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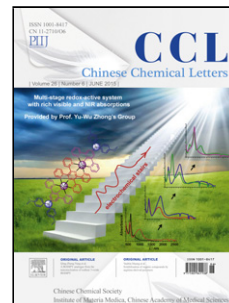
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Original article

Simple and cost-effective determination of ciprofloxacin hydrochloride by electrical micro-titration

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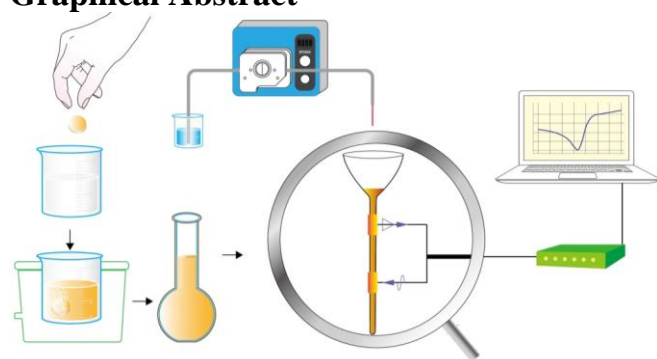
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Graphical Abstract



By employing an electrical micro-titration system, in which a capacitively coupled contactless conductivity detector is used to monitor the reaction process in real time, ciprofloxacin hydrochloride in tablet samples is determined. Because the reaction solutions are isolated from the working electrodes, it shows significant simplicity and cost-effectiveness, by eliminating the requirements for detector exchange/renewal between measurements, and by involving no auxiliary chemicals.

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ABSTRACT

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By employing an electrical micro-titration system, in which a capacitively coupled contactless conductivity detector (C^4D) was used to monitor the reaction process in real time, herein a novel method for determining ciprofloxacin hydrochloride (CIPHC) was developed for the first time. Mode 1: Standard CIPHC solutions at different concentrations were loaded into reaction cells, respectively, and were titrated with standard Ag^+ . Upon the titration, the formation of a precipitate alters the number of ions in the solution, raising the change of conductivity, which was monitored by a special C^4D to construct a titration curve. The endpoint of the titration was located from the peak of the curve. Between the elapsed time and the initial concentration of titrand, a linear relationship was established over the range of 2.0-8.0 mmol/L. Mode 2: Standard Fe^{3+} took the place of Ag^+ , and was used as titrant to recognize ciprofloxacin contributed to the formation of complexation, which also resulting a change of solution conductivity. Under optimized conditions, a working range of 1.0-5.0 mmol/L CIPHC was found. Because the reaction solutions were isolated from the working electrodes, this pioneer work shows significant simplicity and cost-effectiveness, by eliminating the requirements for detector exchange/renewal between different measurements, and by involving no auxiliary chemicals. Both of the two approaches were applied successfully to determine CIPHC in tablet samples. And the results were in good agreement with those obtained by reference method.

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