

# Optimization theories for saline environments

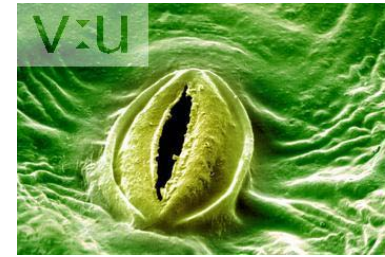
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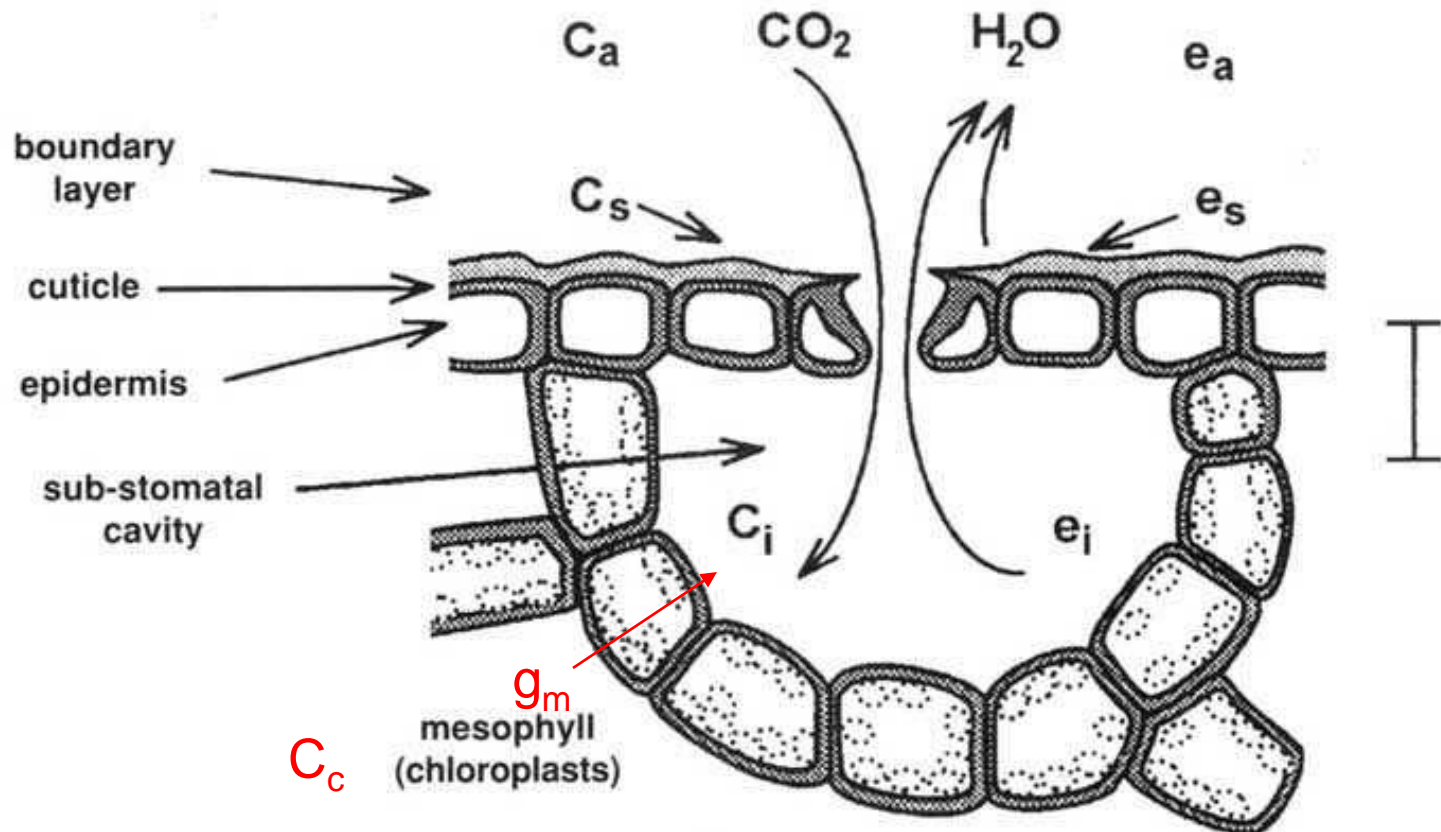
Venice, 18/06/2010

# Motivations

- It has long been suggested that, at the leaf scale, natural selection may have operated to provide increasingly efficient means of controlling the tradeoffs between water vapor loss and carbon gains.
- How does **salinity** modify this picture?

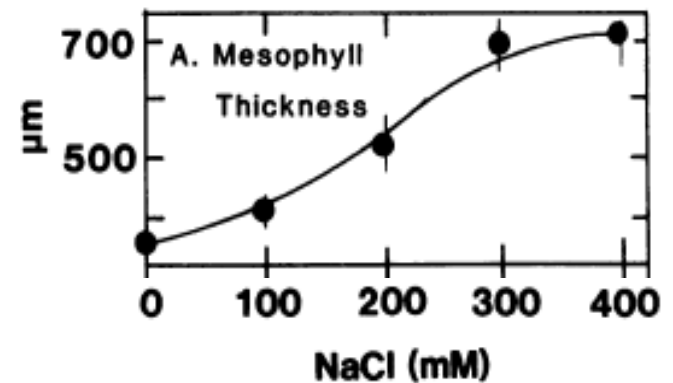
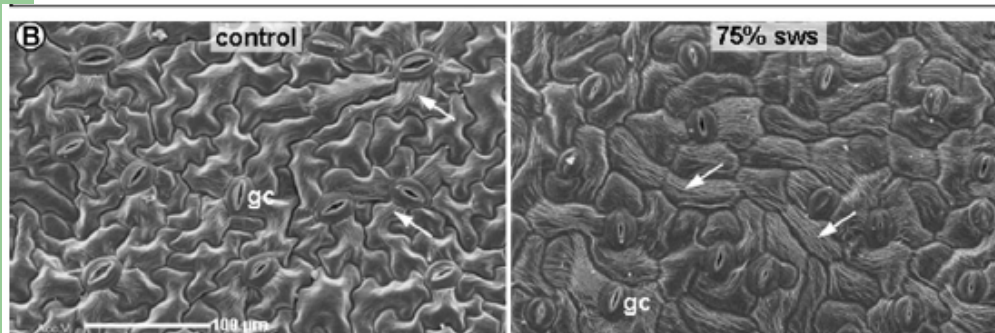


# Photosynthesis - Transpiration



# Structural modifications

	NaCl (molal)					
	0.0	0.05	0.1	0.2	0.3	0.4
Mesophyll thickness ( $\mu\text{m}$ )						
Bean	150	165	260			
Cotton	209	256	329	373	422	
<u>Atriplex</u>	210	--	210	212	260	340



# Optimization model (no mesophyll limitations)

Define the instantaneous:

$$\text{Carbon Gain} = f_c$$

$$\text{Water Loss} = f_e \approx a g_s D$$



John Dalton

OBJECTIVE FUNCTION

$$f(g_s) = f_c(g_s) - \lambda f_e(g_s)$$

Lagrange Multiplier

Leaves are autonomous and attempt to maximize their own carbon gain for a given amount of water loss.



# Basic equations:

Without Salinity

With Salinity

Transport  
Equation:

$$f_c = g_s (C_a - C_i) \quad f_c = g_{eff} (c_a - c_c); \quad g_{eff} = \frac{g_c g_m}{g_c + g_m}$$

Biochemical  
Demand:

$$f_c = \frac{\alpha_1 (C_i - \Gamma^*)}{C_i + \alpha_2} \quad C_i \longrightarrow C_c$$

Optimality Rule:

$$\frac{\partial (f_c(g_s) - \lambda f_e(g_s))}{\partial g_s} = 0$$

Three equations with 4 unknowns ( $f_c$ ,  $C_c$ ,  $g_s$ ,  $g_m$ ) – mathematically unclosed

# Optimization models (linear form) With mesophyll conductance

- Upon differentiating  $f(g_c)$  and setting it to zero:

$$f_c = \frac{a_1 g_m (c_a - c_p)}{a_1 + g_m (a_2 + s c_a)} \left[ 1 - \sqrt{\frac{a \lambda D}{c_a - c_p}} \right]$$

$$g_c = \frac{a_1 g_m}{(a_1 + g_m (a_2 + s c_a))} \left( -1 + \sqrt{\frac{c_a - c_p}{a \lambda D}} \right)$$

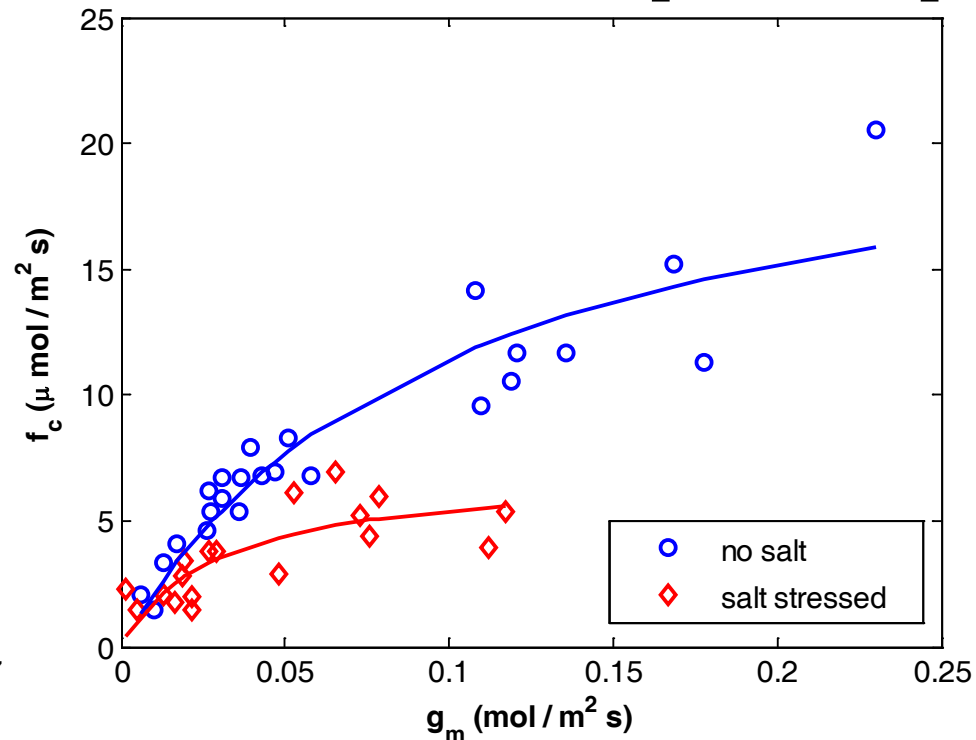
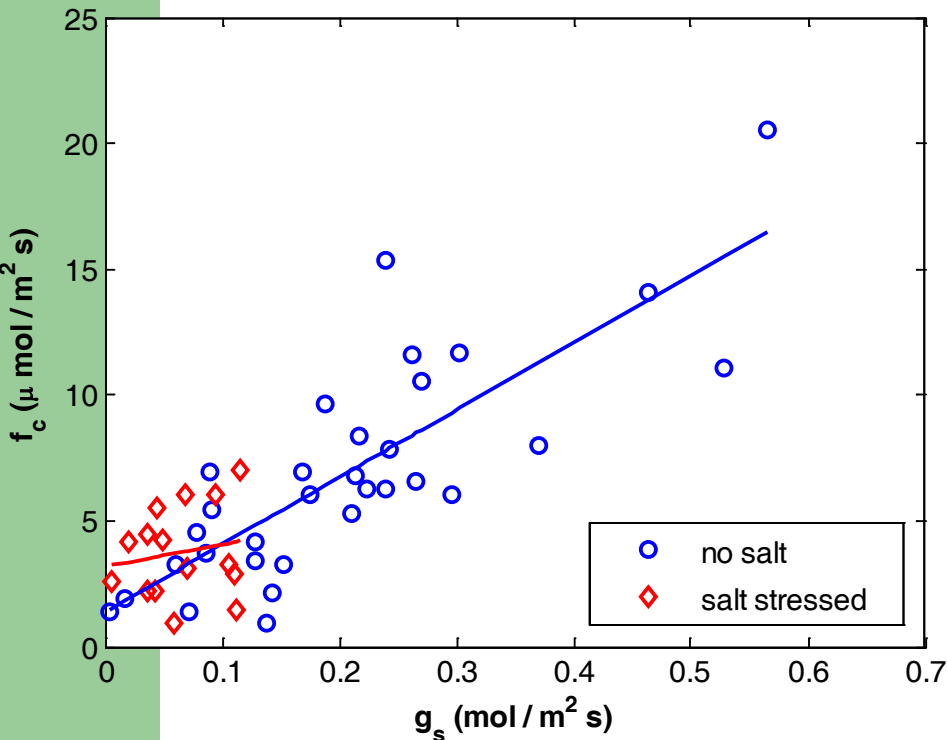
$$\frac{c_i}{c_a} = \frac{c_p}{c_a} + \frac{(c_a - c_p)}{c_a} \left[ 1 - \sqrt{\frac{(a \lambda D)}{(c_a - c_p)}} \right]$$

# Preliminary Results (1)

- Olives (Loreto et al., 2003)

$$f_c = g_c (c_a - c_p) \left[ \sqrt{\frac{a\lambda D}{(c_a - c_p)}} \right] = g_c \left[ \sqrt{a\lambda D (c_a - c_p)} \right]$$

$$f_c = \frac{a_1 g_m (c_a - c_p)}{a_1 + g_m (a_2 + s c_a)} \left[ 1 - \sqrt{\frac{a\lambda D}{c_a - c_p}} \right]$$



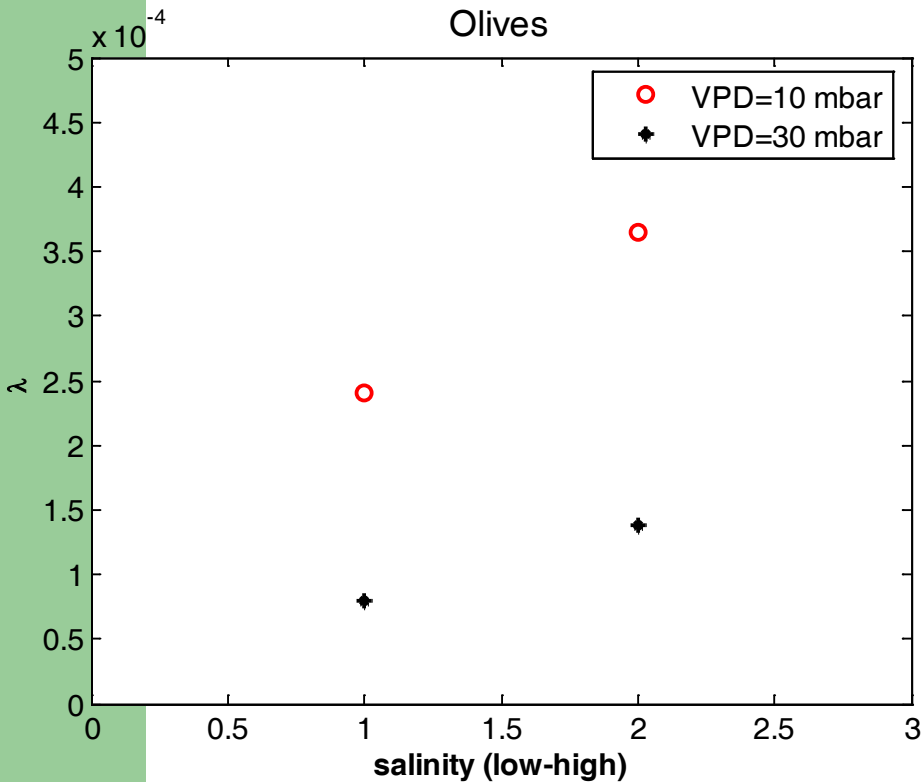


# Preliminary Results (2)

$$\frac{c_i}{c_a} = \frac{c_p}{c_a} + \frac{(c_a - c_p)}{c_a} \left[ 1 - \sqrt{\frac{(a\lambda D)}{(c_a - c_p)}} \right]$$

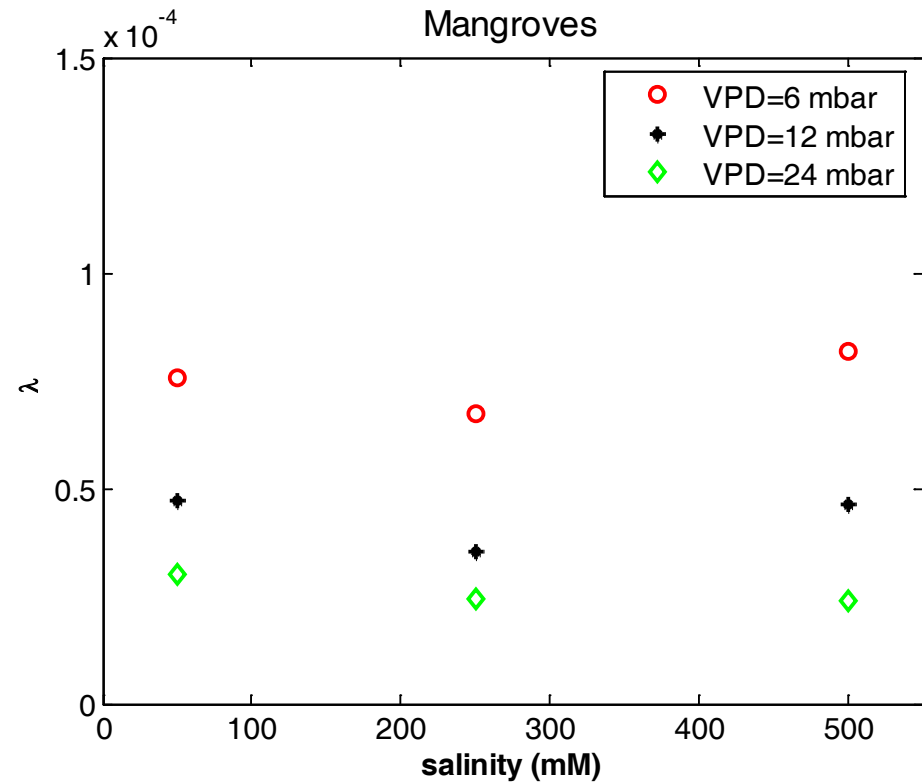
Bongi and Loreto, 1989

Olives



Ball and Farquhar, 1984

Mangroves



# Preliminary conclusions

- 2 time scales of stomata response to increasing salinity:
  - FAST t.s.: stomatal conductance
  - SLOW t.s.: mesophyll conductance
- Optimization theory consistent with datasets analyzed
- Three parameters were used to analyze salt tolerance by different species (olives-mangroves):
  - $g_m$ : changes in all species
  - $\lambda$ : changes only for salt-intolerant species
  - $a_1$ : changes for the olive case

# Futher work

- Short Term: literature review to assess how salinity affects the optimization theory parameters for different species
- Long Term: integrate this results with on-going work on hydrologic models of the soil-plant system to assess how climate change (e.g. CO<sub>2</sub>, VPD, T) affects plant productivity in salt environments

# References

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- Geissler N., Hussin S. and Koyro H.W. (2009). Elevated atmospheric CO<sub>2</sub> concentration ameliorates effects of NaCl salinity on photosynthesis and leaf structure of *Aster tripolium L.*, *Journal of Experimental Botany*, Vol. 60, No. 1, pp. 137–151, 2009
- LORETO F., CENTRITTO M. and CHARTZOULAKIS K. (2003) Photosynthetic limitations in olive cultivars with different sensitivity to salt stress, *Plant, Cell and Environment*, 26, 595–601
- Katul G., Manzoni S., Palmroth S., and Oren R. (2010). A stomatal optimization theory to describe the effects of atmospheric CO<sub>2</sub> on leaf photosynthesis and transpiration. *Annals of Botany* 105: 431–442.