



DETERMINATION OF **PROTEINS CONCENTRATIONS** AND THEIR **MOLECULAR WEIGHTS** BY CAPILLARY GEL ELECTROPHORESIS

INTRODUCTION

The method enables fast separation of proteins according to their molecular weights (Mw) with subsequent protein quantification in protein-containing samples.

MEASUREMENT METHOD

Capillary electrophoresis for the separation and determination of proteins is based on the differential migration of SDS-protein complexes in a narrow fused-silica capillary filled with a low viscous gel, under the influence of the applied electric field. Detection of proteins is performed based on their own absorbance at 220 nm. Due to the presence of a low viscous gel SDS-protein complexes are separated only according to their Mw and thus it enables also to determine Mw of an unknown protein(s).

ADVANTAGES OF THE CAPILLARY ELECTROPHORESIS METHOD

Compared with protein separation with SDS-PAGE, capillary electrophoresis has several important advantages:

- Full automatization
- Direct protein quantification
- Absence of coloring
- Low analysis cost
- Short analysis time

EQUIPMENT AND REAGENTS

The "CAPEL[®]-105/105M" capillary electrophoresis system is used in all measurements.

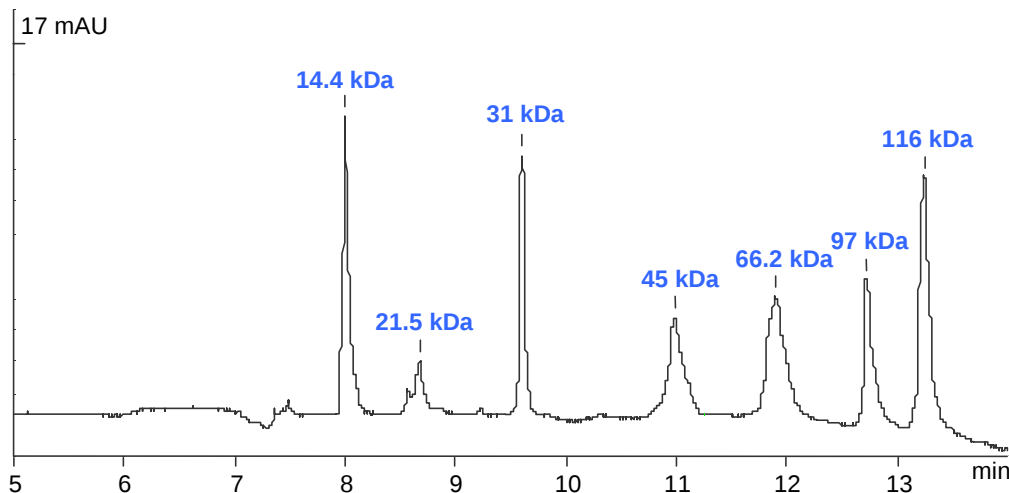
Sample preparation, capillary conditioning and analysis are done according to the detailed protocol, included in the kit.

All reagents must be of analytical grade or higher.

Data acquisition and integration is accomplished with a PC with "Windows[®] 95/98/NT/ME/2000/XP", using chromatographic software "Chrom&Spec[®]" (for "CAPEL[®]-105") or "ELFORUN[®]" software package (LUMEX[®]) for "CAPEL[®]-105M".

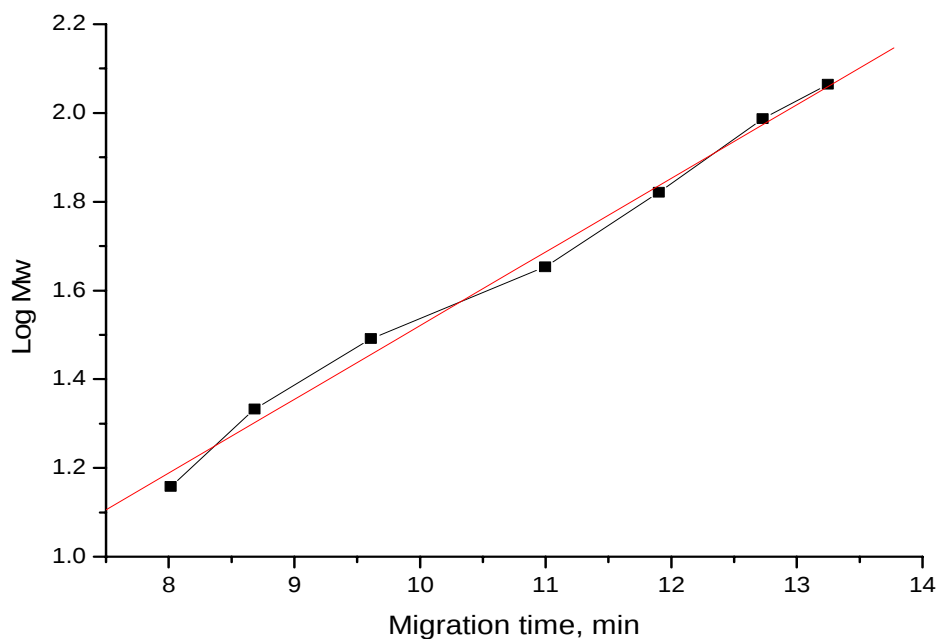
EXAMPLE OF A REAL ANALYSIS

Sample: standard proteins with different Mw (from 14.4 kDa to 116 kDa)





DEPENDENCE OF $\text{Log } M_w$ OF PROTEINS FROM THEIR *MIGRATION TIME*



This dependence enables direct determination of molecular weight of the unknown protein(s) based on its migration time.

The contents on this paper are subject to change without notice.
To get more specific information on this method please contact the developer of this method LUMEX Ltd.:
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