

BANIS, T.Ya.; VEBRA, A.I.; POZHELA, Yu.K.; REPSHAS, K.K. [Repsas, K.];
SHILAL'NIKAS, V.I. [Silalnikas, V.]

Heating of the current carriers in semiconductors in strong
electric fields. Radiotekh. i elektron. 7 no.9:1519-1522 S '62.
(MIRA 15:9)

1. Institut fiziki i matematiki AN Litovskoy SSR.
(Electric fields) (Semiconductors)

STYRO, B.I. [Styra, B.]; VEBRA, E.I.; GHOPAUSKAS, K.K. [Gopauskas, K.]

R: Radioactivity profiles in overcast air. Trudy AN Lit.SSR. Ser.
B. no.1:3-9 '64 (MIRA 17:7)

1. Institut geologii i geografii AN Litovskoy SSR.

STYRO, B.I.; VEBRA, E.I.; SHOPAUSKAS, K.K.; KHUNDZHUA, T.G.

Coagulation of radioactive aerosols with cloud droplets.
Soob. AN Gruz. SSR 33 no.1:61-67 Ju '64. (MIRA 17:7)

1. Institut geofiziki AN Gruzinskoy SSR. Predstavleno chlenom
korrespondentom akada. ii M.M. Miriashvili.

ACCESSION NR: AP4031103

S/0236/64/000/001/0003/0009

AUTHOR: Sty*ro, B.I.; Vebra, E.I.; Shopauskas, K.K.

TITLE: The radioactivity profile in clouded air

SOURCE: AN LitSSR. Trudy*. Seriya B, no. 1, 1964, 3-9

TOPIC TAGS: radioactivity, di tribution in air, distribution in cloud, radon decomposition, coagulation coefficient

ABSTRACT: The distribution of radioactivity in the air in a cloud was studied and the profile obtained was associated with the coagulation coefficient and other structural elements of the cloud. Measurements were made of the radioactivity in the air inside and outside clouds by probing from aircraft; and a series of experimental radioactivity profiles were obtained. There is significantly less radioactivity in cloudy air than outside the cloud zone, but the value never decreases to zero, and the variation of the radioactivity concentration in the cloud is insignificant. This residual radioactivity is closely connected with the microphysical characteristics of clouds and depends on the magnitude of the coefficient of coagulation of cloud droplets with radioactive aerosols. Thus

Card

1/3

ACCESSION NR: AP4031103

it appeared possible to determine this coefficient of coagulation from the radioactivity profile in the cloud. Formulas were derived for determining the change of radioactivity in clouded air ($N_s =$ the number of atoms formed by radon decomposition) according to the altitude:

$$N_s = \left[N_{s,0} - \frac{\lambda_{Rn} N_{Rn,0}}{\lambda_s + kn - \alpha w} \right] e^{-\frac{\lambda_s + kn}{w} z} + \frac{\lambda_{Rn} N_{Rn,0}}{\lambda_s + kn - \alpha w} e^{-\alpha z}$$

and when $\alpha = 0$, i.e., when radon concentration is constant:

$$\dot{N}_s = \left[N_{s,0} - \frac{\lambda_{Rn} N_{Rn,0}}{\lambda_s + kn} \right] e^{-\frac{\lambda_s + kn}{w} z} + \frac{\lambda_{Rn} N_{Rn,0}}{\lambda_s + kn}$$

where λ_s is the isotope decomposition constant; $N_{Rn,z}$ is the concentration of radon in the cloud and $N_{Rn,0}$ at the lower boundary of the cloud; w is the rate of vertical filtration of air through the cloud; z is the altitude; n , the concentration of the drops; λ_{Rn} is radon decomposition, and k is the coefficient of coagulation of drops with radioactive aerosols. The magnitude of the coefficient of coagulating radioactive aerosols with cloud drops is of the order of 10^{-5} 1/sec. Orig. art. has: 4 figures, 1 table and 8 equations.

Card 2/3

ACCESSION NR: AP4031103

ASSOCIATION: Institut geologii i geographii, akademii nauk Litovskoy SSR
(Institute of Geology and Geography, Academy of Sciences, Lithuanian SSR)

SUBMITTED: 20Jul63

ENCL: 00

SUB CODE: ES

NR REF SOV: 005

OTHER: 000

Card

3/3

ACCESSION NR: AP4018352

S/0251/64/033/001/0061/0067

AUTHORS: Sty*ro, B. I.; Vebra, E. I.; Shopauskas, K. K.; Khundzhua, T. G.

TITLE: On the coagulation of radioactive aerosols with cloud drops (Presented by A. M. Mirianashvili, corresponding member of the Academy on May 12, 1963)

SOURCE: AN GruzSSR. Soobshcheniya, v. 33, no. 1, 1964, 61-67

TOPIC TAGS: radioactive aerosol, cloud drop, coagulation coefficient, filtering system D2 O3 27 v, nuclear emulsion A 2, microscope system MBI 2, turbulent mixing, Brownian motion

ABSTRACT: A new experimental method is presented for determining the coagulation of radioactive aerosols with cloud drops. For measuring the radioactivity in the atmosphere an intake nozzle was installed above the overhead port of an aircraft at a distance of 0.5 m from the fuselage along the direction of motion of the aircraft. The air was filtered by a D-2-O3-27v system, using fiber filters. The system was so designed that the drops could not percolate into the filter (this was checked by using erythrozone). During the test flight 1 cubic meter of air was inducted in 6 minutes. The filter was then removed and brought in contact with nuclear photoemulsion of type A-2. After 20 hours of exposure, the system

Card 1/2

ACCESSION NR: AP4018352

was examined under a microscope of type MBI-2. The coefficient of coagulation was computed from the results to be on the order of 10^{-5} to 10^{-4} per second. The half-period of nonradioactive removal of aerosol was computed to be 1 to 2 minutes. Orig. art. has: 3 figures, 1 table, and 7 formulas.

ASSOCIATION: Akademiya nauk Gruzinskoy SSR, Institut geofiziki (Academy of Sciences Georgian SSR, Institute of Geophysics)

SUBMITTED: 12May63

DATE ACQ: 19Mar64

ENCL: 00

SUB CODE: ES

NO REF SOV: 008

OTHER: 002

Card 2/2

STYRO, B.I.; VEBRA, E.Yu.; SHOPAUSKAS, K.K.

Some physical characteristics of hot alpha-radioactive
aerosols. Atom. energ. 16 no.6:528-530 Je '64. (MIRA 17:7)

L 3106-66 EWT(1)/ENT(m)/FCC/EWA(h) GS/G#

ACCESSION NR: AT5023938

UR/0000/65/000/000/0207/0216

AUTHOR: Styro, B. I.; Vebra, E. Yu.; Shopauskas, K. K.

44,55 44,55 44,55

28
3+1

TITLE: Radioactivity, sizes, and composition of α -radiating aerosols

SOURCE: Nauchnaya konferentsiya po yadernoy meteorologii. Obninsk, 1964. Radioaktivnyye izotopy v atmosfere i ikh ispol'zovaniye v meteorologii (Radioactive isotopes in the atmosphere and their use in meteorology); doklady konferentsii. Moscow, Atomizdat, 1965, 207-216

TOPIC TAGS: nuclear meteorology, micrometeorology, atmospheric pollution, radioactive aerosol, hot particle, atmospheric boundary layer, temperature inversion

ABSTRACT: Basically, this paper is an elaboration of an earlier study of atmospheric samples originally collected at altitudes of 0-2 km over the Vilnius area (results published in Atomnaya energiya, no. 16, 1964), in which 20 α -radiating aerosol particles had been discovered. Reexamination of these samples revealed the presence of 42 additional particles of this type. These particles and one particle discovered in studying the radioactive fogs of 19 October 1963 were examined by microphotographic techniques to determine the sizes, composition, and degree of radioactivity. Orig. art. has: 5 figures and 1 table. [ER]

Card 1/2

L 3106-65

ACCESSION NR: AT5023938

8

ASSOCIATION: none

SUBMITTED: 28Apr65

ENCL: 00

SUB CODE: ES, NP

NO REF SOV: 002

OTHER: 001

ATD PRESS: 4101

PC

Card 2/2

ACC NR: AP6034790

SOURCE CODE: UR/0251/66/043/002/0327/0334

AUTHORS: Styro, B. I.; Vebra, E. Yu.; Shopauskas, K. K.; Khundzhua, T. G.

ORG: Institute of Geophysics, Academy of Sciences Georgian SSR (Institut geofiziki Akademiya nauk Gruzinskoy SSR)

TITLE: On the problem of determining the coefficient of turbulent diffusion along vertical concentration profiles of radon decay products

SOURCE: AN GruzSSR. Soobshcheniya, v. 43, no. 2, 1966, 327-334

TOPIC TAGS: atmospheric diffusion, radon, free atmosphere, atmospheric turbulence, alpha particle, nuclear emulsion, aircraft/ A-2 nuclear emulsion, LI-2 aircraft, Yak-12 aircraft

ABSTRACT: An experimental method for determining K_z along radioactivity profiles in the free atmosphere is described. For a layer of free atmosphere, it is assumed that the vertical distribution of the concentration of the i -th element of the radon chain is determined by solving a system of differential equations

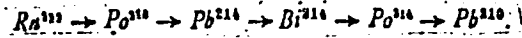
$$\frac{d}{dz} \left(K_z \frac{dN_i}{dz} \right) - \lambda_i N_i = 0,$$

$$\frac{d}{dz} \left(K_z \frac{dN_i}{dz} \right) - \lambda_i N_i + \lambda_{i-1} N_{i-1} = 0.$$

Card 1/3

ACC NR: AP6034790

The following chain of radioactive substances is considered:



Equations describing the profiles of the distribution of radon and three of its decay products are obtained:

$$N_i = \lambda_1 N_{1,h} \sum_{k=1}^i \frac{\prod_{l=1}^{i-1} \lambda_l}{\lambda_k \prod_{l=1}^{i-1} (\lambda_k - \lambda_l) \prod_{l=k+1}^i (\lambda_k - \lambda_l)} \exp \left\{ - \sqrt{\frac{\lambda_k}{K_s}} (\tau - h) \right\}$$

$\gamma = 1, 2, 3, 4$

In the experimental part, the free atmosphere is obtained by filtering air through fibrous materials. A-2 nuclear emulsion is used as the detector. The atmosphere was sounded in the areas of Tbilisi and Vilnius with LI-2 and YaK-12 aircraft. The radioactivity was measured according to the number of alpha tracks/cm² of emulsion (see Fig. 1). The advantages of the method are simplicity and high sensitivity. This paper was presented by Academician F. F. Davitaya on 06 November 1965.

Card 2/3

ACC NR: AP6034790

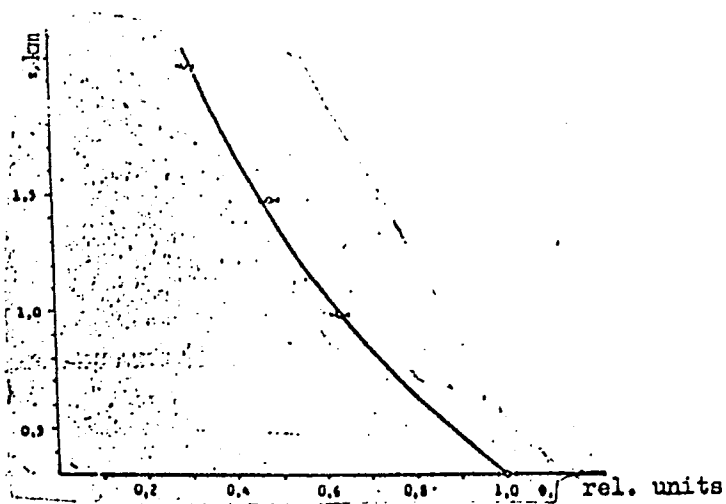


Fig. 1. Experimental curve of decrease in radioactivity with altitude (27 July 1962)

Orig. art. has: 12 formulas, 3 graphs, and 1 table.

SUB CODE: 20, 18, 04/ SUBM DATE: 06Nov65/ ORIG REF: 008/ OTH REF: 003

Card 3/3

STYRO, B.I.; VEBRA, E.Yu.; SHOPAUSKAS, K.K.

Determination of some parameters of the removal of natural
radioactive aerosols from the air. Izv. AN SSSR. Fiz. atm.
i okeana 1 no.12:1299-1309 D '65. (MIRA 19:1)

1. Submitted June 12, 1965.

ACCESSION NR: AP4041454

S/0089/64/016/006/0528/0530

AUTHORS: Sty*ro, B. I.; Vebra, E. Yu.; Shopauskas, K. K.

TITLE: On some physical characteristics of hot Alpha-active aerosol particles

SOURCE: Atomnaya energiya, v. 16, no. 6, 1964, 528-530

TOPIC TAGS: aerosol, fallout, alpha contamination, radon, neptunium

ABSTRACT: The characteristics of about 20 α -active hot aerosol particles found in samples gathered near Vil'nyus at altitudes 0--3 km are described. The particles are characterized by "fans" of alpha tracks. The fan tracks corresponding to the highest energies are probably the daughter products of radon decay or some products of the neptunium family. The activity of the hot particles was determined from the number of tracks, and the dimensions could be determined by making certain assumptions relative to the particle isotopic composi-

Card 1/3

ACCESSION NR: AP4041454

tion. The few actually measured particle sizes lie between the values obtained when the fan is assumed to be produced by U^{235} and Pu^{239} respectively. It can therefore be concluded that the hot aerosol particles are not uniform and consist of isotopes such as U^{235} , U^{238} , Th^{232} , and Pu^{239} . It is concluded that their isotopic composition needs further study. Orig. art. has: 3 figures, 2 formulas, and 1 table.

ASSOCIATION: None

SUBMITTED: 05Aug63

ENCL: 01

SUB CODE: NP, CB

NR REF SOV: 002

OTHER: 002

Cont 2/3

ACCESSION NR: AP4041454

ENCLOSURE: 01

Some properties of hot alpha-active aerosol particles

Legend:

- 1 - particle no.
- 2 - sampling flight altitude
- 3 - exposure, hr
- 4 - number of alpha tracks in fan
- 5 - activity, Cu
- 6 - dia. of act. part., microns
- 7 - measured
- 8 - calc. from U-235 radiation
- 9 - calc. from Pu-239 radiation
- 10 - ground level

1 Номер частицы	2 Высота полета при отборе пробы, м	3 Продолжительность облучения, ч	4 Число треков в секре	5 Активность, кюри	6 Диаметр активной частицы, мкм		
					7 измеренный	8 вычисленный по α-излучению U ²³⁵	9 по β-излучению Pu ²³⁹
1	1000	39,3	688	2,6·10 ⁻¹³	12	22,0	
2	1000	74	161	3,3·10 ⁻¹⁴	8	11,6	
3	1000	74	38	7,8·10 ⁻¹⁵	4	7,0	
4	1000	235	37	2,4·10 ⁻¹⁵	—	4,8	
5	1000	235	26	1,7·10 ⁻¹⁵	—	4,2	
6	1000	235	17	1,1·10 ⁻¹⁵	—	3,7	
7	1000	235	12	7,7·10 ⁻¹⁶	—	3,3	
8	1000	235	10	6,4·10 ⁻¹⁶	—	3,1	
9	1000	235	8	5,1·10 ⁻¹⁶	—	2,9	
10	1000	235	6	3,8·10 ⁻¹⁶	—	2,6	
11	1000	235	10	6,4·10 ⁻¹⁶	—	3,1	
12	1000	235	5	3,2·10 ⁻¹⁶	—	2,5	
13	870	240	400	2,5·10 ⁻¹⁴	7	10,6	
14	870	240	350	2,2·10 ⁻¹⁴	—	10,1	
15	870	240	20	1,8·10 ⁻¹⁵	—	4,4	
16	870	240	22	1,4·10 ⁻¹⁵	—	4,0	
17	870	240	7	4,4·10 ⁻¹⁶	—	2,7	
18		240	16	1,0·10 ⁻¹⁵	—	3,6	
19	на уровне земли	240	400	2,5·10 ⁻¹⁴	—	10,6	
20	10	240	6	3,1·10 ⁻¹⁶	—	2,4	

Card 3/3

1 18999-66 ENT(1)/ENT(m)/FCC/EMA(h) GW
ACC NR: RP6011119

SOURCE CODE: UR/0362/65/001/012/1299/1309

AUTHOR: Styro, B. I.--Styra, B. J.; Vebra, E. Yu.--Vebra, E. J.; Shopauskas, K. K.

ORG: none

34
B

TITLE: Determination of some parameters of removal of natural radioactive aerosols from the air

19

SOURCE: AN SSSR. Izvestiya. Fizika atmosfery i okeana, v. 1, no. 12, 1965, 1299-1309

TOPIC TAGS: atmospheric radioactivity, atmospheric cloud, gas filter, radioactive aerosol

ABSTRACT: The authors describe a method for measuring the radioactivity of air in the free atmosphere and within cloud systems by its filtration through porous filters and the screening of drops. The actual method was described in a previous paper by the author (Tr. AN LitSSR, Seriya B, 1(36), 1964). The authors have developed the theory of the experiment and derived formulas for computing the parameter of nonradioactive removal Λ of radioactive aerosols in cloud droplets. Two methods are proposed for determining Λ : on the basis of the profile of the concentration of radioactive substances in the air in a cloud and outside it and on the basis of disruption of radioactive equilibrium between the daughter products of radon decay in the cloud zone. On the basis of Λ and data in the literature on the drop concentration in a cloud the authors have computed the value of the coagulation coefficient K of radioactive

Card 1/2

UDC: 551.510.721

L 18899-66

ACC NR: AP6011119

aerosols on droplets, which on the average is equal to $3 \cdot 10^{-6}$ cm³/sec.
One shortcoming of the study is the assumption that the processes in the cloud zone are stationary, but at present this formulation is necessary.
Orig. art. has: 4 figures, 21 formulas, and 1 table. [JPRS]

SUB CODE: 18, 04 / SUBM DATE: 12Jun65 / ORIG REF: 009 / OTH REF: 002

Card 2/2 mc.

VEBRAS, E.A.

Scientific work of D.A. Gol'dgammer. Ist. i metod. est. nauk
2:286-297 '63. (MIRA 16:11)

VEERAS, E.A.

From the history of the development in Russia of the electro-
magnetic theory of light (works of D.A.Gol'dammer). Nauch.
trudy Tul.gor.inst. no.3:180-193 '61. (MIRA 16:4)
(Electromagnetic theory)

ACC NR: AP6034772 (A) SOURCE CODE: UR/0362/66/002/010/1055/1063

AUTHOR: Styro, B. I.; Vebrene, B. K.

ORG: none

TITLE: Preliminary results and methods of measuring radioactivity of particles of precipitation

SOURCE: AN SSSR. Izvestiya. Fizika atmosfery i okeana, v. 2, no. 10, 1966, 1055-1063

TOPIC TAGS: radioactivity, ^{measurement} ~~meteorology~~, evaporation rate, ~~particles~~, ~~precipitation~~ *rain, snow, radioactivity fallout, atmospheric radioactivity*

ABSTRACT: This article describes the method of measuring the radioactivity of raindrops and snowflakes by capturing them on polished steel plates or chromatographic filter paper with subsequent contact with an emulsion. The number of nonradioactive raindrops during a continuous rain, based on a computed measurement angle of 2π was found to be equal to 30—40% of the total number of raindrops. The radioactivity of individual raindrops generally increases with size with the rate of increase leveling off. It is shown that the specific radioactivity of raindrops increases exponentially with decreasing

Card 1/2

UDC: 551.510.7

ACC NR: AP6034772

droplet size. Based on this, a computation of the mean value of the relative rate of evaporation of raindrops falling to the earth's surface is made possible. Orig. art. has: 3 figures, 3 tables, and 8 formulas.

SUB CODE: 04/18/ SUBM DATE: 26Feb66/ ORIG REF: 009/ OTH REF: 004

Card 2/2

LAZAROIU, D.F., ing.; UDRESCU, Petre; VECAS, Alexandru; RADU, Emilia; CARACASIAN, E., ing.; BANDI, Fr.; TAIGAR, S., ing.

Present problems on establishing labor norms. Probleme econ 17 no.7: 151-156 J1 '64.

1. Direktor, "Electronica" Plant, Bucharest (for Lazaroiu).
2. Director, Iprofil Bucharest (for Udrescu).
3. Head of the Department of Labor Organization, Iprofil Bucharest (for Vecas).
4. Director, "Rascoala din 1907" Textile Enterprise, Bucharest (for Radu).
5. Chief Engineer, "Rascoala din 1907" Textile Enterprise, Bucharest (for Caracasian).
6. Director, "Bucuresti" Glass Factory (for Bandi).
7. Chairman of the State Committee for Labor and Wage Problems (for Taigar).

VECEK, Ales, inz.

Opening of a heat glazed glass factory in the United States.
Skalr a keramik 14 no.8:236 Ag '64.

VECEK, B.; VOSTATEK, M.

Carbon dioxide. Zvaranie 12 no.1:11-13 Ja '63.

1. Technoplyn, n.p., Praha (for Vecek). 2. Vychodoceske chemicke zavody Synthesia, n.p., Pardubice - Senti (for Vostatek).

VECERA, M.; JURECEK, M.

"Identification of organic substances. III. Studies on alkyl thionium salts", P. 722., (CHEMICKÉ LISTY, Vol. 46, No. 12, Dec. 1952, Praha, Czechoslovakia)

SO: Monthly List of East European Accessions, (EEAL) LC, Vol. 4, No. 6, June 1955, Uncl.

The action of the Grignard agents on the amide group.
 XVIII. Isomerism of pyrrolines. Rudolf Lukes and Miroslav Vozelka (Vysoká škola chem., Prague, Czech.). *Chem. listy* 51(1953); cf. C.A. 47, 11133f. — A series of new 1-methyl-2,5-dialkyl-2-pyrrolines was prepd. from Grignard reagents and 1-methyl-2-alkyl-5-pyrrolidinone. The 3rd possible isomer of 1,2-dimethyl-5-ethylpyrrolin was prepd. by the reduction of the corresponding pyrrolin with Zn and AcOH. The pyrrolines were characterized as perchlorates and binary picrates, the pyrrolidines as normal picrates. MeMgBr (0.5 mole) in 600 ml. ice-cooled Et₂O, stripped of excess MeBr, treated with 0.1 mole 1,2-dimethyl-5-pyrrolidinone in 100 ml. Et₂O started to form a ppt. after 15 min. and a gas was evolved; after 12 hrs., the mixt. was treated with ice and solid Ba(OH)₂, the liberated bases were steam distd., the distillate was neutralized with 85 ml. N HCl, the soln. evaporated to a syrup in vacuo, the residue treated with 9.1 g. NaClO₄ in 15 ml. H₂O giving 5.3 g. HClO₄ salt of 1,2,5-trimethyl-2-pyrrolin (I), m. 217-18° (from EtOH); binary picrate, m. 203.5-4° (from H₂O); H₂PtCl₆ salt, m. 196° (decompn.) (from 80% EtOH acidified with HCl). From the mother liquors after the sepn. of the perchlorate of I was isolated, 1,2,2,5-tetramethylpyrrolidinone picrate, m. 224-6° (decompn.) (from EtOH). Similarly 34 g. 1,2-dimethyl-5-pyrrolidinone and EtMgBr gave 38 g. of the 1,5-dimethyl-2-ethyl-2-pyrrolinone

HClO₄, m. 223-4°; free base (II), liberated from the HClO₄ salt, b. 98-101° (in H stream), d₄ 0.8625, n_D²⁰ 1.4766; binary picrate, m. 170.5-20° (from H₂O); H₂PtCl₆ salt, m. 194° (decompn.) (from 80% EtOH); H₂AuCl₄ salt, m. 83° (from EtOH). 1,5-Dimethyl-2,2-diethylpyrrolidinone, b. 107-8°, d₄ 0.8437, n_D²⁰ 1.4160, was obtained as a liquor after the sepn. of HClO₄, 1,2-Dimethyl-5-pyrrolidinone and PrMgBr gave 1,5-dimethyl-2-propyl-2-pyrrolinone HClO₄ salt, m. 117-18° (from EtOH); binary picrate, m. 105-6° (from H₂O); H₂PtCl₆ salt, m. 183°. The normal method for the prepn. of 1,2-dimethyl-5-ethyl-2-pyrrolin (III) gave a poor yield and was therefore changed. 1-Methyl-2-ethyl-5-pyrrolidinone (0.15 mole) in 100 ml. Et₂O dropped into a boiling soln. of 0.3 mole MeMgBr in 600 ml. Et₂O, the mixt. boiled 2 hrs., decompd. with ice, acidified with HCl, the neutral portions steam distd., the residue alkalinized with 50% NaOH, the bases liberated, steam distd., and the distillate neutralized with 100 ml. N HCl, evaporated in vacuo, and treated with NaClO₄ pptd. 18 g. (63%) III. HClO₄, m. 117.5-118°; free base, b. 85-8°, d₄ 0.8185, n_D²⁰ 1.5007; binary picrate, m. 187°; H₂AuCl₄ salt, m. 123°. 1,2,5-Trimethyl-5-ethylpyrrolidinone (from the mother liquor of the less successful prepn.) gave a picrate, m. 133°. 1-Methyl-2-ethyl-5-pyrrolidinone and EtMgBr gave 1-methyl-2,5-

(over.)

Rudolf Lukes

2/2

diethyl-2-pyrrolino (IV); perchlorate, m. 127-8°; H_2PtCl_6 salt, m. 194° (decompn.). The binary picrate ($C_{11}H_{13}O_6N_3Na$) (IVa) of IV was prepd. 1-Methyl-2,2,5-triethylpyrrolidine (from the mother liquors) picrate m. 95-8°. 1-Methyl-2-ethyl-5-pyrrolidinone and $PbMgBr$ gave only a small amt. of 1-methyl-5-ethyl-2,2-dipropylpyrrolidine; picrate, m. 129-8.5°. 1-Methyl-2-ethyl-5-pyrrolidinone and $PbMgBr$ gave perchlorate of 1-methyl-5-ethyl-2-phenyl-2-pyrrolino- $HClO_4$, m. 136°; picrate, m. 83° (from dil. EtOH). 1-Methyl-2-phenyl-5-pyrrolidinone and $PbMgBr$ gave 1-methyl-2-ethyl-5-phenyl-2-pyrroline; picrate m. 124-5°. 1,2-Dimethyl-5-ethylpyrrole (8.6 g.) was reduced with 10 g. Zn in 30 ml. H_2O to which HCl was added gradually; steam distn. and treatment of the distillate with 10 g. Na picrate gave the normal picrate, m. 147-8° (from EtOH), and binary picrate, m. 180-7° (from EtOH), of 1,2-dimethyl-5-ethyl-3-pyrroline; free base, bp 45-50°, $d_{20} 0.8404$, $n_D^{20} 1.4570$; perchlorate, m. 215°; H_2PtCl_6 salt, m. 173° (decompn.). Similar reduction of 1-methyl-2,5-dieethylpyrrole with Zn and HCl yielded, at 40°, after treatment with Na picrate, a normal picrate, m. 110-11°, and a binary picrate, m. 185°, giving no depression with IVa. Also in *Collection Czechoslov. Chem. Commun.* 19, 263-71 (1954) (in German). M. Hudlický

Večera, Miroslav

6

Identification of organic compounds. V. Reaction rate of halogen derivatives with thiourea. Miroslav Večera and Miroslav Jureček (Výzkumný ústav pro syntezu, Pardu-bice-Rybitví, Czech.). *Chem. Listy* 47, 1342-7 (1953), cf. *C.A.* 48, 3201k. — The rate of interaction of organic halides with thiourea was followed by measuring the cond. of the 0.1M solns. of halides and CS(NH₂)₂ in Me₂CO. Since the reactivity of a halogen depends on the nature and type of linkage of the halogen, and determines the rate of inter-change with CS(NH₂)₂ which causes the increase of cond., it is possible, by means of the cond. measurements, to est. the nature and type of halogens. Specific cond. (in 10⁻¹ D-1 cm. ⁻¹) at 50° after 3 hrs. of the reaction mixt. in 0.95M solns. in acetone of org. halides with 0.05M CS(NH₂)₂ given: MeI 13000, CH₃CH₂Br 4100, PhCH₂Br 3890, CH₃I 3700, Br(CH₂)₂Br 1000, EtBr 1400, C₂H₅Br 1250, PrBr 1240, C₃H₇Br 1200, BuBr 1150, Br(CH₂)₃Br 700, iso-BuBr 680, *tert*-BuBr 540, 2,4-(O₂N)₂C₆H₃Cl 510, *sec*-BuBr 370, *p*-O₂NC₆H₄Cl 40, PhBr 30, *m*-O₂NC₆H₃Cl 20, *o*-O₂NC₆H₃Cl 20. VI. Identification of organic sulfides. M. Jureček, M. Večera, and J. Casparik. *Ibid.* 1410. — A method for the identification of org. sulfides is devised using the reaction of the sulfide with *p*-BrC₆H₄COCH₂Br (I), and the trans-formation of the sulfonium bromide to less sol. HClO₄ salt. Refluxing 0.03 mole EtS (b.p. 62°) and 0.03 mole I in 30 ml. MeOH 1 hr. gave [p-BrC₆H₄COCH₂SEt]⁺Br⁻, m. 114° (from MeOH) which was transformed into the HClO₄ salt, m. 172° (from EtOH). Similarly was prepd. from Bu₂S (b.p. 65°), [p-BrC₆H₄COCH₂SBu]⁺Br⁻, m. 66-67°; HClO₄ salt, m. 129°. M. Hudlický

VECHRA, M.; GASPARIC, J.

Analysis of drugs. Cesk. farm. 2 no.10-11:368-381 Nov 1953.

(CJML 25:5)

1. Of the Research Institute of Organic Syntheses, Pradubice-Rybitvi.

VECERA, M.

LUKES, R.; VECERA, M.

Leukart's reaction of some ketone acids and their derivatives. Part 2.
[in German with summary in Russian]. Sbor. Chekh. khim. rab. 18 no. 2: 243-247
Ap '53. (MLRA 7:6)

1. Institut obshchey organicheskoy khimii Prazhskogo Politekhnicheskogo
instituta. (Ketone acids)

LECERA MIROSLAV

Identification of organic compounds...
 tion of sulfidic Miroslav...
 Chem. List 43, 542-8 (1954)...

again were transformed to...
 suitability of the salts for identification...
 in the order given. Best yields of the bromides were...
 obtained by refluxing 10-30 min. a soln. of...
 H₂COCH₂Br and 0.01... MeOH on a
 steam bath, adding 200 ml. and...
 stand 30 min. in an ice box. The precipitate...
 an aq. soln. of the bromides with satd. aq. soln. of picric
 acid, the HClO₄ salts by pptg. the bromides with 10% aq.
 NaClO₄. The necessary sulfides were prepd. as follows:
 40 g. BuBr, 25.5 g. C₂S(NH₄)₂, and 50 ml. MeOH were re-
 fluxed 6 hrs., the MeOH was distd. off from the steam bath,
 the residue was dild. with H₂O to 80-100 ml., and the soln.
 was heated to the b.p., treated with 20 g. NaOH in 160 ml.
 H₂O, and boiled 15 min. The mercaptan layer deposited

after cooling was dissolved by adding...
 17 g. NaOH; the mix. was heated to...
 50 g. MeOH...
 washed with...
 MeOH...
 from Et...
 105, 119, 105 BuBr-S, 122-...
 BuBr-S, 142-4, 99.5-100, 103, 99.4-5, BuBr-S, 167.8...
 145.1, 145.1, 145.1, 145.1, BuBr-S, 145.1, 145.1, 145.1, 145.1...
 sulfidic...
 M. H. H. H. H.

VECERA, M.

CZECH

Colorimetric determination of small amounts of hydrazo-
benzene. Michoslav Vecera and Jaromir Petrášek (Vyz-
kumny ústav pro svastky, Pardubice-Rybitví, Czech.).
Chem. Listy 48, 1351-3 (1954).—A method for detg. small
quantities of PhNH₂ (I) is based on its rearrangement to
benzidine, and on the colorimetric detn. of the products of
coupling the diazotized benzidine with *N*-1-naphthylethyl-
enediamine (II). A sample contg. 1-5 mg. I is allowed to
stand with 20 ml. EtOH and 10 ml. concd. HCl for 90 min.
at room temp., then diazotized with NaNO₂ in a HCl soln.,
the mixt. treated after 5 min. with glyamic acid, after 2
min. with a soln. contg. 0.272 g. di-HCl salt of II in 250 ml.
H₂O, filled to the mark in a 250-ml. volumetric flask, and
measured after 60 min. M. Kudický

VECERA, M.

95. Identification of organic compounds. III.
~~Alkylthiuronium salts. M. Jureček and M. Vecera.~~
~~(Coll. Czech. Chem. Comm., 1954, 19 (1), 77-84)~~

Various alkylthiuronium salts are investigated for the identification of alkyl and halogenoalkyl groups

attached to O or N. The unknown substance is heated with HI; the alkyl iodide formed is characterized by conversion to the thiuronium iodide and thence to its pierate (*cf. Chem. Listy*, 1951, 46, 149 and 150) or another salt. 3:5-Dinitrobenzoates are now recommended on account of their lower solubility in alcohol. Other salts examined include styphnates, picrolonates, toluene-*p*-sulphonates, oxalates, nitrates and perchlorates. [This is a translation into German of a paper that appeared in *Chem. Listy*, 1952, 46, 722.] D. R. GILSON

AA-25

LUJES, R.; VECERA, M.

Effect of Grignard reagents on the amide group. Part 18. Isomery of
pyrroline [in German with summary in Russian]. Sbor. Chekh.khim.rab.
19 no.2:263-274 Ap '54. (MLRA 7:6)

1. Institut obshchey eksperimental'noy organicheskoy khimii Prazhskogo
Politekhnicheskogo instituta.
(Grignard reagents) (Pyrroline) (Isomerism)

VECERA, M.; JULICEK, M.

Identification of Organic Compounds. II. Alkylthiuronium Salts. p. 77,
(COLLECTION OF CZECHOSLOVAK CHEMICAL COMMUNICATIONS. SERIEM CHERHOSLOVAT-
SKIKH KHIMICHESKIKH RABOT, Vol. 19, No. 1, Feb. 1954, Praha, Czechoslovakia)

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

VECERA, M.; LUKES, R.

"Effect of Grignard Reagents on the Amide group.XVIII. Isomery of Pyrroline." p. 263. (COLLECTION OF CZECHOSLOVAK CHEMICAL COMMUNICATIONS. SBORNIK CHEKOSLOVATSKIKH KHEMICHESKIKH RABOT. Vol. 19, No. 2, Apr. 1954; Praha, Czech.)

So: Monthly List of East European Accessions, (E.EAL), LC, Vol. 4, No. 4, April 1955, Uncl..

VEČERA, M.

CZECH

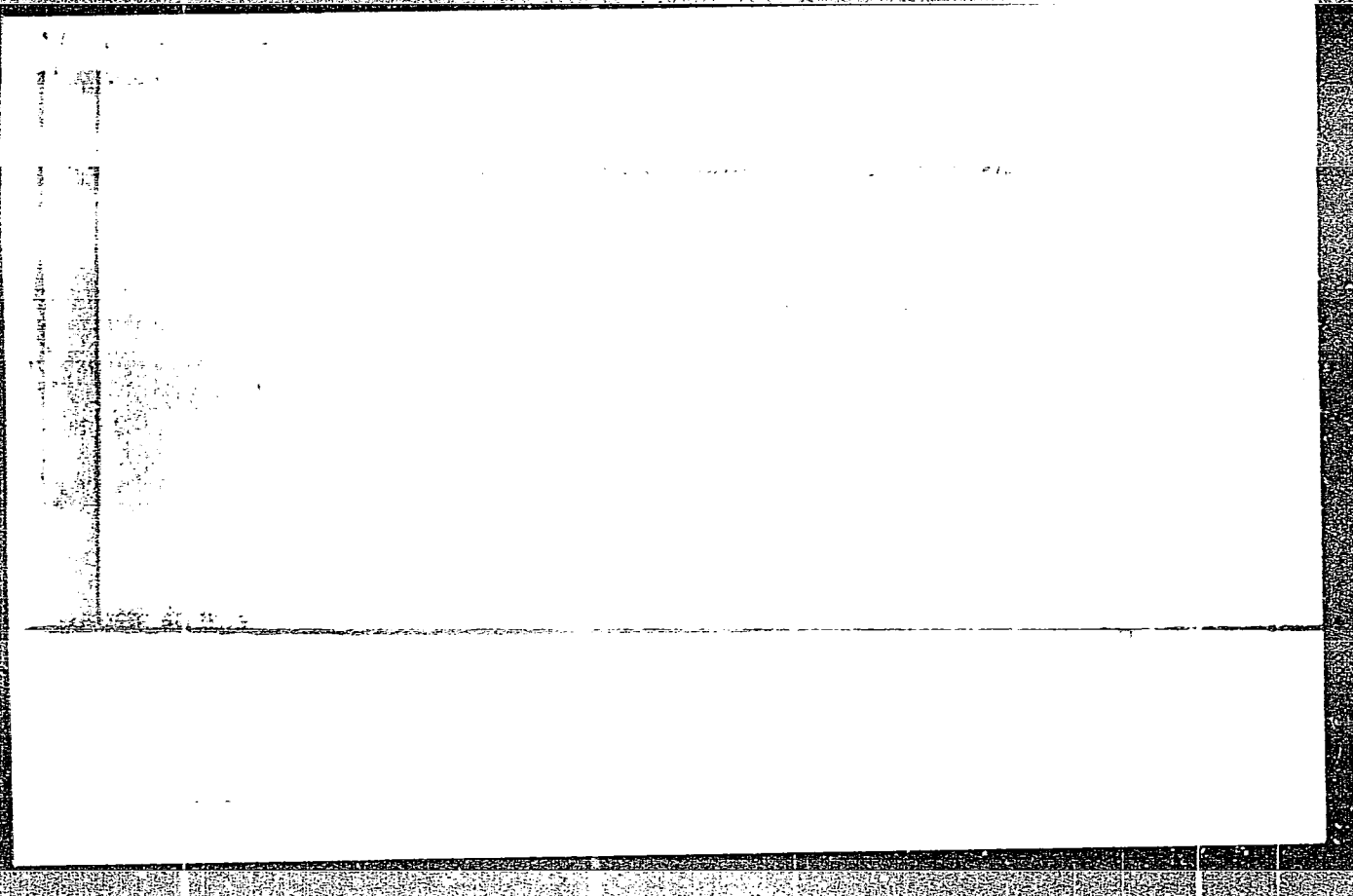
Chromatography of dye intermediates. III. Identifica-
 tion and separation of anthraquinonesulfonic acids by
 paper chromatography. Miroslav Večera, Jiří Gasparič, and
 Jiří Borecký (Vězkumný ústav org. syn., Pardubice-
 Březivka, Czech.). *Chem. Listy* 49, 700-8 (1955); cf. 49,
 619b.—Identification of isomeric anthraquinonesulfonic and
 -disulfonic acids, tests for purity of tech. products, and the
 course of the sulfonation processes are possible by means of
 chromatography on paper (Whatman no. 4) with BuOH-
 NH₄OH-H₂O 2:1:1 (I) or BuOH-C₆H₆-N-H₂O 3:1:1 (II) as
 solvents. Detection was carried out by fluorescence in
 ultraviolet light. R_f values in systems I and II at 21° are
 given for the following anthraquinonesulfonic and -disulfonic
 acids: 1 (0.65, 0.57); 2, (0.75, 0.72); 1,5 (0.04, 0.07); 1,6
 (0.09, 0.15); 1,7 (0.14, 0.19); 1,8 (0.26, 0.35); 2,6 (0.19,
 0.29), 2,7 (0.19, 0.29). M. Hudlícký.

2

BL

"APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001859220003-1



APPROVED FOR RELEASE: 08/31/2001

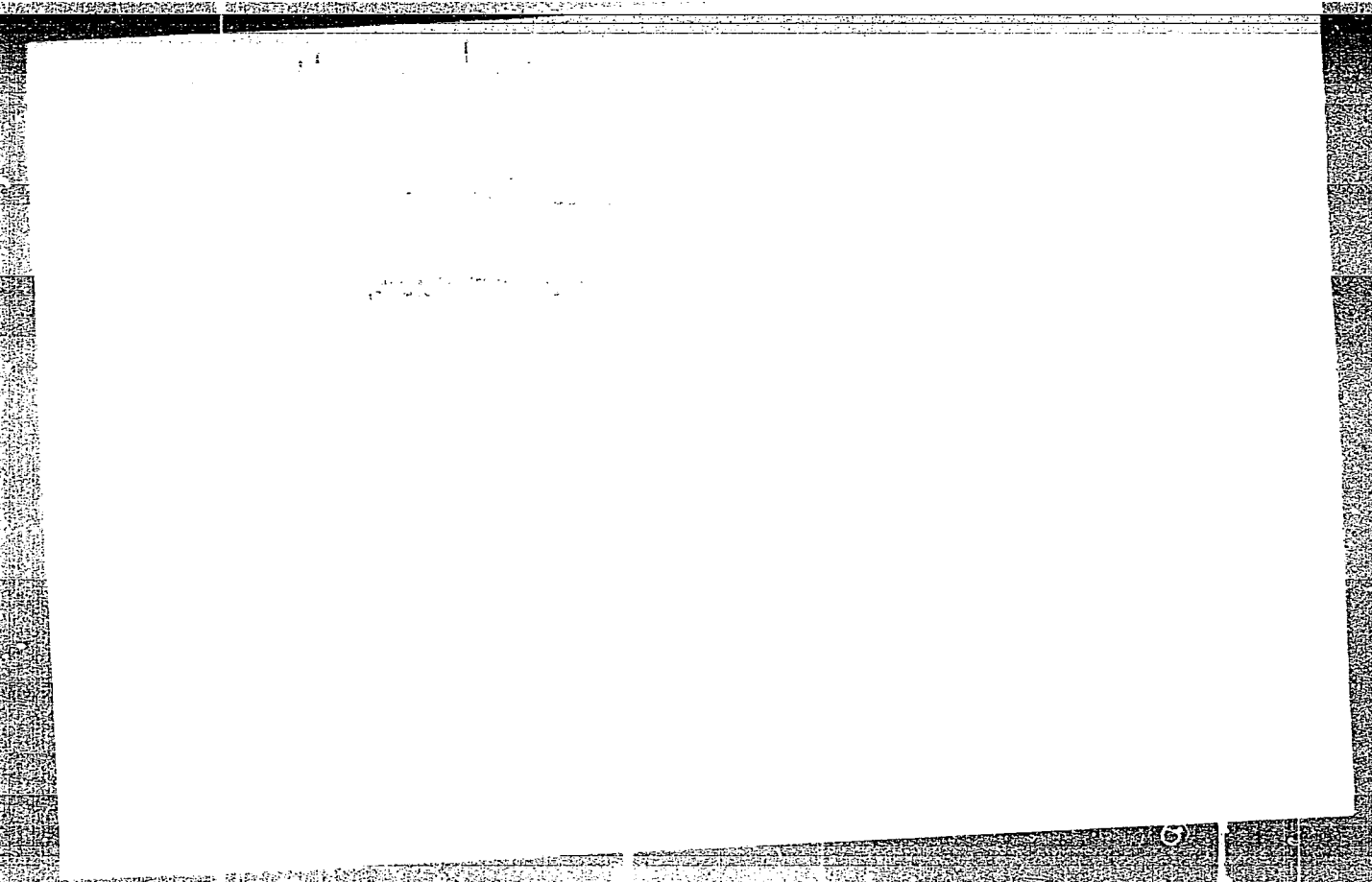
CIA-RDP86-00513R001859220003-1"

"APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001859220003-1

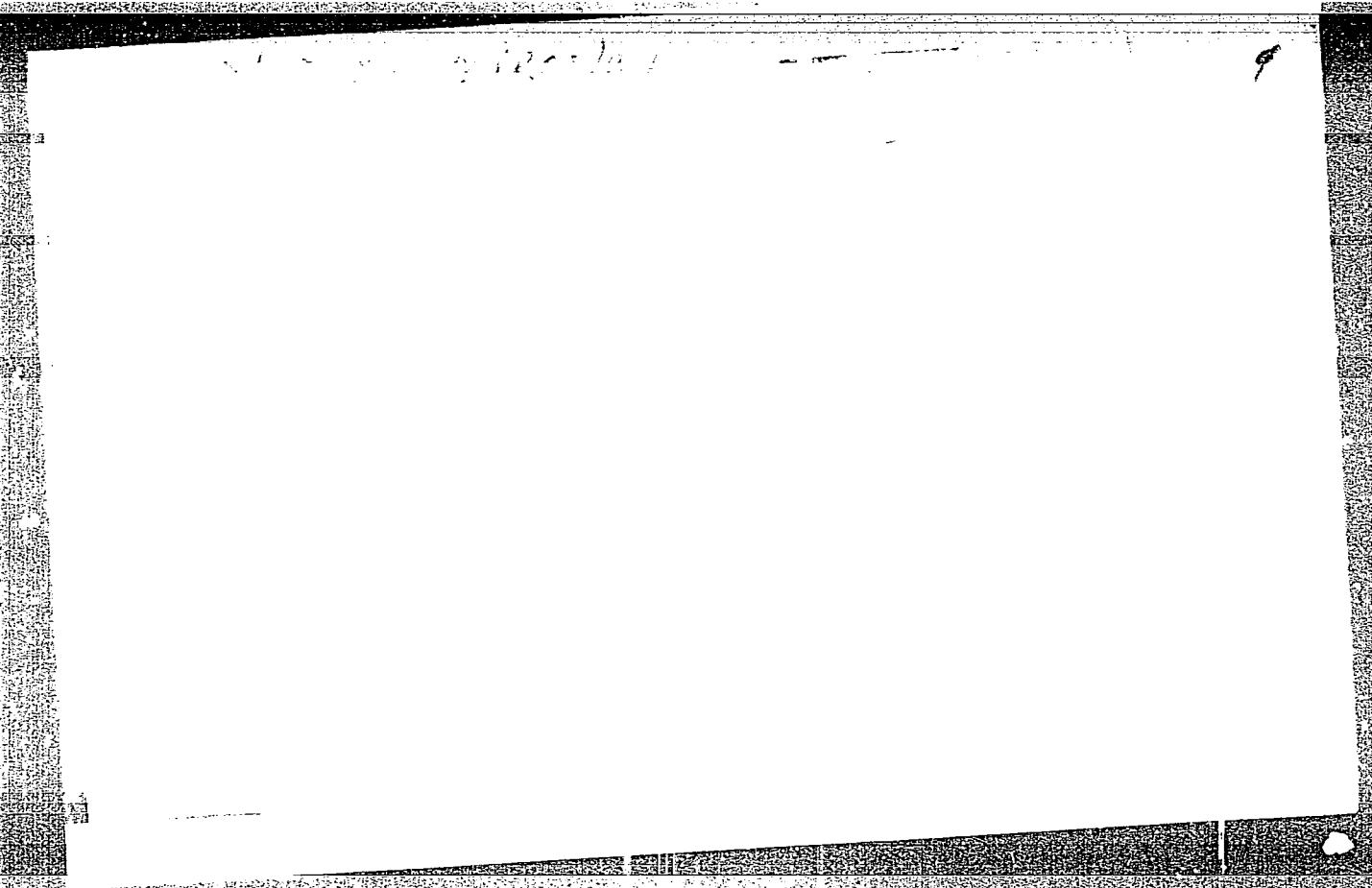
APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001859220003-1"



"APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001859220003-1



APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001859220003-1"

"APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001859220003-1

APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001859220003-1"

MIRDOCLAV VECERA

16.02.1977
Czechoslovakia/Analytical Chemistry - Analysis of Organic Substances, G-3

Abst Journal: Referat Zhur - Khimiya, No 1, 1957, 1280

Author: Vecera, M., and Petranek, J.

Institution: None

Title: Identification of Organic Substances. X. Identification of Sulfides

Original
Periodical: Chem. listy, 1956, Vol 50, No 2, 240-245 (published in Czech); Sb. chekhosl. khim. rabot, 1956, Vol 21, No 4, 912-919 (published in German with a summary in Russian)

Abstract: During the investigation of the derivatives of thioethers, the properties of the sulfolimines (I) prepared by the reaction of organic sulfides (II) with the sodium salt of N-chloro-p-toluenesulfonamide (chloramine T) (III). The effect of the solvent and reaction time on the yield was studied with the type-substances, dibutylsulfide and dibenzylsulfide (IV). The greatest yields of I (75-87%) were achieved as follows: 0.005 M solutions of II and III are mixed in 10 ml CH₃OH, the solvent is distilled off, and the residue washed with 5 ml of

Card 1/3

Czechoslovakia/Analytical Chemistry - Analysis of Organic Substances, G-3

Abst Journal: Referat Zhur - Khimiya, No 1, 1957, 1280

Abstract: 2 N NaOH and 20 ml of water. The product is dissolved in 10 ml toluene over a water bath, filtered, and cyclohexane is added until the solution becomes turbid, after which it is allowed to stand for crystallization. Dibenzylsulfoxide and p-toluenesulfonamide are formed as side products during the reaction of IV with III. Compounds of I have also been prepared from the following II (the melting point and eutectic point (EP) with standard substances such as acetanilide (V), benzil (VI), phenacetin (VII) and cyanoguanidine (VIII), and the index of refraction of molten I (L. Kofler and A. Kofler, Mikromethoden zur Kennzeichnung organischer Stoffe und Stoffgemische, Innsbruck, Universitaetsverlag wagner, 1948) are given). For the determination of the eutectic point the substances were mixed in the ratio 1:1. Di-methylsulfide, mp 158.5-159°, EP with VII 116°, n_D¹⁸²⁻¹⁸⁸ 1.5309; methyl-ethyl, mp 131.5-132.5°, EP with VII 106°, n_D¹⁵⁶⁻¹⁵⁸ 1.5309; diethyl, mp 144.5-145.5°, EP with VII 109°, n_D¹⁶⁵⁻¹⁶⁷ 1.5217; methyl-n-propyl, mp 104-105°, EP with VI 75°, EP with V 74°, n_D¹²⁷⁻¹³⁰ 1.5309; methyl-isopropyl, mp 114-115.5°, EP with V 77°, n_D¹³⁵⁻¹³⁸ 1.5309; methyl-n-butyl, mp 87-88°, EP with VI 68°, n_D¹⁰⁵⁻¹⁰⁷ 1.5309; methylisobutyl, mp 120-121°, EP with V 81°, n_D¹³³⁻¹³⁶

Card 2/3

Czechoslovakia/Analytical Chemistry - Analysis of Organic Substances, G-3

Abst Journal: Referat Zhur - Khimiya, No 1, 1957, 1280

Abstract: D 1.5217; ethyl-n-propyl, mp 104-105°, EP with VI 78°, EP with V 74°,
n110-114° D 1.5309; ethylisopropyl, mp 116.5-117.5°, EP with V 75°,
n140-141° D 1.5217; ethyl-n-butyl, mp 87-88°, EP with V 63°, EP with
VI 67°, n92-95° D 1.5309; ethylisobutyl, mp 108-109°, EP with V 70°,
n120-122° D 1.5217; di-n-propyl, mp 110-111.5°, EP with V 72°,
n120-123° D 1.5217; n-propylisopropyl, mp 107-108°, EP with V 72°,
n128-129° D 1.5217; diisopropyl, mp 117.5-118.5°, EP with V 76°,
n132.5-134.5° D 1.5217; propylisobutyl, mp 99-100°, EP with VI 69°,
EP with V 65°, n105-107.5° D 1.5217; isopropylisobutyl, mp 102-103°,
EP with VI 73°, EP with V 70°, n108.5-110° D 1.5217; isopropyl-s-butyl,
mp 96-97°, EP with VI 71.5°, EP with V 76°, n120-123° D 1.5217; di-
n-butyl, mp 64°, EP with VI 51°, n86-88° D 1.5217; n-butyl-s-butyl,
mp 75-76°, EP with VI 61°, n100-102° D 1.5217; diisobutyl, mp 133.5-
134.5°, EP with VII 105°, n164-165° D 1.4953; isobutyl-s-butyl, mp
95-96°, EP with VII 73°, n112-114° D 1.5151; pentamethylene, mp
148.5-149°, EP with VII 109°, n172-174° D 1.5427; dibenzyl, mp 192-
193°, EP with VIII 168°, n220-222° D 1.5309.

Card 3/3

VECERA, J.

VECERA, M. - Identification of organic substances. X. Identification of sulfides.
p. 240, Vol. 50, no. 2, Feb. 1956
CHEMICKÉ LISTY (Czechoslovak academic ved. Chemický ustav)
Praha, Czech.

SOURCE: East European Accessions List (EEAL) Vol 6 No 4 April 1957

Vecera, M.

CZECHOSLOVAKIA/Analytical Chemistry - Analysis of Organic
Substances

G-3

Abs Jour : Referat Zhur - Khimiya, No 2, 1957, 4830

Author : Vecera, M.
Title : Organic Quantitative Analysis. VI. Micro-Determination of
Sulfur in Organic Substances

Orig Pub : Chem. listy, 1956, 50, No 2, 308-311

Abstract : The previously described method (RZhKhim, 1955, 2305,
49236) has been improved. Use of a quartz combustion
tube (Grote W., Krekeler H., Angew. Chemie, 1933, 46,
106) with a quartz partition and two porous plates
(temperature 700-750°) in lieu of a Pt catalyst, increa-
ses substantially the speed of determination, since the
rate of O₂ feed can be raised to 10-30 ml/minute, and at
the same time combustion is complete with all types of
substances and takes place evenly. Absorption of the
oxides of sulfur is quantitative over the temperature

Card 1/3

- 52 -

CZECHOSLOVAKIA/Analytical Chemistry - Analysis of Organic
Substances

G-3

Abs Jour : Referat Zhur - Khimiya, No 2, 1957, 4830

alum as an indicator, and with Complexon III.
Ag wool also absorbs the halogens which may have a de-
trimental effect. It is recommended to replace a por-
tion of Ag wool by fresh one after 5 analyses of subs-
tances containing halogens. Average error of determi-
nation is 0.15, mean quadratic error is 0.19.

Communication V, see RZhKhim, 1956, 68776.

Card 3/3

- 54 -

VEEPRE M

V. C. C. P. A. D.

1/2

1. 1877 Organic quantitative analysis: VIII. Micro-
estimation of sulphur in organic compounds.
Mechanism of formation of silver sulphate

2.3 sq. dm. combustion time (at 20 mm) and
velocity of the oxygen stream 10 to 20 m per min

MT

"APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001859220003-1

APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001859220003-1"

Večera, Miroslav

Separation and identification of the rearrangement products of hydrazobenzene. Miroslav Večera, Jiří Gasparič, and Jaromír Petránek (Výzkumný ústav org. syntézy, Pardubice-Rybní, Czech.). *Chem. listy* 51, 1690-2 (1957); cf. *C.A.* 51, 14691d.—(PhNH)₂ (1.5 g.) mixed with 30 g. dry sea sand, exposed 72 hrs. to a stream of dry HCl, the mixt. made alk. with dry NH₃, and the bases extd. with 75 ml. C₆H₆ and chromatographed on Al₂O₃ gave 80% of the following bases identified as the Bz derivs. (m.p., n_D): benzidine, m. 308°, —; diphenylene, m. 280-2°, n_D²⁰ 1.5675, o-benzidine, m. 191-1.5°, n_D²⁰ 1.5714; o-semidine, m. 135-0°, n_D²⁰ 1.6004; and p-semidine, m. 166°, n_D²⁰ 1.6111. The mutual ratio of the bases was 1:1.25:1.08:0.84:0.09, resp. In addn., PhNH₂ and (PhN)₂ were found among the rearrangement products. M. Hudlický

5
 4E 4j
 2 Mary
 4E 3d
 4E 2d(j)

RM

~~SECRET~~
V. 10000

Distr: (E2c(j))

~~Organic quantitative analysis. XVII. Determination of small amounts of chlorine and bromine in organic compounds. Miroslav Veselý and Antonín Šolcák (Výzkumný ústav org. syntézy, Pardubice, Říše, Czech). Chem. listy 51, 2037-40 (1957), cl. C.A. 52, 4830. — Combustion of an org. compd. in O₂ is carried out in a quartz tube and the HCl or HBr obtained by absorption of the gaseous products in H₂O is detd. colorimetrically by the reaction with Hg(SCN)₂ and Fe³⁺. The method is suitable for detg. 0.01-2% Cl or Br; as the percentage of halogens increases the accuracy decreases. Ignite a 1-15-mg. sample in O₂ (15 ml./sec.) for 20-30 min.; absorb the exit gases in 2 ml. 5% H₂O₂, add 2 ml. of a solu. contg. 0.3 g. Hg(SCN)₂ in 100 ml. warm MeOH, and 4 ml. of a solu. contg. 0 g. NH₄Fe(SO₄)₂ in 100 ml. 6N HNO₃, dil. to 25 ml., and measure the absorption with a filter with max. transmittancy at 470 mμ.~~

M. Hudlický

6
2-70-10
1

jm

VECERA, M.

Identification of organic compounds. XXIV. Separation and identification of sulfides by paper chromatography. Jaromir Petránek and Miroslav Večeřa (Výzk. ústav org. synth., Pardubice-Rybitví, Czechoslov.). *Chem. listy* 52, 1270-82 (1958); cf. *C.A.* 52, 13544b. — Org. sulfides may be separated and identified in the form of the *p*-nitrobenzenesulfonyl-sepd. and identified in the form of the *p*-nitrobenzenesulfonyl-sulphimines (I) on paper impregnated with HCONH₂ using C₆H₆ or C₆H₁₁-cyclohexane as the moving phase. *p*-O₂NC₆H₄SO₂NH₂ (5 g.) is dissolved in 1 g. NaOH in 10 ml. H₂O, the soln. cooled to 10° and, before the sulfonamide Na salt begins to sep., 17 ml. cool NaOCl soln. added (contg. 1.95 g. NaOCl and prepd. by passing 7 g. Cl into 8 g. NaOH in 10 ml. H₂O and 30 g. ice). The *p*-nitrobenzenesulfochloramide Na salt (II) is filtered off and recrystd. from 5-6 ml. H₂O to yield 6 g. yellow crystals contg. 23.5-5.3% active Cl and forming in H₂O and EtOH intensely yellow colored solns. II (200 mg.) in 5 ml. MeOH is added to 0.5 millimole sulfide in 2 ml. MeOH, the mixt. allowed to stand 15 min., dild. with 20 ml. H₂O, treated with 10 ml. 2*N* NaOH, extd. with CHCl₃, the ext. (contg. 1-10 µg. I) chromatographed on Whatman No. 4 impregnated with 10% HCONH₂ in EtOH, the chromatogram sprayed with a freshly prepd. soln. of 0.7 g. SnCl₄ in 100 ml. 15% aq. HCl, and the spots detected (in a 30 min. interval) by spraying with 1% *p*-Me₂NC₆H₄CHO in EtOH contg. 5% aq. HCl. 1 form yellow spots. The following *R_f* values of the *p*-O₂NC₆H₄SO₂N:SRR' were found in C₆H₆, 3:2 C₆H₁₁:cyclohexane, and 1:1 C₆H₁₁:cyclohexane, resp., (R, R', *R_f* values given): Me, Me, 0.08, 0.02, —; Me, Et, 0.21, 0.08, 0.01; Et, Et, 0.37, 0.19, 0.05; Et, Pr, 0.57, 0.35, 0.11; Pr, Pr, 0.73, 0.55, 0.24; iso-Pr, iso-Pr, 0.72, 0.63, —; Pr, iso-Pr, 0.73, 0.54, —; Et, iso-Bu, 0.72, 0.53, —; Bu, 0.74, 0.54, —; Pr, Bu, 0.83, 0.71, 0.40; Bu, Bu, —, 0.80, 0.57; iso-Am, iso-Am, —, 0.89, 0.75; Me, PhCH₃, —, 0.18, —; Et, PhCH₃, —, 0.33, —; Pr, PhCH₃, —, 0.63, —; Bu, PhCH₃, —, 0.69, —; PhCH₃, PhCH₃, —, 0.0, —; Ph, Ph, —, 0.86, —; Et, Ph, —, 0.66, —; Pr, Ph, —, 0.74, —; Et, *p*-tolyl, —, 0.74, —; Et, *o*-tolyl, —, 0.77, —.

XXV. Identification and separation of aliphatic C₇-C₁₁ alcohols by paper chromatography. Jiří Borecký, Jiří Čáslavský, and Miroslav Večeřa. *Ibid.* 1283-8. — Aliphatic C₇-C₁₁ alcs. and some cyclanols were successfully chromatographed as 3,5-dinitrobenzoates on Whatman No. 3 impregnated with 10% paraffin oil in cyclohexane. New systems of solvents contg. HCONH₂ (I) and HCONMe₂ (II) were used as the mobile phase. For preliminary information is recommended the system 18:4:4 II-MeOH-H₂O where C₇-C₈ alcs. are in the front and C₉-C₁₁ alcs. near the start. The following systems are suitable for given alcs.: 30:70 I-H₂O, C₇-C₈; 50:50 I-H₂O, C₇-C₈; 70:30 I-H₂O, C₇-C₈; 1, C₇-C₁₁; 10:10:1 II-MeOH-H₂O, C₇-C₈; 70:30 I-H₂O, cyclanols C₇-C₈; 10:10:1 II-MeOH-H₂O, cholesterol. The 3,5-dinitrobenzoates are prepd. by dissolving 0.1 g. alc. in 1 ml. C₆H₆, adding 0.5 g. 3,5-(O₂N)₂C₆H₃COCl in 3 ml. C₆H₆, and 1 ml. pyridine, heating 30 min. on a steam bath, letting cool, extg. with 50% KOH with addn. of H₂O to achieve a quick sepn. of the layers, washing the C₆H₆ ext. with portions H₂O, 1:1 HCl, and H₂O, and drying with Na₂SO₄. The C₆H₆ soln. is used directly for chromatography. The method is also suitable for sepn. and identification of higher fatty alcs. used in the textile manuf.

Distr: 4E2c(j)

7
/Rearrangement of aromatic hydrazo compounds. Miroslav
Večeřa (Výzk. ústav org. synth., Pardubice-Rybitví,
Czech.). *Chem. listy* 52, 1373-408(1958).—An extensive
review in 19 chapters with 102 references.

LP

3
7-May

300

1/1

Distr: 4E2c(j)

CZECHOSLOVAKIA/Optics - Optical Technology

K-4

Abs Jour : Ref Zhur - Fizika, No 9, 1958, No 21405

Author : Vecera M.
Inst : ~~Not Given~~
Title : Chronological Study of Anamorphotes

Orig Pub : Jomna mech. a opt., 1957, 2, No 3, 75-78

Abstract : The author examines the properties of surface systems of anamorphote fittings and investigates in particular the dependence of the angular magnification on the angle of incidence of the ray and on the angle of the prism. A graphic method is proposed for determination of optimum angle between the prisms. The condition of parallelness is examined, and the dependence of the coefficient of anamorphotes on the angle of field of view is given.

Card : 1/1

VECERA, H.

Convention of analytic chemists in Gottwaldov.

p. 85 (Chemicky Prumysl. Vol. 7, no. 2, Feb. 1957, Praha, Czechoslovakia)

Monthly Index of East European Accessions (EEAI) LC. Vol. 7, no. 2,
February 1958

VECERA, M.; SNOBL, D.

"Organic quantitative analysis. VIII. Microdetermination of sulfur in organic compounds. Mechanism of the formation of silver sulfate. In German."

p. 986 (Collection of Czechoslovak Chemical Communications. Sbornik Chekhoslovatskikh Khimicheskikh Rabot.) Vol. 22, no. 3, June 1957.
Prague, Czechoslovakia

SO: Monthly Index of East European Accessions (EEAI) LC. Vol. 7, no. 4,
April 1958

MIROSLAV, VECERA

CZECHOSLOVAKIA/Analytical Chemistry - Analysis of Organic
Substances.

E-3

Abs Jour : Ref Zhur - Khimiya, No 8, 1958, 24904

Author : XIII. Vecera Miroslav, Friedrich Kurt
XIV. ~~Gasparic Jiri~~, Vecera Miroslav

Inst : -
Title : Identification of Organic Substances. XIII. Use of Ion-
Exchangers in Organic Analysis. XIV. Separation and
Identification of Aliphatic Aldehydes and Ketones by
Means of Paper Chromatography.

Orig Pub : Chem. listy, 1957, 51, No 2, 283-286, 291; Sb. chakhosl.
khim. robot, 1957, 22, No 5, 1421-1425, 1426-1431

Abstract : XIII. Formation of a solution of an acid on interaction
of the solution of the corresponding salt with a cathio-
nite (K_m) is used for determination of equivalence and
molecular weight of organic acids and bases. First a
difficultly soluble salt of the substance under study is

Card 1/3

CZECHOSLOVAKIA/A analytical Chemistry - Analysis of Organic
Substances.

E-3

Abs Jour : Ref Zhur - Khimiya, No 8, 1958, 24904

the form of solutions of their 2,4-dinitrophenyl hydrazones (DPH) in benzene, chloroform, alcohol or I. The chromatograms are developed with cyclohexane (II) saturated with I. After volatilization of II the chromatogram is sprayed with a 1% solution of NaOH in alcohol; DPH become apparent as brown, red or blue spots. It is possible to identify 1-5 micrograms DPH. R_f increases with the number of C-atoms in the n-chain; A and K with straight and branched chains have same R_f with equal n. The presence of double bonds and OH-groups lowers R_f . Part XII see RZhKhim. 1958, 14240.

Card 3/3

CZECHOSLOVAKIA / Analytical Chemistry. Analysis of Organic
Substances.

E-3

Abs Jour : Ref Zhur - Khim., No 10, 1958, No 32229

tic absorption curves in the visible spectrum range, which permits to detect V first of all, as well as IV by an indirect method. The absorption spectra in the ultraviolet range permit to identify I, II and III as bases and chlorohydrates in solutions. The powder x-ray pictures of I, II, III, IV and V chlorohydrates and aniline can be used for the identification of these substances. They can be identified also by the polarographic curves of their interaction with HNO_2 diazonium salts of K, II and III, with HNO_2 1-phenylbenzotriazole salt of IV and with nitrosodiazonium of V. See report XIV in RZhKhim, 1958, 24904.

Card 2/2

CZECHOSLOVAKIA/Organic Chemistry Theoretical and General
Questions on Organic Chemistry.

G-1

Abs Jour: Ref Zhur-Khim., No 13, 1958, 43212.

Author : Vecera Miroslav, Petranek Jaromir, Gasparic Jiri.
Inst :
Title : Rearrangement of Substituted Aromatic Hydrazo-
Compounds.

Orig Pub: Chem. listy, 1957, 51, No 5, 911-919; Sb chekhosl.
khim. rabot, 1957, 22, No 5, 1603-1612.

Abstract: A study of the rearrangement of hydrazo-benzene
(I), 2- and 4-methyl-hydrazo-benzene (II, III), 2,2'-and
4,4'-dimethyl-hydrazobenzene (IV, V), 4-acetamido-
hydrazo-benzene (VI), N-acetyl-hydrazobenzene (VII)
and 1,1'-hydrazo-naphthalene (VIII), by action of a
solution of HCl in alcohol, or of dry HCl in absence

Card : 1/3

Questions on Organic Chemistry.

G-1

Abs Jour: Ref Zhur-Khim., No 13, 1958, 43212.

of a solvent, at about 20°. The rearrangement products (RP) were isolated by paper chromatography, purified by crystallization and also by chromatography on silica gel impregnated with dimethyl formamide, and were identified by color reactions and fluorescence reactions. Among the RP were found benzidine (IX), diphenylene (X), o-benzidine (XI), o-semidine (XII), p-semidine (XIII), the corresponding azo-compounds (Ia-VIIIa), aniline (XIV). Listing the initial substance and isolated RP: I, IX-XIV, Ia; II, IX-XIII, IIa; III, X-XIII, IIIa; IV, IX-XIII, IVa; V, XI, XII, XIV, Va; VI, XII-XIV, VIa; VII, IX, X; VIII, IX-XIV, VIIIa. Velocity of competing reactions, and proportions of

Card : 2/3

3

V. S. ... M.

1999 Organic quantitative analysis IX. Micro-
determination of sulphur in compounds containing
sulphur and ...

the addition of ... the use of a temp. of
1200° to 1300° have been found suitable.

J. Zika

EM ...

VECEPAM,

1966. Identification of ~~the~~ aromatic aldehydes
and ketones by paper chromatography. ~~Journal of~~
~~Chromatography~~ 2: 1-10

~~Abstract~~
~~Chromatography~~
~~Aldehydes~~
~~Ketones~~
~~Identification~~
~~Paper chromatography~~
~~100 µg of 2,4-dinitrophenylhydrazine~~
~~separated and for the detection of the separated~~
~~spots, a 1% ethanolic solution of NaOH was used.~~

J. ZERA

JM
MT

VECERA, M.

Identification of organic compounds. IV.

Identification of benzidine and its isomers. M

Isotopically labeled benzidine, 1,3,3',5'-tetra-¹⁴C-benzidine, was prepared by the method of Kozlovskii and Kozlovskii (1957, 61 (2): 202).

The derivatives of benzidine, namely 2,6-diaminobenzidine (I), 2,4-diaminobenzidine (II), 2,6-diaminobenzidine-III

and 2,4-diaminobenzidine-IV were prepared by the method of Kozlovskii and Kozlovskii (1957, 61 (2): 202).

The identification of these compounds was carried out by the method of Kozlovskii and Kozlovskii (1957, 61 (2): 202).

The identification of these compounds was carried out by the method of Kozlovskii and Kozlovskii (1957, 61 (2): 202).

The identification of these compounds was carried out by the method of Kozlovskii and Kozlovskii (1957, 61 (2): 202).

The identification of these compounds was carried out by the method of Kozlovskii and Kozlovskii (1957, 61 (2): 202).

The identification of these compounds was carried out by the method of Kozlovskii and Kozlovskii (1957, 61 (2): 202).

The identification of these compounds was carried out by the method of Kozlovskii and Kozlovskii (1957, 61 (2): 202).

The identification of these compounds was carried out by the method of Kozlovskii and Kozlovskii (1957, 61 (2): 202).

The identification of these compounds was carried out by the method of Kozlovskii and Kozlovskii (1957, 61 (2): 202).

The identification of these compounds was carried out by the method of Kozlovskii and Kozlovskii (1957, 61 (2): 202).

The identification of these compounds was carried out by the method of Kozlovskii and Kozlovskii (1957, 61 (2): 202).

The identification of these compounds was carried out by the method of Kozlovskii and Kozlovskii (1957, 61 (2): 202).

The identification of these compounds was carried out by the method of Kozlovskii and Kozlovskii (1957, 61 (2): 202).

The identification of these compounds was carried out by the method of Kozlovskii and Kozlovskii (1957, 61 (2): 202).

The identification of these compounds was carried out by the method of Kozlovskii and Kozlovskii (1957, 61 (2): 202).

The identification of these compounds was carried out by the method of Kozlovskii and Kozlovskii (1957, 61 (2): 202).

The identification of these compounds was carried out by the method of Kozlovskii and Kozlovskii (1957, 61 (2): 202).

M 1957

VECERA, M. : POLSEKY, J.

"Identification of organic compounds. XVII. Identification of anthraquinonesulfonic acids."

p. 974 (Institute of Applied Physics - Czechoslovak Academy of Science)
Vol. 51, No. 5, May 1957

SJ: Monthly Index of East European Accession (EEAI) (LC, Vol. 7, No. 5, May 1958)

CZECHOSLOVAKIA/Analytical Chemistry. Analysis of Organic
Substances.

E-3

Abs Jour: Ref Zhur-Khim , No 13, 1958, 43082.

Author : X. Vecera Miroslav, Dulusek Jaroslav.
XI. Synek Ladislav, Vecera Miroslav.

Inst :

Title : Organic Quantitative Analysis. X. Micro-Determination
of Chlorine and Bromine in Organic Substances. XI.
Use of Cobalto-Cobaltic Oxide as Combustion Catalyst
in Elementary Analysis.

Orig Pub: Chem. listy, 1957, 51, No 8, 1475-1481, 1551-1552;
Collet. czechosl. chem. communs, 1958, 23, No 2, 257-264,
331-333.

Abstract: X. 1-5 mg (determination of Cl) or 4-7 mg (determina-
tion of Br) of the substance are used in the combustion

Card : 1/5

CZECHOSLOVAKIA/Analytical Chemistry. Analysis of Organic Substances.

E-3

Abs Jour: Ref Zhur-Khim., No 13, 1958, 43082.

by the modified method of Grote and Krekeler, in a quartz tube 580 mm long and 9 mm inside diameter containing two porous quartz plates, in a current of O_2 at 750° . If the substance contains alkali metal a tenfold amount of V_2O_5 is added. Liquid substances are weighed in capillaries with NH_4NO_3 . Combustion of the substance is effected within 5-15 minutes after which the tube heated for an additional 5 minutes. The halogen is absorbed as HCl (HBr) in a U-vessel filled with glass beads and containing 5 ml of 5% H_2O_2 . For determination of Cl^- the solution is neutralized first with 0.1 N and then with 0.01 N solution of NaOH, to a mixed indicator (0.1% alcohol solutions of methyl

Card : 2/5

CZECHOSLOVAKIA/Analytical Chemistry. Analysis of Organic Substances.

E-3

Abs Jour: Ref Zhur-Khim., No 13, 1958, 43082.

red and methylene blue 3:1) which has a grey transition shade, 10 ml of freshly neutralized saturated solution of $\text{HgO} \cdot \text{Hg}(\text{CN})_2$ are added, and after 1 minute the titration is carried out with 0.01 N H_2SO_4 to the initial gray shade (Viebock F., Der., 1932, 65, 496). Br^- is determined by titration of exactly neutralized solution with 0.01 N solution of AgNO_3 in the presence of an adsorption indicator (0.1% aqueous solution of Brilliant Yellow) until the yellow-green color changes to orange (RZhKhim, 1955, 46100). A detailed description is given of the selection of optimal conditions of combustion as well as a comparison of the proposed procedure with the

Card : 3/5

CZECHOSLOVAKIA/Analytical Chemistry. Analysis of Organic Substances.

E-3

Abs Jour: Ref Zhur-Khim., No 13, 1958, 43082.

undergo quantitative combustion at about 600° . Catalytic activity of Co_3O_4 is not decreased by the action of halogens or oxides of sulfur, and therefore the same catalyst filling can be used for several months. Preparation of catalyst: to 15 g asbestos and 10 g $Co(NO_3)_2 \cdot 6H_2O$ are added 50 ml water and 5 drops of concentrated NH_4OH , after which the mixture is evaporated to dryness and calcined at about 500° . Analyses are carried out in the usual manner. Combustion of 11-15 mg requires only 5-15 minutes. Halogens and oxides of sulfur are absorbed on Ag-wool at 600° , oxides of nitrogen -- on MnO_2 . Communication IX see RZhKhim, 1958, 4320.

Card : 5/5

CZECHOSLOVAKIA/Analytic Chemistry - Analysis of Organic
Substances.

E-3

Abstr Jour : Ref Zhur - Khimiya, No 14, 1958, 46438

adjust the results of H and N determinations.
The method of linear regression is the most suitable
statistical method for the evaluation of the accuracy;
by that method it is possible not only to describe the
accuracy, but also to make apparent the constant and
regularly variable errors.
See report XI in RZhKhim, 1958, 43082.

Card 2/2

"APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001859220003-1

APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001859220003-1"

VECERA, M. ; PETRANEK, J. ; GASPARIC, J.

"Identification of organic compounds. XVIII. Chromatography of aromatic hydrazo compounds."

p. 1553 (Chemicke Listy, Vol. 51, no. 8, Aug. 1957, Praha, Czechoslovakia.)

Monthly Index of East European Accessions (EEAI) LC, Vol. 7, No.6 June 1958

VECERA, M. ; GASPARIC, J. ; SPEVAK, A.

"Identification of organic compounds. XIX. Microidentification of the lower aliphatic alcohols, O-alkyl and N-alkyl groups by paper chromatography."

p. 1554 (Chemicke Listy, Vol. 51, no. 8, Aug. 1957, Praha, Czechoslovakia.)

Monthly Index of East European Accessions (EMAI) LC, Vol. 7, No.6 June 1958.

VECERA, M).

CZECHOSLOVAKIA / Analytical Chemistry. Analysis of Organic Substances. E

Abs Jour: Ref Zhur-Khimiya, 1958, No 20, 67346.

Author: Petranek J., Vecera M.

Inst: Not given.

Title: Organic Quantitative Analysis. XVI. Calorimetric Determination of Small Quantities of Aldehydes.

Orig Pub: Chem. listy, 1957, 51, No 9, 1686-1689.

Abstract: Aldehydes (A) when condensed with resorcinol (I) or with fluoroglycine (II) in concentrated H₂SO₄ or HCl give very weak and poorly reproducible colorings. In glacial CH₃COOH (III), that contains 10-20 vol. % H₂SO₄, A reacts slowly with I and sufficiently fast with II, even at approx. 20°, resulting in the intensive and stable yellow and orange colorings. Since III should not con-

Card 1/3

30

CZECHOSLOVAKIA / Analytical Chemistry. Analysis of Organic Substances. E

Abs Jour: Ref Zhur-Khimiya, 1958, No 20, 67346.

Abstract: tain A, it is being purified by fractionation with 5% II and 5% concentrated H_2SO_4 . to 5cc of very dilute solution of A in III, 5cc of III, containing 20 vol. % H_2SO_4 is added, the mixture is then cooled to $18-20^{\circ}$ followed by the addition of 1cc of 1% solution of II in III. After 10-30 minutes, the photometric readings are taken using either dark blue or blue-green filters respectively. The Bere's law is being observed for the cases when 5cc solutions contain 26.6 - 265.8 γ of n-toluene A, 4.2 - 29.9 γ of vinylene, or 4.1 - 30.6 γ of piperonal. The greatest intensity of color de-

Card 2/3

CZECHOSLOVAKIA / Analytical Chemistry. Analysis of Organic Substances. E

Abs Jour: Ref Zhur-Khimiya, 1958, No 20, 67346.

Abstract: velops after 10-30 minutes and the color becomes stable after 25-90 minutes. With aliphatic A, II forms a color unstable products. The standard error is $\pm 2.75\%$. Alcohols, carbonic acids and their esters do not interfere, however, in the presence of water the color intensities diminish. This method is applicable in particular for the determination of aromatic A. Some of the latter could be identified by means of spectrophotometrical curves obtained for their respective colored products. For Part XV, refer to Ref. Zhur-Khimiya, 1958, 57232.

Card 3/3

31

VECERA, MIROSLAV

CZECHOSLOVAKIA/Analytic Chemistry. Analysis of Organic
Substances.

E

Abs Jour: Ref Zhur-Khimiya, No 22, 1958, 73760.

Author : Miroslav Vecera, Antonin Spevak.

Inst :

Title : Organic Quantitative Analysis. XVII. Determination
of Little Amounts of Chlorine and Bromine in
Organic Substances.

Orig Pub: Chem. lity, 1957, 51, No 11, 2037-2040.

Abstract: A rapid, simple and universal method of determination
of little amounts (0.01 to 2%) of Cl and Br in
solid and liquid organic substances is proposed.
1 to 15 mg of the substance is burned in a 0.2 flow
(15 ml per min.) in a Pt combustion boat or glass

Card : 1/3

CZECHOSLOVAKIA/Analytic Chemistry. Analysis of Organic
Substances.

E

Abs Jour: Ref Zhur-Khimiya, No 22, 1958, 73760.

capillary placed in a Si tube with diaphragms and heated by an electric stove to 800°. The producing hydrogen halide is absorbed from the combustion gases by 2 ml of 5%-ual H₂O₂ solution. 2 ml of 0.3%-ual Hg (SCN)₂ solution in CH₃OH and 4 ml of 6%-ual NH₄Fe(SO₄)₂ solution in 6 n. HNO₃ are added to the absorbing solution, which is photometered at 470 mμ. The duration of combustion is 10 to 30 min. Liquid substances with small contents of halogens can be burned in calorimetric bombs. The reducing mineralization with metallic Na and combustion in O₂ atmosphere in a flask yields incorrect results. The calibration curve for the photometric determination of

Card : 2/3

VECERA, M.

CZECHOSLOVAKIA/Analytical Chemistry - Analysis of Organic Substances.

E-3

Abs Jour : Ref Zhur - Khimiya, No 2, 1959, 4378

Author : Vecera, M., Synek, L.

Inst : -

Title : Qualitative Organic Analysis. XVIII. Microdetermination of Carbon and Hydrogen Using Cobaltous-Cobaltic Oxide as a Combustion Catalyst.

Orig Pub : Chem Listy, 51, No 12, 2266-2274 (1957) (in Czech)

Abstract : A simple and reliable micromethod has been developed for the determination of C and H in organic substances. The combustion is carried out in a quartz tube of 300 mm length and 9 mm I. D., using CO_3O_4 on asbestos (thickness of catalyst bed 2.5 mm) as catalyst (RZhKhim, 1958, 43082). A 3-5 mg sample is ignited in a Pt-dish in a stream of O_2 for 3-5 min; for substances having high vapor pressures, 5-10 min are required. The high

Card 1/2

Distr: 4E2c

Quantitative organic analysis. XIII. A rapid carbon-hydrogen microdetermination. M. Večeta, D. Šnobl, and L. Synck (Mikroanaly. Lab. VÚBPA, Pardubice-Rybitvi, Czech.) *Mikrochim. Acta* 1973, 9-27; cf. *C.A.* 51, 16207b; 25, 19712b.—This micromethod for the detn. of C and H

27
11
2 may
1

uses Co_2O_3 as a combustion catalyst. The app. for the combustion, that for removing the other products, and the absorption chambers for the H and C is described. The accuracy is equal to that of other methods. XIV. Microdetermination of sulfur. M. Večeta and D. Šnobl. *Ibid.* 28-40.—This is a rapid and accurate micromethod for detg. S. The substance is first burned in a stream of O at 700°, after which the S oxides are trapped in Ag wool. The trap is heated to 450°, the Ag_2SO_4 is extd. with H_2O and the Ag is detd. by potentiometric or visual means. The mechanism of the reaction of the S oxides with Ag is explained. This method is compared statistically with the Zimmermann method (*C.A.* 47, 2638e). Also a rapid colorimetric method for the detn. of 0.005 to 8% S in org. and inorg. material is described. XV. Microdetermination of chlorine and bromine in organic substances. M. Večeta and J. Bulušek. *Ibid.* 41-51.—This is a rapid method for detg. Cl and Br in org. materials. The sample is burned in a quartz tube and the halogen ions are then absorbed in H_2O_2 . The Cl is then detd. by the Vieböck method (*C.A.* 26, 3206) and the Br argentometrically with the aid of an adsorption indicator. The findings were compared with those of Kainz and Resch (*C.A.* 46, 4950f) and Schöniger (*C.A.* 50, 14433d). A method is described for detg. small amts. of Cl and Br in org. materials. H. W. Harvey

5B
1

gjf

VLEBAN, M.

7

/ Identification of organic compounds. XXI. 2. choice of solvent systems for paper-chromatographic separation of organic compounds. J. Gasparic and M. Vleba (Forschungsinst. organische syntheses, Pardubice-Rybitvi, Czech.). *Mikrochim. Acta* 1958, 68-91; cf. C.A. 52, 13544b.—Practical examples are given to show how org. compds. can be sepd. by means of paper chromatography. One is not limited to "tested" solvent systems, but can use new suitable systems as the occasion demands. It has been found best to abide by the rules of soly. of org. compds., provided the compd. to be chromatographed is quite sol. in the stationary phase but less sol. in the mobile phase. By altering the stationary phase (water, nonaq., polar solvent, nonpolar solvent) or the polarity and compn. of the mobile phase, the migration of the stains in the chromatogram can be influenced, selected R_f values can be obtained, and in many cases it is also possible to secure a desired succession of the compds. on the chromatogram. Since the soly. of org. compds. depends on intermol. forces, the problem in connection with structural influences appears very complicated and must be solved individually for each case. Moreover, the soly. characteristics can be affected by using reactive solvents; for instance the compds. can be converted into water-sol. salts. Complications may arise because of the dissocn. and hydrolysis of ionizable compds. The following are among the chief factors, which may make a sepn. possible: functional groups, their number, polarity, relative position, their basicity or acidity, C-atom no. in homologous compds., inter- and intramol. H bonds, steric factors, etc. It then depends on the type of solvent system selected, which of these factors are predominant and which can be neglected or eliminated. If the soly. differences are too slight to permit good sepn., the compds. to be sepd. should, if possible, be converted into derivs. whose structural differences are more pronounced. H. W. Harvey

4
2 May

SB
1/1

ja

6
2. may

Organic quantitative analysis. XIX. Microdetermination of alkoxy groups. Miroslav Večera and Antonín Spěvák (Výzkumný ústav org. syntez, Pardubice-Rybitví, Czech.). *Chem. listy* 52, 1620-6 (1958); cf. *C.A.* 52, 4303g. —The microdetn. of O-bound alkyl groups C₁-C₄ was modified in that the interfering compds. were adsorbed on dry tartar emetic on silica in a modified app. according to Shaw (*C.A.* 41, 7810i) instead of the usual absorption in liquids. The detn. is as precise as the methods of microelementary analysis. XX. Microassay of chlorine in organic compounds by combustion in an empty tube. Miroslav Večera and Jaroslav Buláček. *Ibid.* 1626-36. —In the described method the compd. is heated at 720° in a stream of O in an empty tube (*C.A.* 51, 10207b), Cl is absorbed in dil. H₂O₂, and Cl⁻ titrated with 0.01N AgNO₃ with brilliant yellow as adsorption indicator. The possible sources of errors were investigated, and statistical evaluation of 70 results are presented. I. J. Urhánek

RG JAG

Distr: 4E2c(j)

J Identification of organic compounds. XXVI. X-ray diffraction patterns of *S,S*-dialkyl-*N*-*p*-tolylsulfonylsulfonimines. Dobroslav Šnobl, Věra Kadanková, Jaromír Petránek, and Miroslav Večeřa (Výzkumný ústav org. synt., Pardubice, Czech.). *Chem. listy* 53, 1537-45 (1958); cf. *C.A.* 50, 16859i; 52, 12644b. — X-ray diffraction patterns are suitable for the identification of thioethers after conversion to the title compds., as shown by differentiation of 23 studied derivs. This method is safer than by means of m.ps., eutectical temps., and molten mass ns and the derivs. do not require a high degree of purity. Values of the interplanar spacings *d*, as detd. by the usual technique, in org. compds. having a big elementary cell and low symmetry can be distorted owing to superposition, thus showing considerable deviations from tabulated values. The sources of errors and working technique are discussed. Lower dispersion is recommended. L. J. Urbánek

JW
11

6
2 may
1

Jaf

5
2 May

Distr: 4E3d/4E2c(j)

Identification of organic compounds. XXVII. Use of Friedel-Crafts acylation for the identification of alkyl aryl sulfides. Jiří Gasparík, Miroslav Večera, and Miroslav Jureček (Výzkumný ústav org. syntesy, Pardubice-Rybitví, Czech.) *Chem. listy* 52, 1720-5 (1958); *cf. C.A.* 53, 838c.— Alkyl aryl sulfides are first acylated with AcCl and AlCl₃ and the resulting derivs. of PhAc are transformed to 2,4-dinitrophenylhydrazones, oximes, or are oxidized to sulfones. Me sulfides were prepd. by methylation of thiophenols with Me₂SO₄, and the higher alkyl aryl sulfides are made as follows: 7 g. Na was dissolved in 150 ml. EtOH, 34.5 ml. *o*-MeC₆H₄SH added with stirring and then 40 g. Me₂CHBr, the mixt. refluxed 2 hrs. on the steam bath, the solvent distd., the residue dissolved, washed with 5% aq. NaOH, dried with Na₂SO₄, and distd. to yield 39 g. product. To prep. the ketones, to 0.8 g. AlCl₃ in 5 ml. CHCl₃ was added dropwise 0.5 g. AcCl and, with cooling, 0.5 ml. of the sulfide in 5 ml. CHCl₃, the mixt. allowed to stand 1 hr. at room temp., poured over ice and 5 ml. HCl, the org. layer washed with 5% HCl and 5% NaHCO₃, filtered, the solvent evapd. and the residue treated with the appropriate reagent [2,4-(O₂N)₂C₆H₃NHNH₂ in H₂SO₄ and EtOH, or NH₂OH in NaOH or C₆H₅N (8-10 hrs.)], or oxidized 1 hr. with 5 ml. 30% H₂O₂ in 5 ml. AcOH on the steam bath and dilbd. with 50 ml. H₂O. Alkyl aryl sulfides (b.p./umm., m.p. of the 2,4-dinitrophenylhydrazone (EtOH, AcOH or AcOH), of the oximes (cyclohexane), and of sulfones (H₂O), given): Me, Ph, 53.5°/1.5, 236.5-7.5°, 121°, 128-9°; Et, Ph, 50.3°/2.9, 193°, 90°, 115.5-16.5°; Pr, Ph, 69.1°/2.3, 158.5-60.5°, 75-5.5°, 86.5-7.5°; *iso*-Pr, Ph, 61°/3.1°, 173-4°, 75.5°, 58.5-9.5°; Me, *o*-MeC₆H₄, (I), 55.6°/1.5, 209.5°, 80°; —; Et, I, 61.8°/1.0, 159°, 91°; —; Pr, I, 68.2°/1.1, 171-2°, 91°; —; *iso*-Pr, I, 61.4°/1.2, 162°, —; —; Me, *p*-MeC₆H₄, (II), 61.5°/2.2, 213-15°, 108°; —; Et, II, 62.1°/2, 191-3°, 85°; —; Pr, II, 55.6°/0.8, 191-2°, 98°; —; *iso*-Pr, II, 65.2°/1.4, 159-60°, 91.5°, —. Powder x-ray diagrams were also used for the characterization.

M. Hudlická

LE
11

29

MIROSLAV VECERA
CZECHOSLOVAKIA / Analytical Chemistry. Analysis of
Organic Substances.

E-3

Abs Jour : Ref Zhur - Khim., No 10, 1958, No 32220

Author : Jiri Gasparic, Miroslav Vecera, Miroslav Jurecek

Inst : -

Title : Identification of Organic Substances. XVI. Identifica-
tion of Sulfides.

Orig Pub : Chem. listy, 1957, 51, No 4, 660-666; Collect. czechosl.
chem. communs, 1958, 23, No 1, 97-104.

Abstract : Continuing the similar study of dialkylsulfides (DAS)
containing the alkyls C₁-C₄ with straight chains, bromides,
picrates and perchlorates of dialkyl-n-bromophenacylsulfo-
nia (I) from 17 DAS-s were obtained; they contain one or
two C₃-C₄ alkyls with forked chains (with the exception
of tertiary butyl); their melting points, by which it is
recommended to identify the corresponding DAS-s, were de-

Card 1/3

23

CZECHOSLOVAKIA / Analytical Chemistry. Analysis of
Organic Substances.

E-3

Abs Jour : Ref Zhur - Khim., No 10, 1958, No 32220

terminated (melting points of perchlorates - from 69 to 170°, those of picrates - from 80 to 150°). I is produced by the interaction of the DAS with n-bromophenacylbromide, which takes from 20 min. to 8 hours time; a DAS with forked chains reacts more slowly than a DAS with straight chains. The picrates and perchlorates of higher DAS homologues are produced from bromides without separating the latter from the reaction mixture. It is recommended for a more complete identification of the DAS-s to use the x-ray pictures of powdered I perchlorates and the eutectic temperatures of their mixtures with some standard preparations. The melting points of bromides, perchlorates and picrates of I, as well as x-ray picture characteristics and eutectic temperatures of I perchlorates obtained from the above mentioned DAS-s are presented. See report XV in RZhKhim, 1958, 32229.

Card 2/2

VECERA, MIROSLAV

CZECHOSLOVAKIA/Analytic Chemistry - Analysis of Organic Substances.

E-3

Abs Jour : Ref Zhur - Khimiya, No 14, 1958, 46452

Author : XVII - Miroslav Vecera, Jiri Borecky. XVIII - Miroslav Vecera, Jaromir Petranek, Jiri Gasparic. XIX - Miroslav Vecera, Jiri Gasparic, Antonin Spevak.

Inst : -

Title : Identification of Organic Substances. XVII. Identification of Anthraquinone Sulfoacids. XVIII. Chromatography of Aromatic Hydrazo Compounds. XIX. Microidentification of Lower Aliphatic Alcohols and O-Alkyl and N-Alkyl Groups by Paper Chromatography.

Orig Pub : Chem. listy, 1957, 51, No 5, 974-976; No 8, 1553-1554; 1554-1556; reports VII, VIII, Collect. czechosl. chem. commun., 1958, 23, No 1, 130-133; No 2, 333-335.

Abstract : XVII. The benzylthiuronic (I) and 1-naphthylmethylthiuronic (II) salts of mono- and disulfo acids of

Card 1/6

25

CZECHOSLOVAKIA/Analytic Chemistry - Analysis of Organic
Substances.

E-3

Abs Jour : Ref Zhur - Khimiya, No 14, 1958,46452

anthraquinone are suitable for the identification of the latter. The ultraviolet spectra of these salts are close, but their melting points and eutectic temperatures in mixtures with dicyanamide are different. The salts are precipitated by mixing aqueous solutions of reagents and in the majority of cases, they are recrystallized from 30 to 80%-ual C_2H_5OH , and some from $CH_3OH + (CH_3)_2CO$. Melting points measured under microscope or in a capillary and the eutectic temperatures of mixtures with dicyandiamide (about 1 : 1) of I and II of all isomer anthraquinone sulfoacids are presented. I is more suitable for the identification than II, because it melts more distinctly.

XVIII. The aromatic hydrazo compounds (III) are separated chromatographically and identified on Watman paper No 4 treated with 25%-ual dimethylformamide solution in

Card 2/6

CZECHOSLOVAKIA/Analytic Chemistry - Analysis of Organic Substances.

E-3

Abs Jour : Ref Zhur - Khimiya, No 14, 1958, 46452

in alcohol (IV) or 10% - ual solution of formamide in alcohol (V) and dried. From 0.5 to 200 γ of the sample in 0.1 to 1% - ual alcohol or ether solution is put on the paper. It is developed by the descending method at $21 \pm 1^{\circ}$ with cyclohexane or benzene. The dried chromatogram is sprayed with 1% - ual solution of n-dimethylaminobenzaldehyde (VI) in 95 parts of alcohol and 5 parts of concentrated HCl. At this occasions the III-s regroup into corresponding diamines, which together with VI yield products of characteristic color and fluorescence presented in the report. The values of R_f of III-s are also presented. This method permits to chromatograph several hundreds of γ of the substance and to identify 0.5 γ of a III.; it can be used for the control of the benzidine regroupation at industrial scale. For that purpose, 10 ml of the reaction solution is

Card 3/4

26

CZECHOSLOVAKIA/Analytic Chemistry - Analysis of Organic Substances.

E-3

Abs Jour : Ref Zhur - Khimiya, No 14, 1958, 46452

alkalized with 5 ml of 50% aq KOH solution and shaken with 5 ml of C_6H_6 . From 10 to 30 ml of the extract is put on the paper soaked in V, it is developed in a test tube with cyclohexane by the ascending method and treated with VI solution. The chromatographic method is recommended also for the purification of III: 60 g of silica gel (VII) is saturated with 24 ml of IV, suspended in petroleum ether (VIII) and a column 25 mm in diameter is prepared. 2 g of the substance is dissolved in 1 ml of IV, VII is added and the mixture is transferred on the column, which is developed first with 200 ml of VIII and with the mixture VIII - C_6H_6 (4 : 1) after that. XIX. The lower alkyl groups ($C_1 - C_3$) are identified in the shape of corresponding alkyl-3,5-dinitrobenzoates (IX). In order to convert alcohols (X) into IX-s, 0.1 ml of pyridine (XI) and 1 ml of C_6H_6 are added to 10 ml

Card 4/6

CZECHOSLOVAMIA/Analytic Chemistry - Analysis of Organic
Substances.

E-3

Abs Jour : Ref Zhur - Khimiya, No 14, 1958, 46452

of aqueous solution of 5 to 50 mg of X, after which 11 g of K_2CO_3 and the solution of 0.5 g of 3,5-dinitrobenzoyl chloride (XII) in 2 ml of C_6H_6 are added to it at cooling. After having shaken it 3 minutes, IX is extracted with ether, the extract is washed with 1%-ual H_2SO_4 and water, and ether is distilled off. In the case of water-free X, 5 to 50 mg of X are dissolved in 5 ml of C_6H_6 , 50 mg of XII and 0.3 ml of XI are added, all is boiled 1 hour, the benzene solution is cooled, washed with 20%-ual NaOH solution, with water, with 5%-ual H_2SO_4 , and again with water, and benzene is distilled off. In order to split the alkyl groups off the alkoxy compounds and alkylamines and to obtain IX-s, 1 to 2 (2 to 5 correspondingly) mg of the substance is boiled 1 hour with HI solution in N_2 flow; the alkyl iodides are absorbed while forming by the suspension of 3 to 4 mg of Ag-3,5-dinitrobenzoate

Card 5/6

37

Card 6/6

VEČERA
CZECHOSLOVAKIA / Analytic Chemistry. Analysis of In-organic Substances. E

Abs Jour: Ref Zhur-Khimiya, No 18, 1958, 60643.

Author : M. Vecera, L. Synek.

Inst :

Title : Quantitative Organic Analysis. IX. Microdetermination of Sulfur and Substances Containing Barium, Lead, Silver and Magnium.

Orig Pub: Collect. czechosl. chem. communs, 1958, 23, No 1, 133-135.

Abstract: See RZhKhim, 1958, 4320.

Card 1/1

81