SPECTROSCOPY OF PHOTOSYNTHETIC PIGMENT-PROTEIN COMPLEX LHCII

States 6

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Outline:

1. What is LHCII?

2. Why study LHCII?

3. Spectroscopy of LHCII





Photodegradation







Illumination during a day





Whole plants





Oxalis oregana









Excitation quenching

Fluorescence spectroscopy of

LHCII







Absorption spectra of pigments bound to LHCII





Fluorescence lifetime chlorophyll *a* in LHCII





FLIM single molecule of LHCII

Ex 470 nm





Ex 635 nm





W.I. Gruszecki et al., J. Plant Physiol. 167 (2010) 69-73.



FLIM single molecule of LHCII



W.I. Gruszecki et al., J. Plant Physiol. 167 (2010) 69-73.





FLIM single LHCII trimer

Ex 470 nm





Ex 635 nm





W.I. Gruszecki et al., J. Plant Physiol. 167 (2010) 69-73.





Fluorescence spectra of single LHCII particles







Molecular mechanisms

Raman spectroscopy of LHCII

Pigment absorption spectra



Neoxanthin 488 nm * Violaxanthin 492 nm Lutein 489 nm Lutein 495 nm

according to R. Croce et al., Photosynth. Res. 64 (2000) 221-231.



Raman spectra of carotenoids







Carotenoid fluorescence in LHCII



W.I. Gruszecki et al., J. Phys. Chem. B 113 (2009) 2506-2512.









Molecular mechanisms

FTIR spectroscopy

of LHCII





Protein structure

 α -helix





















Blue-light-induced reorganization of LHCII







FLIM LHCII aggregated structures

Ex 470 nm Ex 635 nm













Conclusions:

1. Illumination of LHCII drives molecular configuration changes of xanthophylls

2. Xanthophyll configuration changes drive reorganization of LHCII

3. Reorganization of LHCII leads to excitation quenching



Thank you for attention!