

ACC NR: AP7001224

SOURCE CODE: UR/0066/66/000/012/0034/0036

AUTHOR: Mekler, V. Ya.; Zyk, S. L.

ORG: none

TITLE: Air conditioning for computers

SOURCE: Kholodil'naya tekhnika, no. 12, 1966, 34-36

TOPIC TAGS: electronic computer, air conditioning equipment

ABSTRACT: An air-conditioning system for computers is described (see Fig. 1) in which surface air coolers with direct evaporation are used. The air being forced into a computer is maintained at a constant temperature by bridge circuit 13 which, in winter and during periods of changing temperatures, acts on valve 18 for the first recirculation. The ratio of atmospheric to recirculated air is thereby varied. When a relative humidity of 40% is reached, humidity sensor 8, mounted in the air conduct, opens solenoid valve 7, and water starts to flow to the six sprayer nozzles (diameter, 1 mm). When 55% relative humidity is reached, the solenoid valve closes. When the atmospheric air temperature reaches 17C, the first cooling unit is switched on; at 17.5C the second is switched on, at 18C the third, and at 18.5C the fourth. If during summer the relative humidity of the incoming air is 70%, humidity sensor 9 gradually opens the valve for the second recirculation. At 65% humidity this valve closes. When the heat content of the atmospheric air exceeds that of the recirculated air, the valve for the first recirculation is fully opened by resistance

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ACC NR: AP7001224

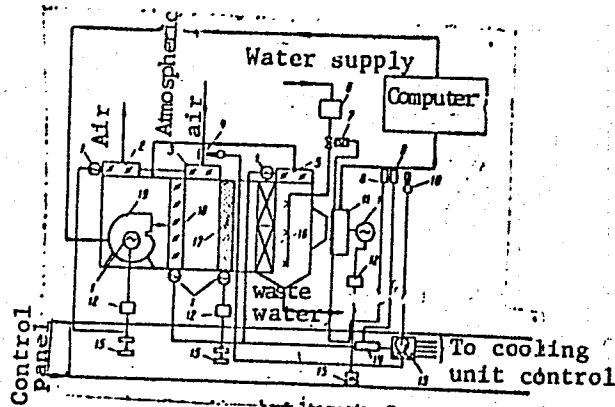


Fig. 1. Air-conditioning system with surface air coolers (schematic drawing)

- 1 - Electric drive; 2 - release vent;
- 3 - atmospheric air vent; 4 - resistance thermometer; 5 - valve for the second recirculation; 6 - water filter;
- 7 - solenoid valve; 8 - winter humidity sensor; 9 - summer humidity sensor;
- 10 - resistance thermometer; 11 - inflow ventilator; 12 - magnetic starter;
- 13 - recording hydrometer bridge;
- 14 - step interrupter; 15 - electric motor starting switch; 16 - surface air coolers; 17 - recording oil filter;
- 18 - valve for the first recirculation;
- 19 - recirculation fan.

thermometer 4 and the system operates in a closed air cycle. The four cooling units, mounted on a common condenser, have a capacity of 15,000 kcal/hr each. The main advantage of this air-conditioning system is its relatively small size and low electricity consumption. Orig. art. has: 3 figures.

SUB CODE: 09, 13/ SUBM DATE: none/ ATD PRESS: 5110

[JR]

Card 2/2

ZYK, W.

Some remarks concerning the starting and operation of a watertube boiler.  
p. 33, (GOSPODARKA CIEPLNA. ENERGETYKA PRZEMYSLOWA, Vol. 1, No. 6, Dec.  
1953, Warszawa, Poland)

SO: Monthly List of East European Accessions, (EEAL), LC, Vol. 4, No. 5  
May 1955, Uncl.

ZYK, Witold, mgr inz.

Significance and possibilities of limiting compressed air  
consumption in mines. Gosp paliw 11 no.6:205-207 Je '63.

ZYK, W.

Construction installations for hydraulic and sand pillars in deep mines, p. 55.  
(PRZEGLAD GORNICZY, Stalinogrod, Vol. 11, no. 2, Feb. 1955.)

SO: Monthly List of East European Accessions, (EEAL), LC, Vol. 4, No. <sup>6</sup>~~7~~, Jan. 1955,  
Uncl.

ZYKA, F.

Cervenka, K.; Rybar, F. Drawing and production of broaches, p. 409.  
STROJIRENSKA VYROBA, Prague, Vol. 3, no. 10, Oct. 1955.

SO: Monthly List of East European Accessions, (EEAL), LC, Vol. 5, No. 6,  
June 1956, Uncl.

METYS, Rene; ZYKA, Ivan

Chronic volvulus of the stomach with spontaneous recovery. Cesk.  
rentg. 13 no.2:101-103 Apr 59.

1. Rentgenologicke oddeleni nemocnice OUM v Susici, zast, pred-  
nosta dr. Rene Matys.

(STOMACH, dis.

torsion, spontaneous recovery, x-ray (Cs))

CZECHOSLOVAKIA

HERMAN, M; SULCEK, Z; ZYKA, J

1. Central Geological Institute (Geologisches Zentralinstitut) (for Herman ?); 2. Institute for Analytical Chemistry, Karlova University (Institut für analytische Chemie, Karlsuniversität), Prague (for ?)

Prague, Collection of Czechoslovak Chemical Communications, No 5, May 1966, pp 2005-2013

"Oxidimetric determination and identification of cobalt and manganese, using titration of a ferricyanide solution in a medium of certain aminoalcohols."



ZYKA, VACLAV

Czechoslovakia/Cosmochemistry - Geochemistry. Hydrochemistry,  $\mu$ D

Abst Journal: Referat Zhur - Khimiya, No 19, 1956, 61339

~~\_\_\_\_\_~~ Vaclav

Institution: None

Title: On Genesis and Hydrochemistry of Mineral Waters in the Prerov District

Original

Periodical: Ke generaci a hydro-geochemii mineralnich vod na Prerovsku. Sbor. sluko, 1951-1953 (1954), A1, 89-97; Czech; Russian and German resumé

Abstract: In Prerov District are found 2 types of mineral waters: (a) of sodium bicarbonate type and (b) of calcium-bicarbonate type. Of greatest interest are waters from Khrapyné of sodium bicarbonate type having the following composition (in mg/l): Cl 1,087.0, SO<sub>4</sub> 4.1, I 0.72, Br 2.9. Temperature of water 20.5°, air temperature 8.5°, pH 7.3, rH 22.7 (-10 mv), free CO<sub>2</sub> 30.8 mg/l, free H<sub>2</sub>S traces. Other waters are typically surface waters but their

Card 1/2

Czechoslovakia/Cosmochemistry - Geochemistry. Hydrochemistry, D

Abst Journal: Referat Zhur - Khimiya, No 19, 1956, 61339

Abstract: composition reveals influence of seepage of petroleum water although to a lesser extent than in the case of Khropyne water. In waters of sodium bicarbonate type predominates  $\text{CH}_4$  in the other  $\text{CO}_2$ .

Card 2/2

Geochemistry of mineral waters in the region Svitavy: Lucéne-Lovice Václav Zýka (Masaryk Univ., Brno, Czech.). *Geol. Práce* 3, 76-126 (1963) (German summary). Chem. analyses are given of 42 waters. These include NaCl-rich waters, NaHCO<sub>3</sub> waters derived from them by biogenic reduction of sulfate in the presence of org. resins, and Ca bicarbonate waters. Qual. tests showed the presence of Cu, Hg, V, Co, and As. Some of the waters are high in Br and iodine and may be related to oil-field waters. Michael Mebscher

27.11.61.  
CZECHOSLOVAKIA / Cosmochemistry. Geochemistry. Hydro- D  
chemistry.

Abs Jour: Ref Zhur-Khimiya, 1958, No 20, 67229.

Author : Zyka V., Juranek J.

Inst : Not given.

Title : On the Problem of Geochemistry of Mineral Waters  
of the Northern and North-Western Parts of Presov-  
skiy Kray.

Orig Pub: Sbirka praci vyzkumn. ust., 1956, E, No 17-21, 81-  
117.

Abstract: Based on the data developed by the geochemical sur-  
vey the major mineral water bearing localities are  
classified as follows: 1) sodium-bicarbonate type,

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ZYKA, V.

Hydrogeochemical zones in central Europe.

p. 383 , (ACTA GEOLOGICA), Vol 4, no. 3/4, 1957, in German  
Budapest, Hungary

SC: Monthly Index of East European Accessions (EEAI) LC. Vol. 7, No. 3,  
March 1958

ZYKA, VACLAV

CZECHOSLOVAKIA/Cosmochemistry. Geochemistry. Hydrochemistry. D.

Abs Jour : Ref Zhur - Khimiya, No 11, 1958, 35820

Author : Zyka, Vaclav

Inst : -

Title : The Distribution of Microelements in the Mineral Waters of Moravia.

Orig Pub : Rudy, 1957, 5, No 11, Prace Vyzkumn. ustavu, 1-6

Abstract : The mineral waters of Moravia and Slovakia were studied. It is shown that the majority of the waters studied belong, according to their mineral composition, to the group of calcium and sodium waters of the hydrocarbonate class. A content of Bi, Ag, Pb, Cu, Zn, and Fe microelements characteristic of ore deposits was determined spectrally.

Card 1/1

E N D  
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ZYKA, V.

GEOGRAPHY & GEOLOGY

Periodicals CASOPIS PRO MINERALOGII A GEOLOGII Vol. 3, no. 1, 1958

ZYKA, V. Geochemical types of mineral waters in Bohemia and their genesis. p. 103.

Monthly List of East European Accessions (EEAI) LC, Vol. 8, No. 5,  
May 1959, Unclass.

Country : Czechoslovakia  
Category : Cosmochemistry. Geochemistry. Hydrochemistry. D  
Abs. Jour. : Ref Zhur-Khimiya, No 6, 1959 19022  
Author : Zyka, V.  
Institut. :  
Title : Hydrochemistry and Genesis of Hydrogen-Sulfide Springs of the Gottwaldovska Region.  
Orig Pub. : Geol. sbor., 1958, 9, No 1, 129-157

Abstract : A hydrochemical study has made it possible to differentiate two types of waters: I. Sodium-hydrocarbonate type of high mineralization (M). II. Calcium-hydrocarbonate type of lower M. Data of 18 chemical analyses are included. Group I (mg/liter): dry residue at 180° 1113.6-4429.6, Cl 0.3-2.0, Br 0.7-4.1, H<sub>2</sub>S 0.7-28.5, pH > 7. Group II: dry residue 288.2-1052.6, pH > 7; H<sub>2</sub>S is present sometimes in small amounts. Waters of group I are waters of petroleum deposits, the chemism of which is affected to some extent by waters of infiltration. Group II -- their chemical composition is correlated with the presence of bitumen in the sedimentary rocks. Distribution of these 2 types of waters  
Card: 1/2



Country : Czechoslovakia D  
Category : Cosmochemistry. Geochemistry. Hydrochemistry.  
Abs. Jour. : Ref Zhur-Khimiya, No 6, 1959 19022  
Author :  
Institut. :  
Title :  
Orig. Pub. :  
Abstract : is shown on the appended map. About 20 analyses  
of water are listed. --- L. Flerova.

Card: 2/2

D-13

TRELJCKA, Zdenek, inz.; ZYKA, Václav, inz.

Contribution to the geochemistry of gold. Sbor VSB Ostrava 10  
n. 1/2:11-22 '64.

1. Institute of Mineral Raw Materials, Kutná Hora. Submitted  
February 1961.

ZYKA, Vaclav, RNDr., kandidat geologicko-mineralogickych ved

Do not look for difficulties, but for ways of performing  
the tasks. Geol pruzkum 6 no. 3:74-76 Mr '64.

1. Institute of Mineral Raw Materials, Kutna Hora.

NOVOTNY, J.; ZYKA, V.; KUDELASEK, Vl., dr.

Contribution to the chemism of Algonkian schists. Sbor VSB  
Ostrava 8 no.4:445-462 '62.

1. Ustav nerostnych surovin, Kutna Hora; Vysoka skola banska,  
Ostrava.

ZYKA, Vaclav, RNDr., kandidat geologicko-mineralogickych ved

Anomaly of the zinc content in mineral waters in the Bohemian limestone formations. Geolog pruzkum 5 no.2:49-50 F '63.

1. Ustav nerostnych surovin, Kutna Hora.

ZÝKA, V.

Czechoslovakia

Institute of Raw Materials -- Kutna Hora (Ústav  
nerostných surovin -- Kutná Hora)

Prague, Věstník ústředního ústavu Geologického,  
No 6, 1962, pp 455-457

"Interesting trace elements in the magnesium  
carbonate mineral waters in the Mariánské  
Lazně area (western Bohemia)."

COUNTRY : Czechoslovakia D  
CATEGORY :  
ABS. JOUR. : RZKhim., No. 1959, No. 85929  
AUTHOR : Zyka, V.  
INST. :  
TITLE : Geochemical Zoning of Mineral Waters of  
Central Europe  
ORIG. PUB. : Geol. sb., 1958, 9, No 2, 265-299

ABSTRACT : There have been determined 9 principal hydro-chemical types of mineral waters of Central Europe: HCO<sub>3</sub> - Ca, SO<sub>4</sub> - Ca, SO<sub>4</sub> - Fe, HCO<sub>3</sub> - Na, Cl - Na, Cl - Ca, Cl - Mg, SO<sub>4</sub> - Na, SO<sub>4</sub> - Mg. Their occurrence in connection with geological environment is reviewed. Bibliography 108 references. -- V. Konshin.

CARD:

COUNTRY : Hungary  
CATEGORY :

ABS. JOUR. : RZKhim., No. 22 1959, No. 78247

AUTHOR :  
INST. :  
TITLE :

ORIG. PUB. :

ABSTRACT : is the cause for the appearance of polymetallic and sulfide deposits which up to the present have been considered to be of hydrothermal origin. The majority of known native sulfur deposits were formed by the biogenic reduction of sulfates in oil field waters. The mixing of sodium-bicarbonate oil field waters with calcium-sulfate waters has led to the formation of tremendous travertine deposits at many points.

G. Volkov

CARD: 2/2

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ADAM, I.; DOLEZHAL, Ya.; ZYKA, Ya.

Use of hydroxy acids in polarographic analysis. Report 2:  
Determination of manganese in a sulfosalicylate medium. Zhur.anal.  
khim. 16 no.5:592-595 S-0 '61. (MIRA 14:9)

1. Karlov University, Prague, Czechoslovakia.  
(Manganese--Analysis) (Salicylic acid) (Polarography)

ZYKA, Ya.

Hydroquinone as a reagent in redoxometric titration. Zav.lab.  
27 no.9:1075-1079 '61. (MIRA 14:9)

1. Kafedra analiticheskoy khimii Karlova universiteta, Praga.  
(Hydroquinone) (Electrochemical analysis)

RYBACEK, J.; DOLEZAL, J.; ZYKA, J., prof. dr. mr. (Praha 2, Albertov 2030).

Reductometric determination of aromatic nitro compounds with  
ferrous sulfate in alkaline triethanclamine. Gesk. farm. 14  
no. 2:59-64 F '65.

1. Katedra analytické chemie přírodovědecké fakulty Karlovy  
University, Praha.

BERKA, A.; JIROVEC, J.; ZYKA, J., prof. dr. mr. (Praha 2, Albertov 2030)

Determination of organic compounds by oxidation with permanganate.  
I. Determination of some monosaccharides. Cesk. farm. 14 no.2:  
64-67 F '65.

1. Katedra analytické chemie přírodovědecké fakulty Karlovy  
University, Praha.

JANATA, J.; ZYKA, J.

Application of coulometry in constant current to measure velocity constants. Coll Cz Chem 30 no.5:1723-1727 My '65.

1. Institut für analytische Chemie, Karlsuniversität, Prague.  
Submitted April 10, 1964.

BUZKOVA, V.; MOLDAN, B.; ZYKA, J.

Mass analytic determination of iodide and bromide through lead (IV) acetate solutions. Coll Cz Chem 30 no.1:28-33 Ja '65.

1. Institut für analytische Chemie, Karlsuniversität und Zentralinstitut für Geologie, Prague. Submitted December 3, 1963.

BERKA, A.; DOLEZAL, J.; ZYKA, J.

Analytical examination of the reaction between hexacyanoferrates (III) and hydroquinone. Coll Cz Chem 30 no.2:695-607 F '65.

1. Institute fur analytische Chemie, Karls-Universitat, Prague.  
Submitted April 6, 1964.

ZYKA, J.

"Analysis of kalium" by Hans Tollert. Reviewed by J. Zyka. Coll  
Cz Chem 29 no.5:1332-1333 My '64.



ADAMEK, P.; DOLEZAL, J.; ZYKA, J.

Contribution to the theory of polarometric titrations based on oxidation-reduction reactions. Coll Cz Chem 28 no.8:2131-2137 Ag '63.

1. Abteilung für Instrumentalanalyse, Technische Hochschule für Chemie, und Institut für analytische Chemie, Karlsuniversität, Prag.

CZECHOSLOVAKIA

HORAK, P.; ZYKA, J.; Research Institute for Botanical Drugs, Chair of Analytical Chemistry at Charles University [Vyzkumny Ustav Prirodnich Leciv, Katedra Analyticke Chemie Karlovy University], Prague.

"Indirect Photometric Determination of Alkaloids after Chromatographic Separation. IV. The Chromatographic Separation of Tropic Alkaloids."

Prague, Ceskoslovenska Farmacie, Vol 12, No 8, 1963, pp 394-398

Abstract: Octanol was used as stationary and ammonia as mobile phase. 1-2% admixture may be determined, there is no loss of alkaloids. Hyosciamine cannot be separated from atropine. Detail instructions for the analysis are given, using amounts of 10-50 micrograms of the analyzed substance. The authors used the method for a successful separation of 14 opium alkaloids. 3 Figures, 16 Western, 2 Czech references.

BERKA, A.; FARA, M.; ZYKA, J.

Determination of glycerine in pharmaceutical preparations.  
Cesk. farm. 12 no.7:366-367 S '63.

1. Katedra analytické chemie Karlovy university, Praha.  
(GLYCERINE) (CHEMISTRY, PHARMACEUTICAL)

ZYKA, J.

"Analysis of potassium; classic and modern separation and determination methods with critical comparison of their efficiency" by Hans Tollert. Reviewed by J. Zyka.  
Chem listy 57 no.11:1197-1198 N '63.

HORAK, P.; ZYKA, J.

Indirect photometric determination of alkaloids after chromatographic separation. I. Precipitation of alkaloids using thallium complexes. Cesk. farm. 17 no.6:286-288 JI '63.

1. Vyzkumny ustav prirodnich leciv, Praha, - Katedra analyticky chemie Karlovy University, Praha.

(ALKALOIDS) (THALLIUM) (CHROMATOGRAPHY)  
(ATROPINE) (BELLADONNA) (SCOPOLAMINE)  
(COCAINE) (PHOTOMETRY)

HORAK, P.; ZYKA, J.

Indirect photometric determination of alkaloids after chromatographic separation. II. Photometric determination of thallium with crystal violet. Cesk. farm. 17 no.6:289-293 JI '63.

1. Vyzkumny ustav prirodnich leciv, Praha - Katedra analyticke chemie Karlovy university, Praha.

(THALLIUM) (GENTIAN VIOLET) (CHROMATOGRAPHY)  
(PHOTOMETRY)

CZECHOSLOVAKIA

ADANEK, P; DOLEZAL, J; ZYKA, J.

1. Department of Instrumental Analysis of the Technical Higher School of Chemistry, Prague; 2. Institute of Analytic Chemistry of Charles University, Prague

Prague, Collection of Czechoslovak Chemical Communications,  
Vol 8, 1963, pp 2131-2137

"Report on the Theory upon which the Oxydation-Reduction  
Reaction-Polarometric Titration is Based."

HORAK, P., Research Institute for Natural Drugs (Výzkumný ústav přírodních léčiv), Prague, and ŽYKA, J., Chair of Analytic Chemistry (Katedra analytické chemie), Charles University, Prague.

"Indirect Photometric Determination After Chromatographic Separation. III. Detection and Extraction of Alkaloids."

Prague, Ceskoslovenska Farmacie, Vol XII, No 1, September 63, pp 359-362.

Abstract [Authors' English summary, modified]: A complex of thallium salt with iodine and iodide (thallium concentration being 0.005 M) was used for detecting alkaloids on paper chromatograms. Results showed a low variation and the alkaloid precipitation was quantitative. The technique of the method is described. The method may be used for determining 10 to 60  $\mu$ g of tropane alkaloids on paper (photometric determination of the thallium components by means of crystal violet after oxidation with bromide). Three references, including 2 Czech.

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2

KOTOUCEK, M.; DOLEZAL, J.; ZYKA, J.

CSSR

Institute of Organic, Analytical and Physical Chemistry, Palacky University,  
Olomouc, and Institute of Analytical Chemistry, Charles University,  
Prague (for all)

Prague, Collection of Czechoslovak Chemical Communications, No 2, 1963  
pp 521-524

"Selective Proof and Volumetric Determination of Gold"

(3)

KOTOUCEK, M.; DOLEZAL, J.; ZYKA, J.

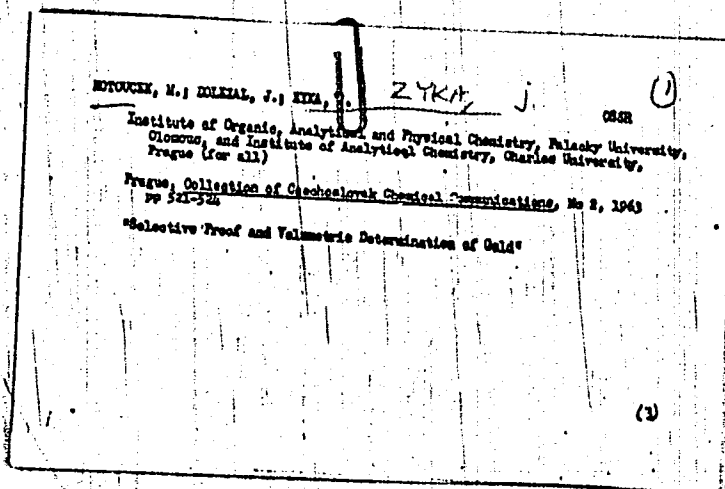
Selective test and volumetric gold determination.  
Coll Cz Chem 28 no.2:521-524 F '63.

1. Institut für organische, analytische und physikalische  
Chemie, Palcky-Universität, Olomouc und Institut für  
analytische Chemie, Karlsuniversität, Prag.

KRACMAR, J.; ZYKA, J.

Analytical study of anticholinergics of the carboxylic acid ester group.  
II. Other methods of determination of 2-(diisopropylamino)-ethyl  
xanthrene-O-carboxylate methobromide and 2-diethylaminoethyl xanthrene-  
9-carboxylate methobromide. Cesk. farm. 11 no.9:459-463 N '62.  
(METHANTHELINE) (CHEMISTRY, PHARMACEUTICAL)

ZYKA, J.



CZECHOSLOVAKIA

HORAK, P., Magister of Pharmacy, Candidate of Sciences, Research Institute for Natural Medicines (Vyzkumny ustav prirodnich leziv), Prague, and ZYKA, J., Chair of Analytic Chemistry (Katedra analyticko chemie), Charles University, Prague.

"Indirect Photometric Determination of Alkaloids after Chromatographic Separation. I. Precipitation of Alkaloids by Means of Thallium Complexes"

Prague, Ceskoslovenska Farmacie, Vol XII, No 6, July 1963, pp 286-289.

Abstract [Authors' English summary, modified]: A precipitation reagent of optimal composition can be prepared from one part of 0.1 N thallium sulfate and three parts of 0.1 N iodine in a 0.15 M potassium iodine solution. It precipitates alkaloids in a neutral or acid medium. Precipitates of the tropane alkaloids, which can be prepared in a crystalline form, are relatively water-soluble, and the sensitivity of the reagent is in conformity with the sensitivity of other alkaloids precipitation reagents. In view of the properties of alkaloid precipitates and the sensitive color reaction of thallium it is possible to use this precipitation reagent for an indirect determination of small quantities of alkaloids. Seven references, including 2 Russian.

ZYKA, J.

"Qualitative analytic chemistry" by A. Okac. Reviewed by J. Zyka.  
Coll Cz chem 27 no.10:2472-2473 0 '62.

DOLEZAL J.; NOVOZAMSKY, I.; ZYKA, J.

Indirect complexometric determination of sodium. Coll Cz Chem 27 no.8:  
1830-1834 Ag '62.

1. Institut für analytische Chemie, Karls-Universität, Prag.

CHRONOLOGIA

J. PRACMAN and J. SYMA, State Drug Control Institute, and Department of Analytical Chemistry, Charles University (Katedra analytické chemie Karlovy University) Prague.

"Analytical Study of Anticholinergics of the Carboxylic Acid Esters Group. Part 2. Other Methods of Determination of Propentheline and Methantheline."

Prague, Ceskoslovenska Farmacie, Vol 11, No 9, Sept 1962; pp 459-463.

Abstract [English summary modified]: Solution in ion exchange resin, with distilled water; potentiometric titration; UV spectrophotometry; latter is best for determination of the 2 drugs in tablets; titration in non-aqueous medium best for substances. Three graphs, 3 Czech periodical and 7 pharmacopoeial references.

1/1



KRACMAR, J.; ZYKA, J.

Analytical studies on anticholinergics from the group of carboxylic acid esters. I. Ultraviolet spectrophotometry of 2-di-isopropylamino-ethylxanthene-9-carboxylate methobromide and 2-di-ethylaminoethyl-xanthene-9-carboxylate methobromide. Cesk. farm. 10 no.9:440-455 '61.

1. Státní ústav pro kontrolu léčiv, Praha Katedra analytické chemie  
Karlovy university, Praha.  
(METHANTHELINE chem)

ZYKA, J.

dy

Program, Collection of Chemicals, Chemical Communications, no. 41, 1961, April 1962 (continued)

24. "A. AMERIKALIS of the Institute of Chemical Technology at A. Mikheev University in Perm; Poland; pp 979-986.

25. "In Japanese, Part CXXX, Composition of the Oil from the Leaves of *Madura* *sp.* (Linn.); Institute of Organic Chemistry and Biochemistry at the Chinese Academy of Sciences, Peking; pp 987-993 (English article); 6 NOV and V. UNKJ.

26. "In Polish, Part LXXX, The Primary Structure of Spun Beads of Polyacrylonitrile in the Presence of Organic Solvents; Institute of Polymer Chemistry and Technology of the Polish Academy of Sciences, Peking; pp 994-1000 (English article).

27. "The Primary Structure of the Primary Structure of Benzene and Acetylene, Part LXX, Institute of Organic Chemistry and Biochemistry at the Czechoslovak Academy of Sciences, Peking; pp 1001-1008 (English article).

28. "Dependence of IR Spectra, by T. H. H. on the Structure of Polymers; J. H. H. and E. S. S. of the Czechoslovak Academy of Sciences, Peking; pp 1009-1013 (English article).

29. "Contributions to the Chemistry of Amino Acids at Higher Temperatures," A. H. H. and V. K. K. from the Institute of Organic Chemistry and Technology of the Czechoslovak Academy of Sciences, Peking; pp 1014-1018.

30. "The Pyrolysis of Methyl Glycidyl Ether," J. V. V. and J. G. G. from the Institute of Organic Chemistry and Technology of the Czechoslovak Academy of Sciences, Peking; pp 1019-1023.

31. "The Polymers of the Acrylate and Methacrylate of Methyl Methacrylate," J. H. H. and J. G. G. from the Institute of Organic Chemistry and Technology of the Czechoslovak Academy of Sciences, Peking; pp 1024-1030.

32. "The Polymers of the Acrylate and Methacrylate of Methyl Methacrylate," J. H. H. and J. G. G. from the Institute of Organic Chemistry and Technology of the Czechoslovak Academy of Sciences, Peking; pp 1031-1038.

33. "The Condensation Polymerization of Styrene in Emulsion," J. H. H. and J. G. G. from the Institute of Organic Chemistry and Technology of the Czechoslovak Academy of Sciences, Peking; pp 1039-1045.

34. "The Direct Nitration of Nitrobenzene with Nitric Acid," A. H. H. and J. G. G. from the Institute of Organic Chemistry and Technology of the Czechoslovak Academy of Sciences, Peking; pp 1046-1052.

27

1/6



ZYKA, J.

27

Prague, Collection of Czechoslovak Chemical Communications, Vol. 87, No. 6, April 1964 (continued)

37. "Reductive Determination of Inorganic Oxidation States with Periodate Ions," B. ZEMEK and J. ZEMEK, *Journal of Analytical Chemistry at Charles University, Prague*, pp. 1031-1033.

38. "Organic Qualitative Analysis, Part XXII. The Mono-Determination of Carbon in Organic Substances by Means of Measuring the Electric Conductivity and by Using  $\text{CO}_2$  as a Combustion Catalyst," M. ZEMEK, J. ZEMEK and L. ZEMEK, *Journal of Analytical Chemistry at Charles University, Prague*, pp. 1033-1037.

39. "Method of Separating Natural Substances, Part V. The Determination of Nitrogen in Extracts from Peppercorns," P. HUBER, J. BUDAK, M. V. ZEMEK and L. ZEMEK, *Research Institute for Natural Drugs, Prague*, pp. 1037-1044.

40. "Spectrophotometric Determination of Heparin with the Modified Coumal and Sulfuric Acids," J. ZEMEK, *Journal of Analytical Chemistry at Charles University, Prague*, pp. 1045-1049.

41. "Gas-Field Chromatography. The Relation between the Delayed Elution Volume and the Molecular Structure of Organic Compounds," L. S. VILKIN, *Journal of Analytical Chemistry at the Chemical-Technological Institute in Prague*, pp. 1049-1054.

42. "Purification of an Unidentified Component of Wood Acetone. Part II. Determination of the Nature of the Isomers of Coproporphyrin I and III, Following Paper-Chromatographic Separation," V. BUDAK, *Research Institute for Natural Drugs, Prague*, pp. 1055-1061.

43. "Sulfonic Acid Compounds and their Analogues, Part XVII. Reaction of the Ethyl end of the Azo Analogue with Allylic Carbons," M. ZEMEK and J. ZEMEK, *Institute of Organic Chemistry and Biochemistry of the Czechoslovak Academy of Sciences, Prague*, pp. 1061-1068 (English article).

44. "Synthesis of 2-Phenylpyridine," I. SIK, *Department of Organic Synthesis at the Institute of Organic Chemistry, Czechoslovak Academy of Sciences, Prague*, pp. 1068-1075.

45. "Plant Substances. Part XIII. Quercetin, the Bitter Principles of *Quercus pedunculata* L.," M. SIK, *Institute of Organic Chemistry and Biochemistry, Czechoslovak Academy of Sciences, Prague*, pp. 1075-1083 (English article).

ADAM, I.; DOLEZHAL, Ya. [Doležal, J.]; ~~Zyka, Ya.~~ [Zyka, J.]

Use of hydroxy acids in polarographic analysis. Report No. 1: Half-wave potentials of certain ions in solutions of sodium salts of lactic, malic, and salicylic acids. Zhur. anal. khim. 16 no. 4:395-398 J1-Ag '61.  
(MIRA 14:7)

1. Charles University, Prague, Czechoslovakia.  
(Polarography) (Acids, Organic)

CHANG YE-SIYA; DOLEZHALL, Yan [Dolezal, J.]; ZYKA, Yaroslav, [Zyka, J.]

Potentiometric determination of cobalt with ferricyanide in a  
glutamic acid medium. Zhur.anal.khim. 16 no.3:308-312 My-Je '61.  
(MIRA 14:6)

1. Karlov universitet, Praga (Chekhoslovaskiya)  
(Cobalt--Analysis)  
(Potentiometric analysis)

CANG JE-SIA; DOLEZAL, J.; ZYKA, J.

Use of amino compounds in the polarography of inorganic substances.  
Part 9: Polarographic behavior of zinc in the environment of glutamic  
acid. Coll Cz Chem 26 no.7:1768-1774 J1 '61.

1. Institut fur analytische Chemie, Karlsuniversitat, Prag.

(Amino compounds) (Zinc) (Glutamic acid)

CHANG JE-SIA; DOLEZAL, J.; ZYKA, J.

Use of amino compounds in the polarography of inorganic compounds.  
VIII. Polarographic behavior of bi- and trivalent cobalt in glutamic-  
acid medium, Coll Cz Chem 25 no.12:3143-3152 D '60.  
(EEAI 10:9)

1. Institut für analytische Chemie, Karlsuniversität, Prag.

(Amines) (Polarigraph and polarography) (Cobalt)  
(Glutamic acid)



CZECHOSLOVAKIA

CUAN, Sun-Pao; DOLEZAL, J.; KALVODA, R.; ZYKA, J.

1. Institute for Analytic Chemistry, Karlova Univ. (Institut für analytische Chemie, Karlsuniversität); 2. J. Heyrovsky Polarographic Institute, Czechoslovak Academy of Sciences (Polarographisches Institut J. Heyrovsky, Tschechoslowakische Akademie der Wissenschaften), Prague

Prague, Collection of Czechoslovak Chemical Communications, No 12, Dec 1965,

"Use of oscillographic polarography in quantitative analysis. Part 23:  
Experiments in melting."

4

ZYKA, VACLAV

Mineralni prameny Gottwaldovskeho kraje.

Gottwaldov, Czechoslovakia, Krajske museum, 1957, 77p.

Monthly List of East European Accessions (MEAI), LC, Vol. 8, No. 9, September 1959.

Unclassified.

CATEGORY : Chemical Technology. Chemical Products and Their Applications. Chemical Processing of Natural Gases\*

ABS. JOUR. : RZhKhim., No 19, 1959, No. 69235

AUTHOR : Kudelasek, V.; Zyka, V.

INSTITUTE : -

TITLE : Microelements in Czechoslovakian Crude Oils

ORIG. PUB. : Shor. vedec. prací Vysoké školy báňské Ostrava, 1958, 4, No 4, 358-359

ABSTRACT : In addition to C, H, O, S, N, P, Br, I and Mo, the presence of which in the Czechoslovakian crudes could be logically presupposed, mineral ash samples, derived from these crudes, were analyzed spectrometrically revealing the presence of Na, K, Li, Ca, Sr, Ba, Mg, Al, Ti, Si, Fe, Mn, Au, Ag, Cu, Pb, Zn, Sn, Cr, Co, Sb, In, Bi, As, Ge, Ni, V, Pd, Be, Pt and Ga. Contents of the above metals in crude oils as well as in the Czechoslovakian asphalts

\*and Petroleum. Motor and Rocket Fuels. Lubricants.

Card: 1/2

ZYKA, V.

"Role of oil-field waters in the accumulation and distribution of Chemical elements." In German, p. 435.

ACTA GEOLOGICA. (Magyar Tudomanyos Akademia) Budapest, Hungary, Vol. 5, No. 3/4, 1958.

Monthly List of East European Accessions (EFAI) LC, Vol. 6, No. 6, June 1959.  
Uncl.

ZYKA, V.

"Hydrogeochemistry and genesis of the hydrogen sulfide springs in the Gottwaldov region."

p. 129 (Geologicky Sbornik, Vol. 9, no. 1, 1958, Praha, Czechoslovakia)

Monthly Index of East European Accessions (EEAI) LC, Vol. 7, no. 9,  
September 1958

C.A.

Use of polarometric titrations to determine small quantities of morphine. R. Kalvoda and J. Zjka (Charles Univ., Prague). *Časopis Českého Lékařského Spolku* 82, 134-40 (1949). Morphine can be detd. by oxidation to pseudomorphine with  $K_2Cr_2O_7$  and then titrating the unreacted  $K_2Cr_2O_7$  with  $Pb(NO_3)_2$ . This method allows the detn. of morphine over a broader concn. range than does the visual or potentiometric titration. Cf. C.A. 43, 5550A. Otilich Sebek

15

6382\* Some New Polarometric Titrations. (In English.) R. Kulvoda and J. Zyka. *Collection of Czechoslovak Chemical Communications*, v. 15, nos. 10-17, 1950, p. 630-638. Polarometric titrations for Cu<sup>2+</sup>, Tl<sup>+</sup>, Ag<sup>+</sup>, and WO<sub>3</sub> were worked out, using analytical precipitation reactions. A dropping-mercury electrode was applied and a saturated calomel electrode was used as a reference. Results are plotted. 21 ref.

ASAC-SLA METALLURGICAL LITERATURE CLASSIFICATION

GROUP	1	2	3	4	5	6	7	8	9	10	11	12	13	14	15	16	17	18	19	20	21	22	23	24	25	26	27	28	29	30	31	32	33	34	35	36	37	38	39	40	41	42	43	44	45	46	47	48	49	50	51	52	53	54	55	56	57	58	59	60	61	62	63	64	65	66	67	68	69	70	71	72	73	74	75	76	77	78	79	80	81	82	83	84	85	86	87	88	89	90	91	92	93	94	95	96	97	98	99	00
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CA

17

/ The polarometric determination of hypnotics as their mercury salts. R. Kalvoda and J. Zjka. *Inst. of Pharma-*

*ceutical Chem., Charles Univ., Prague, Czech.). Casopis Ceskeho Lékarskeho 63, 36-41(1950).—A study was made of the quant. detn. of barbituric acid derivs. (I) by conversion to the Hg salts. Initial work involved back-titrating excess Hg with NH<sub>4</sub> rhodanide soln. Subsequent work showed that direct titration of I with Hg salts was more satisfactory particularly when a recording polarograph was used to detect the equivalence point. Analyses were made successfully on the following compds., either pure or in pharmaceutical preps.: Phenylethyl-, diethyl-, isopropyl-(2-bromoallyl)-, *sec*-butyl(2-bromoallyl)-, cyclohexenylethyl-, diallyl-, 1-methylethylphenylbarbituric acids, Na isoamylethylarbiturate, and diphenylhydantoin.*

James L. Jeyl



ZYKA, J.

JINDRA, A.; KALVODA, R.; ZYKA, J.

Polarometric determination of certain pharmaceutically important substances with para-diazobenzene-sulphonic acid. Cas. cesk. lek. Ved. priloha. 63 no.7-8:106-110 1950. (CML 20:4)

ZYKA J.

CZECHOSLOVAKIA/Chemistry - Acridine Derivatives

Dec 50

"Setermination of Some Pharmaceutically Used Acridine Derivatives," A. Blazek, R. Kalvoda, J. Zyka, Inst Anal Chem and Inst Phar Chem, Charles U, Prague.

"Casopis Ceskeho Lekarnictva" Vol LXIII, No 9-12, pp 138-145

Developed polarometric detn methods for these substances in pure state and in tablet form, using 0.05 mol  $K_2Cr_2O_7$  pptg soln mercury drop cathode and satd calomel anode with equal parts of water and acetic acid buffer soln (pH 4.8). Detd Atebrin, Rivanol, and Trypaflavin in this manner. Ploarographic examn of Atebrin, Trypaflavin, Proflavin, and Rivanol in pH 4.8 buffer soln provided reproducible waves for quant detn in concn of  $10^{-5}$  to  $10^{-3}$  mol in pure and in tablet form.

181T12

ZYKA J.

CZECHOSLOVAKIA/Chemistry - Analytical; Alkaloids Dec 50

"Polarometric Titrations for Determination of Alkaloid Salts," P. Kalvoda, J. Zyka, Inst Phar Chem, Charles U, Prague.

"Casopis Ceskeho Lekarnictva" Vol LXIII, No 9-12, pp 219-221

Developed polarometric method for detn of mixt of alkaloid sulfates and chlorides. Used silver nitrate and lead nitrate solns for titration. Employed mercury drop electrode. Method is particularly suitable

181715

CZECHOSLOVAKIA/Chemistry - Analytical; Alkaloids (Contd) Dec 50

for detn of mixts of sulfate and chloride of same alkaloid and of anaesthetics in therapeutic preps.

181715

ZyKA, J.

KALVODA, R.; ZYKA, J.

Possibilities of determination of salol in combination with thymol  
and chloride of pilocarpine in combination with chloride of novocaine.  
Cas.cesk.lek. 63 no.11:123-125 15 June 50. (CJML 19:4)

1. Of the Institute of Pharmaceutical Chemistry, Charles University.

ZYKA, J.

New antibiotics, chloromycetin, aureomycin, and iridomyrmecin.  
Cas. cesk. lek. 63 no.15:172-173 15 Aug. 1950. (CIML 20:1)

CA

7

Polarographic determination of the purity of *p*-aminosalicylic acid. R. Kalvoda and I. Zika (Charles Univ., Prague). *Radiometer Polarographic* 1, 73-6(1951).--Mix 1 ml. of an 0.1 to 0.2% soln. of *p*-aminosalicylic acid with 1 ml. of 1% H<sub>2</sub>SO<sub>4</sub> and 8 ml. of water. Cool and diazotize with 0.5 ml. of a 1% soln. of KNO<sub>2</sub>. After 3 min, add 5 ml. of a 10% soln. of K<sub>2</sub>CO<sub>3</sub>, and after another 2 min. add 10 drops of 0.5% soln. of gelatin. Polarograph after removing dissolved O with N<sub>2</sub>. The diazotized *m*-aminophenol forms a compd. whose polarographic wave ( $\tau/2 = -0.09$  v. vs. the satd. calomel electrode) is proportional to the content of *m*-aminophenol. *p*-Aminosalicylic acid is converted to  $\beta$ -resorcylic acid and a wave appears which may be due to nitroso- $\beta$ -resorcylic acid (prior to the *m*-aminophenol wave,  $\tau/2$  not given). *p*-Aminosalicylic acid yields two readily measured waves with half-wave potentials =  $-0.149$  and  $-0.772$  vs. satd. calomel electrode. Gerakl Reed

ZYKA, J.

Czechoslovakia

CA:47:11663

with R. KALVODA

Charles Univ., Prague

"Polarometric determination of hypnotics (barbituric acid derivatives)  
by means of mercury salts."

Sbornik Mezinarod. Polarog. Sjezdu Praze, 1st Congr. 1951, Pt. III,  
Proc., 550-4 (in Czech)

CA

7

New polarometric titrations. II. Determination of thallium with potassium iodide and potassium dichromate. R. Kalvoda and J. Zřka (Charles Univ., Prague, Czech.). *Chem. Listy* 45, 82-3 (1951); cf. C.A. 45, 5550h. --Tl was detd. by the polarometric titration with KI (a) or  $K_2Cr_2O_7$  (b) with e.m.f.  $-0.0$  to  $-0.7$  v. as follows: (a) Mix the  $0.01-0.001$  M Tl soln. with  $KNO_3$  and acetone to make a vol. of 10 ml. which is  $0.1$  N in  $KNO_3$  and contains 30% acetone. Add 0.2 ml. of 0.5% gelatin soln., introduce  $N_2$  for 10 min. through the sample and then titrate with  $0.1-0.05$  M KI. (b) Shake up the sample which is  $0.01-0.001$  M in Tl to 50 ml., add 10% acetone, and  $0.5$  N  $KNO_3$ . III. Determination of silver with thiocyanate, ferrocyanate, and nitroprusside. *Ibid.* 83-4. --The polarometric titration of Ag was carried out with  $0.1$  M  $KSCN$ ,  $K_4Fe(CN)_6$ , and  $Na_2[Fe(CN)_5NO]$  at  $-0.3$  to  $-0.5$  v. The samples were made up to 50 ml. with the resulting concn.  $0.05-0.005$  M Ag and  $0.1$  N  $KNO_3$ . IV. Determination of tungstate with lead nitrate. *Ibid.* 84-5.  $WO_4^{2-}$  was detd. by the polarometric titration with  $0.1-0.001$  M  $Pb(NO_3)_2$  at  $-0.1$  v. The concn. of  $WO_4^{2-}$  in the sample contg. 50% EtOH and 50%  $0.1$  M  $KNO_3$  was  $0.05-0.0001$  M.

M. Hudlíky

7957



CA

17

Application of dead stop titrations in pharmaceutical analysis. I. Titration with potassium bromate. R. Kalvoda and J. Zýka (Charles Univ., Prague, Czech). *Chem. Listy* 45: 401-2 (1951).—Electrometric titration with 0.1 N  $KBrO_3$  was carried out with Pt electrodes and an elec. potential of 10 mv. Increase of current at the end of the titration was indicated by a galvanometer. Content of  $As_2O_3$  in Fowler's soln., and of Sb in K Sb tartrate were detd. in HCl soln. Ascorbic acid and dihydroxycodeinone were successfully titrated, whereas the method failed in the case of  $p$ -aminosalicylic acid, resorcinol, and other phenols.  
M. Huillek

CA

19

Polarographic estimation of purity of *p*-aminosalicylic acid. R. Kalvoda and J. Zylka (Univ. Prague). *Chem. Abstr.* 48: 21-6 (1952).—A method for the detn. of *m*-aminophenol (I), a toxic product produced by the decarboxylation of *p*-aminosalicylic acid (II), is worked out. By diazotization, II is converted to *p*-resorcylic acid (III); and diazotized I in alk. medium is converted to a complex, the diffusion current of which can be measured. The height of this current corresponds to the concn. of I and is not influenced by the presence of III. Another polarographic method is described for the detection of the therapeutic inactive *p*-aminosalicylic acid. Dagmar Hubáková

CA

17

The use of polarometric titrations in pharmaceutical analysis. III. The determination of glycerophosphates by means of lead nitrate. R. Kalvoda and J. Zřeka (Univ. Prague). *Ceskoslov. farm.* 1, 98-100(1952); *Ch. C.A.* 46, 4171c.—The described method is based on the pptn. of glycerophosphate with a 0.1 M soln. of  $Pb(NO_3)_2$  in 30% acetone made alk. to thymolphthalein. Dagmar Hubíková

JINDRA, A.; PALKOVA, M.; ZYKA, J.

Electrophotometric studies on certain antipyretics; determination of antipyrine. Cesk. farm. 1 no.8:350-355 Sept 1952. (CML 23:2)

1. Of the Institute of Pharmaceutical Chemistry of Charles University, Prague.

KALVODA, R.; ZYKA, J.

Use of polarometric titration in pharmacological analysis; mercurimetric determination of salicylates. Cesk. farm. 1 no.9:515-518 1952.

(CIML 23:4)

1. Of the Institute of Pharmacological Chemistry of Charles University, Prague.

ZYKA, JAROSLAW

Czechoslovakia

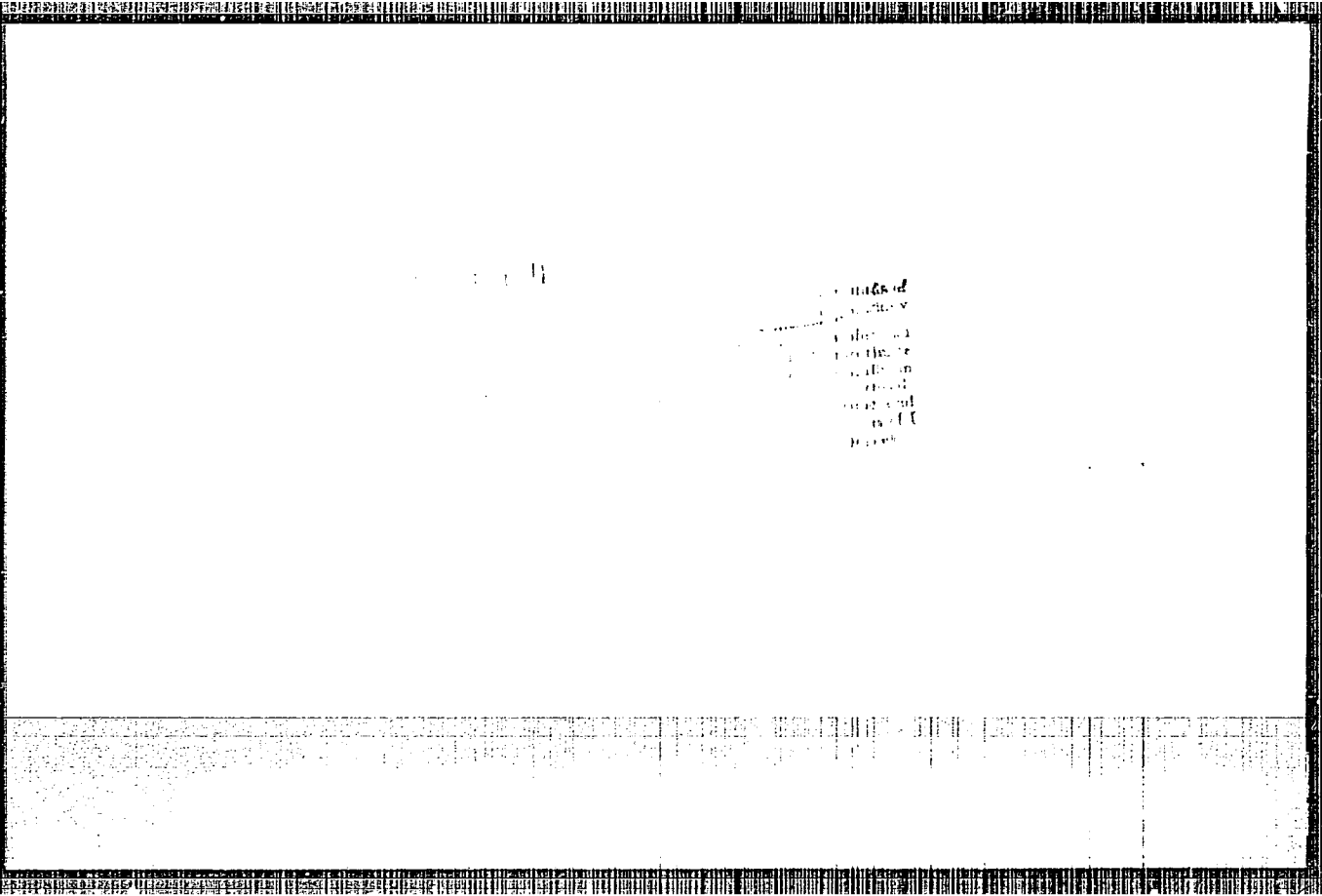
CA:47:11664

with ROBERT KALVODA

Karls-Univ., Prague.

"Polarographic titrations in pharmaceutical analysis and control."

Pharmazie 7, 535-42 (1952)



CA

17

Application of polarometric titrations in pharmaceutical analysis. I. Mercurimetric estimation of antipyrine. R. Kalvoda and J. Zilka (Charles Univ., Prague, Czech.). *Chem. Listy* 46, 64 (1952).--The estn. of antipyrine is based on measurement of a diffusion current produced by excess of  $\text{Hg}(\text{ClO}_4)_2$  during the titration with 0.05 N  $\text{Hg}(\text{ClO}_4)_2$  of antipyrine. The current is measured on the polarograph with a dropping-Hg electrode. The method is suitable for analysis of pharmaceuticals. II. Estimation of 4-butyl-1,2-diphenyl-3,5-pyrazolidinedione in pharmaceuticals. *Ibid.* 57.--A sample 0.05-0.3 g. of 4-butyl-1,2-diphenyl-3,5-pyrazolidinedione (Irgapyrine) is neutralized with  $\text{NaOH}$ , dil. to 40-50 ml., and titrated with 0.05 N  $\text{Hg}(\text{ClO}_4)_2$ . Excess of  $\text{Hg}(\text{ClO}_4)_2$  is indicated by the diffusion current measured between dropping-Hg cathode and satd. calomel anode. M. Hudlický



2/21/68

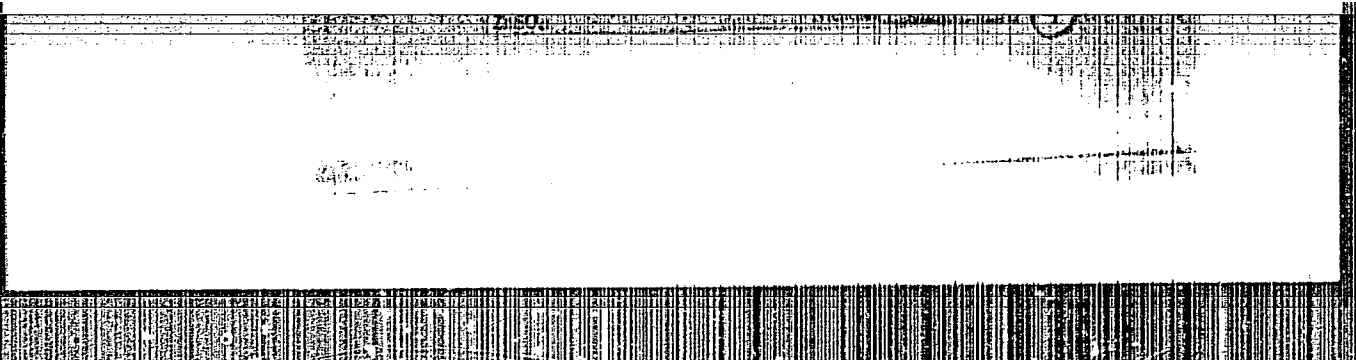
10

51

Comprehensive information in pharmaceutical literature  
is available in the following sources: 1. 2011

**"APPROVED FOR RELEASE: 09/01/2001**

**CIA-RDP86-00513R002065810002-7**



**APPROVED FOR RELEASE: 09/01/2001**

**CIA-RDP86-00513R002065810002-7"**

ZYKA, J.

Member of the Czechoslovakian Academy of Science prof. Oldrich Tomicek,  
Cesk. farm. 2 no.1:3-4 Jan 1953. (CIML 25:1)

3

218A, J.  
Chemical Abstracts  
Vol. 48 No. 5  
Mar. 10, 1954  
Pharmaceuticals, Cosmetics, and  
Perfumes

The use of polarimetric titrations in pharmaceutical analysis. VI. The possibility of determination of tartrates by the titration with lead salts. R. Kalvoda and J. Zýka (Univ. Prague). *Českoslov. farm.* 2, 14-15(1953); cf. C.A. 47, 5070i.—Tartaric acid and some tartrates were polarimetrically titrated with 0.1N  $Pb(NO_3)_2$  with a dropping Hg electrode as a cathode. D. Hubíková

CIHALIK, J.;DOLEZAL, J.;SIMON, V.;ZYKA, J.

Determination of thiopental with silver nitrate solution. Cesk. farm.  
2 no.2:43-47 Feb 1953. (GIML 24:4)

1. Of the Institute of Analytical Chemistry of Charles University, Prague.

BULTAS, Z.; JINDRA, A.; ZYKA, J.

Electrophotometric determination of 8-quinolinol and its pharmaceutical derivatives. Cesk. farm. 2 no.3:80-84 Mar 1953. (CLML 24:4)

1. Of the Institute of Pharmaceutical Chemistry of Charles University, Prague.

... complexometric determination of magnesium  
analysis. V. Determination of magnesium  
Prchal, J., Chvalik, J., Holodaj, V., Simpa, and J. Zyska  
*Czechoslov. Farmac.* 1963, 2:76, 184-185  
*Zh. Khim.* 1974, Abstr. No. 20359i. Magnesium  
sulphate or chloride is detected by a color reaction  
the addition of 5 to 8 ml of 10% sodium hydroxide  
solution to the test solution. The color reaction  
is observed in 10 to 15 minutes.  
A color reaction is observed in 10 to 15 minutes  
with 10% sodium hydroxide solution. The color  
reaction is observed in 10 to 15 minutes and the  
reaction is observed in 10 to 15 minutes. Magnesium  
maximum with  $\text{Na}_2\text{S}_2\text{O}_8$  and  $\text{Na}_2\text{S}_2\text{O}_8$  is  
observed. The color reaction is observed in 10 to 15 minutes.





JAROSLAV, ZYKA

ZYKA, Jaroslav

Chemical Abst.  
Vol. 48 No. 6  
Mar. 25, 1954  
Inorganic Chemistry

6  
Polarographic and polarometric study of some noble metals. III. Complexes formed by palladium and gold in solutions of certain amines. ~~Oldrich Tomkoc, Jaroslav Cihalik, Jan Dolezal, Vladimir Simon, and Jaroslav Zyka (Charles Univ., Prague, Czech). Chem. Listy 47, 383-6 (1953); cf. C.A. 47, 243a.~~—The behavior of the Au<sup>+++</sup> and Pd<sup>++</sup> ions in H<sub>2</sub>NH, H<sub>2</sub>NCH<sub>2</sub>CH<sub>2</sub>NH<sub>2</sub>, NH(CH<sub>2</sub>CH<sub>2</sub>NH<sub>2</sub>), N(CH<sub>2</sub>CH<sub>2</sub>OH)<sub>2</sub>, C<sub>4</sub>H<sub>9</sub>N<sub>3</sub>, and H<sub>2</sub>NCH<sub>2</sub>CH<sub>2</sub>NH<sub>2</sub>.HOOC(CH<sub>2</sub>OH)<sub>2</sub>.COOH. (D) solns. has been studied. For the polarographic detn. of Pd in the presence of Au, the 1 M soln. of I is recommended. B. Erdos

11-5-54

ZYKA, Jaroslav

Chemical Abst.  
Vol. 48  
Apr. 10, 1954  
Electrochemistry

13  
6

Polarographic and polarometric study of some noble metals. IV. ~~Polarographic~~ behavior of gold and palladium in solutions of ethylenediamine tartrate. Přemysl Betan, Jaroslav Čihák, Jan Doležal, Václav Šimon, and Jaroslav Zýka (Karlůva univ., Přírod. vědy, Chem. ústř., Praha, Czech). *Chem. Listy* 47, 189-191 (1953); cf. C.A. 48, 3182a. — The half-wave potential of complex Au<sup>3+</sup> ions depends on the concn. of ethylenediamine tartrate. The tartrate anion does not take part in the Au<sup>3+</sup> complex (D) formation. It is suppressed by the presence of Cl<sup>-</sup> ions. The *E*<sub>1/2</sub> of complex Pd<sup>2+</sup> ions is -0.56 v, against the satd. Hg<sub>2</sub>Cl<sub>2</sub> electrode. The formation of the Pd<sup>2+</sup> complex is substantially faster than that of I. V. Polarographic behavior of gold, palladium, and other metals in complex-forming electrolytes. *Ibid.* 1314-23. — The polarographic behavior of Au, Pd, Pb, Cd, Cu, Bi, As, Sb, Sn, W, Mo, U, Fe, Cr, Co, Ni, Mn, and Zn in various mixts. of ethylenediamine tartrate with the complexons I, II, and IV (C.A. 49, 10060s) is summarized in a table of half-wave potentials and in a chart of polarographic spectra. Au, Pd, and other components of dental alloys can be detd. simultaneously. E. Erdős



Z Y N A, U

CIHALIK, J.; DOLEZAL, J.; SIMON, V.; SERNY, V.; ZYKA, J.

Polarometric titration in pharmaceutical analysis. 7. Determination of cyanides in aqua laurocerasi. Cesk. farm. 3 no.4:136-137 Ap '54.

1. Z Ustavu pro chemii analytickou Karlovy university v Praze.  
(CYANIDES, determination,  
\*polarometric titration, in aqua laurocerasi)

3<sup>b</sup>

2184  
FRIBIL, R.; CIHALIK, J.; DOLAZAL, J.; SIMON, V.; ZYKA, J.

Complexometric titration in pharmaceutical analysis. VII. Determination of insulin zinc. Cesk. farm. 3 no.7:242-244 Sept 54.

1. Z Ustavu pro chemii analytickou Karlovy university v Praze.  
Z Vyzkumneho ustavu pro farmacii a biochemii v Praze.  
(INSULIN, determination,  
zinc insulin, complex titration)

226. Colorimetric determination of Conliten  
(thiazotazone). F. Ichn, R. Kalvoda and J. Eysa  
(Ceskosl. Farmaz., 1964, 3 (7), 244-246. *Neprilennyi*  
Za kaim, 1955, Abstr. No. 1600). This thione  
aldehyde formed a colored adduct with  
phenylhydrazine in methanol. The hydrazone form  
is more alkaline with respect to thiazotazone. It  
is on the present state of knowledge.



*Z YKH, H*

### CZECH

Colorimetric determination of (acetyl)levodopa and vanillo.  
R. Kocourek and J. Zeman. *Chem. Průmysl (Prague)* 1964, 4, 111.  
and vanillin the reaction with MeOH soln of 2,4-dinitro-  
phenylhydrazine (I) has been used. 0.1 ml. sample in  
MeOH (MeOH purified with  $H_2SO_4$ ) and 1 ml. MeOH, 1 ml.  
satd. MeOH soln of I, and 2 drops concd. HCl. Heat the  
soln. at 60-80° for an hr. After cooling, add 5 ml. 10%  
KOH soln. in MeOH soln. and 15 ml. MeOH. Measure  
the resulting wine-red color at 525 m $\mu$ . Lambert-Beer's  
law holds for  $10^{-5}$  to  $3 \times 10^{-4}$  M concn.  $\epsilon = 1.8 \times 10^4$  M $^{-1}$ cm $^{-1}$ .

*201-244*

JAKOSLAV ZYKA

10  
⑤

~~Polarographic and polarometric study of some noble metals. VI. Selective polarographic determination of gold.~~ Jaroslav Cihálek, Jan Doležal, Vladimír Šimon, and Jaroslav Zýka (Karlova Univ., Prague, Czech.). *Czech. Listy* 48, 28-31(1954); cf. *C.A.* 48, 3413f. — The 0.6M-ethylenediamine tartrate and 0.1M- $\text{Na}_2\text{C}_2\text{O}_4$  soln. is a suitable electrolyte for the selective detn. of Au. Some other metals can be detd. simultaneously. A graph of polarographic spectra and a table of half-wave potentials of 22 metals in this soln. are given. E. Erlösa

ME



ZYKA, J.

8

②

Volumetric determinations in strongly alkaline media.  
VIII. Titration of the peroxide with potassium ferricyanide. J. Vultejin and J. ZYKA (Karlova Univ., Prague, Czech. J. Chem. Listy 48, 919-921(1954); cf. C.A. 44, 930c; 46, 9782g.—The volumetric detn. of  $H_2O_2$  is based on its oxidation to  $O$  with  $K_3Fe(CN)_6$  in strongly alk. soln. Approx. 2% soln. of  $H_2O_2$  (0.5 ml.) is added to 5 ml. with 30% KOH and titrated with 0.1N  $K_3Fe(CN)_6$  with potentiometric detn. of the end point. The potential drop is approx. 400 mv., potential of inflexion  $\pm$  50 mv.

M. Fudický

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mlg

ZYKA, J.; MICHAL, J.

"Tetraethylthiuram Disulfide As An Analytic Reagent. I. New Specific Reaction for Copper", P. 915, (CHEMICKÉ LISTY, Vol. 48, No. 6, June 1954, Praha, Czech.)

SO: Monthly List of East European Accessions (EEAL), LC, Vol. 4, No. 3, March 1955, Uncl.

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Tetraethylthiuram disulfide is an analytical reagent. Photometric determination of cobalt. Mohr and Liska, *J. Chem. Ed.*, 1958, 35 (1), 1011-1020. The basis of the present method for the photometric determination of Co in the presence of other metals is the formation of an intense yellow-brown color with an absorption max. at 445 m $\mu$ , when Co reacts with tetraethylthiuram disulfide (I). In the presence of Hg<sup>2+</sup> an excess of the reagent should be used, for Hg<sup>2+</sup> also forms a stable, though colorless, complex with I. Coloured cations, if present in high concentration, interfere. The following procedure is recommended for the determination of Co in the presence of an excess of Fe<sup>3+</sup>. Remove all Fe by the precipitate with 10% NaOH, and filtering. The filtrate is diluted with water to 100 ml and a 25 ml aliquot with 10% NaOH is added. The solution is adjusted to pH 4.0 and a 0.05 g aliquot is used for the determination of Co. The absorbance is measured at 445 m $\mu$  in a 1 cm cell.

Handwritten signature or initials, possibly "J. Mohr", with a date "1958".

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