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MICROCHIP CAPILLARY ELECTROPHORESIS WITH AMPEROMETRIC DETECTION FOR METAL ION ANALYSIS

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งานวิจัยนี้เป็นการพัฒนาเพื่อการตรวจวิเคราะห์โลหะหนักหลายชนิดพร้อมกันและตรวจวิเคราะห์ แบบรวคเร็วซึ่งได้แก่โลหะตะกั่ว. แคคเมียม. และทองแคง โคยใช้เทคนิคไมโครซิพคะปีลารีอิเล็คโทรฟอริ ซิสร่วมกับตัวตรวจวัดทางเกมีไฟฟ้า ระบบการตรวจวิเกราะห์โดยตรงแบบแอมเพอโรเมทรีในไมโกรชิพ คะปีลารีอิเล็คโทรฟอริซิสถูกนำมาใช้ในการตรวจวิเคราะห์ไอออนโลหะเหล่านี้ได้เป็นอย่างคื อิทธิพลจาก ศักย์ไฟฟ้าที่ให้กับระบบ สักย์ไฟฟ้าในการตรวจวัด. ความเข้มข้นและพีเอชของบัฟเฟอร์ที่ใช้ในการ วิเคราะห์ต่อค่าการตอบสนองของตัวตรวจวัดซึ่งได้ทำการตรวจสอบและหาค่าที่ทำให้เกิดประสิทธิภาพการ จากผลการทดลองแสดงให้เห็นว่าการวิเคราะห์โดยใช้โซนอิเล็กโทรฟอริซิสในการแยกโลหะ ตรวจวัด ์ ตะกั่ว แกดเมียม และทองแดงได้ในเวลาน้อยกว่า 3 นาที ใช้บัฟเฟอร์เอ็มอีเอส (MES) (พีเอช 7.0. 25 มิลลิโม ลาร์) และแอลฮีสทีดีน (L-histidine), ให้ศักย์ไฟฟ้าในการแยก 1.2 กิโลโวลต์ และศักย์ไฟฟ้าในการตรวจวัด ที่ -0.8 โวลต์ ค่าขีดจำกัดต่ำสุดของการตรวจวัดโลหะตะกั่ว, แคดเมียม, และทองแดงเป็น 1.74, 0.73, และ 0.13 ใมโครโมลาร์ (ค่าสัญญาณกระแสต่อสัญญาณรบกวนมีค่ามากกว่า 3) ค่าเบี่ยงเบนมาตรฐานสัมพัทธ์ ของสัญญาณกระแสไม่เกิน 6 เปอร์เซ็นต์ และของเวลาในการเคลื่อนที่ของสารในชันแนลไม่เกิน 2 ในงานวิจัยนี้ได้แสดงให้เห็นหลักการในการให้ศักย์ไฟฟ้ากับระบบไมโครชิพซีอีซึ่งสามารถ เปอร์เซ็บต์ ้นำไปใช้ได้ในอนาคต นอกจากนี้ยังแสคงให้เห็นการวิเคราะห์ไอออนโลหะในตัวอย่าง จากผลการวิเคราะห์ ทำให้ได้ความมุ่งหมายว่าไมโครชิพซีอีร่วมกับตัวตรวงวัดทางเคมีไฟฟ้านี้จะเป็นอีกวิธีที่ใช้ในระบบการ วิเคราะห์ระดับไมโครสำหรับตรวจวิเคราะห์อาหาร

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SARAWADEE KORSRISAKUL : MICROCHIP CAPILLARY ELECTROPHORESIS WITH AMPEROMETRIC DETECTION FOR METAL ION ANALYSIS. THESIS ADVISOR : ASSOC.PROF. ORAWON CHAILAPAKUL, Ph.D., THESIS COADVISOR : LUXSANA DUBAS, Ph.D., 127 pp.

This thesis demonstrates the fast and simultaneous detection of prominent heavy metals including lead, cadmium and copper using a microchip capillary electrophoresis with electrochemical detection. Direct amperometric detection mode for microchip capillary electrophoresis was successfully applied to analytes, the heavy metal ions. The influences of the separation voltage, detection potential, concentration and pH value of running buffer on the response of the detector were carefully investigated and optimized. The zone electrophoretic separation of lead, cadmium and copper is less than 3 min using a MES buffer and L-Histidine as background electrolyte (pH 7.0, 25 mM), employing 1.2 kV as the separation voltage and -0.8 V as the detection potential. The detection limits for Pb²⁺, Cd²⁺, and Cu²⁺ were 1.74, 0.73 and 0.13 μ M (S/N = 3), respectively. The %RSD of peak current was < 6 % and the %RSD of migration times <2% for prolong operation. To demonstrate the potential and future role of microchip CE, a new route in the real sample analysis was presented. The results obtained allow the proposed microchip capillary electrophoresis-Electrochemical detection as a real gateway to microanalysis in foods.

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ABBREVIATIONS AND SYMBOLS

i current (A) anodic peak current (A) i_{pa} cathodic peak current (A) ipc -Ep peak potential (V) - E_{pa} anodic peak potential (V) -E_{pc} cathodic peak potential (V) -Faraday constant (96,484.6 C equiv⁻¹) F _ area of electrode (cm^2) Α diffusion coefficient ($cm^2 s^{-1}$) D kinematic viscosity of the liquid $(cm^2 s^{-1})$ ν scan rate (V sec^{-1}) υ ~ angular velocity of the disk (radians per second) ω -С solution concentration (mol dm^{-3}) part per million ppm part per billion ppb mL milliter microliter μL g gram microgram μg μA microamp nA nanoamp μm micrometer micromolar μM nm nanometer i.d. internal diameter r^2 _ correlation coefficient MES -2-Morpholinoethanesulfonic acid L-his (s)-2-Amiono-3-(4-imidazyl)propionic acid -V volt