

photometric determination of Nickel (II) and Cobalt (II) ions. The reagent (CMTSC) gives intense yellow colour water soluble complexes with Nickel (II) and Cobalt (II) in presence of micellar medium (Triton-X-100 (5%) at pH 9.0. The maximum absorbance was observed at λmax 440 and 400 nm for Nickel (II) and Cobalt (II) respectively. The molar absorptivity and sandell's sensitivity for Nickel (II) and Cobalt (II) complexes are found to be 4.77x10⁴, 4.70x10⁴ L.mol⁻¹.cm⁻¹ and 0.0012, 0.0012 µg/cm² respectively. The reagent (CMTSC) forms 1:1 [M:L] complexes with Nickel (II) and Cobalt (II) with stability constants 8.17x 10⁵ and 10.10x10⁶ respectively. The First order derivative spectrophotometric technique has been developed for the simultaneous determination of Nickel (II) and Cobalt (II) at λmax 460 and 420 nm.

Introduction:

Hydrazones are important organic analytical reagents for the spectrophotometric determination of metal ions in microgram quantities. They react with metal ions forming colour complexes and act as chelating agents. The potential applications of hydrazone derivatives for the spectrophotometric determination of metal ions have been reviewed by Singh et.al^[1].

The great interest towards derivative spectrophotometry was due to the increased resolution of spectral bands, that means it is resolving two overlapping spectra and eliminating matrix interferences in the assay of two component mixtures using zero-crossing technique.²⁻³ In the absence of zero-crossing point, two simultaneous equations can be solved to determine the components in such a mixture^[4-5] Hydrazone reagents are widely used in our laboratories for the derivative spectrophotometric determination of metal ions^{[5-8].} Derivative spectrophotometric analysis of two-component mixtures was also carried out without need to solve simultaneous equations^[6-7] In the light of good analytical characteristics of hydrazones, here in we report the simultaneous first order derivative spectrophotometric determination of Nickel (II) and Cobalt (II) using Cinnamaldehyde semithicarbazone(CMTSC). The proposed simultaneous method involves the use of peak-to-base line measurement technique.

Experimental part-Materials & methods: Apparatus:

Shimadzu 16A microcomputer based U.V-Visible spectrophotometer equipped with 1.0 cm quartz cells used for all absorbance studies and amplitude measurements in derivative spectrophotometery. An ELICO LI-120 digital pH mater was used in pH adjustments.

Recommended procedure for Simultaneous determination of Nickel (II) and Cobalt (II)

For simultaneous determination of Nickel (II) and Cobalt (II), a solution containing both Nickel (II) and Cobalt (II) in the optimum concentration range 0.176-0.727 μ g/mL of Nickel (II) and 0.142-0.637 μ g/mL of Cobalt (II) are used. The absorption spectra indicates that Nickel (II) and Cobalt (II) complexes shows maximum amplitude in derivative spectrophotometric technique at Imax 460 and 420 nm respectively. Calibration curves are prepared between amplitude and amount of metal ion at their respective wavelength. The amount of metal ion [Ni (II) or Co (II)] was calculated from calibration graphs.

At λ max 420 nm Cobalt (II) complex has a zero crossing point, which is independent of the metal ion concentration and at this wavelength Nickel (II) has appreciable amplitude shown in Figue.2. Hence, simultaneous first order derivative spectrophotometric determination of Ni (II) and Co (II) has been carried out by measuring peak-base line technique at 460 nm and 420 nm respectively without employing the simultaneous equations.

Calibration plots were constructed at 460 nm and 420 nm by plotting the derivative amplitudes against the corresponding amounts of Ni (II) and Co (II) respectively. The plots thus obtained are linear obeying the relationship.

Ni (II) A 460 nm = 0.66846X+2.66667x10⁻⁴

Co (II) A 420 nm = 0.68153X+0.00125



Figure.1: Zero order Absorption Spectra



Fig.2: Simultaneous first order Derivative Spectra (a) Reagent CMTSC Vs water blank, (b) Co (II)-CMTSC Complex Vs Reagent blank,

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(c) Ni (II)-CMHBH Complex Vs Reagent blank. (D) Co (II)-CMTSC (E) Ni (II)-CMTSC [Ni (II) &Co (II)] = 1.5x10⁻⁵, [CMTSC] = 3×10⁻³ M, pH = 9.0 Triton X-100 (5%) = 0.5 mL



Figure.3: Typical first order derivative spectra

[Ni (II)]	$= 1.5 \times 10^{-6}$
[CMHBH]	= 3X10 ⁻³
рН	= 9.0
Triton-X-100 (5%) = 0.5 mL
λmax	= 460 nm



Figure.4: Typical first order derivative of Ni (II)-CMTSC Co (II)-CMTSC

[Co (II)]	= 1.5x10-6
[CMHBH]	= 3X10 ⁻³
pН	= 9.0
Triton-X-100 (5	5%) = 0.5 mL
λmax	= 420 nm

Results and discussion:

Cinnamaldehyde semithiocarbazone (CMTSC) was a novel type of reagent. The above reagent was easily synthesized as any other Schiff base. Aldehyde-4-hydroxybenzoylhydrazone reagents are not exploited much for spectrophotometric determination of metal ions. Moreover, the derivative methods using this class of reagents are scare.



Structure of CMTSC

Interference:

The effect of foreign ions was studied to know the selectivity of the derivative methods. The amount of foreign ions, which brings about change in amplitude by \pm 2%, was taken as the tolerance limit. Interference of various ions which often accompany with Nickel (II) and Cobalt (II) has been studied in the determination of 0.440 μ g/mL of Nickel (II) and 0.441 μ g/mL of Cobalt (II).

The data obtained in derivative method is also incorporated.

It is noticed that all the ions that do not interfere in the zeroorder determination of metal ions also do not interfere in the first derivative analysis. The tolerance limit values, for many anions and cations are higher in the first order derivative methods than those in the zero-order determination of metal ions. The interference of associated metal ion such as Fe (III) was decreased by adding the masking agent fluoride.

Applications:

The simultaneous first derivative spectrophotometric determination of Cobalt applied in biological samples, Nickel in plant samples, some alloy steels and soil sample. The results are presented in Table 1, 2 &3 respectively.

Table.1: Simultaneous	determination	of Ni (II)	and	Co	(II)
in Alloy steel samples					

Alloy sample	Certified (%)		Amount found (%)		Error (%)	
	Ni (II)	Co (II)	Ni (II)	Co (II)	Ni (II)	Co (II)
(a) Eligiloy M-1712	15.00	40.00	14.95	39.89	-0.27	-0.15
(b) BCS 406/1	0.14	0.016	0.138	0.015	-1.40	+0.63
(c) Alloy steel	11.22	23.72	11.23	23.67	+0.08	-0.04

*Average of three determinations; (a) 20% Cr, 40% Co, 15% Ni, 0.15% C, 15% Fe, 2% Mn, 7% Mo, 0.05% Be. (b) 0.066% Mn, 1.06% Cr, 0.05% Mo, 0.14% Ni, 0.016% Co, 0.091% Cu, 0.19% V^z. (c) 51.15% Fe, 11.22% Ni, 5.09% Cu, 23.72% Co, 6.9% Al, 0.79% Ti, 0.235% Mn, 0.57% Si.

Table.2: Simultaneous first-order derivative determination of Co (II) and Ni (II) in synthetic binary mixtures

Amount	taken (µg/	Amount f	ound (µg/	Error (%)		
Ni (II)	Co (II)	Ni (II)	Co (II)	Ni (II)	Co (II)	
0.2347	0.2357	0.2345	0.2356	-0.08	+0.42	
0.4694	0.2357	0.4689	0.2349	-0.10	-0.34	
0.7041	0.2357	0.7045	0.2352	+0.05	-0.21	
0.9388	0.2357	0.9389	0.2356	+0.01	-0.04	
1.1785	0.2357	1.1789	0.2353	0.03	-0.16	
0.2347	0.2357	0.2345	0.2356	-0.08	-0.04	
0.2347	0.4714	0.2345	0.4712	-0.08	-0.04	

*Average of three determinations.

Table.3: Determination of Ni (II) and Co (II) in Soil samples

Sample and	Amount in ppm				Relativ	e error
(ppm)	Certified Amount found (GSI) value by present (ppm) method*(ppm)			1(%)		
	Ni (II)	Co (II)	Ni (II)	Co (II)	Ni	Co (II)
S-18 20.20 Pb (II) 20.30 Zn (II) 88.85 Cu (II)	50.50	40.35	50.56	40.23	+0.11	-0.29

* Average of best three among five determinations.

Conclusions:

The present method, using HDMBHBH as spectrophotometric reagent for the simultaneous determination of Ni (II) and Co (II) in presence of Triton X-100 is simple, rapid, reasonably sensitive and selective. The present method is not laborious and there is no need of heating the components or pre-extraction. Large amounts Bromide, Iodide, Tartarate and Nitrate do not interfere in the present method.

This method was also free from the interference of strontium, arsenic, tungsten, lanthanum and other rare earth metals. Triton X-100 enhances the stability of the metal complexes. Further, the simultaneous determination of Ni (II) and Co (II)

carried out without the need of employing the simultaneous equation.

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[1]. Singh, R. B; Jain, P.; Singh, R. P. Talanta, 29 (2) (1982) 77. | [2]. Wahbi, A.M.; Abdine, H.; Balaiah. S. J. Assoc. Off. Anal. Chem., 60 (1977) 1175. | [3]. Bedair M.; Korany M.A.; El-Yazbi. Sci. F. A. Pharma., 54 (1986) 31. | [4]. Korany, M.A.; Wahbi, A.M.; Elasayed, M.A.; Mandour, S. Anal Lett., 17 (1984) 1373. | [5]. Hussain Reddy, K.; Chandrasekhar, K.B. Indian J. Chem., 40 (A) (2001) 727. | [7]. Chandrasekhar K.B.; Reddy K.H.; Sreenivasula Reddy, T. J. Indian Chem. Soc., 80(2003) 930. | [8]. Chandrasekhar K.B.; Reddy K.H. Indian J. Chem., 41(A (2002) 1643. | [9]. Reddy KH, Reddy DV. Q. chem., Rev. 1 (1985) 47. | [10]. Buseve AJ, Ivanov VM. Zh. Annal. Chim. 18 (1963) 208. | [11]. Piper CS. Soil and Plant Analysis. Bombay, India: Hans Publishers; 1966. | [12]. Reddy BK, Reddy KJ, Kumar JR, Kumar AK, Reddy AV. Highly sensitive extractive | [13]. Bertrand G, Nakamure H. Bulletin de la Société de Chimie Biologique. 16 (1934)1366. | [14]. Welch RM. The biological significance of nickel. Journal of Plant Nutrition.(1981) 345–356. | [15]. Heilmayer L. Spectrophotometry in Medicine. London, UK: Adam Hilger; 1943. |