Simultaneous Estimation of Metformin and Glimipride by Ultraviolet Spectrophotometry Using Absorbance Ratio Method.

*Suruse S.D., Mundhe D.B.

Dr. V. V. Patil college of Pharmacy Viladghat, Ahmednagar. Maharashtra, India.

Abstract

The simple, accurate, economical and reproducible spectrophotometric method for simultaneous estimation of two-component drug mixture of metformin and Glimipride in combined tablet dosage form has been developed. Developed method is based on direct estimation of metformin and Glimipride at 234nm and 232nm. For estimation absorbance ratio method was developed .The developed methods obey Beer's law in the concentration ranges 5-25µg/ml. The results of analysis were validated statistically.

Key Words

Absorbance Ratio Spectrophotometric Analysis, Metformin, Glimepiride.

Introduction

Combined tablets of metformin and Glimipride are available containing 500 mg of metformin and 1 mg of glimepiride. Metformin hydrochloride chemically 1,1- dimethlybiguanide hydrochloride is white crystalline powder, hygroscopic and freely soluble in water Because metformin's insulin-sensitizing effect occurs mainly at the liver, combination with thiazolidinediones (TZDs), which mainly sensitize muscle to insulin-mediated glucose uptake, is a rational therapeutic strategy. The aim of this paper was to explore the possibility of using techniques of absorbance ratio or Q-analysis method and derivative spectrophotometry methods for quantifying metformin and pioglitazone simultaneously in their mixture form. Glimipride is a medium-to-long acting sulfonylurea anti-diabetic drug and is chemically 3-ethyl-4-methyl-N-(4-[N-((1*r*,4*r*)4methylcyclohexylcarbamoyl)sulfamoyl]phe nethyl)-2-oxo-2,5-dihydro-1H-pyrrole-1carboxamide (Figure 2). Glimipride acts as a secretagogue. It lowers blood sugar by stimulating the release of insulin by pancreatic beta cells and by inducing increased activity of intracellular insulin receptors. It is used for the treatment of diabetes mellitus type 2 Glimipride is not yet official in any of the pharmacopoeia.

*Corresponding Author:

amoldsuruse@gmail.com



Materials and Methods Instrument

A Jasco UV-630 UV/VIS Spectrophotometer was used with 1 cm matches quartz cell.

Solvent used

0.1N NaOH solution.

Preparation of stock Solution

GLM (10mg) and MET(10mg) were accurately weighed and transferred to two separate 100 ml volumetric flask, dissolved in 0.1N NaOH solvent to obtained stock solution of 100 μ g/ml each. The stock solutions of both the drugs were further diluted separately with solvent to obtain 5-25 μ g/ml solution each and scanned in spectrum mode from 400-200 nm. The overlain spectra of both the drug obtained to determine the λ max. GLM has λ max 239nm while MET has λ max 234nm.

Method (Absorbance ratio Method)

From the stock solution, working standard solution of drugs were prepared by appropriate dilution and were scanned in the entire U.V. range. Two wavelengths selected for the method are 234 nm is absorption maxima of MET and 232 nm is the Isobestic point. A series of dilutions were prepared of standard solutions GLM and MET 5-25 µg/ml for both. For the estimation of drugs in the commercial formulations, twenty tablets containing 1 mg of GLM and 500 mg of MET were weighed and average weight was calculated. The tablets were crushed and powdered in glass mortar. Quantity of powder equivalent to 10 mg of MET was transferred to 100ml volumetric flask , dissolved in sufficient quantity of 0.1N NaOH solvent, sonicated and volume was adjust up to mark with solvent to obtain a stock solution of 100 µg/ml of MET and further diluted to get 10 μ g/ml. This solution was then filtered Whatmann filter through paper # 41. Absorbances of these solutions were measured at appropriate wavelengths, and values were substituted in the respective formula to obtain their respective concentrations.

$$Cx = \frac{Qm-Qy}{Qx-Qy} X \frac{A}{ax}$$
$$Cy = \frac{Qm-Qx}{Qy-Qx} X \frac{A}{ay}$$

Validation

The methods were validated with respect to accuracy, linearity, and Limit of detection (LOD) and limit of quantification (LOQ).

Accuracy (Recovery Test)

To ascertain the accuracy of proposed methods, recovery studies were carried out by standard addition method at three different levels (80%, 100% and 120%).

Linearity

The linearity of measurement was evaluated by analyzing different concentration of the standard solution of GLM and MET. For both the method the Beer-Lambert's concentration range was found to be5-25 μ g/ml.

Limit of Detection (LOD) and Limit of Quantification (LOQ)

The LOD and LOQ of GLM and MET were determined by using standard deviation of the response and slope approach as defined in International Conference on Harmonization (ICH) guidelines.

Result and Discussion

The method discussed in the present work provides a convenient and accurate way for simultaneous analysis of MET and GLM. In this method, wavelengths selected for analysis were 234.0 nm $(\lambda \max \text{ of Metformin})$ and 232.0 nm (Isobestic point). In this method linearity for detector response was observed in the concentration range of 5-25 $\mu\text{g/ml}$ for both MET and GLM. Absorptivity coefficient were Calculated for both the drugs at selected wavelengths and substituted in equations for determining concentration of MET and GLM in tablet sample solution. Low values of LOD and LOQ indicated good sensitivity of proposed method. Accuracy of proposed methods was ascertained by recovery studies. The percent recovery for MET and GLM, by this method, was found 99.43 and 99.01. The proposed method could be employed for routine quality control of Metformin and Glimipride in combined dose tablet formulation.

References

- Indian Pharmacopoeia. Govt. of India, Ministry of Health and Family Welfare, Delhi Publication by Controller of Publication, Vol-2, 1996.
- 2. British Pharmacopoeia. Her Majesty Stationery Office, London, Vol-I, 2002
- Analytical Method Validation; Michael E. Swartz, Ira S. Krull, Marcel Dekker, Inc. Ist edition, 2009; 56-65.
- 4. Indian Journal of Pharmaceutical Sciences-2008 May-Jun; 70(3): 383–386.
- 5. Indian Journal of Pharmaceutical Sciences 2010 Jul-Aug; 72(4): 508–510.



Fig. 1: Overlain Spectra of Metformin (MET) and

Glimipride (GLM)



Fig. 2: Calibration curve of Metformin at 234 nm.



Fig. 3: Calibration curve of Glimipride at 239 nm.

Table 1:	Validation	Parameters.
----------	------------	-------------

Parameters	234nm.		232nm.	
	MET	GLM	MET	GLM
Beers law	5-25	5-25	5-25	5-25
limit(µg/ml)				
Regression equation	y = 0.09x - 0.0586	y = 0.0801x + 0.0193	Y=0.0847x - 0.0146	y = 0.0283x + 0.0511
Slope (m)	0.09	0.0801	0.0847	0.0283
Intercept(c)	0.0586	0.0193	0.0146	0.0511
S.D.	0.1396	0.49	0.79	0.2241
LOD	5.12	16.73	9.72	26.13
LOQ	15.52	50.0	27.12	79.20
Correlation	0.9941	0.9998	0.9996	0.9934
coefficient (r2)				

Table 2: Accuracy by percentage recovery method.

Sr. No.	Tablet Formulation (µg/ml)	Level of addition (%)	% estimated (mean ± SD)	
			Metformin	Glimipride
1.	5	80	99.78 ± 0.45	99.20 ± 0.79
2.	5	100	99.69 ± 0.24	98.65 ± 1.01
3.	5	120	98.82 ± 0.42	99.45 ± 1.02