Spectrophotometric Determination of Antidepressant Drug Duloxetine in Pharmaceutical Preparations using π-Acceptors

SIDIKA ERTÜRK Toker* and ARMAĞAN ÖNAL

Istanbul University, Faculty of Pharmacy
Department of Analytical Chemistry, 34116, Beyazit, Istanbul – Turkey
serturk@yahoo.com

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Abstract: In this study, two simple and accurate spectrophotometric methods were presented for the determination of duloxetine hydrochloride (DLX) in pharmaceutical preparations. The methods were based on the reaction of DLX as \( n \)-electron donor with 2,3-dichloro-5,6-dicyano-1,4-benzoquinone (DDQ) and 7,7,8,8-tetracyanoquinodimethane (TCNQ) as \( \pi \)-acceptors to give highly colored complex species. The colored products were quantitated spectrophotometrically at 477 and 841 for DDQ, TCNQ, respectively. All variables were studied in order to optimize the reaction conditions. Beer’s law was obeyed in the concentration ranges 10.0-50.0 and 15-60 \( \mu \)g mL\(^{-1}\) for DDQ and TCNQ method, respectively. The proposed methods have been successfully applied to the pharmaceutical analysis without any interference from excipients. The suggested procedures could be used for the determination of DLX in pharmaceutical preparations being sensitive, simple and selective.

Keywords: Duloxetine, Charge-transfer Reaction, DDQ, TCNQ, Pharmaceutical preparations

Introduction

Duloxetine hydrochloride (DLX) [(+)-(S)-N-methyl-3-(1-naphthalenylxoy)-2-thiophene-propanamine hydrochloride, Figure 1, acts as a dual serotonin and norepinephrine reuptake inhibitor. It has been clinically used for the treatment of major depressive disorder, with or without melancholic symptoms, pain due to diabetic peripheral neuropathy and urinary incontinence\(^{1-6}\).

To the best of our knowledge, only three papers can be found in the literature about determination of duloxetine at pharmaceutical preparations, which were based on UV spectrophotometric, high-performance thin layer chromatographic and high-performance liquid chromatographic analysis\(^{7,9}\). There is no visible spectrophotometric method available
for determination of DLX in tablets; therefore two visible spectrophotometric analyses were
developed for analysing of DLX in tablets. The methods were based on charge transfer reaction
with 2,3-dichloro-5,6-dicyano-1,4-benzoquinone (DDQ) and 7,7,8,8-tetracyanoquinodimethane
(TCNQ). A significant advantage of spectrophotometric methods unlike HPTLC or HPLC
methods, the instrument is simple and is not of high cost. The sensitivity in terms of molar
absorptivity and the precision of the proposed methods are better than UV spectrophotometric
method. Because of the wide applicability of proposed methods, these are suggested for routine
quality control assay of the drugs in pure form and its pharmaceutical preparations.

![Chemical structure of duloxetine](image)

**Figure 1.** Chemical structure of duloxetine

**Experimental**
Spectrophotometric measurements were carried out using a Shimadzu UV-160A
spectrophotometer with 1 cm glass cells.

**Reagents and solutions**
Duloxetine hydrochloride was kindly supplied by boehringer ingelheim pharma GmbH
&Co. KG (Germany). Its pharmaceutical preparation cymbalta capsul®, containing 60 mg
of duloxetine (DLX) per capsul and were obtained from local drugstore. All chemicals and
reagents were of analytical-reagent grade.

Stock solutions were prepared by dissolving equivalent 25 mg of duloxetine base in a
100 mL volumetric flask to give a concentration of 250 µg mL\(^{-1}\) of DLX. For this purpose,
10 mL aliquot of the stock solution was transferred to stoppered tube and 1 mL 1 N NaOH
solution was added. Three 5 mL portion of chloroform was added for the extraction. Combined
extracts were dried using anhydrous Na\(_2\)SO\(_4\) and diluted with 20 mL with chloroform. All of
this solution was evaporated to dryness under nitrogen with mild heating. The residue was
dissolved with acetonitrile (in 20 mL volumetric flask) using an ultrasonic bath. Acetonitrile
was added to the mark (125 µg mL\(^{-1}\) as the base).

2,3-Dichloro-5,6-dicyano-p-benzoquinone (DDQ) (Merck-Schuchardt, Munich, Germany)
was freshly prepared as (0.2%(w/v)) solutions in acetonitrile; the solution was stable for 1
week at 4 °C. TCNQ was obtained from Merck (Darmstadt, Germany). 0.2% TCNQ
solution was prepared in acetonitrile.

**General procedure**
Into 10 mL calibrated flasks were placed 0.80-4.0 mL and 1.2-4.8 mL aliquots of DLX
standart solution for DDQ and TCNQ method, respectively. The volume was bring to 4.0 mL
(for DDQ) and 4.8 mL, (for TCNQ) with acetonitrile and 3.5 mL of DDQ or 3 mL of TCNQ
reagent solution was added. The reaction mixture was heated for 20 min at 60 °C for DDQ
and allowed to stand for 5 min at room temperature for TCNQ method. After cooling and
diluting to volume 10 mL with acetonitrile the absorbance was measured at 477 and 841 nm
for DDQ and TCNQ, respectively, against a reagent blank prepared in the same manner. The
calibration graph was prepared by plotting absorbance vs. concentration of pregabalin.
**Assay procedure for pharmaceutical preparations**

The contents of 20 capsules were mixed and accurately weighed amount of the contents equivalent to 10 mg of DLX was transferred into a 100 mL volumetric flask. About 50 mL of water was added and then extraction was performed mechanically for 20 minutes and sonicated for 20 more minutes. The volume was brought to 100 mL with water and final solution was filtered. Aliquots of filtrate were studied as described under preparation of General procedure. The nominal contents of the capsules were calculated using either the calibration graph or the corresponding regression equation.

**Results and Discussion**

As to our best knowledge, visible spectrophotometric determination of DLX in tablets has not been yet described. Therefore, two visible spectrophotometric analyses were developed for the determination of DLX in tablets. The developed methods are based on the reaction of DLX as n-electron donor with 2,3-dichloro-5,6-dicyano-1,4-benzoquinone (DDQ) and 7,7,8,8-tetracyanoquinodimethane (TCNQ) as a π-acceptor to give highly colored complex species. π-acceptors are known to yield charge transfer complexes and radical anions with a variety of electron donors\textsuperscript{10-15}.

The interaction of DLX with DDQ and TCNQ, in acetonitrile yielded intense orange red colored chromogen for DDQ, a bluish-green colored chromogen for TCNQ and absorbing maximally at wavelengths 477 and 841 nm, respectively (Figure 2). The influence of different parameters on the color development was studied to determine optimum conditions.

![Figure 2(A)](image1.png) Absorption spectrum of 50.0 μg mL\(^{-1}\) DLX complex with DDQ reagent against reagent blank. (B) Absorption spectrum of 60.0 μg mL\(^{-1}\) DLX complex with TCNQ reagent against reagent blank.
Choice of solvent

Different solvents included chloroform, acetonitrile, acetone, ethanol, 1,4-dioxan, methanol, and methylene chloride were investigated in order to select the suitable solvent. Acetonitrile is considered to be an ideal solvent for the colour reaction as it offers solvent capacity and gives the highest yield of the radical as indicated by high ε values.

Reagent concentration

The effect of DDQ and TCNQ concentration (by volume) on its reaction with the DLX were investigated. It is found that, when various concentrations (by volume) of DDQ and TCNQ solution added to a fixed concentration of DLX, 2.0 mL of TCNQ and 2.5 mL of DDQ solutions 0.2% (w/v) are found to be sufficient for quantitative determination of DLX (Figure 3).

Figure 3. Effect of volume of DDQ, (0.2%, w/v) and TCNQ (0.2%, w/v) on the development of the reaction product of DLX with these reagents at (▲) DDQ and (■) TCNQ.

Reaction time

The optimum reaction time was determined by following the color development at room temperature and 60-80 °C. Complete color development was attained after 5 min for TCNQ at room temperature, whereas for DDQ, complete color development was obtained after heating on a water bath at 60 °C for 20 min (Figure 4). The resultant complexes were stable up to 24 h at room temperature in the dark.

Figure 4. The effect of temperature and time of heating on the reaction between DDQ (at 60 °C) and TCNQ (at room temperature) reagents with DLX at (▲) DDQ, (♦) TCNQ.
Stoichiometry of the reaction

The molar ratio of DDQ and TCNQ to DLX in the reaction mixture was studied according to Job's method of continuous variation. Utilizing equimolar solution of DLX and DDQ and TCNQ, the reaction stoichiometry was found to be a good approximation 1:1 ratio (drug/reagent), confirming that one molecule of DLX reacts with one molecule of DDQ or TCNQ.

Method validation

Linear relationship was found with between the absorbance at $\lambda_{\text{max}}$ and the concentration of the drug in the ranges 10.0-50.0 and 15.0-60.0 µg mL$^{-1}$ for DDQ and TCNQ method, respectively. Molar absorptivity value of DDQ and TCNQ method was found as $1.60 \times 10^4$ and $1.16 \times 10^4$ L/mol.cm, respectively. The DDQ method is more sensitive compared with that of the TCNQ method, with a high $\varepsilon$ value. Sandell's sensitivity ($S$) represents the number of micrograms of the determinant per milliliter of a solution having an absorbance (A) of 0.001 for a path length (l) of 1 cm. Thus, $S = 10^{-3}/a = \mu$g cm$^{-2}$ where, $a$ is the specific absorptivity and its value (in mL g$^{-1}$ cm$^{-1}$) corresponds to the determinant in a cuvette with an optical length of 1 cm. Also, $a = (b/\text{molecular weight of duloxetine base}) \times 1000$, where $b =$ molar absorptivity = $A/CI$, where $C =$ the molar concentration of the determinant and $l = 1$ cm path length. Sandell sensitivity was found as 0.022 and 0.030 µg cm$^{-2}$ for DDQ and TCNQ method, respectively. The limits of detection (LOD) and limits of quantitation (LOQ) were determined using the formula: LOD or LOQ = $\kappa$SD$a/b$, where $k =$ 3 for LOD and 10 for LOQ. SD$a$ is the standard deviation of the intercept and $b$ is the slope. The inter- and intra-day precision were examined by analysis of drugs for same day and five consecutive days (each $n =$5). The RSD values for intra-day precision 0.56, 0.78% and inter-day precision 0.93, 1.07% were found for DDQ and TCNQ methods, respectively which were found indicating good precision.

The obtained results are summarized in Table 1.

Table 1. Results of validation parameters for proposed methods

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Using DDQ reagent</th>
<th>Using TCNQ reagent</th>
</tr>
</thead>
<tbody>
<tr>
<td>Linearity range$^a$, µg mL$^{-1}$</td>
<td>10.0-50.0</td>
<td>15.0-60.0</td>
</tr>
<tr>
<td>Intra-day$^b$, RSD %</td>
<td>0.56</td>
<td>0.78</td>
</tr>
<tr>
<td>Inter-day$^c$, RSD %</td>
<td>0.93</td>
<td>1.07</td>
</tr>
<tr>
<td>Regression equation$^d$</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Slope ± SD</td>
<td>0.018 ± 0.000216</td>
<td>0.0167 ± 0.0015</td>
</tr>
<tr>
<td>Intercept ± SD</td>
<td>-0.0836 ± 0.000911</td>
<td>-0.173 ± 0.00013</td>
</tr>
<tr>
<td>Correlation coefficient, $r$</td>
<td>0.9999</td>
<td>0.9994</td>
</tr>
<tr>
<td>LOD, µg mL$^{-1}$</td>
<td>0.199</td>
<td>0.294</td>
</tr>
<tr>
<td>LOQ, µg mL$^{-1}$</td>
<td>0.663</td>
<td>0.980</td>
</tr>
</tbody>
</table>

$^a$ Average of six determinations, $^b$n=5 correspond to replicate analysis for each level, $^c$Results of five different days, $^d$A = a + bC (where C is the concentration of drug in µg ml$^{-1}$, A is the absorbance at $\lambda_{\text{max}}$)

To check accuracy of the proposed methods, the standard addition technique was applied. A different amount of pure sample solution was added to three different concentrations of the standard drug solution and assayed. The percent recovery of the added standard to the assay samples was calculated from:

$$\text{Recovery} \% = \left(\frac{(C_t-C_a)}{C_a}\right) \times 100$$
Where $C_t$ is the total concentration of the analyte found; $C_u$ is the concentration of the analyte present in the formulation and $C_a$ is the concentration of the pure analyte added to the formulation. The results of analysis of the commercial dosage forms and the recovery study as shown in Table 2. The average percent recoveries obtained were quantitative (99.92-100.05%), indicating good accuracy of the methods.

Table 2. Results of recovery studies by standard addition method

<table>
<thead>
<tr>
<th>Method</th>
<th>Amount taken, µg mL$^{-1}$</th>
<th>Amount added, µg mL$^{-1}$</th>
<th>Total amount found$^b$ µg mL$^{-1}$ (Mean± S.D)$^c$</th>
<th>Recovery, %</th>
<th>RSD, %</th>
</tr>
</thead>
<tbody>
<tr>
<td>Using DDQ reagent</td>
<td>10</td>
<td>10.0</td>
<td>19.950±0.093</td>
<td>99.50</td>
<td>0.47</td>
</tr>
<tr>
<td></td>
<td>30.0</td>
<td>40.1422±0.26</td>
<td>99.47</td>
<td>100.47</td>
<td>0.65</td>
</tr>
<tr>
<td></td>
<td>40.0</td>
<td>49.910±0.512</td>
<td>99.78</td>
<td>100.08</td>
<td>0.72</td>
</tr>
<tr>
<td></td>
<td>15.0</td>
<td>25.012±0.18</td>
<td>99.83</td>
<td>100.25</td>
<td>0.53</td>
</tr>
<tr>
<td>Using TCNQ reagent</td>
<td>10</td>
<td>30.0</td>
<td>39.950±0.345</td>
<td>99.83</td>
<td>0.86</td>
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<tr>
<td></td>
<td>50.0</td>
<td>60.123±0.321</td>
<td>100.25</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

$^a$Cymbalta capsule®, containing 60 mg of duloxetine per tablets, $^b$Five independent analyses. $^c$Standard deviation

The applicability of the proposed method was tested by the determination of drugs in their pharmaceutical preparations. The results obtained are satisfactorily accurate and precise as indicated by the excellent % recovery and RSD<2 (Table 3). Experiments showed that there was no interference from the additions and excipients, e.g. lactose, glucose, fructose, magnesium stearate and starch.

Table 3. Analysis of DLX in capsules (n=5)

<table>
<thead>
<tr>
<th>Method 1$^a$</th>
<th>Using DDQ reagent</th>
<th>Using TCNQ reagent</th>
</tr>
</thead>
<tbody>
<tr>
<td>Mean± S.D</td>
<td>60.47±0.47</td>
<td>60.49±0.51</td>
</tr>
<tr>
<td>Recovery, %</td>
<td>100.78</td>
<td>100.82</td>
</tr>
<tr>
<td>RSD, %</td>
<td>0.78</td>
<td>0.85</td>
</tr>
</tbody>
</table>

$^a$Cymbalta capsule®, containing 60 mg of duloxetine per tablets, $^b$five independent analyses

Conclusion

Two simple and less time consuming visible spectrophotometric methods using DDQ and TCNQ reagents were described for analysing of DLX in tablets. DDQ method was the more sensitive than TCNQ method, with a high ε value (1.60×10$^4$ L/mol cm). The proposed visible spectrophotometric methods are simple, accurate and highly cost-effective and suitable for routine determination of the duloxetine in capsules.

References
