A new stability-indicating liquid chromatographic method was developed and validated for the estimation of glycopyrrolate in pharmaceutical formulations. A contemporary approach to analytical life-cycle management was followed to develop a robust and reliable chromatographic method. Scouted method variables such as % methanol, the strength of tetra butyl ammonium hydrogen sulfate and mobile phase flow rate were optimized using the design of experiment approach and their effect on critical quality attributes was studied. The critical quality attributes viz. retention time, theoretical plate count and symmetry factor were highly influenced by the three critical method variables. Optimum chromatography was attained on a C-18 column with a mobile phase methanol: 10 mM tetra butyl ammonium hydrogen sulfate (80:20, v/v) flowing at 1.0 mL.min$^{-1}$. Chromatographic method specificity was ensured by degrading the drug forcefully. Validation studies postulated method acceptability and suitability for estimating glycopyrrolate in both bulk as well as injection formulation. Results for parameters viz. linearity (5-250 µg.mL$^{-1}$), accuracy (>99%) and precision (<2%) advocated method reliability. Overall the method was reliable and of optimum quality and, possess the potential of application in routine and bio-analytical purposes.

**Keywords:** Glycopyrrolate. Stability-indicating. Liquid chromatography. Life cycle management. Validation.

**INTRODUCTION**

Analytical lifecycle management (ALM) is a novel approach which derives its basic principle from the combination of ICH guidelines Q8, Q9 and Q10 (USP, 2013). It has several benefits over the traditional approach as it integrates validation, transfer and verification of procedure (Parr, Schmidt, 2018). This approach is divided into three stages starting with procedure design, which includes defining the analytical target profile (ATP) and critical quality attributes (CQAs). Once the ATP and CQAs are defined, the quality risk management (QRM) tools like fish-bone diagram, control-noise-experimental (C-N-X) approaches are best utilized to identify the critical method variables (CMVs) demanding further studies. The identified variables are investigated using the design of experiments (DoE) to minimize the risks and optimize the experimental conditions. The second stage is known as procedure performance qualification which includes experimentation based on optimized conditions and suitable analytical control strategies are derived. In the conclusive stage i.e. procedure verification the compliance with analytical control strategy is monitored continuously to improve the method performance. Considering the advantages over the traditional approach and lack of analytical methods with the ALM framework, the authors decided to implement the same for the present research.
Another important aspect of developing a quality analytical method is selecting the proper analytical technique. Ultrafast liquid chromatography (UFLC) is a better option for rapid chromatographic studies and is widely used by analysts throughout the world (Panda et al., 2013a; Panda et al., 2013b; Panda et al., 2014). Low mobile phase usage and faster reproducible analysis are the two advantages of UFLC over conventional HPLC, advocating its suitability for use in routine applications.

In the present study glycopyrrolate(GLP), 3-[[cyclopentyl(hydroxy)phenylacetyl]oxy]-1,1-dimethylpyrrolidin-l-ium bromide (Figure 1) is an anti-cholinergic antispasmodic drug and also used in treating chronic severe drooling (O’Neil, 2006; Evatt, 2011). Based on literature review it was found that few HPLC methods are reported for analysis of GLP in various sample matrixes (Nebiu et al., 2007; Rumpler, Sams, Colahan, 2011; Gandala, Pabba, Akula, 2011; Kusuma, Rao, Rameshraju, 2016; Misra, Arora, 2016). However, the reported methods possessed different drawbacks like using complex and corrosive mobile phases, need of column oven, high mobile phase flow rate, lower sensitivity, lack of reliability etc. Moreover, none of the reported methods is ALM supported ensuring the reliability of data obtained. Hence, lack of reliability, as well as robustness, were the two critical aspects basing upon which the authors attempted to develop a new reversed-phase UFLC method for estimation of GLP present in bulk as well as injections. Further, method validation studies were carried out as per ICH Q2 (R1) guidance (ICH, 2005). Forced degradation revealed the stability nature of GLP. Injection formulation of GLP containing 0.2mg.mL⁻¹ of the drug was analyzed by the developed method.

MATERIAL AND METHODS

Material

Standard GLP (purity > 98%) was kindly provided by Roland Institute of Pharmaceutical Sciences, Berhampur. Melting point depression of GLP was measured by Shimadzu DSC-60 (Differential Scanning Calorimeter) Thermal Analyzer. Purity assessment of organic and pharmaceutical compounds by measuring the melting point employing DSC is an age-old established technique(Brennan et al., 1984; Deangelis, Papariello, 1968; van Dooren, Muller, 1984). The melting point of GLP was found to be 195.36 °C which is within the prescribed values of 193-198 °C (US FDA, 2010). The compendial specifications for GLP consider the ICH Q3B (R2) guidance and establish the overall purity of drug being not less than 98% with an individual impurity threshold of not more than 0.15% for the impurities, except the erythro isomer where the threshold is up to 0.4% (USP, 2017). HPLC grade methanol and analytical reagent grade tetra butyl ammonium hydrogen sulfate (TBAHS), hydrochloric acid, sodium hydroxide, hydrogen peroxide were used for the purpose. HPLC grade water prepared by TKA GenPure Ultra-Purification System, Germany was used as the aqueous phase. The commercial injection containing 0.2mg.mL⁻¹ of GLP in water for injection along with preservative benzyl alcohol (0.9%) and hydrochloric acid and/or sodium hydroxide for pH maintenance was analyzed by the UFLC method.

Instrumentation and chromatographic condition

A SHIMADZU Prominence Series UFLC equipped with binary pumps and a PDA detector was used for the purpose. A reversed-phase column (250×4.6 mm, 5 µm) was used for chromatography. The mobile phase contained methanol and 10 mM TBAHS in a composition of 80:20%, v/v with a flow rate of 1 mL.min⁻¹, and detection at 204 nm. Prior use both the mobile phase components were sonicated for degassing.

Procedure Design

Setting ATP and CQAs

The ATP for the present study was to develop and validate a rapid, robust and reliable chromatographic method for estimation of GLP in bulk and injection.
formulation. Based on the ATP a UFLC was selected for the purpose. The selected CQAs were retention time, plate number and symmetry factor.

**Risk management**

Risk assessment is a constitutive part of effective risk minimization process. Ishikawa fish-bone diagram acts as a preliminary tool for risk assessment and addresses all potential method variables that can affect the CQAs adversely. From the various variables, a critical few are selected after being subjected to a C-N-X analysis. Method variables with scores higher than 200 were selected as the critical few and were further subjected to DoE investigation. Three variables viz. % methanol, the strength of TBAHS solution and flow rate of the mobile phase were identified as CMVs for DoE based investigation.

**Design of experiment and method optimization**

Based on the number of variables to be studied it was found suitable to choose a Box-Behnken design (BBD) for the robustness investigation and optimization study. Minimal 15 runs were performed in a randomized manner having 3 centre points. The obtained experimental data was analyzed mathematically to study any possible interaction effect among the variables. Further statistical data analysis was carried out to evaluate the aptness of the study model. Various model parameters along with polynomial equations, 2-dimensional and 3-dimensional plots were assessed for checking the model suitability. Based on the results of the above assessment analytical control space was developed for procedure performance qualification purpose.

**Procedure Performance Qualification**

**Analytical control strategy**

Control strategies were developed based on the optimized method conditions and control charts were prepared for the CQAs over a period of 6 days (n=3). It helped to derive robust working limits for the method variables to obtain desired CQAs.

**Procedure Verification**

It consisted of continuous monitoring of the newly optimized analytical method in order to reduce any incident of risks arising from time to time and to improve the method performance in terms of specified CQAs.

**Method Validation**

**Specificity**

Specificity capability of this current method was confirmed by forcefully degrading GLP in an ICH compliant framework. The ICH guidance advocates for any suitable set of experimental conditions which can result in degradation of the analyte to a considerable level and generate degradation products, if any. Hence, to attain the best of both of these objectives, stress conditions were developed utilizing minimal time and chemicals. Stress conditions like acid, alkali, peroxide, temperature and UV light helped in the degradation of GLP. An aliquot of 400 µL of GLP was taken in five different 5 mL volumetric flasks. GLP was exposed to 500 µL of 0.1M HCl, 0.1M NaOH, 3% H$_2$O$_2$, 80 ºC and UV light (365 nm) for a specified time period (30min for all the conditions except photolysis where the exposure time was up to 3 h). Afterwards, the acid and alkali stressed samples were neutralized using respective counterparts, and the other samples were diluted to 5 mL with mobile phase. The amount of drug degraded was calculated by comparing the results with the untreated drug solution. GLP at 80 µg.mL$^{-1}$ was chosen for the forced degradation study purpose.

**Preparation of Calibration Curve**

Around 10mg of accurately weighed GLP was dissolved in mobile phase placed in a 10 mL volumetric flask. Further, this solution was used to prepare working standard solutions within 5-250 µg.mL$^{-1}$ of GLP. The average peak areas (n=3) for each concentration were used to prepare the curve.

**Accuracy**

A fixed concentration (80 µg.mL$^{-1}$) solution of GLP present in injection solution was chosen and was spiked with 80%, 100% and, 120% of standard GLP. Recovery of standard GLP was indicative of method trueness.

**Precision**

Method precision was assessed in terms of repeatability, intermediate and instrument precision at
80 µg.mL⁻¹. The precision of the UFLC was determined based on hexaplicate injection of aforementioned concentration of GLP, whereas the repeatability and intermediate precision were determined from injection of six separately prepared solution of the analyte at the same concentration. A % relative standard deviation (%RSD) value less than 2.0 was the acceptance limit.

**Sensitivity**

Signals of the lowest concentration of samples were measured and compared to that of blank to establish the effective concentrations of the analyte as the limit of detection (Signal/Noise=3) and quantitation (Signal/Noise=10). Analyte concentrations producing the desired signal-to-noise ratio were designated as LOD and LOQ, respectively.

**Assay of a commercial formulation**

An injection volume equivalent to 5mg of GLP was taken and ultrasonicated for 30 min in the presence of mobile phase. Once the final volume was made up this solution was filtered using a membrane filter (0.45 µ). Thereafter, it was utilized to prepare a sample solution for LC analysis. Further, solutions of formulation components such as hydrochloric acid and sodium hydroxide mixture (acts as pH balancing agent) and benzyl alcohol (preservative) were also prepared separately and injected onto the UFLC to assess any possible interference at analytes retention time.

**RESULTS AND DISCUSSION**

**Preliminary method development using a lifecycle management approach**

According to the physicochemical property of analyte, a reversed-phase LC method was found befitting for chromatography of GLP. GLP is a highly polar compound with a greater tendency for early chromatographic elution. This typical problem was overcome by choosing a superior ion-pair reagent like tetra butyl ammonium hydrogen sulfate (TBAHS) which has the capability of enhancing method selectivity as well as the solubility of GLP and also may reduce adsorptive interaction on the stationary phase. The literature revealed that other researchers have used complex and corrosive mobile phases, column oven, and high mobile phase flow rate. Moreover, the methods were less sensitive and lacked reliability. Addressing these demerits in the present paper 10 mM TBAHS solution was used as the aqueous phase avoiding any blockage issue due to high amounts of buffer salts. Further, none of the reported methods utilized 10 mM TBAHS solution as the mobile phase for estimating GLP. Hence, considering the above facts TBAHS solution was chosen as the aqueous phase. The typical mobile phase constituted of methanol: 10 mM TBAHS. Various compositions of mobile phase and flow rates such as 1.0 mL, 1.1 mL and 1.2 mL were tested to check the response of GLP at 204 nm. A composition of methanol: 10 mM TBAHS at 80:20, v/v flowing at 1.0 mL.min⁻¹ gave optimum peak performance.GLP eluted at 3.134minwith a run time of 7 with Further, considering the above conditions as initial method development start point, life cycle management approach was followed for establishing a reliable stability-indicating analytical method for estimation of GLP in pharmaceuticals.

With the intent of accomplishing ATP preliminary risk assessment and its minimization was carried out. The typical fish-bone based causal-effect relationship diagram (Figure 2) depicted numerous method variables with a possible threat to method performance quality. Out of the different risk, causes were assessed for the level of risk using a traffic light risk analysis matrix. In the mentioned matrix variables were designated to high, medium and low risk through colours viz. red, yellow and green, respectively (Table I). Certain method variables were found to possess a possible risk to the method performance i.e. CQAs. For enhanced discernment of the criticality of these variables a controlled noise experimental (C-N-X) approach was initiated and the results are displayed in Table II. This approach concluded that the method variables such as % methanol composition, the strength of TBAHS solution and flow rate had the highest risk potential based on their CNX score. Further, these three variables were named as CMVs and subjected to DoE investigation in order to enhance method robustness, optimize the method and minimize the incidence of any risks to method performance.

**Design of experiments based investigation**

The BBD domain proved to be worthy, as it unearthed the influence of CMVs on the CQAs. Experiments performed on randomized manner provided bias-free results (Table III).
The optimization data was subjected to appropriate mathematical models for analysis. Polynomial equations (Eq. 1, 2 and 3) consisting of model terms for both main effects and interaction effects were generated for the CQAs. It helped to unearth the connection among the CMVs and CQAs, for a sound understanding of the various effects. Synergistic and antagonistic effects were sorted from the polynomial equations based on positive and negative signs of each coefficient.

Assessment of ANOVA (P<0.05) along with satisfactory values of $r^2$ ($r^2>0.9$) advocated for the appropriacy of the selected mathematical model for obtaining optimum values of CQAs. Significant nature of the studied models was established by evaluating the predicted F-values obtained (4435.78, 282.3, and 79.17 for retention time, theoretical plate count and symmetry factor, respectively). Model appropriacy for design space analysis was further ascertained by obtaining adequate precision (signal-to-noise ratio) values greater than 4.0 for all the three CQAs.

Response surface evaluation was performed employing 3-D plots (Figure 3-(A-I)). Figure 3(A) depicts a slight decrease in retention time at high levels of % methanol and strength of TBAHS. A very similar interpretation was noticed for the CMVs % methanol and flow rate (Figure 3(B)). A typical “hat” shaped response surface was obtained depicting complex interaction among the CMVs % methanol and strength of TBAHS (Figure 3(C)). Increased retention times were seen at low levels of both the CMVs. However, the retention of GLP decreased gradually with the increasing flow of mobile phase and with low levels of strength of TBAHS.
### TABLE I - Traffic light risk analysis matrix

| Symmetry factor | Theoretical Plate Count | Retention Time | CQAs                      |
|-----------------|--------------------------|----------------|---------------------------|
| HIGH            | HIGH                     | HIGH           | Mobile Phase              |
| HIGH            | HIGH                     | HIGH           | Strength of TBAHS         |
| LOW             | LOW                      | MEDIUM         | Injection volume          |
| LOW             | MEDIUM                   | HIGH           | Solvent grade             |
| LOW             | LOW                      | HIGH           | Sample Purity             |
| LOW             | LOW                      | MEDIUM         | Reagent Purity            |
| LOW             | MEDIUM                   | LOW            | Humidity                  |
| LOW             | LOW                      | LOW            | Temperature               |
| MEDIUM          | LOW                      | LOW            | Peak integration          |
| MEDIUM          | MEDIUM                   | LOW            | Peak Purity               |
| HIGH            | HIGH                     | HIGH           | UFLC                      |
| LOW             | LOW                      | LOW            | Flow rate                 |
| MEDIUM          | MEDIUM                   | MEDIUM         | Sonicator                 |
| LOW             | MEDIUM                   | LOW            | Calculation Error         |
| LOW             | MEDIUM                   | LOW            | Dilution Error            |
| MEDIUM          | LOW                      | MEDIUM         | Glassware error           |
| LOW             | MEDIUM                   | HIGH           | Equilibration Time        |
| LOW             | MEDIUM                   | HIGH           | Stationary Phase          |
TABLE II - Typical control-noise-experimental (C-N-X) based identification of CMVs

| Method Variables         | Risk Level* on CQAs       | Score | C,N,X | Action Plan     |
|--------------------------|---------------------------|-------|-------|-----------------|
|                          | Retention time | Theoretical Plate Count | Symmetry factor |        |     |
| % methanol               | 10             | 10   | 10    | 300    | X       | DoE   |
| Strength of TBAHS        | 10             | 9    | 8     | 270    | X       | DoE   |
| Flow rate                | 10             | 8    | 7     | 250    | X       | DoE   |
| Sample Purity            | 6              | 3    | 3     | 120    | N       | Quality Assessed |
| Solvent grade            | 5              | 3    | 3     | 110    | C       | Controlled |
| Stationary Phase         | 4              | 2    | 3     | 90     | C       | Controlled |
| UFLC                     | 3              | 3    | 2     | 80     | C       | Controlled |
| Calculation error        | 2              | 2    | 2     | 60     | N       | Careful Measurement |

*Risk Level: 1-Negligible, 5-Low, 10-High; Final Score = (Risk Level of 1st CQA×10) + (Risk Level of 2nd CQA×10) + (Risk Level of 3rd CQA×10)
### Table III - Results for robustness-cum-optimization study by Box-Behnken experimental domain

| Run | % Methanol | Strength of TBAHS (mM) | Flow rate (mL.min⁻¹) | Retention Time (min) | Theoretical Plate Count | Symmetry factor |
|-----|------------|-------------------------|----------------------|----------------------|-------------------------|-----------------|
| 1   | 80         | 10                      | 1.0                  | 3.157                | 2732                    | 1.628           |
| 2   | 82         | 10                      | 0.9                  | 3.231                | 2576                    | 1.546           |
| 3   | 80         | 10                      | 1.0                  | 3.155                | 2739                    | 1.621           |
| 4   | 82         | 9.5                     | 1.0                  | 3.086                | 2387                    | 1.492           |
| 5   | 80         | 9.5                     | 0.9                  | 3.41                 | 2643                    | 1.621           |
| 6   | 80         | 9.5                     | 1.1                  | 2.933                | 2584                    | 1.623           |
| 7   | 78         | 10                      | 1.1                  | 3.182                | 2625                    | 1.601           |
| 8   | 78         | 9.5                     | 1.0                  | 3.185                | 2678                    | 1.78            |
| 9   | 80         | 10                      | 1.0                  | 3.155                | 2742                    | 1.624           |
| 10  | 78         | 10.5                    | 1.0                  | 3.218                | 2599                    | 1.574           |
| 11  | 82         | 10.5                    | 1.0                  | 2.948                | 2487                    | 1.651           |
| 12  | 80         | 10.5                    | 1.1                  | 3.111                | 2523                    | 1.542           |
| 13  | 82         | 10                      | 1.1                  | 2.824                | 2501                    | 1.492           |
| 14  | 80         | 10.5                    | 0.9                  | 3.114                | 2727                    | 1.611           |
| 15  | 78         | 10                      | 0.9                  | 3.251                | 2805                    | 1.656           |

Coded Levels

| Low  | 78 | 9.5 | 0.9 |
| Mid  | 80 | 10  | 1.0 |
| High | 82 | 10.5| 1.1 |

Where A= Methanol (%), B= Strength of TBAHS (mM) and C= Flow rate (mL.min⁻¹)
A “maxima” system depicting maximum response was obtained around the lower-mid level of % methanol and mid-levels of the strength of TBAHS. The theoretical plate count was found to decrease with increasing values of methanol content at all levels of strength of TBAHS (Figure 3(D)). The theoretical plate values were found decreasing in a linear pattern at low and high levels of methanol content and around all the levels of strength of TBAHS. However, an increase in theoretical plates around mid-levels of % methanol was noticed (Figure 3(E)). A curvilinear response surface was obtained for theoretical plates with increasing strength of TBAHS throughout all levels of flow rate (Figure 3(F)).

A typical “hat” shaped response surface was obtained depicting complex interaction among the CMVs % methanol and strength of TBAHS (Figure 3(G)). Increased tailing was observed at low levels of both the CMVs. However, the tailing of GLP decreased gradually with increasing methanol content and at low
levels of strength of TBAHS. No significant change in
tailing was noticed with a change in % methanol and
flow rate of mobile phase (Figure 3(H)). Tailing was
found to decrease to a small extent with higher strength
of TBAHS and high levels of flow rate (Figure 3(I)).
Analogous information was drawn by interpreting the
2-dimensional contours (not shown in figures) for all
the respective CMVs.
Response sensitiveness was evaluated by
perturbation plots (not shown in figures), which
advocated and reinforced the results obtained during
response surface interpretations. Intersecting curved
and steep lines were indicative of the presence of
interaction among studied CMVs. The desirability,
as well as overlay plot, represented chromatographic
conditions for obtaining optimum values of all the
three CQAs. Relying on the above-obtained conditions
the method was subjected to validation studies.

Method validation studies

Specificity

Forced degradation studies were performed with
the dual intention of ascertaining method specificity
as well as establishing stability behaviour of the drug.
Visual evaluation of the chromatograms of stressed
drug solutions advocated method specificity (Figure
4). The forced degradation experiments were carried
out with an objective of obtaining degradation products
with minimal use of chemicals along with a study
protocol consuming less time. As a general start point
based on prior knowledge and practice 0.1M HCl, 0.1M
NaOH, 3% H₂O₂, 80 °C temperature and UV light at
365 nm were chosen to stress the analyte. The chemical
reagents were utilized at a very low volume of 500 µL
for a minimal time period of 30 min. In the case of
UV radiation, the exposure time was up to 3 h with
the intent of providing sufficient exposure to UV light
for generating degradation products. GLP degraded
to a smaller extent in applied peroxide, thermal and
photolytic conditions. However, extensive degradation
was observed in the acid and alkali stress conditions.
GLP was completely degraded upon exposure to 0.1 M
NaOH. Hence, the strength of NaOH was reduced and
a 0.01M NaOH was used to assess stability. The GLP
degraded more than 50% of its initial content indicating
its susceptibility towards the new alkaline condition
than compared to the acid condition. Two additional
peaks were noticed at 5.0 min (Degradation product-I;
DP-I) and 6.3 min (Degradation product-II; DP-II), in
both the acid and alkaline stress conditions, which may
be attributed to the possible break down of the ester
linkage between α-cyclopentyl mandelic acid and
3-hydroxy-1, 1-dimethylpyrrolidinium bromide during
exposure to both acid and alkaline stress. Further,
according to USP, the DP-I i.e. α-cyclopentyl mandelic
acid is an official related compound C of GLP (USP,
2017) which was generated during the stress studies
of GLP. Moreover, the compendial monograph reports
two separate procedures for determining the organic
impurities of GLP and in both of them cyclopentyl
mandelic acid is a reported related substance. Method
specificity was found intact as both the degradation
product peaks were well resolved from GLP peak
(resolution >5) as well as among them (resolution>3).
The analyte was found stable to exposure to UV light
up to 3 h suggesting suitability of the process for
regular routine use. The results endorsed the study
intent (Table IV).
FIGURE 4 - Typical chromatograms depicting GLP pure drug (A) and after treatment with 0.1 M HCl (B), 0.01 M NaOH (C), 3% \( \text{H}_2\text{O}_2 \) (D), 80 °C temperature (E) and UV light at 365 nm (F).

Table IV - Results of forced degradation study

| Stress Condition | Time Period | Retention time(min)          | Degradation/No Degradation |
|------------------|-------------|------------------------------|----------------------------|
| 0.1M HCl         | 30 min      | Drug: 3.189
Unknown DP-I: 5.060
Unknown DP-II: 6.360 | Degradation |
| 0.01 M NaOH      | 30 min      | Drug: 3.154
Unknown DP-I: 5.078
Unknown DP-II: 6.356 | Degradation |
| 3% \( \text{H}_2\text{O}_2 \) | 30 min      | Drug: 3.164
Unknown DP: 4.464 | Degradation |
| 80 °C            | 30 min      | Drug: 3.148
Unknown DP: 5.244 | Degradation |
| UV light (365 nm)| 3 h         | Drug: 3.129 | No Degradation |
Linearity

The method was found linear over the concentration range of 5-250 µg.mL\(^{-1}\) (\(r^2 =0.999\)). Further, satisfactory results obtained through regression analysis and ANOVA of linearity data indicated the goodness of fit.

Accuracy

Satisfactory recoveries of GLP between 99.64-100.35%, advocated for optimum method accuracy and reliability.

Precision

The precision study revealed acceptable values of % RSD (<2%). The values were 0.63%, 0.74% and 0.37% for intraday, inter-day and system precision, respectively.

Limit of detection (LOD) & limit of quantitation (LOQ)

The LOD and LOQ values were 2 and 5 µg.mL\(^{-1}\), respectively.

Analytical control strategy

Preparation of control charts (Table V) helped to develop analytical control strategies. Reproducible results for CQAs were obtained by working within the analytical control space. The control space was defined to be within limits such as methanol proportion (±2%), the strength of TBAHS (±0.5 mM) and, flow rate (±0.1 mL.min\(^{-1}\)).

| Parameter | Retention Time (min) | Theoretical Plate Count | Symmetry Factor |
|-----------|----------------------|-------------------------|-----------------|
| Mean      | 3.153                | 2738.88                 | 1.626           |
| S.D.      | 0.0012               | 6.614                   | 0.0012          |
| R.S.D. (%)| 0.038                | 0.241                   | 0.073           |

(continuing)

Table V - Results of control charts obtained for CQAs

| Parameter | Retention Time (min) | Theoretical Plate Count | Symmetry Factor |
|-----------|----------------------|-------------------------|-----------------|
| LCL       | 5.111                | 4227.57                 | 1.381           |
| UCL       | 5.114                | 4262.43                 | 1.383           |

Assay of a commercial formulation

The visual evaluation of chromatograms obtained for injections indicated method selectivity due to the non-interference of any of the formulation components (Figure 5). The mean (n= 3) content of GLP was found to be 98.33% (SD = ±0.29). Further, a visual comparison of chromatograms of standard GLP and GLP in injection preparation was carried out to establish method selectivity (Figure 6).

CONCLUSIONS

The present research explains the optimization and development of a UFLC method for estimating GLP in bulk and dosage forms. To achieve the objective a systematized analytical lifecycle management approach was followed. Utilizing the ALM approach, not only ensured increased method robustness but also presented an option for continuous improvement in the performance of CQAs. It helped to discover three CMVs and their effects on the CQAs. Based on the result of control charts, control strategies were outlined to obtain desired UFLC method performance. Overall, this analytical method is suitable and trustworthy for estimating GLP. Results of the validation study were found compliant with ICH guidelines. Forced degradation studies revealed that GLP is highly susceptible to the applied acid and alkaline stress conditions and produced two degradation products DP-I and DP-II, which calls for adequate preventive measures to be developed by the formulation scientist to protect therapeutic safety and efficacy of the pharmaceutical product. The forced degradation conditions were satisfactory as a known related compound C was generated as impurity implying appropriateness of the selected stress conditions. The optimum system suitability and satisfactory assay results depicting non-interference of the formulation
FIGURE 5 - Typical chromatograms depicting non-interference of formulation components at analytes retention time when blank mobile phase was injected with (A) hydrochloric acid and sodium hydroxide mixed solution, (B) benzyl alcohol solution.
components advocated for the accomplishment of ATP. Hence, this method is acceptable for estimating GLP in bulk and injection formulation. Further, this method has the potential for determining GLP in biomatrixes.

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