

Mitochondria – Chloroplast interactions: Acclimation of a Nicotiana sylvestris mitochondrial mutant to growth light intensity.

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Mitochondria - Chloroplast interactions:

Acclimation of a Nicotiana sylvestris mitochondrial mutant to



growth light intensity.



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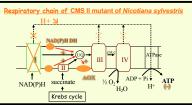
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The CMS II mutant of Nicotiana sylvestris is functionally defect in complex I of the mitochondrial electron transport chain (1) but has a slightly higher dark respiration rate. Furthermore, enzymes of the alternative respiratory pathway (NAD(P)H DH, AOX) are elevated in CMS as compared to the WT (see figure).

As mitochondrial function interacts with photosynthetic metabolism, this mutant was used as a tool to investigate the importance of mitochondria in photosynthetic acclimation to light.



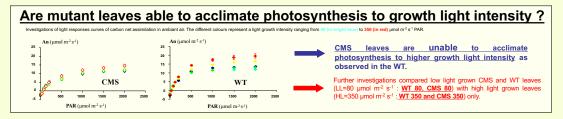
Spectrophotometric assays

RubisCO / NADP-MDH / SPS Chl a/b

Acclimation to growth light intensity

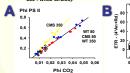
oCO₂/H₂O exchanges Chl Fluorescence

(Licor 6400-40, Mini PAM, PAM 2000)



Are CMS leaves affected in primary photochemistry which might limit carbon assimilation?

ren quantum yields of PSII and of CO_2 assimilation calculated from light responses curves performed under 21% O_2 and 380 tresponse curve of excess electron transport not used for carbon assimilation calculated as ETR – $(4^+\Lambda_1+R_2)$ assuming that 4 assamy to assimilate one molecule of CO_2 (WT 80 : black squares ; CMS 80 : black circles ; WT 350 : white squares ; CMS





(A) Approximately 15 photons are used for the assimilation of one molecule of CO2 in WT 80, WT 350 and CMS 80 but 23 photons are

(B) Excess ETR is higher in CMS 350 than in WT 350 requiring additional electrons sinks.

Are CMS leaves affected in secondary photosynthetic reactions limiting carbon assimilation?

Absolute and relative stimulation of initial Rubisco and SPS activities by high light (350 µmol m² s¹) as compared to low light growth (80 µmol m² s¹) in WT and CMS leaves measured either after one hour at growth light intensity or after 3h illumination at 100 µmol m² s² i PAR

Rubisco			
growth light	μmol m ⁻² s ⁻¹	13.9	3.3
	% stimulation	78	20
3h 1000 μmol m ⁻² s ⁻¹ PAR	μmol m ⁻² s ⁻¹	36.8	14.4
	% stimulation	183	58
SPS			
growth light	μmol m ⁻² s ⁻¹	0.59	0.12
	% stimulation	166	20

- Stimulation of Rubisco and SPS initial activity by a higher growth light intensity is impaired in CMS
- Maximal activities are very similar between CMS 80 and WT 80 and increased in both genotypes in response to higher growth light intensity (data not

Is the acclimation response of CMS leaves affected by different redox homeostasis possibly regulating gene expression?

ightharpoonupInvestigation of $q\underline{P}$ as redox parameter of the ETR chain and $\underline{NADP-MDH}$ as redox parameter of the chloroplast stroma.

A: Light response curve of the photochemical quenching (qP) reflecting the redox state of the plastoquinone pool (=1-qP). B: nihial and total activities of NADP-MDH (µm) mf x² s²) measured at growth light intensity after 1 h in the light period and after 3h illumination at 1000 µmol m² s² l*PAR and 25 °C.

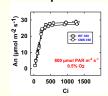




•Redox parameters are not affected in CMS as compared to the WT.

The mismatch of ETR, carbon assimilation and initial activity of Rubisco is further 4 investigated in CMS 350 leaves in order to characterize alternative electron sinks such as:

(a) Photorespiration:



 Under non photorespiratory conditions, the difference between WT 350 and CMS 350

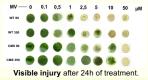
- 1. CMS and WT leaves have the same internal leaf CO₂ concentration (Ci) suggesting
- 2. Glycolate oxidase activity is about 15% higher in the CMS 350 than in the WT 350 after 1h of growth light.
- 3. WT 350 and CMS 350 have a similar specificity factor of Rubisco.

(estimated from determination of Γ^*)

4. In accordance with calculations from models, first experimental determinations of chloroplastic CO₂ concentration (Cc) by mass spectroscopy revealed a lowe in CMS 350 than in the WT 350.

(b) oxidative metabolism:

- → wolecular biology and exidative stress physiology are investigate to see whether excess electrons in the mutant induce higher oxidative stress tolerance.
- Chl content, Fv/Fm and Catalase activity are less affected by MV treatment in CMS plants than in WT plants independent of
- The inhibition of electron transport and carbon net assimilation is delayed in CMS plants as compared to WT plants when



These results show that mitochondrial function is necessary for acclimation of plants to growth light intensity and for oxidative stress response.

CMS plants do not acclimate to higher growth light intensity as the WT. This is shown by a lower quantum yield of CO2 assimilation under ambient air conditions and lower activation state of photosynthetic enzymes. Currently, the fate of excess chloroplastic electrons in CMS plants grown at 350 µmol m⁻² s⁻¹ as compared to the WT is investigated and tend to favour an higher photorespiration in the CMS 350.

CMS leaves showed higher resistance to MV-induced oxidative stress as indicated by lower: chlorophyll bleaching, photoinhibition of PS II and catalase activity. Furthermore, CMS leaves maintained higher rates of total chloroplastic electron transport and CO2 assimilation after the same time of incubation and the same concentration of MV, suggesting that altered transcript and/or protein contents of some antioxidative enzymes (SOD, APX, GR and CAT) are involved in oxidative stress resistance, by confering a more elevated protection in CMS than in the WT.