UV and two derivative spectrometric methods for determination of racedotril in tablet formulation

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ABSTRACT

UV, second and third derivative spectrophotometric methods have been developed for the determination of Racacdotril in pharmaceutical formulation. The solutions of standard and sample were prepared in Methanol. For the first method, UV spectrophotometry, the quantitative determination of the drug was carried at 231 nm and the linearity range was found to be 8-100ug/ml. For the second and third derivative spectrophotometric methods the drug was determined at 250nm and 240nm with linearity ranges for both 8-100ug/ml. The calibration graphs constructed at their wavelength of determination were found to be linear for UV and derivative spectrophotometric methods. All the proposed methods have been extensively validated. The described methods can be readily utilized for the analysis of pharmaceutical formulation.

Key words: UV spectrophotometry, derivative spectrophotometry, Racacdotril.

INTRODUCTION

Racacdotril is an effective and safe drug for acute diarrhea in adults and children. Chemically racacdotril is N-[(R,S)-3-acetylmercapto-2-benzylpropanoyl]-glycine benzy]ester. The drug controller General of India approved it as an antidiarrheal in October 2001. It is not yet official in any pharmacopoeia. A survey of literature survey reveals that a few HPLC and UV3,4,7 method were reported for the estimation of racacdotril in Pharmaceutical formulation and biological fluids2,5,6. In the present report the authors purpose a simple, sensitive and economical UV and two derivative spectrophotometric methods for the determination of Racacdotril in tablet formulation. The method is based on the measurement of light absorption in UV region in methanol.

EXPERIMENTAL

Pharmaceutical grade Racacdotril was a generous gift from Dr.Reddy’s laboratories Pvt.Ltd (Hyderabad, India). All analytical grade chemicals were purchased from Merck (India). Racacdotril sachets (sachactredotil) DR. Reddys laboratories Pvt. Ltd; Hyd, India) containing 10mg of Racacdotril per sachet was assayed. A shmadzu UV/VIS 1601 Spectrophotometer with data processing system was used. UV and derivative spectra of the solution were recorded in 1 cm quartz cells at a scan speed of 2800nm/min, a scan range of 200-400nm for UV, 221-350nm for second and third derivative.

Preparation of standard solutions

Stock standard solution was prepared by
dissolving 10 mg of Racecadotril in 10 ml methanol. The standard solutions were prepared by dilution of the stock solution with methanol in a concentration range of 8-100 µg/ml for UV, 2nd and 3rd derivative spectrophotometric methods, respectively.

**Market sample analysis**

A total of 10 sachets of Racecadotril were accurately weighed and powdered. An amount of sachet equivalent to 100 mg of Racecadotril was weighed and transferred in 100 ml calibrated volumetric flask, containing about 60 ml methanol. The contents were sonicated for 10-15 minutes with shaking to ensure the complete solubility of the drug and then volume made up with methanol. This solution was filtered to remove any insoluble matter. The filtrate was collected in a clean flask. Appropriate dilution were made to obtain with methanol from stock solution for UV and derivative spectrophotometric methods at 231, 250 and 240 nm respectively.

**Method validation**

**Linearity**

Under the experimental conditions described the graph obtained for UV, Second and third derivative spectra showed linear relationship (Fig 1, 2, 3). Regression analysis using the method of least squares was made for the slope, intercept and correlation co-efficient values. The regression equations of calibration curves were \( Y = 0.0093X + 0.0014, (R=0.9998) \) for the UV, \( Y = 0.0003X + 0.0018, (R=0.9998) \) for the Second, \( Y = 0.0001X - 0.000018(R=1) \) for Three derivative spectrophotometric methods, respectively (Fig 1A, 2A, 3A). The range was found to be 8-100 µg/ml for UV, second and third derivative spectrophotometric methods. The statistical parameters given are the regression equation calculated from the calibration graphs, along with standard deviation of the slope and intercept. The results are presented in Table 1.

| Table 1: Statistical data for calibration curves for determination of racecadotril |
|---------------------------------|-----------------|-----------------|-----------------|
| Parameters                      | UV              | Second Derivative | Third Derivative |
| Absorption Maxima(nm)           | 231             | 250              | 240             |
| Beer's law limits(ug/ml)        | 8-100           | 8-100            | 8-100           |
| Molar extinction co-efficient   | 0.009275        | -                | -               |
| Sandal's sensitivity            | 0.0927921       | -                | -               |
| Regression equation(y)          | 0.9998          | 0.9998           | 1               |
| Slope(b)                        | 0.0093          | 0.0003           | 0.0001          |
| Intercept(a)                    | 0.0014          | 0.0018           | 0.00018         |
| Co-efficient of variance        | 0.9111          | -                | -               |
| Standard deviation              | 0.00334         | 0.00953          | 0.00015         |
| LOD(ug/ml)                      | 0.8874          | 0.07874          | 0.04315         |
| LOQ(ug/ml)                      | 0.268938        | 0.19782          | 0.1262          |

| Table 2: Results of analysis of REDOTIL Sachets |
|-----------------------------------------------|---------------|-----------------|-----------------|
| Statistical value                             | UV            | Second derivative | Third derivative |
| X*                                            | 9.98          | 9.96            | 9.89            |
| SD                                            | 0.0033        | 0.0953          | 0.0673          |

* X* mean of six readings, SD is the Standard Deviation. REDOTIL are sachets containing 10 mg sof Racecadotril, n=6
Fig. 1: Zero-Order spectrum of RACE

Fig. 1(A): Zero-Order linearity plot of RACE

Fig. 2: Spectrum of second-derivative method spectra at 250 nm
y = 0.0001x - 4E-18
R^2 = 1

Fig. 3(A): Calibration plot of 3rd derivative spectrum at 240 nm

y = 0.0003x + 0.0018
R^2 = 0.9998

Fig. 2(A): Calibration plot of second-derivative spectrum at 250 nm

Fig. 3: Spectrum of RACE in third derivative spectroscopy spectra at 240 nm

linearity of racec:

Fig. 3(A): Calibration plot of 3rd derivative spectrum at 240 nm
Second and third derivative spectra of Racecadotril in standard and drug formulation solutions showed that the wavelength of maximum absorbance did not change. According to the result obtained by recovery study, the derivative spectrophotometric method is able to access the analyte in presence of excipients and hence, it can be considered specific limit of detection (LOD) and limit of Quantitation (LOQ) were determined by using the formula based on the standard deviation of response and the slope. The limit of detection (LOD) and limit of Quantitation (LOQ) were calculated by using the equation LOD = 3Xσ/s and LOQ = 10Xσ/S, Where σ is the standard deviation of intercepts is the slope (Table 1).

**Accuracy**

To study the accuracy of the proposed methods and to check the interference from excipients used in the dosage forms, recovery experiments were carried out by the standard addition method. This study was performed by addition of known amounts of Racecadotril to preanalysed solution of commercial tablet. The mean recoveries were found to be 100.51±0.987, 100.38±0.0953, 100.60±0.678 respectively for UV, second and third derivative spectroscopy.

**Precision**

To determine the precision of the method, Racecadotril solutions at a concentration of 8, 20, 40, 60, 80 μg/ml were analyzed each in triplicate. Solutions for the standard curves were prepared fresh everyday. The methods were found to be precise. The % RSD values for intra day precision studies were found to be 0.82, 0.90 and 0.77 for UV, Second and Third derivative spectroscopy respectively.

For robustness and regudness of analytical methods the tests mentioned below were carried out. The robustness of developed methods was tested by changing parameters such as degree of derivation, wavelength range and N value and the optimum parameters were chosen for this study. The UV and derivative spectrophotometric determinations of Racecadotris were carried out by two different analysts on the same standard. The results showed no statistical differences suggesting that the developed methods were robust and druged (Table 2). The developed methods are accurate, sensitive and precise and can be easily applied to be pharmaceutical formulation.

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**REFERENCES**

