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MODULATION OF EXTRA VIRGIN OLIVE OIL DIGESTIBILITY THROUGH OLEOGELATION

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Background



The development of a **SUSTAINABLE FOOD SYSTEM**, to favor the **TRANSITION TO HEALTHIER AND MORE SUSTAINABLE DIETS**, is one of the major challenges of the modern food industry [1].



The use of extra virgin olive oil (**EVOO**) as a **FUNCTIONAL INGREDIENT** would be particularly interesting due to its recognized health-promoting capacity [2]. However, the direct addition of EVOO to food is challenging due to its liquid state.



Liquid oil conversion into a solid-like material through **OLEOGELATION** could enlarge its possible applications, increasing the technological performances, while reducing hard stock fat content in food [3].

AIM

To understand the digestibility of EVOO triglycerides in oleogels obtained by different oleogelators

Material & Methods



Monoglycerides (MG)

Rice Bran Wax (RW)

Sunflower Wax (sw)

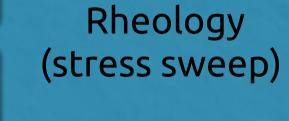
β-sitosterol/ γ-oryzanol mixture (PS)

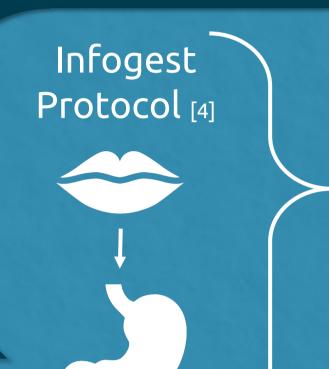
10% (w/w)



Polarized Light
Microscopy
(PLM)







IN-VITRO DIGESTION

After the intestinal phase pH-stat protocol was performed to assess Free Fatty Acid release (FFA) [5]

$$FFA (\%) = \frac{V_{NaOH \ over \ time}}{V_{total \ teoric \ NaOH}} * 100$$

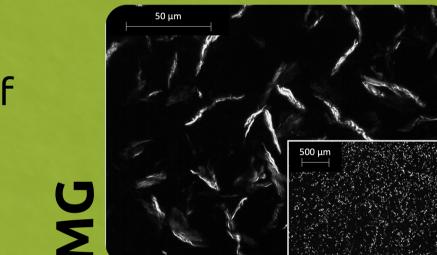
Confocal Light Scattering Microscopy (CLSM)

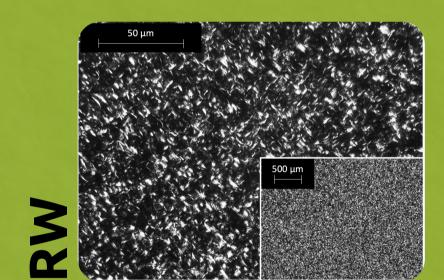
OLEOGEL PREPARATION and PHYSICAL CHARACTERIZATION

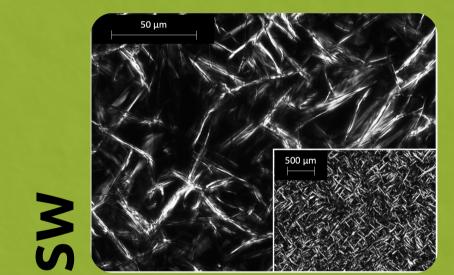
MICROSCOPIC APPEARANCE

PLM showed **needle-like crystals** in MG, SW, and RW of different sizes.

PS is not recordable with PLM being a fibrillar network.

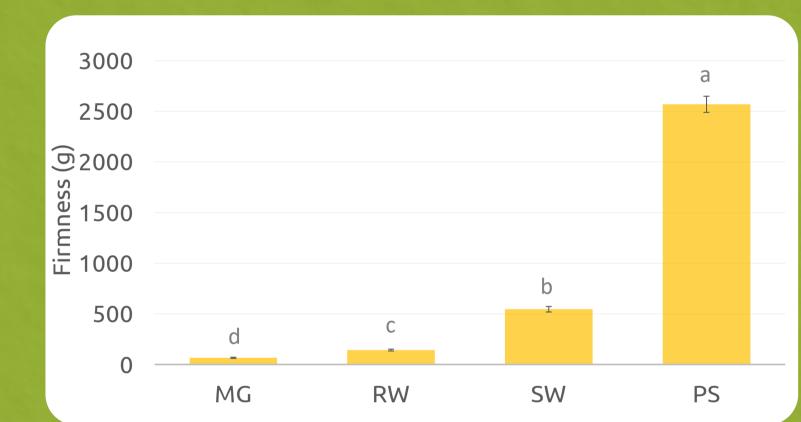


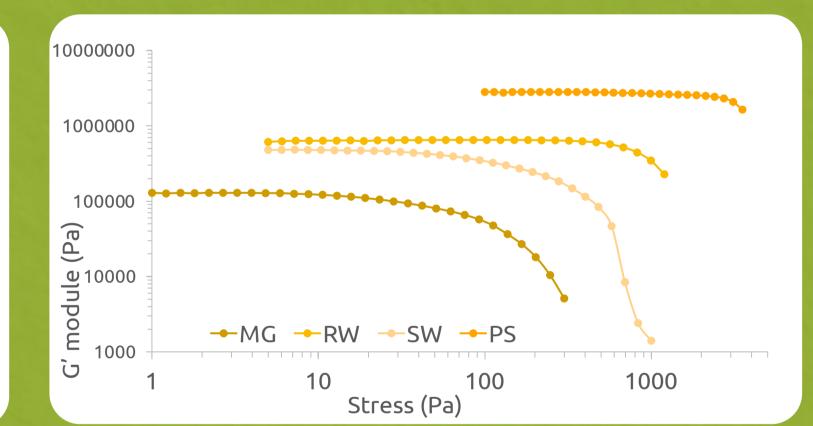




MECHANICAL BEHAVIOUR

Both firmness and critical stress indicated that PS was the strongest gel, followed by SW, RW, and MG. These results were attributed to **the microstructure and the nature of the network**, i.e., fibrillar (PS) or crystalline (MG, RW, and SW).





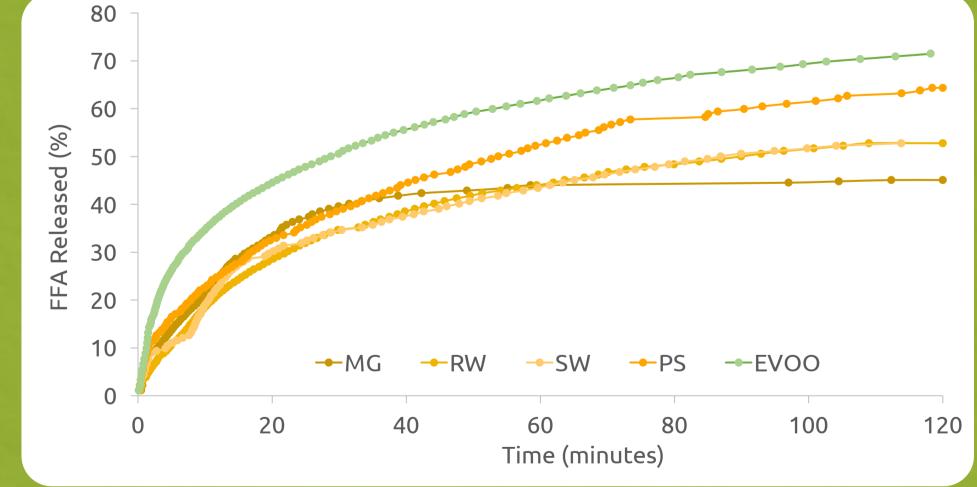
Results & Discussion

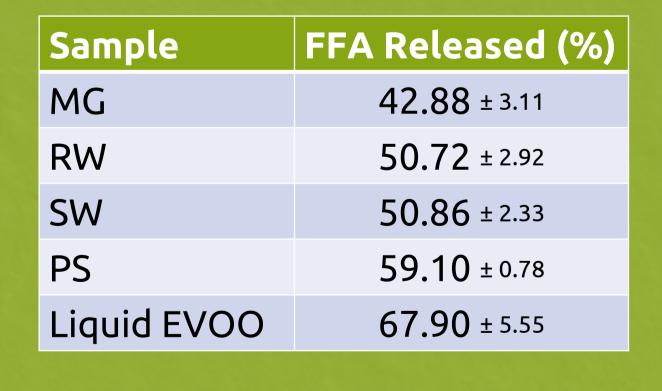
DIGESTION BEHAVIOUR

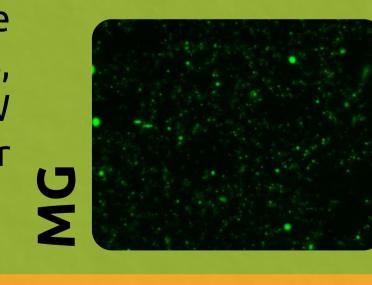
The kinetics of FFAs released differed among samples. All oleogels presented a **lower lipid digestibility** compared to unstructured EVOO. Different structures led to changes in lipid digestibility: MG had the lowest while PS had the highest FFA release values.

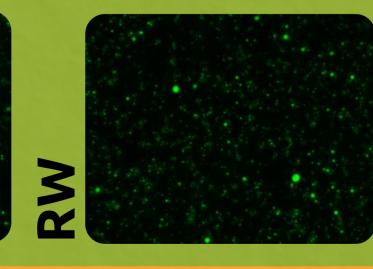
CLSM highlighted the effect of different gelators on the **formation of mixed micelles**

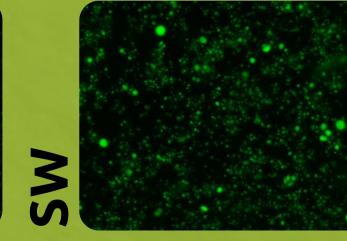
upon intestinal in-vitro digestion.
Larger oil droplets were observed in EVOO and PS, whereas MG, RW, and SW enabled the formation of smaller and more dispersed micelles.

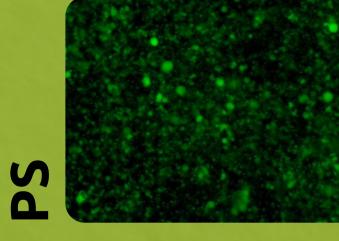


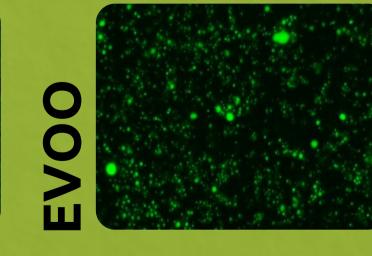












Conclusions

All **gelators** (MG, RW, SW, and PS) **successfully structured EVOO into oleogels** with peculiar physical characteristics. This can enlarge the possible applications of **EVOO in food formulations**.

The entrapment of liquid oil into oleogel networks based on different microstructures allowed modulating FFA release during in vitro digestion. Oleogels can thus represent a promising strategy to tailor lipid digestibility.

References

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