

Article

A Robust Liquid Chromatographic Method for the Analysis of Azithromycin in Bulk Samples, Tablets and Suspensions

Alex O. Okaru ^{1,*}, Kennedy O. Abuga ¹, Franco N. Kamau ¹, Stanely N. Ndwigah ¹
and Dirk W. Lachenmeier ²

¹ Department of Pharmaceutical Chemistry, University of Nairobi, Off Ngong Road, Nairobi, P.O. Box 19676-00202 Nairobi, Kenya; koabuga@e-mail.com (K.O.A.); franco.kamau@uonbi.ac.ke (F.N.K.); snndwigah@uonbi.ac.ke (S.N.N.)

² Chemisches und Veterinäruntersuchungsamt (CVUA) Karlsruhe, Weissenburger Straße 3, 76187 Karlsruhe, Germany; lachenmeier@web.de

* Correspondence: alex.okaru@gmail.com; Tel.: +254-726-53-9193

Abstract: A simple, isocratic and robust RP-HPLC method for the analysis of azithromycin was developed, validated and applied for the analysis of bulk samples, tablets and suspensions. The optimum chromatographic conditions for separation were established as mobile phase comprising of acetonitrile-0.1M KH₂PO₄ pH 6.5-0.1M tetrabutyl ammonium hydroxide pH 6.5-water (25:15:1:59% v/v/v/v) delivered at a flow rate of 1.0 ml/min. The stationary phase consisted of reverse-phase XTerra® (250 mm× 4.6 mm i.d., 5 µm particle size) maintained at a temperature of 43 °C with a UV detection at 215 nm. The method was found to be linear in the range 50-150% (r²=0.997). The limits of detection and quantification were found to be 0.02% (20 µg) and 0.078% (78 µg) respectively with a 100.7% recovery of azithromycin. Degradation products of azithromycin in acidic and oxidative environments at 37 °C were resolved from the active pharmaceutical ingredient and thus the method is fit for the purpose.

Keywords: azithromycin; HPLC; robust; isocratic; degradation products

1. Introduction

Azithromycin is a semi-synthetic macrolide antibiotic used clinically for a wide range of bacterial infections [1,2]. Azithromycin is listed in the World Health Organization and the Ministry of Health, Kenya essential drugs list [3,4] where it is indicated for the management of atypical infections. Suitable analytical methods are required to monitor the quality of azithromycin (AZT) products during manufacture, batch-release and post market surveillance. The analytical method ought to be selective for AZT in the presence of related substances, namely the synthetic intermediates azithromycin (AZA), erythromycin A oxime (EAOX) and erythromycin A imino ether (EAIE) and the degradation products *N*-demethylazithromycin (NDMAZT) and decladinosylazithromycin (DAZT) (Figure 1).

There are several published methods for the analysis of azithromycin such as biological [5], electroanalytical [6], spectrophotometric [7], capillary electrophoresis [8], thin layer chromatography [9] and liquid chromatography [10–13]. Liquid chromatography is the method of choice for the analysis of azithromycin in bulk samples and formulations since it is able to determine azithromycin in the presence of related substances. The current pharmacopoeial liquid chromatographic methods for the analysis of azithromycin utilize relatively expensive columns, unstable detectors and high pH and/or temperature limiting the use of conventional silica based columns [12,13]. The United States

Pharmacopoeia [13] employs a mobile phase of pH 11 with electrochemical detection while the British Pharmacopoeia [12] utilizes a polymer column with a mobile phase of pH 8. The electrochemical detector is not commonly available and produces unstable response compared to the UV detector since the electrodes have direct contact with column effluent and may react with sample matrix while polymer columns are relatively more expensive compared to silica-based columns. This calls for use of pH and temperature stable strategies to improve on column longevity, stability and peak parameters. Such a method can be used for routine quality control of azithromycin bulk samples and formulations as well as market surveillance of samples. A reverse phase high performance liquid chromatographic method for the estimation of azithromycin suspension described by Sachin *et al.* [14] did not exhibit the required specificity regarding the related substances, NDMAZT, AZA, DAZT, EAOX and EAIE. This paper describes the development, validation and application of a selective, isocratic and robust liquid chromatographic method for the analysis of azithromycin in bulk samples and oral dosage forms.

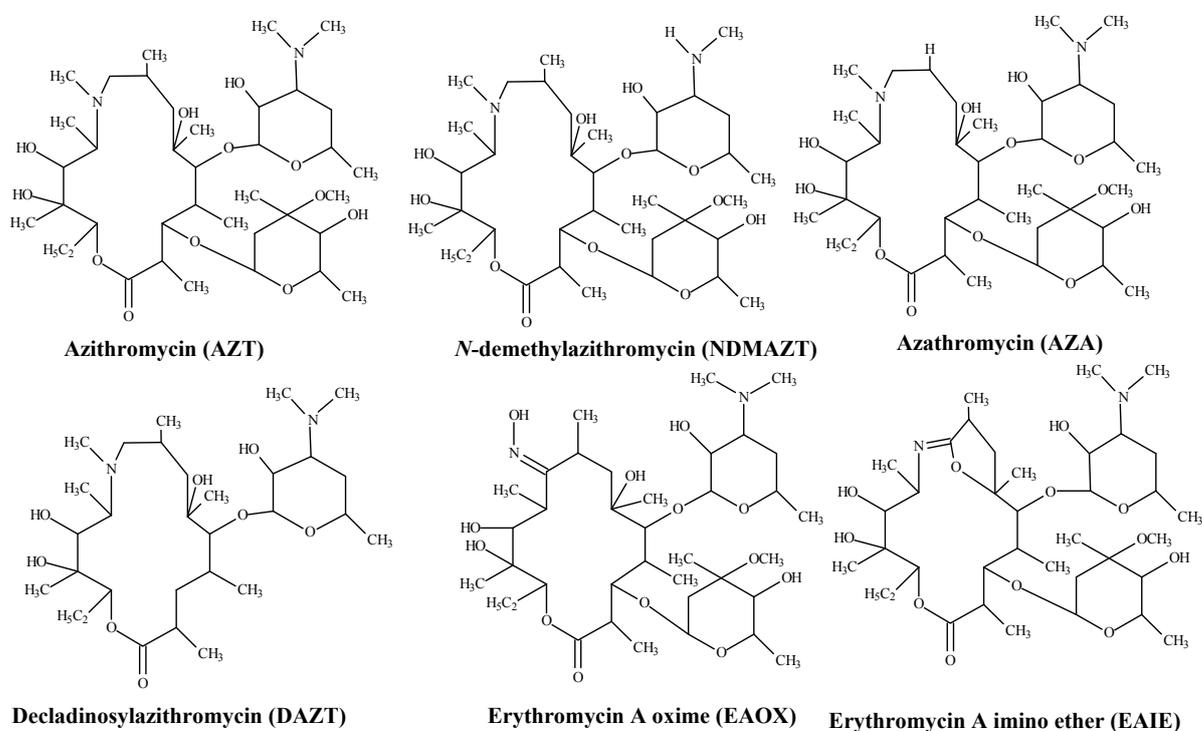


Figure 1. Chemical structures of azithromycin and some related substances.

2. Materials and Methods

2.1. Chemicals and reagents

Analytical grade KH_2PO_4 , K_2HPO_4 and H_3PO_4 were from Loba Chemie Pvt. Ltd. (Mumbai, India) while triethylamine and triethanolamine were from BDH Laboratory Supplies (Poole, UK). Tert-butylammonium hydroxide was from Sigma-Aldrich (Steinheim, Germany). HPLC grade acetonitrile (Scharlau, Barcelona, Spain) was used as organic modifier. Purified water was prepared using distillation by GFL distillation machine type 2001/4 (Gesellschaft für Labortechnik mbH, Burgwedel, Germany). Azithromycin dihydrate working standard was donated by the Drug Analysis and Research Unit (DARU), University of Nairobi, Kenya. The working standards of erythromycin A oxime, erythromycin A imino ether, N-demethylazithromycin,

decladinosylazithromycin and azithromycin standards were purchased from Pfizer Inc. (Groton, CT, U.S.A.). Azithromycin tablets and suspensions were randomly purchased from various pharmacies in the Central Business District of the City of Nairobi. The innovator brand, Zithromax®, was from Pfizer Laboratories Ltd. (Industrial Area, Nairobi, Kenya). Azithromycin bulk sample was from SkyLight Chemicals (Nairobi, Kenya).

2.2. HPLC instruments and analytical conditions

A Shimadzu high performance liquid chromatographic (HPLC) system (Shimadzu Corp., Kyoto, Japan) was used during the study. It was supported by a CBM-20A Prominence communication bus module as the system controller and an LCSolutions software Ver. 1.22, SP1 and equipped with an SPD-M20A Prominence UV/Visible photo array diode detector operating with a deuterium and a tungsten lamps as UV and Visible light sources. A LC-20AD Prominence solvent delivery system with a dual-plunger tandem-flow solvent delivery system and a SIL-20AC Prominence autosampler were a part of the HPLC system. The temperature was controlled using a CTO-M20AC column oven with a block heating thermostatic chamber. Mobile phases were degassed using a Power sonic 410 bench-top ultrasonic bath (Daihan Labtech Ltd., Kyonggi-Do, Korea). A Waters XTerra®, a hybrid, reverse-phase C18 column (250 mm ×4.6 mm, 5 µm) (Waters Corporation, Milford, MA, USA) was used for separations. The detection wavelength was set at 215 nm while the sample injection volume was at 20 µl and the flow rate at 1 ml/min.

2.3. Preparation of solutions

2.3.1. Working standard mixture

The working standard mixture was prepared by dissolving the working standards in acetonitrile-water (50:50 % v/v) to concentrations of azithromycin 5.0 mg, 1.5 mg azithromycin, 0.64 mg demethylazithromycin, 0.75 mg decladinosylazithromycin, 0.304 mg erythromycin A oxime and 0.31 mg erythromycin A imino ether in 1 ml of solution. These concentrations were preferred since they yielded comparable peak heights among the related substances for accurate derivation of chromatographic parameters. The related substances have varying response factors compared to AZT. Thus erythromycin A imino ether was incorporated at 2.5% level while decladinosylazithromycin and azithromycin were at 10% level while maintaining azithromycin at 100% (5 mg/ml). The other related substances were incorporated into the working mixture at levels ranging between 2.5% to 10% for ease of monitoring of chromatographic parameters of all components.

2.3.2. Mobile phases

Mobile phases were prepared by mixing appropriate volumes 0.1M KH₂PO₄ (stock solution) with water and acetonitrile. The pH of 0.1M KH₂PO₄ had been preadjusted with an equimolar solution of K₂HPO₄. Mobile phases were degassed in an ultra-sonic water bath for 30 min before use.

2.4. Peak purity analysis

Peak purity analysis was performed over a wavelength range of 200-900 nm using the Shimadzu LCSolutions software Ver. 1.22 SP1 (Shimadzu, Kyoto, Japan) [15]. Although peak purity

analysis does not completely prove that the peak observed is due to one component, it gives confidence in the method [16].

2.5. Method validation

2.5.1. Linearity

Linearity was determined on solutions of azithromycin working standard within the 50-150% range. For this purpose, a 5.0 mg/ml solution was taken as 100%. Solutions were made by dissolving azithromycin in acetonitrile-water (50:50% v/v) and run in triplicate. The data obtained from the linearity experiments was subjected to linear regression analysis with the concentration of injected AZT standard being plotted against the peak areas obtained.

2.5.2. Sensitivity

The detection limit and limit of quantitation were determined by serially diluting 5 mg/ml AZT solution to yield signal to noise ratios of 3 and 10 respectively.

2.5.3. Accuracy

Accuracy was evaluated by analyzing synthetic mixtures spiked with known quantities of azithromycin. Triplicate determinations at three concentration levels (80, 100 and 120 %) were used. The mean recoveries of the assays were assessed for compliance with ICH guidelines [17].

2.5.4. Robustness

The Box-Wilson Design (BWD) was used to determine the optimum range of the chromatographic factors within which the method was robust. A three level (-1, 0 and +1) experimental design was applied out as shown in Table 1. The experiments were randomized with the aid of the Statgraphics Centurion XVI (StatPoint Technologies Inc., VA, USA) software. The interactive effects of the three variables of temperature, acetonitrile concentration and buffer pH on the selectivity of the critical peak pair of EAIE and AZT were evaluated as a measure of robustness. Pareto analysis was used to identify the interactive effects that had the greatest impact ($p=0.05$) on the selectivity of the critical pair of EAIE and AZT.

Table 1. Nominal values corresponding to low, central and high levels.

Factor	Low level (-1)	Central level (0)	High level (+1)
pH	6.0	6.5	7.0
Acetonitrile	22	25	28
Temperature	41	43	45

2.6. Assay of azithromycin in bulk samples, tablets and suspensions

Twenty tablets were weighed and pulverized using a mortar and pestle and powder equivalent to 250 mg azithromycin was transferred into a 50 ml volumetric flask and dissolved in 25 ml acetonitrile under sonication for 15 min before making up to volume with distilled water. The samples were filtered through a 0.45 μm membrane filter before chromatography.

Suspensions equivalent to 40 mg/ml were weighed into 5 ml volumetric flasks and dissolved in 2.5 ml of acetonitrile under sonication for 15 min before making up to volume with distilled water. The sample was filtered through a 0.45 μm membrane filter before chromatographic analysis. Azithromycin bulk sample was prepared by dissolving 50 mg in 25 ml of acetonitrile under sonication before diluting to 50 ml with distilled water. The test and standard solutions were run under the optimum HPLC conditions and the peak areas obtained were used for the determination of the content of AZT in the samples.

2.7. Forced degradation and stability-indicating study

Solutions of azithromycin working standard were subjected to stress testing under oxidative and acidic conditions. The stability indicating ability of the method was tested under these two conditions.

2.7.1. Oxidative degradation of azithromycin

Oxidation was carried out using 0.0005% v/v H_2O_2 as described by Abuga *et al.* for clarithromycin [18] and the solutions sampled at defined intervals for HPLC analysis. For this purpose, azithromycin 5 mg/ml solution was incubated in 0.0005% v/v H_2O_2 at 37 °C and the peak areas of AZT were monitored over a period of 7 h. To determine the order of reaction, the data was fitted into zero, first and second order kinetic models whereby the r^2 value of the linear plots was used as a measure of goodness of fit.

2.7.2. Degradation of azithromycin in acid

Acid degradation was carried out by incubating a 5 mg/ml solution of AZT in 0.1M H_3PO_4 (pH 1.60), 0.05M H_3PO_4 (pH 1.68), 0.025M H_3PO_4 (pH 1.92) and 0.01M H_3PO_4 (pH 2.37), respectively, at 37 °C using the procedure previously described in literature [18]. The peak area of AZT was monitored by HPLC under the optimized conditions.

3. Results and Discussion

3.1. Method development and optimization

The development of the HPLC method followed a systematic manipulation of the chromatographic factors of pH, temperature, organic modifier concentration and buffer concentration. The process involved selection of appropriate conditions and their optimization. These conditions included type of column packing, column dimensions, mobile phase composition and flow rate, oven temperature, sample amount and detection wavelength. A detection wavelength of 215 nm was selected on account of better baseline stability while maintaining acceptable sensitivity.

The UV spectra obtained for the impurities used in this study did not show appreciable absorption differences. Acetonitrile was selected as organic modifier because it is commonly available in the market, has a higher eluting power and a lower UV cut-off compared to methanol. A Waters XTerra® hybrid, reverse-phase C18 column (250 mm \times 4.6 mm, 5 μm) was chosen for this study since it gave the highest efficiency compared to Phenomenex Luna® and Phenomenex Gemini® of similar dimensions.

The effect of adjusting pH of the phosphate buffer on the separation and symmetry of components was investigated using mobile phases at pH 5.0, 6.0, 6.5 and 7.0. A plot of capacity factor versus pH (Figure 2) revealed that an increase in buffer pH improved selectivity and also increased retention times of all the components. Furthermore, improved peak shape was observed with increase in pH. However at pH 7.0 there was co-elution of AZT and EAIE while at pH 6.0 there was poor resolution between AZT and NDMAZT. Through several experiments AZT and EAIE was identified as the critical peak pair (CPP) and was used as an indicator of method performance.

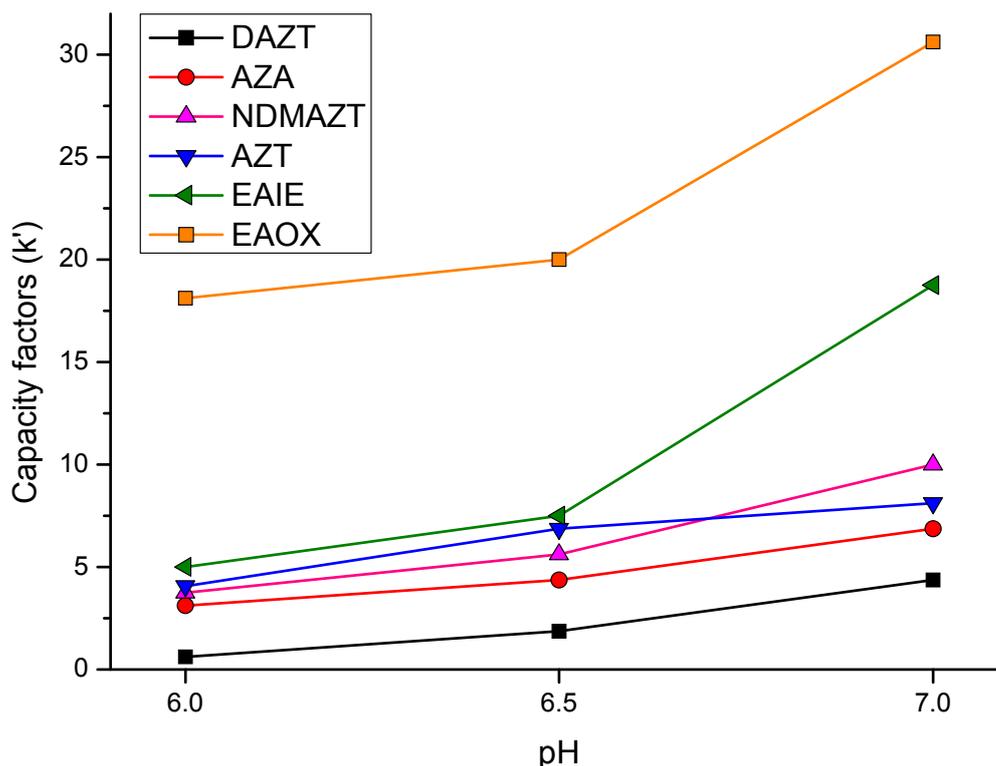


Figure 2. Effect of chromatographic factors of buffer pH on capacity factors of components of the working mixture.

To improve the separation between AZT and EAIE, the proportion of the organic modifier was reduced to 20% while maintaining pH 6.5. Under these chromatographic conditions, partial resolution was achieved although the run time increased from 60 min to 100 min. To achieve baseline resolution, the incorporation of tetrabutylammonium hydroxide at a concentration of 1% led to shorter run time (45 min) but with decreased resolution of decladinosylazithromycin (DAZT) with the solvent front. To improve the resolution and peak symmetry factor of DAZT, the effect of buffer concentration was investigated in the range 10-25% at 5% intervals. A buffer concentration of 15% was found to be optimal. In order to reduce the run time with baseline resolution of all components, the concentration of acetonitrile and temperature were optimized to 25% v/v and 43 °C respectively. Under these conditions, the peaks of all separated components had peak purities of unity.

The optimum chromatographic conditions for separation of all components were established as mobile phase comprising of acetonitrile-0.1M KH₂PO₄ pH 6.5-0.1M tetrabutyl ammonium hydroxide pH 6.5-water (25:15:1:59% v/v/v/v). Figure 3 shows a typical chromatogram obtained under the optimum conditions.

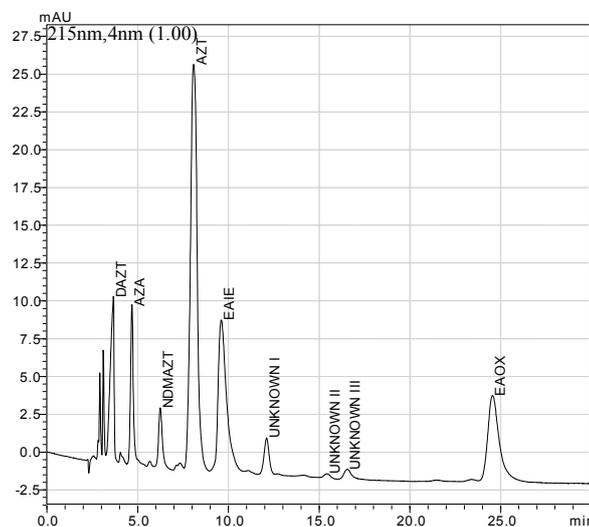


Figure 3. Typical chromatogram of the working mixture obtained under optimized chromatographic conditions.

3.2. Method validation

3.2.1. Linearity

The linear regression equation was $y = 9948x - 40457$ with a coefficient of determination (r^2) of 0.997, where y = peak area and x = concentration of solution. The coefficient of variation (CV) of the peak areas obtained ranged between 0.02-0.12%. The developed method therefore met the ICH guidelines for linearity over the concentration range investigated [17].

3.2.2. Sensitivity

When calculated against a sample solution of 5.0 mg/ml, the LOD was 0.02% (20 μ g) ($n = 6$) at a S/N ratio of 3 while the LOQ was 0.078% (78 μ g) ($n = 6$) at a S/N ratio of 10. The LOD and LOQ results obtained demonstrated that the developed method was adequately sensitive for the determination of AZT and its related substances at the 0.1% level [17,19].

3.2.3. Accuracy

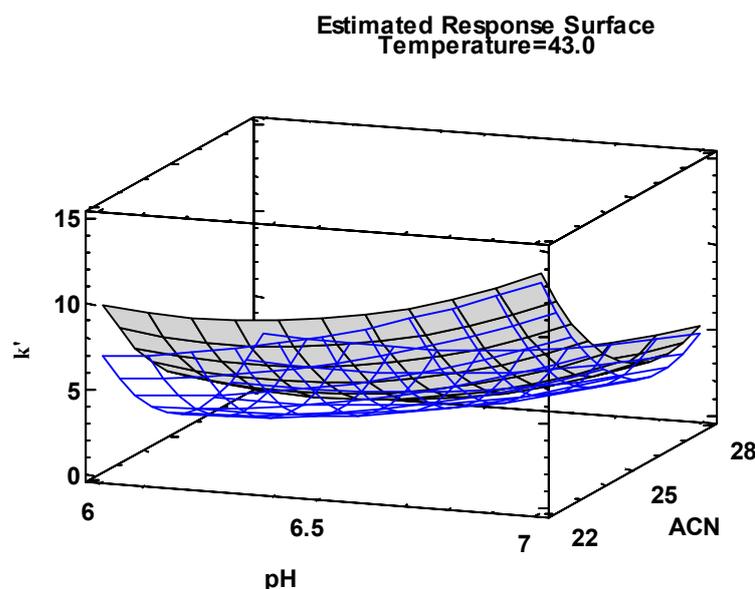
The results for the recovery of azithromycin from spiked samples are shown in Table 2. The ICH guidelines for the validation of analytical procedures specify a mean recovery of $100 \pm 2.0\%$ at each concentration over the range of 80-120% of nominal concentration [17]. The results shown in Table 2 indicate that the developed method is accurate with a mean recovery of 100.7%.

Table 2. Recovery of azithromycin from spiked samples. CV; coefficient of variation

Target concentration (%)	Azithromycin				Mean recovery (%)
	Amount added (mg)	Amount recovered (mg)	Recovery (Absolute %)	Recovery (CV %)	
80	4.98	4.975	99.9	0.16	100.7
100	5.08	5.171	101.8	0.20	
120	5.13	5.152	100.4	0.50	

3.2.4. Robustness

The robustness ranges for pH were found to be pH 6-7, 22-28% v/v acetonitrile and 41-45 °C for column temperature. Pareto analysis revealed that acetonitrile concentration and buffer pH had the biggest impact on the separation of AZT and EAIE ($p = 0.05$). The response surface plot (Figure 4) showed non-intersection of response surface plots at all nominal values investigated indicating that there was no co-elution of this critical peak.

**Figure 4.** Overlaid response surfaces of AZT and EAIE.

3.3. Forced degradation of azithromycin in acidic and oxidative environments

The data generated from oxidative degradation of azithromycin fitted into the second order rate equation with linear regression equation $y = 0.0106x + 0.1377$ ($r^2 = 0.994$), a half life of 13 min and a rate constant (k) of 1.06×10^{-2} area units $^{-1}$ min $^{-1}$. This oxidative study may be indicative of the possible nature of degradation of AZT in the presence of oxidants and exposure to air during storage.

Acid degradation data showed that AZT degrades rapidly by about 90% within 10 min in 0.1M H₃PO₄ and 0.05M H₃PO₄ solutions maintained at 37 °C. However at 0.025M H₃PO₄ at 37 °C, the kinetics of AZT decomposition were derivable. The data generated fitted into the first order rate equation with linear regression equation $y = -0.0002x + 13.651$ ($r^2 = 0.9907$), a half life of 57.8 h (3465

min) and a rate constant (k) of $2.0 \times 10^{-4} \text{ min}^{-1}$. In all instances, the degradation products in acid were well separated from AZT peak thus providing further evidence on the stability-indicating ability of the method. The degradation of AZT under acidic conditions is likely to affect oral bioavailability and efficacy of the drug due to gastric acid. For instance, with a mean gastric residence time for AZT base as low as 10 min in gastric fluid of pH 1.6, 93% of the administered dose may be lost due to degradation. The degradation of AZT in gastric fluid may differ since there is a significant concentration gradient of the drug from gastric tissue to gastric juice [20].

3.4. Assay of azithromycin tablets and suspensions

The developed and validated liquid chromatographic method was applied for the assay of azithromycin bulk samples and the oral dosage formulations of tablets and suspensions. The product excipients did not interfere with the determination of AZT in all the formulations analysed. The peak purity index of AZT was confirmed to be unity by photo diode array analysis.

The USP (2102) limits for assay (90.0-110.0% label claim) were adopted for quality assessment of the samples[13]. The assay test failure rates were 23% (n=13) for tablet brands, 60% (n=5) for premixed suspensions and 25% (n=4) for dry powders (Table 3). There was a relatively higher assay test failure rate for premixed suspensions compared to dry powders for reconstitution. These findings underscore the need for generic manufacturers to carry out comprehensive formulation studies to justify any formulation aspects that significantly differ from those of the innovator products. The Drug Regulatory Authorities ought to evaluate supporting data for products that deviate from the innovator formulation to ensure that generics registered are pharmaceutically equivalent and bioequivalent to the innovator brand. Where significant changes in formulation occur, the generic manufacturer should carry out full clinical studies to support the rationale for such deviation [21].

The assay test failure rates observed underscore the need for continuous market surveillance on products available in the Kenyan market to ensure circulation of good quality medicines. This will help curb the emergence of macrolide-resistant strains thus preserving the antibiotic arsenal to treat susceptible microorganisms. Batch to batch variation of AZT content was not observed where different batches of the same product were analysed (Table 3).

Table 3. Assay results for selected commercial samples of azithromycin. The figures in parentheses represent coefficient of variation; P, Powder for reconstitution; M, Premixed; a and b represent different batches of each product

Product code	Formulation	AZT content (%)	Remarks
I	a) Tablet	93.6 (1.6)	Complies
	b) Tablet	94.8 (1.0)	Complies
II	a) Tablet	110.3 (0.2)	Complies
	b) Tablet	109.1 (0.1)	Complies
III	a) Tablet	98.5 (0.5)	Complies
	b) Tablet	99.2 (0.3)	Complies
IV	a) Tablet	97.2 (0.2)	Complies
	b) Tablet	97.6 (0.2)	Complies

V	Tablet	102.0 (1.5)	Complies
VI	a) Tablet	109.9 (0.3)	Complies
	b) Tablet	107.3 (0.3)	Complies
VII	a) Tablet	105.1 (1.6)	Complies
	b) Tablet	104.3 (1.3)	Complies
VIII	a) Suspension (M)	110.0 (0.6)	Complies
	b) Suspension (M)	109.3 (0.5)	Complies
IX	a) Suspension (M)	96.4 (0.7)	Complies
	b) Suspension (M)	96.1 (0.2)	Complies
X	Suspension (M)	69.3 (1.7)	Did not comply
XI	Suspension (M)	89.3 (0.4)	Did not comply
XII	Tablet	81.4 (0.5)	Did not comply
XIII	Tablet	99.8 (0.3)	Complies
XIV	Tablet	83.3 (0.2)	Did not comply
XV	Tablet	87.0 (1.1)	Did not comply
XVI	Tablet	103.3 (1.3)	Complies
XVII	Suspension (P)	88.6 (0.2)	Did not comply
XVIII	Suspension (P)	105.5 (0.4)	Complies
XIX	Suspension (M)	87.8 (1.3)	Did not comply
XX	Suspension (P)	98.7 (0.7)	Complies
Bulk sample	Powder	94.7 (0.1)	Complies
Zithromax®	Suspension (P)	99.6 (0.7)	Complies
Zithromax®	Tablet	96.4 (1.5)	Complies

5. Conclusions

A selective, simple, specific, robust and isocratic liquid chromatographic method for the analysis of azithromycin in bulk samples and oral dosage forms was developed. The method met ICH guidelines for validation parameters. With a run time of 25 min, the method allows a relatively high sample throughput. Therefore, the method may be employed in the routine laboratory analysis of samples and stability studies. The presence of products that do not meet assay specifications in the market is a matter of public health concern and it underscores the need for sustained post market surveillance by the Drug Regulatory Authority (DRA). This study did not encounter any counterfeits of AZT in the market.

Supplementary Materials: None

Acknowledgments: The authors wish to thank the management of the National Quality Control Laboratory for granting permission to use equipment, as well as the Drug Analysis and Research Unit of the University of Nairobi for logistical support and providing instruments to undertake this research.

Author Contributions: AOO, KOA, FNK and SNN conceived the study. AOO, KOA and DWL helped to draft the manuscript, organized data and revised the final version of the manuscript. All authors contributed to the interpretation of findings, read and approved the final manuscript.

Conflicts of Interest: The authors declare no conflict of interest.

References

1. Piscitelli, S. C.; Danziger, L. H.; Rodvold, K. A. Clarithromycin and azithromycin: new macrolide antibiotics. *Clin. Pharm.* **1992**, *11*, 137–52.
2. Stott, G. A. New macrolide antibiotics: Clarithromycin and azithromycin. *Am. Fam. Physician* **1992**, *46*, 863–869.
3. World Health Organization (WHO) *WHO Model Formulary 2008*; World Health Organization: Geneva, Switzerland., 2009.
4. Ministry of Health *Kenya Essential Medicines List 2010*; Ministry of Health: Nairobi, Kenya, 2016.
5. Breier, A. R.; Garcia, C. V.; Oppe, T. P.; Steppe, M.; Schapoval, E. E. S. Microbiological assay for azithromycin in pharmaceutical formulations. *J. Pharm. Biomed. Anal.* **2002**, *29*, 957–961.
6. Nigović, B.; Šimunić, B. Voltammetric assay of azithromycin in pharmaceutical dosage forms. *J. Pharm. Biomed. Anal.* **2003**, *32*, 197–202.
7. Walsh, M. I.; Rizk, M. S.; Eid, M. I.; Fathy, M. E. Spectrophotometric determination of four macrolide antibiotics in pharmaceutical formulations and biological fluids via binary complex formation with eosin and spectrophotometry. *J. AOAC Int.* **2007**, *90*, 1579–1587.
8. Zhang, S.; Huang, X.; Yao, N.; Horváth, C. Preparation of monodisperse porous polymethacrylate microspheres and their application in the capillary electrochromatography of macrolide antibiotics. *J. Chromatogr. A* **2002**, *948*, 193–201.
9. Khedr, A.; Sheha, M. Quantitative thin-Layer chromatographic method of analysis of azithromycin in pure and capsule forms. *J. Chromatogr. Sci.* **2003**, *41*, 10–16.
10. Kamau, F. N.; Chepkwony, H. K.; Ngugi, J. K.; Debremaeker, D.; Roets, E.; Hoogmartens, J. Isocratic liquid chromatographic method for the analysis of azithromycin and its structurally related substances in bulk samples. *J. Chromatogr. Sci.* **2002**, *40*, 529–533.
11. Debremaeker, D.; Visky, D.; Chepkwony, H. K.; Van Schepdael, A.; Roets, E.; Hoogmartens, J. Analysis of unknown compounds in azithromycin bulk samples with liquid chromatography coupled to ion trap mass spectrometry. *Rapid Commun. Mass Spectrom.* **2003**, *17*, 342–350.
12. *British Pharmacopoeia 2015*; The Stationery Office Ltd: Norwich, UK, 2014.
13. *United States Pharmacopoeia 38: The National Formulary 33*; United States Pharmacopoeial Convention: Rockville MD, USA, 2015.
14. Sachin, K. S.; Prakash, D.; Tamizh, M. T. Development and validation of stability indicating UV spectrophotometric method for the estimation of azithromycin suspension. *Int. J. ChemTech Res.* **2010**, *2*, 1939–1944.
15. Stahl, M. Peak purity analysis in HPLC and CE using diode-array technology. Available online: <http://www.agilent.com/cs/library/applications/5988-8647EN.pdf> (Accessed on 24 April 2016).
16. Wiberg, K.; Andersson, M.; Hagman, A.; Jacobsson, S. P. Peak purity determination with principal component analysis of high-performance liquid chromatography-diode array detection data. *J. Chromatogr. A* **2004**, *1029*, 13–20.
17. ICH Q2 (R1) Validation of Analytical Procedures: Text and Methodology. Available online: http://www.ich.org/fileadmin/Public_Web_Site/ICH_Products/Guidelines/Quality/Q2_R1/Step4/Q2_R1_Guideline.pdf (Accessed on 10 April 2016).
18. Abuga, K. O.; Chepkwony, H. K.; Roets, E.; Hoogmartens, J. A Stability-indicating HPLC method for the separation of clarithromycin and its related substances in bulk Samples. *J. Sep. Sci.* **2001**, *24*, 849–855.

19. ICH Q6A Specifications: Test Procedures and Acceptance Criteria for New Drug Substance and New Drug Products: Chemical Substances. Available online: <http://www.fda.gov/Drugs/GuidanceComplianceRegulatoryInformation/Guidances/ucm134966.htm> (Accessed on 10 April 2016).

20. Harrison, J. D.; Jones, J. A.; Morris, D. L. Azithromycin levels in plasma and gastric tissue, juice and mucus. *Eur. J. Clin. Microbiol. Infect. Dis.* **1991**, *10*, 862–4.

21. FDA Guidance for Industry Bioavailability and Bioequivalence Studies for Orally Administered Drug Products - General Guidance for Industry Bioavailability and Bioequivalence. Available online: <http://www.fda.gov/downloads/Drugs/DevelopmentApprovalProcess/HowDrugsareDevelopedandApproved/ApprovalApplications/AbbreviatedNewDrugApplicationANDAGenerics/UCM154838.pdf> (Accessed 12 April 2016).



© 2016 by the authors; licensee *Preprints*, Basel, Switzerland. This article is an open access article distributed under the terms and conditions of the Creative Commons by Attribution (CC-BY) license (<http://creativecommons.org/licenses/by/4.0/>).