

Multifunctionality of rapeseed meal protein isolates prepared by sequential isoelectric precipitation

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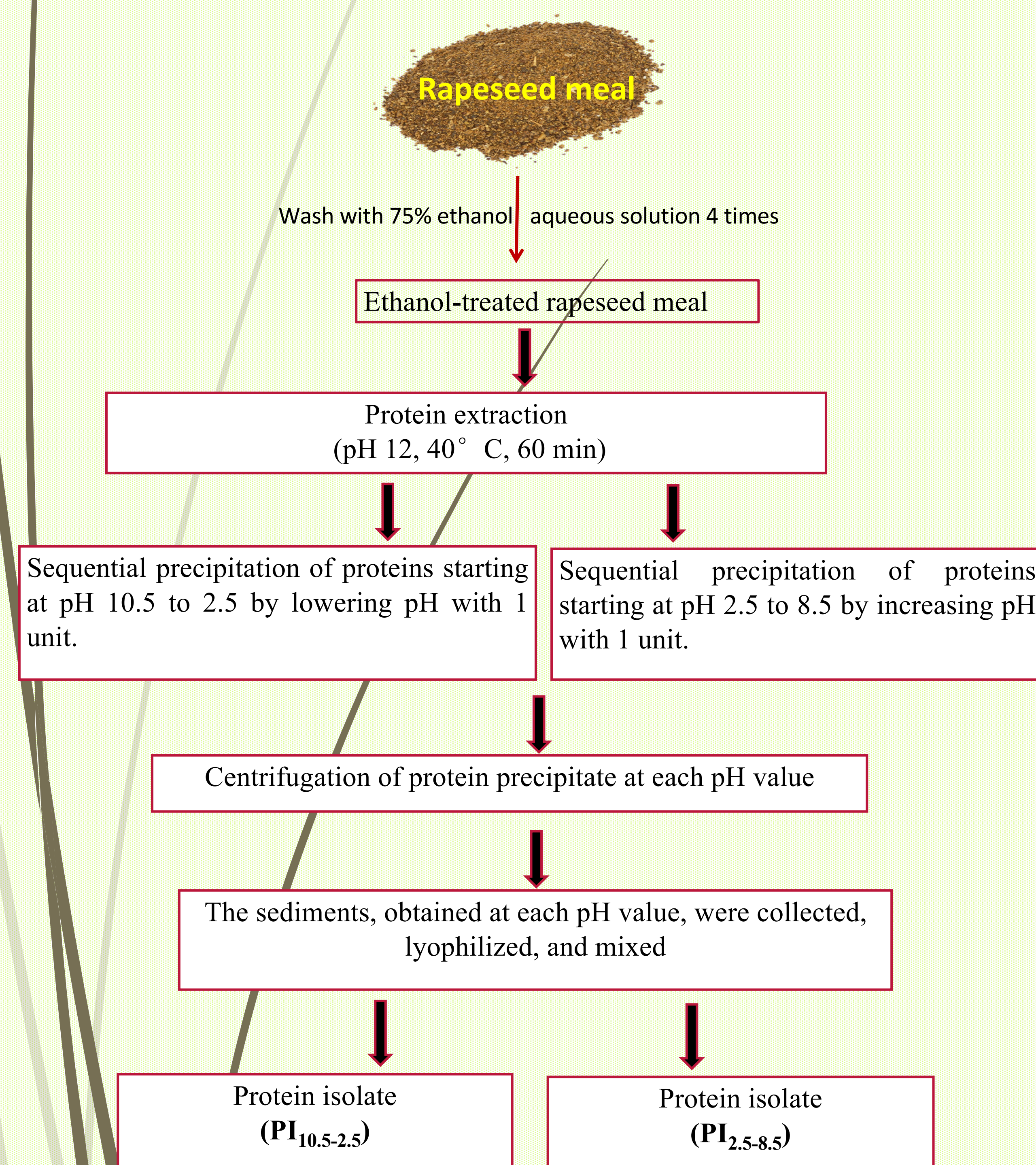
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Background and objectives: Rapeseed oil is a valuable commodity that is widely used in the food industry or other industrial sectors for the production of biofuel, paper, textile, plastics, lubricants, and surfactants. The enhanced demand for this vegetable oil world-wide results in the generation of high amounts of rapeseed meal as a byproduct which reached 38.8 million tons in 2018. In addition to being used as an inexpensive protein-rich ingredient in feed formulation, the rapeseed meal is an appropriate source for preparation of protein isolates for the food industry. **THE AIM OF THE STUDY** was to generate rapeseed meal protein isolates with enhanced solubility and evaluate their functional properties.

METHODS

Preparation of protein isolates



Antioxidant properties of PI_{2.5-8.5} and PI_{10.5-2.5}.

- DPPH method
- FRAP method
- Hydroxyl radical scavenging assay (2-deoxy-D-ribose method)

Emulsifying properties of PI_{2.5-8.5} and PI_{10.5-2.5}.

- Emulsifying activity
- Emulsion stability

Acknowledgment: This study was financially supported by the Bulgarian National Science Fund, project □ KP-06-H37/21.

RESULTS

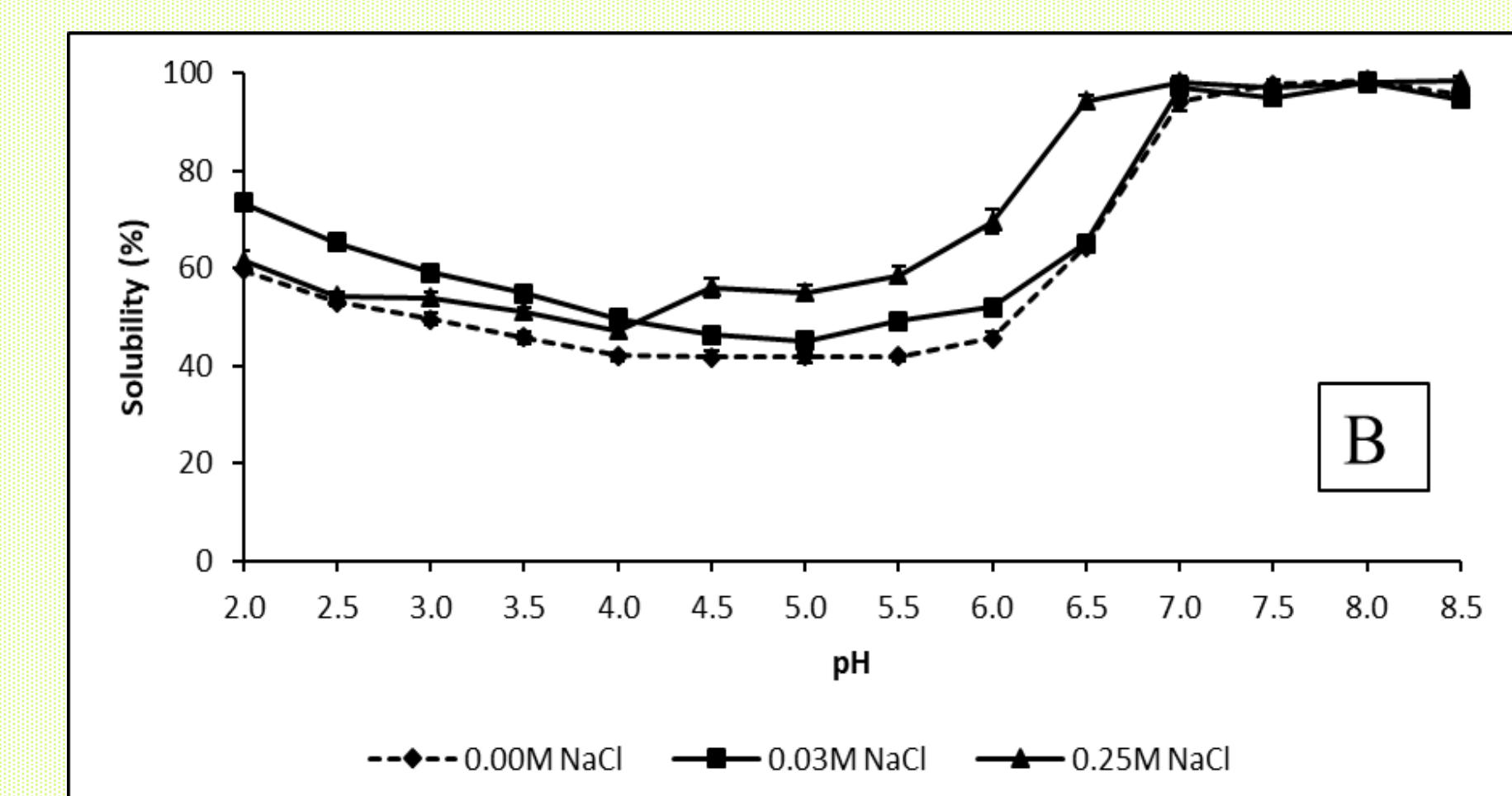
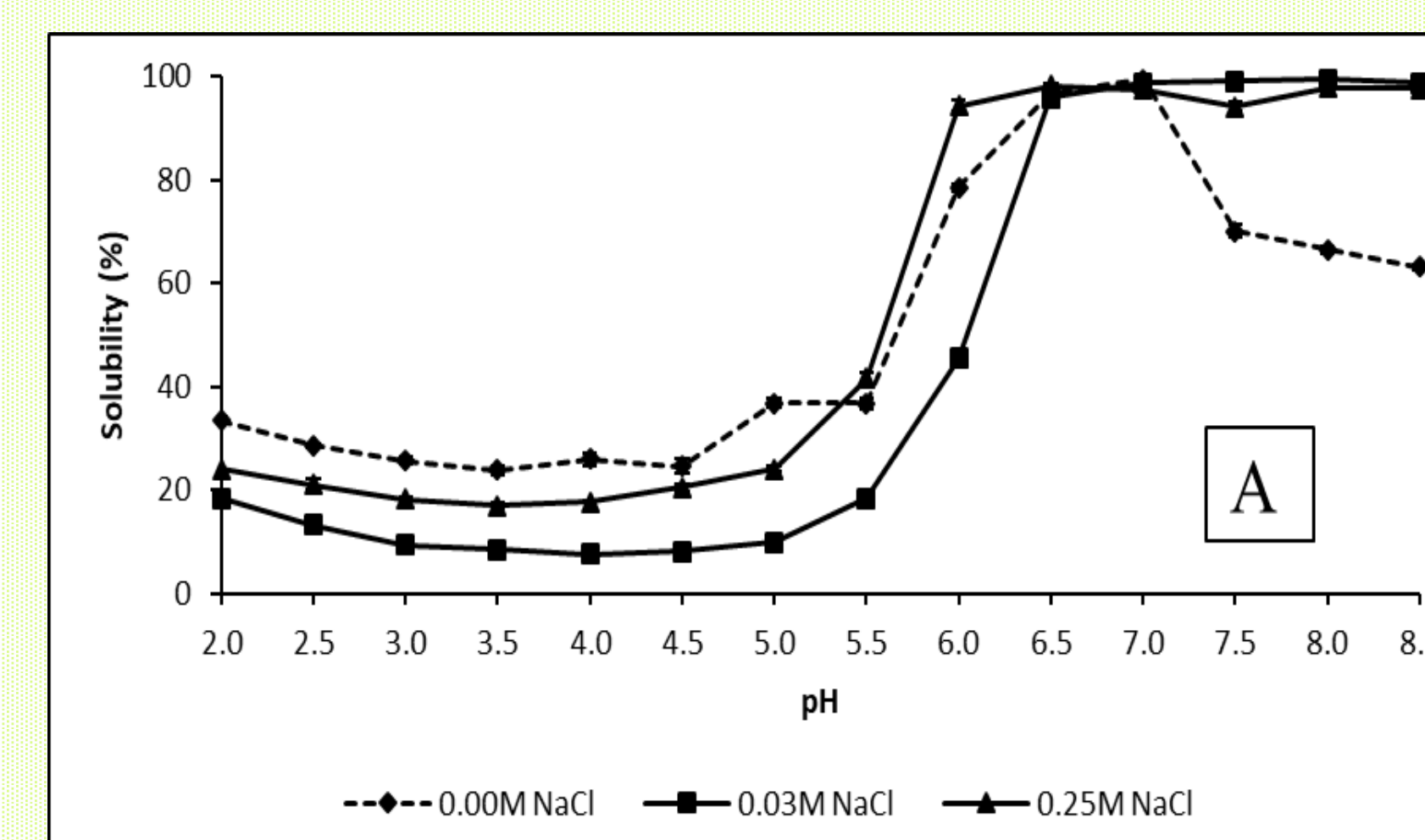


Figure 1. Solubility of protein isolates PI_{2.5-8.5} (A) and PI_{10.5-2.5} (B) at different pH values and NaCl concentrations.

Antioxidant properties

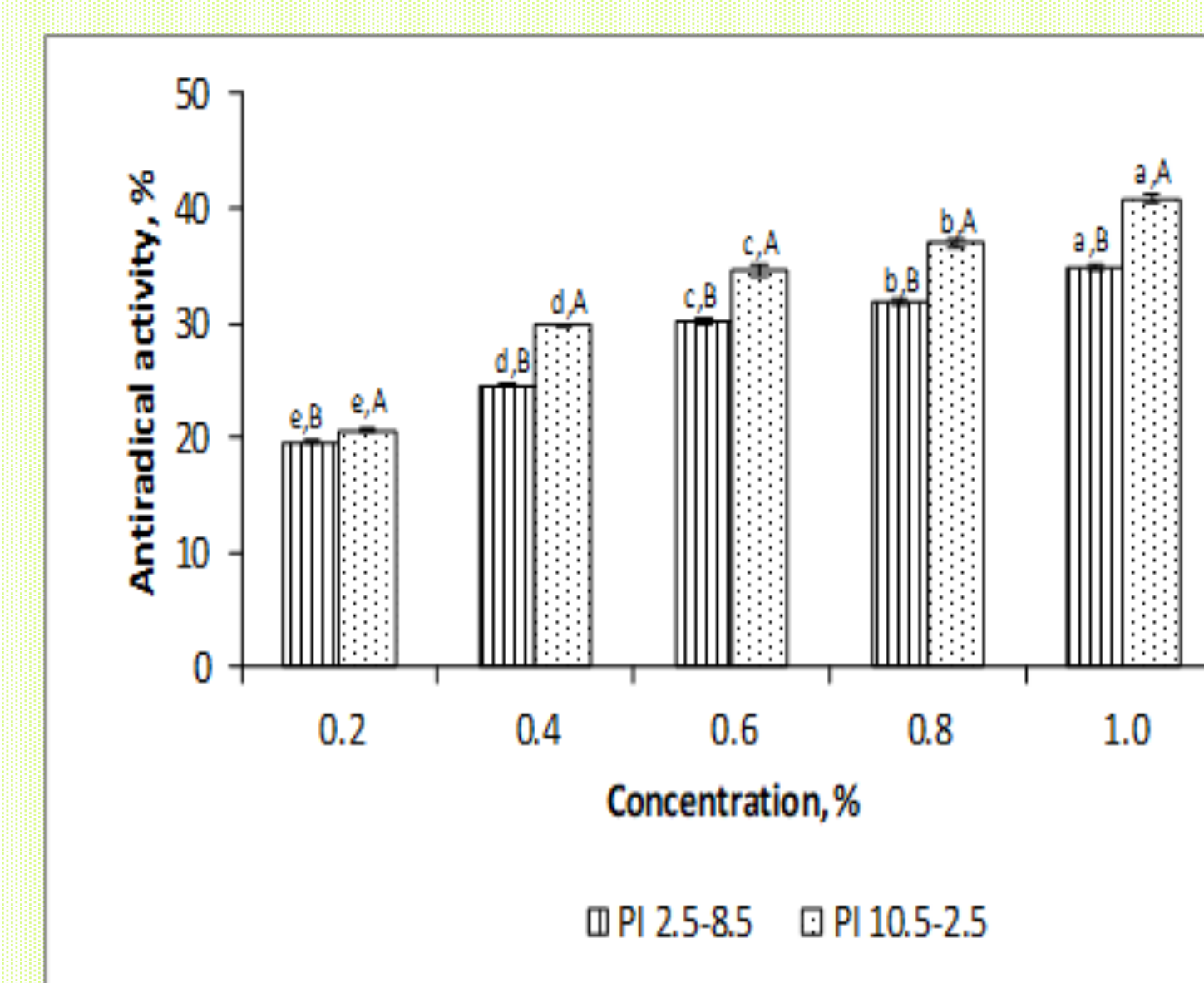


Figure 2. Scavenging effect of ethanol-treated rapeseed meal protein isolates, PI_{2.5-8.5} and PI_{10.5-2.5}, on 2,2-diphenyl-1-picryl hydrazyl (DPPH) radical. ^{a-c} Means of one sample with different concentrations without a common letter differ significantly ($p < 0.05$). ^{A-B} Means of the samples with one and the same concentration without a common letter differ significantly ($p < 0.05$).

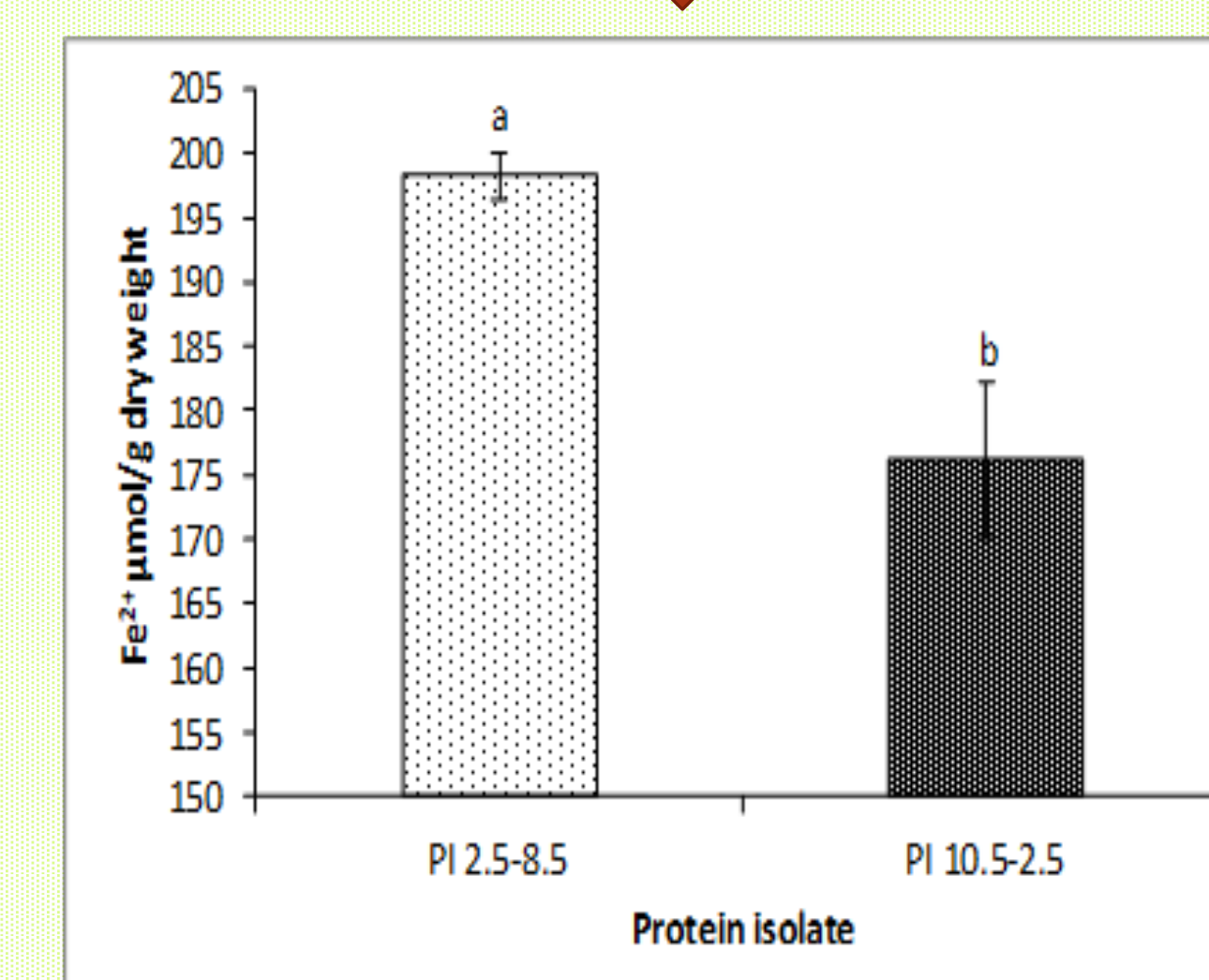


Figure 3. Ferric reducing antioxidant power (FRAP) of ethanol-treated rapeseed meal protein isolates, PI_{2.5-8.5} and PI_{10.5-2.5}. ^{a-b} Means with different superscripts differ significantly ($p < 0.05$).

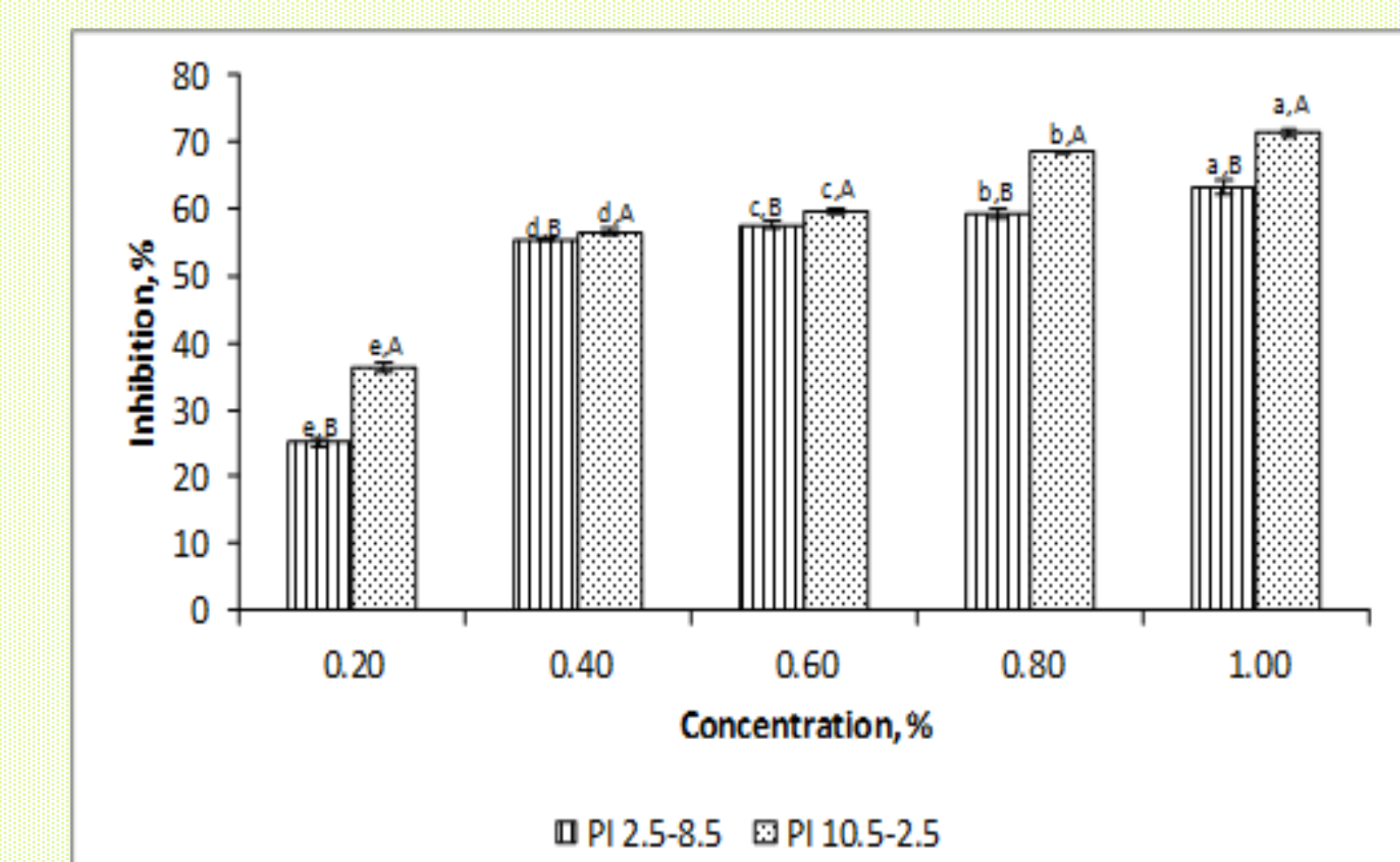


Figure 4. Hydroxyl radical scavenging activity of ethanol-treated rapeseed meal protein isolates, PI_{2.5-8.5} and PI_{10.5-2.5}. ^{a-c} Means of one sample with different concentrations without a common letter differ significantly ($p < 0.05$). ^{A-B} Means of the samples with one and the same concentration without a common letter differ significantly ($p < 0.05$).

Emulsifying properties

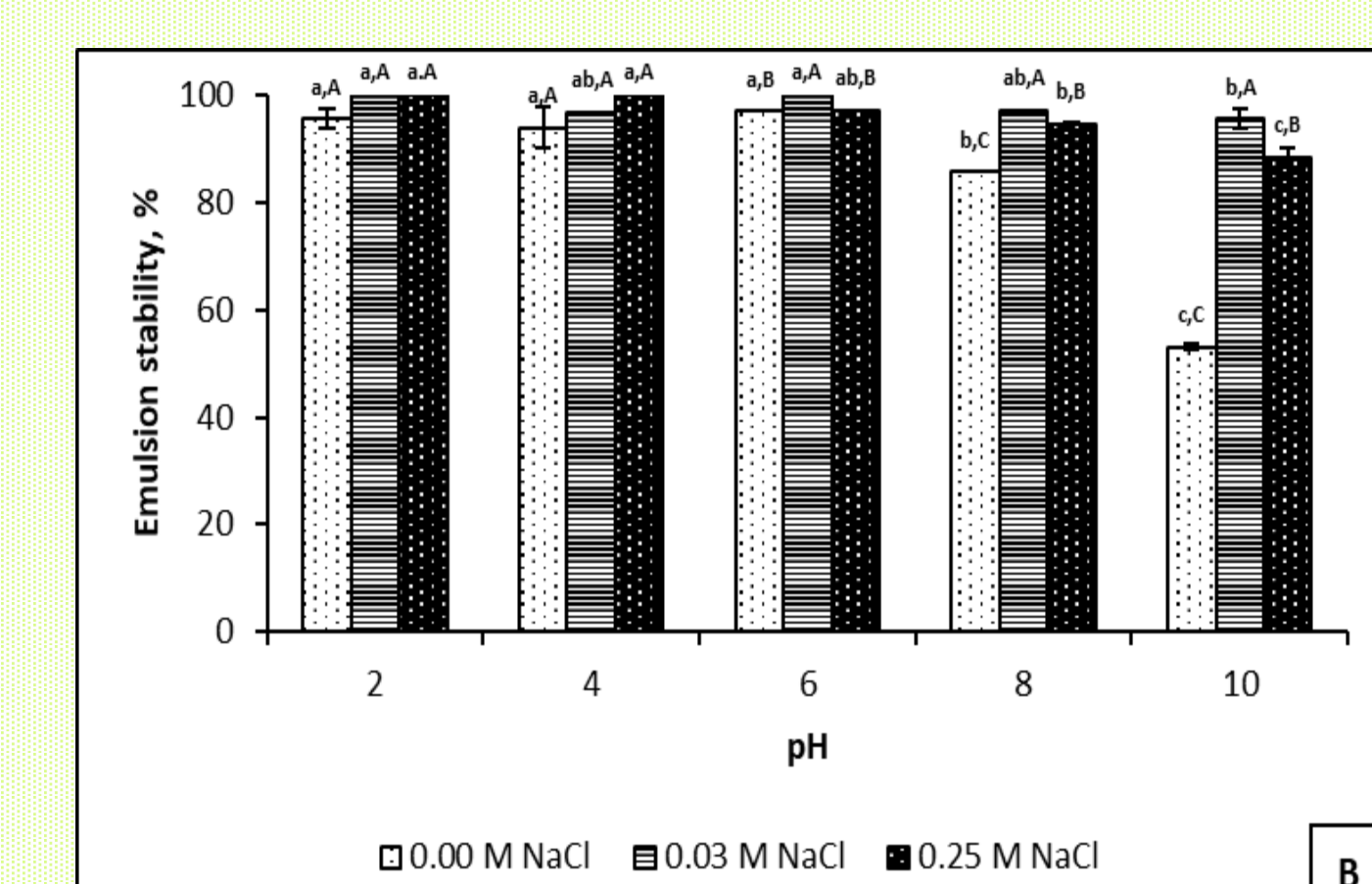
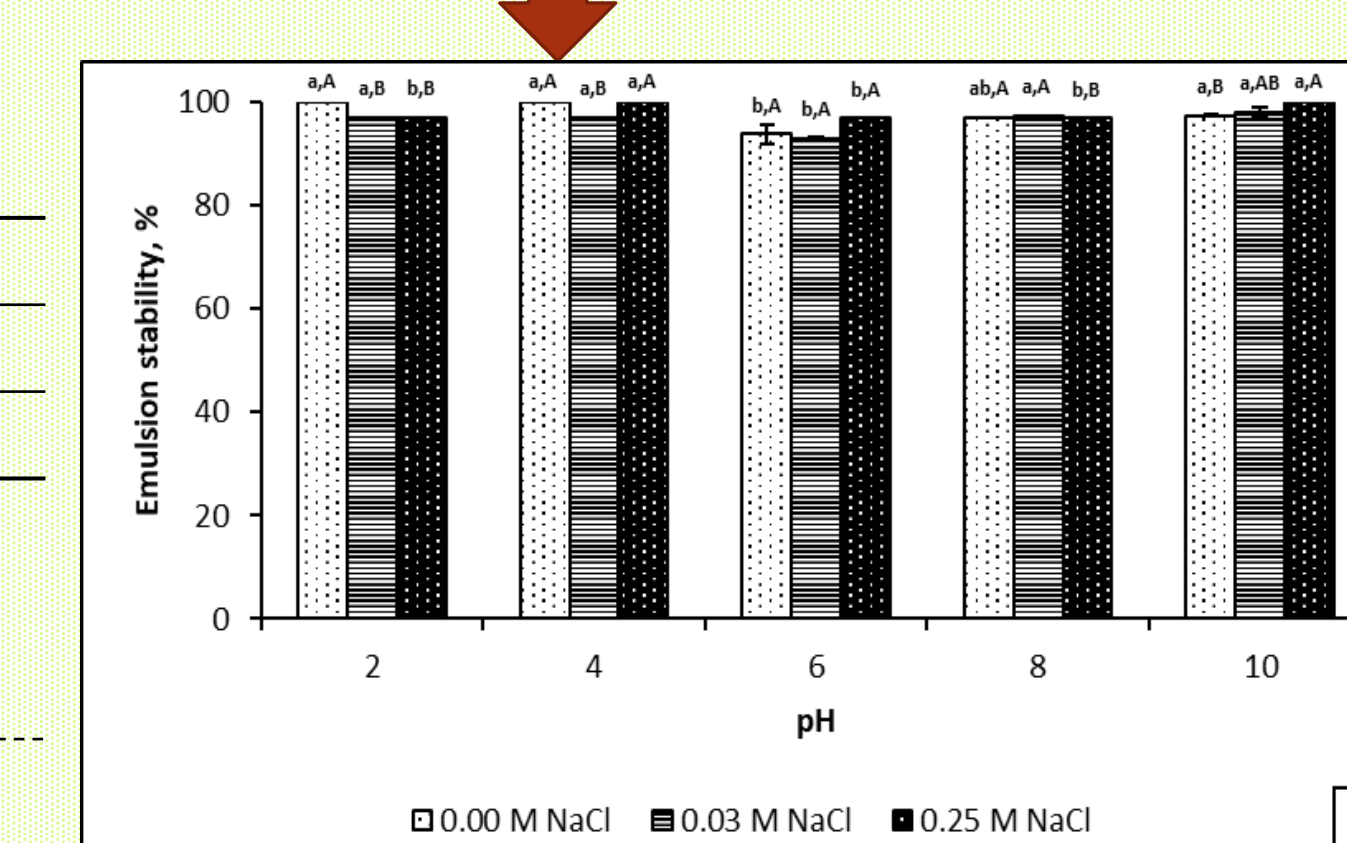


Figure 5. Emulsion stability of PI_{2.5-8.5} (A) and PI_{10.5-2.5} (B) at different pH and NaCl concentrations. ^{a-b} Means without the same lowercase letter for a particular NaCl concentration differ significantly ($p < 0.05$). ^{A-B} Means without the same capital letter for a particular pH differ significantly ($p < 0.05$).

Conclusions

- PI_{10.5-2.5} is more soluble than PI_{2.5-8.5}.
- PI_{10.5-2.5} has higher scavenging capacity on DPPH and hydroxyl radicals than PI_{2.5-8.5}.
- PI_{2.5-8.5} exhibits high emulsion stability which is weakly responsive to pH changes from 2 to 10.