

COMPENSATION OF OXYGEN PARTIAL PRESSURE VARIATIONS DURING OPTICAL ENZYMATIC GLUCOSE SENSING IN ADIPOSE TISSUE

M. Rumpler¹, M. Hajnsek¹, I. Klimant², F. Sinner^{1,3}, T.R. Pieber^{1,3}

CONTACT

¹
JOANNEUM RESEARCH
Forschungsgesellschaft mbH

HEALTH
Institute for
Biomedicine and
Health Sciences

Martin Hajnsek

Neue Stiftingtalstrasse 2
8010 Graz

Phone +43 316 876-4000
Fax +43 316 8769-4000

martin.hajnsek@joanneum.at

health@joanneum.at
www.joanneum.at/health



Graz University of Technology

²

Graz University of Technology

Institute of Analytical Chemistry
and Food Chemistry,
Graz, Austria



³

Medical University of Graz

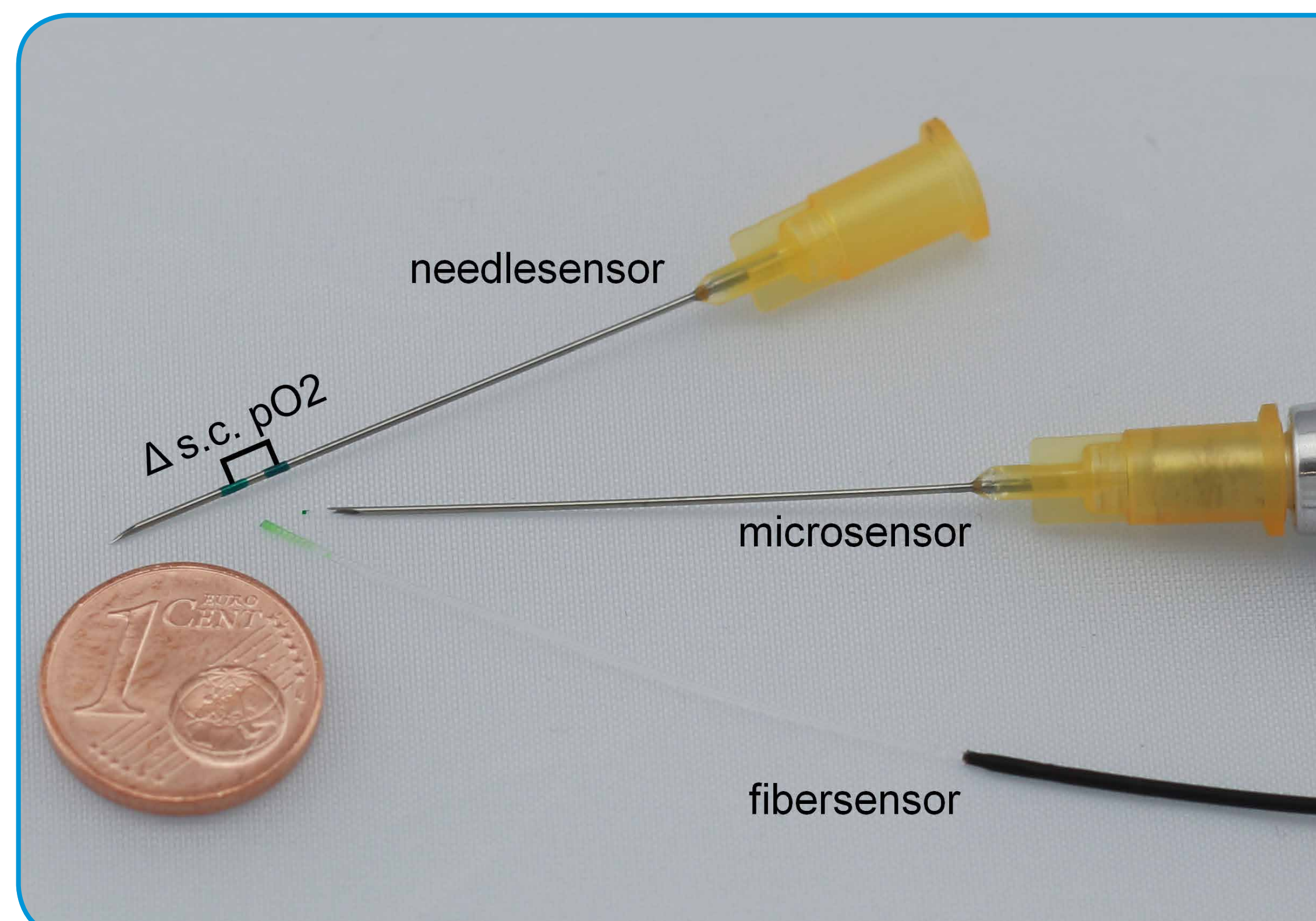
Department of Internal Medicine
division of endocrinology and
metabolism

Graz, Austria

This single-port concept combines an optical enzymatic glucose measurement and an insulin delivery system placed at the same position in subcutaneous adipose tissue (AT). Changes in oxygen partial pressure (pO₂) in the AT influence the glucose

Methods

During in-vivo experiment in domestic pigs, the blood pO₂ level During in-vivo experiments in domestic pigs, the blood pO₂ level was modified to different steady state plateaus. During each plateau phase, three different sensor systems were used to measure tissue pO₂. Fiber-sensors and micro-sensors continuously recorded tissue pO₂ profiles over a distance of 21 mm. Six stationary needle-sensors, each with two 2 mm long sensor elements, were randomly placed across the abdominal area and recorded simultaneously the tissue pO₂ for 30 min per steady state plateau. Fiber- and micro-sensors were read-out with light-guiding and needle-sensors were read-out transcutaneously by a phase fluorimeter.



Conclusion

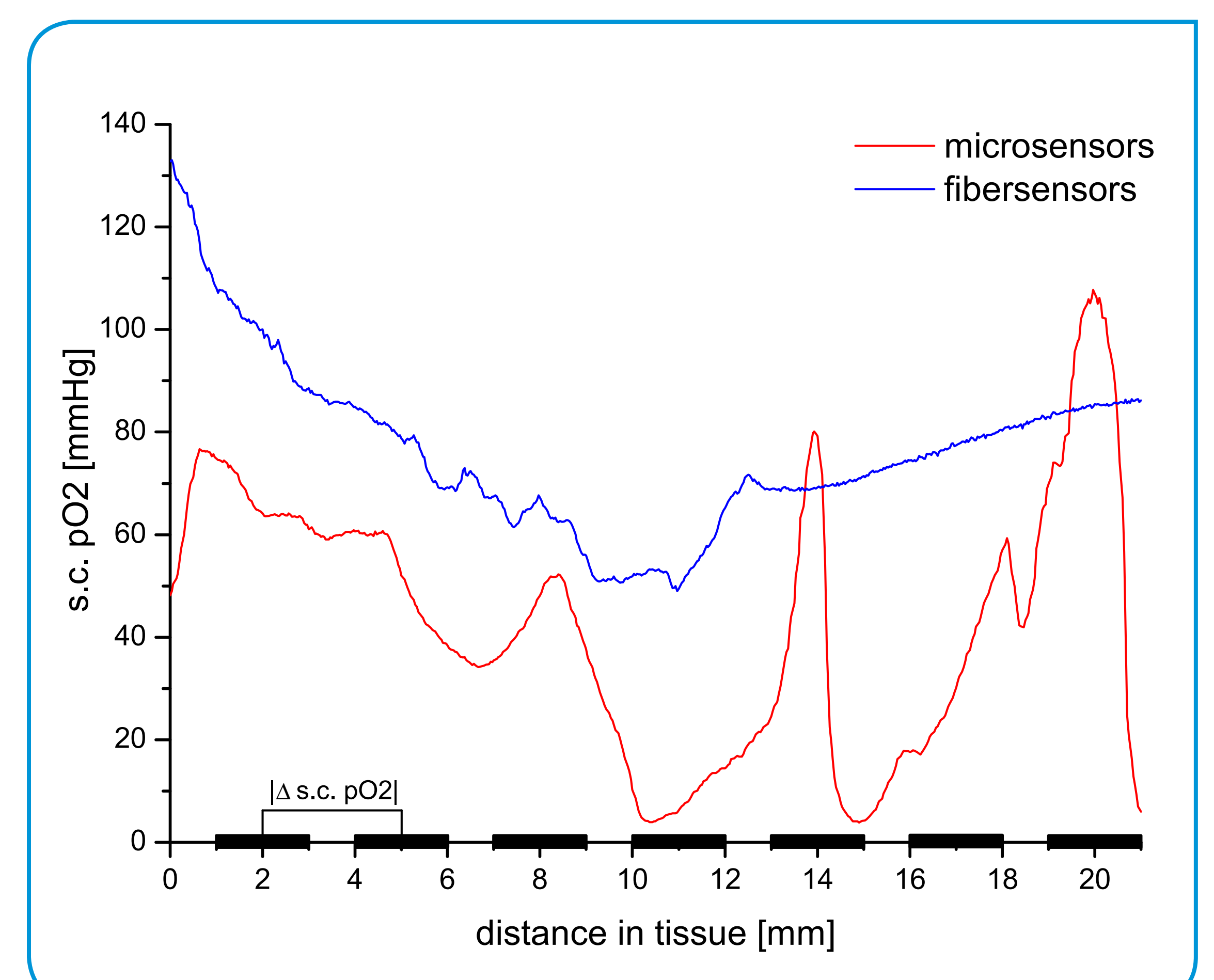
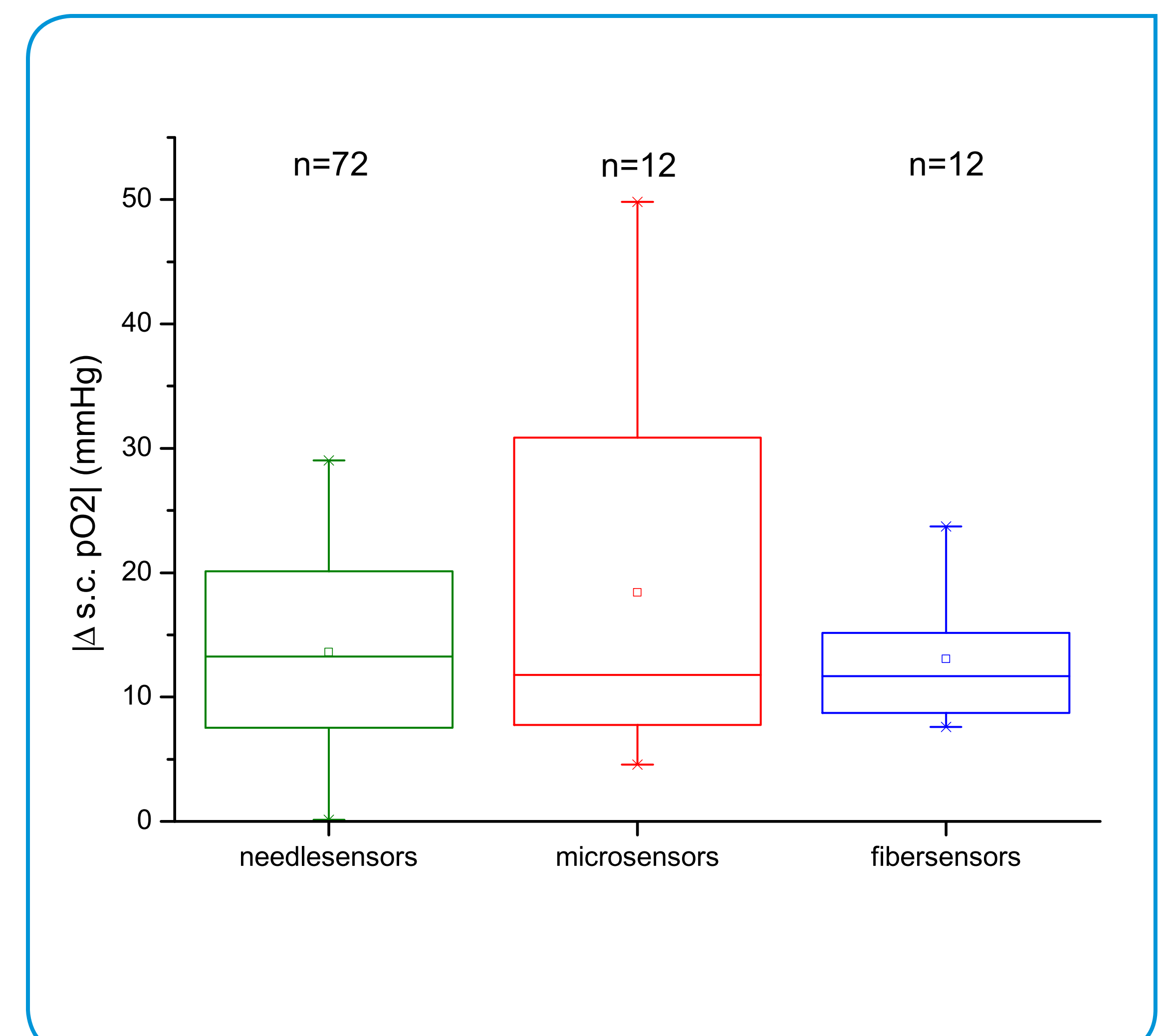
- We were able to show that tissue pO₂ varies considerably along the distance of 21 mm (micro-sensors).
- An increased sensing area (such as fiber- and needle-sensors) can reduce these heterogeneities, so that
- a mathematical correction of the glucose measurement should be possible.

Background

measurement and thus require direct pO₂ measurement in AT to mathematically correct the measured glucose values. The aim of this work was to assess the homogeneity of pO₂ in AT and a possible mathematical correction of tissue pO₂.

Results

All measured tissue pO₂ values were similar to values found in literature. Medians of the absolute differences of the pO₂ values (Δ s.c. pO₂) had a maximum intersystem deviation of 1.6 mmHg. The micro-sensors showed the largest interquartile range of all used sensor systems.



Acknowledgement

This project is funded by the
EU Framework 7 Programme,
contract no 305343.

