

# **Empirical algorithm for global chlorophyll: Is it really that “bad”?**

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# Acknowledgement:

NASA

NRL

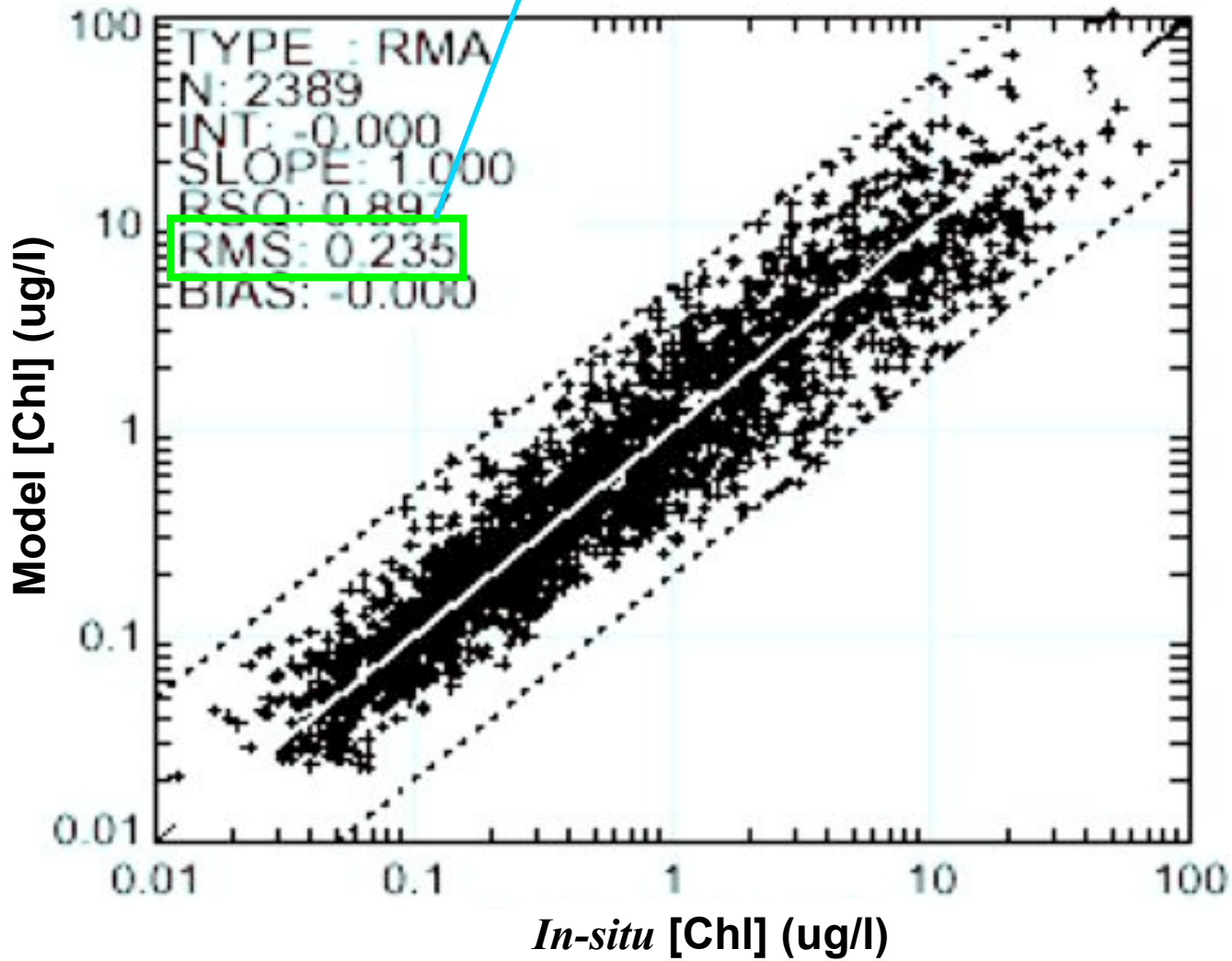
Jeremy Werdell (OBPG)

Janet Campbell (UNH)

# Motivation:

Model is ~1.7 times higher or lower than *in-situ*.

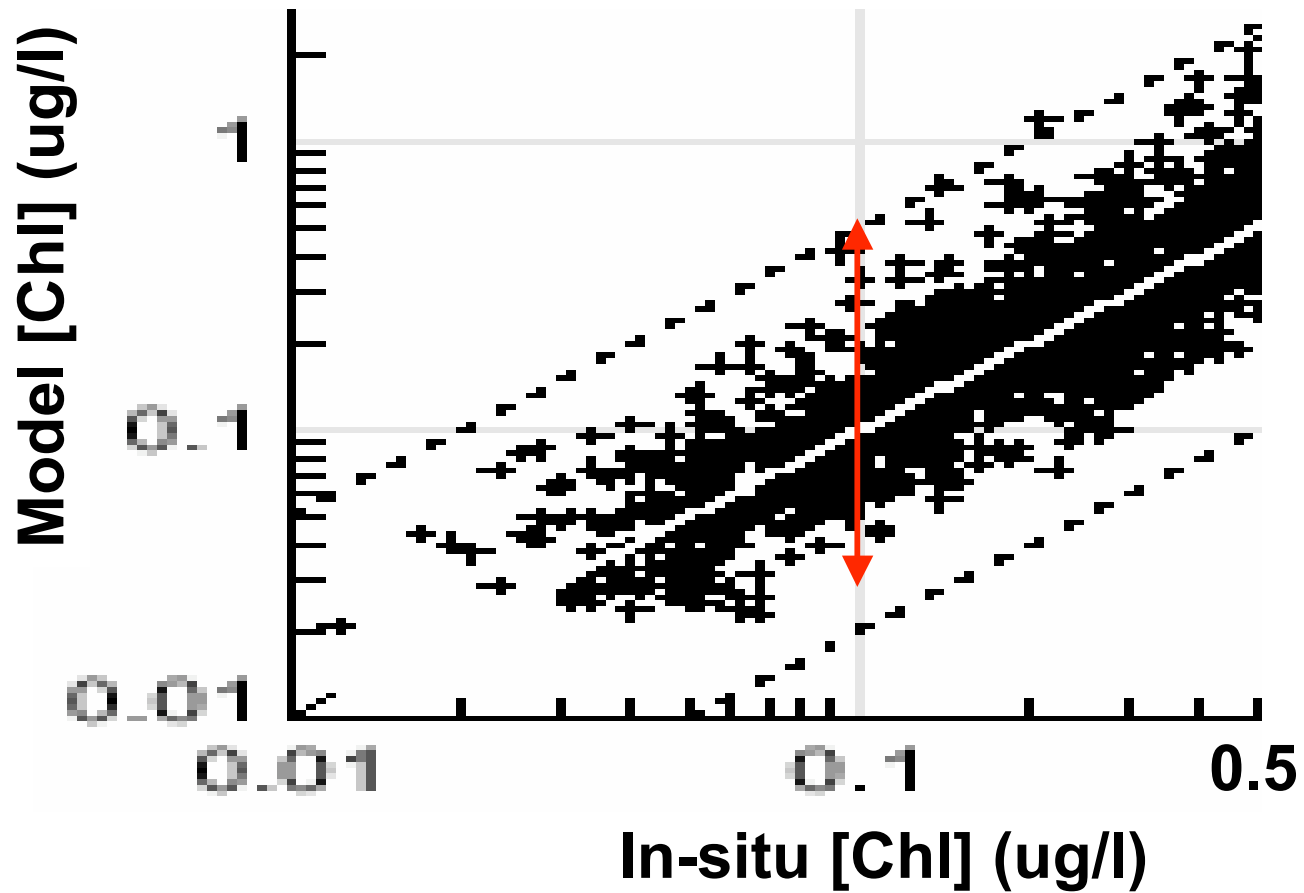
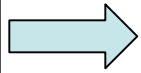
Model [Chl] = Fun(Rrs( $\lambda_1$ )/Rrs( $\lambda_2$ ))



(O'Reilly's OC4v5)

# Oceanic waters

Nonlinear scaling of total absorption



**Q: How much of these dispersion are natural?**

[Chl]

38.19  
 35.01  
 26.91  
 47.96  
 23.55  
 23.97  
 19.15  
 33.46  
 17.32  
 10.12  
 0.401  
 0.515  
 0.539  
 0.091  
 0.079  
 0.461  
 0.132  
 0.296  
 0.4  
 0.314  
 3.287  
 0.919  
 1.741

[Chl]  
 Rrs443 Rrs489 Rrs510 Rrs555

0.001686 0.003293 0.004036 0.007479  
 0.001384 0.002173 0.002499 0.004152  
 0.001185 0.001843 0.002288 0.004246

log([Chl])

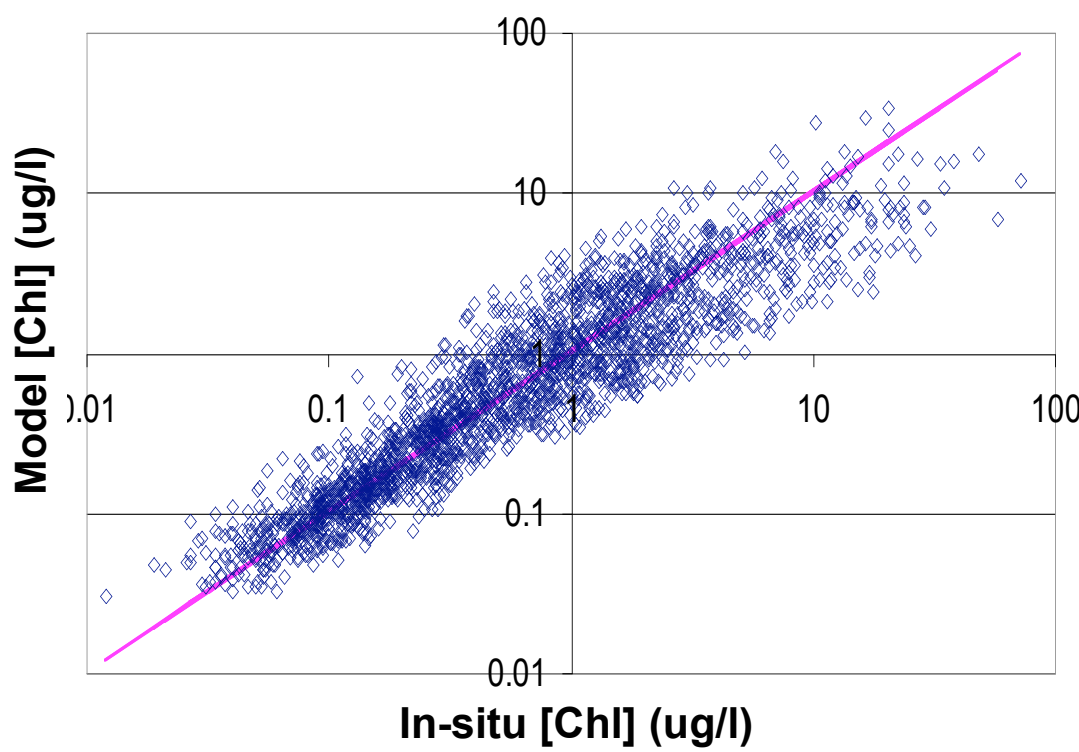
1.58195  
 1.544192  
 1.420014

log(rr\_max)

-0.26789  
 -0.22049  
 0.36052

rr\_max

0.539644  
 0.601879  
 0.53886  
 0.524778  
 0.6794  
 0.715184  
 0.600039  
 0.548027  
 0.689655  
 0.717385  
 1.646516  
 1.284376  
 1.325207  
 3.242857  
 3.592568  
 3.271292  
 5.874929  
 2.909639  
 3.29818  
 3.846339  
 1.284128  
 2.05341  
 1.583873



0.00247 0.002691 0.00234 0.001699  
 1.741

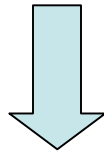
0.240799

0.19972

(OBPG NOMAD subset)

## Assumptions:

1. Data has '**enough**' representation
2. Data has '**good**' quality
3. '**Normal**' distribution of errors



## Impression:

Empirical algorithm for global chlorophyll produces '**bad**' results, even for oceanic waters.

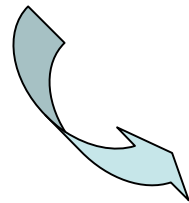
Explicitly or implicitly, the large dispersion is viewed as algorithm “error”.

## Sources of “error”:

1. Natural (optical, bio-optical) variations
2. Algorithm
- 3. Data imperfection**

## Source of imperfection:

Measurement methodology  
In-homogeneity (horizontal and vertical)  
Sample handling  
Data processing  
etc



[Chl]

Consistent?



Rrs

What is the algorithm error if data *inconsistency* is removed?

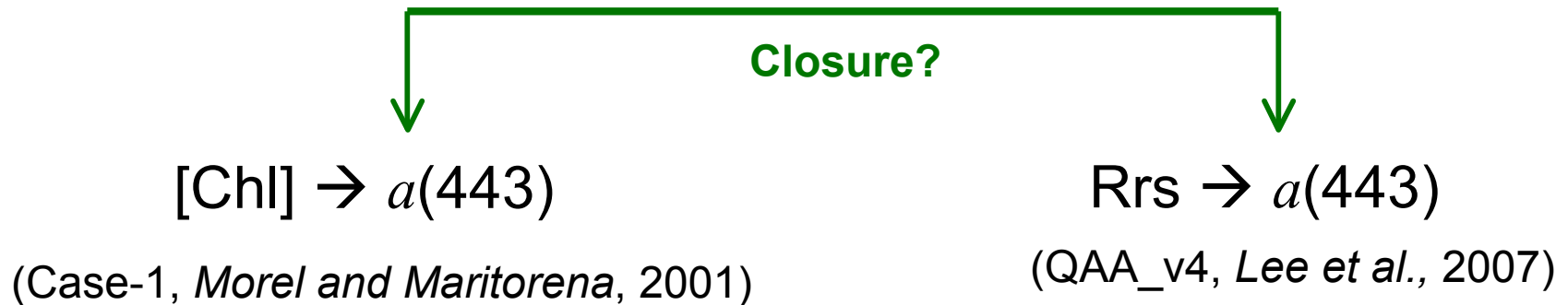
Filter out 'bad' data Easy for obvious or extremes; but difficult for all others.



# Re-analyze existing data (NOMAD)

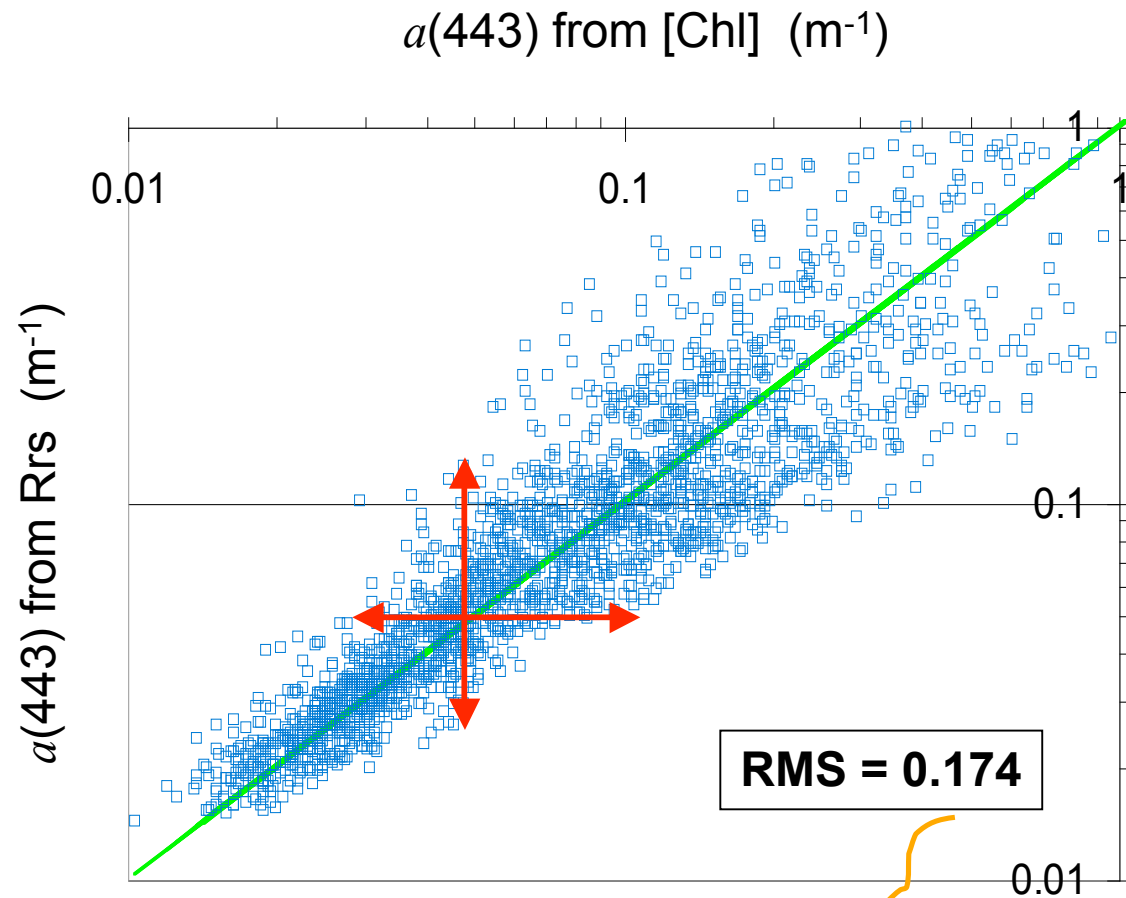
Between [Chl] and Rrs, how mutually consistent are they?

Only when [Chl] and Rrs are consistent, can it possible to use Rrs ratio to infer [Chl].



**Scenario 1:**

Both  $\{[Chl]\}$  and  $\{Rrs\}$  were measured perfectly and are assumed mutually consistent.



How much can the dispersion be explained with a **minimum** “correction” on data imperfection?

1.5 times higher or lower

# Environment/procedure of *in-situ* measurements

“[Chl] is accurate, to the best, within  $\pm 35\%$ .”

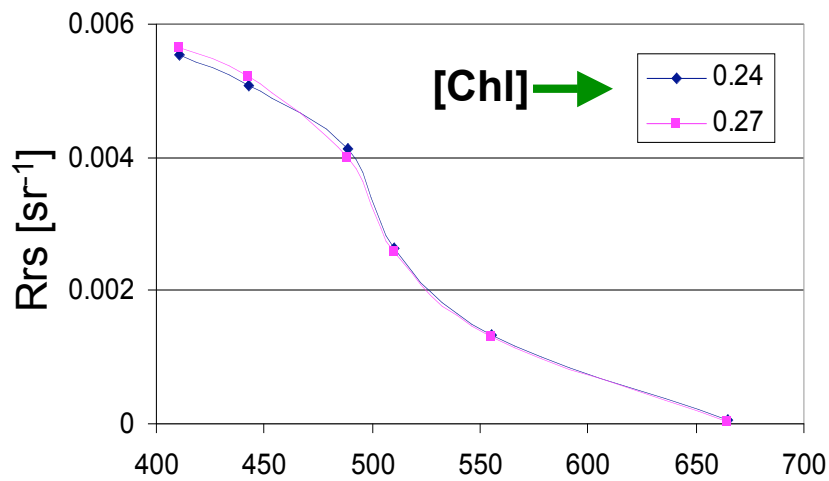
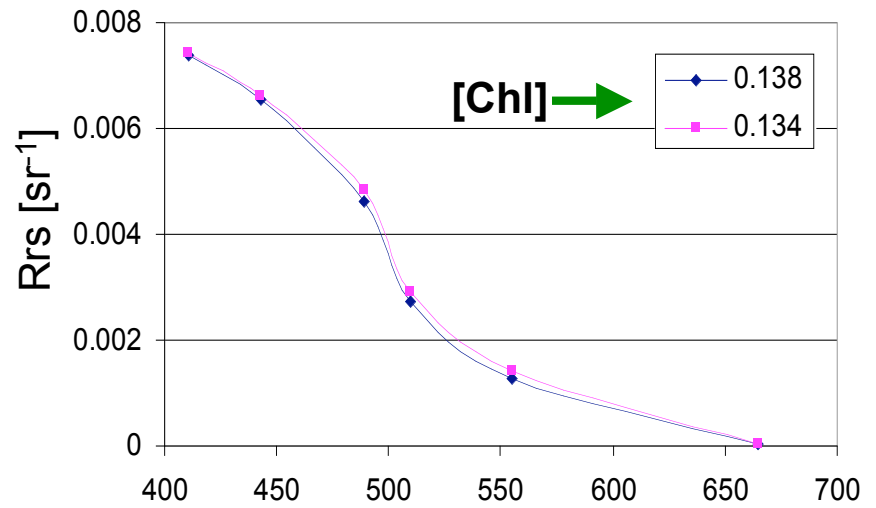
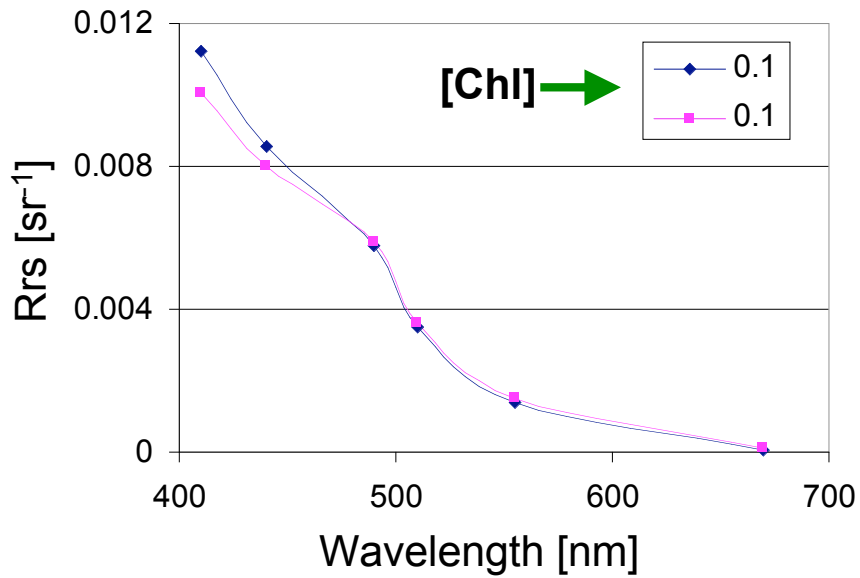
[Chl]

Rrs is easy to be off by  $\pm 10\%$ .

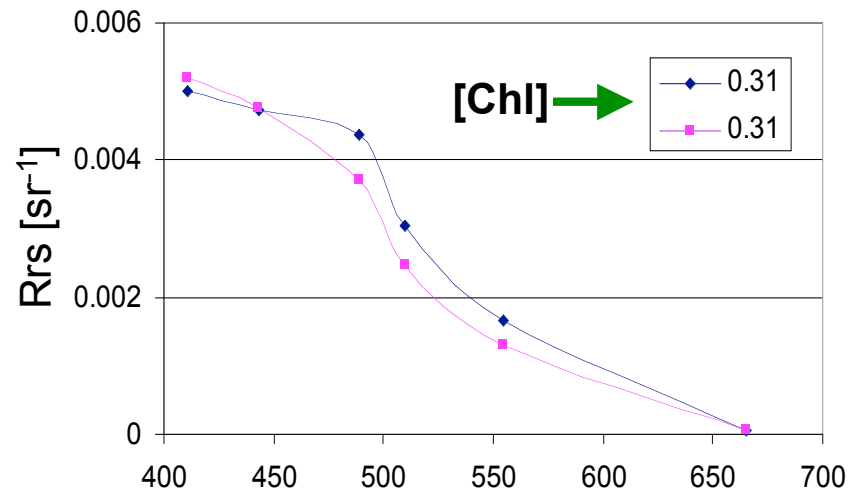
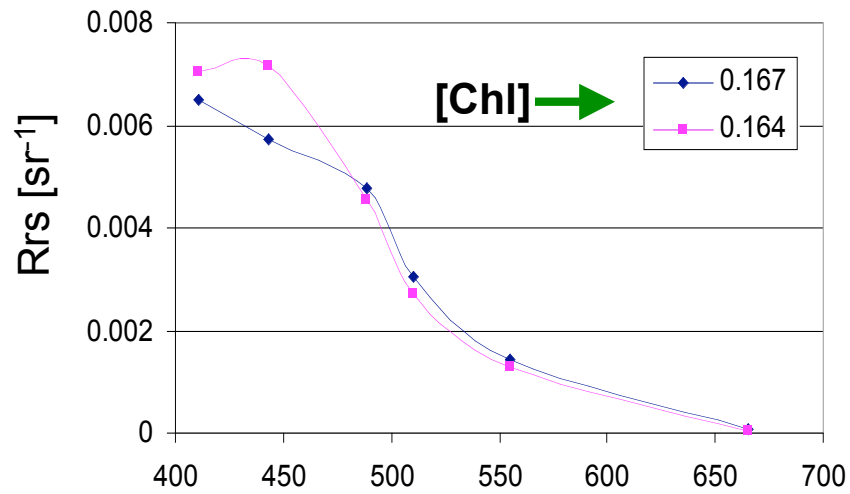
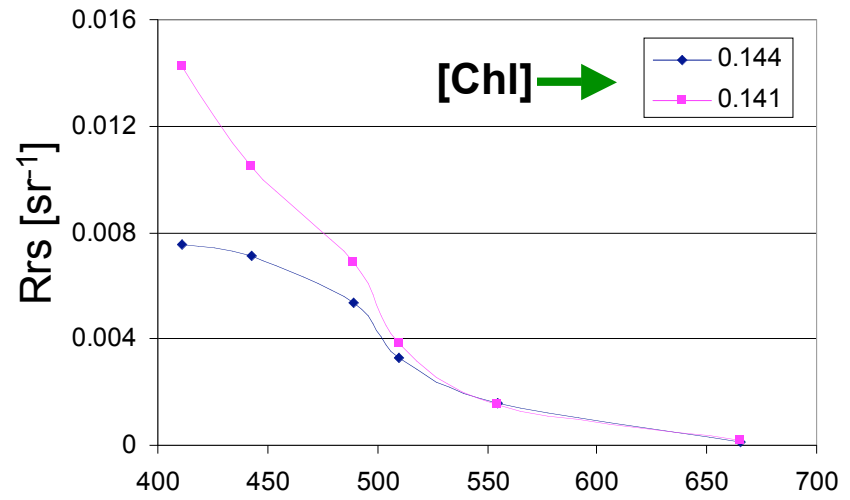
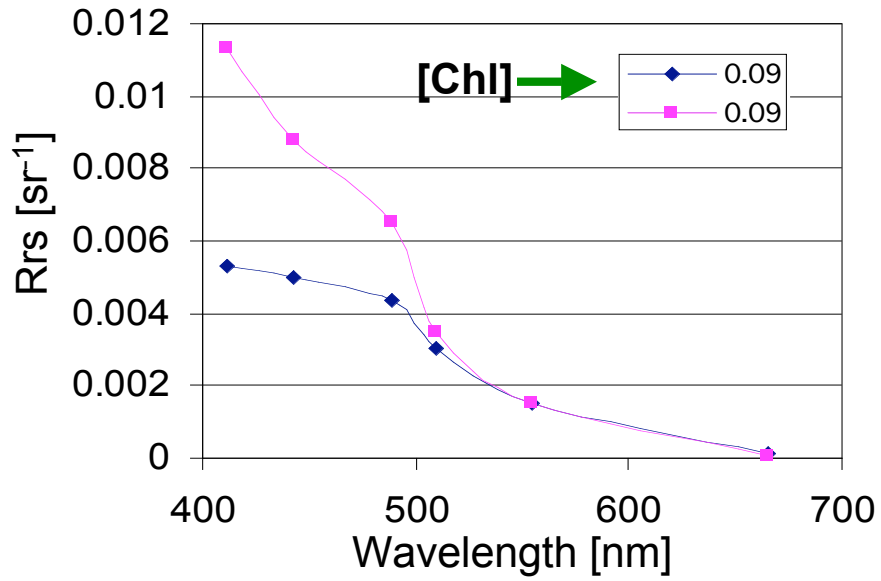
Rrs



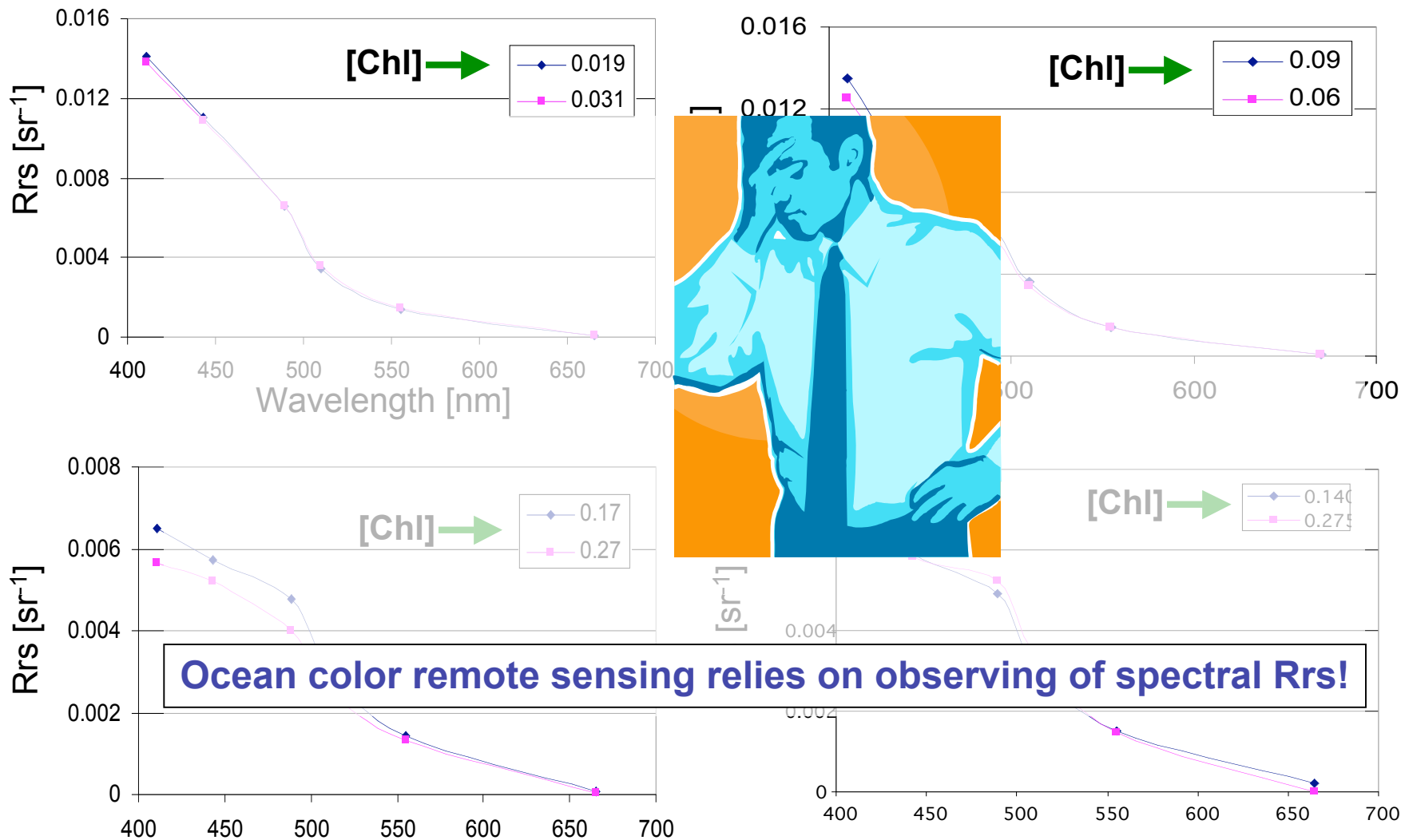
# Mutually consistent data



# Similar [Chl], quite different Rrs



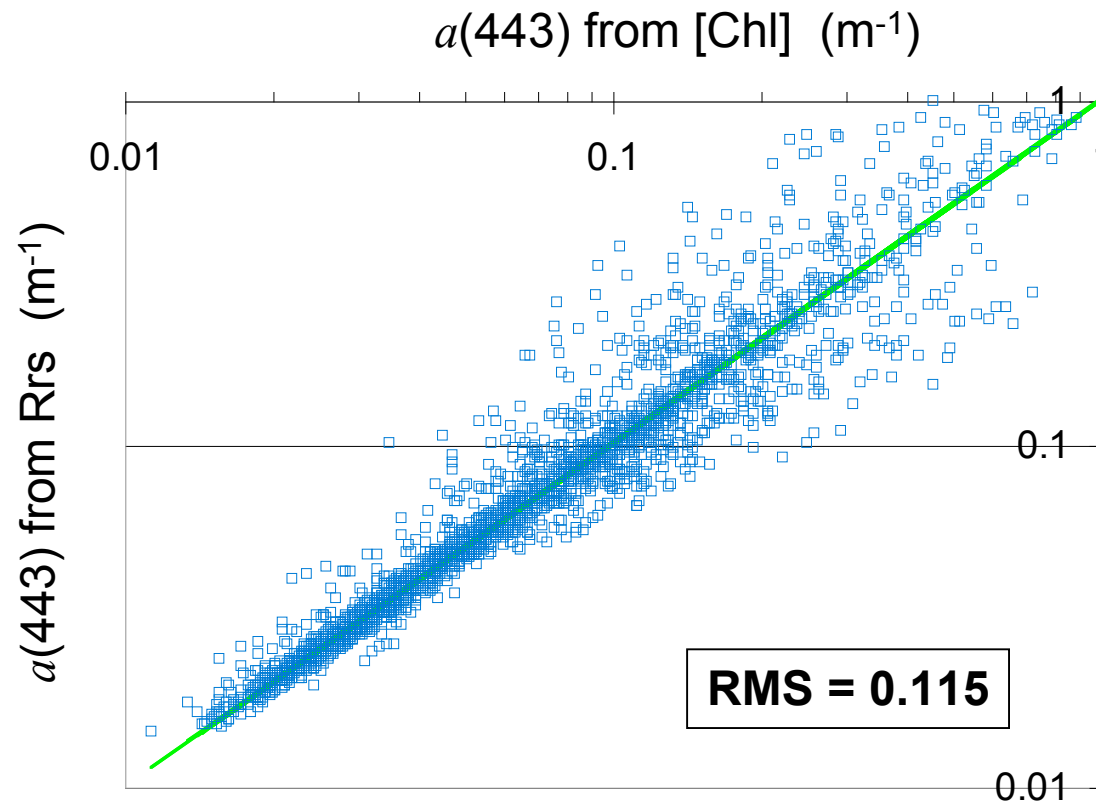
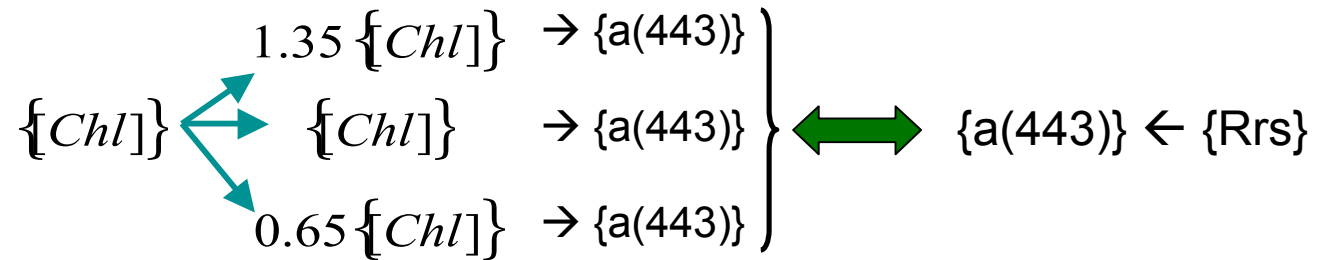
# Similar Rrs, quite different [Chl]



**Ocean color remote sensing relies on observing of spectral Rrs!**

**Scenario 2: {Rrs} are ok, but {[Chl]} is imperfect.**

Rrs no change,  
[Chl] with  $\pm 35\%$



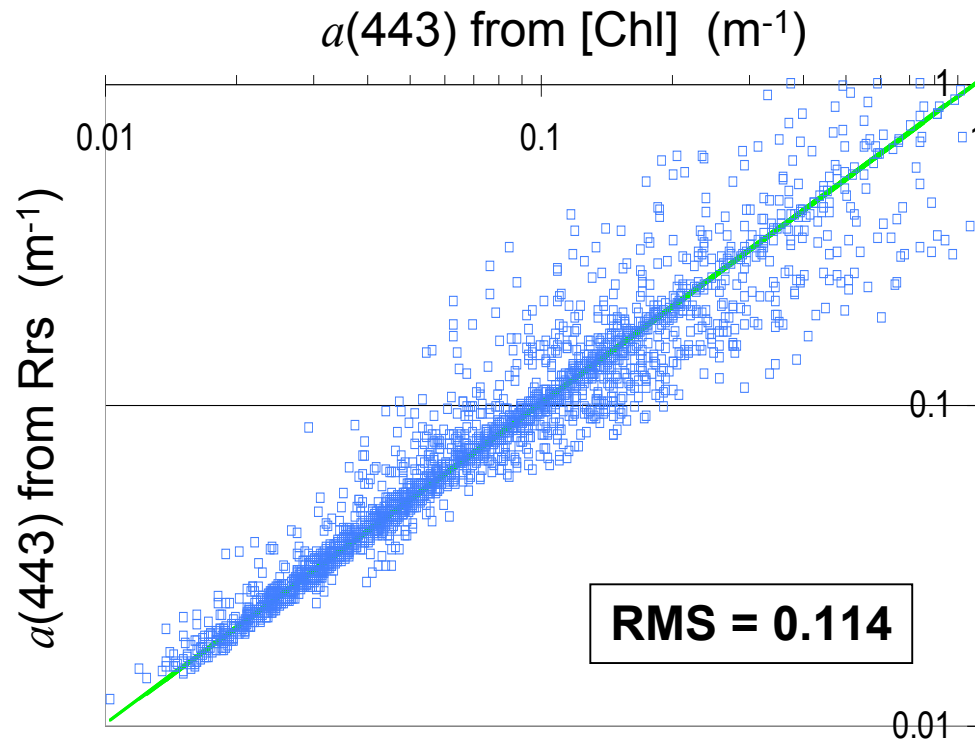
### Scenario 3: {[Chl]} in ok, but {Rrs} is imperfect.

[Chl] no change,  
Rrs with  $\pm 10\%$

{Chl}  $\rightarrow$  {a(443)}  $\longleftrightarrow$

Factors applied to Rrs at

	443	490	555
{a(443)} $\leftarrow$	1.1	1.1	0.9
{a(443)} $\leftarrow$	1.1	0.9	0.9
{a(443)} $\leftarrow$	1	1	1
{a(443)} $\leftarrow$	0.9	0.9	1.1
{a(443)} $\leftarrow$	0.9	1.1	1.1

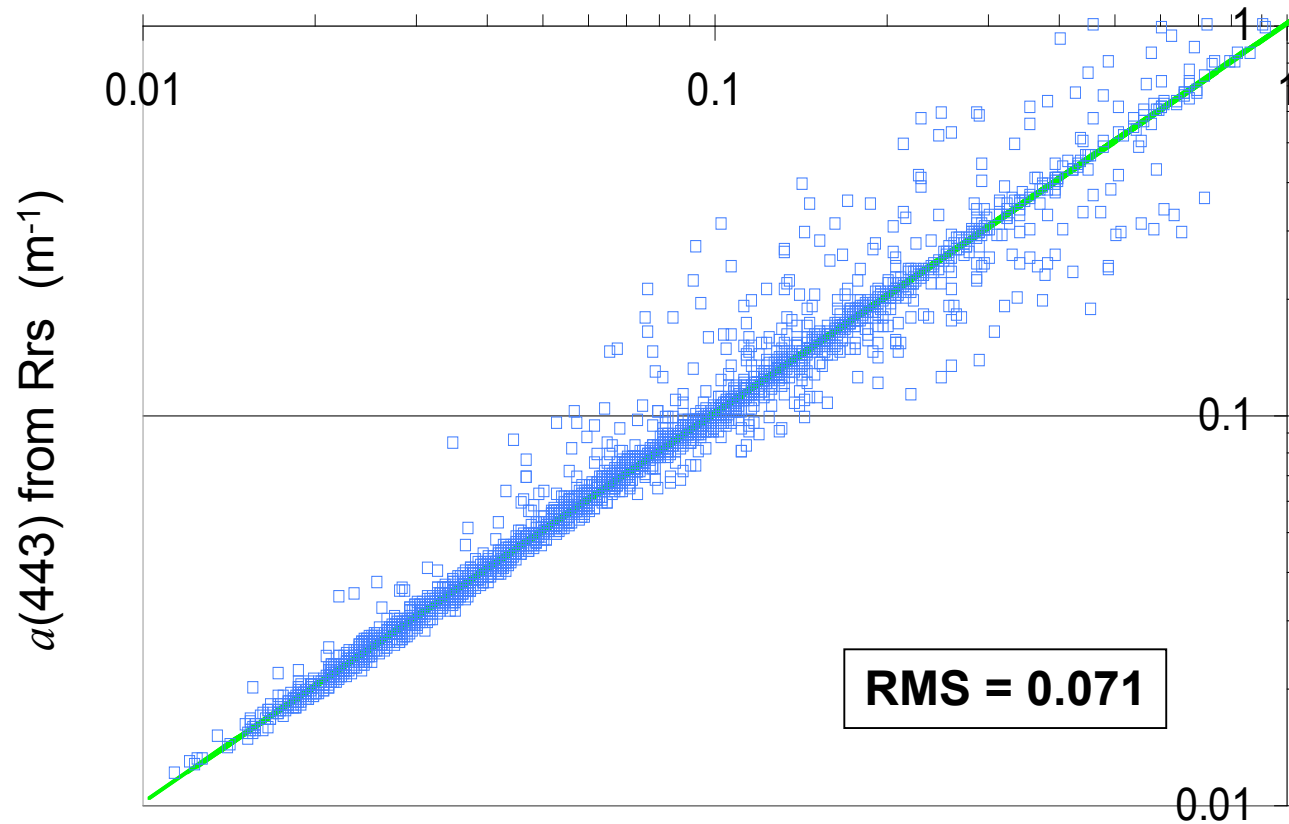




**Scenario 4: Both  $\{[Chl]\}$  and  $\{Rrs\}$  are imperfect.**

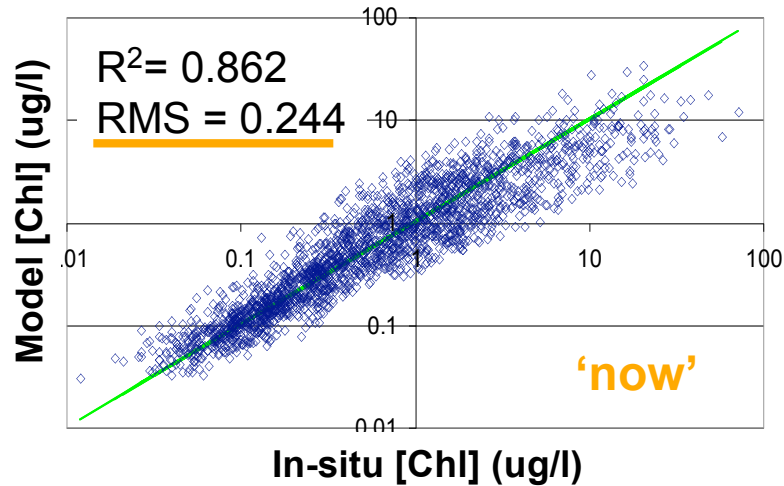
$Rrs$  with  $\pm 10\%$ ,  $[Chl]$  with  $\pm 35\%$

$a(443)$  from  $[Chl]$  ( $m^{-1}$ )

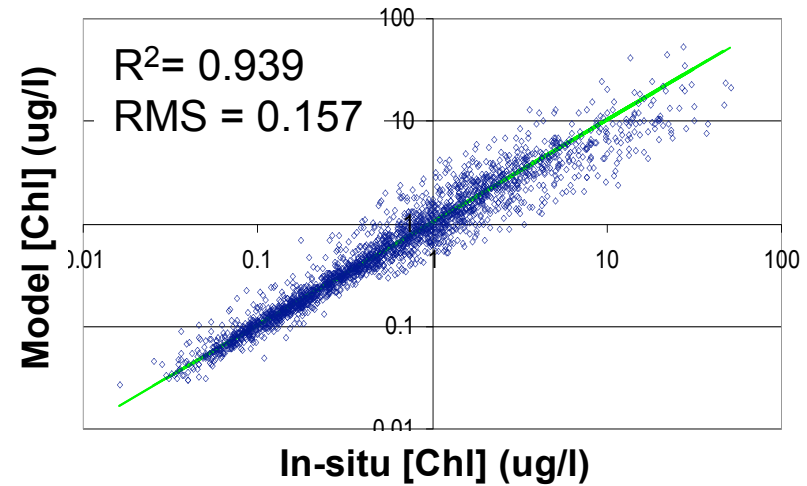


# Effects to empirical algorithm for [Chl]

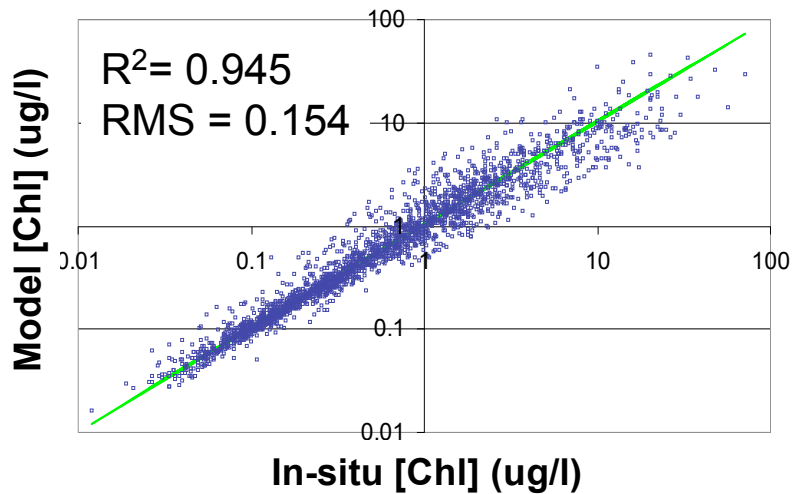
S1: Original data



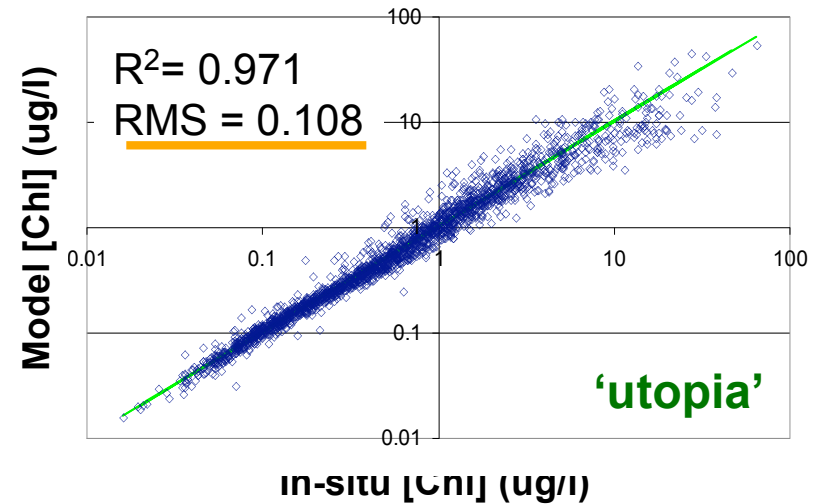
S2: Rrs no change, [Chl] with  $\pm 35\%$



S3: [Chl] no change, Rrs with  $\pm 10\%$



S4: Rrs with  $\pm 10\%$ , [Chl] with  $\pm 35\%$



(OBPG NOMAD subset)

# Summary of RMS

	$a(443)$	[Chl]
<b>Scenario 1</b>	<b>0.174</b>	<b>0.244</b>
<b>Scenario 2</b>	0.115	0.157
<b>Scenario 3</b>	0.114	0.154
<b>Scenario 4</b>	<b>0.071</b>	<b>0.108</b>

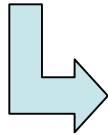
overall dispersion

More than a factor of 2  
reduction in RMS.

~ best algorithm contribution

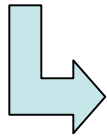
# The message:

1. “Bad” or **less-consistent** data could contribute ~**half** of the overall **dispersion** between sample-measured [Chl] and Rrs-derived [Chl].



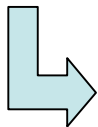
This conclusion may be applicable to IOP products also.

2. For **consistent** measurements, simple ratio derived [Chl] is not as ‘bad’ as it is appeared now, especially for oceanic waters with low [Chl] (< ~0.5 mg/m<sup>3</sup>).

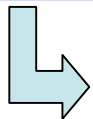


**Semi-analytically derived [Chl] will be even better as other optically active constituents are separated.**

3. Improvement on measurement accuracy and data **consistency** are critical for validating and understanding algorithm (either for [Chl] or IOPs, etc) performance.



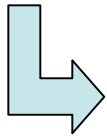
To understand/quantify error propagation, it is necessary to separate algorithm error from the overall dispersion.



Data-based algorithm will then be difficult.

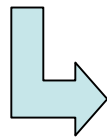
# The message cont.:

4. For waters with high [Chl] ( $> \sim 0.5 \text{ mg/m}^3$ ), larger gap still exist between sample-measured and ratio-derived [Chl] even after “correcting” measurement “errors”. Separation of the active optical components with semi-analytical models is required for improvement of accuracy and for understanding of error propagation.



**Demands better understanding on the spatial and temporal variations of bio-optical properties.**

5. [Chl] is an important ocean-color product, but **not ideal** for validating a remote-sensing system *if the  $\pm 35\%$  minimum uncertainty is still exist.*



A property with less uncertainty from in situ measurements is desired for such purpose.

6. IOPs are not only important ocean-color products, can also serve as a quality check regarding consistency between [Chl] and  $R_{rs}$  values.