

Průběh obhajoby diplomové práce:

- 1/ Outline of the thesis given by F.K.: introduction CP43
intrinsic - binds PSII STSD
PsaD; PsaB chlorophyll
6 transmembrane α -helices
- 2/ Preparation of CP43 protein - recombinant CP43ect
cloning
overexpression: design of the CP43ect construct
→ recombinant protein GST HIS TCT CP43ect,
Protein expression into Incl. broth from protein library etc
- 3/ Refolding of misfolded protein it did work whereas urea/guanidinium
chloride
→ amyloid sarcosine
- 4/ His-tag affinity chromatography upon trouble cleavage
Protein purification
- 5/ MALDI TOF: confirmation of the sequence integrity
- 6/ Stability testing; Unstable N-term

Body:

Klasifikace: EXCELLENT

Celková klasifikace:

Datum obhajoby: 29/3/2019

- 7/ NMR measurements
preparation of samples
• 2D spectra
• 3D spectra

W. Zurek
podpis předsedy

Hodnocení ústních zkoušek:

- 1/ Resonance assignments
↳ in CARS
- 2/ Crystal-FT structure
predicted α -helix
- 3/ Summary
Extrinsic domain CP43
↳ stable sample for NMR
NMR monitored titrations
with the PsaB
- Discussion: the lowest concentration of
protein for NMR? (≈ 0.1 mol/L)
• MALDI TOF fragmentation?