

COMPARATIVE EFFECTS OF MICRO vs. NANO FILLER EMULSIONS ON TEXTURAL PROPERTIES OF MYOFIBRILLAR PROTEIN COMPOSITE GELS

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I. INTRODUCTION

The interactions between protein and oil to form a composite gel matrix play an important role in textural properties [1]. Commercial comminuted and restructured meat products typically comprise a continuous protein matrix filled with emulsified fat droplets, and in the composite gel system, myofibrillar protein (MP) plays a principal structural role [2]. The aim of this research is to test the effects of nano-emulsions vs. micro-emulsions on the rheological, water-binding, and structural properties of emulsion gels. MP-stabilized O/W emulsions in nano- vs. micro-scale particles have different effects on the physical properties of emulsion gels.

II. MATERIALS AND METHODS

MP-based oil-in-water micro-emulsions and nano-emulsions were compared with each other and with lecithin-stabilized micro-emulsions and nano-emulsions. For emulsion preparation, MP (1 mg/mL) was the testing emulsifier, and lecithin (1 mg/mL) was the comparative surfactant in canola oil (10% v/v)-based water-in-oil (O/W) emulsions. High-pressure homogenization (40 MPa) was applied to produce nano-emulsions and high-speed homogenization (17,000 rpm) was used to prepare micro-emulsions. The physicochemical properties of emulsions and emulsion gels were analyzed. Emulsion particle size, ζ -potential, and morphological properties (transmission and confocal microscopies) were analyzed; dynamic rheological behaviors (storage and loss modulus), mechanical strength, water-holding capacity, water mobility, and protein secondary structures of the emulsion gels (2.5% protein, 5% oil) were measured. Statistical analyses were performed using the SPSS software. All experimental values were measured in triplicates, and the results are presented as the mean \pm standard deviation. Means within groups were compared using a one-way analysis of variance followed by Duncan's multiple range test ($p < 0.05$).

III. RESULTS AND DISCUSSION

The gel strength of pure MP gel (control) was significantly lower than that of emulsion gels due to induce the interaction between protein-oil in the gel matrix [3]. The MP micro-emulsion gel (~500 nm) was slightly stronger than the MP nano-emulsion gel (~2 μ m), consistent with the trend seen in moduli tests. However, emulsion gel embedded with MP-nano showed the highest water-holding capacity and water mobility, and almost no cooking loss when compared with all other composite gels. Partial structural unfolding protein suggests the improved interfacial activity of MP in nano-emulsions, and this is supported by the confocal imaging results where MP-nano displayed numerous smaller droplets distributed within the MP matrix. Moreover, depending on the emulsion type, the secondary structure was affected. Overall, emulsion gels tended to be stronger than oil-free control gels, and MP-based emulsions were more effective than lecithin-stabilized emulsions for modifying the gelling properties due to a visible interfacial protein film formed that prevented oil droplet aggregation.

Table 1 Characterization of MP solutions, micro-emulsions, and nano-emulsions

Treatments	Particle size (nm)	[-] ζ -potential (mV)	Hydrophobicity	DSC 1 st peak		DSC 2 nd peak	
				Enthalpy (mJ/g)	Peak ($^{\circ}$ C)	Enthalpy (mJ/g)	Peak ($^{\circ}$ C)
5% MP			776.1	569.6	57.6	12.4	67.4
Non-treated HPH			982.9	348.4	56.2	11.6	68.5
Treated HPH			1143.2	239.3	59.0	5.8	66.8
MP Micro	2090.7 \pm 587.4 ^a	10.3 \pm 3.3 ^a					
MP Nano	522.1 \pm 190.5 ^c	10.3 \pm 3.3 ^a					
Lec Micro	1330.0 \pm 229.1 ^b	12.6 \pm 0.7 ^a					
Lec Nano	543.5 \pm 84.7 ^c	11.5 \pm 0.5 ^a					

5% MP, 5% myofibrillar protein in sodium phosphate buffer (0.6 M NaCl, pH 6.25); Non-treated HPH, treated high speed homogenizer; Treated HPH, treated high speed homogenizer and high-pressure homogenizer. MP, myofibrillar protein; Lec, lecithin; micro, micro-emulsion; nano, nano-emulsion. ^{a-c} Different letters represent significant differences at $p < 0.05$.

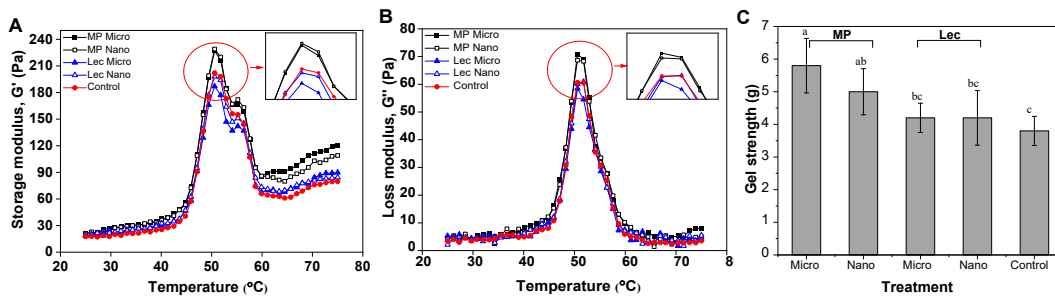


Figure 1. Storage (A), loss (B) modulus and gel strength (C) of emulsion gel. MP, myofibrillar protein; Lec, lecithin; micro, micro-emulsion; nano, nano-emulsion; control, 2.5% MP gel. ^{a-c} Different letters represent significant differences at $p < 0.05$.

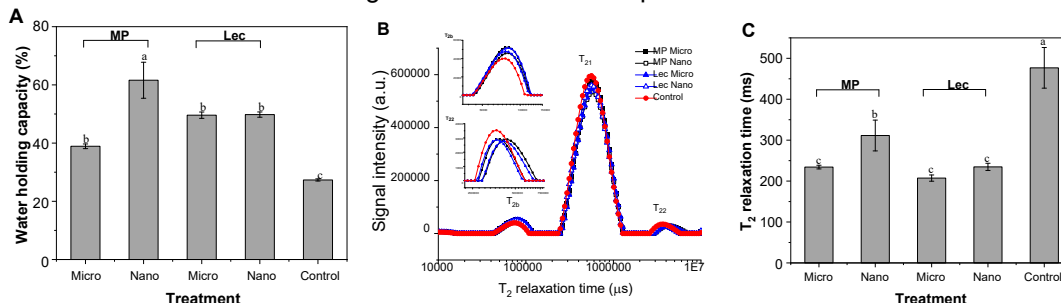


Figure 2. Water-holding capacity (A), transverse relaxation curves (B), and T_2 relaxation time (C) of emulsion gel. MP, myofibrillar protein; Lec, lecithin; micro, micro-emulsion; nano, nano-emulsion; control, 2.5% MP gel. T_{2b} , T_{21} , and T_{22} indicate bound, immobilized, and free water, respectively. ^{a-c} Different letters represent significant differences at $p < 0.05$.

IV. CONCLUSION

Based on the results, protein-based emulsions were preferred over lecithin-based emulsions, and MP nano-emulsions improved moisture retention in cooked MP gels.

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