

RESEARCH INTERESTS

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My research interests center around the so-called lipophilic pigments. Specifically, these are the chlorophylls, chlorophyll derivatives, carotenoids, certain polyaromatic hydrocarbons (perylene) and a dimeric indole-phenol (scytonemin).

An examination of my vita reveals that I have had certain periods of research. The early seventies included studies on *de novo* carotenogenesis in various species of the genus *Mycobacterium*. Several of the mycobacteria are photochromogenic, producing pigments only when induced to do so by light - and light of the proper wavelength. This was my introduction into the fascinating world of biological pigments. I extended my interest in carotenoids with my master's thesis topic - carotenoids metabolism in Crustacea, namely the blue-crab (*Callinectes sapidus* Rathbun 1895). From the late seventies through the early nineties, I was involved in organic geochemical studies of deep sea, lake and coastal sediments as well as sediment traps and selected experimental (senescence / death) studies. These investigations traced chlorophyll and its derivatives from living biota into the geosphere to eventual destruction (recycle) or preservation as a variety of free-base and metallo-geoporphyrins, notably in oil shales and petroleum crudes. For the two decades I have been involved in two main aspects of the biogeochemistry of chlorophylls and carotenoids, namely the changes that occur during the senescence / death of microalgal cells and the use of pigment distributions as "chemotaxonomic" markers for the assessment of microalgal community structure, productivity and overall dynamics. Most of these studies recently have dealt directly or indirectly with the ecosystems affected

by the Comprehensive Everglades Research Plan (CERP) and have expanded over to the Bahamas.

For the past decade, I have also devoted much time and energy to the investigation of environmental water quality, namely primary plant nutrients (nitrogen, phosphorus, iron etc.). This facet of my studies includes investigating the impact of phosphorus and nitrogen pollution from the indiscriminant over-use of equestrian wastes in agri-business. Namely horse manure on nurseries with runoff into the canals and other surface waters of south Florida. Lately (2015+) this also includes investigating all sources of nutrient pollution as it applies to harmful algal / cyanobacterial blooms in Lake Okeechobee and all waters downstream.

Following I discuss a few points pertinent to my present research efforts. I will be glad to delve deeper into any portion of my studies and potential collaborative research, especially with prospective students in the Department of Chemistry and Biochemistry and the Environmental Sciences Program.

CHLOROPHYLLS and CAROTENOIDS

Pigment-based chemotaxonomy, algal blooms & community dynamics:

The chlorophylls are the main pigments of photosynthesis. (see <http://life.uiuc.edu/govindjee/paper/gov.html> or <http://photoscience.la.asu.edu/photosyn>)

They process trapped solar energy into 'reductant' (NADPH, e^-), which can then reduce ('fix') the carbon in carbon dioxide (CO_2) to *organic carbon*. This is the so-called *carbon fixation* which yields the organic compounds (sugars, proteins, fats *etc.*), fuels and builds the majority of life on earth.

The most common, abundant and best-studied chlorophyll is chlorophyll-*a*, as shown here.

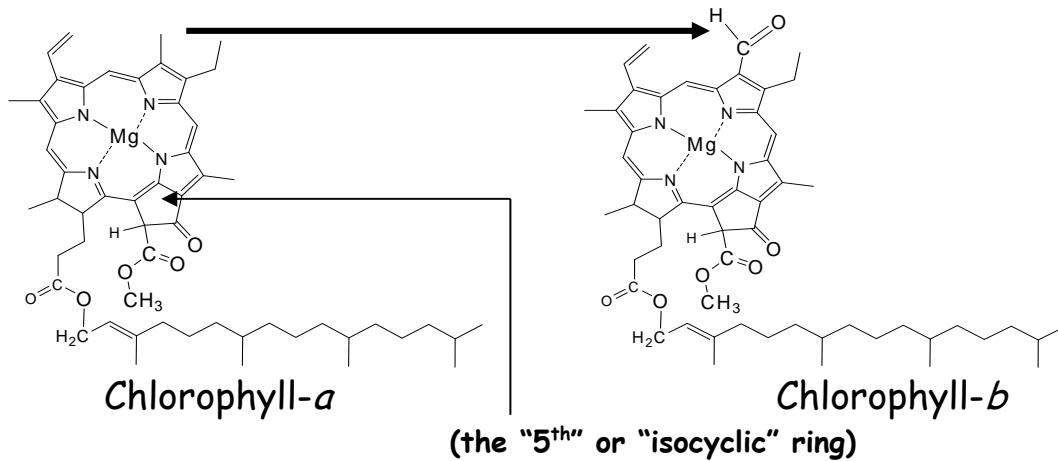


FIGURE R1: Structures of chlorophylls-*a* and -*b*.

Chlorophyll-*a* can either be part of the solar energy-gathering complex or serve as the hinge point for the conversion of that energy to chemical energy for carbon fixation. That is, special chlorophyll-*a* molecules exist in the so-called antenna array. Non-antenna chlorophyll-*a* molecules plus a wide variety of 'accessory' pigments pass 'solar energy', as electron flow to the antenna pigments where photosynthesis is initiated.

Throughout the history of life on Earth, numerous 'accessory' pigments have evolved as bacteria, algae and plants have undergone adaptive radiation into their specific niches. These accessory pigments are able to either absorb parts of the electromagnetic spectrum (~ light for our purposes) which chlorophyll-*a* cannot and/or do so in a more efficient manner. A single chemical change in chlorophyll-*a*, the conversion of a methyl to formyl group (Fig.R1, arrow), gives chlorophyll-*b*, an accessory pigment in green algae (Chlorophyta) and all higher plants.

Photoautotrophs that use water (H_2O) to generate the reductants ($NADP \rightarrow NADPH$) yield molecular oxygen (O_2) as a by-product. This is called **oxygenic photosynthesis**. The presence of **chlorophyll-*a*** reveals the existence of oxygenic photosynthesis and,

even though it can be related ('estimated') to overall photosynthetic biomass, it tells us nothing about the taxonomic makeup of that community. Photoautotrophs that utilize hydrogen sulfide (H_2S) as a source of reductant, giving various forms of sulfur (S_8^0 , *etc.*) as by products, are involved in ***anoxygenic photosynthesis***. These anoxygenic species contain a variety of **bacteriochlorophylls** (Fig.R2; -a in purple-S

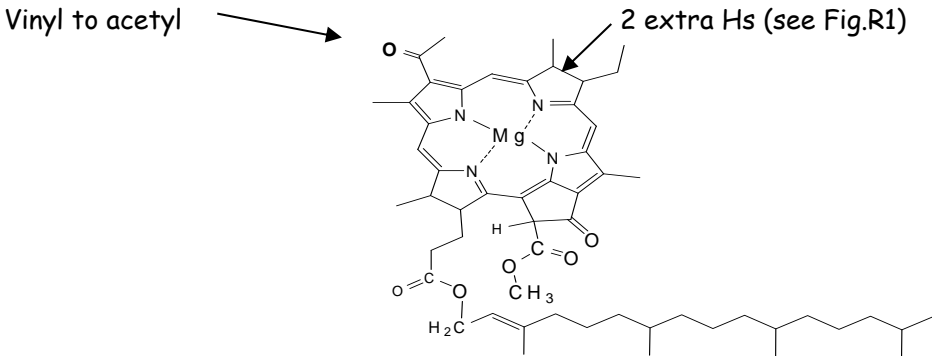


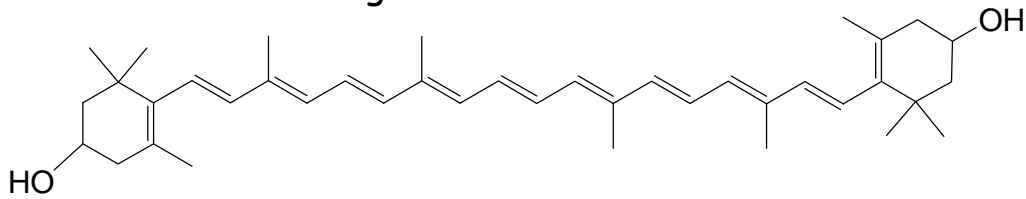
Figure R2: Bacteriochlorophyll-*a* (note extra site of reduction and vinyl group replaced with an acetyl moiety: see Fig.R1)

bacteria, -c/-d/-e/-f/-g in the green and brown sulfur bacteria). Thus, the presence and abundance of chlorophyll-*a* and the bacteriochlorophylls can be utilized to *estimate* the relative importance of oxygenic and anoxygenic photoautotrophs and their organic matter (*e.g.* protein biomass) in an ecosystem.

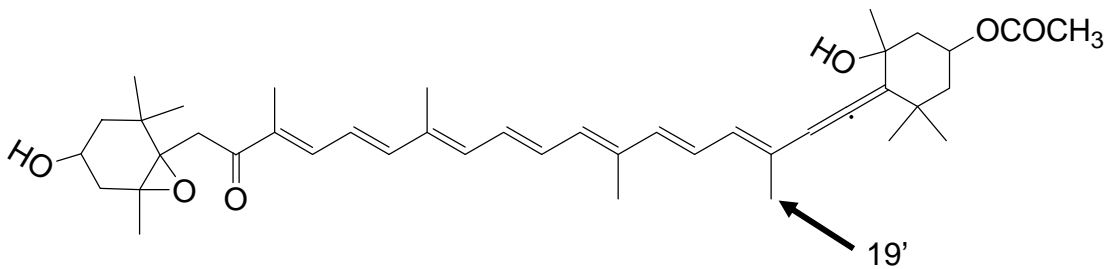
Likewise, one can determine the presence and estimate the abundance of the various taxonomic groups of the oxygenic photoautotrophs through total pigment analysis.

As stated, there are several ***photosynthetic accessory pigments (PAPs)***. This group includes chlorophyll-*b* (chlorophytes, higher plants), fucoxanthin plus the chlorophylls-*c* (chrysophytes, diatoms and relatives), gyroxanthin diester (Florida Red Tide, *Karenia brevis*) peridinin (dinoflagellates), and the divinyl chlorophylls-*a* / -*b* (prochlorophytes). Additionally, there are many taxon specific (or

abundant, "zea") **photoprotectorant pigments (PPPs)**, such as zeaxanthin ("zea", cyanobacteria), myxoxanthophyll (cyanobacteria), keto-carotenoids (echinenone, canthaxanthin: cyanobacteria), lutein (chlorophytes), alloxanthin (cryptophytes), and others. The structures of but a few of these are given below.

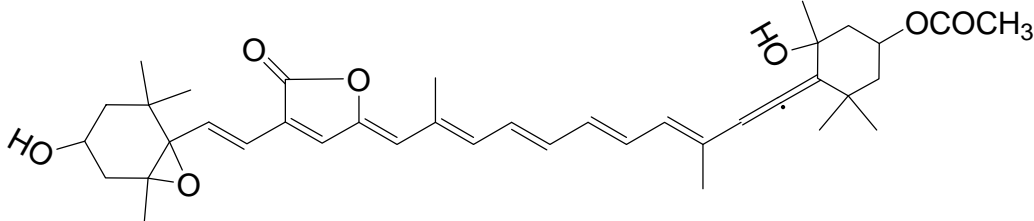


Zeaxanthin (cyanobacteria and a 'bit' in Chlorophytes)

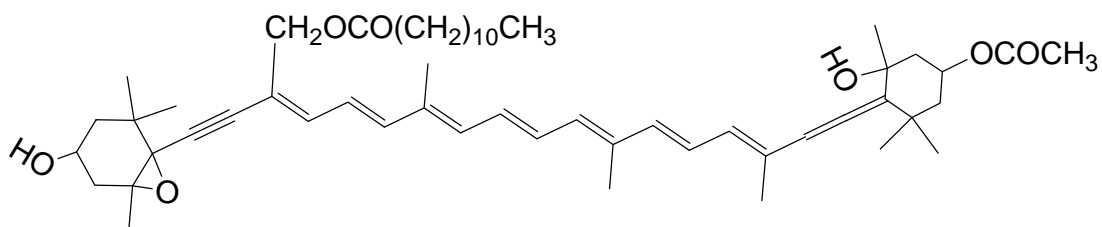


Fucoxanthin (Chrysophytes)

19'-butanoyloxy- and 19'-hexanoyloxy-fucoxanthins (prymnesiophytes)



Peridinin (Pyrrhophyta, Dinoflagellates)



Gyroxanthin diester (Florida Red Tide, *Karenia brevis*: Pyrrhophyta)

Figure R3: Selected chemotaxonomic biomarker carotenoids.

Therefore, through pigment analysis and the relation of the abundance of these 'biomarkers' to chlorophyll-*a*, one is able to estimate the (chemo-) taxonomic structure of microalgal communities. This methodology, pigment-based chemotaxonomy, has gained increasing favor for rapid temporal and spatial investigations of microalgal communities, such as phytoplankton distributions in lakes and oceans. The main premise rests with our ability to assign a numerical relationship between the marker pigment and chlorophyll-*a*, the biomass marker. As an example, the analyses of a great many diatoms reveals that the chlorophyll-*a* to fucoxanthin (molar) ratio is 1.1 : 1. Hence, for every mole of fucoxanthin that we find in a sample, we should have 1.1 moles of diatom-contributed chlorophyll-*a*. This same process is then extended to all of the marker pigments that we isolate and identify from a sample. The percent composition of the community is then calculated by the relative abundances of the taxon-specific chlorophyll-*a*.

An example of the pigment analysis of a very simple phytoplankton community (cyanobacteria / diatom >95% / < 5%) is given below (Fig. R4). This sample, from Snake Bight in north-central Florida Bay was undergoing a cyanobacterial bloom (Total CHL-*a* = 13.3 $\mu\text{g} / \text{L}$) at the time of collection. The great dominance of cyanobacteria was indicated by the abundance of zeaxanthin and only a trace of fucoxanthin (diatoms) could be found.

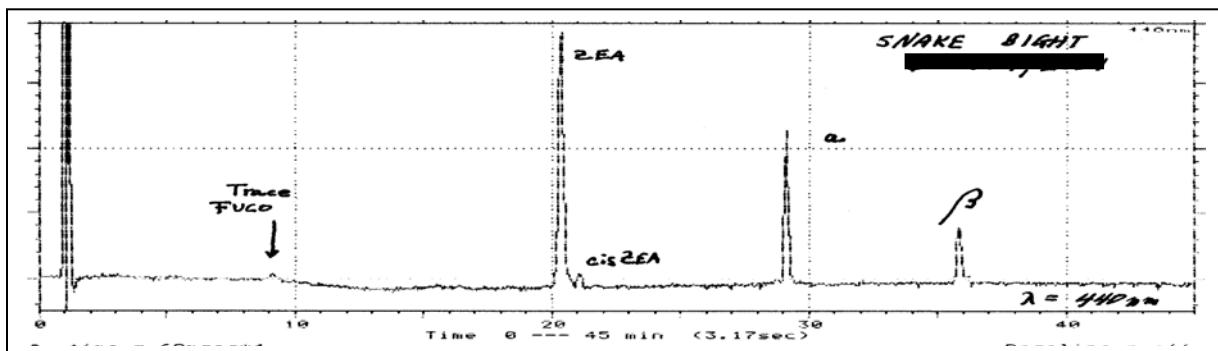


Figure 4: HPLC chromatogram of Snake Bight phytoplankton. Common cyanobacterial bloom in Florida Bay.

A more complicated, but also more interesting, pigment distribution is shown as Figure R5. Here, a mixed community (~ 21/20/44/0/15 % as cyanobacteria / chlorophytes / diatoms / dinoflagellates / cryptophytes) was indicated by the relative abundances of zeaxanthin, chlorophyll-*b*, fucoxanthin, peridinin (absent) and alloxanthin, respectively (Total CHL-*a* ~ 1.5 $\mu\text{g} / \text{L}$).

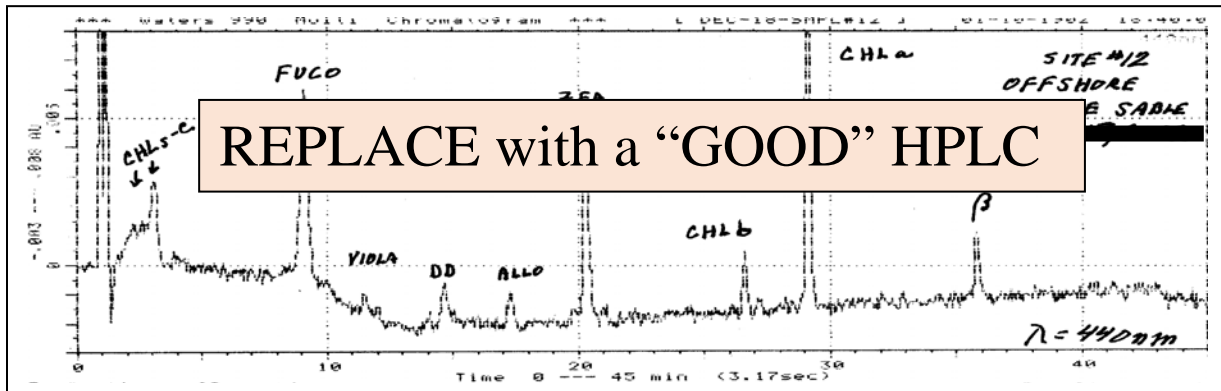


Figure R5: HPLC-PDA chromatogram of phytoplankton pigments from the Gulf of Mexico about 1 mile west of Mid Cape Sable.

Open ocean deep water phytoplankton assemblages, such as the example below (Fig. R6), often contain a mixture of prymnesiophytes and prochlorophytes. These taxa are detected based upon the presence of the 19'-butanoyloxy- and 19'-hexanoyloxy-fucoxanthins or divinyl chlorophyll-*a*, respectively.

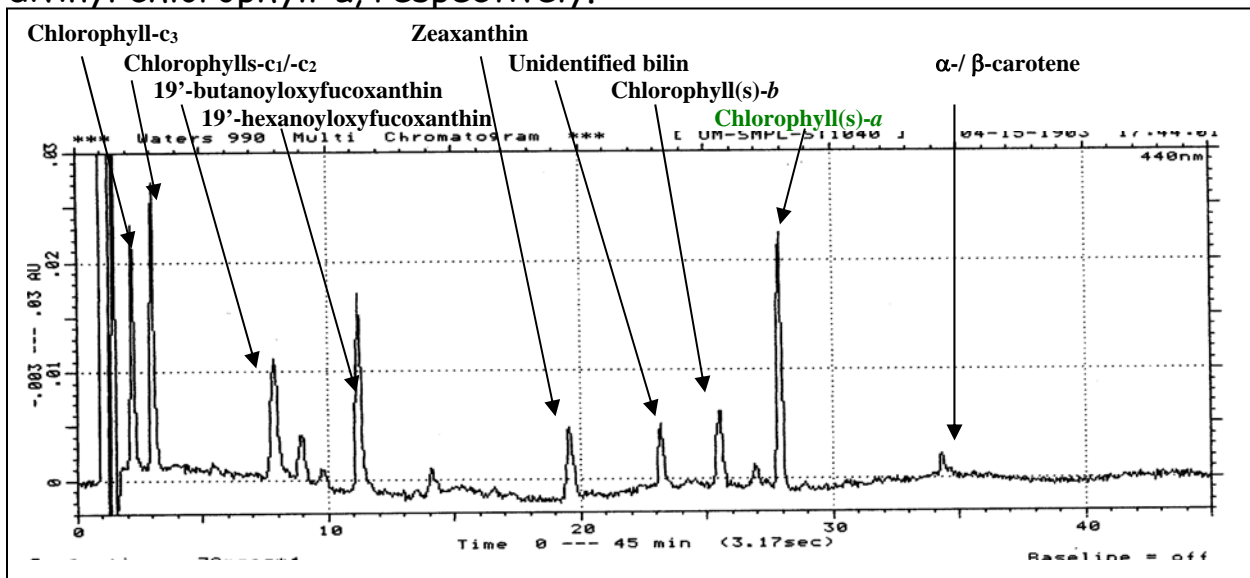


Figure R6: A C-18 Reverse Phase HPLC chromatogram of phytoplankton pigments from a sub-tropical western Atlantic deep water chlorophyll maximum.

As you may have noticed, I mentioned 2 forms of chlorophyll-*a* yet only 1 is seen in the C-18 RP-HPLC chromatogram (Fig.R6) above. This is because C-18 columns fail to resolve 'regular' (MonoVinyl "MV") CHL*a* from divinylchlorophyll-*a* (DV-CHL*a*). However, by running these pigment extracts in C-8 reverse phase columns (Fig. R7), we can easily separate MV and DV CHL-*a*.

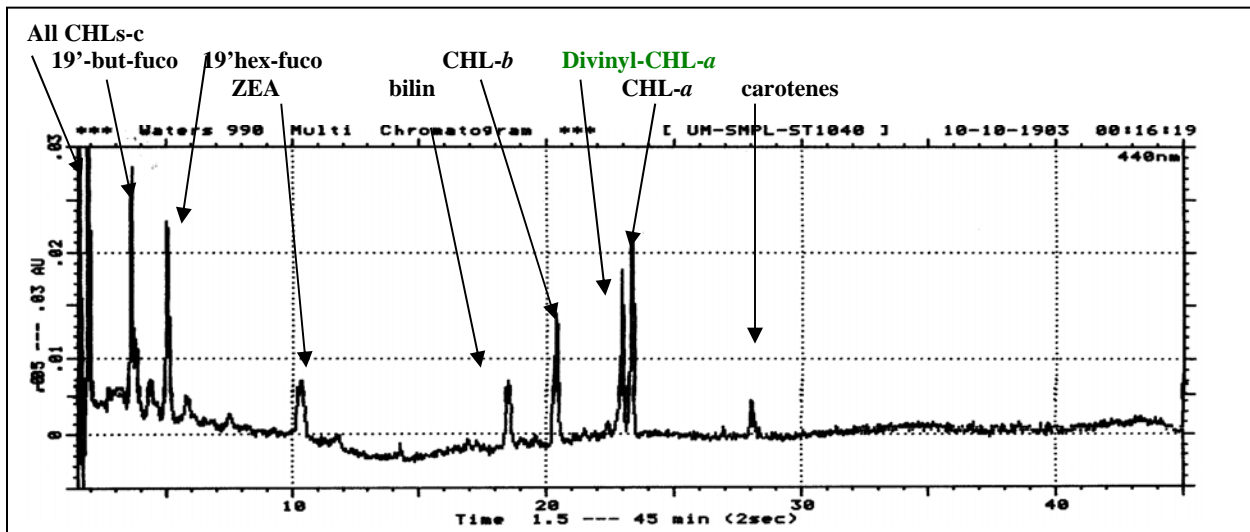


Figure R7: C-8 (octyl-silane) RP-HPLC chromatogram of the same sample given in Figure R6.

Aside from chromatographic retention time, the first dimension of analytical identification, all 'peaks' are also analyzed for their absorption spectrum using full spectral (190-800 nm) photodiode array

or 'PDA' detection. In this manner, the difference between divinyl and 'regular' (monovinyl) chlorophyll-*a* is easily noted (Fig. R8).

Compared to chlorophyll-*a* (Fig. R8: maxima at 430 and 662 nm), divinyl chlorophyll-*a* (maxima at 438 and 662 nm) exhibits an 8 nm red shift in the larger, so-called Soret, absorption band. Thus, the 2 dimensional identification of retention time, on C-8 columns) and the absorption spectrum allows for the identification of DV-CHL*a* and, by chemotaxonomic principles, the Prochlorophyta.

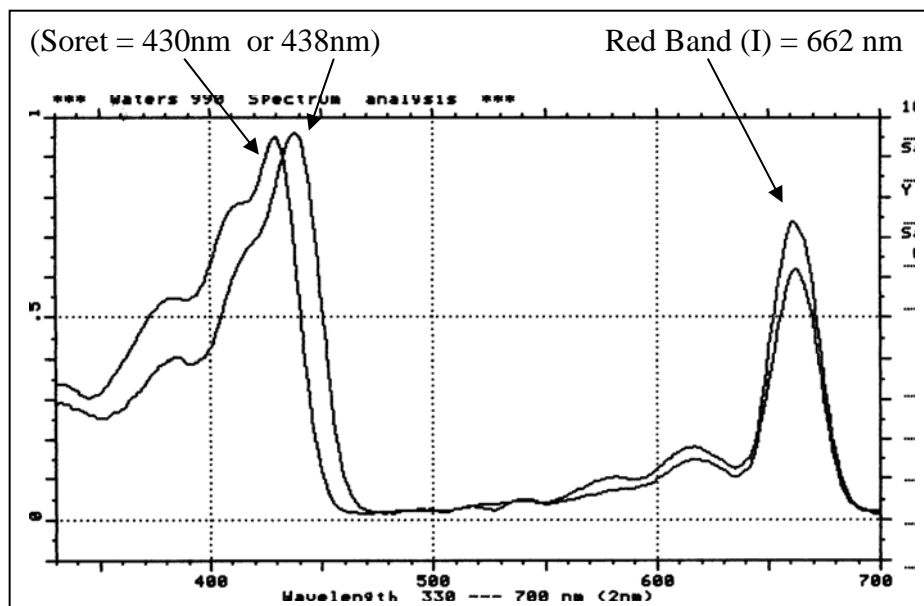


Figure R8: The absorption (PDA) spectra of 'regular'(monovinyl) chlorophyll-*a* (S= 430nm) and of divinyl chlorophyll-*a* (S= 438nm).

Pigment-based chemotaxonomy, while it will *never* replace microscopic examination by a highly skilled taxonomic expert or the newer yet tedious biomolecular methods. **However**, is extremely useful in the rapid spatial and temporal monitoring of ecosystems. That is, the microscopist can take identification to the species level whereas the chemotaxonomic methodology is good only to the Division or Class level. However, pigment analyses are an objective technique whereas

microscopy can be subjective in relation to the number of fields counted, bias/knowledge of the microscopist and other parameters.

Pigment-based chemotaxonomy can be extremely advantageous and cost effective in large ecosystem-scale research and monitoring programs, such as the Comprehensive Everglades Restoration Plan covering the south Florida Ecosystems. Much research remains to be done regarding the effects of light, nutrients, and their synergies on the pigment ratios we decide to use for these (chemo-)taxonomic estimations of microalgal communities. However, as this basic research continues, utilization of these methods for ecosystem surveys is ongoing. Both the basic studies as well as the monitoring efforts are active portions of the research in my laboratory.

Pigment-based chemotaxonomy studies of phytoplankton and epiphytes are being performed in the Atlantic Ocean, Florida Bay other lagoons/ estuaries, Lake Okeechobee, the Bahamas and on periphyton in the Everglades in association with the South Florida Water Management District.

Microalgal culture: In order to study pigment ratios of living microalgae as well as provide living cells with which to begin senescence-death studies (below), we grow pure cultures of algae, as well as purchase, beg, borrow and ~~steal~~ from other sources. By growing our own cells, we can provide a nutrient and light history and one can then begin to ascertain the effects of these parameters and their synergy have upon pigment quantities and ratios in various species. These data will eventually be used to 'fine-tune' pigment-based chemotaxonomy.

Senescence-death related alteration of pigments:

Once algal cells begin to undergo senescence and death, their pigments undergo alterations and eventual recycle. However, the fates and rates of these reactions are different not only starting structure

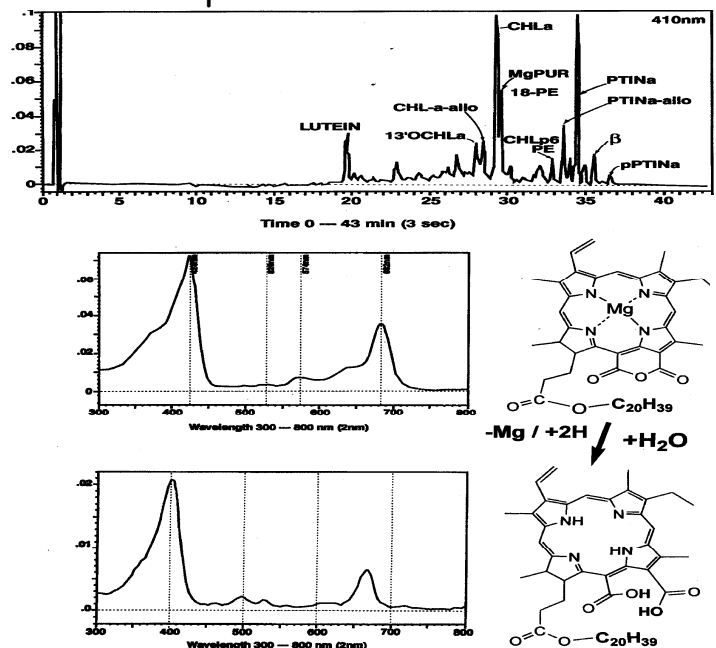
but by the environment (oxygen fugacity, pH, light/dark, pE, grazing etc.) to which they are exposed during these initial phases of recycle.

My interest in the senescence / death related alteration of chlorophylls and carotenoids comes from 2 separate foci.

First, hardly any natural microalgal population consists entirely of log phase perfectly "healthy" cells. That is, certain cells in senescence, dead cells, heterotrophically processed cells (fecal pellets) and even resuspended dead-older cells (sediments) can all be admixed with the healthy population. Thus, to begin to estimate the effects which these degradation processes have on the pigment ratios utilized in our chemotaxonomic estimates of the viable algae, we need information on these changes and relative rates. For example, if fucoxanthin is degraded faster than chlorophyll-*a* in a population of diatoms then the overall diatom chlorophyll-*a* contribution (to total OM, 'living' plus 'dead') will be underestimated if dead diatoms are present.

Second, my studies on the entire geochemistry continuum of chlorophyll, often including the metalloporphyrins in oils and coals, requires knowledge of the very early stages of organic diagenesis. That is, the 'geochemical' reactions which overlap and succeed senescence / death phenomena.

As an example, Figure R9 is an HPLC chromatogram of a culture of the chlorophyte *Closterium sp.* Which was aged (dark with oxygen) for 5 y8m. The phytol esters of Mg-purpurin-18 and chlorin-*p*₆ were found to have been produced from Chlorophyll-*a*. These pigments offer substantial proof for an early loss of the 5th (isocyclic) ring of CHLa



during death -'diagenesis'.

Figure R9: Generation of Purpurin-18.

These *in vitro* studies
 Allow us to trace many of
 The reactions required or
 proposed to occur during
 'real world' geochemical
 processing of natural
 pigment assemblages.

Therefore, laboratory
 experimentation with algal
 cells and also with pure
 compounds afford us
 with a glimpse into the
 workings of nature and
 allow us to extend these
 results to the interpretation
 of biogeochemical processes.

Here, Figure R10, the
 oxidation of chlorophyll-*a*,
 giving purpurin-18 can be traced
 to the eventual production of an "etioporphyrin" (see Louda, 2017 *in*
vita). Note the lack of the "5th" ring (see Figure R1).

There are many many more experiments to do, both with
 microalgal cultures and/or with pure compounds.

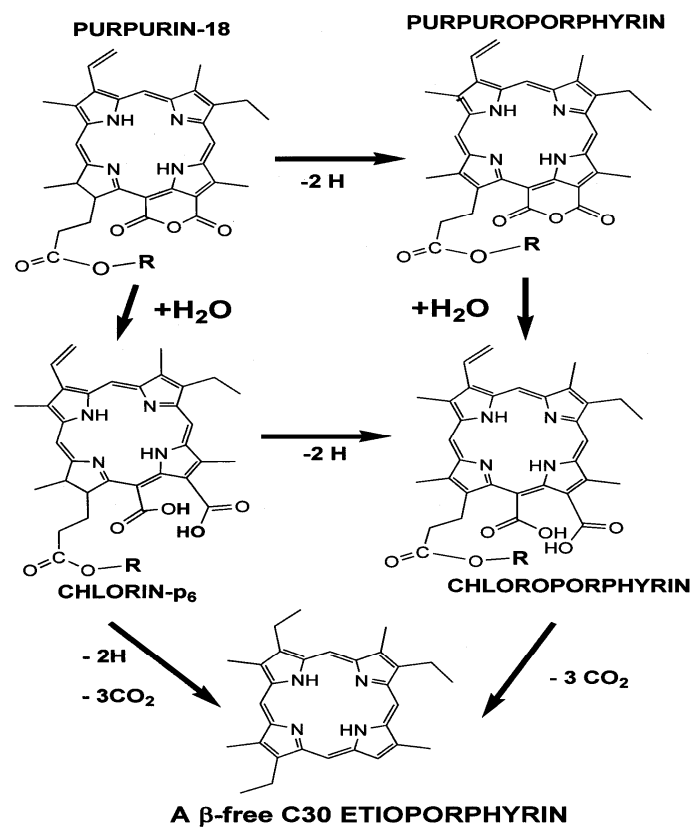


Figure R10: Oxidative changes.

SUNSCREEN PIGMENTS: Many algae and cyanobacteria grow in very high light conditions and can and do experience photoinhibition of their photosynthetic processes. In these cases, many species will generate photoprotectorant pigments. Sometimes, this involves on the increase in the amount of a pigment already present, such as increasing amounts of zeaxanthin in many cyanobacteria. In other cases, such as other cyanobacteria, the synthesis of canthaxanthin (4,4'-diketo-b-carotene) will occur (see Grant and Louda, 2010 in vita.). The cyanobacteria that live in intense solar radiation also generate dimeric indole phenolic pigments known as the scytonemins. Most of the scytonemin nuclei containing sunscreen pigments absorb mainly ultraviolet radiation (UVR). Recently, we have isolated and identified a scytonemin-based pigment that also absorbs strongly in the visible range. This pigment, scytonemin-3a-imine, 'may' be protecting the cytochrome electron transport system (grant and Louda, 2013 in vita). This remains to be proven but the possibility of this is highly intriguing as cells in intense light and desiccating conditions would be destroyed if the cytochromes were excited in non-normal ways and electron flow created oxygen free radicals.

~~~~~CLOSING~~~~~

As you may tell, my research interests span the realm of biological pigments from primary productivity / community dynamics, Everglades and coastal ecosystem restoration / monitoring, the geochemistry of pigments and relation to petroleum / coal generation, to the pure biochemistry and chemistry of the chlorophylls and carotenoids.