

Method Development And Validation Of Forced Degradation Studies Of Pioglitazone Hydrochloride By Using UV Spectroscopy

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Abstract: Pioglitazone hydrochloride is an oral anti diabetic agent used in the treatment of type 2 diabetes mellitus and also known as non-insulin dependent diabetes mellitus (NIDDM) or adult onset diabetes. A simple, an accurate and economic, precise and reproducible UV Spectroscopy method has been developed for the estimation of Pioglitazone hydrochloride tablet dosage form and validated by ICH guidelines. The standard (10 µg/ml) was scanned between 200-400 nm and maximum absorption was recorded at 270 nm. The assay results are found to be 99.7 ± 0.352 . The linearity range of 10-50 µg/ml proved that it obeyed Beer's Law and the correlation coefficient (r^2) was found to be 0.99986 at 270 nm with an intercept of 0.008 and a slope of 0.011 with RSD 0.187 complied ICH. The pH degradation studies of tablet formulation were found to be less at pH 6-8. The force degradation studies of pioglitazone tablet formulation was done on Stress degradation by hydrolysis under alkaline condition by using 0.1N NaOH was found to be 13.07% for 60min, 17.95% for 90min. Stress degradation by hydrolysis under acidic condition by using 3N HCl and product degradation was found to be 19.75% for 60min and 21.79% for 90 min. Dry heat induced degradation was done by using 70°C temperature was found to be 0.14% for 48 hrs. Oxidative degradation was done by using hydrogen peroxide and product degradation was found to be 12.65% for 15 min. Photolytic degradation was found to be 12.53% for 3hrs and 18.36% for 6hrs.

Keywords: Pioglitazone Hydrochloride, UV Spectroscopy, P^H degradation, Forced Degradation.

INTRODUCTION

Pioglitazone is an oral anti diabetic agent belonging to the class of thiazolidinediones that acts primarily by decreasing insulin resistance. It is used in the management of type 2 diabetes mellitus. It improves sensitivity to insulin in muscle and adipose tissue and inhibits hepatic gluconeogenesis also improves glycemic control while reducing circulating insulin levels. Pioglitazone [(±)-5-[[4-[2-(5-ethyl-2-pyridinyl)ethoxy] phenyl] methyl]-2,4-thiazolidinedione

monohydrochloride belongs to a different chemical class and has a different pharmacological action than the sulfonylureas, metformin, or the -glycosidase inhibitors⁽¹⁾. The chemical structure for Pioglitazone Hydrochloride was shown in Fig.1.

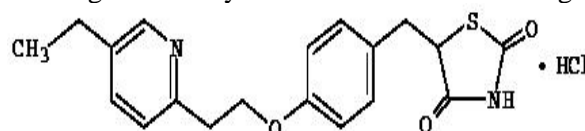


FIGURE 1: Chemical structure of Pioglitazone HCl

Determination of pioglitazone by various analytical methods like Spectrophotometric method⁽²⁾ and HPLC and MECK method⁽³⁾ in tablet dosage form, HPLC and solid phase extraction method in human serum⁽⁴⁾ and in dog serum⁽⁵⁾, HPLC and LC MS in human plasma^{(6), (7)} have been reported. Pioglitazone is not official in any pharmacopoeia. There is a need for a simple, rapid, cost effective and reproducible method for assay of pioglitazone in its dosage forms But there was no reported method for the Forced degradation studies of pioglitazone hydrochloride by using UV spectroscopy. So the present work is to carry out the force degradation studies along with its pH degradation studies. The method was validated according to the ICH (Q2A1995) guidelines⁽⁸⁾ Forced degradation studies may help facilitate pharmaceutical development as well in areas such as formulation development, manufacturing, and packaging, in which knowledge of chemical behaviour can be used to improve a drug product. The available regulatory guidance provides useful definitions and general comments about degradation studies⁽⁹⁾. The International Conference on Harmonization (ICH) guidelines⁽¹⁰⁻¹¹⁾ indicates that stress testing is designed to determine the intrinsic stability of the molecule by establishing degradation pathway in order to identify the likely degradation products and to validate the stability indicating power of the analytical procedure used. ICH guidelines stability testing of new drug substances and products' Q1A(R2)⁽¹⁰⁾ and (Q1B)⁽¹¹⁾ requires that stress testing should be carried out to elucidate the substance. It suggests that the degradation products that are formed under the variety of condition should include the effect of temperature, appropriate oxidation, photolysis and susceptibility

to hydrolysis across a wide range of pH value. In the guideline, the study of effect of temperature is suggested to be done in 10 °C increment above the accelerated temperature (50 °C, 60 °C etc.) and that of humidity at a level of 75 % or greater. Exact details are however provided for the study of oxidation, photolysis and hydrolysis at different pH values⁽¹²⁾

EXPERIMENTAL

Instrument

Absorption spectral measurements were carried out with a UV – Visible spectrophotometer (Shimadzu Model 1700) using UV Probe software version 2 was employed with spectral bandwidth of 1 nm and wavelength accuracy of 0.3 nm (with automatic wavelength correction with a pair of 5 cm matched quartz cells.

Chemicals

Water used for dilution was distilled in the laboratory. A double beam UV spectrophotometer (Shimadzu UV-1800) was used with 1 cm matched quartz cell. Tablet formulation [Pioz 15, USV Ltd., Baddi, Solan- Dist, Himachal pradesh] were procured from a local pharmacy with labeled amount 15 mg per tablet.

Preliminary solubility study of drug

Solubility of the drug was determined at 28±1 °C. A small quantity of standard drugs were dissolved in different solvents like distilled water, methanol, ethanol, acetonitrile, isopropyl alcohol, dimethyl sulfoxide, dimethyl formamide, 0.1 N HCl, chloroform, acetonitrile and pH 4, 7, 9.2 buffer solutions. The results are reported in table 1.

Table 1: Solubility data for Pioglitazone hydrochloride

S.No.	Solvent	Solubility status
1.	Distilled water (HPLC and spectroscopic grade)	In soluble
2.	Ethanol	Slightly soluble
3	Methanol	Freely soluble
4.	Acetonitrile	Soluble
5.	Di methyl sulfoxide	Freely soluble
6.	Di methyl formamide	Soluble
7.	Chloroform	Soluble
8.	Iso propyl alcohol	Insoluble
9.	0.1NaoH	Soluble
10.	Buffer P ^H 4 solution	In soluble
11.	Buffer P ^H 7 solution	In soluble

Solvent Selection:

Various solvents were selected for the solubility studies and it was found that Pioglitazone was soluble in the following solvents; dimethyl sulfoxide, dimethyl formamide, methanol, 0.1 N HCl, chloroform, acetonitrile, *etc.* In the present investigation methanol was selected as a solvent.

Selection of analytical wavelength and absorption maxima:

Appropriate 10 μ g/ml dilutions were prepared for drug from the standard stock solution and the solutions were scanned in the wavelength range of 200-400 nm. The absorption spectra thus obtained was derivatized for zero order spectroscopy. This zero order spectrum was selected for the analysis of the drugs. The absorption maximum was found at 270 nm which can be further used for analysis as shown in Fig.2.

Preparation of stock solutions:

Standard Pioglitazone 100mg was weighed and transformed to a 100 ml volumetric flask and dissolved in 25 ml of methanol. The flask was shaken and volume was made up to the mark with methanol to give a solution containing 1000 μ g/ml (Stock solution A). From this stock solution A, pipette out 5 ml and place into 50 ml volumetric

flask. The volume was made up to the mark with methanol to give a solution containing 100 μ g/ml (Stock solution B)

Selection of analytical concentration range:

From the standard stock solution B of Pioglitazone, appropriate aliquots 1, 2, 3, 4 and 5 ml were pipetted out in 10 ml volumetric flasks and dilutions were made with methanol to obtain working standard solutions of concentrations from 1-50 μ g/ml. Absorbance for these solutions were measured at 270 nm. For standard solution analytical concentration range was found to be 1-50 μ g/ml and overlain spectra was obtained and optical characteristic and linearity data was reported in table 2.

Calibration curve for the Pioglitazone:

Appropriate volumes of aliquots from standard Pioglitazone stock solution B were transferred to different volumetric flasks of 10 ml capacity. The volume was adjusted to the mark with methanol to obtain concentrations of 10, 20, 30, 40 and 50 μ g/ml. Absorbance value of each solution against methanol as a blank were measured at 270 nm. From that absorbance value, regression equation and correlation coefficient (r^2) are determined and presented Fig.3.

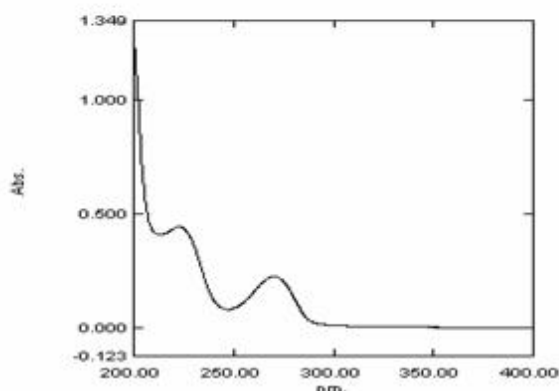
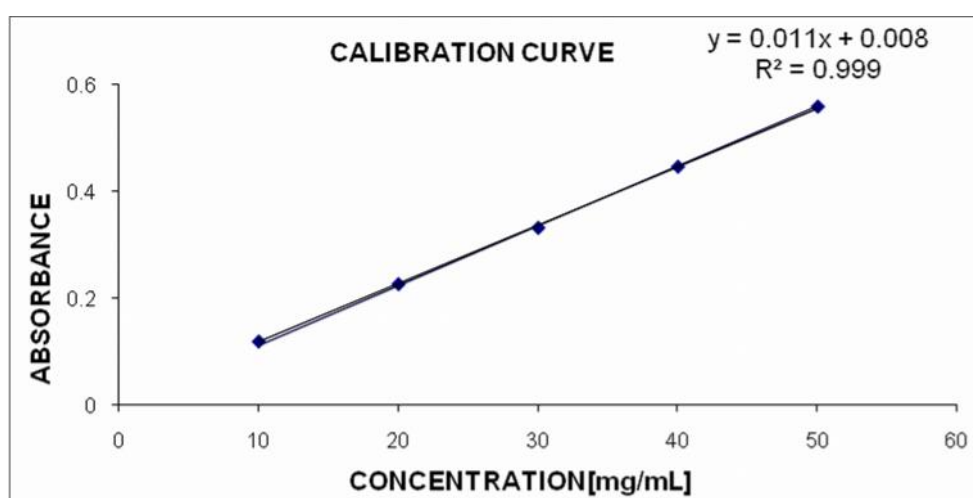


FIGURE 2: UV Spectra of Pioglitazone Standard

Table 2: UV Optical characteristic and linearity data

Parameters	Pioglitazone
max (nm)	270
Beer's law limits in µg/ml	1-50
Correlation coefficient	0.99986
Regression equation $Y=mx+c$	$Y=0.011x+0.008$
Intercept(c)	0.008
Slope	0.011
Std error	0.003302
Molar Abs	4620.504
Sandee's	0.091324

**FIGURE 3: UV Linearity graph of Pioglitazone****Analysis of Pioglitazone from Tablet Dosage form:**

Twenty tablets of formulation were weighed and finely powdered. The powder equivalent to 100 mg of Pioglitazone was accurately weighed. It was then transferred to volumetric flask of 100 ml capacity containing 25 ml of methanol and sonicated for 30 min. The flask was shaken and the solution was filtered through Whatmann filter paper (No. 41) into 100 ml volumetric flask.

Volume was made up to the mark with methanol to give a solution of 1000 µg/ml (Stock solution A). From this solution 5 ml was taken and placed in 50 ml volumetric flask. The volume was made up to the mark using methanol to give a solution of 100 µg/ml (Stock solution B). From the stock solution B, 2.0 ml was taken and diluted to 10 ml to give 20 µg/ml and it was further used for the estimation of Pioglitazone. The result was reported in Table 3.

Table 3: Analysis data of Tablet formulation by UV

Drug	Label claim mg/tab	Amount found mg/tab	Label claim (%)	S.D.*	% COV	S.E*
PIO	15	15.1	100.6	0.115	0.066	0.005

PIO: Pioglitazone, S.D: Standard deviation, COV: Coefficient of variation, S.E: Standard error

METHOD VALIDATION

Validation parameters:

The method was validated with reference to accuracy, precision, and ruggedness.

Accuracy

The accuracy of the proposed methods was assessed by recovery studies at three different levels i.e. 80%, 100%, 120%. The recovery studies were carried out by adding known amount of standard solution of the drug to preanalysed tablet solutions. The resulting solutions were then re-analysed by proposed methods; the results are reported in Table 4.

Precision

Precision of the methods was studied as intra-day, interday and repeatability. Intra-day study was

performed by analyzing, the three different concentration of drug for three times in the same day. Inter-day precision was performed by analyzing three different concentration of the drug for three days in a week. Repeatability was performed by analyzing same concentration of drugs for six times. The results are reported in Table 5.

Ruggedness

Ruggedness of the proposed method is determined by analysis of aliquots from homogenous slot by different analysts using similar operational and environmental conditions. The results are reported in Table 6.

Table 4: Results of Accuracy studies by UV spectroscopy

Level of recovery	Amount of sample (µg/ml)	Amount of drug added (µg/ml)**	Amount of drug recovered (µg/ml)**	% Recovery ± S.D**
80%	10	8.0	7.98	99.7±0.352
100%	10	10.1	9.92	99.2±0.207
120%	10	12.0	11.97	99.7±0.215

** is average of six determinations

Table 5: Precision study data of Pioglitazone by UV spectroscopy

Concentration (µg/ml)	Inter-day Absorbance mean ± SD**	% RSD	Intra-day Absorbance mean ± SD**	% RSD
10	0.120±0.00041	0.342	0.120±0.000117	0.975
20	0.227±0.00052	0.224	0.227±0.00052	0.225
30	0.332±0.00063	0.192	0.332±0.00041	0.124
40	0.446±0.00041	0.089	0.446±0.00052	0.113
50	0.558±0.00052	0.092	0.558±0.00117	0.209

** is average of six determinations.

Table 6: Ruggedness study data of Pioglitazone by UV

Sample	Label claim(mg)	Analyst 1		Analyst 2	
		Amount found (mg)	% Recovery ±SD**	Amount found (mg)	% Recovery ±SD**
Pioz	15	14.89	99.31 ±0.1636	14.95	99.67 ±0.5953

Pioz-Pioglitazone ** is average of six determinations

pH DEGRADATION STUDIES:

The pH effect on the drug was carried out by using 0.1N Hydrochloric acid, 2N Hydrochloric acid, 0.1N Sodium Hydroxide and 2N Sodium Hydroxide solution. The drug solutions (20µg/ml) from pH 0-14 were prepared in the manner as shown in table 7 and these were allowed to stand for 4 hours. Finally the absorbances were

measured at 270nm. The K value for 1st order kinetics was determined by using the formula:

$$K = (2.303/t) \log (C_0/C)$$

Where K= 1st order rate constant,

C₀=initial drug concentration,

C=final drug concentration

The results were reported in table 8.

Table 7: Preparation of sample solution of pH 0-14 for pH stability

pH	Amount of Drug solution added(100 µg/ml) in ml	Amount of 0.1N NaOH solution added(ml)	Amount of HCl added in ml
0	2.5	2.5	12.5 of 2N HCl
1	2.5	2.5	1.25 of 2N HCl
2	2.5	2.5	1.25 of 0.2N HCl
3	2.5	2.5	1.25 of 0.02N HCl
4	2.5	2.5	1.25 of 0.002N HCl
5	2.5	2.5	1.25 of 0.002N HCl
6	2.5	2.5	1.25 of 0.002N HCl
7	2.5	2.5	
8	2.5	2.5	12.5 of 2N NaOH
9	2.5	2.5	1.25 of 2N NaOH
10	2.5	2.5	1.25 of 0.2N NaOH
11	2.5	2.5	1.25 of 0.02N NaOH
12	2.5	2.5	1.25 of 0.002N NaOH
13	2.5	2.5	1.25 of 0.0002N NaOH
14	2.5	2.5	1.25 of 0.00002N NaOH

Table 8: pH Degradation Results:

P ^H	ABSORBANCE (at 270 nm)	CONCENTRATION (µg/ml)	% DRUG DEGRADED	k VALUE	Log k
0	0.45	17.1103	14.45	0.039	-1.4089
1	0.452	17.1863	14.07	0.0379	-1.4214
2	0.449	17.0722	14.64	0.0396	-1.4023
3	0.458	17.4144	12.93	0.0346	-1.4609
4	0.458	17.4144	12.93	0.0346	-1.4609
5	0.502	19.0875	4.56	0.0117	-1.9318
6	0.521	19.8099	0.95	0.0024	-2.6198
7	0.515	19.5817	2.09	0.0053	-2.2757
8	0.508	19.3156	3.42	0.0087	-2.0605
9	0.499	17.0722	14.64	0.0396	-1.4203
10	0.422	16.0456	19.77	0.0551	-1.2588
11	0.392	14.9049	25.48	0.0735	-1.1337
12	0.378	14.3726	28.14	0.0826	-1.083
13	0.342	13.0038	34.98	0.1076	-0.9682
14	0.331	12.5856	37.07	0.1158	-0.9363

Degradation Studies:

The International Conference on Harmonization (ICH) guideline entitled stability testing of new drug substances and products requires that stress testing be carried out to elucidate the inherent stability characteristics of the active substance. The aim of this work was to perform the stress degradation studies on the Pioglitazone hydrochloride using the method developed.

Stress degradation by hydrolysis under acidic condition:

To 3 ml of stock solution (1000 µg/ml) of Pioglitazone, 1 ml of 3 N HCl was added in 10 ml of volumetric flask and the volume was made up to the mark with methanol. Then, the volumetric flask was kept at normal condition for 90 minutes. After 60 min. time interval, 1 ml of solution was pipette out from this flask, neutralised and diluted with methanol in order to make the volume up to 10 ml and the dilution was carried out to achieve the appropriate concentration (30 µg/ml). This solution was taken in cuvette. For the blank, 0.5 ml solution of 3N HCl and 0.5 ml solution of 3N NaOH were diluted with methanol in 10 ml of volumetric flask. After 90 minutes, again 1ml of the solution was pipetted out from the flask and the above procedure was repeated

Stress degradation by hydrolysis under alkaline condition:

To 3 ml of stock solution of Pioglitazone 1 ml of 0.1 N NaOH was added in 10 ml of volumetric flask and made up the volume to the mark with methanol. Volumetric flask was kept at normal condition for 90 min. After 60 min time interval, 1 ml of solution was pipette out from this flask, neutralized and diluted with methanol in order to make the volume up to 10 ml and the dilutions were carried out to achieve the appropriate concentration (20 µg/ml). The solution was then taken in cuvette. For the blank, 0.5 ml solution of 0.1N HCl and 0.5 ml solution of 0.1N NaOH

diluted with methanol in 10 ml of volumetric flask. After, 90 minutes 1ml of solution was again pipetted out from the flask and the above procedure was repeated.

Dry heat induced degradation:

Pioglitazone sample was taken in a petriplate and exposed to a temperature of 70°C for 48 hours in an oven. After 48 hours, 10 mg of the sample was diluted with methanol in order to make the volume up to 10 ml. From this solution, dilutions were carried out to achieve the appropriate concentration (20 µg/ml) and the solution was taken in cuvette for the UV-VIS Analysis.

Oxidative degradation:

To 1.5 ml of the stock solution of Pioglitazone (1000 µg/ml), 1 ml of 30% w/v of hydrogen peroxide added in 10 ml of volumetric flask and the volume was made up to the mark with methanol. The volumetric flask was then kept at room temperature for 15 min. For the blank, 1 ml of the 30 % w/v of hydrogen peroxide was kept at normal condition for overnight in 10 ml of volumetric flask. Both solutions were heated on boiling water bath to remove the excess of hydrogen peroxide. Finally, after 15 minutes dilutions were made from the stock solution to achieve the required concentration (30 µg/ml). The solution was then taken in a cuvette and analysed in UV.

Photolytic degradation:

Sample of Pioglitazone was exposed to near ultraviolet lamp in photostability chamber providing illumination of not less than 1.2 million lux hours. Ten milligrams sample was dissolved in methanol and volume made up to 10 ml. From this solution appropriate dilution (30 µg/ml) was made using methanol and taken in cuvette for the U.V analysis.

The results were reported in table 9.

Table 9: Results of Stress Degradation Studies

Condition	Time	%Degradation
0.1N NaOH(1ml)	60min	13.07%
	90min	17.95%
3N HCl(1ml)	60min	19.75%
	90min	21.79%
30% Hydrogen Peroxide(1ml)	15min	12.65%
Dry Heat 70°	48hr	0.14%
Photolytic	3hr	12.53%
	6hr	18.36%

RESULTS AND DISCUSSION

Pioglitazone was freely soluble in Methanol and Di methyl sulfoxide. Methanol was chosen as a solvent. The drug has maximum absorbance at 270nm. The optical characteristic of drug was found to be Beer's law limits 1-50 µg/ml, Correlation coefficient is 0.99986, Std error is 0.003302, Molar Absorbance is 4620.504. The drug sample was analyzed by UV spectroscopy using methanol as solvent and the average content of drug present in the formulation was found to be 100.6%. The % RSD of accuracy studies was found to be 99.7±0.352. The %RSD of precision was found to be 0.0209 to 0.975. The % recovery of ruggedness was found to be 99.31±0.1636 and

99.67±0.5953. The pH degradation studies of tablet formulation were found to be less at pH 6-8. The force degradation studies of pioglitazone tablet formulation was done on Stress degradation by hydrolysis under alkaline condition by using 0.1N NaOH was found to be 13.07% for 60min, 17.95% for 90min. Stress degradation by hydrolysis under acidic condition by using 3N HCl and product degradation was found to be 19.75% for 60min and 21.79% for 90 min. Dry heat induced degradation was done by using 70°C temperature was found to be 0.14% for 48 hrs. Oxidative degradation was done by using hydrogen peroxide and product degradation was found to be 12.65% for 15 min. Photolytic degradation was found to be 12.53% for 3hrs and 18.36% for 6hrs.

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