A Modified and Credible Methods to Estimate Nitrofurantoin In the Standard of Substances and Pharmaceutical Dosage

KHAWLA SALMAN ABD ALRASSOL¹, QUTAIBA A. QASIM², GHASSAN SALAH AHMED³, H. N. K. AL-SALMAN^{*4}

^{1,4}Department Pharmaceutical Chemistry, College of Pharmacy, University of Basrah, Iraq
²Department of Clinical Laboratory Sciences, Collage of pharmacy, Basrah University, Iraq
³Department of Clinical Pharmacy, Collage of pharmacy, Basrah University, Iraq
*Corresponding Author
Email ID: hsennaserh@yahoo.com
Received: 22.09.19, Revised: 02.10.19, Accepted: 29.11.19

ABSTRACT

For identifying the nitrofurantoin drug, the four selective, sensitive and simple methods were developed. And these methods are then proved as well as validated in our particular research work. These methods are dependent of the nitrofurantoin reactions, performed by utilizing ZN/CI, as well as iron mixture (II)and neutral medium is used for ferric chloride reduction through this drug along with I, method A- I0- phenanthroline or 2, method B- bisperdyl or blue chromogen is formed when this particular drug binds with the oxidized ferric chloride and potassium ferritic cyanide reagent(method D). A colourful product is produced with the ninhydrin and nitrofurantoin reagent interaction and also method D is also dependent on this. By using method A, B, C and D the measurement of resulted red chromosomes is found to 500nm, 515nm, 735nm, and 575nmrespectively.Method A, B, C and D uses the concentration ranges of 0.20-8.0 mg/ml 0.25-40 mg/ml, 0.50-30 mg/ml and 0.50-50 mg/ml respectively and in such optimal conditionsBers law is applied along with molar's absorption values are also estimated.The statistical comparability of the suggested methods resulted with all those acquired by the reference technique which proved outstanding agreement as well as also shown there does not exist any interference through typical excipients in pharmaceutical formulations.

Keywords: Chelating agents, Ferric chloride, Nitrofurination drug.

INTRODUCTION

One of the nitrofuran's drug derivative namely nitrofurantoin (NTF) is essential during the disease of urinary area and possesses a chemical structurenamely (1-((5-nitro-2-furfurylidene)-1amino) hydantoin) and is presented in Figure 1. Additionally, a few gram-positive organisms likecorynebacterium, viridians streptococci, group D streptococci,S. agalactiae,enterococcus faecalis,S. saprophyticus,S. epidermidis,andS. aureus induces activity in NTF.

The activity spectrum for gram negative organisms that are shigella, salmonella, Neisseria,Enterobacter, andE. coli, [1,2].Furthermore, from the bacterial resistancedevelopment this drug is highly stable and this characteristic is result of the mechanismswork multiplicity [3].

In addition to, it would once deal with bladder infections through antibiotic that is provided through theoral cavity. Therefore, it is essential for preventing as well as treatingthe infections of urinary tract infection by opposing the bacteria's growth. Although, in case of kidney infections the drug is inactive [4].

In the pure drug and pharmaceutical preparation the nitrofurantoinevaluation is registered by some of the analytical techniques. Furthermore, the drug can be estimated in the form of tablet dosage through the HPLC technique, additionally, spectrophotometric method verified that evaluation of the nitrofurantoinelectroanalytical techniquewas posted to drugdetermination [5,6].