

Spectrophotometric Determination of Benzocaine by Azo-Dye Formation Reaction

Dawood Habbo Mohammeda*

Lamya Adnan Sarsamb**



* Mosul University- Education College for women

** Mosul University- College of Science.

ARTICLE INFO

Received: 16 / 6 /2010
Accepted: 9 / 2 /2011
Available online: 14/6/2012
DOI: 10.37652/juaps.2011.15433

Keywords:

Spectrophotometric ,
Benzocaine ,
Azo-Dye Formation Reaction.

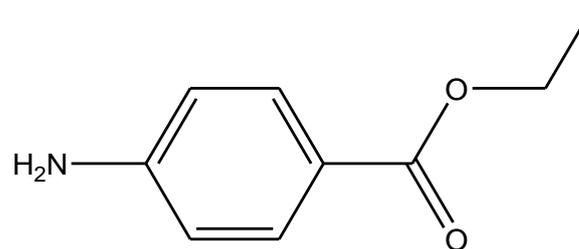
ABSTRACT

A spectrophotometric method for the assay of trace amounts of benzocaine was based on the reaction of benzocaine with nitrite ion to form the corresponding diazonium salt followed by coupling reaction with ethyl cyano acetate to form a stable and a soluble yellow azo dye with maximum absorption at 405 nm. Beer's law was obeyed over the range 5-250 μ g/25ml, i.e, (0.2-10ppm) of benzocaine and correlation coefficient 0.998 with a molar absorptivity of 3.1×10^4 L.mol⁻¹.cm⁻¹, a relative error of 0.0 to 0.25 % and a relative standard deviation of ± 0.03 to ± 0.38 %, depending on the concentration level. The method has been successfully applied for the assay of benzocaine in one pharmaceutical preparation (Lozenges).

Introduction:

Benzocaine (4-aminobenzoic acid ethyl ester)(synonym: ethyl aminobenzoic acid , is a local anaesthetic of the ester type with a poor solubility in water which is used for superficial anesthesia, for the local and temporal relief of pain ,among other disorders, to buccal effects (1). For such reasons , it is a drug extensively used in odontology(2-3). It is used in Cattle , sheep ,swine and horses for local and prolonged low epidural anaesthesia . Benzocaine acts on the central nervous system ,cardiovascular system , neuromuscular junctions and ganglion synapse .Its mechanism of action is to prevent the generation and conduction of the nerve impulse. It has been proposed that the drug penetrates cell membranes in its uncharged form and binds to putative intracellular receptors. Various spectrophotometric (4-10) and chromatographic(11-16) methods for determination of benzocaine have been reported. In the present work an

attempt was made to develop a rapid and sensitive method for the determination of benzocaine in pharmaceutical formulation.



4-amino benzoic ethyl ester

Experimental

Instruments

All spectrophotometric measurements were performed on Shimadzu UV-Visible Recording Spectrophotometer UV-160 using 1 cm silica cell, pH meter type Philips PW 9420 was used for pH measurements.

Reagents

All chemicals used in this investigation are of analytical – reagent grade, benzocaine standard material is provided from general establishment for

* Corresponding author at: Mosul University- Education College for girls.E-mail address:

medical appliance and drugs / NDI – Mosul / Iraq.

Solutions

Benzocaine (100 µg/ml): 0.01g was dissolved in ethanol solution transferred into a 100 ml volumetric flask, and diluted to the mark with distilled water.

Ethyl cyano acetate, 2% (v/v), was prepared freshly daily by dissolving 2 ml of ethyl cyano acetate in 100 ml distilled water.

Sodium nitrite solution, 1% (w/v), was prepared by dissolving 1 g of sodium nitrite (BDH) in 100 ml distilled water.

Sulphamic acid solution, 3% (w/v), was prepared by dissolving 3 g of sulphamic acid (Fluka) in 100 ml distilled water.

Hydrochloric acid solution, 1N. was prepared by diluting 8.47ml of concentrated acid (11.8 N) to 100 ml with distilled water.

General procedure

To a series of 25-ml calibrated flasks, transfer 0.05 – 2.5 ml of benzocaine solution, then 1 ml of 1M hydrochloric acid and 0.3 ml of 1% (w/v) sodium nitrite solution were added and the mixture was allowed to stand for 2 minute and then 1.0 ml of 3% (w/v) sulphamic acid solution was added with occasional shaking for 3 minutes. After that a 1.0 ml of 2% (v/v) ethyl cyano acetate was added. Then the solutions was let to stand for 1 minute at room temperature before adding 2ml of 1M ammonium hydroxide then the volumes were completed to the mark with distilled water, The absorbance was read at 405 nm against the reagent blank. A linear calibration graph was obtained over the concentration range of 5 – 250 µg benzocaine / 25 ml (0.2-10 ppm) and a concentration above 250 µg / 25 ml gave a negative deviation (Fig. 1). The molar absorptivity has been found to be 3.1×10^4 L. mol⁻¹. cm⁻¹.

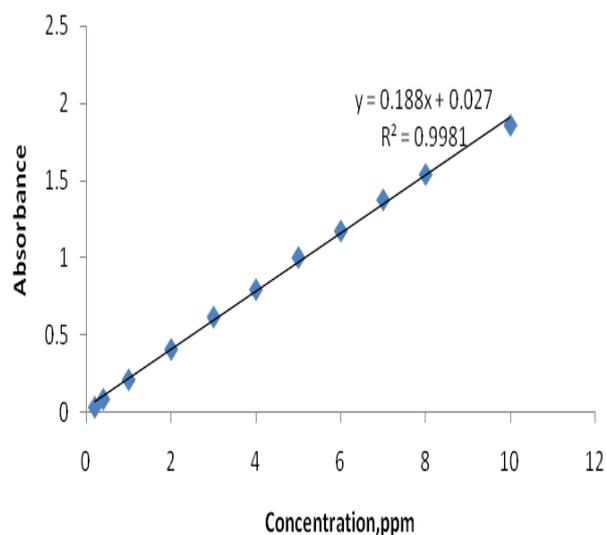


Fig. 1. Calibration graph of benzocaine determination at 405nm.

Results and Discussion

Study of the optimum reaction conditions

Effect of acid

Different amounts and types of acids have been used in diazotization of benzocaine; the results showed that 1 ml of 1 M HCl has been selected for subsequent experiments (Table 1).

Table 1. Effect of acids on absorbance and colour contrast

1M Acid solution used	Absorbance/ml of acid added				
	0	0.5	1.0	1.5	2
HCl	0.339	0.387	0.406	0.387	0.380
HNO ₃	0.339	0.128	0.109	0.084	0.070
H ₂ SO ₄	0.339	0.402	0.261	0.084	0.071
CH ₃ COOH	0.339	0.380	0.377	0.095	0.098

Effect of sodium nitrite amount and time

The maximum absorbance reading was obtained by adding 0.3 ml of 1% sodium nitrite for 2 minutes of reaction time (Table2).

Table 2. Effect of sodium nitrite amounts and time on the absorbance of benzocaine

ml of 1% NaNO ₂ solution	Absorbance / minute standing time					
	0	1	2	3	4	5
0.1	0.387	0.371	0.376	0.377	0.391	0.388
0.3	0.375	0.389	0.402	0.358	0.380	0.385
0.5	0.379	0.387	0.384	0.369	0.382	0.389

0.7	0.378	0.377	0.387	0.384	0.379	0.398
1.0	0.393	0.378	0.390	0.381	0.354	0.383

Effect of sulphamic acid amounts and time

The excess of nitrite can be removed by the addition of sulphamic acid solution. The effect of sulphamic acid amount and time has been studied. (Table3).

Table 3. Effect of sulphamic acid amounts and time on the absorbance of benzocaine

ml of 3% Sulphamic acid solution	Variable	Absorbance/minute standing time					
		0	1	2	3	4	5
0.1	Sample = S	0.383	0.390	0.387	0.380	0.395	0.377
	Blank = B	0.00	-0.010	-0.008	-0.006	-0.004	-0.018
0.3	S	0.389	0.387	0.408	0.379	0.383	0.395
	B	-0.003	-0.012	-0.001	0.009	0.000	-0.014
0.5	S	0.385	0.394	0.387	0.384	0.387	0.372
	B	-0.007	-0.010	-0.008	0.000	0.008	-0.006
0.7	S	0.377	0.391	0.408	0.374	0.376	0.387
	B	0.001	-0.007	-0.005	-0.003	0.000	-0.008
1.0	S	0.365	0.368	0.409	0.379	0.388	0.370
	B	0.007	-0.014	-0.006	0.001	-0.006	-0.013

The results in the table 3 indicated that 1.0 ml of sulphamic acid solution (3%, w/v) with 2 minutes as

standing time for the reaction gave the most suitable effect on the intensity of the azo-dye.

Effect of ethyl cyano acetate amount on absorbance

The effect of ethyl cyano acetate amount on the absorbance of the dye has been studied. From the results, it can be observed that 1.0 ml of 2% ethyl cyano acetate with 1 minute of reaction time was the more suitable which gave the highest value of intensity for the azo-dye (Table 4).

Table 4. Effect of coupling agent amount on absorbance

MI of ethyl cyano acetate solution (2%)	Absorbance/min. standing time		
	0	1	3
0.5	0.368	0.393	0.376
1.0	0.392	0.406	0.373
3.0	0.375	0.385	0.362
5.0	0.385	0.357	0.361
6.0	0.366	0.361	0.377

Effect of time

The coloured azo dye developed rapidly after addition of ethyl cyano acetate and the stability period (within the first hour of stability) was sufficient to perform several measurements and the results are given in table 5. It is shown from table (5) that the maximum absorbance was obtained during the first five minutes and declined gradually.

Table 5. The effect of time and benzocaine amount on absorbance

µg of benzocaine present	Absorbance / minute standing time									
	0	5	10	15	20	25	35	45	55	60
5	0.792	0.051	0.060	0.061	0.057	0.060	0.058	0.058	0.065	0.060
	0.790	0.211	0.213	0.221	0.206	0.219	0.212	0.211	0.216	0.208
	0.790	0.406	0.406	0.405	0.406	0.404	0.406	0.406	0.406	0.396
25	0.792	0.211	0.213	0.221	0.206	0.219	0.212	0.211	0.216	0.208
	0.790	0.406	0.406	0.405	0.406	0.404	0.406	0.406	0.406	0.396
	0.792	0.792	0.790	0.790	0.792	0.793	0.792	0.792	0.792	0.784
50	0.792	0.211	0.213	0.221	0.206	0.219	0.212	0.211	0.216	0.208
	0.790	0.406	0.406	0.405	0.406	0.404	0.406	0.406	0.406	0.396
	0.792	0.792	0.790	0.790	0.792	0.793	0.792	0.792	0.792	0.784
100	0.792	0.211	0.213	0.221	0.206	0.219	0.212	0.211	0.216	0.208
	0.790	0.406	0.406	0.405	0.406	0.404	0.406	0.406	0.406	0.396
	0.792	0.792	0.790	0.790	0.792	0.793	0.792	0.792	0.792	0.784

Table 6. The effect of amount and type of base

Base used (1M)	Absorbance / ml of based used		
	1	2	3
NaOH	0.349	0.354	0.356
pH	12.10	12.56	12.71
Na ₂ CO ₃	0.380	0.371	0.364
pH	9.57	10.02	10.17
NH ₄ OH	0.388	0.406	0.402
pH	8.92	9.47	9.74

Effect of amount and type of base

The preliminary experiments have shown that the azo-dye developed only completely in alkaline medium. Different amounts of bases (strong and weak) have been used (table 6).

The experimental data showed that ammonium hydroxide gave better sensitivity than sodium hydroxide and sodium carbonate. So that 2.0 ml of 1M NH₄OH is recommended for the subsequent experiments

Effect of order of additions

To obtain optimum results the order of additions of reagents should be followed as given under the general procedure, otherwise a loss in colour intensity was observed.

absorption spectra

The absorption spectra of the yellow azo dye formed by coupling of diazotised benzocaine with ethyl cyano acetate shows a maximum absorption at 405 nm. The reagent blank gives very weak absorption at this wavelength (Fig. 2).

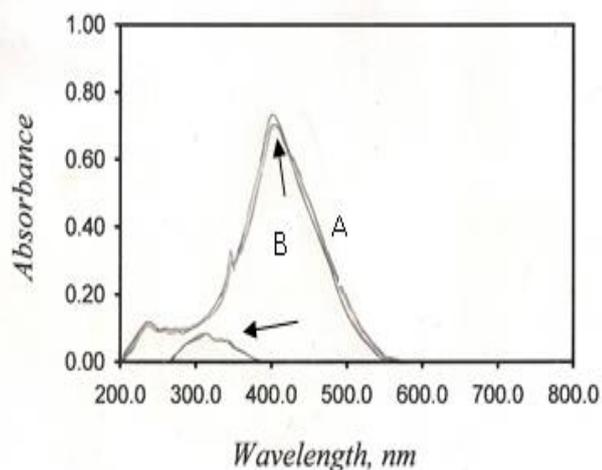


Fig.2: Absorption spectra of 100µg benzocaine / 25ml were treated according to the recommended procedure and measured against (A) reagent blank, (B) distilled water and (C) reagent blank measured against distilled water.

Nature of the dye

The stoichiometry of the azo dye thus formed by reaction of diazotised benzocaine with ethyl cyano acetate was investigated by applying the continuous variations method (Job's method). The results indicated that the azo-dye was formed in the ratio of 1:1 diazotised benzocaine to ethyl cyano acetate (Fig.3).

Interference

The effect of some foreign compounds which often accompanied pharmaceutical preparations were studied by adding three different amounts (50, 100 and 200µg) to 100µg benzocaine in a final volume 25ml (Table 7).

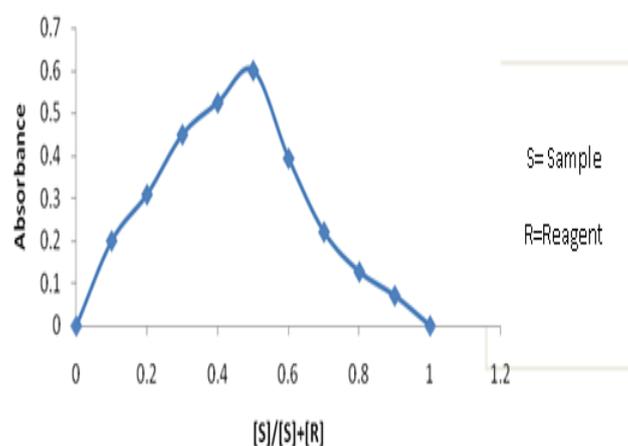


Fig.3: Job's plot for benzocaine - ethyl cyano acetate

Therefore the azo-dye may have the following suggested structure:

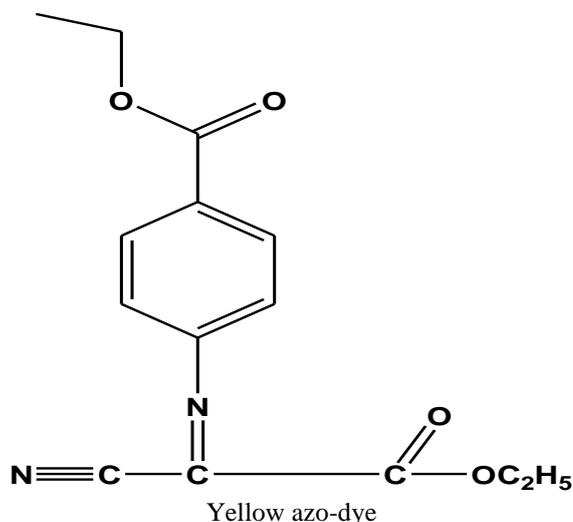


Table 7: Effect of foreign compounds for assay of benzocaine

Interferent	Recovery (%) of 100 µg benzocaine / µg of interferent added		
	50	100	200
	Acaccia	98.28	99.70
Cetyl pyridenium chloride	99.60	99.26	95.7
Glucose	96.40	98.30	99.76
Lactose	98.52	97.05	100.1
Lindocaine	97.30	97.54	100.0
menthol	98.54	99.24	100.1
Starch	97.50	98.51	98.76

The results in table7 indicated that the studied foreign compounds did not interfere in determination of benzocaine using the proposed method.

Accuracy and precision.

To check the accuracy and precision of the method, benzocaine was determined at four different concentrations. The relative error% and relative standard deviation% results indicated the high accuracy and precision of the proposed method (Table 8).

Table 8. Accuracy and precision of determination of benzocaine using spectrophotometric method.

Amount of benzocaine taken, µg	Relative error, %*	Relative standard deviation, %*
25	0.00	± 0.23
50	+0.25	± 0.38
100	0.00	± 0.06
200	0.00	± 0.03

* Average of five determinations

Analytical application

The proposed method was applied to assay benzocaine in two Synthetic Pharmaceutical Preparations lozenges of benzocaine and throat lozenges solutions (E.Y. Hassen 2005). On applying proposed procedure, good recovery was obtained for lozenges of benzocaine solution only as shown in table 9.

Table9. Analytical applications of the spectrophotometric method in determination of benzocaine in Lozenges

Pharmaceutic al preparation	µg benzocaine present/25ml	Recovery* (%)
-----------------------------	----------------------------	---------------

Lozenges of benzocaine	50	97.3
	100	98.0
	150	99.0

*Average for five determinations

The proposed method for the determination of benzocaine in pharmaceutical preparation was simple and sensitive. The azo-dye formed was fairly soluble in aqueous solution. The statistical analysis of the results indicated that the method has good accuracy and good precision.

References

1. British Pharmacopoeia ,2000, Version 4.0 CD , The Stationery office Ltd , London , May.
2. M.I.Arufe-Martinaz, J.L.Romero-Palanco and M.A.Vizcaya-Rojas, (1989),"Application of derivative spectrophotometry for the simultaneous determination of cocaine and other local anesthetics , II Cocaine and benzocaine mixture" Spainal J.Anal. Toxicol , 13(6) , 349-353 ; Anal.Abst. (1990) , 52 , 11E9 , p.734.
3. A.M.Casas-Hernandez, M.P.Aguilar-Caballos and A.Gomez-Hens , (2002), "Application of time-resolved luminescence to dry reagent chemical technology" Anal.Chem.Acta, 452(2) , 169-175.
4. L.R.Paschoal and W.A.Ferreira,(2000), "Simultaneous determination of benzocaine and cetylpyridinium chloride in tablets by first-derivative spectrophotometric method" IIFarmaco, 55(11) , 687-693.
5. N.S. Othman,(2001) , " A continued investigation of spectrophotometric disometry" Ph.D. Thesis ,University of Mousal , Cellege of Sciences, p. 75-95.
6. N.D.Dinesh, P.Nagaraja, and K.S.Rangappa, (2002), "Sensitive spectrophotometric method for the analysis of some anesthetic drug" Indian J.Pharm.Scie. , 64(5), 485-488 .

7. A.S.Amin and A.M.El-Didamony,(2003) ,
"Colorimetric determination of benzocaine ,
lignocaine and procaine hydrochlorides in pure
form and in pharmaceutical formulation using p-
benzoquinone" Anal.Scie. , 19 , 1457-1459.
8. R.A.A.Zakaria, (2004)," Spectrophotometric
determination of benzocaine and Salbutamol
sulphate using diazotisation coupling method-
application to some drugs preparation "
M.Sc.,Thesis, University of Mousl, College of
Sciences, p. 5-34.
- 9.E.Y.Hassen" ,(2005),Development of
Spectrophotometric Methods for the
Determination of Benzocaine in Two Synthetic
Pharmaceutical Preparations
"M.Sc.Thesis, University of Mosul, College of
Sciences.
10. D.H.Mohammed, H.H.Ahmed and H.A.
Mohammed",(2009), pectrophotometric
determination of Benzocaine by formation azo dye
with acetyl acetone" J. Nat.Chem.,35,383-395.
11. P.Linares, M.C.Gutierrez, F.Lazaro, M.D.Luque
De Castro and M.Valcarcel ,
(1991),"Determination of benzocaine
,dextromethorphan and cetylpyridinium ion by
high-performance liquid chromatographic with UV
detection " J.Chro. A , 558(1) , 147-153.
12. Gigante, A.M.V.Barros, A.Teixeira and
M.J.Marcelo-Curto, (1991) , "Separation and
simultaneous high-performance liquid
chromatographic determination of benzocaine and
benzyl benzoate in pharmaceutical preparation"
J.Chro.A , 549 , 217- 220.
13. G.S.Sadana and A.B.Ghogare, (1991) ,
"Simultaneous determination of chloramphenical
and benzocaine in topical formulation by high-
performance liquid chromatographic" J.Chro. , 542
, 515-520.
14. T. A. Biemer, N. Asral and J. A. Albanese,(1992),
"Simultaneous stability- indicating capillary gas
chromatographic assay for benzocaine and the two
principal benzyl esters of Balsam Peru formulated
in a topical ointment", J. Chro. A., 623(2), 395-
398.
15. J.A.Bernardy, K.S.Coleman, G.R.Stehly and
W.H.Gingerich,(1996), "Determination of
benzocaine in rainbow trout plasma" J.AOAC
International , 79(3) , 623-627.
16. J.Joseph-Chartes, M.Montaqut, M.H.Lanalis,
C.Boyer and J.P.Dubost, (2001),"Simultaneous
determination of rutine and benzocaine in
suppositories by reversed-phase HPLC" Anal.Lett.
, 34 , 2685-2692.

التقدير الطيفي للبنزوكائين بواسطة تفاعلات تكوين صبغة الأزو

داود حيو محمد لمياء عدنان سرسم

الخلاصة

تضمن البحث طريقة طيفية لتقدير كميات متناهية في الصغر من البنزوكائين. وأعدت الطريقة على أزوتة البنزوكائين وذلك بمفاعله مع ايون النتريت بوجود حامض الهيدروكلوريك ثم اقتران ملح الدايازونيوم الناتج مع كاشف الاقتران اثيل سيانو اسيتيت لتكوين صبغة آزوية صفراء مستقره وذائبة في الماء، تم قياس شدة الامتصاص للصبغة الناتجة عند الطول الموجي 405 نانوميتر وكانت حدود قانون بير في مدى التركيز من 5 إلى 250 مايكروغرام من البنزوكائين في حجم نهائي 25 مل (0.2 - 10 جزء /مليون) ويمعامل ارتباط 0.998 وبلغت قيمة الامتصاصية المولارية 3.1×10^4 لتر. مول⁻¹. سم⁻¹، والخطأ النسبي تراوح بين 0.0 و 0.25 % والانحراف القياسي النسبي بين $0.03 \pm$ و $0.38 \pm$ % اعتمادا على مستوى تركيز البنزوكائين. تم تطبيق الطريقة بنجاح لتقدير البنزوكائين في احدى مستحضراته الصيدلانية المحضرة (Lozenges).