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Batch and flow injection spectrophotometric methods for determination of Ceftazidime in pharmaceutical formulations

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Abstract:

It is generally accepted that there are two spectrophotometric techniques for quantifying ceftazidime (CFT) in bulk medications and pharmaceutical formulations. The methods are described as simple, sensitive, selective, accurate and efficient techniques. The first method used an alkaline medium to convert ceftazidime to its diazonium salt, which is then combined with the 1-Naphthol (1-NPT) and 2-Naphthol (2-NPT) reagents. The azo dye that was produced brown and red in color with absorption intensities of λ_{\max} 585 and 545nm respectively. Beer's law was followed in terms of concentration ranging from (3-40) $\mu\text{g} \cdot \text{ml}^{-1}$ For (CFT-1-NPT) and (CFT-2-NPT), the detection limits were 1.0096 and 0.8017 $\mu\text{g} \cdot \text{ml}^{-1}$, respectively, and the molar absorptivity was 0.7926×10^4 and $0.5466 \times 10^4 \text{ L} \cdot \text{mol}^{-1} \cdot \text{cm}^{-1}$. The Flow Injection Analysis (FIA) method is used to estimate ceftazidime and in the second procedure record measurements using the UV-Visible approach. The Flow injection allows for exact drug estimation under ideal experimental conditions. The concentrations were in the range of (3-50) $\mu\text{g} \cdot \text{ml}^{-1}$ For (CFT-1-NPT) and (CFT-2-NPT), the detection limits were 0.8102, 1.2809 $\mu\text{g} \cdot \text{ml}^{-1}$, and the molar absorptivity was 0.9565×10^4 , $0.7106 \times 10^4 \text{ L} \cdot \text{mol}^{-1} \cdot \text{cm}^{-1}$, respectively. The proposed two methods for determination Ceftazidime in Pharmaceutical formulation were successfully applied, as these methods were characterized by simplicity, speed, accuracy, and low cost.

Keywords: 1-Naphthol , 2-Naphthol Spectrophotometric, Azo – aye , Batch , Ceftazidime , Diazotization, Flow injection.

Introduction

Antibiotics are substances or compounds that kill or inhibit the growth of bacteria¹. And the antibiotics are antimicrobial substances used to treat infections caused by bacteria, fungus, and parasites.². Ceftazidime is a third-generation semi-synthetic cephalosporin that inhibits Gram-negative and Gram-positive bacteria, including pseudomonas aeruginosa.³ Chemically, Ceftazidime (CFT) is [7-(2-(2-aminothiazol-4-yl)-2-(6R,7R,Z)-carboxypropan-2-yl)oxymino)acetamido]-8-oxo-3-(pyridinium-1-ylmethyl)-5-thia-1-azabicyclo[4.2.0]oct-ene-2-carboxylate .Fig .1

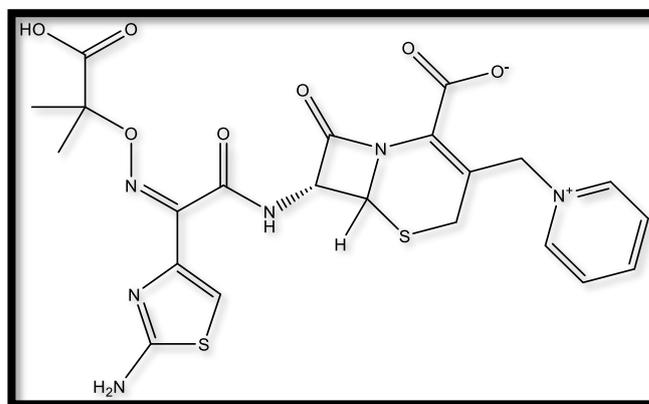


Figure 1. Structural formula for ceftazidime

(Fortaz, Tazicef)⁴ is the trade name. • It is used to treat infections caused by Pseudomonas aeruginosa,

which include infections of the bones and joints, cystic fibrosis (respiratory tract infections), and cystic fibrosis gastroenteritis (intestinal infections) ^{5,6}

Many researches have been conducted to estimate the presence of this medication in pharmaceutical formulations such as : Two Oxidative Visible Spectrophotometric Methods⁷ RP-UPLC Method ⁸ Poly Diphenylamine Partially Oxidized for Sensitive Electrochemical Applications ⁹ Using resorcinol and 2-methyl phenol, spectrophotometric detection of ceftazidime in pharmaceuticals ¹⁰ The utility of a fluorescamine-based technique for very sensitive spectrofluorimetric measurement of Ceftazidime and Vancomycin in pharmaceuticals and real human plasma was investigated using flow injection analysis (FIA). ¹¹ Flow injection analysis is one of many analytical chemistry techniques that have been widely embraced and adopted in a variety of analytical fields. Diazotization and coupling processes have been widely used to determine a wide range of pharmaceuticals and pharmaceutical preparations, including the determination of the ceftazidime compound employing the reaction between the diazotized drug and the coupling

reagent 4-tert-butylphenol ¹². Many papers have been written about diazotization reactions and their connection to the flow injection system ¹³⁻¹⁷. The proliferation of these domains is owing to the ease with which they can be linked or coupled with a variety of techniques such as chromatic Flame atomic absorption ^{18,19}, electrochemical detection ²⁰, chemical fluorescence ²¹. Through the (FIA) technology, which was characterized by high repetition and conformity, as well as its simplicity, speed, and low cost, the (FIA) technology has demonstrated its high efficiency. ^{22,23}. (FIA) is a quantitative method for assessing or estimating minerals and materials using injection flow such as Manganese in plants ²⁴, cholesterol ²⁵ and Mebeverine Hydrochloride²⁶ In this paper, new methods for determining ceftazidime in its pure and pharmaceutical formulations have been developed using diazotization and coupling reactions with 1-naphthol and 2-naphthol reagents, and then using these reactions with the flow injection technique in the presence of a visible and ultraviolet detector, and their applications in the scientific field to estimate these drugs in their pure and pharmaceutical formulations have been developed.

Flow injection configuration a three-channels manifold was employed with, Peristaltic pump (AILITEA, C4, made in Sweden) with polyvinyl chloride tube (0.8) internal diameter as in fig.2.

Materials and Methods

Apparatus;

All spectral and absorbance measurements were executed on an Advanced microprocessor UV-VIS spectrophotometer single beam LI-295 recording spectrophotometer using 1 cm quartz cells.

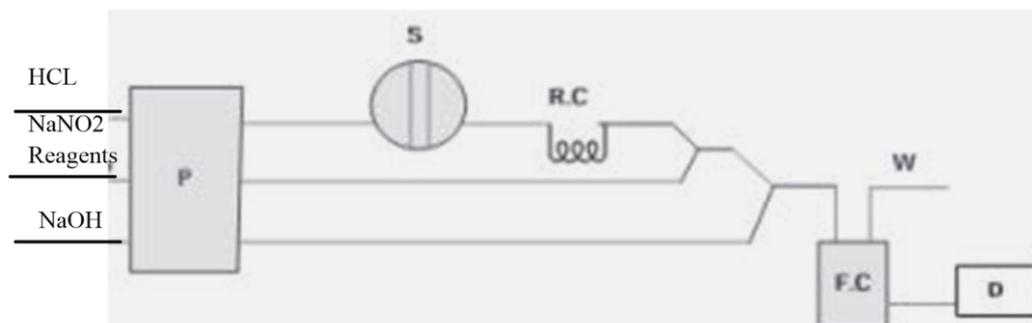


Figure 2. System of the employed flow system, P :peristaltic pump, R.C: reaction coil, S: sample injection ,W: waste, FC: flow ce:

Reagents;

Pharmaceutical preparations: Ceftazidime Pharmaceutical preparations were purchased from commercial sources, and Reagents were all analytical reagent grade chemicals.

Ceftazidime (CFT) (1000 µg/ml) ; (Laboratorios TORLAN Barcelona - Spain) A stock solution (1000 µg/ml) 9.1478×10^{-5} M of ceftazidime was prepared by dissolving 0.1 gm of CFT in distilled water. Then transfer it to a conical flask 100ml and complete the volume to the mark.

1-Naphthol (1-NPT); 1-Naphthol (50µg/ml) 3.471×10^{-4} M was prepared by dissolving 0.1 g (BDH) distilled water with a small amount of 25% NaOH were added to ensue complete dissolution . Then transfer it to a conical flask 50ml and complete the volume to the mark.

2-Naphthol (2-NPT): 2-Naphthol (50µg/ml) 3.471×10^{-4} M was prepared by dissolving 0.1 g (AR) distilled water with a small amount of 25% NaOH were added to ensue complete dissolution . Then transfer it to a conical flask 50ml and complete the volume to the mark.

Different concentrations of reagents from (0.0694×10^{-3} to 0.832×10^{-3}) M (300 -1200) µg/ml used in the injection method were prepared by drawing different volume (1-12) ml from 2000µg/ml for reagents completing the volume of the marker in a 20 ml conical flask .

Sodium nitrite NaNO₂ 1% (0.1449 M) 1g sodium nitrite (BDH) was dissolved in distilled water and diluted to the desired concentration in a 100 mL conical flask.

HCl Solution 1:1 (36.46 M) In a 100 ml conical flask, 50 mL of 11.64 M concentrated hydrochloric acid (ADR) was diluted with distilled water

NaOH 25% solution (6.2505 M) 25 g sodium hydroxide (BDH) was dissolved in distilled water and diluted to the desired concentration in a 100 mL conical flask.

Pharmaceutical Ceftazidime;

Commercially available pharmaceutical preparations were used

Ceftazidime Roth (Pharma Roth GmbH 65189 Wiesbaden Germany) , YENIZidime (yeni-pharma company Turkey –Ankara) and TOTTIZIM (Made in ITALY) (1000µg/ml) 0.1 g dissolved in distilled water and fill to the mark in a 100 ml conical flask

General procedure of diazotization reaction;

An aliquot of a sample solution containing CFT (1ml) from 1000µg/ml was transferred into 20.0 mL of conical flasks and then cooled in an ice bath maintained at -5.0 °C to this solution, 1ml of HCl followed by 1.0 mL from NaNO₂ 1%(w/v), were added to the mixture and left to stand for 10 min . Then, 1 mL of 1-naphthol and 2-naphthol was added to flask, then 1ml of NaOH 25% solution was added to the mixture, and diluted to the mark with double distilled water(D.W). The azo dye that formed was monitored at λ_{max} 585 and 545nm respectively

General procedure of Flow Injection method;

Using a three-channels peristaltic pump, the first channel contains a mixture of hydrochloric acid and sodium nitrite This mixture is passed through the injection valve, containing 100 µL of ceftazidime, to form the diazonium ion, which is passed into a 100 cm reaction coil. the second channel contains the reagent (1-NPT or 2-NPT) that coupled with diazonium ion to form diazonium salt, the third channel contains 5% of NaOH which is added to the product, here the colored product (Brown and red respectively) appears which will pass to the UV-Visible reagent for absorption and is measured the resulting brown and red colors were measured at λ_{max} 585 and 545 nm for the two detectors, respectively.

Result and discussion

Absorption spectra

Diazotized CFT was coupled with 1-Naphthol and 2-Naphthol in an alkaline solution to produce brown and red colored chromophore. The absorbance of the azo dyes were recorded at wavelengths of 585 and 545nm against a blank solution that which is prepared in the same way as sample – the same additions - except for drugs, and were obtained with a maximum absorbance under optimal conditions. Figs. 3, 4.

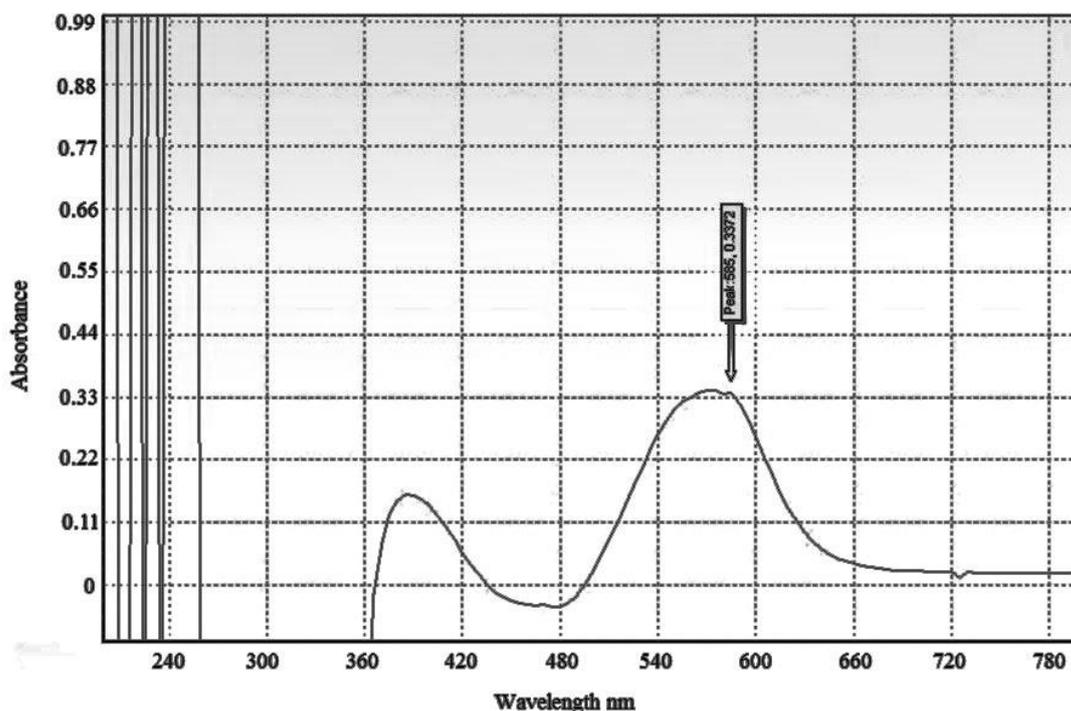


Figure 3. Absorption spectrum of 40ppm of CFT-1-NPT against reagent blank.

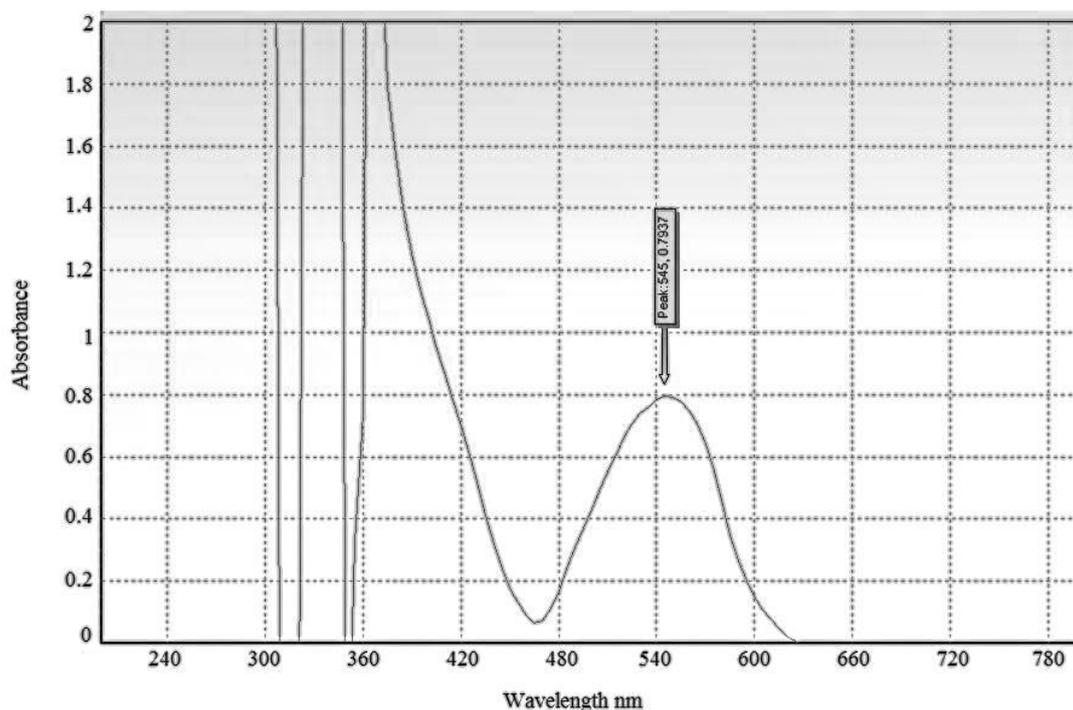


Figure 4. Absorption spectrum for 70ppm of CFT-2-NPT against reagent blank.

Part 1: Optimization of experimental conditions

To achieve the optimum detection limit and highest sensitivity, the many aspects impacting the spectrum characteristics of the dyes generated by

coupling nitrogenous drug with organic reagent were examined. All of these tests were carried out in 20ml volumetric flasks with (1000 μ g/ml) 9.1478×10^{-5} M of CFT standard solution.

The effect of the acid type on the diazonium salt:
Acids such as HCL, H3PO4, HNO3, and H2SO4 diluted (1:1) have been examined. The results demonstrate that HCl (1:1) is the best acid for this procedure because it offers the maximum

absorbance signal for both dyes. Although H2SO4 in reagent 2-NPT gave the highest absorbance, the solution was turbid, thus HCl was employed, as shown in Table 1.

Table .1 Effect of acid type on absorbance signal of CFT.

Type of acid (1:1)	Abs of CFT-1-NPT at 585nm	Abs of CFT-2-NPT at 545nm
HCL	0.864	0.984
HNO3	0.180	1.198
H2SO4	0.089	0.339
H3PO4	0.220	0.284

Effect of the hydrochloric acid volume on diazonium salt reaction;

Various volumes of HCl ranging from 0.5ml to 2ml were used to investigate the effect of acidity on the

colored solution. Although 1.5 and 2ml in reagent 2-NPT had the maximum absorbance, the solution was unstable and contained a precipitate, thus 1ml was chosen instead, Fig.5.

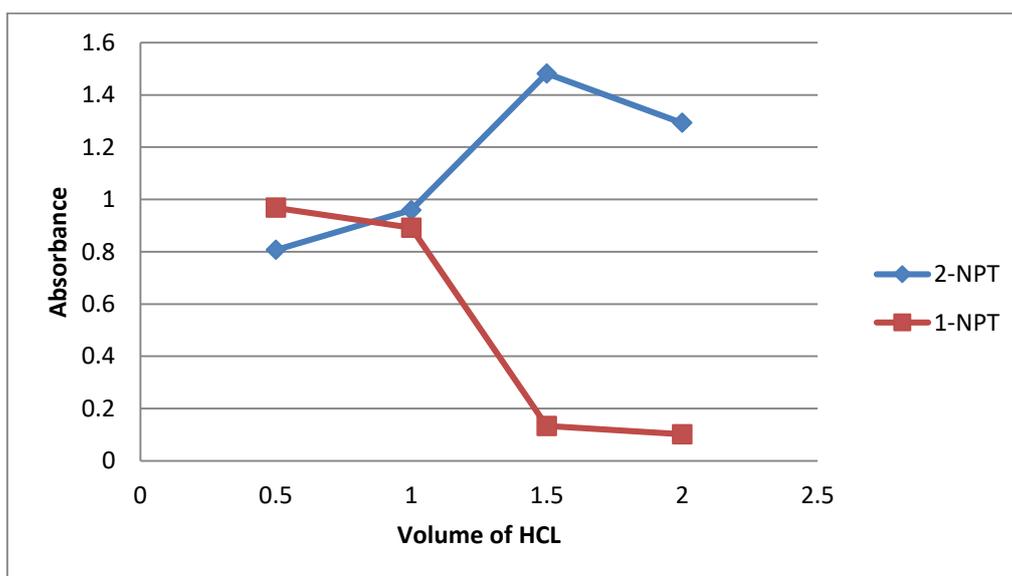


Figure 5. Effect volume of (1:1) HCL

Effect of the NaNO₂ 1% volume

The effect of the volume of the NaNO₂ solution was investigated by using various volumes of NaNO₂

ranging from 0.5 to 2ml, with the result that 0.5 showed better absorption for both reagents Fig.6.

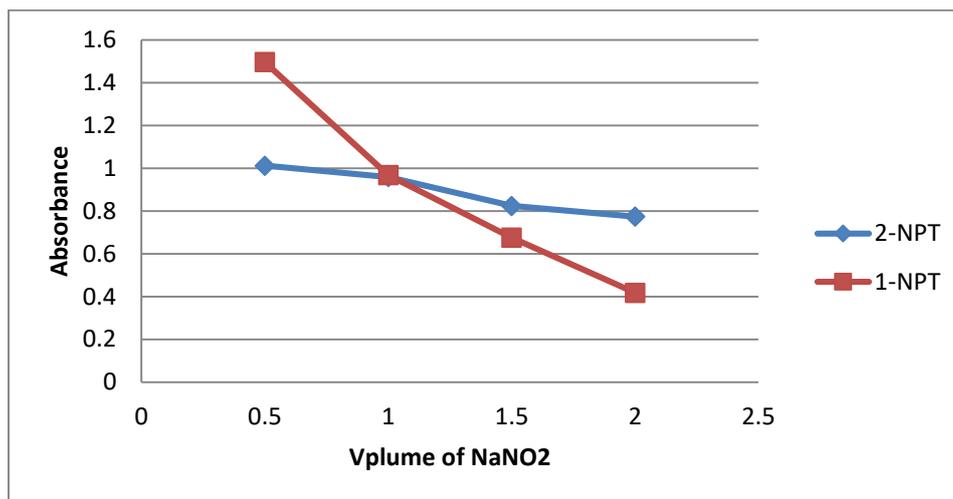


Figure 6. Effect volume of 1% NaNO₂

Effect of reaction time after addition of NaNO₂
The effect of reaction time was investigated by utilizing different intervals ranging from 5 to 30

minutes, 15 min gave highest absorption for CFT-1-NPT and CFT-2-NPT Fig.7.

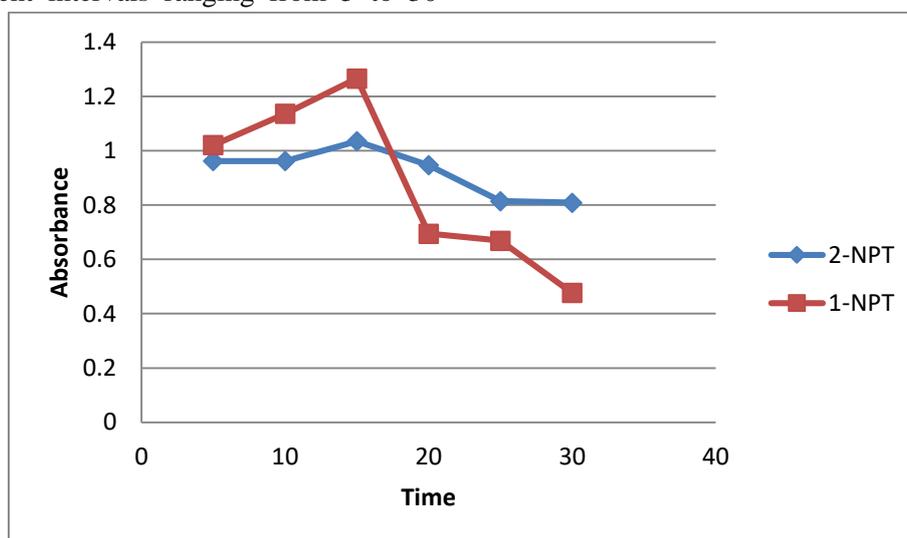


Figure 7. Effect time after the addition of NaNO₂

The impact of different types of bases on absorption;

The effect of different bases, 25 percent w/v [KOH, NaOH, Na₂CO₃], on the absorption of azo dye has been tested, and it was found that the KOH solution

showed the highest absorption for both the reagent and the CFT-2-NPT, but the solution was unstable and continuation of bubbles, therefore NaOH was used. as shown in Table 2.

Table 2. Effect type of base on the diazonium salt reaction

Type of base 25%	Absorbance of CFT with	
	1-NPT at 585nm	2-NPT at 545nm
NaOH	1.496	1.010
KOH	1.487	1.428
Na ₂ CO ₃	0.047	1.318

Effect of the sodium hydroxide volume;
The effect of varied sodium hydroxide volumes on absorbance were tested for the volume range (0.5-

2ml), with 1,0.5ml gave the maximum absorption for CFT-1-NPT and CFT-2-NPT, respectively, Fig.8.

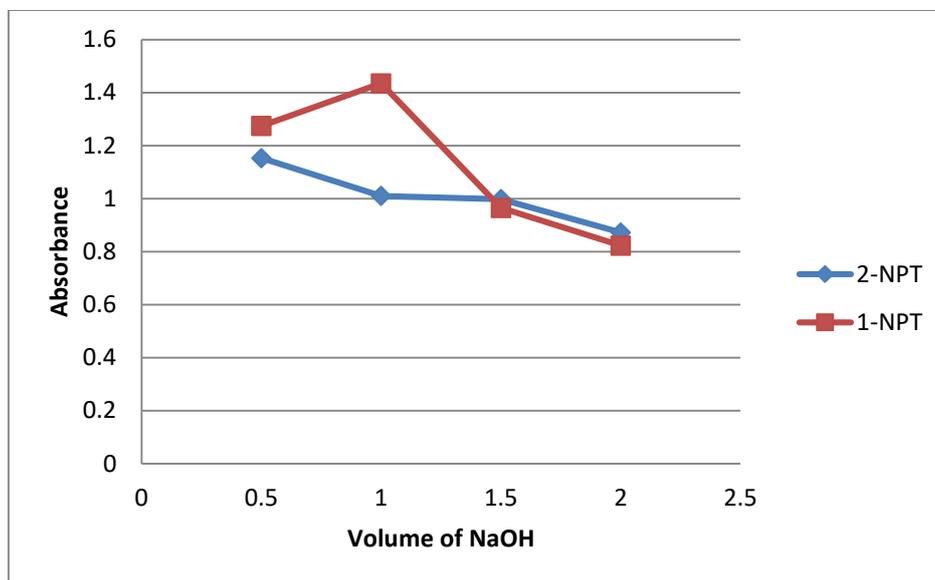


Figure 8. Effect of the NaOH 25%

The Effect of reagent volume on diazonium salt:
With 1 ml of 1000µg/ml ceftazidime solution, several quantities of (50µg/ml) 3.4710^{-4} M of reagents 1-NPT and 2-NPT were examined (0.5-

1.00ml). The reagent with the highest absorbance was 0.5ml. The absorbance signal remained steady after that. Fig.9.

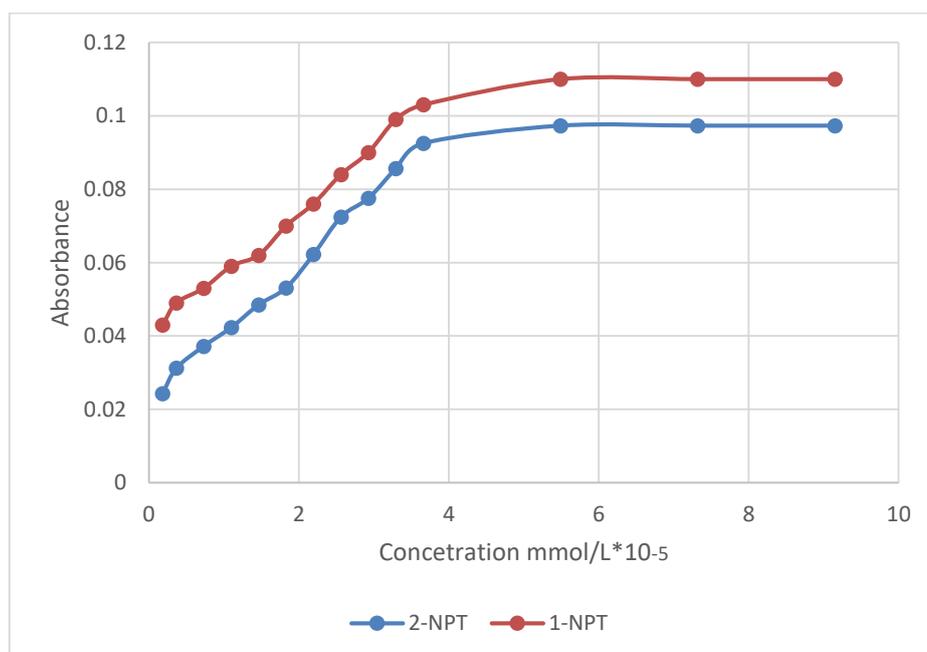


Figure 9. The mole-ratio plot for diazotized ceftazidime to 1-NPT and 2-NPT

The possible reaction mechanism can be written as in the Fig.10.

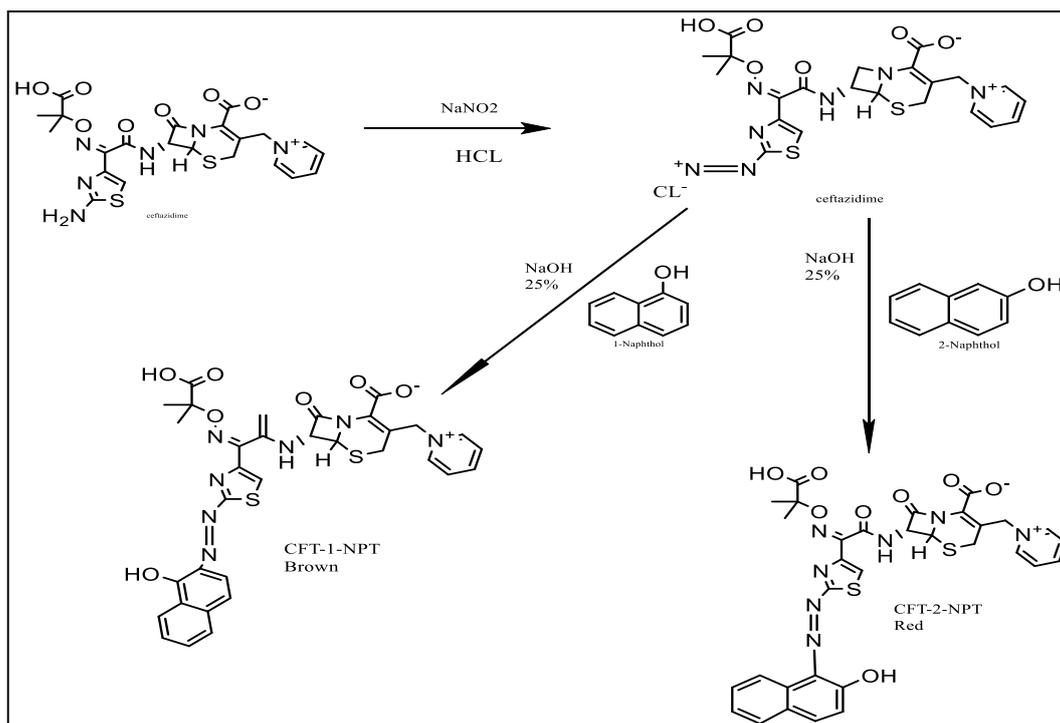


Figure 10. The Diazotize salt reaction's proposed mechanism

Data analysis

The absorption signals were measured against a series of ceftazidime concentrations to plot a

calibration curve. Fig 11 and Table 3 exhibit the developed method's calibration curve and analytical figures.

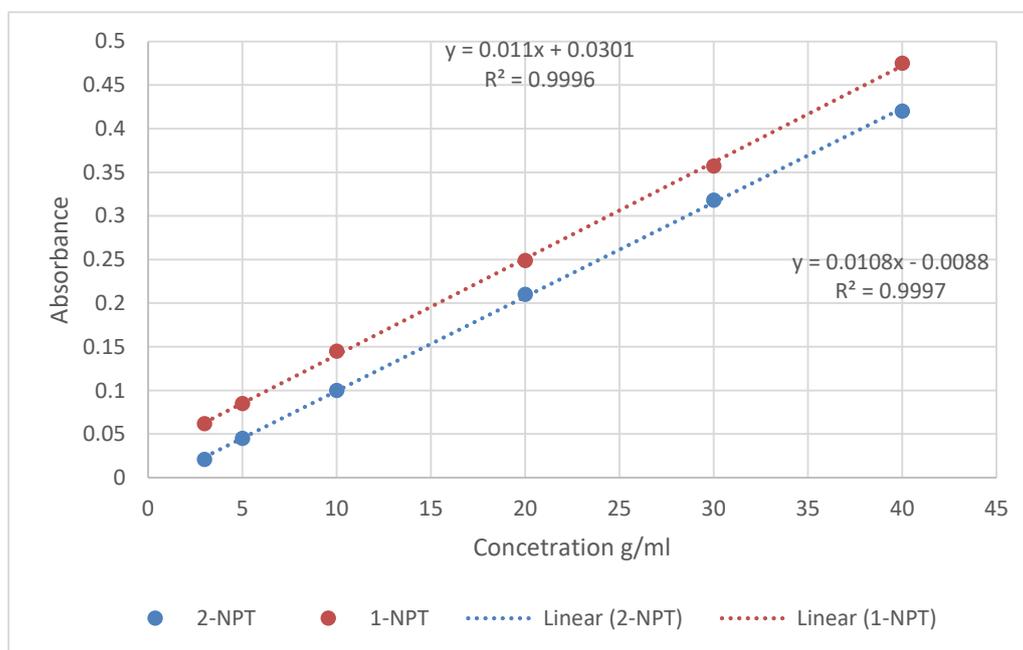


Figure 11. Calibration curve of CFT-1-NPT and CFT-2-NPT

Table 3.The proposed diazonium salt techniques' distinctive parameters.

Parameter	CFT-1-NPT	CFT-2-NPT
Color of product	Brown	Red
λ max (nm)	585	545
Linear range ($\mu\text{g.ml}^{-1}$)	(3-40)	(3-40)
Molar absorptivity, ϵ ($\text{L.mol}^{-1}.\text{cm}^{-1}$)	0.7926×10^4	0.5466×10^4
Regression equation	$y=0.011x+.0301$	$y=0.0108x-0.0088$
Sandell sensitivity, $S (\mu\text{g} .\text{cm}^{-2})/0.001\text{A.U}$	0.0689	0.0952
Intercept (a)	+0.0301	-0.0088
Slope (b) ($\text{L.mg}^{-1}.\text{cm}^{-1}$)	0.011	0.0108
Coefficient of determination % R^2	99.96	99.97
Correlation coefficient (r)	0.9997	0.9998
Limit of detection ($\mu\text{g.mL}^{-1}$)	1.2809	0.8017
Limit of quantification ($\mu\text{g.mL}^{-1}$)	4.293	2.6722
Standard error for regression line (Sy/x)	0.004722	0.002886

Part 2: optimization reaction of FIA for CFT

The effect of HCl concentration

The effect of the concentration of HCl was studied in the range (1.16 to 5.8) M with constant CFT

concentration of (100 $\mu\text{g/ml}$). The concentration (5.8 and 4.64) M gave highest absorbance for reagent (1-NPT ,2-NPT) respectively Fig.12.

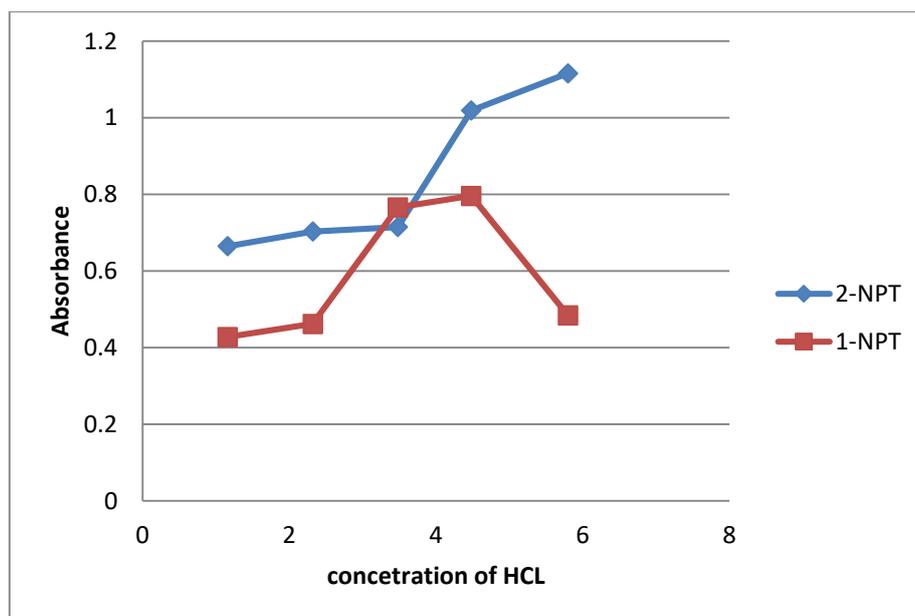


Figure 12. Effect concentration of HCl

The effect of NaNO₂ 1% volume;

The effect of the volume of the sodium nitrite on the absorbance of the colored products Fig .13. The effect of different volumes of sodium nitrite was

studied that range from 25 to 5 ml, 20 and 15 ml that was gave the highest absorption for reagents respectively.

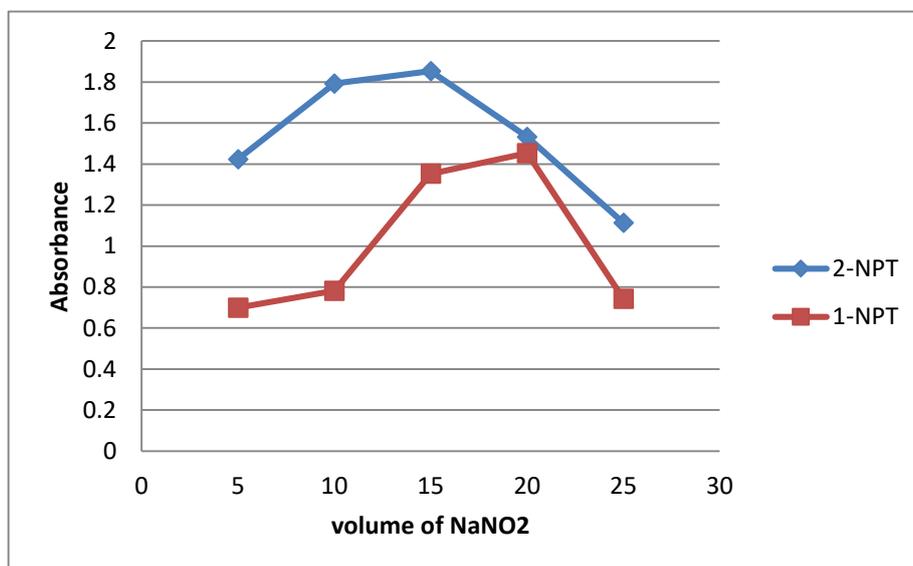


Figure 13. Effect volume of NaNO₂

The effect of reagents concentration;

At constant CFT concentrations, the effects of 1-NPT and 2-NPT concentrations on absorbance were investigated. CFT was injected into the HCl stream in fixed volumes (100 mL) at 1000 µg/mL. Fig. 14 shows the effect of changing the reagent

concentrations in the range from 0.0694×10^{-3} to 0.832×10^{-3} M (300 -1200) µg/ml on the absorption peak. The figure shows that when the reagent concentration reached 0.832×10^{-3} M, a maximum analytical signal was obtained, and it was chosen for further use

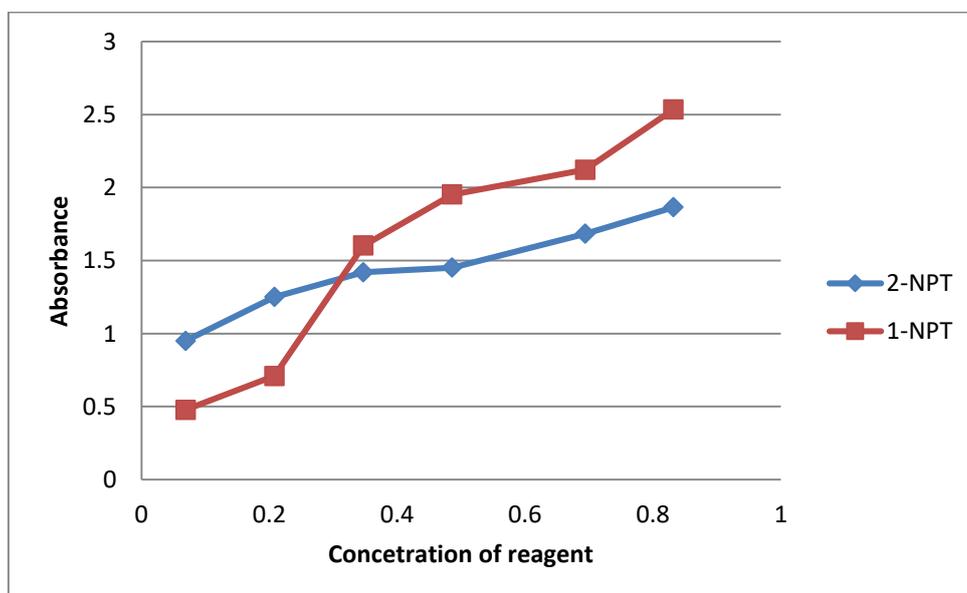


Figure 14. Effect concentration of reagents.

The effect of different concentration of NaOH.

The effect of different concentration of NaOH was studied, the range from (5 to 25%), with a fixed

concentration of acid and drug, and it 5% showed the highest absorption intensity to both of reagents. Fig. 15.

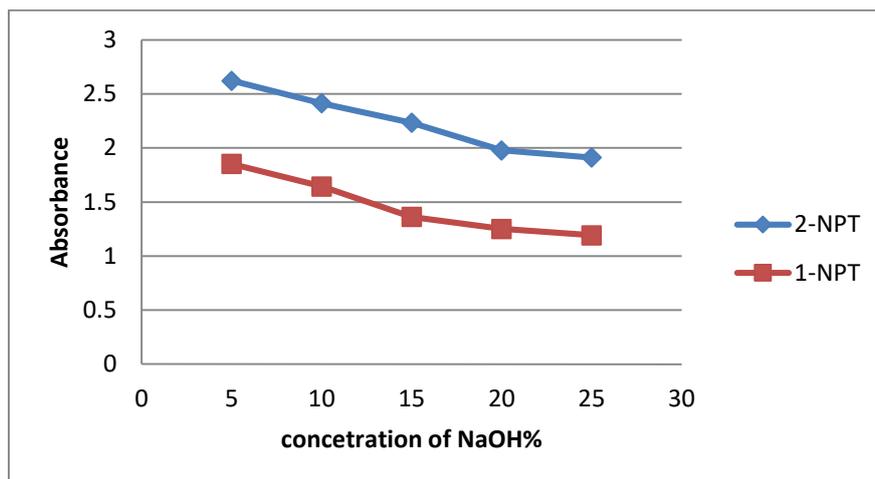


Figure 15. Effect concentration of NaOH%

Analytical data for Flow injection.

After determining the optimal experimental conditions, the calibration curve was plotted as it is

shown in Fig. 16 and Table 4 calibration curve and analytical information, respectively

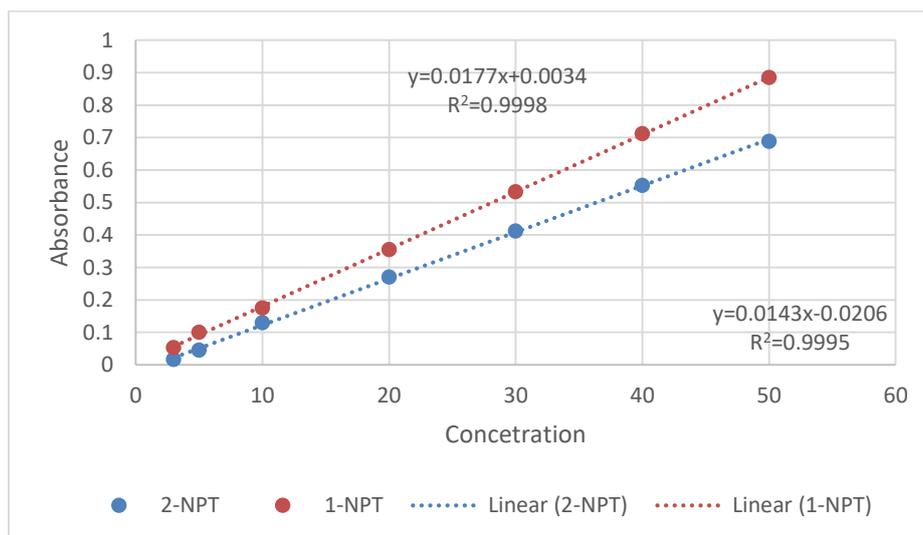


Figure 16. Calibration curve of the FLA method.

Table 4. characteristic parameters of proposed Flow injection methods.

Parameter	CFT-1-NPT	CFT-2-NPT
Color of product	Brown	Red
λ max (nm)	585	545
Dynamic range ($\mu\text{g}\cdot\text{ml}^{-1}$)	(3-50)	(3-50)
Molar absorptivity, ϵ ($\text{L}\cdot\text{mol}^{-1}\cdot\text{cm}^{-1}$)	0.957×10^4	0.7106×10^4
Regression equation	$y=0.0177x+0.0034$	$y=0.0143x-0.0206$
Sandell sensitivity, S ($\mu\text{g}\cdot\text{cm}^{-2}$)/0.001A.U	0.05714	0.07692
Intercept (a)	+0.0034	-0.0206
Slope (b) ($\text{L}\cdot\text{mg}^{-1}\cdot\text{cm}^{-1}$)	0.0177	0.0143
Coefficient of determination %	99.98	99.95

R ²		
Correlation coefficient (r)	0.9999	0.9997
Limit of detection (µg.mL ⁻¹)	0.8108	1.281
Limit of quantification (µg.mL ⁻¹)	2.703	4.27
Standard error for regression line (Sy/x)	0.004784	0.006106

Analytical Application:

Three types of pharmaceuticals were used to examine the proposed procedures (Diazotization,

Flow injection) for the analysis of pharmaceutical formulations. Table of contents (5, 6) provides all of the results that were obtained, as both approaches were accurate and precise.

Table 5. Application of the proposed Batch method for the determination of Ceftazidime in different brands of drugs.

Ceftazidime YENIZIM made in Turkey Ankara	Amount of CFT(µg/ml)		E%	Recovery%	Average%	RSD (n=5)
	Taken	Found				
1-NPT	5	4.99	-0.18	99.81	100.05	2.48
	20	19.81	-0.95	99.05		0.737
	40	40.53	1.34	101.3		0.486
2-NPT	5	4.88	-2.22	97.77	99.67	0.495
	20	20.35	1.76	101.76		0.614
	40	39.79	-0.5	99.49		0.431
Ceftazidime TOTTIZIM made in ITALY	Amount of CFT(µg/ml)		E%	Recovery%	Average%	RSD (n=5)
	Taken	Found				
1-NPT	5	4.9	-2	98	99.16	1.57
	20	19.72	-1.41	98.59		0.813
	40	40.35	0.88	100.88		0.472
2-NPT	5	4.79	-4.07	95.93	99.17	0.598
	20	20.37	1.85	101.85		0.699
	40	39.88	-0.28	99.72		0.374
Ceftazidime ROTH made in Wiesbaden Germany	Amount of CFT(µg/ml)		E%	Recovery%	Average%	RSD (n=5)
	Taken	Found				
1-NPT	5	4.81	-3.82	96.18	98.32	1.367
	20	19.63	-1.86	98.13		0.533
	40	40.26	0.66	100.66		0.495
2-NPT	5	5.03	5.03	100.56	100.76	0.504
	20	20.44	2.22	102.22		0.613
	40	39.79	-0.51	99.49		0.456

Table 6. Application of the proposed Flow injection method for the determination of Ceftazidime in different brands of drugs.

Ceftazidime YENIZIM made in Turkey Ankara	Amount CFT($\mu\text{g/ml}$)		of E%	Recovery%	Average%	RSD (n=5)
	Taken	Found				
1-NPT	10	9.75	-2.49	97.5	99.05	1.027
	30	29.98	-0.08	99.92		0.431
	50	49.86	-0.27	99.73		0.343
2-NPT	10	9.97	-0.28	99.72	99.93	0.932
	30	30.25	0.839	100.84		0.276
	50	49.62	-0.76	99.24		0.4889
Ceftazidime TOTTIZIM made in ITALY	Amount CFT($\mu\text{g/ml}$)		of E%	Recovery%	Average%	RSD (n=5)
	Taken	Found				
1-NPT	10	9.81	-1.92	98.08	99.34	1.189
	30	30.03	0.113	100.11		0.339
	50	49.92	-0.16	99.84		0.375
2-NPT	10	9.90	-0.98	99.02	99.73	1.067
	30	30.32	1.072	101.07		2.011
	50	49.55	-0.89	98.10		0.455
Ceftazidime ROTH made in Wiesbaden Germany	Amount CFT($\mu\text{g/ml}$)		of E%	Recovery%	Average%	RSD (n=5)
	Taken	Found				
1-NPT	10	9.86	-1.4	98.6	99.62	1.782
	30	30.09	0.301	100.30		0.431
	50	49.98	-0.05	99.95		0.375
2-NPT	10	10.04	0.419	100.42	100.23	1.296
	30	30.39	1.305	101.3		0.216
	50	49.48	-1.04	98.96		0.457

Conclusion:

Because the 1-NPT and 2-NPT reagents are methods that are available and inexpensive, the research includes two simple, sensitive, fast, and inexpensive methods for estimating ceftazidime. The first (diazotization-coupling) entails converting ceftazidime into a colored dye that can be measured using a UV-method Vis spectrophotometer. The second step involves using flow injection to pre-concentrate colored dye. This is the first method for extracting CFT from a flow injection. These methods have been used to estimate ceftazidime in pharmaceuticals with great success.

Author's declaration:

- Conflicts of Interest: None.
- We hereby confirm that all the Figures and Tables in the manuscript are ours. Besides, the Figures and images, which are not ours, have been given the permission for re-publication attached with the manuscript
- Ethical Clearance: The project was approved by the local ethical committee in University of Mustansiriyah.

Authors Contribution:

M J M H: Suggesting a research project, preparing a research plan, providing all laboratory supplies and chemicals

H Ab : Executing and conducting all laboratory experiments and obtaining the results and presenting them to the supervisor.

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طرق القياس الطيفي للحقن الدفعي والتدفق لتقدير السيفتازيديم في التركيبات الصيدلانية

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الخلاصة :

تم وصف تقنيتين طيفيتين لقياس كمية السيفتازيديم (CFT) في الأدوية والتركيبات الصيدلانية. فهي بسيطة وحساسة وانتقائية ودقيقة وفعالة. يستخدم الأسلوب الأول وسيطاً قلوياً لتحويل السيفتازيديم إلى ملح الديازونيوم ، والذي يتم دمج بعد ذلك مع كواشف 1-NPT-Naphthol و 2-NPT-Naphthol. تم إنتاج صبغة الأزو باللونين البني والأحمر ، مع كثافة امتصاص تصل إلى λ_{max} 585nm and max 545nm ، على التوالي. تم اتباع قانون بير من حيث مدى التركيز (3-40) لـ (CFT-1-NPT) و (CFT-2 NPT) ، كانت حدود الكشف 1.0096 g.ml-1 and 0.8017 g.ml-1 على التوالي ، وكانت الامتصاصية المولية 104×0.7926 and 0.5466×104 L.mol⁻¹cm⁻¹. تُستخدم طريقة تحليل حقن التدفق (FIA) لتقدير السيفتازيديم وتسجيل القياسات باستخدام نهج الأشعة فوق البنفسجية المرئية في الإجراء الثاني. يسمح حقن التدفق بتقدير دقيق للعقار في ظل ظروف تجريبية مثالية. كانت التركيزات في نطاق (3-50) لـ (CFT-1-NPT) و (CFT-2-NPT) ، وكانت حدود الكشف 0.8102, 0.8102 g.ml-1 and 11.2809, 0.8102 g.ml-1 ، والامتصاصية المولية 104×0.9565 and 104×0.7106 L.mol⁻¹cm⁻¹. على التوالي تم استخدام الأسلوبين المقترحين لتقدير السيفتازيديم في المستحضرات الصيدلانية بنجاح حيث تميزت هذه الطريقة بالبساطة والسرعة والدقة والتكلفة المنخفضة.

الكلمات المفتاحية : صبغة الأزو, الدفعات, سيفتازيديم, الأزوتة, الحقن الجرياني, 1-نفثول, 2-نفثول, الطيفية.