

Biomonitoring of Urban Habitat Quality by Anatomical and Chemical Characteristics of Plant Foliage

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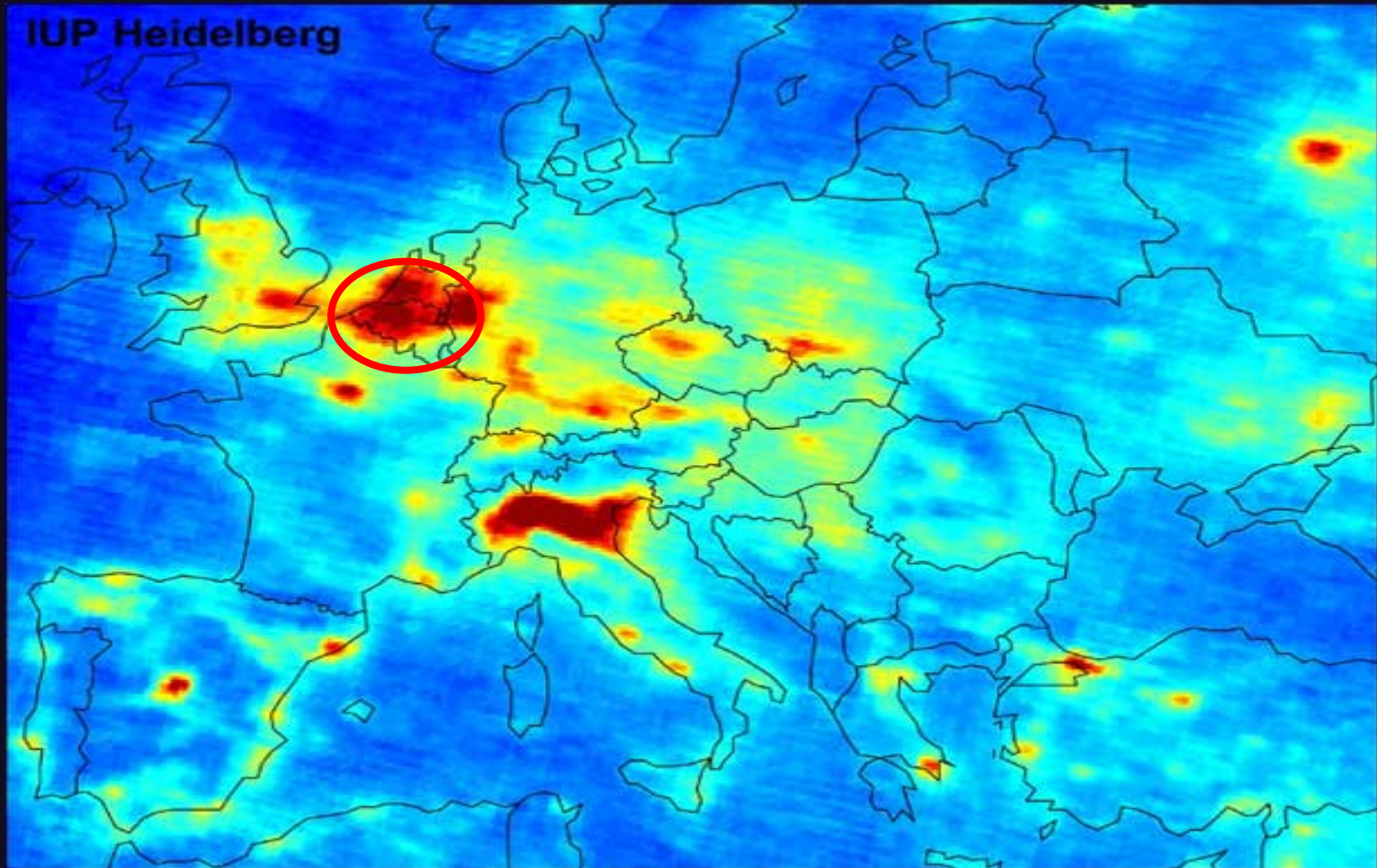


Introduction

- Urban Habitat Quality
 - Air/ Soil/ Water quality
- Physico-Chemical Monitoring
 - High-tech equipment and methods
 - High cost
 - Based on single pollutants:
 - SO_2 , NO_x , O_3 , CO_2 , PM_{10} ,...
 - Eg: Flemish Environmental Agency (VMM)

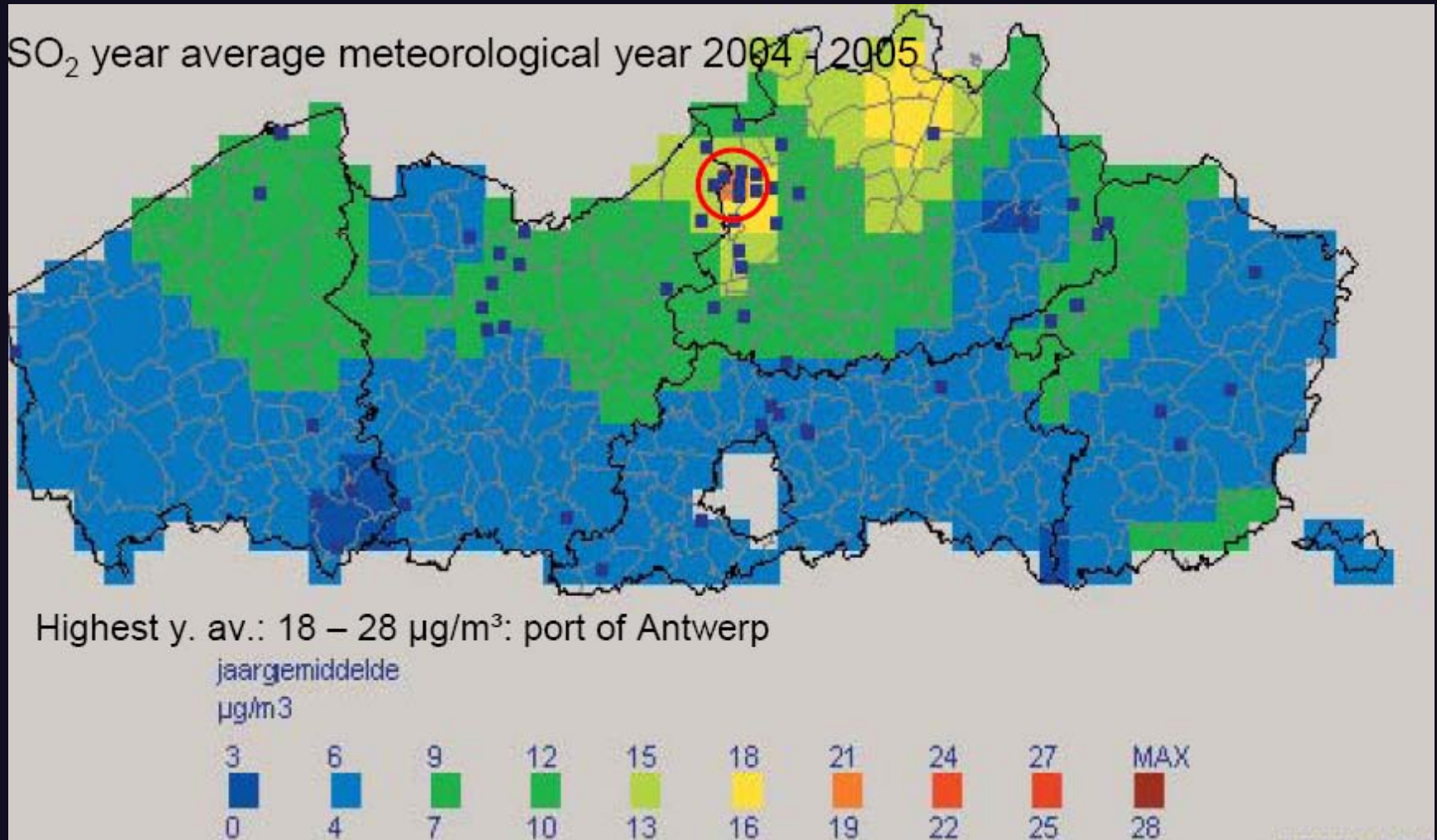
Introduction contd..

NO₂ ☹️



Introduction contd..

SO₂ ☹️

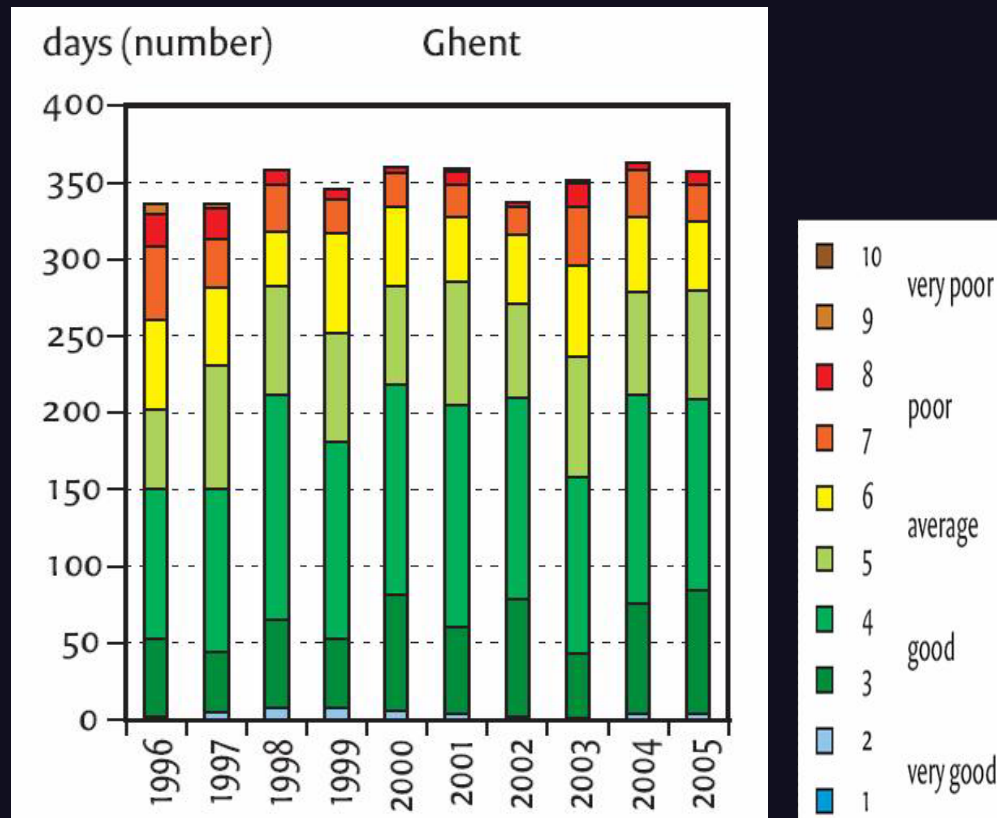


Source: IRCEL CELIN

Introduction contd..

Air Quality Index

-Calculated to integrate several pollutants



Source:IRCEL, WWW.vmm.be/mira

Introduction contd..

Biomonitoring

- combined effect – additive/synergistic/antagonistic
- low-tech equipment and methods
- less cost – large data base

- Bioaccumulation

Trace element accumulation : Lichens, *Quercus illex*

- Plant responses

Physiological responses

eg: **photosynthesis and stomatal conductance**

Morphological and anatomical characteristics

- **Foliar injuries, leaf area**
- **Specific leaf area, stomatal density, stomatal pore surface**

Objectives

To evaluate the potential of

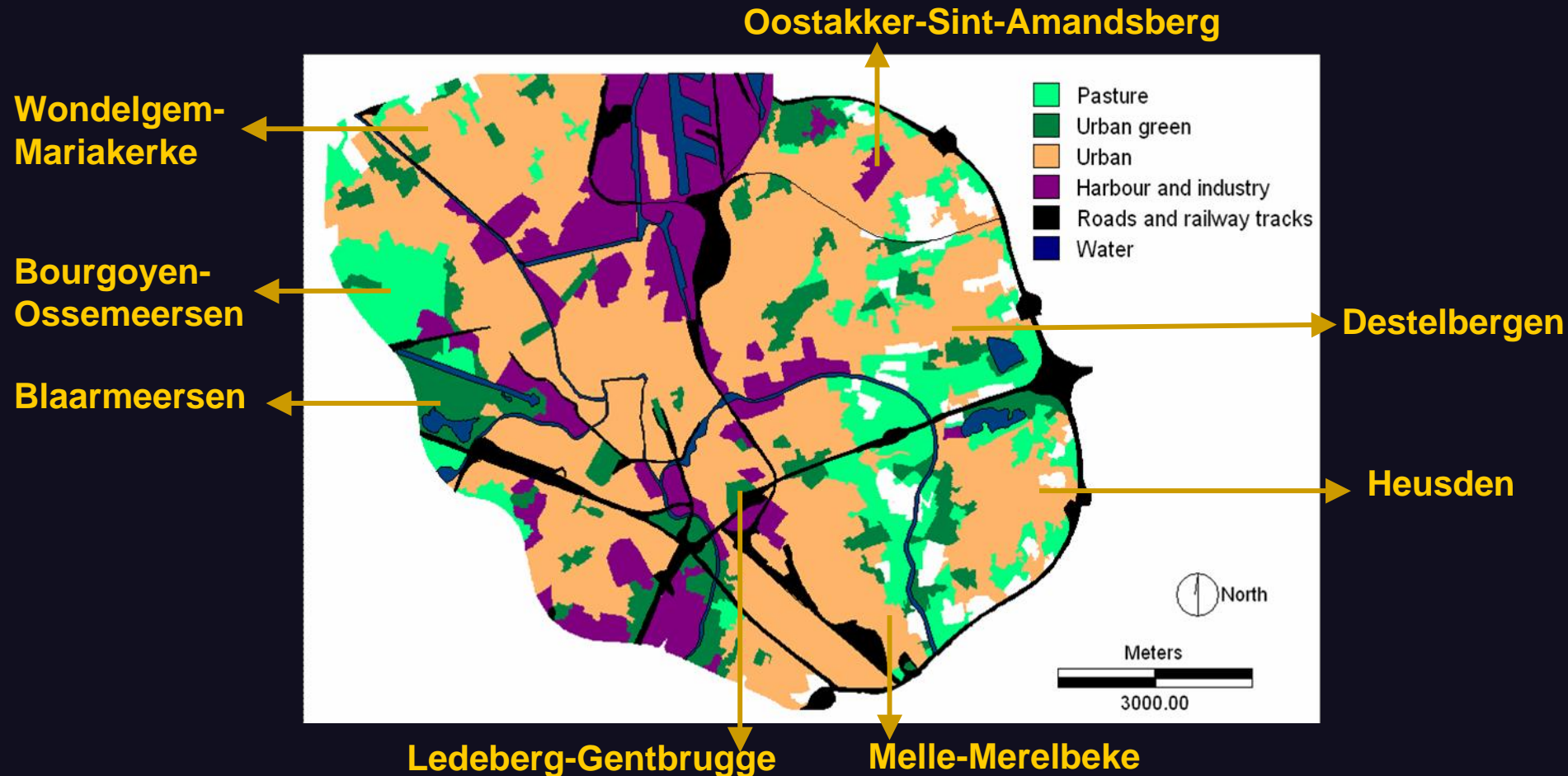
- Specific leaf area
- Stomatal density and
- Stomatal pore surface

as parameters in biomonitoring of urban habitat quality

Materials and Methods

Study area

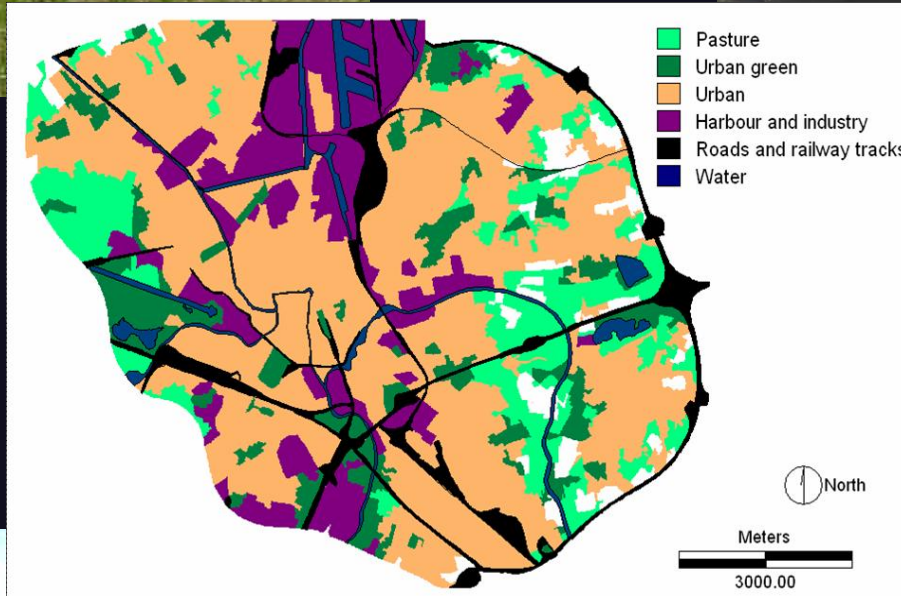
- City of Gent
- 81.5 km² area
- 4 land use classes



Bourgoyen-Ossemeersen



Gent-harbour



Blaarmeersen



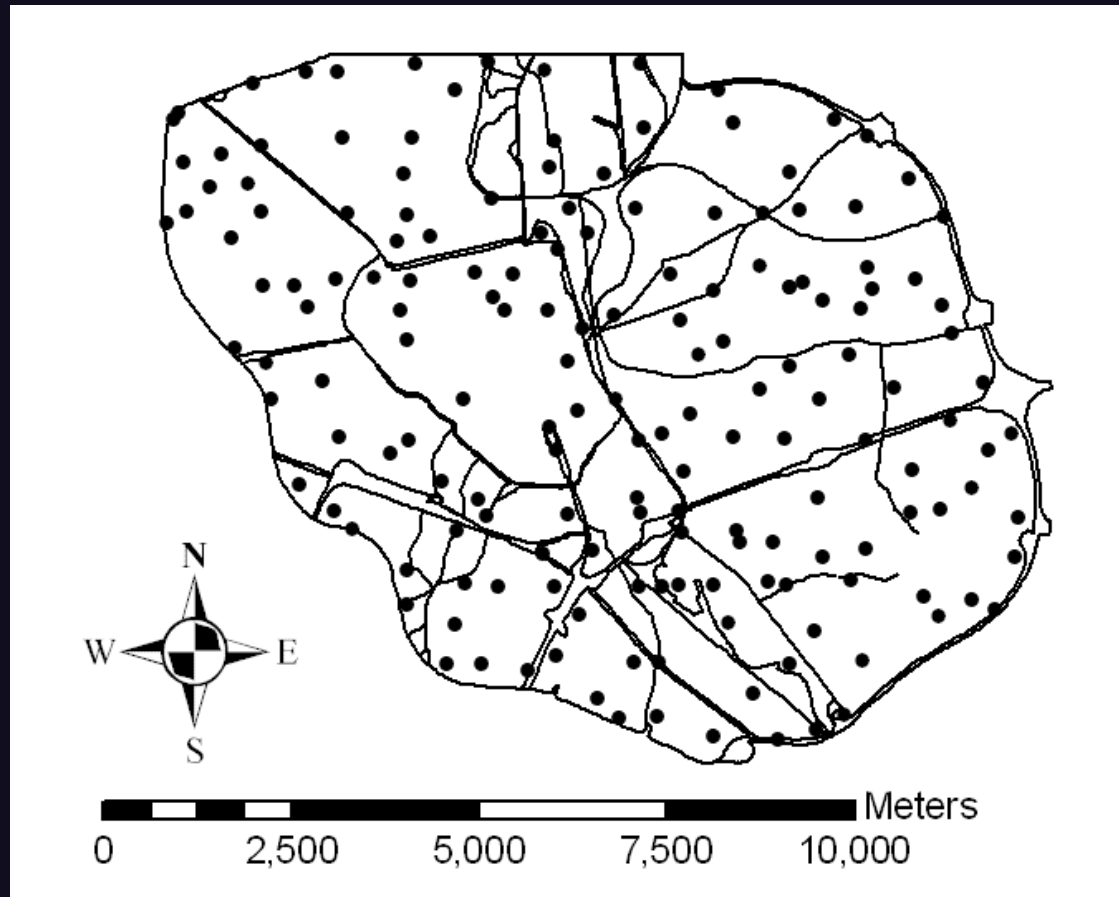
City Centre



Materials and Methods contd..

Sampling Survey:

- Partially random & grid sampling design
- 169 sampling locations
- During 2005-2007



Materials and Methods contd..

Selection of species

- Widely distributed
- Perennial
- Easy to sample
- Sensitive/ tolerant to pollutants (literature)



Materials and Methods contd..

At each sampling location

- 4 sub-sampling points
- Leaf samples
- Stomatal imprints

Determination of bio-chemical parameters (only for *T. officinalis*)

- Chlorophyll a and b
- C and N content
- $\delta^{13}\text{C}$ and $\delta^{15}\text{N}$ content

Determination of SLA (SLA, $\text{m}^2 \text{g}^{-1}$)

- Measurement of leaf area and dry weight
- $\text{SLA} = \text{LA}/\text{DW}$



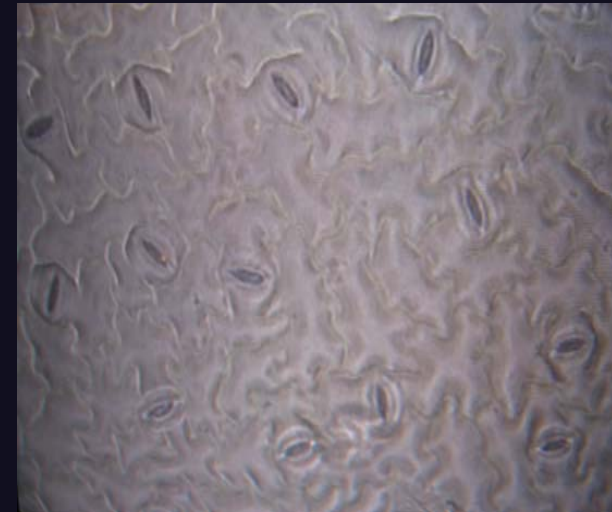
Materials and Methods contd..

Determination of stomatal characteristics

- 40 x 10 magnification under light microscope
- **Stomatal density (SD, mm⁻²)** = No. of stomata per leaf area
- Measurement of pore length and width and calculation of
 - Stomatal pore surface (SPS, μm²)**
 - Theoretical stomatal resistance (Rs, sm⁻¹)**

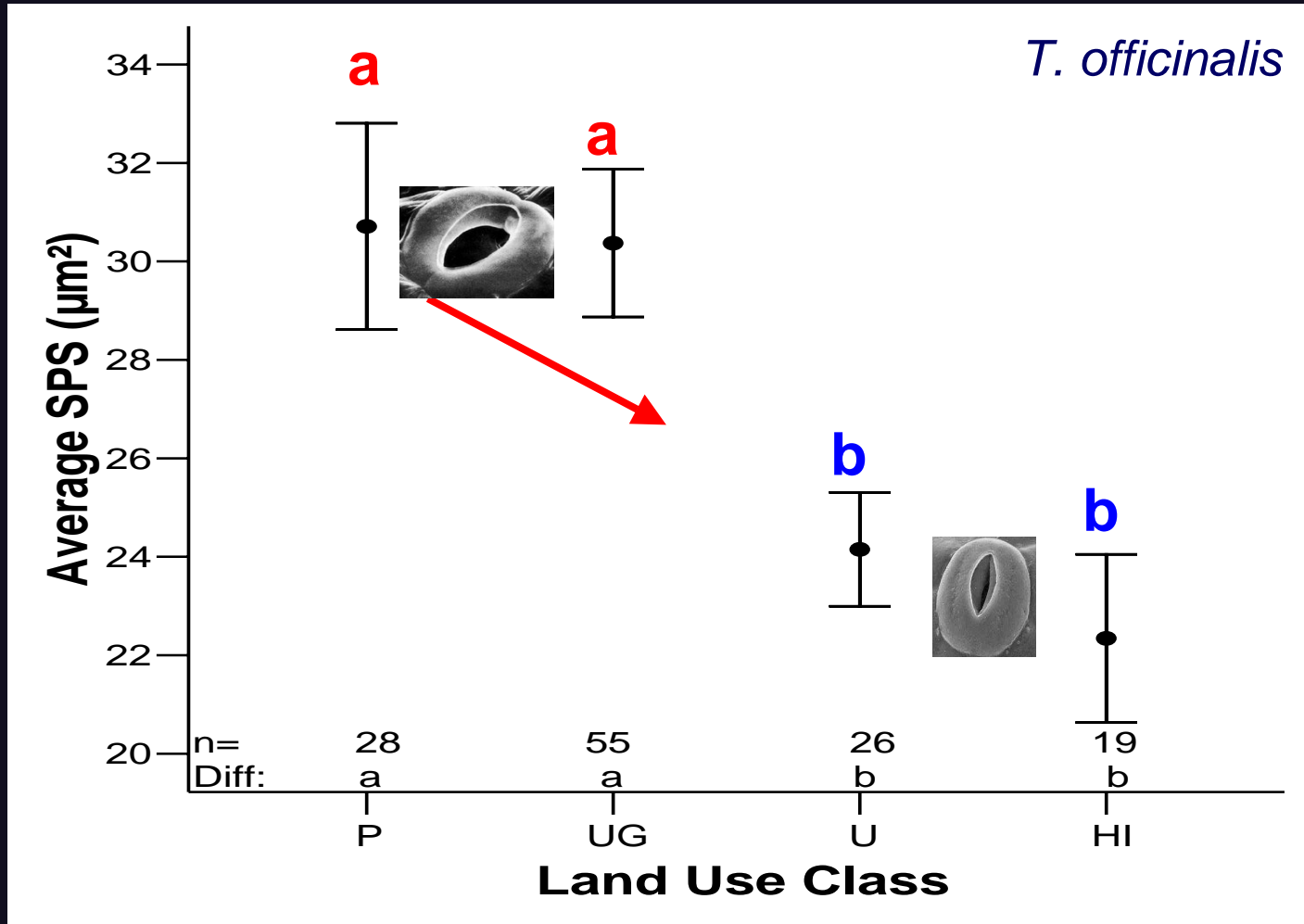
Data analysis

- ANOVA with Duncan test for differences between land use classes
- Geostatistical analysis - Ordinary Kriging and Simple Kriging with varying Local Means (SKLM).



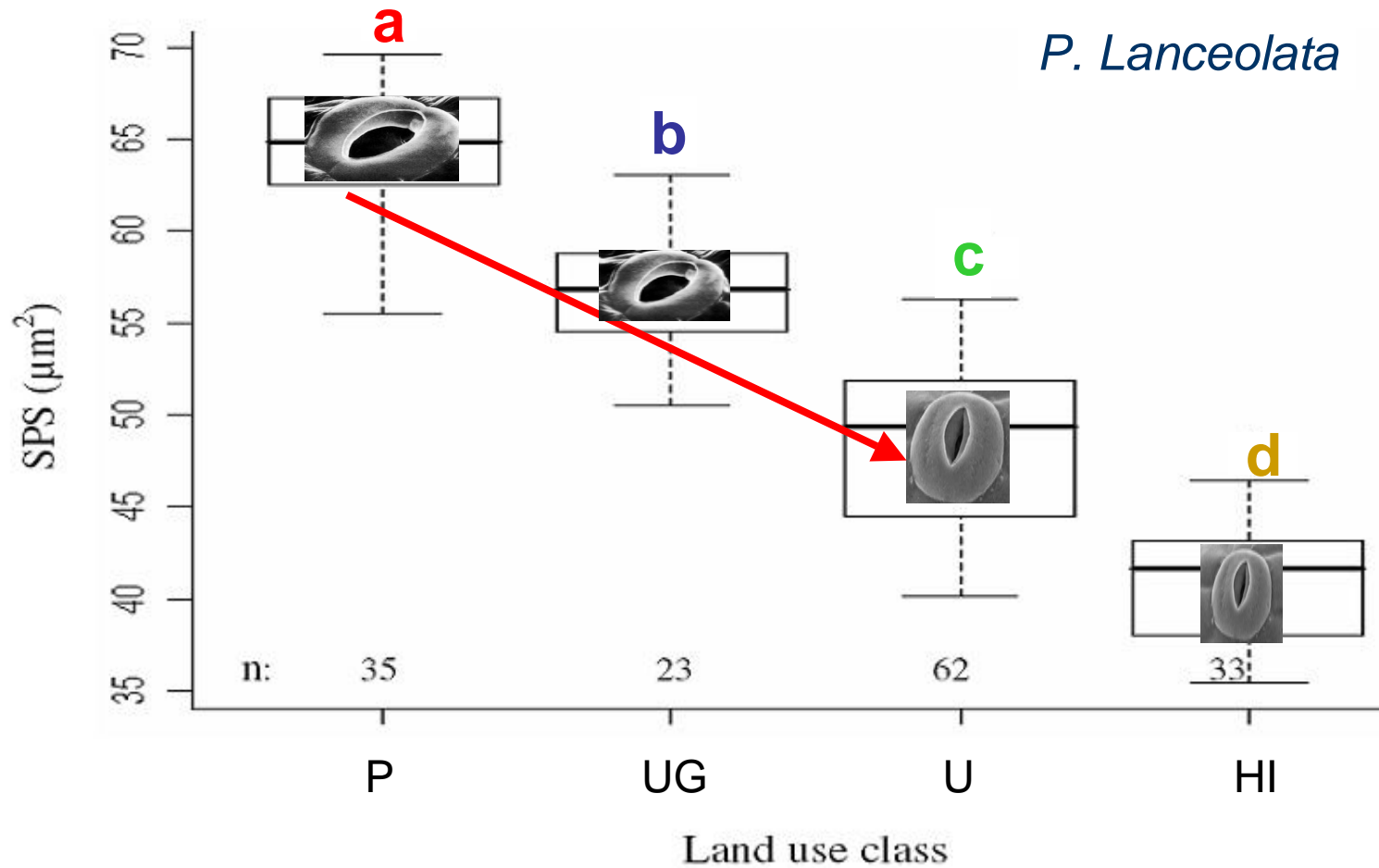
Results

SPS



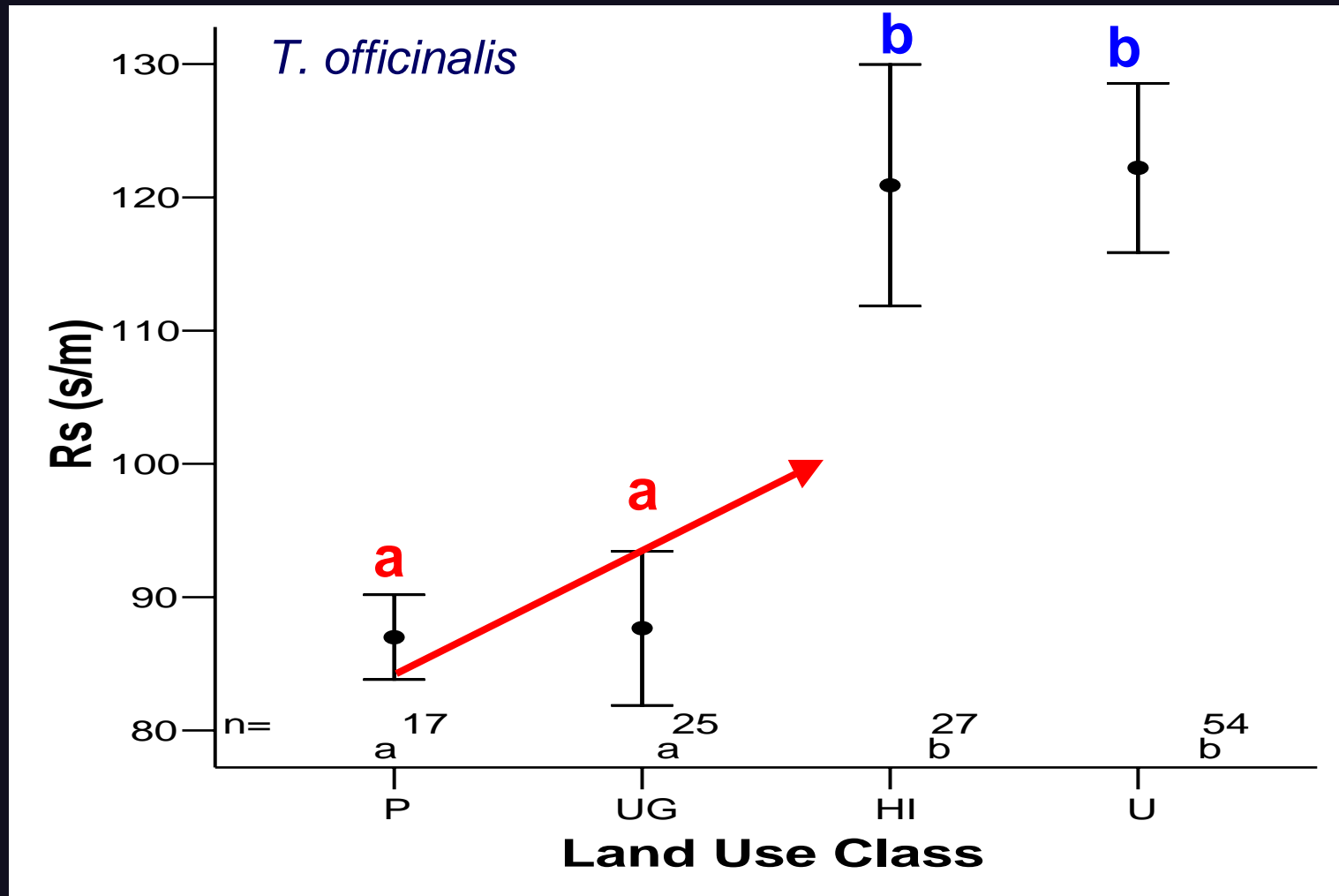
Results

SPS



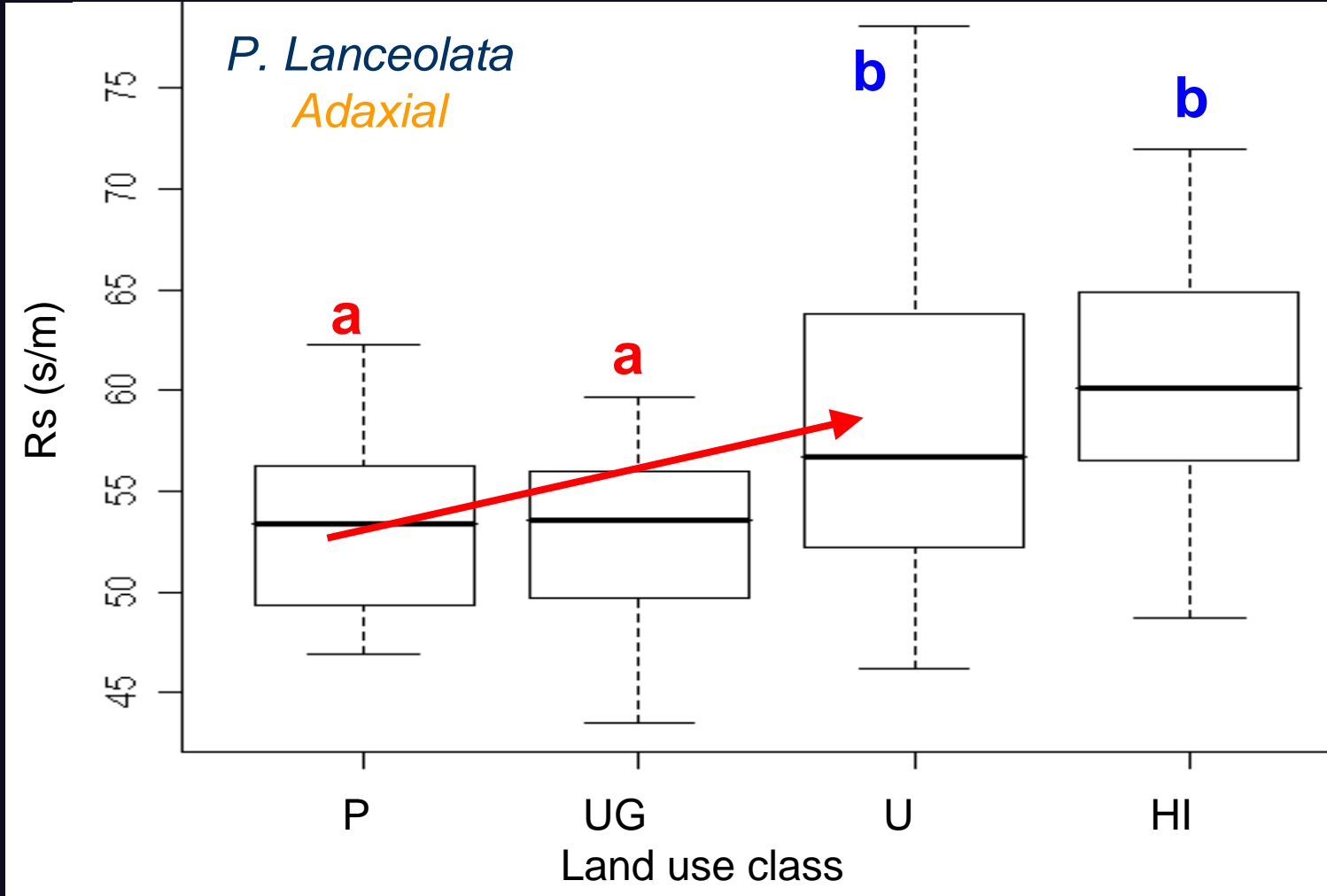
Results

R_s



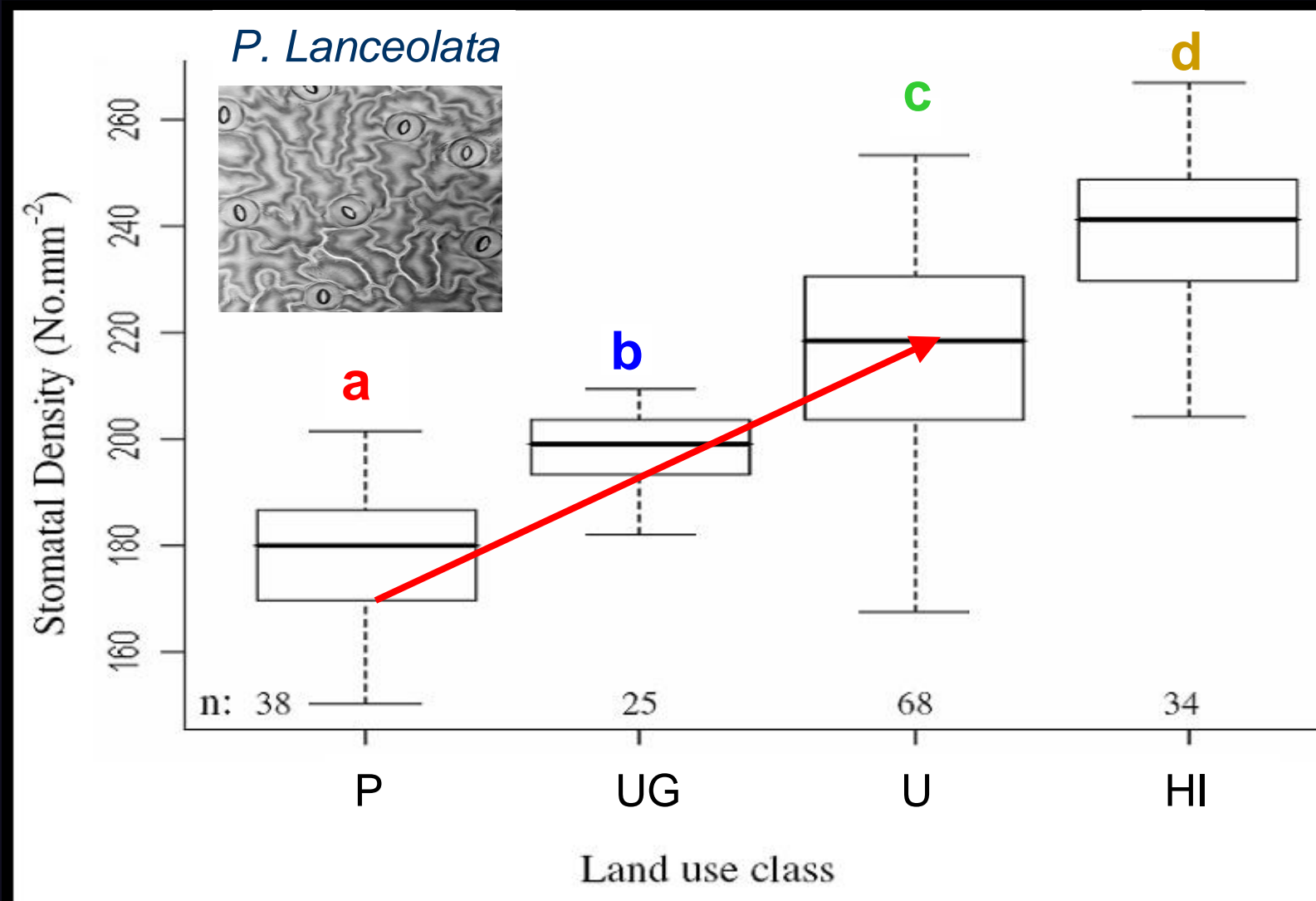
Results

Rs



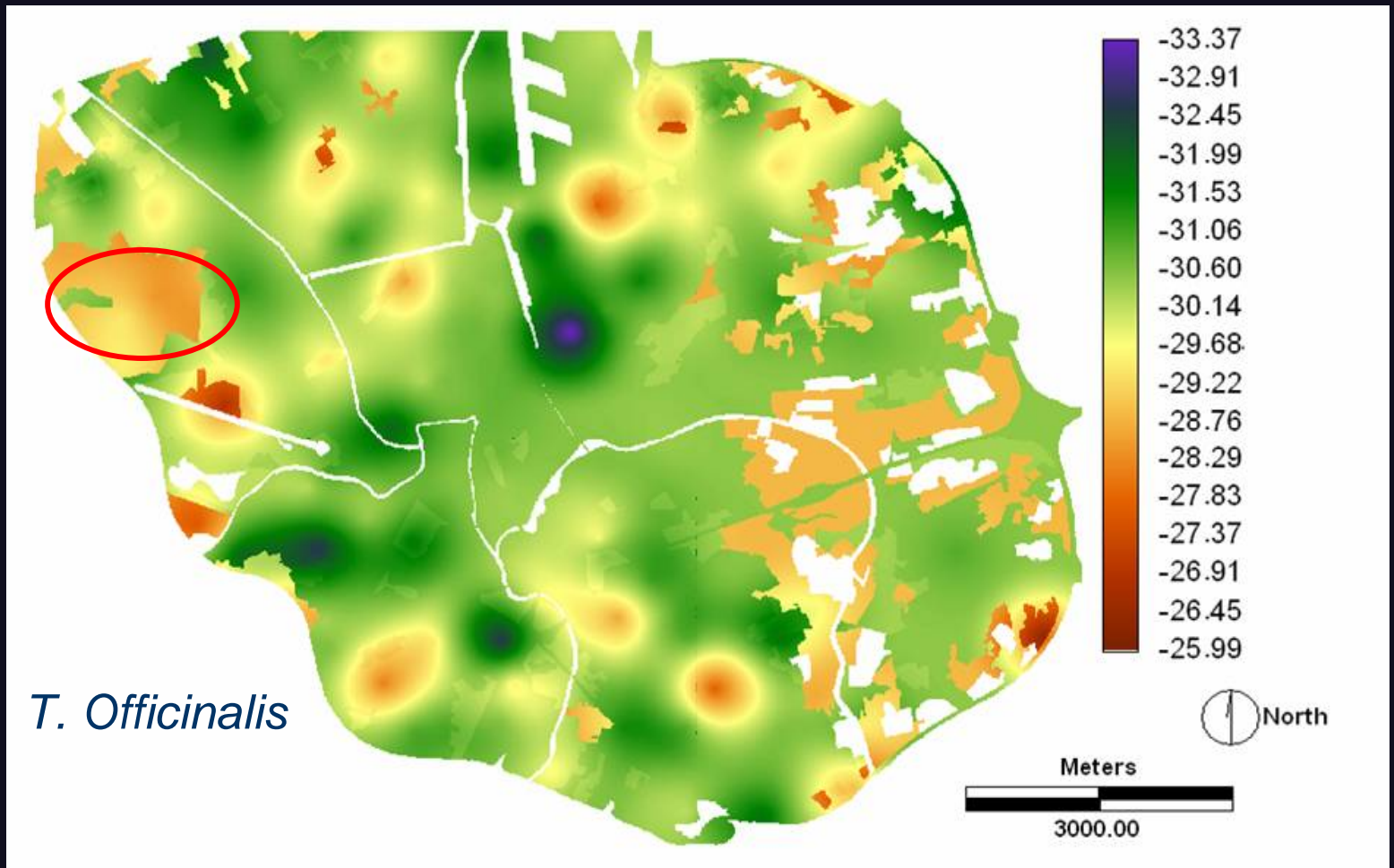
Results

SD

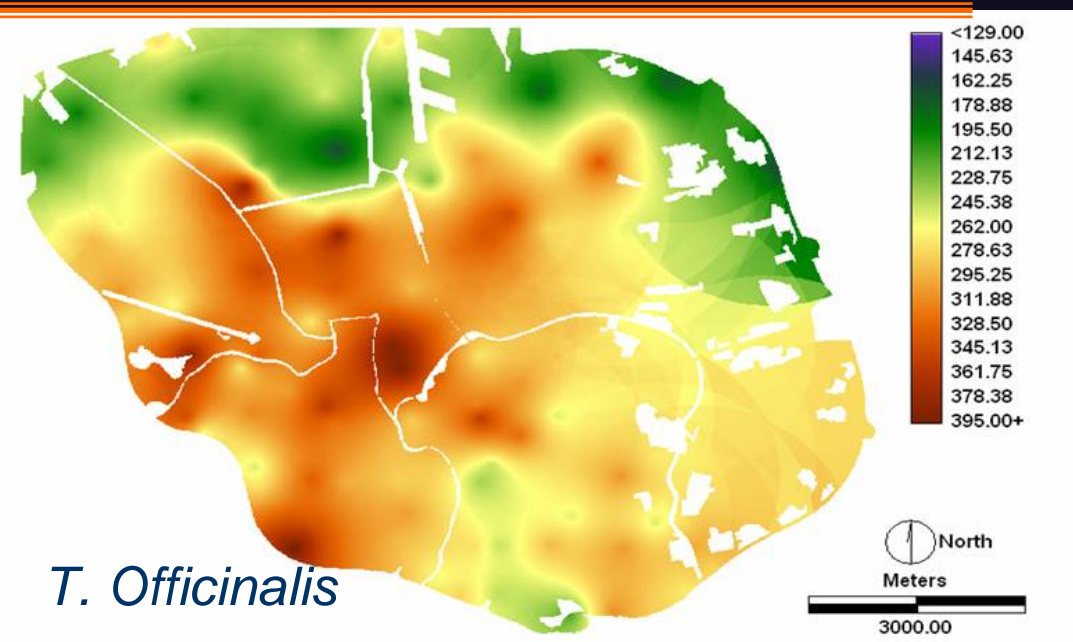


Results

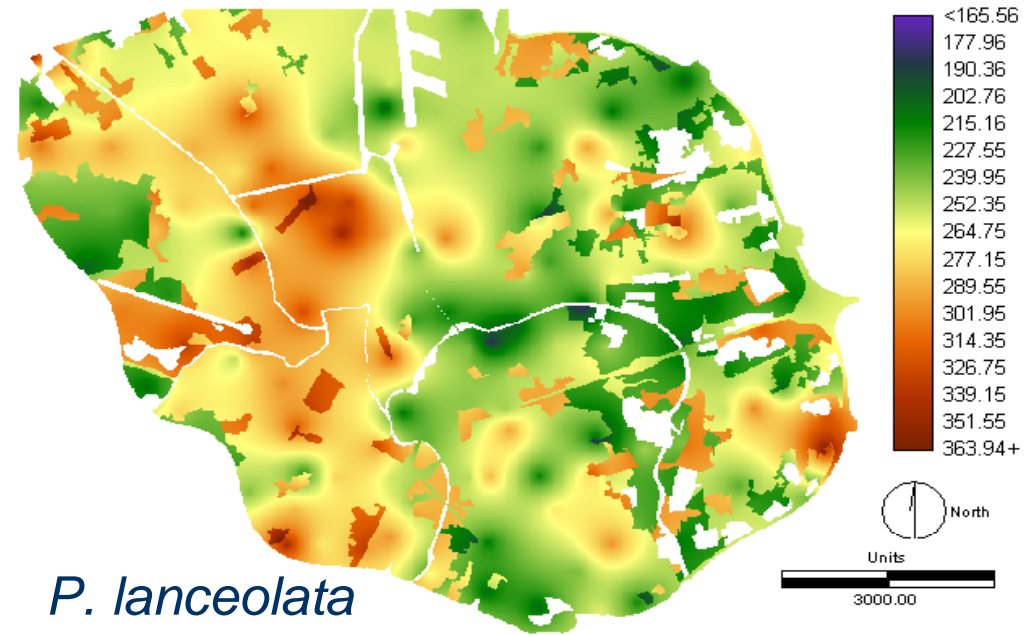
C13 ↓



Results

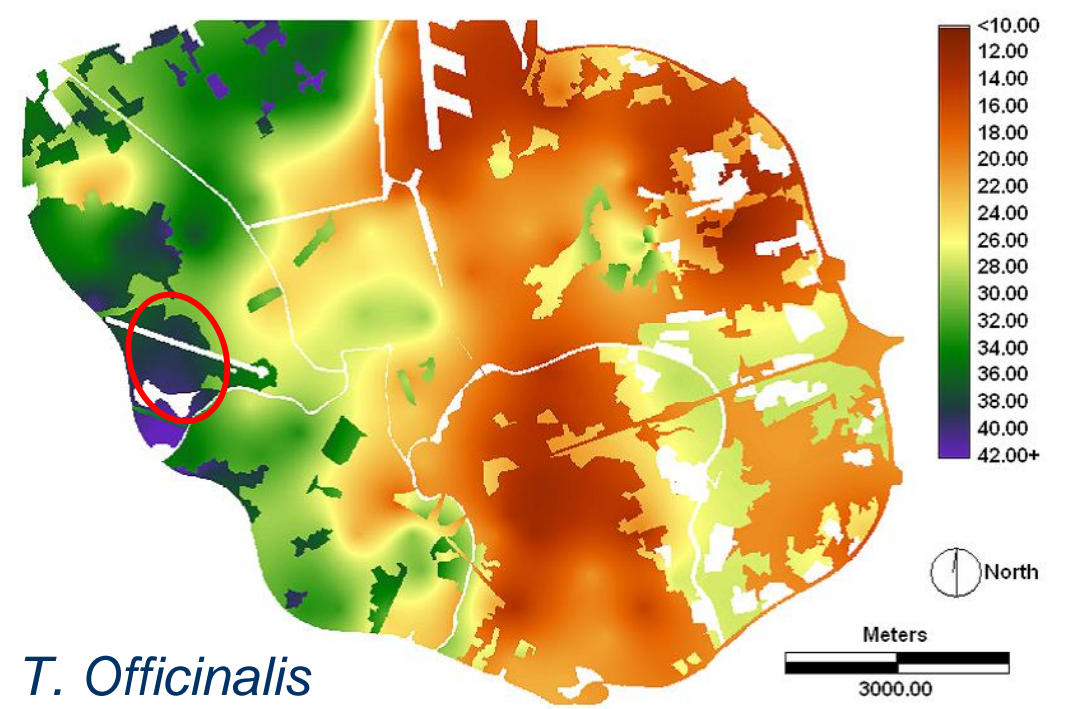
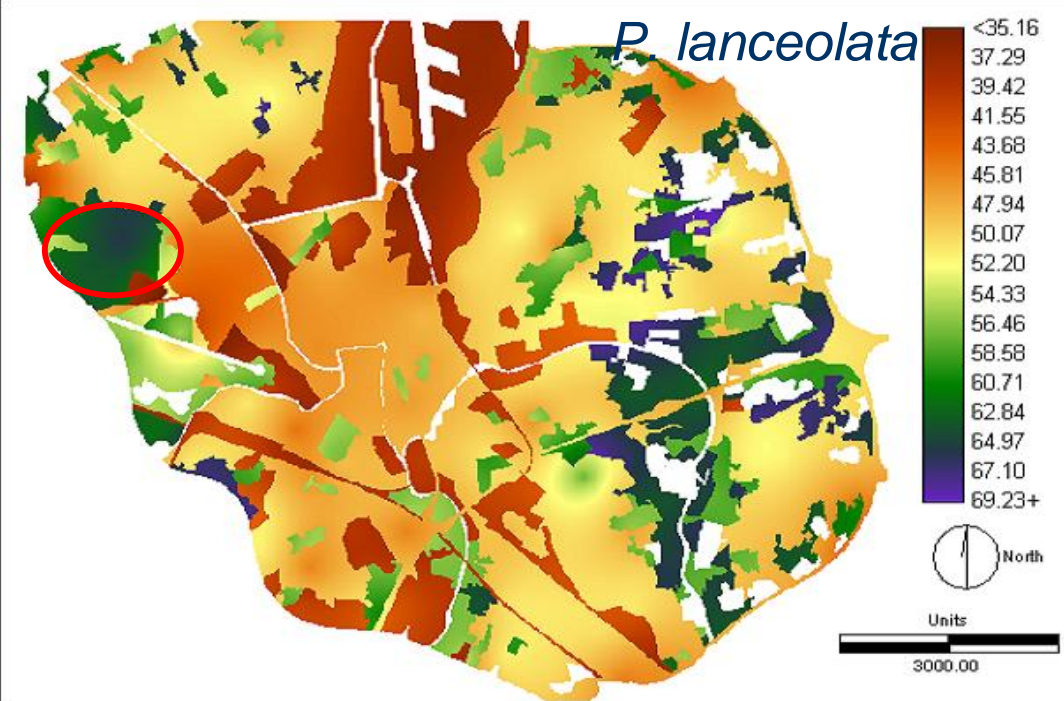


SLA

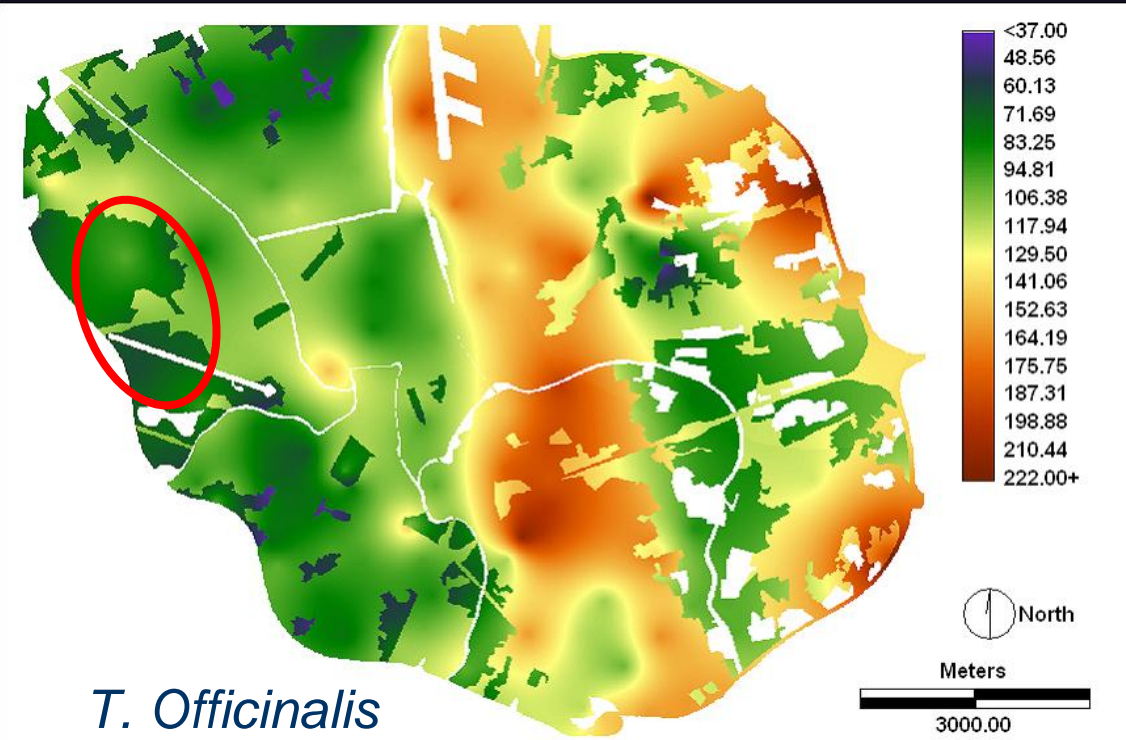
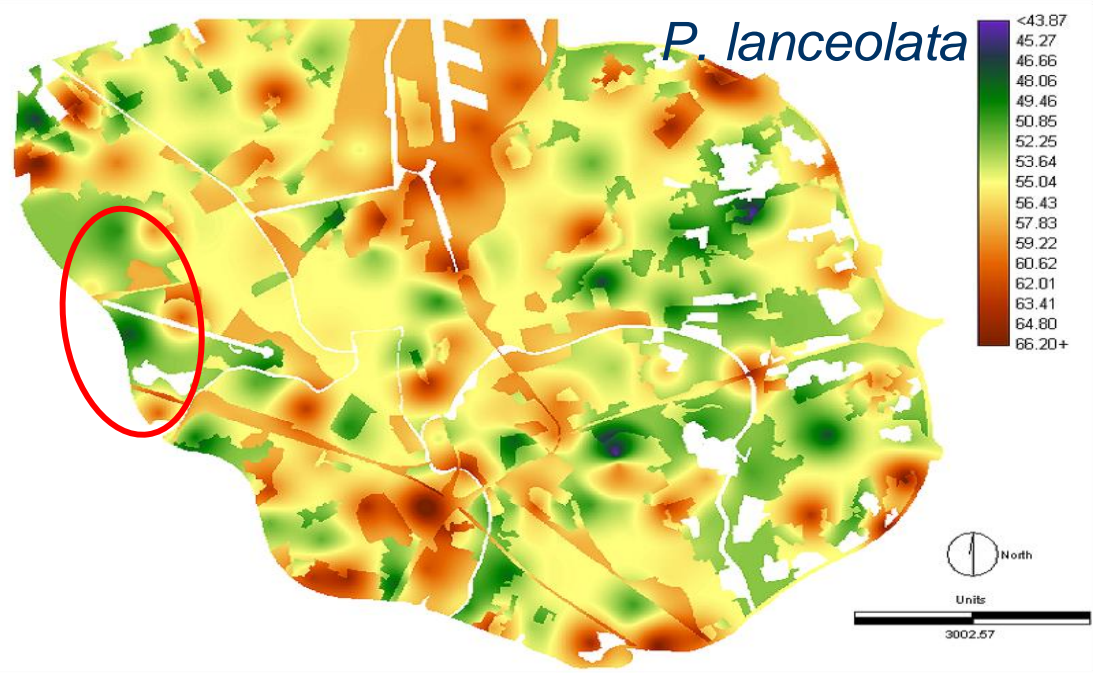


SPS ↓

Bourgoyen-Ossemeersen

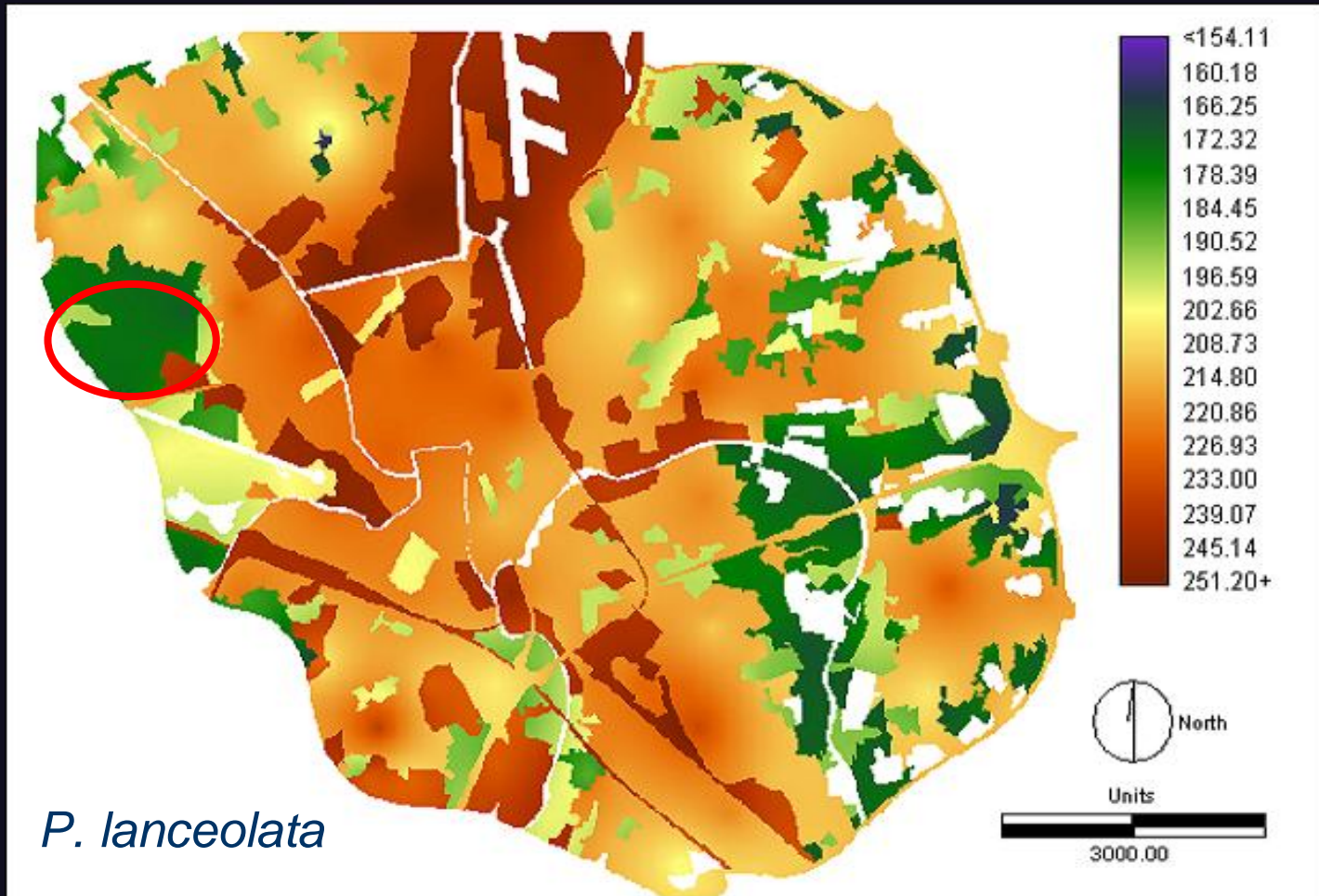


R_s ↑



Results

SD ↑



Conclusions

- SPS significantly decreased towards more polluted areas
- R_s was higher in more polluted areas
- SD of *P. lanceolata* increased significantly towards more polluted areas
- Stomatal characteristics of *T. officinalis* & *P. lanceolata* are suitable bioindicators of urban habitat quality

Suggestions for future research

- Test in the different climatic regions
- A comparison of the different methodologies
- Use of image analysis software to replace the laborious microscopic analysis

Thank You