

ABSTRACT

Capillary isotachopheresis (ITP) with conductivity detection has been used for separating and determining of glucosamine (GA) from the SYSADOA group (symptomatic slow acting drugs of osteoarthritis). The optimised ITP electrolyte system consisted of 10 mM potassium picolinate and 10 mM picolinic acid (pH 5.4) as the leading electrolyte (LE) and 10 mM formic acid (pH 3.0) as the terminating electrolyte (TE). The driving and the detection currents were 50 μA (for 340 s) and 10 μA , respectively. A single analysis took < 10 min. Under the optimum conditions the effective mobility of GA was determined as $24,7 \times 10^{-9} \text{ m}^2 \cdot \text{V}^{-1} \text{ s}^{-1}$ when using tetraethylammonium-iodide as the standard of the mobility. The calibration graph was rectilinear in the range 50 – 200 $\text{mg} \cdot \text{l}^{-1}$ of GA with $r^2 = 0,99747$. The RSD was 0,52 % (n=6) when determining 100 $\text{mg} \cdot \text{l}^{-1}$ of GA. The ITP method was applied to the assay of GA content in various pharmaceuticals and nutraceuticals. Acceptable accuracy of the method was confirmed by the added/found GA recovery experiments (recoveries from 96.2-100.3%).