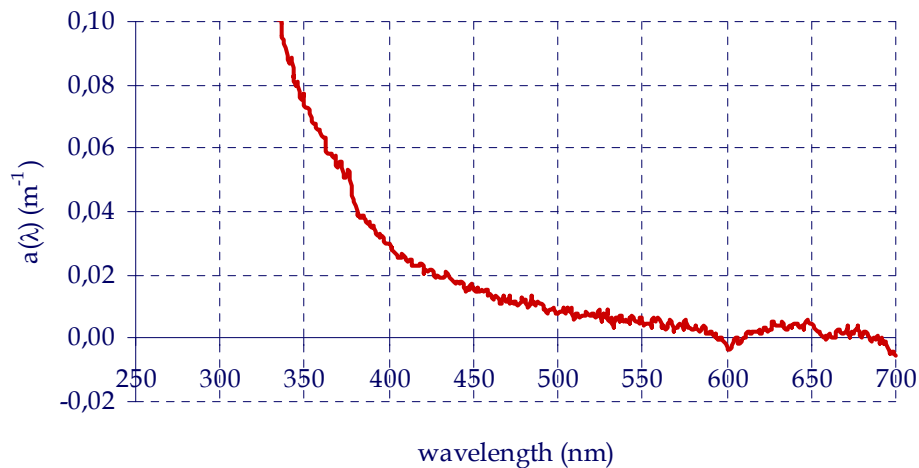
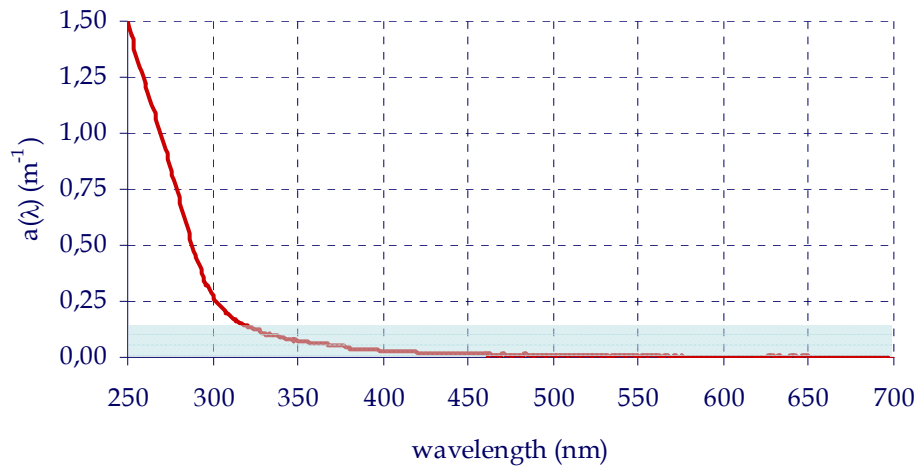


International Training Workshop on CDOM Characterization  
using Spectroscopic Techniques

Granada, 19-21 May 2010

Corrections and Compatibility  
discussion group

## ABSORBANCE



### ► filtration

- GF/F (0.7  $\mu\text{m}$ ) ?
- polycarbonate (0.2  $\mu\text{m}$ )?
- nitrocelulose (0.2  $\mu\text{m}$ )?
- supor (0.2  $\mu\text{m}$ )?

### ► cell path length

- 1 cm ( $\pm 0.2 \text{ m}^{-1}$ )
- 5 cm ( $\pm 0.05 \text{ m}^{-1}$ )
- 10 cm ( $\pm 0.02 \text{ m}^{-1}$ )
- 1-5 m ( $\pm 0.002$  to ( $\pm 0.0005 \text{ m}^{-1}$ ))

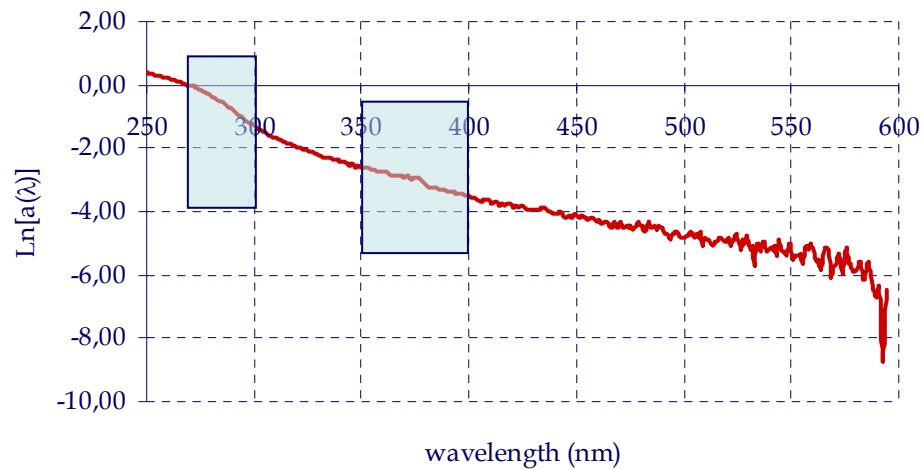
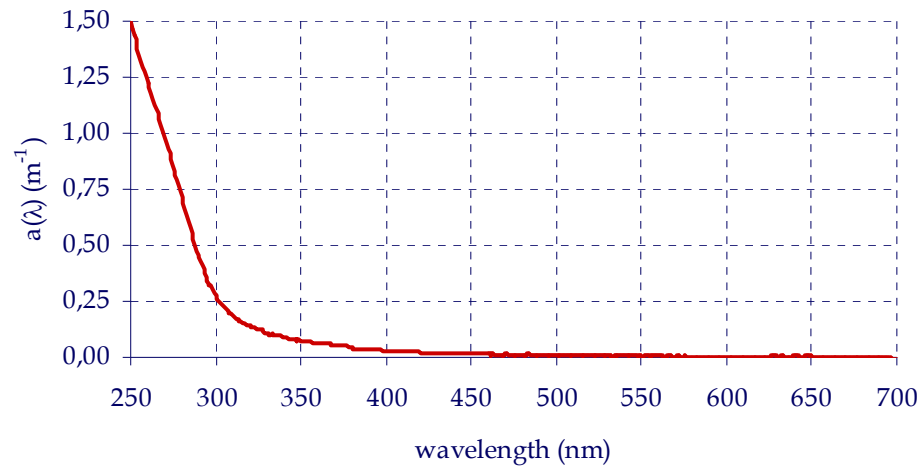
► scan speed (250 nm/min)

► slit width (1 nm)

► Wavelength collection range

- 250 nm – 800 nm

# ABSORBANCE



## ► spectral corrections

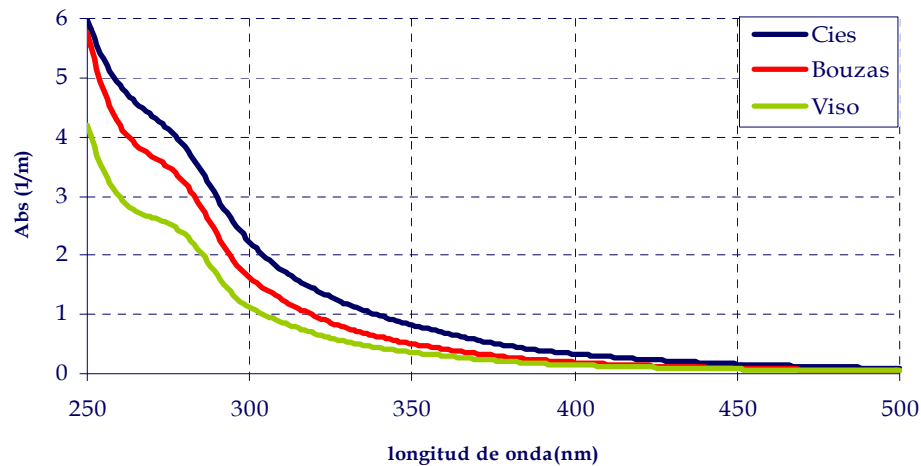
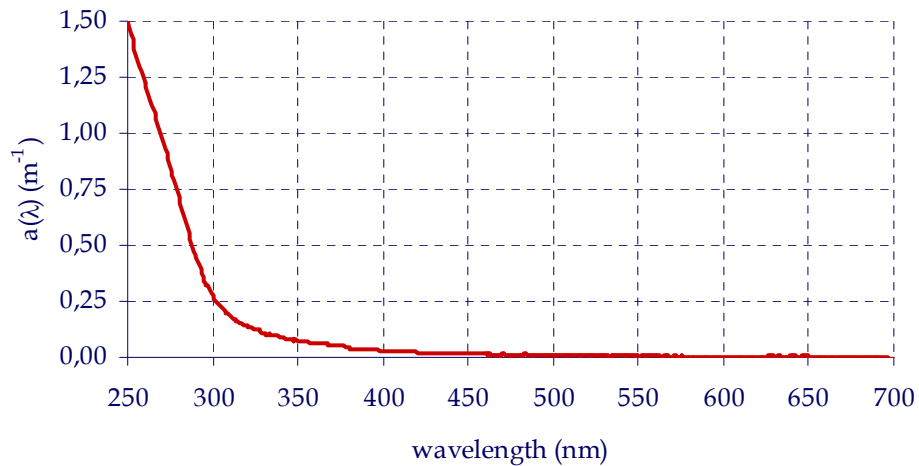
$$\blacktriangleright a(\lambda) = 2.303 \cdot \frac{\text{Abs}(\lambda) - \text{Abs}(600 - 700)}{L}$$

## ► Spectral slope

$$\blacktriangleright a(\lambda) = a(\lambda_0) \cdot \exp[-S \cdot (\lambda - \lambda_0)]$$

$$\blacktriangleright \log[a(\lambda)] = \text{Ln}[a(\lambda_0)] - S \cdot (\lambda - \lambda_0)$$

# ABSORBANCE



## ► spectral corrections

$$\blacktriangleright a(\lambda) = 2.303 \cdot \frac{\text{Abs}(\lambda) - \text{Abs}(650 - 750)}{L}$$

## ► Spectral slope

$$\blacktriangleright a(\lambda) = a(\lambda_0) \cdot \exp[-S \cdot (\lambda - \lambda_0)]$$

$$\blacktriangleright \log[a(\lambda)] = \text{Ln}[a(\lambda_0)] - S \cdot (\lambda - \lambda_0)$$

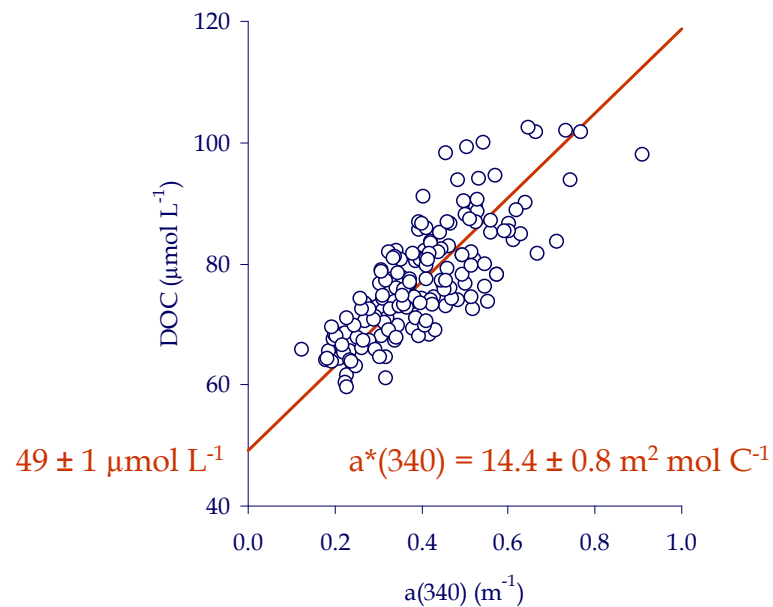
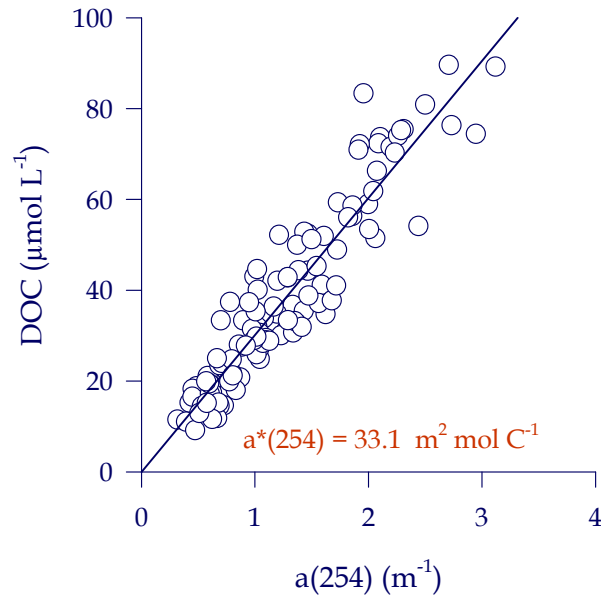
## ► Non exponential decay

## ► Spectral indices

$$\blacktriangleright a(\lambda), \text{ with } \lambda = 254, 325, 340, 350, 355, 443 \text{ nm}, \dots$$

$$\blacktriangleright a^*(\lambda), \text{ e.g. SUVA} = a^*(254)$$

# ABSORBANCE



## ► spectral corrections

$$\blacktriangleright a(\lambda) = 2.303 \cdot \frac{\text{Abs}(\lambda) - \text{Abs}(600 - 700)}{L}$$

## ► Spectral slope

$$\blacktriangleright a(\lambda) = a(\lambda_0) \cdot \exp[-S \cdot (\lambda - \lambda_0)]$$

$$\blacktriangleright \log[a(\lambda)] = \text{Ln}[a(\lambda_0)] - S \cdot (\lambda - \lambda_0)$$

## ► Non exponential decay

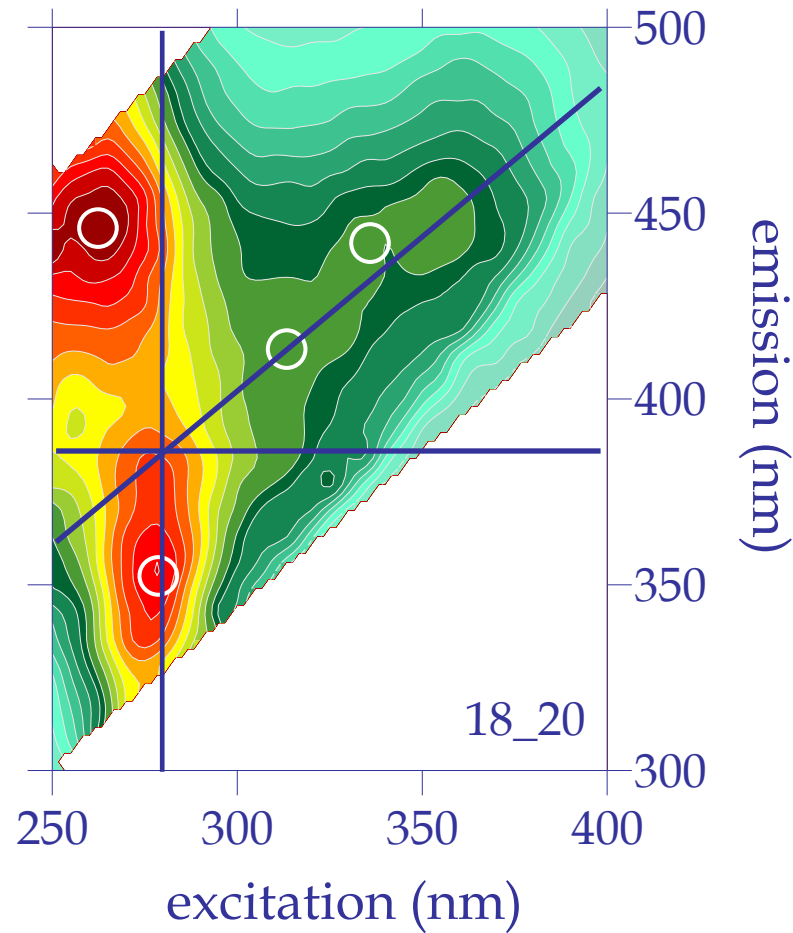
## ► Spectral indices

$$\blacktriangleright a(\lambda), \text{ with } \lambda = 254, 325, 340, 350, 355, 443 \text{ nm}, \dots$$

$$\blacktriangleright a^*(\lambda), \text{ e.g. SUVA} = a^*(254)$$

$$\blacktriangleright a(254)/a(365)$$

# FLUORESCENCE



▶ filtration/preservation

▶ cell path length (1 cm)

▶ integration time (0.25-0.50 s)

▶ slit width

▶ excitation 1.5 -10 nm

▶ emission: 1.5 -10 nm

▶ data acquisition

▶ excitation/emission pairs

▶ Exc/Emm indices (e.g. F.I.)

▶ excitation spectra

▶ emission spectra

▶ synchronous spectra

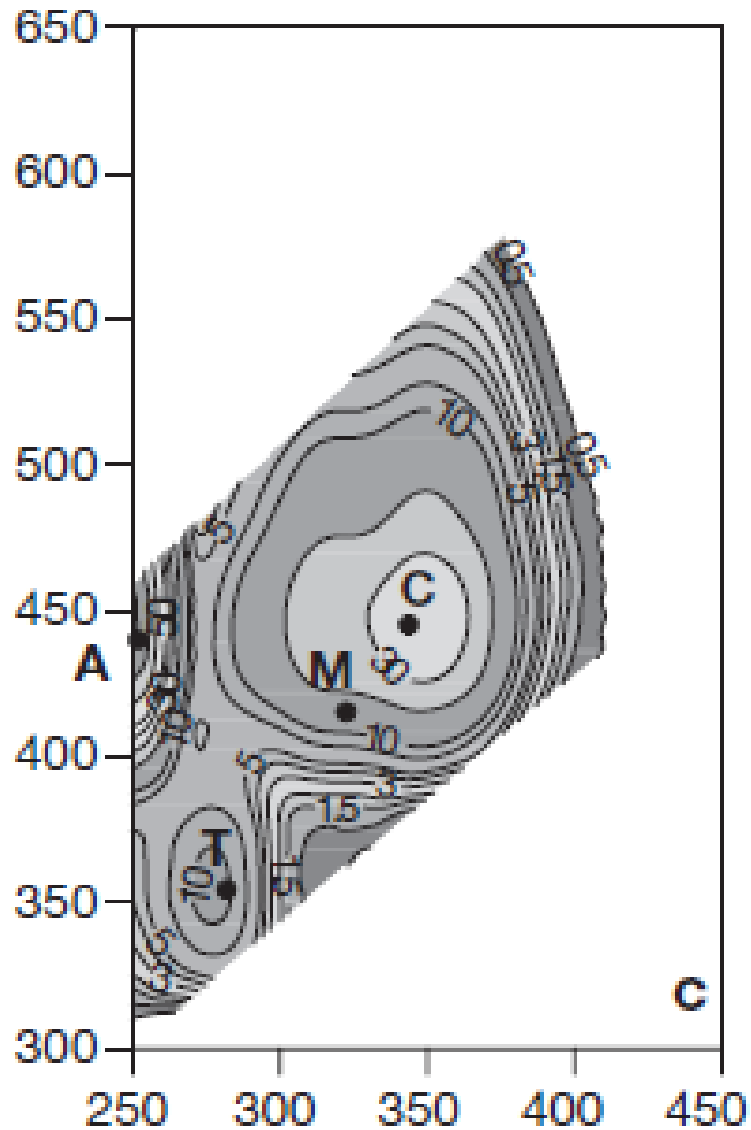
▶ excitation/emission matrices

▶ excitation

▶ emission

▶ synchronous

## FLUORESCENCE



### ▶ spectral corrections

- ▶ Rayleigh scatter
- ▶ Raman scatter
- ▶ inner filter effect ( $> 10 \text{ m}^{-1}$ )
- ▶ excitation spectral correction
- ▶ emission spectral correction

### ▶ calibration

- ▶ Raman peak height/area
- ▶ QS units (Exc/Em = 350/450)
- ▶ QS units (specific Exc/Em)
  - ▶ QS units / Trp units
- ▶ natural CDOM (IHSS?)