

**Opinion Article** 

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## Description

Capillary Electrophoresis (CE) has emerged as a powerful analytical technique for the separation and quantification of biomolecules. This versatile method offers high resolution, rapid analysis, and minimal sample requirements, making it particularly valuable in the field of bioanalytical chemistry. In this article, we will explore the principles of capillary electrophoresis and its applications in the quantitative analysis of biomolecules.

Capillary electrophoresis is a separation technique based on the differential migration of charged species in an electric field within a narrow capillary. The capillary, typically made of fused silica, is filled with an electrolyte solution, and an electric field is applied across it. Biomolecules, such as proteins, nucleic acids, and peptides, are separated based on their size, charge, and shape as they migrate through the capillary.

One of the key advantages of CE is its ability to achieve high resolution in a short analysis time. This is attributed to the high efficiency of the separation process in the narrow capillary, reducing the band broadening effects seen in traditional gel electrophoresis.

Quantitative analysis in capillary electrophoresis involves determining the concentration of specific biomolecules in a given sample. This is achieved by relating the peak area or height of the separated analytes to known standards of the same or similar compounds. Calibration curves are constructed by analyzing standards with known concentrations, and these curves are then used to quantify the target biomolecules in unknown samples.

Capillary electrophoresis is widely used for the separation and quantification of proteins and peptides. It offers high resolution and sensitivity, making it suitable for applications such as the analysis of complex protein mixtures, monitoring protein purity, and assessing post-translational modifications.

CE is also employed for the analysis of nucleic acids, including DNA

and RNA. It enables the separation of DNA fragments with high rtgekukqp." o cmkpi"kv"xcnwcdng"kp"FPC"ugswgpekpi."igpqv{rkpi."cpf"vjg fgygevkqp"qh" o wvcvkqpu0

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