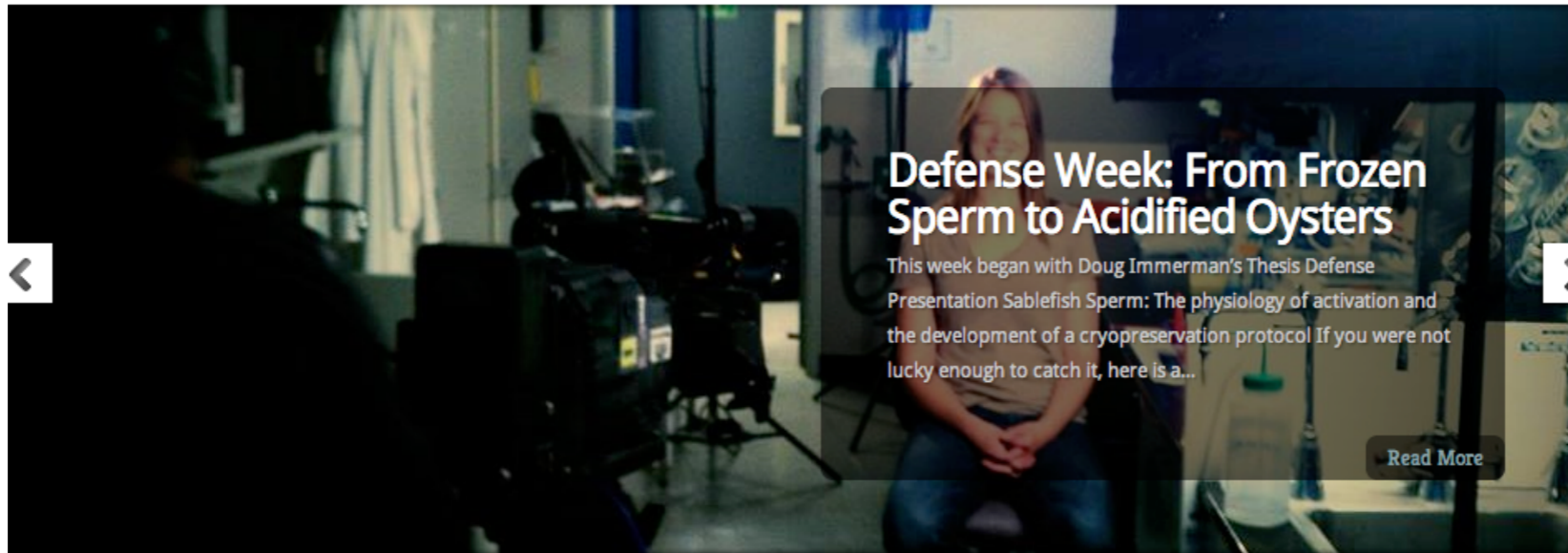
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### Research

Research in our lab focuses on characterizing physiological responses of marine organisms to environmental change. Using integrative approaches we strive to examine impacts and adaptive potential from the nucleotide to organismal level. A core component of this includes investigating the *functional relationship of genetics, epigenetics, and transcription.*

### Open Science

We practice open science with lab members maintaining online electronic lab notebooks. In addition we have developed a Data and Resource Sharing Plan and are in the process of providing a list of resources we use for open science. To learn more about what we are doing, be sure to view our weekly, live lab meetings on YouTube, follow us on Facebook, or check out one or more of our blogs.

### News and Notes

- Congrats to Emma Timmins-Schiffman! Successfully Passed her PhD Final Exam! [2/14]
- Congrats to Doug Immerman! Successfully Passed his Masters Final Exam! [2/14]
- Archive

### Labcam



### Notebook Updates

- Sam's Working Notebook Jan - July 2014 (kubu4)
- sr320 Notebook (sr320)
- sr320 Notebook (sr320)
- sr320 Notebook (sr320)

### Fresh from the lab...

Big Data!  
Exploring the Biology of Oysters  
*a few million base pairs at a time*

### Connect

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 Roberts Lab  
 @sr320  
 206.685.3742

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# Open Science Philosophy

Transparency with limited effort

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# Open Science Philosophy

Transparency with limited effort

*will try just about anything*

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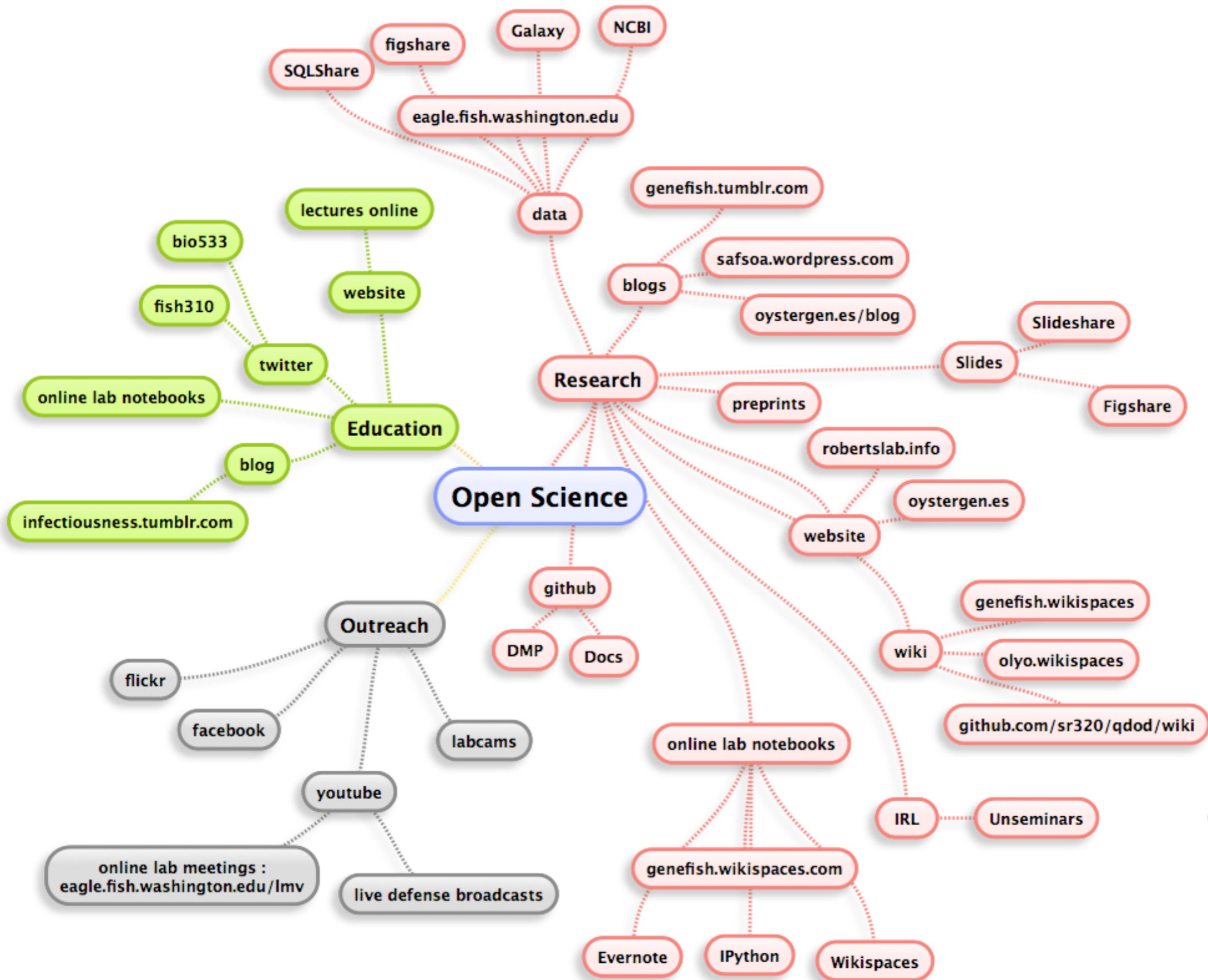
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# BioCode's Notes

Computational Proteomics & Bioinformatics

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- Software & Tools
- Programming Tips
- Core Manuscript
- Polls
- E

Wednesday, 19 February 2014

In the ERA of science communication, Why you need Twitter, Professional Blog and ImpactStory?

Yasset Perez-Riverol on Wednesday, February 19, 2014

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Male  
20.026  
Female

Start them early



# Acknowledgements

Emma Timmins-Schiffman



*acidification*  
Saltonstall-Kennedy

Mackenzie Gavery  
Claire Olson

*DNA methylation*



Sam White  
Brent Vadopalas  
Jake Heare



Aquaculture Program



Bill Howe  
Dan Halperin

[oystergen.es/data](http://oystergen.es/data)



