

7. Pries AR, Secomb TW, Gaehtgens P. Biophysical aspects of blood flow in the microvasculature. *Cardiovasc Res* 1996; 32: 654-67.
8. Artmann GM, Kelemen C, Porst D, Büldt G, Chien S. *Biophys J.* 1998 Dec; 75(6):3179-83.
9. Singel DJ, Stamler JS. Chemical physiology of blood flow regulation by red blood cells: the role of nitric oxide and S-nitrosohemoglobin. *Annu Rev Physiol* 2005; 67: 99-145.
10. Jensen FB. The dual roles of red blood cells in tissue oxygen delivery: oxygen carriers and regulators of local blood flow. *J Exp Biol* 2009; 212: 3387-93.
11. Bergfeld GR, Forrester T. Release of ATP from human erythrocytes in response to a brief period of hypoxia and hypercapnia. *Cardiovasc Res* 1992; 26: 40-7.
12. Jia L, Bonaventura C, Bonaventura J, Stamler JS. S-nitroso hemoglobin: a dynamic activity of blood involved in vascular control. *Nature* 1996; 380: 221-6.
13. Cosby K, Partovi KS, Crawford JH, et al. Nitrite reduction to nitric oxide by deoxyhemoglobin vasodilates the human circulation. *Nat Med* 2003; 9:1498-505.
14. Maher AR, Milsom AB, Gunaruwan P, et al. Hypoxic modulation of exogenous nitrite-induced vasodilation in humans. *Circulation* 2008; 117: 670-7.
15. Dern RJ, Brewer GJ, Wiorkowski JJ. Studies on the preservation of human blood. II. The relationship of erythrocyte adenosine triphosphate levels and other in vitro measures to red cell storageability. *J Lab Clin Med* 1967; 69: 968-78.
16. Low FM, Hampton MB, Winterbourn CC. Antioxid Redox Signal. 2008 Sep;10(9):1621-30. PMID: 18479207.
17. Hanson MS, Ellsworth ML, Achilleus D, et al. Insulin inhibits low oxygen-induced ATP release from human erythrocytes: implication for vascular control. *Microcirculation* 2009; 16: 424-33.
18. d'Almeida MS, Jagger J, Duggan M, et al. A comparison of biochemical and functional alterations of rat and human erythrocytes stored in CPDA-1 for 29 days: implications for animal models of transfusion.
19. Matthes G, Strunk S, Siems W, Grune T. Posttransfusional changes of 2,3-diphosphoglycerate and nucleotides in CPD-SAGM-preserved erythrocytes. *Infusionsther Transfusionsmed* 1993; 20: 89-92.
20. K.M. Faulkner, A.L. Crumbliss, C. Bonaventura, A spectroelectrochemical method for differentiation of steric and electronic effects in hemoglobins and myoglobins, *J.Biol. Chem.* 270 (1995) 13604–13612.
21. A.Arnone, X-ray diffraction study of binding of 2,3- diphosphoglycerate to human deoxyhaemoglobin, *Nature* 237 (1972) 146–149.
22. J. Kister, C.Poyart, S.J.Edelstein, An expanded twostate allosteric model for interactions of human hemoglobin A with nonsaturating concentrations of 2,3-diphosphoglycerate, *J.Biol. Chem.*262 (1987) 12085–12091.
23. C.H. Taboy, K.M. Faulkner, D. Kraiter, C. Bonaventura, A.L. Crumbliss, Concentration-dependent effects of anions on the anaerobic oxidation of hemoglobin and myoglobin, *J.Biol. Chem.* 275 (2000) 39048–39054.
24. C.F.S. Bonafe, A.Y. Matsukuma, M.S.A. Matsuura, ATP-induced tetramerization and cooperativity in hemoglobin of lower vertebrates, *J.Biol. Chem.* 274 (1999) 1196–1198.
25. R.Benesch, R.E.Benesch, Intracellular organic phosphates as regulators of oxygen release by hemoglobin, *Nature* 221 (1969) 618–622.
26. M.F. Perutz, Stereochemistry of cooperative effects in haemoglobin, *Nature* 228 (1970) 726–734.
27. Rao S, Rossmann M (1973). Comparison of super-secondary structures in proteins. *J Mol Biol* 76 (2): 241–56.
28. Scheeff E, Bourne P (2005). Structural evolution of the protein kinase-like superfamily. *PLoS Comput Biol* 1 (5): e49.
29. K. Zerlin, I. Digel, A. Stadler, G. Büldt, G. Zaccai, G.M. Artmann. Dynamics and interactions of hemoglobin in human red blood cells and concentrated hemoglobin solutions. *Regenerative Medicine* September 2007. - Vol 2. - No 5. - pp. 573.

**Preiß C<sup>1</sup>., LinderP<sup>1</sup>., Wendt K<sup>1</sup>., Krystek M<sup>1</sup>., Digel I<sup>2</sup>., Gossmann M<sup>3</sup>., Artmann<sup>3</sup>A.T.,  
Porst D<sup>3</sup>., Kayser P<sup>3</sup>., Bassam R<sup>2</sup>., Artmann G.M<sup>1</sup>.**

## **ENGINEERING TECHNOLOGY FOR PLANT PHYSIOLOGY AND PLANT STRESS RESEARCH**

*Institute of Bioengineering (IFB), FH Aachen University of Applied Sciences, <sup>1</sup>Laboratory for  
Cellular Biophysics, <sup>2</sup>Laboratory of Cell- and Microbiology, and <sup>3</sup>Laboratory of Medical &  
Molecular Biology, Juelich, Germany, Linder@fh-aachen.de*

Plant physiology and plant stress: Plant physiology will be much more important for human mankind because of yield and cultivation limits of crops determined by their resistance to stress. To assess and counteract various stress factors it is necessary to conduct plant research to gain information and results on plant physiology. Especially for agriculture this is of great significance, because stress is very harmful to plants resulting in reduction of biomass production of crops. Stress

is defined as a tension state, describing the effect of a load of the organism caused by external factors which impair the metabolism or growth. Not only humans but also plants and animals are exposed to a so-called stress. There are many different types of plant stress. Major abiotic stress factors in plants are:

- mechanical stress
- water stress
- aridity stress
- salt stress
- heat stress
- frost stress
- oxygen stress
- light stress
- UV radiation stress

**C3-Plants:** The C3-photosynthesis is the best known and most common type of photosynthesis in nature. It is the most effective process for the synthesis of biomass. In hot climates, the stomata of plants close to keep the water-loss due to transpiration as low as possible. It follows a reduction in CO<sub>2</sub>-uptake necessary for photosynthesis. Examples of C3-plants are: Wheat, rice, sugar-beet and potatoes.

**C4-Plants:** During evolution these plants, developed from the C3-plant-type, pursue a special kind of photosynthesis. These botanical species are able to pursue photosynthesis with low CO<sub>2</sub>-concentrations and they also lose very little water under strong solar irradiation. Thus these plants are adapted well to hot climates. Examples of C4-plants are: Millet, maize, sugarcane, china-reed and tumbleweed.

**Aim of the study:** Climate change and growing population lead to food shortages in the near future. A major objective in plant research is to combine the characteristics of C4 plants to the species of C3 plants in order to protect them against climate change and the increased use of agricultural areas for C3 plants in hot areas of the world too. Therefore it is important to exactly quantify the biomass production of individual plants at regular conditions as well as at defined plant stress. The determination process has to be quick and accurate. Therefore we combined our skills in technical engineering with knowledge in plant biology to built an automated system for analyzing the shape and cross section of individual plant leaves at high accuracy.

**Experiments:** To reinforce the idea of a mobile plant-scanner, there were two experiments conducted with a stationary test-stand-construction. A plant leaf of a potted plant was fixed between two laser sensors, one coming from below and one from above the leaf (Figure 1).



Figure 1: fixed plant leaf in an experimental setup

**1. aridity stress:** Cross-sectional area changes during slow drying of the plant and subsequent one-time irrigation were detected (Figure 2). The graphic shows a fast decrease of the normalized cross-section of the leaf to 20 % at day three. After the irrigation the cross-section recovers and increases. However, never completely but up to only about 75% of the value before irrigation.

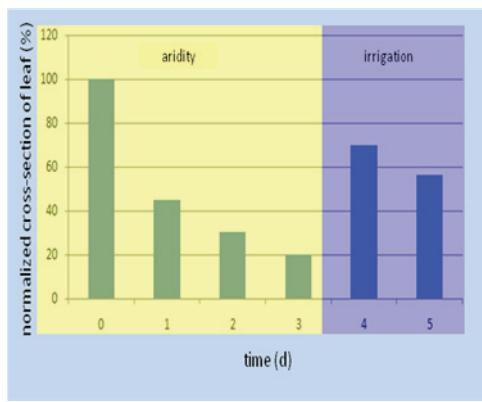


Figure 2: aridity stress

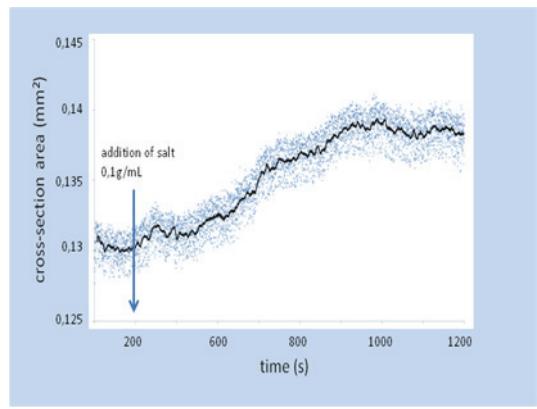


Figure 3: salt stress

2. *salt stress*: It was also possible to detect very fast cross-sectional area changes of the leaf caused by salt stress. Similarly to the first experiment, a plant leaf of a second potted plant was fixed to the test-stand. After 200 seconds a salt solution (0.1g/ml) was added to the pot (Figure 3). Contrary to the expected decrease of the cross-section the volume of the leaf increased. In literature this is described as direct admission of nutrients of the cells.

PhytoScan Alpha 70: After successful preliminary experiments the plant-scanner PhytoScan Alpha 70 (Phytos: greek plant, Alpha: first Version, 70 mm scan width,

Figure 6) was developed. It is a mobile device to be used in greenhouses and plant research institutes to analyze the shape and cross-section in real time (Figure 4 and Figure 5). It is possible to create a 3D-model of the plant leaves.



Figure 4: analyzed plant leaf

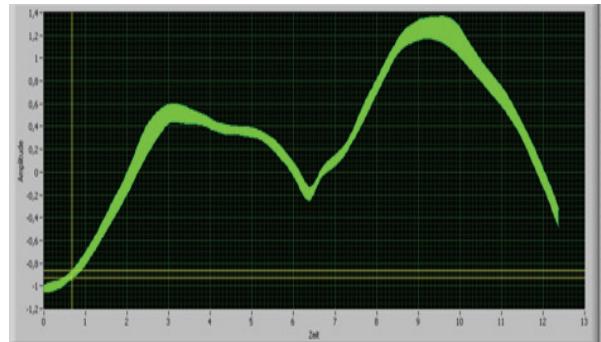


Figure 5: computer computation of a cross-section of a leaf



Figure 6: PhytoScan Alpha 70