Determination of L-Cystine in Tablets Using the Chemiluminescence Method

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Aim. To develop the method for the quantitative determination of L-cystine in sublingual tablets Elthacin™ using the method of chemiluminescence inhibition in the $H_2L$ ($l$uminol) – $H_2O_2$ – $Hb$ (hemoglobin) system.

Materials and methods. The study objects were the pure substance of L-cystine of pharmaceutical grade and sublingual tablets Elthacin™ produced by the Medical Scientific-Production Complex “Biotics Ltd” (Russia) containing 70 mg of glycine, L-glutaminic acid and L-cystine in their composition. The intensity of chemiluminescence was measured on the device with a FEU-84-A photoelectric multiplier using an IMT-0.5 measuring instrument of low currents and a quick-acting automatic potentiometer.

Results and discussion. The method of the L-cystine quantitative determination in tablets based on the inhibition of chemiluminescence in the $H_2L$ – $H_2O_2$ – $Hb$ system has been developed. The calibration graph was linear over the concentration range from 7 to 70 μg · mL$^{-1}$. No interferences were observed in the presence of common components of the tablets, such as glycine and L-glutaminic acid. RSD = ±2.35 %, (δ = +1.13 %), LOD (3S) = 4 μg · mL$^{-1}$, LOQ (10S) = 13 μg · mL$^{-1}$ for the sublingual tablets Elthacin™.

Conclusions. The method proposed is promising for further research on the subject of its application for the determination of L-cystine in drugs.

Key words: L-cystine; inhibition; luminol; chemiluminescence; hemoglobin

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Визначення Л-цистину з використанням методу хемілюмінесценції

Мета. Метою даної роботи було розроблення методики кількісного визначення L-цистину в сублінгвахних таблетках Елтацин® за ефектом інгібування хемілюмінесценції системи $H_2L$ ($l$умінол) – $H_2O_2$ – $Hb$ (гемоглобін).

Матеріали та методи. Об’єктом дослідження були пігулки під’язикові Елтацин® виробництва ТОВ Медичний науково-виробничий комплекс “Біотики” (Росія) складу: гліцину, кислоти L-глутамінової та L-цистину у лікарських препаратах.

Результати та їх обговорення. Описано методику та показана можливість кількісного визначення L-цистину в пігулках під’язкових Елтацин® по 70 мг методом інгібування хемілюмінесценції системи $H_2L$ – $H_2O_2$ – $Hb$. Калібрувальний графік лінійний в діапазоні концентрацій від 7 до 70 мкг · мл$^{-1}$. Присутність гліцину та L-глутамінової кислоти не чинала жодного впливу на інтенсивність хемілюмінесценції. RSD = ± 2.35 %, (δ = + 1.13 %), LOD (3S) = 4 мкг · мл$^{-1}$, LOQ (10S) = 13 мкг · мл$^{-1}$ для сублінгвахних таблеток Елтацин®.

Висновки. Запропонований метод може бути використаний для визначення L-цистину у лікарських препаратах.

Ключові слова: L-цистин; інгібування; люмінол; хемілюмінесценція; гемоглобін

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Определение L-цистина в таблетках с использованием метода хемилюминесценции

Цель. Целью данной работы была разработана методика количественного определения L-цистина в сублингвахальных таблетках Элтацин® по эффекту ингибирования хемилюминесценции системы $H_2L$ (люминол) – $H_2O_2$ – $Hb$ (гемоглобин).

Материалы и методы. Объектом исследования были сублингвальные таблетки Элтацин® производства ООО Медицинский научно-производственный комплекс "Биотики" (Россия) состава: глутаминовая кислота и L-цистин по 70 мг и субстанция - L-цистину по эффекту ингибирования хемилюминесценции. RSD = ±2.35 %, (δ = + 1.13 %), LOD (3S) = 4 мкг · мл$^{-1}$, LOQ (10S) = 13 мкг · мл$^{-1}$ для сублингвальных таблеток Элтацин®.

Выводы. Предложенный метод может быть использован для определения L-цистина в препаратах.

Ключевые слова: L-цистин; ингибирование; люминол; хемилюминесценция; гемоглобин
Introduction. L-Cystine (3,3'-dithio-bis-2-amino-propionic acid, dicystein) is one of the well-known substituted of α-aminoacids, a stable (oxidized) form of the amino-acid of cysteine. It participates in the formation of peptides and other proteins during the formation of their structure. L-cystine provides elasticity of keratin; therefore, it is a part of complexes of vitamins for improvement of appearance of the skin and hair, biologically active additives and shampoos.

A wide range of medicinal products based on L-cystine can be used in cases of intoxication with heavy metals, as well as antioxidants, hepatoprotectors, detoxicants, immunomodulators, mucolytics, etc. Often L-cystine is also used as part of the complex therapy for the treatment of diabetes, Alzheimer’s disease, bronchitis, as well as rehabilitation after surgery and joint diseases.

The drug is available in the form of capsules and ampoules, and is also a part of the combined medicaments, such as the tablets Elthacin® (the main composition: L-cystine, 70 mg; glycine, 70 mg; L-glutaminic acid, fillers: methylcellulose, 7.8 mg; magnesium stearate, 2.2 mg).

Several methods, such as those based on isotopic titration using diperoxoyacidic acid, voltamperometry, polarography, capillary electrophoresis method, flow-injection analysis after its reduction to cysteine with the subsequent spectrophotometric detection, thin-layer chromatography, liquid chromatography in combination with mass-spectrometry and also using Raman microscopy combined chemometrics of the PCA (Principal Component Analysis) and HCA (Hierarchical Cluster Analysis) have been reported for its determination in different ranges of concentrations [1-9].

The advantages of the iodometric determination using diperoxoyacid as a reagent are selectivity: other amino acids being the constituents of the combined dosage forms and excipients do not interfere the L-cystine determination, but there is a need for the synthesis of the analytical reagent [1]. All these methods require specialized equipment and expertise and can be laborious for the routine analysis. The European Pharmacopoeia (EPh 8.0) recommends the method of inverse bromatometry for the quantitative determination of L-cystine. The sensitivity of the recommended method is limited by the relatively high concentration of the titrant (0.1 mol · L⁻¹) [10, 11].

The aim of our work was to develop a selective, sufficiently sensitive and easy-to-perform procedure for the routine analysis of L-cystine in the drug Elthacin®.

Materials and methods. The study objects were the pure substance of L-cystine of pharmaceutical grade (the content of the main substance was 98.5 %) and sublingual tablets Elthacin® produced by the Medical Scientific-Production Complex “Biotics Ltd” (Russia) containing 70 mg of glycine, L-glutaminic acid and L-cystine in their composition [10].

Solutions were prepared by the volume-weight method at 293 K. Double distilled water (DDW) was used to prepare the solutions in all cases.

Standard 1 · 10⁻³ mol · L⁻¹ solution of Luminol (5-amino-2.3-dihydro-1.4-flialazindion, H₂L, RPF Synbias, Ukraine) was used. 0.1772 g of 3-Aminophthaldehydezide with qualification “R” was transferred in a 100 mL volumetric flask, dissolved in 10 mL of 0.01 mol · L⁻¹ sodium hydroxide solution and diluted to the volume with DDW. The solution was kept in a dark place.

0.1 mol · L⁻¹ solution of sodium hydroxide was used for the medium pH stabilization, the solution pH was controlled by an Ionomer I-130 laboratory potentiometer with an ESL-43-07 glass electrode and silver-chloride electrode and a laboratory Ionomer I-130 (ZIP, Gomel, Belarus).

Hydrogen peroxide (H₂O₂) 5.8 % (wt.) solution was prepared from 58 % high pure preparation (produced by LTD Inter-Syntes, Boryslav, Ukraine) by its 10 time dilution with DDW: 10 mL was transferred into a 100 mL volumetric flask and diluted to the volume at 293 K. This solution was stored at reduced temperature of +8-10 °C. The content of hydrogen peroxide in solution was controlled by permanganometric titration.

The working solution of H₂O₂ 0.058 % (wt.) (1.7 · 10⁻² mol · L⁻¹) was obtained by the appropriate dilution of the original solution exactly 100 times. The working solution can be used throughout the day.

Human blood hemoglobin (Hb) produced by Simko Ltd, Lviv, Ukraine was used as a catalyst. Hemoglobin solution 100 μg · mL⁻¹ was prepared by dissolving in a 100 mL volumetric flask of 10 mg hemoglobin in 50 mL DDW by heating to 313…323 K and adding 1 mL of 1.0 mol · L⁻¹ sodium hydroxide solution. It was diluted to the volume with DDW at 293 K and stirred. The working solution of hemoglobin was prepared by dilution with DDW of the initial one exactly 100 times. The working solution can be used throughout the day.

The intensity of chemiluminescence was measured in relative units on the device with a FEU-84-A photonic multiplier using an IMT-0.5 measuring instrument of low currents and a quick-acting (time constant 0.1 s) automatic potentiometer. The reaction accompanying CL was performed in a cylindrical quartz cell of 30 mm in diameter with the working volume of 10 mL. The following order of mixing reagents was performed: to the mixture of luminol indicator in the alkali solution and H₂O₂, with the presence or absence of L-cystine solution in the control experiment, 0.50 mL of Hb was added with the help of a dosage pipette P-1, and the kinetic curve of chemiluminescence intensity (I) in relative units (I – time (s)) was registered. The dosage pipette is built in to the mobile keeper that isolates photocathode of a photoelectric multiplier from outside light, and further allows working at the common lighting. All experiments were performed at 293 K.

For the optimal conditions for the chemiluminescence occurrence in the H₂L – H₂O₂ – Hb analytical system the conditions determined earlier as a result of detailed studies by one of the authors were used. They are given in the monograph [12].

For the analytical signal, the value ΔI (rel. un.) was chosen. It is the difference between the maximum values of the CL intensities in the control (in the absence of an inhibitor) and working experiments.
In the course of studies, it was found that under the optimum conditions \((c(\text{NaOH}) = 0.05 \text{ mol} \cdot \text{L}^{-1}, c(H_2O_2) = 8.53 \cdot 10^{-4} \text{ mol} \cdot \text{L}^{-1}, c(H_2L) = 10^{-4} \text{ mol} \cdot \text{L}^{-1}, C(Hb) = 5 \cdot 10^{-2} \mu\text{g mL}^{-1})\) L-cystine exhibited the inhibitory effect on the intensity of the CL in \(H_2L - H_2O_2 - Hb\) system [12]. This phenomenon was used by us to develop a new method for the quantitative determination of L-cystine in solutions of tablets Elthacin®.

The standard solution sample (SSS) of L-cystine 0.7 μg · mL⁻¹ was prepared as follows: 0.0710 g of L-cystine was transferred in a 100 mL volumetric flask and dissolved in 10 mL of 1 mol · L⁻¹ sodium hydroxide solution and diluted to the volume with DDW. The working solution of L-cystine was obtained by the appropriate dilution of the original solution in the required number of times for analysis.

**Procedure of determination.** Solutions were added to a chemiluminescent quartz cell consistently as follows: 0.50 mL of \(1 \cdot 10^{-3} \text{ mol} \cdot \text{L}^{-1} H_2L, 0.50 \text{ mL} \) of 1 mol · L⁻¹ sodium hydroxide solution, \((10 - x) \text{ mL}\) of DDW where \(x\) was the total volume of all reagents and samples, (mL), 0.50 mL of \(1.7 \cdot 10^{-2} \text{ mol} \cdot \text{L}^{-1} H_2O_2\) and 0.5 mL of a dilute working-standard solution of L-cystine. A cell with the mixture was placed in a chemiluminometer, and 0.5 mL of the working solution of hemoglobin with the concentration of 1 μg·mL⁻¹ was added.

Similarly, the control experiment with DDW instead of the use of dilute working-standard solution of L-cystine in the same amount was performed. The content of L-cystine was calculated by Equation 1.

**Procedure of the quantitative determination of L-cystine in tablets Elthacin®.** Approximately 0.22 g of powdered tablets (accurately weighed) was transferred into a 100 mL volumetric flask, dissolved in 10 mL of 1 mol·L⁻¹ sodium hydroxide solution and diluted to the volume with DDW. Similarly, L-cystine SSS was prepared by the volume-weighted method with the concentration of 0.710 μg · mL⁻¹. L-cystine working solutions were prepared by the appropriate dilution immediately before the analysis.

When performing the experiment a certain order of adding solutions was observed in accordance with the above mentioned procedure.

The content of L-cystine in \(X\) (g to one tablet) was calculated by the Equation:

\[
X = \frac{m_o \times (\Delta I) \times \bar{m} \times w}{(\Delta I_r) \times m_r \times 100 \%},
\]

where: \(m_o\) – is the weight of L-cystine in SSS, g; \((\Delta I)\) – is the maximum value of \(\Delta I\) in the working experiment, relative units; \((\Delta I_r)\) – is the maximum value of \(\Delta I\) in SSS, relative units; \(\bar{m}\) – is the average tablet weight \((n = 20), \text{g}\); \(m_r\) – is the mass of pounded tablets in the batches used for analysis, g; \(w\) – is the content of the basic substance in the standard sample, %.

**Results and discussion.** The presence of L-cystine in the \(H_2L - H_2O_2 - Hb\) system leads to a decrease in the maximum intensity of CL, indicating inhibition of the CL reaction. This effect increases with increasing of the inhibitor of the process concentration (Fig. 1).
cate the absence of a negative effect of excipients on the results of the determination of API in a dosage pharmaceutical form.

The results of these preliminary studies allowed us to perform the analysis of the tablets using a simpler standard method. These results of the quantitative determination of L-cystine in tablets Elthacin® 70 mg are given in Tab. 2.

The analysis of the data showed that the means relative standard deviation did not exceed the permissible value. Since δ ≤ RSD, the results of the determination of L-cystine were considered to be correct (systematic error is absent). The sensitivity for the L-cystine quantitative determination was calculated: LOD (3S) = 4 μg · mL⁻¹, LOQ (10S) = 13 μg · mL⁻¹.

Conclusions. The new method has been developed, and the possibility for the quantitative determination of L-cystine in sublingual tablets Elthacin®, 70 mg, using the method of chemiluminescence inhibition in the H₂L–H₂O₂–Hb system has been shown. RSD = ± 2.35 %, (δ = ± 1.13 %), LOD (3S) = 4 μg · mL⁻¹, LOQ (10S) = 13 μg · mL⁻¹.

Conflict of interests: authors have no conflict of interests to declare.
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