SPECTROPHOTOMETRIC DETERMINATION OF PIZOTEFIN MALEATE IN PURE FORM AND TABLETS USING $\pi$-ACCEPTORS

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ABSTRACT. Two simple, quick and sensitive methods were described for the spectrophotometric determination of pizotefin maleate (PZT) either in pure form or pharmaceutical form (tablets). The methods were based on the reaction of pizotefin maleate as ‘n’ electron donor with chloranilic acid ($p$-CLA) and 7,7,8,8-tetracyanoquinodimethane (TCNQ) as $\pi$-acceptors to give highly coloured radical anion species which were quantitated spectrophotometrically at 533 and 839 nm for chloranilic acid and 7,7,8,8-tetracyanoquinodimethane, respectively. Different experimental conditions for the two methods were optimized. Beer’s law was obeyed in the concentration ranges of 10-160 and 2.0-20 $\mu$g mL$^{-1}$ PZT for $p$-CLA and TCNQ, respectively with good correlation coefficient of $\geq$0.9993 and a relative standard deviation (RSD) of $\leq$1.31%. The molar absorptivity, Sandell sensitivity, detection and quantification limits were also reported. The proposed methods were successfully applied for determination of PZT in tablets with good accuracy and precision and without interferences from common additives by applying the standard addition technique. The developed methods have been validated statistically for their accuracy, precision, sensitivity, selectivity, robustness and ruggedness as per ICH guidelines and the results compared favourably with those obtained using the reported method.

KEY WORDS: Pizotefin maleate, Charge transfer complexes, Spectrophotometry, Tablets

INTRODUCTION

Pizotifen maleate (PZT) is a sedating antihistamine that has strong serotonin antagonist and weak antimuscarinic properties. It is chemically known as 9,10-dihydro-4-(1-methylpiperidin-4-yldene)-4H-benzo[4,5]cyclohepta[1,2-b]thiophene hydrogen maleate (Figure 1) [1]. It also antagonises the action of tryptamine. Pizotifen is used, usually as the maleate, for the prophylaxis of migraine and for the prevention of headache attacks during cluster periods. It is not effective in treating an acute attack [2]. The drug is official in British Pharmacopoeia [1], which describes a potentiometric titration with perchloric acid in anhydrous acetic acid medium.

Several analytical methods have been reported for the determination of PZT in pure drug, pharmaceutical dosage forms and biological samples using high performance liquid chromatography [3–7], spectrofluorimetry [8] and two spectrophotometric methods [8, 9]. All the above reported methods developed for the determination of PZT employed complex analytical techniques, complicated, expensive, require careful control of conditions or skilled personnel and are inaccessible to many laboratories. In contrast, visible spectrophotometry is considered as the most convenient analytical technique in most quality control and clinical laboratories. Spectrophotometric methods have several advantages, such as low interference level, good analytical selectivity, and they are easier, less expensive, and less time consuming compared with most of the other methods.

The molecular interactions between electron donors and acceptors are generally associated with the formation of intensely colored charge transfer complexes, which absorb radiation in the visible region [10]. A variety of electron donating compounds have been reported to yield charge-transfer complexes with various $\pi$-acceptors [11-14].
The aim of the present work was directed to investigate direct, sensitive, simple and precise spectrophotometric methods for determination of PZT as a good n-electron donor via charge transfer complexation with two π-acceptors: chloranilic acid (p-CLA) and 7,7,8,8-tetracyanoquinodimethane (TCNQ) as chromogenic reagents in pure form and its dosage forms (tablets). The reaction conditions of the methods have been established. In addition, the stoichiometric ratio of reactants was determined. No interference was observed in the determination of PZT from common excipients in levels found in dosage forms. These methods are validated by the statistical data.

**EXPERIMENTAL**

**Apparatus**

All the absorption spectra were made using double beam Unikon 930 spectrophotometer (Kontron Instruments, Munchen, Germany) with a scanning speed of 200 nm/min and a band width of 2.0 nm, equipped with 10 mm matched quartz cells.

**Materials and reagents**

Pharmaceutical grade pizotefin maleate (PZT) was received from (Novartis, Egypt). Mosegor tablets, labeled to contain (0.5 mg PZT per tablet) were purchased from local commercial markets.

**Standard PZT solution**

A stock standard solution of PZT (500 µg mL⁻¹) was prepared by dissolving an exact weight (50 mg) of pure PZT in about 10 mL methanol and the volume was diluted to the mark in a 100 mL calibrated flask with acetonitrile. Stock solutions of PZT (100 µg mL⁻¹) and (5.0 x 10⁻³ M) were prepared from suitable dilution of the stock standard solution. The stock solutions of PZT are stable for a period of 3.0 days when kept in the refrigerator.

**Reagents**

All the reagents and chemicals used were of analytical or pharmaceutical grade and all solutions were prepared fresh daily. Chloranilic acid (p-CLA) (Fluka, Switzerland) and 7,7,8,8-tetracyanoquinodimethane (TCNQ) (Aldrich Chem. Co., Milwaukee, USA) and stock solution (1.0 x 10⁻³ M) of each reagent were freshly prepared in acetonitrile. The solutions were stable for at least one week at 4 °C and kept in the dark when not use.
Recommended procedures

Two sets of experiments were carried out. In first set, aliquots (0.2-3.2 mL) of standard PZT (500 µg mL\(^{-1}\)) solution were accurately transferred into 10-mL calibrated flasks in case of using \(p\)-CLA; whereas aliquots of (0.1-1.0 mL) of standard PZT (200 µg mL\(^{-1}\)) solution were transferred into 10-mL calibrated flasks in the second set in case of using TCNQ; then added to each flask 2.0 and 1.0 mL of (1.0 x 10\(^{-3}\) M) \(p\)-CLA and TCNQ reagents in the first and second set, respectively. The reaction mixtures were heated in a water-bath at 60±5 ºC for 10 min. The mixtures were cooled and then diluted to volume up to 10 mL with acetonitrile and the absorbance was measured at 533 and 839 nm for \(p\)-CLA and TCNQ, respectively, against a reagent blanks prepared in the same manner.

Procedure for pharmaceutical formulations (tablets)

The contents of twenty Mosegor tablets (0.5 mg PZT per tablet) were crushed and finely powdered. An accurate weight equivalent to 5.0 mg PZT were transferred into a 100-mL calibrated flask, dissolved in least volume of methanol with shaking for 5.0 min and filtered through a sintered glass crucible (G\(_4\)) to remove excipient in the powdered tablets. The filtrate was diluted to 25 mL with acetonitrile in a 25 mL measuring flask to give 200 µg mL\(^{-1}\) stock solution of PZT. For the proposed methods, the content of tablets was calculated using the corresponding regression equation of the appropriate calibration graph. The method of standard addition was used for the accurate determination of PZT contents.

Stoichiometric relationship

Job's method of continuous variations [15] of equimolar solutions was employed to establish the stoichiometry of the coloured products. Standard solution of PZT (1.0 x 10\(^{-3}\) mol L\(^{-1}\)) and (1.0 x 10\(^{-3}\) mol L\(^{-1}\)) solution of \(p\)-CLA or TCNQ were prepared. A series of solutions was prepared in which the total volume of PZT and reagent was kept constant (2.0 mL). The solutions were mixed in various proportions and the volume was diluted in a 10-mL volumetric flask with acetonitrile. The absorbance of the resulting solutions was measured at optimum wavelengths (\(\lambda_{\text{max}}\)) at best time and temperature against the reagent blank under the same conditions.

RESULTS AND DISCUSSION

This is the first report on the application of charge-transfer complexation reaction for spectrophotometric determination of PZT in pure form and tablets. The methods are based on well-characterized charge-transfer complex formation of PZT as electron-donor with selected \(\pi\)-acceptors (\(p\)-CLA or TCNQ) in acetonitrile and have the advantages of simplicity, rapid, accuracy and precision, and use of inexpensive equipment compared to the reported HPLC methods.

Absorption spectra

The reaction of \(p\)-CLA and TCNQ as \(\pi\)-acceptor with PZT as n-electron donor results in the formation of an intense orange-red and bluish-green colored radical anions (\(p\)-CLA\(^-\) and TCNQ\(^-\)) in acetonitrile, which exhibits strong absorption maxima at wavelengths 533 and 839 nm, respectively (Figures 2 and 3), most probably due to the formation of charge-transfer complexes between PZT acting as n-donor (D) or Lewis base, and \(p\)-CLA or TCNQ, as \(\pi\)-acceptors(A) or Lewis acids [10]:

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The dissociation of DA complex is promoted by the high dielectric constant of acetonitrile (37.5). Further support for the assignment was provided by the comparison of the absorption bands with those of the \( p \)-CLA\(^{-}\) and TCNQ\(^{-}\) radical anions produced by the iodide reduction method [16].

Figure 2. Absorption spectra of reaction products of 160 \( \mu g \text{ mL}^{-1} \) PZT with \( p \)-CLA \( (1.0 \times 10^{-3} \text{ M}) \) against blank solution.

Figure 3. Absorption spectra of reaction products of 30 \( \mu g \text{ mL}^{-1} \) PZT with TCNQ \( (1.0 \times 10^{-3} \text{ M}) \) against blank solution.
Optimization of reaction conditions

Effect of solvents

In order to select the suitable solvent for charge transfer complex formation, the reaction of PZT with p-CLA and TCNQ was carried out in different solvents such as acetone, methanol, ethanol, dichloromethane, 1,2-dichloroethane, acetonitrile and chloroform. Acetonitrile was found to be an ideal solvent for all the reagents, because it afforded the maximum sensitivity when compared with all other solvents and it possesses the highest dielectric constant of all solvents examined, a property which is known to promote the dissociation of the original charge-transfer complexes to the CLA\(^-\) and TCNQ\(^-\) radical anions (Figure 4).

![Absorption as a function of solvent](image)

Figure 4. Effect of solvent on the absorbance of charge transfer complex of PZT with p-CLA and TCNQ acceptors.

Effect of reagent concentration

The effect of the reagent concentration on the intensity of the color developed at the selected wavelengths was ascertained by adding different volumes of p-CLA and TCNQ reagents to fixed concentrations of PZT. It was found that 2.0 and 1.0 mL of \((1.0 \times 10^{-3}\) M\) p-CLA and TCNQ solution, respectively, were sufficient for the production of maximum and reproducible color intensity and the highest absorbance remained unaffected by further addition of both reagents (Figure 5).

Effect of reaction time and temperature

The optimum reaction time was determined by following the colour development at room temperature \((25 \pm 2\) °C). Complete colour development was attained after 10 min on raising the temperature on a water-bath to 60±5 °C using both p-CLA and TCNQ methods. The absorbance of these radical anions remained stable for at least 4.0 h for p-CLA and TCNQ methods.
Figure 5. Effect of reagent concentration on the absorbance of charge transfer complexes formed between PZT and \((1.0 \times 10^{-3} \text{ M}) p\)-CLA and TCNQ acceptors.

**Molar ratio of the reaction**

The stoichiometric ratio was determined by Job’s method \([15]\) of continuous variation for the reaction between PZT and \(p\)-CLA and TCNQ reagents, which shows that the interaction occurs between an equimolar solution of the PZT and the reagents. The result indicated that the charge transfer complex was formed in the ratio of 1:1 (Figure 6). On the basis of the literature data and our experimental results, tentative reaction mechanisms for PZT-TCNQ complex is proposed and given in Scheme 1.

Figure 6. Continuous variation plots for the reaction of PZT with \(p\)-CLA and TCNQ. \(\lambda = 533\) and 839 nm, respectively. Total molar concentration = \(1.0 \times 10^{-4} \text{ mol L}^{-1}\).
Scheme 1. Proposed reaction pathway for the formation of charge transfer complex between PZT and TCNQ.

Method validation

Validation of the described methods for assay of bulk PZT was examined via linearity, sensitivity, precision, accuracy, repeatability, reproducibility, selectivity and robustness according to ICH guidelines [17].

Linearity

Under the optimum conditions a linear correlation was found between absorbance at $\lambda_{\text{max}}$ and concentration of PZT in the ranges of 10-160 $\mu$g mL$^{-1}$ and 2.0-20 $\mu$g mL$^{-1}$ using p-CLA and TCNQ method, respectively. The calibration graph is described by the regression equation:

$$A = a + b \, C$$

(1)

(where $A = \text{absorbance}$, $a = \text{intercept}$, $b = \text{slope}$ and $C = \text{concentration in } \mu\text{g mL}^{-1}$) obtained by the method of least squares [18]. Correlation coefficient, intercept and slope for the calibration data are summarized in Table 1. For accurate determination, Ringbom concentration range [19] was calculated by plotting log concentration of drug in $\mu$g mL$^{-1}$ against transmittance % from which the linear portion of the curve gives accurate range of micro determination of PZT and TCNQ.
The linearity of the calibration graphs was demonstrated by the high values of the correlation coefficient ($r$) and the small values of the y-intercepts of the regression equations.

Sensitivity parameters such as apparent molar absorptivity ($\varepsilon$) and Sandell’s sensitivity ($S_s$) values, the limits of detection and quantification were calculated as per the current ICH guidelines (ICH) [17], are illustrated in Table 1. The high molar absorptivity and lower Sandell sensitivity values reflect the good and high sensitivity of the proposed methods. The validity of the proposed methods was evaluated by statistical analysis [20] between the results achieved from the proposed methods and that of the reported methods. Regarding the calculated Student’s $t$-test and variance ratio $F$-test (Table 1), there is no significant difference between the proposed and reported method [8] regarding accuracy and precision.

Table 1. Analytical and regression parameters of proposed oxidation spectrophotometric methods for determination of PZT.

| Parameters                          | $p$-CLA | TCNQ |
|-------------------------------------|---------|------|
| **Wavelengths $\lambda_{\text{max}}$ (nm)** | 533     | 839  |
| **Color stability**                 | 4.0     | 3.0  |
| **Beer’s law limits (µg mL$^{-1}$)** | 10-160  | 2.0-20 |
| **Ringbom optimum concentration range (µg mL$^{-1}$)** | 20-140 | 4.0-18 |
| **Molar absorptivity $\varepsilon$, (L/mol cm$^{-1}$) x 10$^4$** | 0.1044 | 0.6145 |
| **Sandell’s sensitivity (ng cm$^{-2}$)** | 282.95 | 48.07 |
| **Regression equation**             |         |      |
| Slope ($b$)                         | 0.0037  | 0.0199 |
| Intercept ($a$)                     | -0.0033 | 0.0033 |
| Correlation coefficient ($r$)       | 0.9993  | 0.9994 |
| Mean ± SD                           | 100.20 ± 1.31 | 99.40 ± 0.96 |
| Relative standard deviation, RSD%   | 1.31    | 0.96  |
| Relative error, RE%                | 1.37    | 1.01  |
| LOD, (µg mL$^{-1}$)                 | 2.83    | 0.54  |
| LOQ, (µg mL$^{-1}$)                 | 9.43    | 1.80  |
| Calculated $t$-value               | 0.75    | 0.47  |
| Calculated $F$-value               | 2.27    | 1.22  |

$A = a + bC$, where $C$ is the concentration in µg mL$^{-1}$, $A$ is the absorbance units, $a$ is the intercept, $b$ is the slope.

Accuracy and precision

The accuracy and precision of the proposed methods (within-day and between-days) were evaluated by performing six replicate analyses on pure drug solution at three different

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concentrations of PZT (within the working range). Percentage relative standard deviation (% RSD) as precision and percentage relative error (% RE) as accuracy of the proposed spectrophotometric methods were calculated. The relative standard deviation (RSD) values were \( \leq 1.1\% \) in all cases, indicating good repeatability of the suggested methods. This level of precision of the proposed methods was adequate for the quality control analysis of the studied drugs. % RE, an indicator of accuracy, is in the range (-0.7-0.6) also indicating high accuracy and repeatability of the methods. The intra-day and inter-day precision and accuracy results show that the proposed methods have good repeatability and reproducibility reflecting the usefulness of the methods in routine analysis (Table 2).

Table 2. Evaluation of intra-day and inter-day accuracy and precision for PZT obtained by the proposed methods.

| Methods | Added (μg mL\(^{-1}\)) | Recovery % | Precision RSD %\(^a\) | Accuracy RE % | Confidence limit\(^b\) |
|---------|---------------------|------------|------------------------|--------------|----------------------|
|         | Intra-day            |            |                        |              |                      |
| p-CLA   | 40                  | 99.6       | 0.57                   | -0.40        | 39.9 ± 0.24          |
|         | 80                  | 99.9       | 0.91                   | -0.10        | 79.9 ± 0.76          |
|         | 160                 | 99.3       | 1.08                   | -0.70        | 159 ± 1.80           |
| TCNQ    | 5.0                 | 99.4       | 0.64                   | -0.65        | 4.97 ±0.01           |
|         | 10                  | 100        | 0.86                   | 0.25         | 10.0±0.08            |
|         | 15                  | 99.9       | 1.07                   | -0.10        | 15.0±0.15            |
|         | Inter-day            |            |                        |              |                      |
| p-CLA   | 40                  | 99.7       | 0.48                   | -0.30        | 39.9±0.20            |
|         | 80                  | 100        | 0.85                   | 0.10         | 80.1±0.71            |
|         | 160                 | 101        | 0.99                   | 0.60         | 161±1.67             |
|         | 5.0                 | 99.8       | 0.54                   | -0.20        | 4.99±0.03            |
|         | 10                  | 100        | 0.86                   | 0.25         | 10.0±0.08            |
|         | 15                  | 99.5       | 0.70                   | -0.55        | 9.95±0.07            |

\(^a\)Mean of six determination, RSD%, percentage relative standard deviation; RE%, percentage relative error. \(^b\)Mean ± standard error.

**Robustness and ruggedness**

The robustness of the methods was evaluated by making small incremental changes in the volume of reagent (± 0.2 mL) and time (±1.0 min), and the effect of the changes were studied on the absorbance of the charge transfer complex. The changes had negligible influence on the results as revealed by small intermediate precision values expressed as RSD% (≤ 2.60%). Method ruggedness was demonstrated having the analysis done by three analysts, and also by a single analyst performing analysis using three different spectrophotometer instruments in the same laboratory. Intermediate precision values (RSD%) in both instances were (≤ 2.0%) indicating acceptable ruggedness.

**Recovery studies by standard addition technique**

The accuracy and validity of the proposed methods were further ascertained by performing recovery studies. In this study, pre-analyzed tablet powder was spiked with pure PZT at different concentration levels and the total was determined by the proposed methods using standard addition technique. The percent recovery of pure PZT added was in the range 98.5–101% with relative standard deviation of 0.65–0.74 (Table 3) indicating that the recovery was good, and revealed that the co-formulated substances did not interfere in the determination. The results of recovery study are compiled in Table 3.

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Table 3. Application of the standard addition technique for the determination of PZT in Mosegor tablets using the proposed methods.

| Sample                 | Taken (µg mL⁻¹) | p-CLA Added (µg mL⁻¹) | Recovery (%) | Taken (µg mL⁻¹) | TCNQ Added (µg mL⁻¹) | Recovery (%) |
|------------------------|-----------------|-----------------------|--------------|-----------------|----------------------|--------------|
| Mosegor tablets        | 10              | -                     | 99.5         | 2.0             | -                    | 99.1         |
|                        | 10              | 10                    | 99.1         | 2.0             | 10                   | 99.5         |
|                        | 30              | 90                    | 98.5         | 8.0             | 30                   | 99.7         |
|                        | 50              | 95                    | 98.5         | 8.0             | 50                   | 99.7         |
|                        | 100             | 95                    | 99.5         | 12              | 100                  | 101          |
|                        | 150             | 101                   | 101          | 16              | 150                  | 98.6         |
| Mean ± SD              |                 | 99.4±0.7              |              | 99.5±0.7        |                      |              |
| RSD%                   |                 | 0.75                  |              | 0.65            |                      |              |
| V                      |                 | 0.55                  |              | 0.42            |                      |              |
| SE                     |                 | 0.30                  |              | 0.26            |                      |              |

*Average of six determinations.

Selectivity

The studied drug (PZT) was determined in the presence of possible excipients and additives such as lactose, microcrystalline cellulose, sodium starch glycolate and magnesium stearate. Under the experimental conditions employed, to a known amount of drug, excipients in different concentrations were added and studied. Excipients do not interfere with the assay indicating the high selectivity of the proposed methods and applicability to use for routine determination in pure and in dosage forms. In addition, recoveries in most cases were around 100% and the lower values of the RSD (≤ 2.0%) indicate the good precision of the methods.

Application of the proposed methods to analysis of tablets

In order to evaluate the analytical applicability of the proposed methods to the quantification of PZT in commercial tablets, the results obtained by the proposed methods were compared with those of the reference method [8] by means of Student’s t-test and F-test for accuracy and precision. The results (Table 4) show that the Student’s t- and F-values at 95% confidence level are less than the theoretical values, indicating that there is a good agreement between the results obtained by the proposed methods and the reference method with respect to accuracy and precision.

Table 4. Application of the proposed methods for the determination of PZT in Mosegor tablets using the proposed charge transfer methods.

| Samples         | Proposed methods | Reported method [8] |
|-----------------|------------------|---------------------|
|                 | p-CLA            | TCNQ                |
| Recovery ± RSDf | 99.4±0.7         | 99.5±0.7            |
| t-valueg        | 0.15             | 0.46                |
| F-valueh        | 1.51             | 1.96                |

*Average of six determinations. fThe theoretical values of t and F are 2.57 and 5.05, respectively, at confidence limit at 95% confidence level and five degrees of freedom (p = 0.05).

CONCLUSION

The present study describes the first report on the application of charge-transfer complexation reaction for spectrophotometric determination of PZT in pure form and tablets using two π
acceptors (p-CLA and TCNQ) as analytical reagents and validated as per the current ICH guidelines. The present spectrophotometric methods are characterized by simplicity since they do not involve any critical experimental variable and are free from tedious, time-consuming, extraction steps and do not need expensive sophisticated apparatus unlike many previous methods. The proposed methods have additional advantages of accuracy, precision, ease of operation and possibility of carrying them out with a common laboratory instrument unlike many other instrumental methods reported for PZT. The methods are characterized by high selectivity and comparable sensitivity with respect to the existing methods. The highlight of the proposed methods is their ability to quantify PZT in tablets. Thus, the methods are useful for the quality control and routine analysis of PZT in pharmaceuticals since there is no interference from the common excipients that might be found in commercial formulations.

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