ANION-EXCHANGE SEPARATION OF WHEY PROTEINS

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Three strong anion–exchange membranes (CIM QA, Q100 and HiTrap Q) and reversed–phase highperformance liquid chromatography(15 μ m particle with a pore size of 300 Å) were investigated for the separation of the major proteins, which were contained in whey, such as σ -Lactalbumin, BSA and β -Lactoglobulin contained. Experiments were performed to determine the optimum mobile phase composition for separating the whey proteins using the standard chemicals of the proteins. For strong anion–exchange membranes, the mobile phase was buffer A (20 mM piperazine–HCl pH 6.4) and buffer B (buffer A + 1 M NaCl) and the linear gradient elution changes of salt concentration were applied. For Reversed–phase high–performance liquid chromatography, the mobil phase were consisted of a linear gradient of the two mobile phases of 0.1% trifluoroacetic acid in water and 0.1% trifluoroacetic acid in acetonitrile. The standard chemicals of the proteins were used to investigate the optimal mobile phase compositions with the three anion–exchange membranes. It was experimentally confirmed that HiTrap Q was the most effective to resolve the whey proteins.

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