

Dairy proteins' effect on metabolism in obese non-diabetics

Jan Stanstrup^a, Daniela Rago^a, Jens Holmer-Jensen^b, Kjeld Hermansen^b, Lars O. Dragsted^a

^aDepartment of Human Nutrition, Faculty of Life Sciences, University of Copenhagen, Frederiksberg C, Denmark.

^bDepartment of Endocrinology and Metabolism MEA, Aarhus University Hospital, Aarhus, Denmark.

Objectives

- ☐ To investigate time-dependent changes in the metabolome following ingestion of different proteins from whey
- ☐ To identify protein specific markers
- ☐ To automatically group mother compounds, adducts and fragments

Intervention study

- Randomized single-blinded full crossover study
- 11 obese non-diabetic subjects
- Meals containing 80 g of fat, 45 g of carbohydrate and 45 g of protein
- The protein source was α-lactalbumin (ALPH), whey isolate (WI), whey hydrolysate (WH) or caseinoglycomacropeptide (CGMP)
- Blood samples drawn at five time points during 8-h postprandial period

Compound	T _r	[M+H] ⁺	Fragments, adducts and unknown features
	0.60		164.0
	0.60		180.0
	0.61	188.0	210.0
Proline	0.66	116.1	70.1
Methionine	0.95	150.1	133.0, 104.1, 87.0, 61.0
Tyrosine	1.11	182.1	165.1, 147.0, 136.1, 123.0, 119.1
	1.35		218.1
Isoleucine	1.36	132.1	263.2, 86.1, 69.1
	1.40		199.1
Mevalonolactone	1.42	131.1	
Phenylalanine	1.43	166.1	149. 1, 103.1, 120.1, 107.1, 93.1
	1.51		201.1
Tryptophan	1.52	205.1	409.2, 188.1, 170.1, 159.1, 146.1, 144.1, 132.1, 118.1, 100.1
γ-Glu Leu / γ-Glu lle	1.61	261.1	244.1, 132.1
	2.37		247.1

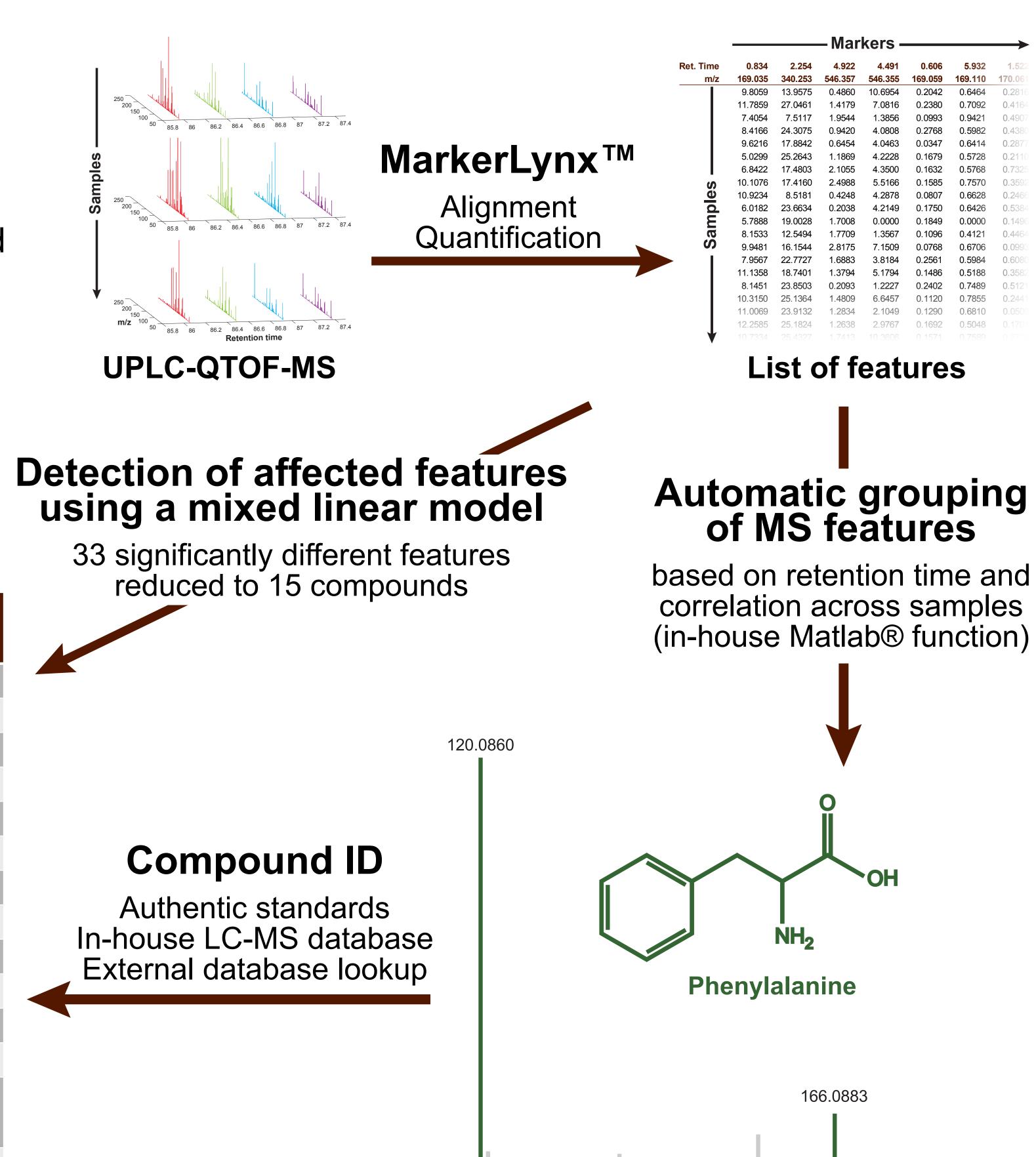
List of features significantly different between meals

Results

- The features were dominated by amino acids (AAs)
- The AA composition in the protein sources was largely reflected in the blood levels of the individual AAs
- Approximately a 30 40 % change in intake of individual AAs could be detected with this method
- The aromatic AAs (Phe, Tyr, Trp) and mevalonolactone were found at a lower level for the CGMP group whereas the group had higher levels of Pro and Ile
- The different whey groups also showed different levels of AAs (as exemplified by phenylalanine to the right)
- Two markers with masses of 188 and 199 Da were 20 fold higher in plasma after ingestion of WH compared to the other protein sources

Conclusions

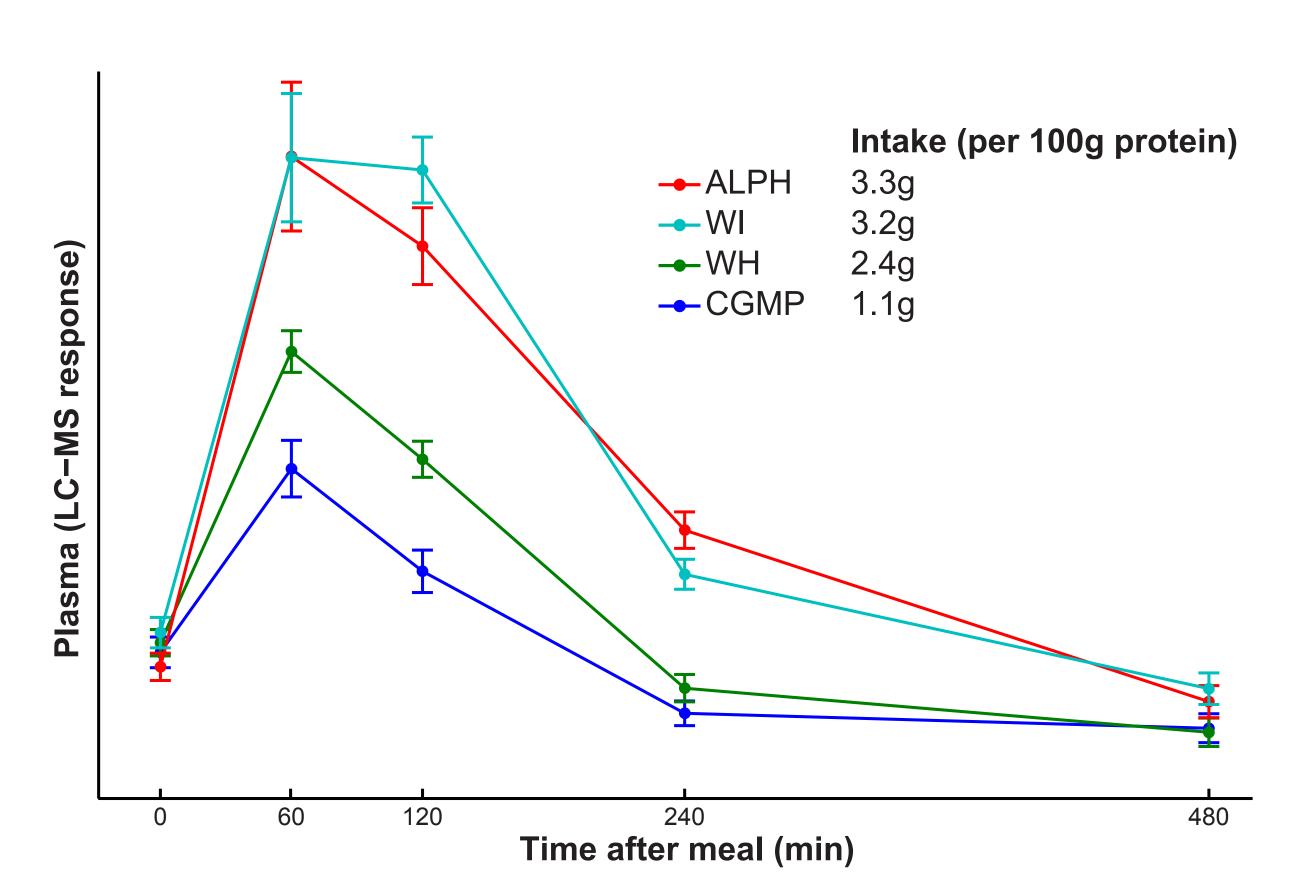
- ✓ Protein sources were differentiated based mainly on amino acid composition
- ✓ Strong markers of whey hydrolysate have been distinguished but remain to be identified
- ✓ Fragments, polymers and adducts were automatically grouped using a simple Matlab® algorithm



85 90 95 100 105 110 115 120 125 130 135 140 145 150 155 160 Masses corresponding to phenylalanine

103.0501

93.0503



149.0757

Kinetic profile for Phe extracted from metabolomics profiles. Error bars indicate the standard error of the mean