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Adult and older adult *in vitro* digestion of α -tocopherol fortified yogurt using DIDGI®

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Introduction

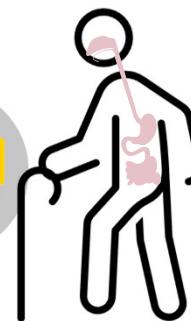
Older adult population is ageing

- Decreased metabolism
- Reduced ability to digest and adsorb food nutrients
- Nutritional deficiencies and malnutrition



Aiming to improve nutraceutical and bioactive compounds delivery extent and functionality

How do these products digest by older adults?



Objectives

- Adapt *in vitro* dynamic digestion protocols to better suit older adult' gastrointestinal function parameters (Méard et al., 2023)
- Understand how adult and older adult digestive parameters modulate the digestion of α -tocopherol fortified yogurts

Methods

Fortified yogurt production

- Fat-free natural stirred yogurt
- Oil-in-Water nanoemulsions (w/ lecithin and sunflower oil) encapsulating α -tocopherol



Dynamic *in vitro* digestion - DIDGI® for adults and older adults

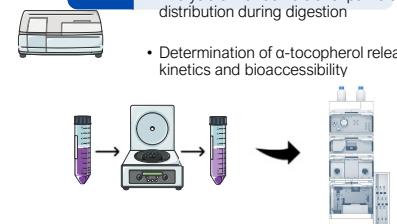


Figure 1. The DIDGI® dynamic *in vitro* system.

	Adult	Older adult
Gastric emptying $t_{1/2}$	70 min	98 min
Pepsin (U/mL)	2000	1200
Pancreatin (%)	7	5.5
Bile salts (%)	4 (0 - 30 min) 30 min - end)	2.7 (0 - 30 min) 1.3 (30 min - end)

Influence of protocol on digestibility and release kinetics

- Analysis of nanoemulsions' particle size distribution during digestion
- Determination of α -tocopherol release kinetics and bioaccessibility



Results

Adult *in vitro* digestion (gastric phase)

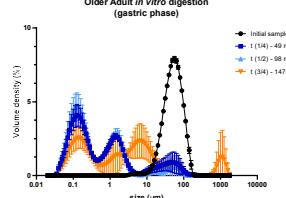
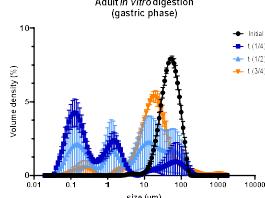
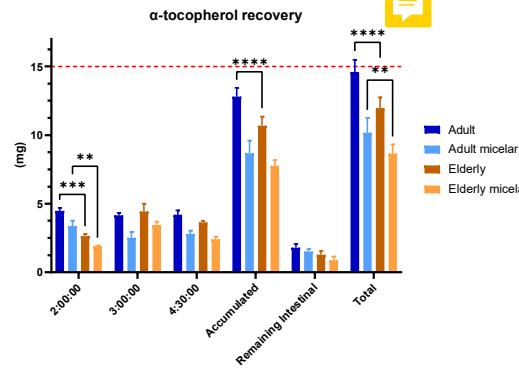


Figure 2. Particle size distribution stability during the gastric phase of the *in vitro* digestion using the adult and the older adult protocol. Samples were diluted in SDS to avoid aggregation effects.

- Samples' stability was similar between protocols until $t_{1/4}$.
- Aggregation effects were more pronounced in the adult protocol after $t_{1/2}$.
- Adult protocol was substantially more effective in degrading and homogenizing the gastric content.

- Superior α -tocopherol concentrations at initial stages of the intestinal phase were obtained using the older adult protocol.
- Gastric digestion by-products on adult protocol (i.e., higher content of peptides and amino acids) may have hindered initial α -tocopherol release.
- α -tocopherol release kinetics were greatly affected by the application of different digestion parameters, affecting its release extent and profile.

α -tocopherol recovery



- α -tocopherol was entirely recovered using the adult protocol (i.e., 14.60 ± 0.89 mg), whereas in the older adult it was not (i.e., 11.96 ± 0.7 mg).
- O/W nanoemulsions produced and the selected food matrix granted high stability to α -tocopherol (i.e., 97.3 ± 5.9 and 79.8 ± 5.2 % for the adult and older adult protocol, respectively)
- α -Tocopherol release balance was greatly affected by the protocol applied until 2 h, however release balance attained at the end of the digestion was similar in both protocols.
- Bioaccessibility of the delivered samples throughout the intestinal phase of the *in vitro* digestion ranged between 60.54 ± 7.38 and 78.90 ± 8.88 %

Figure 4. α -Tocopherol recovery from the digesta and micellar phase of samples delivered from the intestinal reactor. Red dotted line indicates the total amount of α -tocopherol added into the fortified yogurt. Statistically significant differences are indicated by ** ($p < 0.01$), *** ($p < 0.001$) and **** ($p < 0.0001$).

α -tocopherol concentration per meal percentage (intestinal phase)

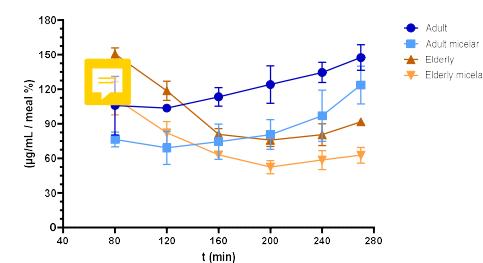


Figure 5. α -Tocopherol release kinetics during the intestinal phase of the *in vitro* digestion using adult and older adult protocols..

α -tocopherol release balance

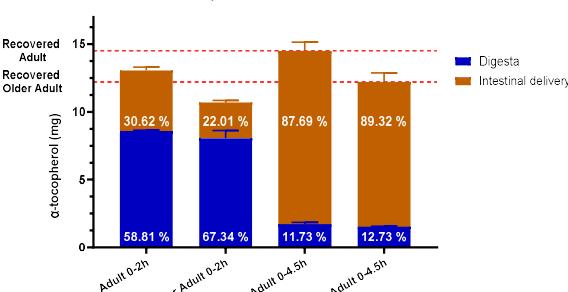


Figure 6. α -Tocopherol release balance of adult and older adult models at 0 and 4.5 h. Blue bar indicates the amount of α -tocopherol within the intestinal reactor, whereas orange bar indicates the amount of α -tocopherol delivered from the intestinal reactor. Red dotted lines indicate the total amount of α -tocopherol recovered when using the adult or older adult model.

- The adult protocol was substantially more effective in degrading and homogenizing the gastric content, despite the similar degradation of the fortified yogurt at earlier stages of digestion.
- Gastric phase duration and the obtained by-products from gastric digestion interfere with the consequent intestinal degradation and with the release of the bioactive compound of interest.
- Bioaccessibility of the delivered samples throughout the intestinal phase of the *in vitro* digestion ranged between 60.54 ± 7.38 and 78.90 ± 8.88 , only presenting statistically significant differences at 3 hours which could be attributed to the prolonged gastric emptying on the older adult protocol.
- The adaptations to the *in vitro* dynamic protocol caused significant impact on the fortified yogurt proteins' hydrolysis during the gastric phase and on the α -tocopherol release rate and extent.