### SAREMCO DENTAL



Zurich Universit / Applied Science



# In vitro characterization of a new composite material for biomedical applications and 3D (bio)printing

Epifania Bono<sup>1</sup>, Christoph Evers<sup>2</sup>, Franca Schmid<sup>2</sup>, Ursula Graf-Hausner<sup>3</sup>, Markus Rimann<sup>1</sup>

<sup>1</sup>Zurich University of Applied Sciences ZHAW, Institute of Chemistry and Biotechnology ICBT, Einsiedlerstrasse 31, 8820 Waedenswil, Switzerland <sup>2</sup>Saremco Dental AG, Gewerbestrasse 4, 9445 Rebstein, Switzerland <sup>3</sup>graf3dcellculture, Bühlackerweg 5, 8405 Winterthur, Switzerland

## **Study goal**

**Project data** 

The present project aims at evaluating the cytocompatibility and printability of a new composite material, based on a mixture of a new methacrylate-based monomer developed within a CTI project (18514.1 PFLS-LS) and glass-ceramic powder supplemented with co- and photo-initiators (patent in preparation). This study is the basis to demonstrate the suitability of the biomaterial, for biomedical applications, such as stent, orthopedic implants and hearing aid components, as well as for 3D (bio)printing

## **Key findings**

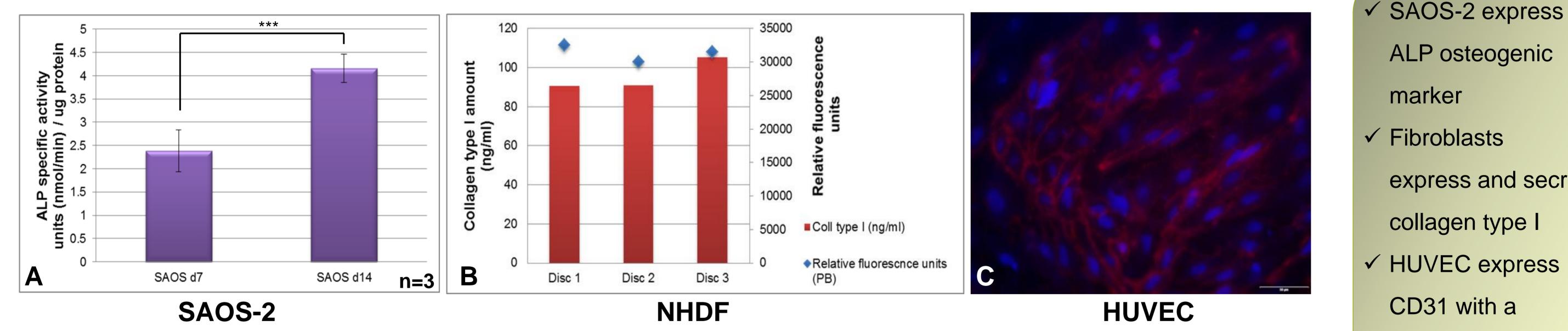
- > Cultivation, proliferation and differentiation of three different human cell types were successfully established on composite material discs (1cm diameter, 1mm height). Biological activity was shown
- $\succ$  The material is suitable for 3D (bio)printing, printing protocols were established
- > The new composite material is suitable for cell and tissue interaction in **biomedical applications**

#### **Cells adhesion and proliferation** SAOS-2 HUVEC NHDF $\checkmark$ All cell types 35000units Bay1 30000-adhere, spread Day3 25000-Day7 and proliferate on 20000-15000 the biomaterial Day 1 Day 1 10000over time Relative 5000-✓ Cells are able to SAOSil HUNE form a monolayer at day 7 on top of Day 7 Day 7 Day 7 **n=3** each discs

Figure 1: SAOS-2 (osteoblasts), NHDF (fibroblasts) and HUVEC (endothelial cells) seeded onto discs and cultivated up to 7 days before staining for F-actin (green signal, fluorescent phallacidin) and nuclei (blue signal, DAPI). Scale bar, 100µm

*Figure 2*: Metabolic cell activity detected with PrestoBlue at day1, 3 and 7. Unpaired T-test, p-values: \*\* p<0.01; \*\*\* p<0.001; \*\*\*\* *p*<0.0001

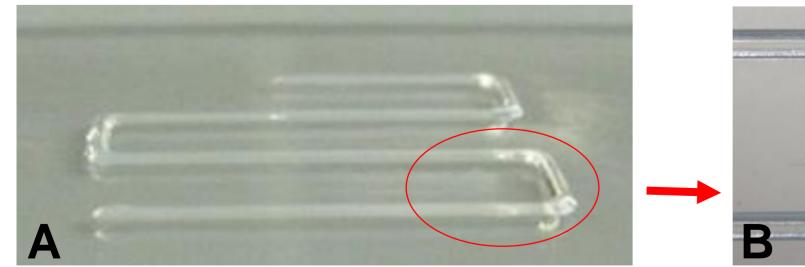
### **Biological cell activity**



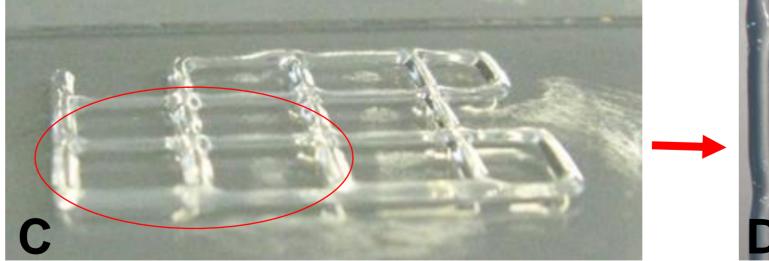
#### Figure 3:

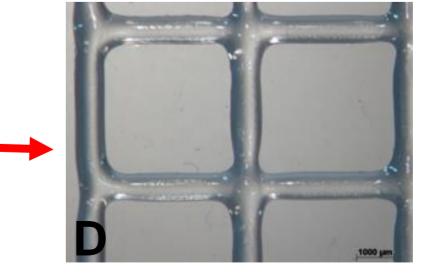
- A, ALP activity normalized to total protein amount on SAOS-2 culture. Unpaired T-test, \*\*\* p<0.001
- **B**, Coll I expression in fibroblasts cultures vs metabolic activity
- C, CD31 immunocytochemistry analysis in HUVEC cells cultivated up to 7 days (red signal). Nuclei in blue (DAPI). Scale bar, 50µm

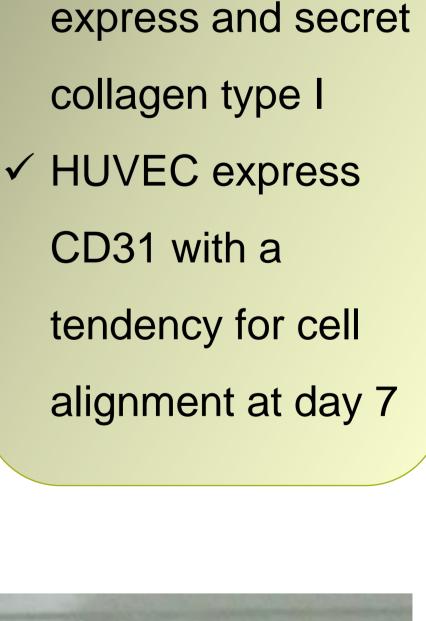
### **Biomaterial printability**















✓ The composite biomaterial is printable in a precise manner

✓ Well defined structures/models with high resolution can be obtained

✓ 3D printed models tested are stable showing rigid shape

Figure 4: 3D structures printed with 3DDiscovery (regenHU) with direct dispensing in a layer-by-layer fashion. **A-D**, 3 layers models; *E*, 10 layers model. Scale bar, 1000µm

## Conclusions

The new composite material is not only cytocompatible (DIN EN Iso 10993-5), but also allows the biological activity of human cells (spreading, proliferation and differentiation). Moreover, preliminary studies demonstrate the suitability of the material for (bio)printing. The composite shows high potential as biomaterial for 3D printing in different biomedical applications

### Acknowledgments

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