

TOROPOV, N.; SKVORTSOV, A.

All-out participation in fire prevention. Pozh.delo
6 no.8:11 Ag '60. (MIRA 13:8)

1."Zamestitel' nachal'nika pozharnoy okhrany kombinata
"Krasnyy Perekop (for Toropov). 2. Starshiy inspektor
Upravleniya pozharnoy okhrany, Yaroslavl' (for Skvortsov).
(Yaroslavl--Factories--Fires and fire prevention)

ACCESSION NR: AR4035562

S/0271/64/000/003/B010/B010

SOURCE: Ref. zh. Avtomat., telemekh. i vy*chisl. tekhn. Av. t., Abs. 3B48

AUTHOR: Butakov, Ye. A.; Toropov, N. R.

TITLE: Synthesis of the schemes realizing combinatorial operators

CITED SOURCE: Tr. Sibirsk. fiz.-tekhn. in-ta, vy*p. 42, 1963, 44-55

TOPIC TAGS: Gray code counter, shift register, Gray code counter synthesizing, combinatorial operator

TRANSLATION: Based on the Boolean algebra, a synthesis of a Gray-code counter with a digit blocking and a synthesis of a generator of combinations of n things, k at a time, are presented. These devices can be used in a control system of a specialized computer for investigating communication and control systems. Two methods are analyzed in synthesizing the Gray-code counter. In the first method, the conventional binary positional counter is used for obtaining Gray-code numbers (by a definite algorithm). Owing to the drawbacks of the first method (lower counting rate and need for additional equipment when symmetrical outputs in each counter digit are realized), the second method is used in the synthesizing in

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ACCESSION NR: AR4035562

which the Gray-code natural numbers appear directly in the counter. A Gray-code counter synthesis with blocking any k digits is presented. A functional scheme of odd and even cells is given. An algorithm underlying the generator of combinations of n things, k at a time, is described. This algorithm is also illustrated by a table presenting all combinations of 6, three at a time. An optimum scheme for realizing the algorithm consisting of k shift registers is presented. A scheme of 2-cycle magnetic-core shift register is analyzed. Six illustrations, two tables. Bibliography: 5 titles.

DATE ACQ: 14Apr64

SUB CODE: DP

ENCL: 00

Card 2/2

KASHIROV, V. I.; BUTAKOV, Ye. A.; POTTOSIN, Yu. V.; TOROPOV, N. R.; TSVETNITSKAYA, S. A.

"Problems in Realizing the L-Machine."

r report presented at the Symp on Relay Systems Theory & Finite Automata, Moscow,
24 Sep-2 Oct 62.

TOROPOV, P.I.

Complete filling for the preservation of industrial buildings and structures in the Donets Basin. Ugol' 38 no.12:
13-14 '63. (MIRA 17:5)

1. Nachal'nik tekhnicheskogo otdela Gosudarstvennogo komiteta pri Sovete Ministrov UkrSSR po nadzoru za bezopasnym vedeniyem rabot v promyshlennosti i gornomu nadzoru.

TOROPOV, P.I., inzh.; NOVIKOV, N.V.

Boring and blasting in mines under construction. Bezop. truda v prom.
2 no.9:10-11 S '58. (MIRA 11:9)

1.Kombinat L'uganskshakhtostroy (for Toropov). 2.Trest Kadiyevshakhto-
stroy (for Novikov).
(Mining engineering)

LOCKEFLEY, F. V.

41-100

REVIEWS:

Yazilizov, Yu. A., Zvezdin, Yu. S., Litvin, Yu. I.,
Bogoliubov, I., Troppov, F., Tsvetkov, N.5/06/66/CU03/03/02/033
SAC/SCATITLE: Measurement of Spectra and the Average Reaction Number in the Fusion of U^{235} and U^{238} by 14.1-Mev NeutronsPERIODICAL: Journal of Experimental and Theoretical Physics, 1960,
Vol. 30, No. 3, pp. 671-684

TERM: The present article deals in detail with the experimental investigation made in the energy range 0.4 - 5 Mev by means of the time-of-flight technique and a pulsed neutron source. The experimental arrangement is schematically shown in Fig. 1. The reaction $\text{T}(\text{d},\text{n})\text{He}$ served as primary neutron source in the target of an accelerator. The target was bombarded with 150-kev deuterons. The time-of-flight determination was carried out electronically by measuring the time interval between the pulses in the detector. The primary deuterons were obtained by modulation, i.e., by means of a sinusoidal

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electric field ($f = 200/\text{sec}$). The pulses of the 14.1-Mev neutrons lasted 20-30 nanoseconds and had a frequency of 4 K/sec. On the average, 4 neutrons were obtained per pulse. Two fusion chambers were used (with U^{235} (90 per cent) and U^{238} (natural isotope composition)), the chamber were filled with a mixture of argon and CO₂ (1% per cent), at 150 torr. A silicon crystal (diameter 20 mm, thickness 20 mil) with a photomultiplier of the type REN-35 served as neutron detector. The efficiency of the detector was determined according to Baier. Fig. 2 shows the efficiency as a function of the energy of various threshold energies 0.1, 0.2, and 0.3 Mev. The electronic apparatus used to measure the pulse distribution in the detector with respect to time is described in detail. Fig. 3 illustrates block scheme. Fig. 4 recorded pulse waveform diagram. Fig. 5 shows the time distribution of the pulses recorded with the measurement of the neutron spectrum of the U^{238} fission. Besides neutrons and γ -rays of the fission the following were also recorded: 14-Mev protons and parts of the apparatus, radiations of the activated primary neutrons and parts of the apparatus, radiations of the activated

Card 2/4

substances, neutrons, and γ -quanta due to primary neutron scattering, and 2.5-Mev neutrons from the accelerator. Details and accuracy of the comparison of the measured values from the background are discussed. The neutron spectra of U^{235} and U^{238} fission are shown in Figs. 6 and 7. All curves show a similar course: a steep ascent, a peak, and an asymptotic in the coordinates $\ln(\text{Y}(\text{E})/\text{E})$ and E . The spectrum may be approximately represented by

$$\text{Y}(\text{E}) \propto \frac{1}{\text{E}^2} \exp(-\text{E}/T) + (1-\alpha) \exp(-\text{E}/T_0) \exp(-\text{E}/T_1) \text{ on } \frac{\ln(\text{Y}(\text{E})/\text{E})}{\text{E}}$$

The analytical results are listed in Table 1. The following parameter values are indicated for U^{235} : $T = (1.06 \pm 0.03)$ Mev; $T_0 = (0.37 \pm 0.04)$ Mev;

α (fraction of evaporated neutrons) = (0.16 ± 0.02) ; for U^{238} : $T = (0.15 \pm 0.03)$ Mev; $T_0 = (0.21 \pm 0.03)$ Mev; $\alpha = (0.21 \pm 0.03)$. The average number of neutrons emitted in the fission is 4.17 ± 0.50 (U^{235}) and 4.17 ± 0.50 (U^{238}), of which are Soviet.

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4.28 ± 0.39 (U^{238}), the ratio $\bar{Y}(\text{U}^{238})/\bar{Y}(\text{U}^{235}) = 1.03 \pm 0.03$. The following data were obtained: $\bar{Y}^{235} dY/dE_n = 0.112 \pm 0.011$ and $\bar{Y}^{238} dY/dE_n = 0.115 \pm 0.011$ (E = neutron energy). In conclusion, the authors thank Yu. Yu. Olshevsky, A. V. Malyshev, Yu. A. Kudryavtsev, Yu. A. Pashkov, A. I. Reznichenko, T. S. Korobkina, and V. V. Ushakov for their assistance, L. A. Kostyleva for computer calculations, G. M. Klimov, V. A. Kostylev, Yu. S. Litvin, Yu. I. Tsvetkov, Yu. A. Belyaev, and L. P. Kudryavtsev. There are 9 figures, 2 tables, and 21 ref-

SUBMITTED: August 5, 1959

AUTHORS: Vasil'yev, Yu. A., Zamyatnin, Yu. S., Toropov, P. V., 89-12-9/29
Fomushkin, E. F.

TITLE: Measurement of the Neutron Spectrum in the Area below 0,5 MeV by
Means of the Time of Flight Method (Izmereniye spektrov neytronov
v oblasti energiy nizhe 0,5 MeV metodom vremeni proleta)

PERIODICAL: Atomnaya Energiya, 1957, Vol. 3, Nr 12, pp. 542-544 (USSR)

ABSTRACT: By applying an impulse source of neutrons the secondary neutron spectrum is measured, which develops, if 14 MeV neutrons pass through layers of uranium. A fission chamber, which was connected with a 50 channel analyzer, was used as a neutron detector. The distance between source and detector was 6 m.

The energy spectra for the following samples were shown by a graph:

- a) U²³⁵ : 2,7 cm thick ($\sim 1/3 \lambda_{in}$)
- b) U²³⁸ : 2,5 cm thick ($\sim 1/3 \lambda_{in}$)
- c) U²³⁸ : 8 cm thick ($\sim in$)

The spectra obtained from a) and b) originate from a simple interaction between 14 MeV neutrons and the uranium nuclei: It can be assumed that in the measured area of energy the development of the secondary neutrons originate from evaporation from

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Measurement of the Neutron Spectrum in the Area 0,5 MeV by Means of the Time of Flight Method. 89-12-9/29

the stimulated conditions of the compound core. For the case c) the development of a higher number of slow neutrons was ascertained. These are the consequence of a multiple-inelastic interaction which confirms the existence of low situated levels in the U₂₃₈ nucleus. There are 3 figures and 3 references, 2 of which are Slavic.

SUBMITTED: July 20, 1957

AVAILABLE: Library of Congress

Card 2/2

VASIL'YEV, Yu.A.; ZAMYATNIN, Yu.S.; IL'IN, Yu.I.; SIROTININ, Ye.I.;
TOROPOV, P.V.; FOMUSHKIN, E.F.

Measuring the spectra and average number of neutrons in the
fission of U²³⁵ and U²³⁸ induced by 14.3 Mev neutrons.

Zhur.eksp.i teor.fiz. 38 no.3:670-684 Mr '60.

(MIRA 13:7)

(Neutrons) (Nuclear fission) (Uranium--Isotopes)

L 2435h-66 EdF(e)/EMT(m) WH

ACC NR: AP6007259

SOURCE CODE: UR/0363/66/002/002/0357/0362

AUTHOR: Toropov, N.A.; Zhukauskas, R.-S.M.; Aleynikov, F.K.

33

B

ORG: Institute for Chemistry and Chemical Technology AN LitSSR
(Institut khimii i khimicheskoy tekhnologii AN LitSSR)TITLE: The structural transformations of synthetic cordieriteSOURCE: AN SSSR. Izvestiya. Neorganicheskiye materialy, v. 2, no. 2,
1966, 357-362

TOPIC TAGS: cordierite, crystal structure, silicate

ABSTRACT: The test samples were of cordierite synthesized from glass in a heat treatment of from 0.5 to 120 hours, at temperatures from 1100 to 1460°C. The heat treatment was done in a Silit furnace in platinum crucibles, with subsequent air cooling. Glasses of three composition were investigated: a stoichiometric cordierite composition, a composition with 10 weight % more silicon dioxide, and a composition with 10 weight % less silicon dioxide. X-ray investigations were carried out on a URS-50I unit. Results are shown in graphic and tabular form. As the result of prolonged heat treatment at 1400°C a lower rhombic form was obtained from the higher hexagonal cordierite. On raising the temperature up to 1460°C, the reverse transition was

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UDC:548.19

L 24354-66

AP6007259

ACC NR: obtained with ordering of the structure of the cordierite. The process of transition from the high to the lower cordierite was observed electromicroscopically. It was established that the polymorphous transition with formation of rhombic cordierite is accompanied by partial amorphization of the crystal structure which proceeds at a high rate in a narrow temperature interval. The rhombic modification of cordierite is stable in the temperature range up to 1440°C. Orig. art. has: 4 figures and 2 tables.

SUB CODE: 07,11/ SUBM DATE: 29Jun65/ ORIG REF: 008/ OTH REF: 005

Card 2/2 *pla*

L 24279-66 EWP(e)/EWT(m)/EWP(j)/T/EWP(t)/ETC(m)-6 IJP(c) DS/JD/WW/JG/RM/WH

ACC NR: AP6009792

SOURCE CODE: UR/0062/66/000/002/0212/0217

70

13

AUTHOR: Bondar', I. A.; Toropov, N. A.

ORG: Institute of Silicate Chemistry im. I. V. Gribenshchikov, Academy
of Sciences, SSSR (Institut khimii silikatov Akademii nauk SSSR)

TITLE: Phase equilibria in the ytterbium oxide-alumina system and their
comparison with equilibria in other Ln_2O_3 - Al_2O_3 systems

SOURCE: AN SSSR. Izvestiya. Seriya khimicheskaya, no. 2, 1966,
212-217

TOPIC TAGS: phase diagram, phase equilibrium, alumina, aluminum oxide,
aluminate, samarium compound, ytterbium compound, hardness, density,
melting point, chemical resistant material, rare earth compound

ABSTRACT: Phase diagrams were constructed for the Yb_2O_3 - Al_2O_3 and
 Sm_2O_3 - Al_2O_3 systems. X-ray studies confirmed the compounds $2\text{Yb}_2\text{O}_3 \cdot \text{Al}_2\text{O}_3$
and $3\text{Yb}_2\text{O}_3 \cdot 5\text{Al}_2\text{O}_3$. The physical properties of the La, Er, Sm, Y, Tb and
Yb aluminates were compared. The invisibility, hardness, density and
chemical resistance of the rare earth aluminates helps make them
suitable for high frequency ceramics, luminophores, materials for
absorption of neutrons and protection against gamma radiation, and

Card 1/2

UDC: 539.26+546.65

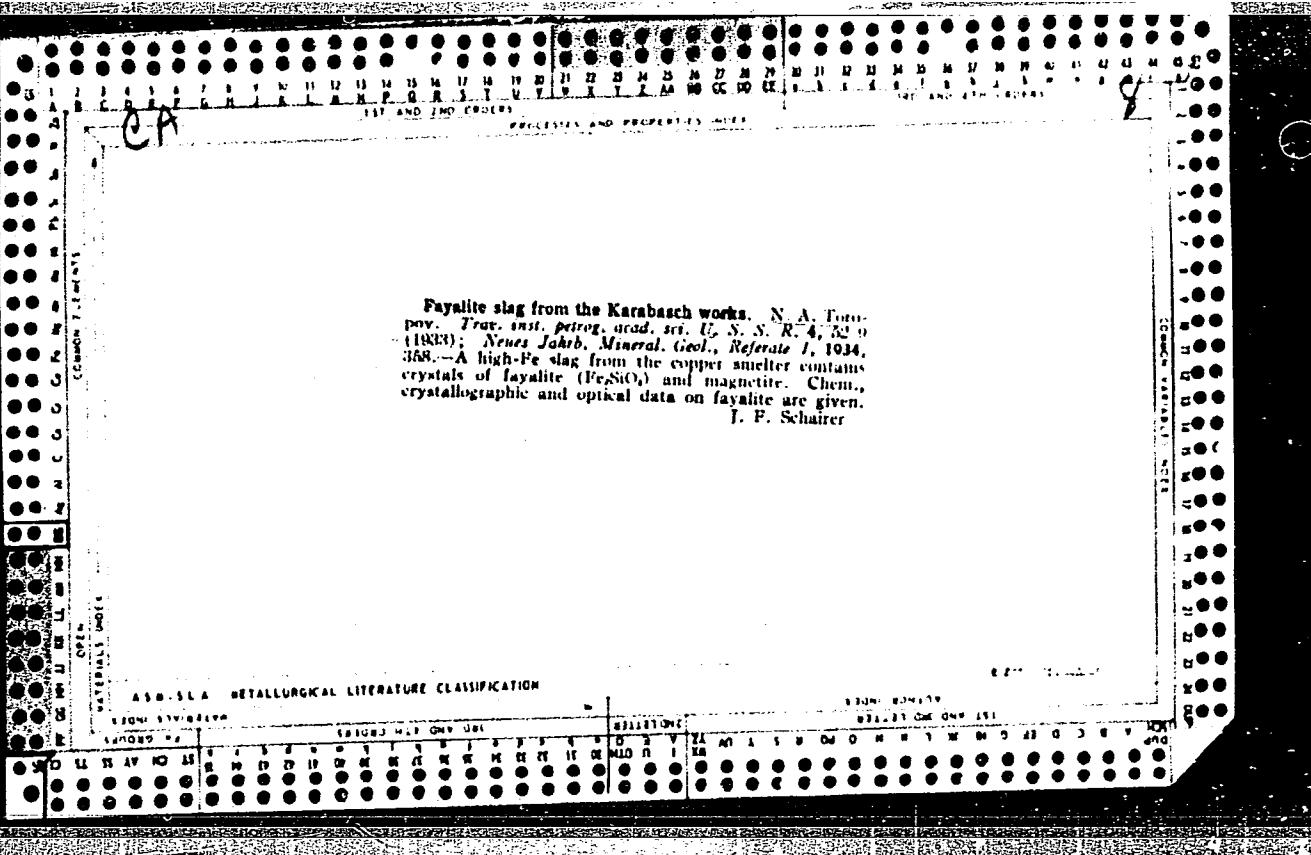
L 24279-66

ACC NR: AP6009792

materials resistant to high temperature and aggressive media. Orig. art.
has: 1 table and 5 figures.

SUB CODE: 11, 07/ SUBM DATE: 02Sep63/ ORIG REF: 003/ OTH REF: 001

Card 2/2 ✓

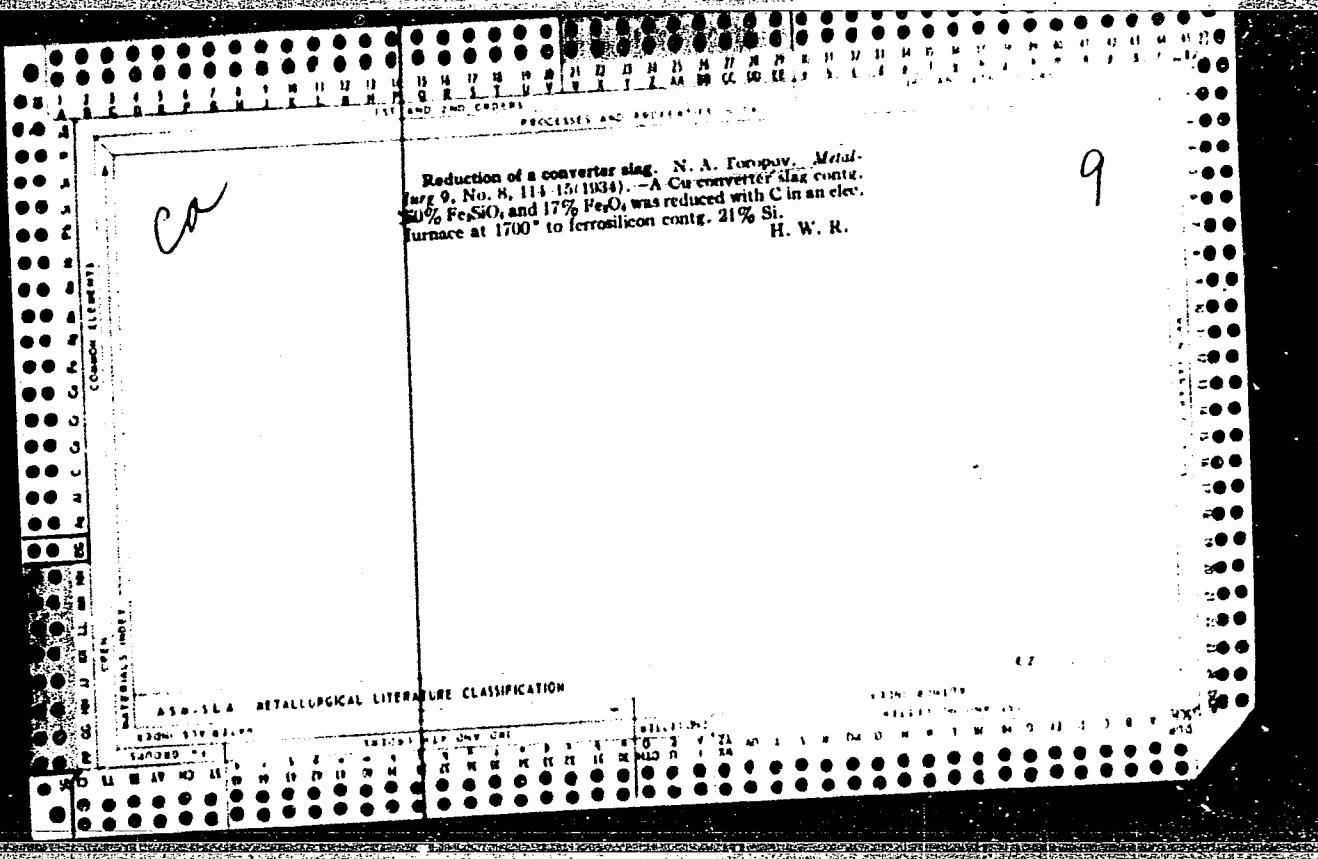


Petrographic investigations of some ferrochromite slags and of fused magnesite. N. A. Finsen, *Ind. and Engin. Acad. us., U. R.*, S. S. G., *Edelstahl-Lesung-Festband* 437-42 (1934); *Chem. Zentral.* 1937, II, 1247-8. Chem. and mineralogical investigations of slags from basic and acid fusions in the elec. furnace indicate that slags from the ferrochromite process consisted chiefly of $MgO\cdot Al_2O_3$ minerals of the forsterite-monticellite group of the formula $Mg(Ca, Fe)_SiO_4$, and glass; in the acid process more glass is present in the slag. Cr_2O_3 occurs in the form of a spinel or as a silicate. A spinel with 4.17% Cr_2O_3 shows a refraction of $n = 1.731 + 0.002$. The refraction is somewhat higher for monticellite from the basic process. Crystals of periclase and slight amounts of forsterite are present in the fused magnesite. The optical properties of fused and of causticized magnesite are similar. The suitability of slags from the production of ferrochromite for the manuf. of abrasives depends upon the spinel content. M. O. Moore

M. G. Minge

APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001756330003-3"



CA

2002-03-20 2002-03-20

Chemical and mineralogical investigation of barium humates. N. A. Tropov. *Compt. rend. acad. sci. U. R. S. S. I. No. 2-3, 445-456* (Berlin 150-1) (1935).—Three compds. were found, all apparently with congruent m. pt.: $\beta\text{Ba}_2\text{O}_3\text{Al}_2\text{O}_5$ (green), $n = 1.735$, double refraction weak, $d_{100} = 4.64 \pm .01$, hygroscopic and very sol. in water. $\text{BaO}_2\text{Al}_2\text{O}_5$, cubic crystals, $n = 1.693$, $d_{100} = 3.99 \pm .03$, m. p. = $1820^\circ \pm 20^\circ$, readily sol. in water, not hygroscopic. $\text{BaO}_2\text{Al}_2\text{O}_5$, hexagonal crystals

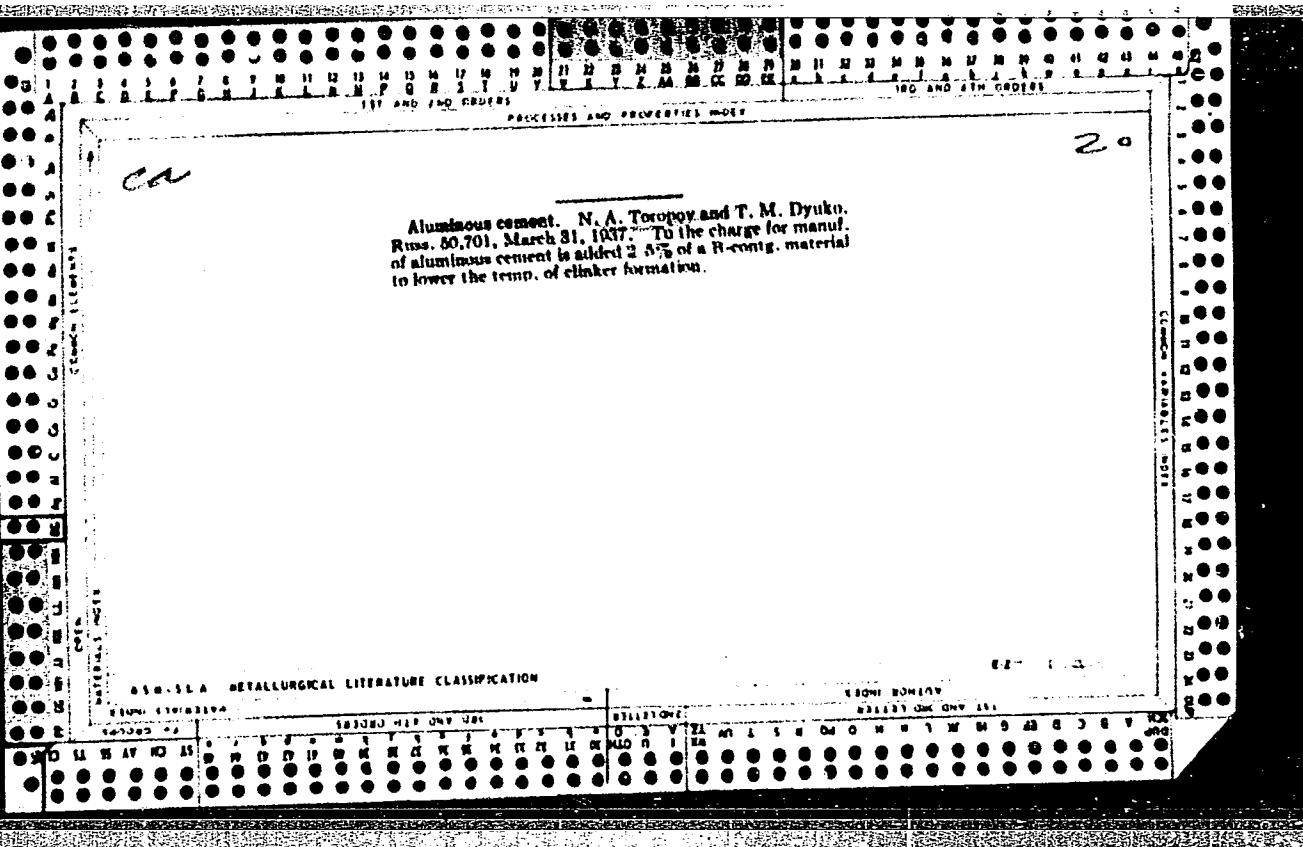
with perfect basal cleavage, $a_0 = 1.694$, $a_{\infty} = 1.702$, $d_{\infty} = 3.69 \pm .01$. These properties are close to those reported for β -Al₂O₃ and the compd. may therefore be a solid soln. of BaO in Al₂O₃. The results do not agree with those of v. Wartenberg and Reusch (*C. A.* 20, 6230).

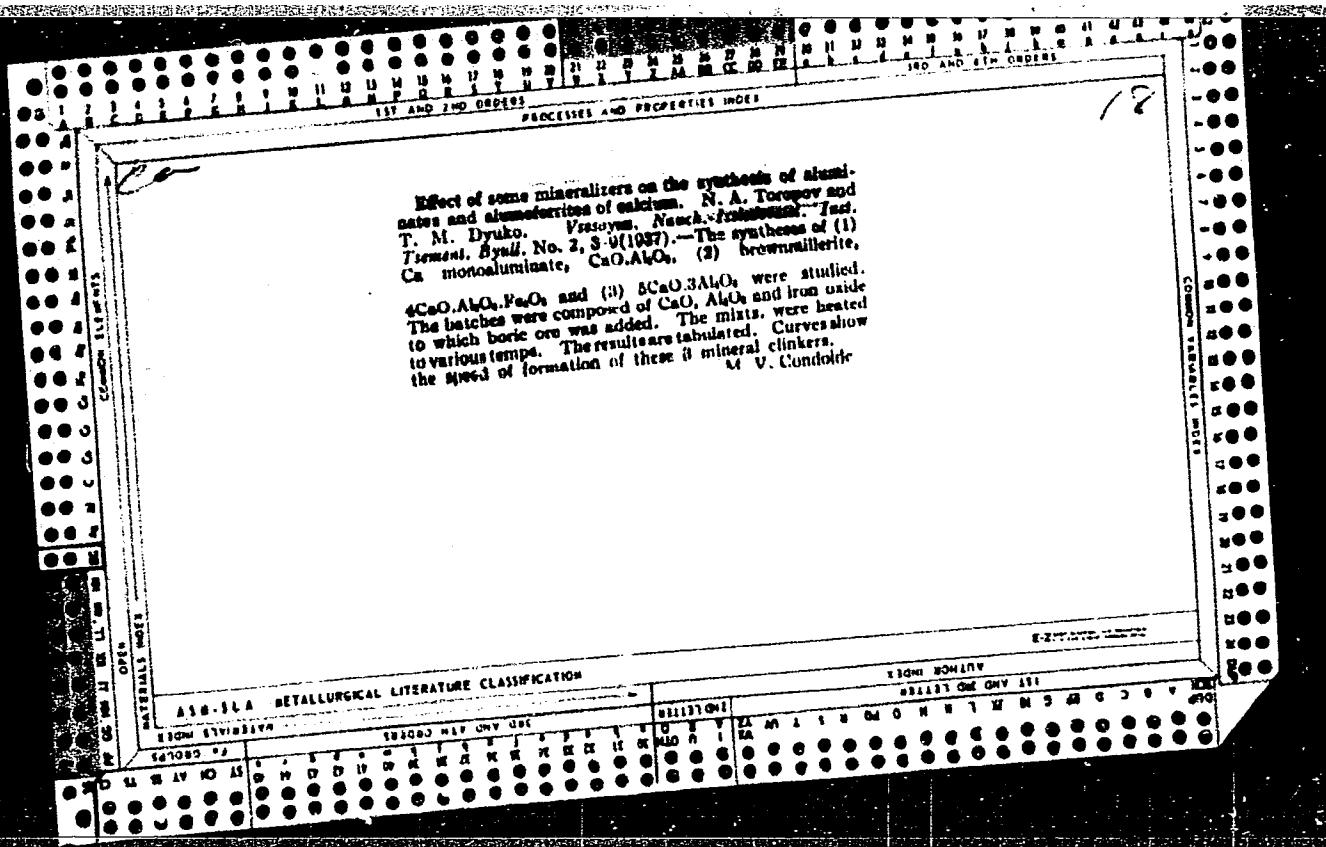
(S. A. 20, 1990).
Michael Fleischer

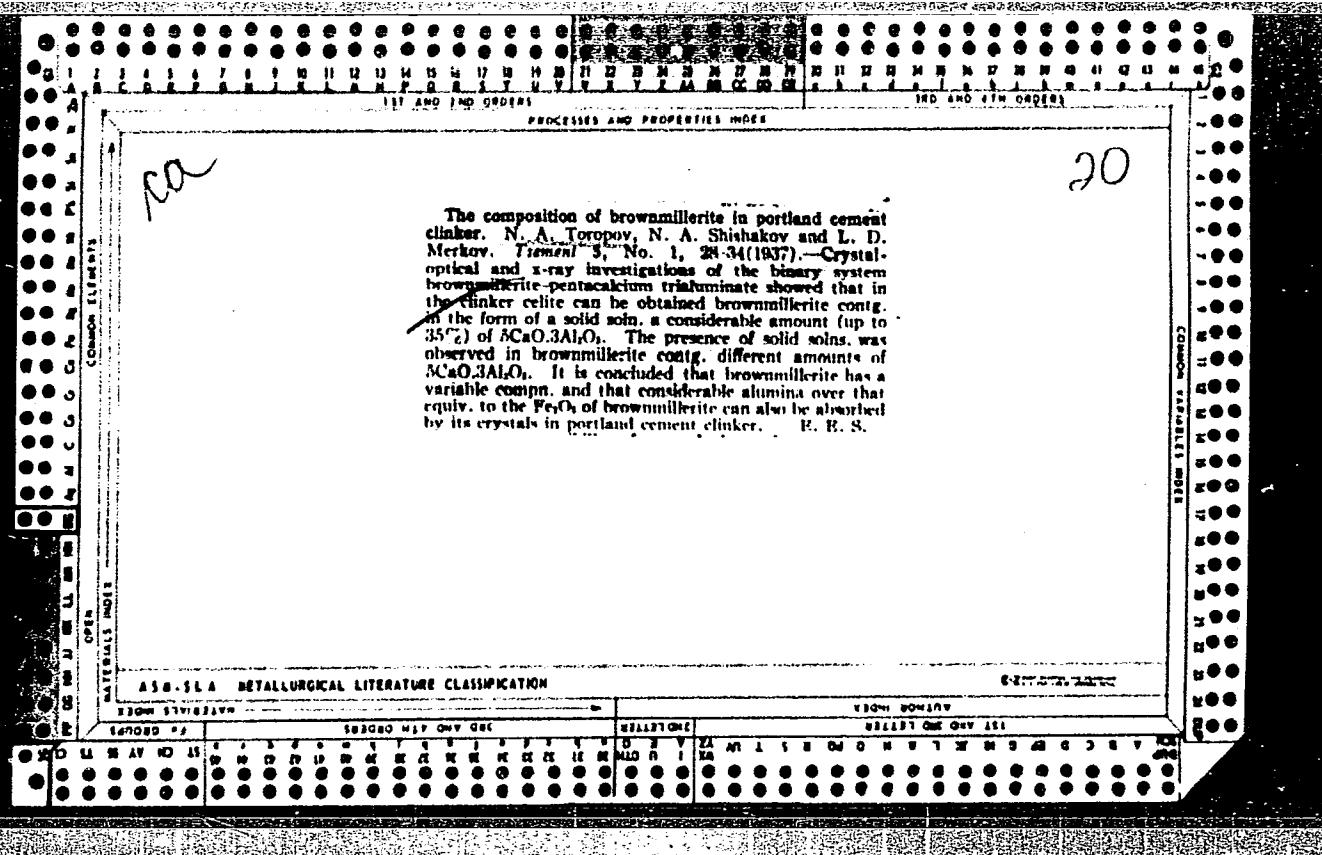
ASH-SLA METALLURGICAL LITERATURE CLASSIFICATION

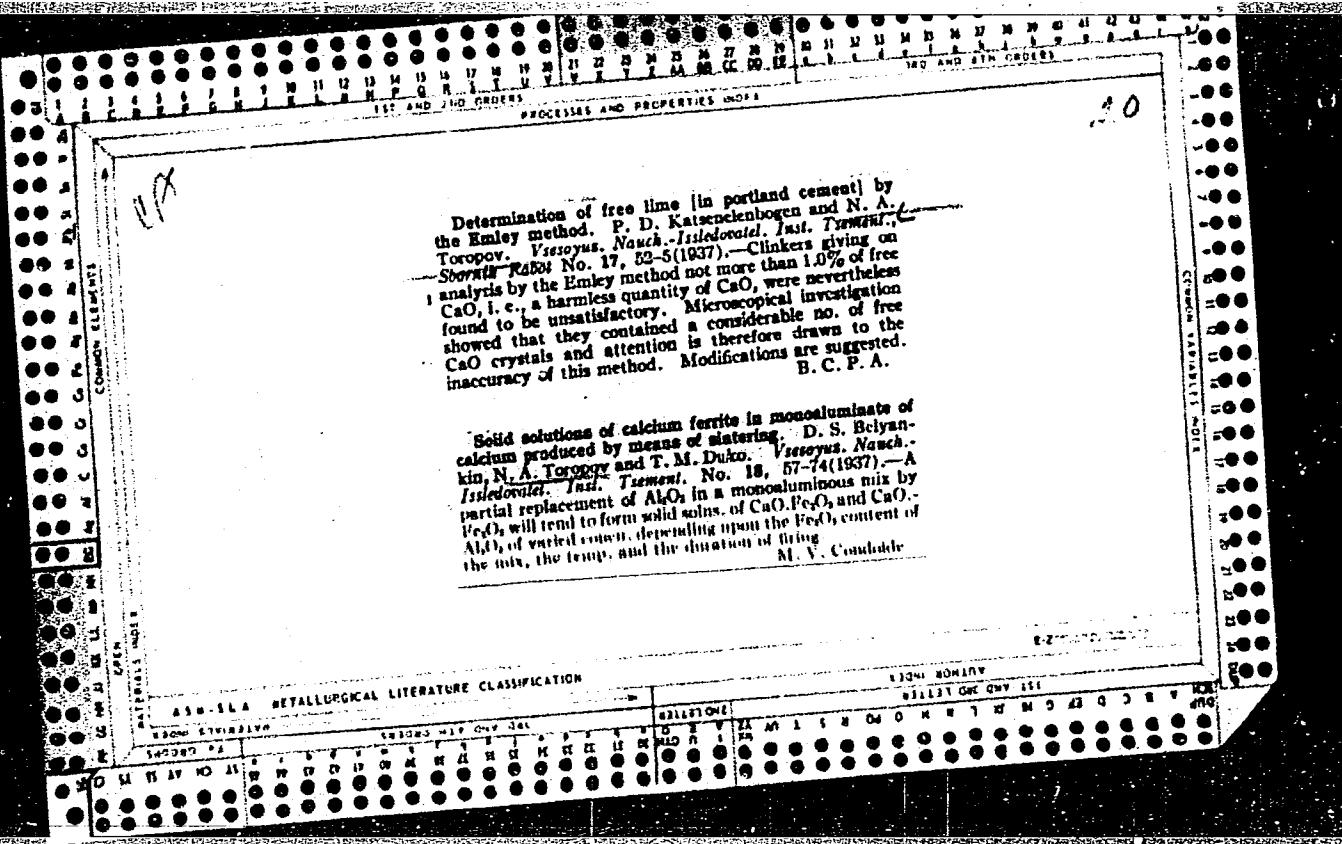
APPROVED FOR RELEASE: 08/31/2001

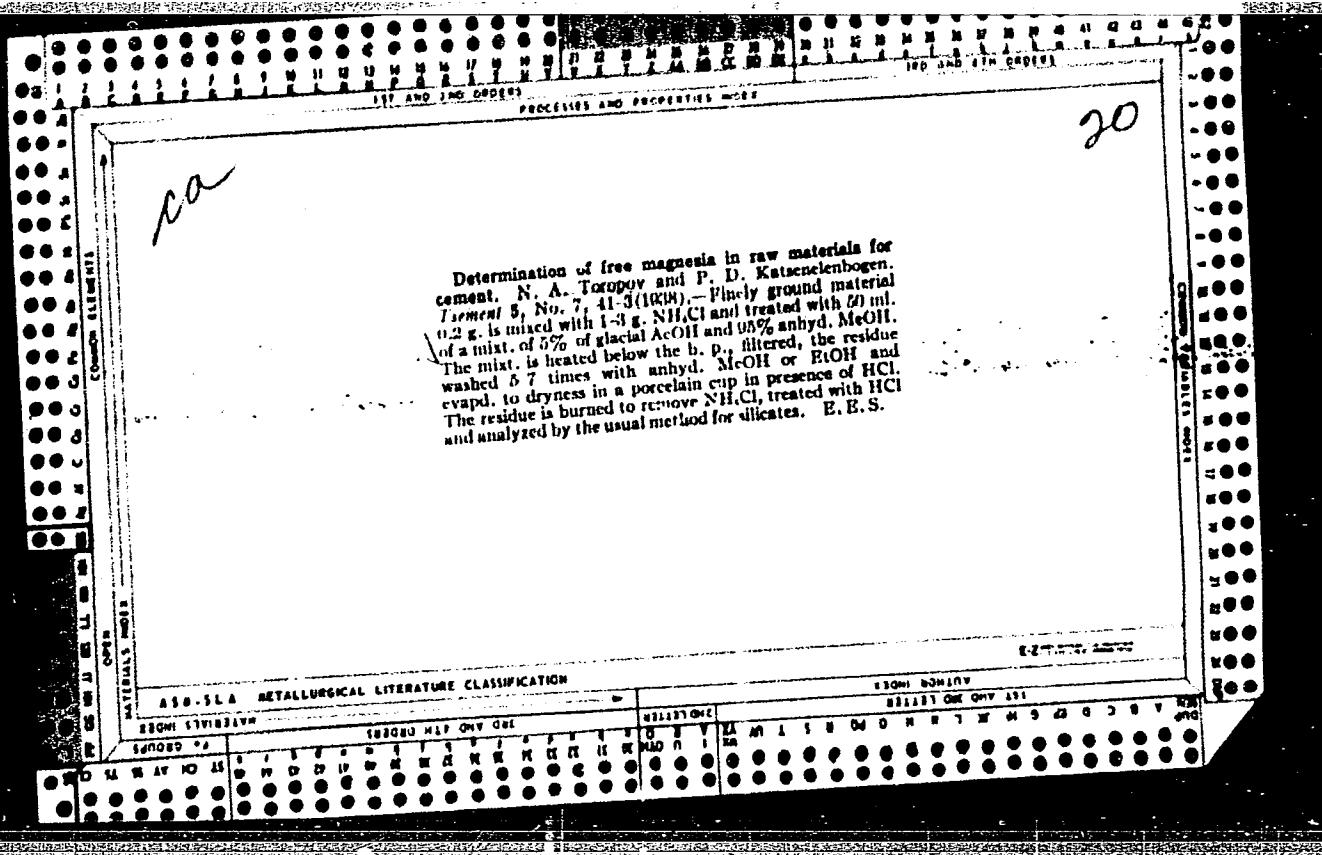
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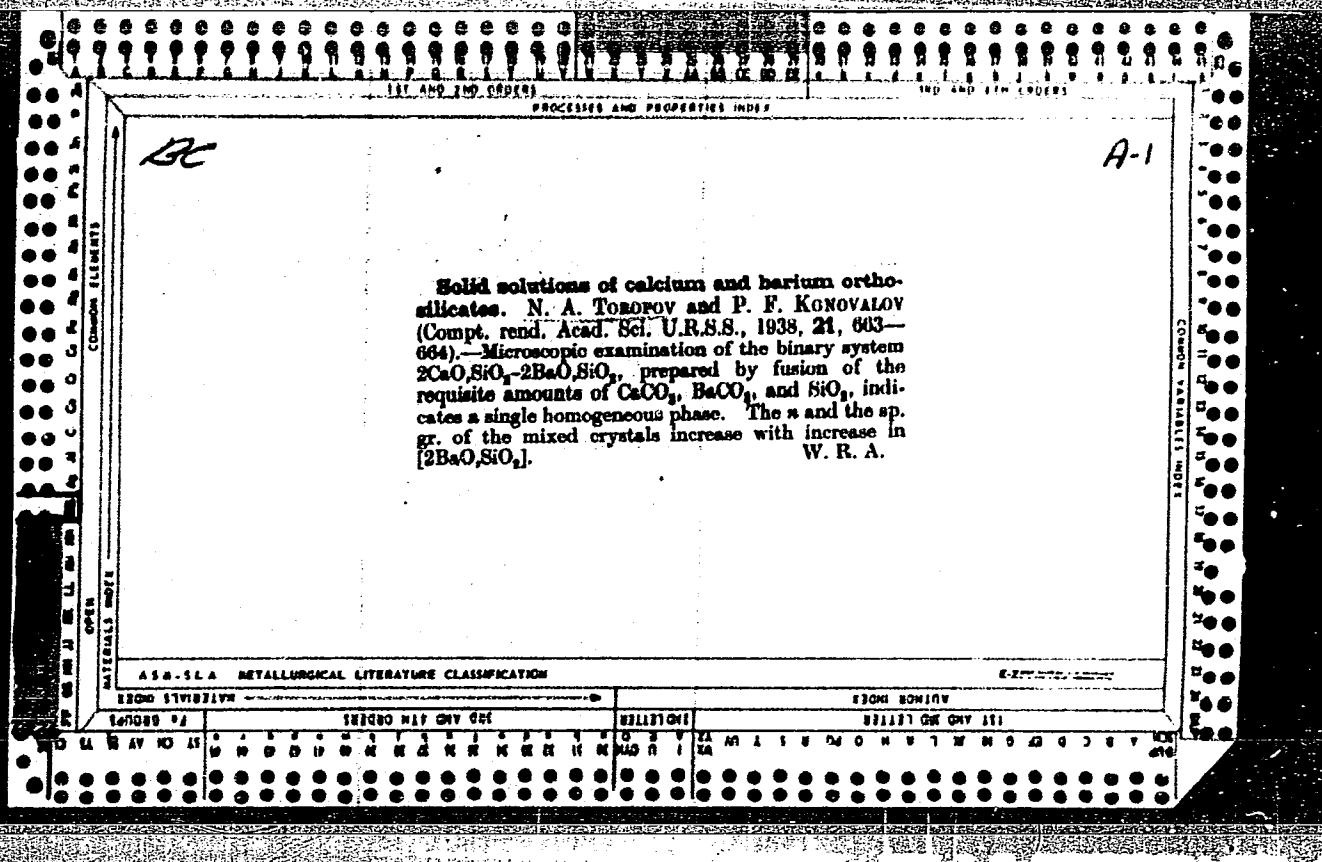


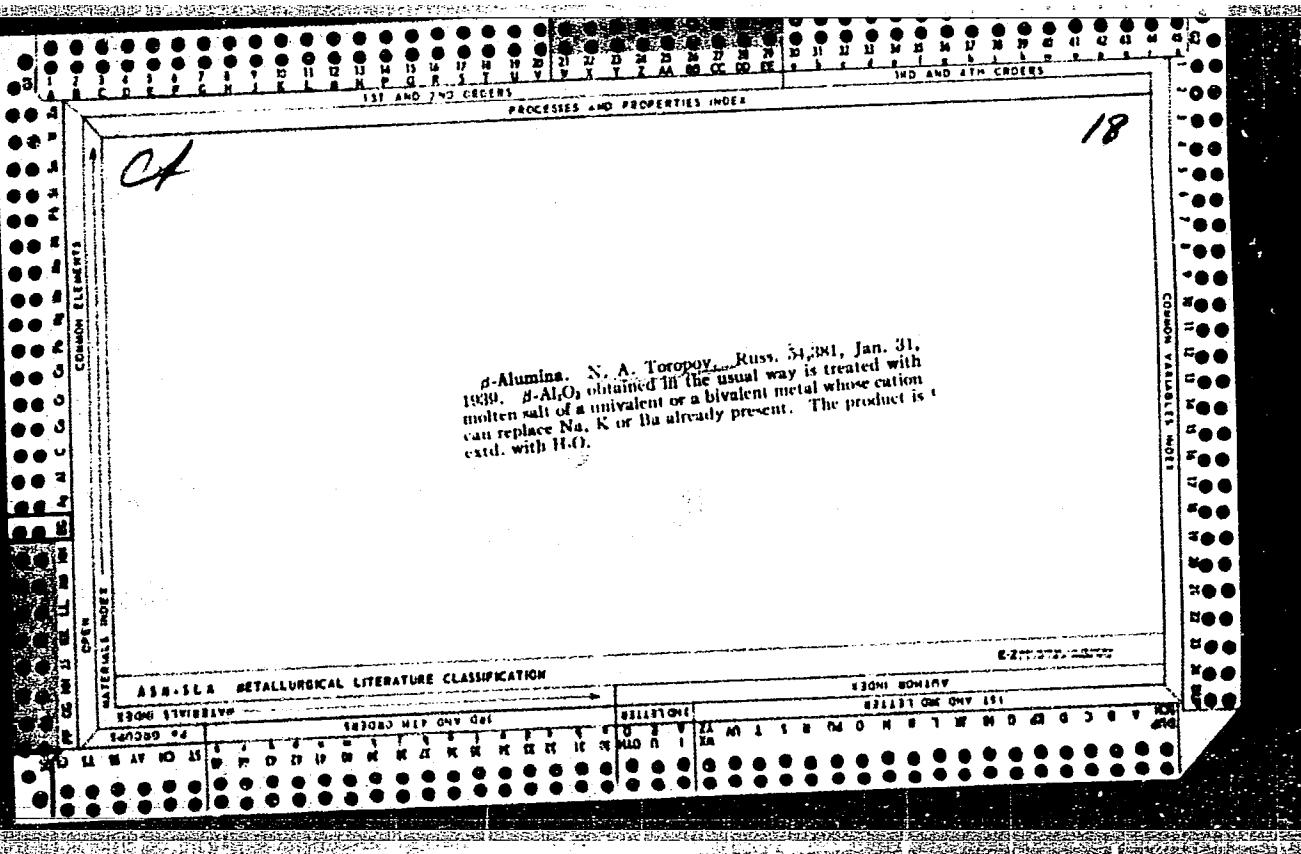


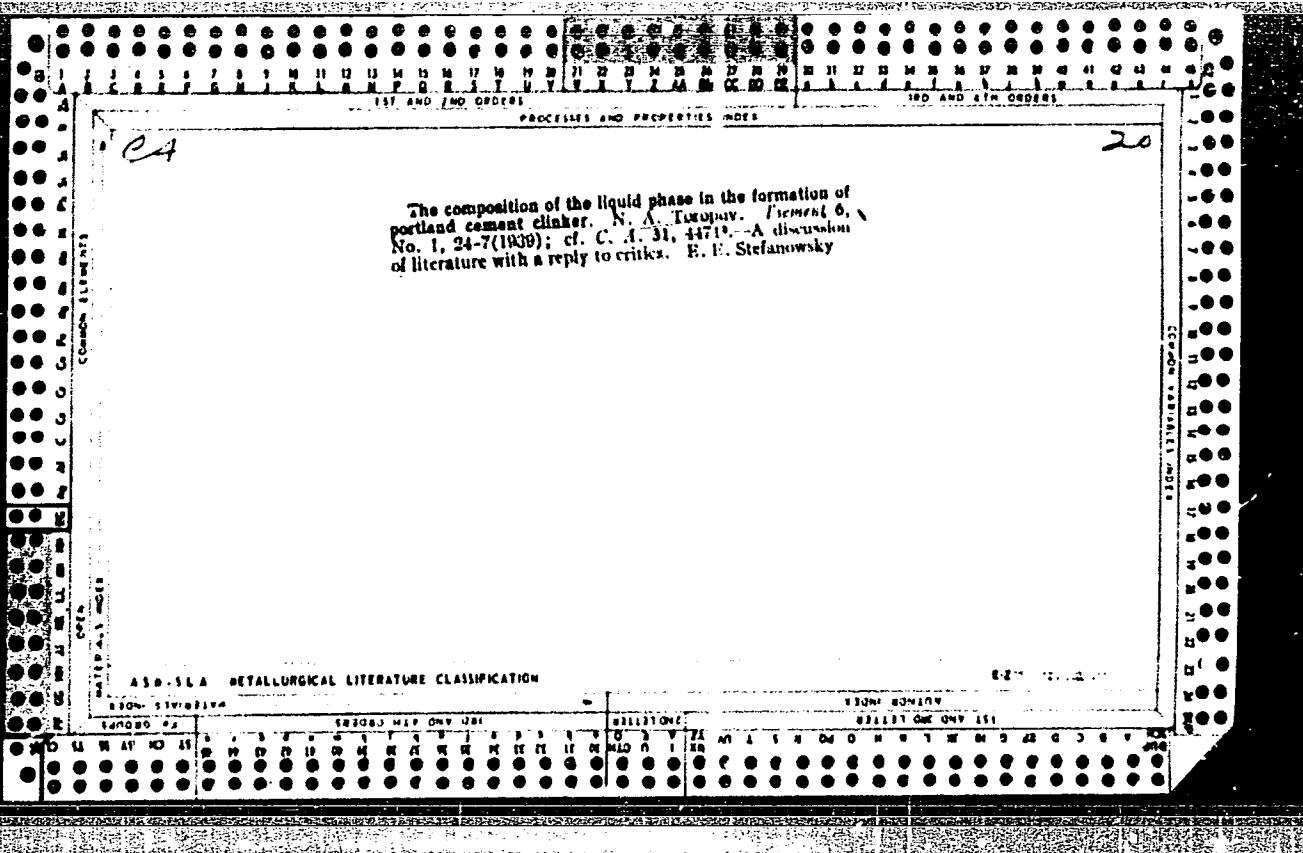










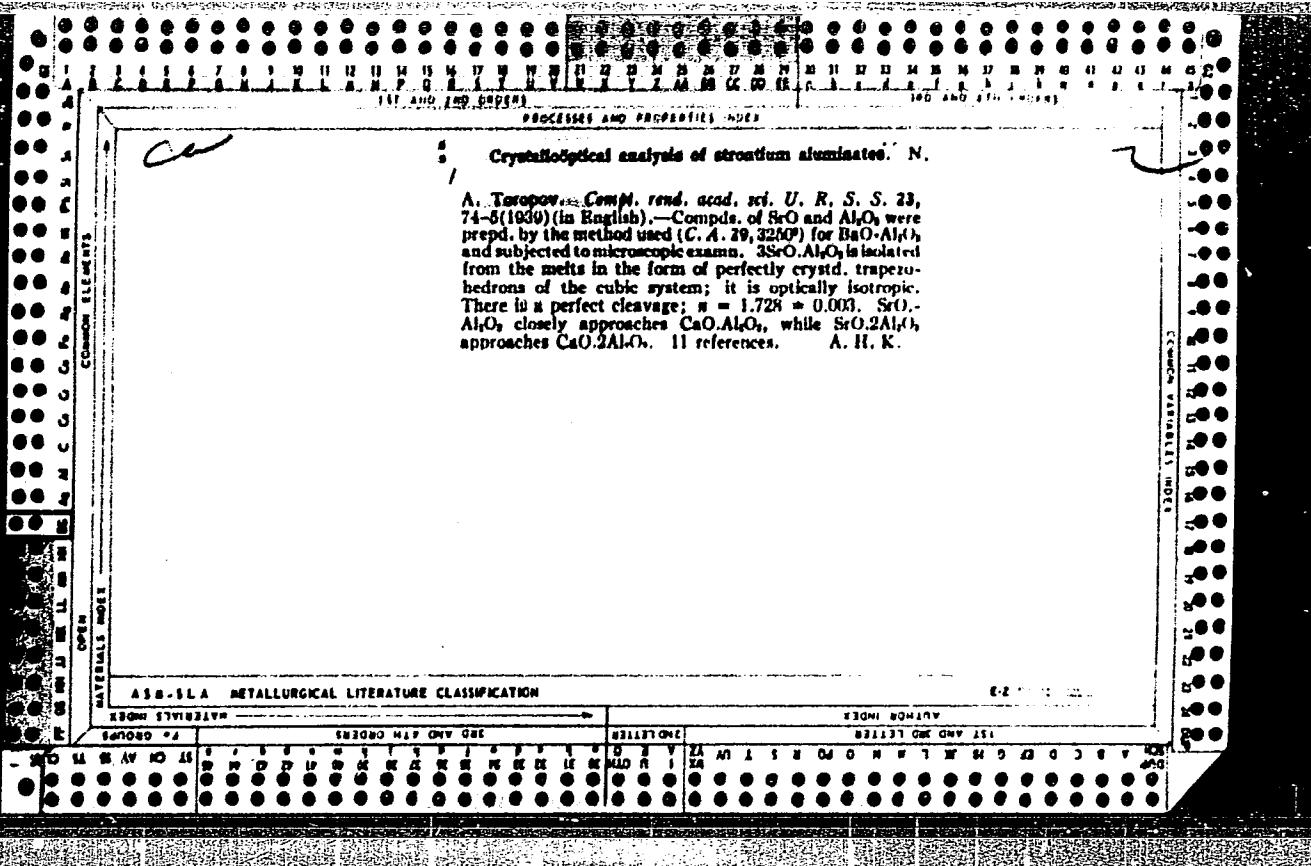


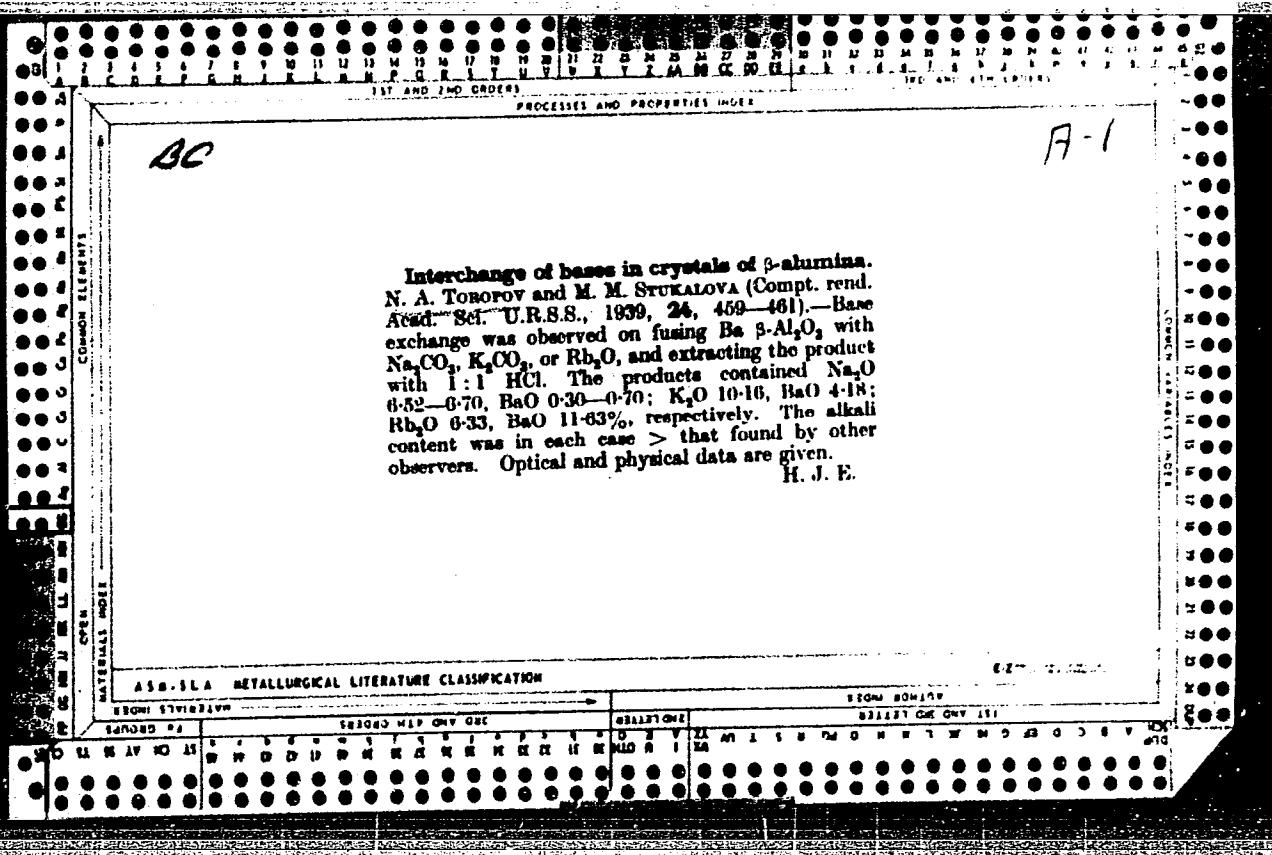
Binary system sodium ferrite-sodium aluminate. N. A. Terenov and N. A. Shishakov. *Acta Physicochim. U. R. S. S.* 11, 277-86 (1938) (in English).—Microscopic details of the π and Debye x-ray photographs show that the system $\text{Na}_2\text{O} \cdot \text{Al}_2\text{O}_5 - \text{Na}_2\text{O} \cdot \text{Fe}_2\text{O}_3$ forms a continuous series of solid solns. The sep. components have very similar patterns, and both are thermally stable at 1500° . Data obtained for $\text{Na}_2\text{O} \cdot \text{Al}_2\text{O}_5$ agree with those of Brownmiller and Boggs (*C. A.* 24, 4223) but those for $\text{Na}_2\text{O} \cdot \text{Fe}_2\text{O}_3$ do not agree with those of Goldztaub (*C. A.* 27, 2690). E. H. Bathman.

(C. A. 37, 2000).

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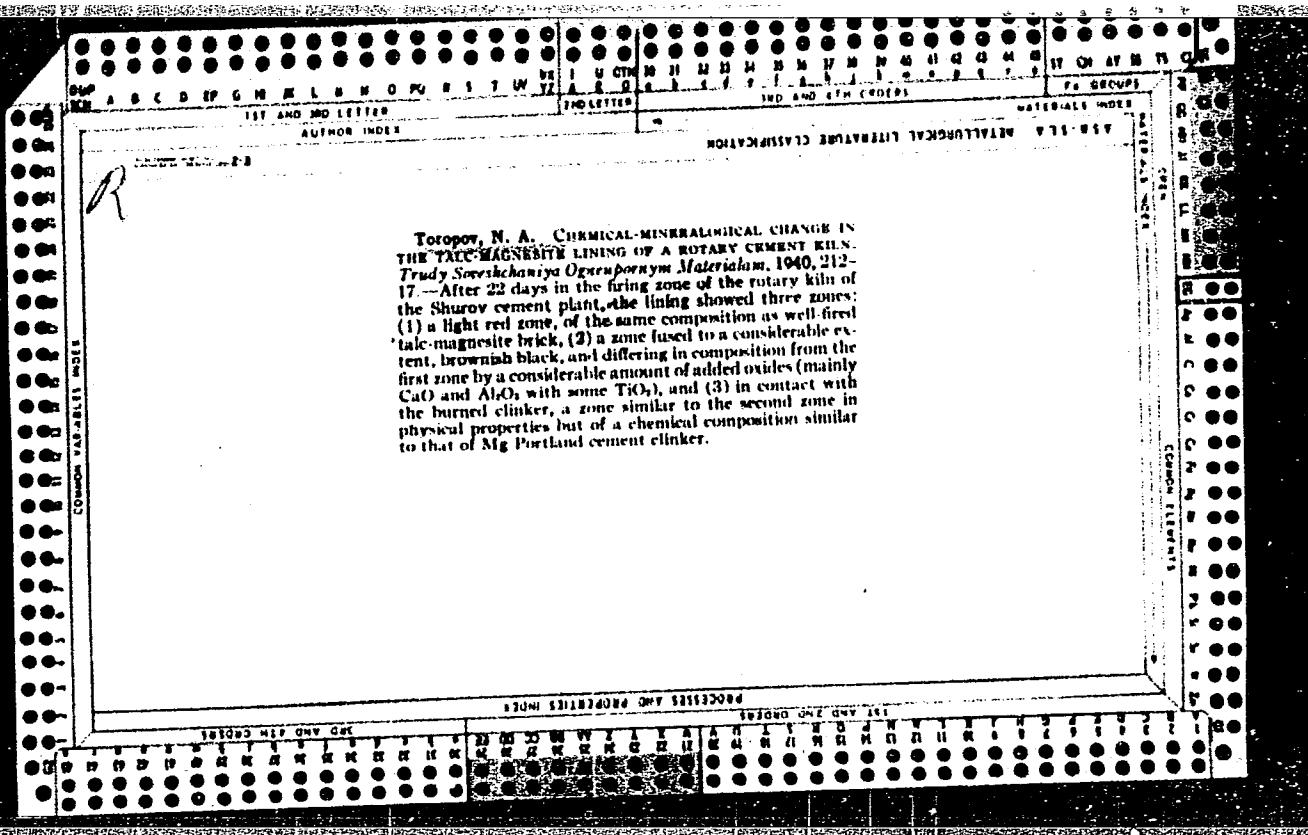
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A.C.S. *10/17/61*

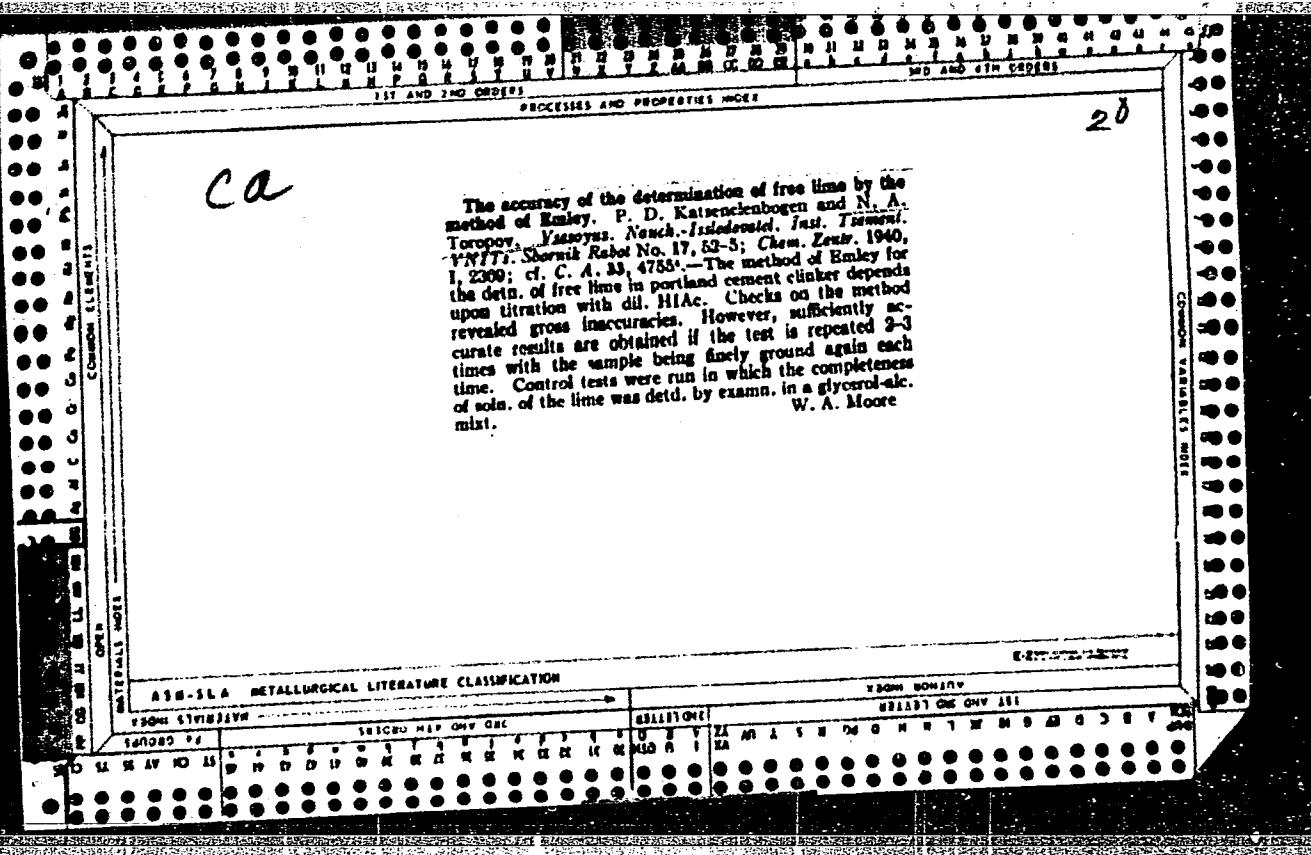
Hydraulic cement of high density. N. A. TOKOVY AND
P. F. KONOVALOV. Russ. 58,561, Dec. 31, 1940; Chem.
Abs., 20, 1033 (1945).—To the usual cement ingredients,
are added Ba compounds, e.g., BaO.



PC

A-1

Binary systems magnesium oxide-boron anhydride. N. A. Toropov and P. F. Konovalov (*J. Phys. Chem. Russ.*, 1949, 16, 1102-1110). Mixtures containing <1 MgO per 1 B_2O_3 form two layers below 1165°, and their cooling curves show an endothermal transformation at 1080° and an exothermal one at 780°. $MgO.B_2O_3$ melts at 1191°, and $Mg_2O.B_2O_3$ at 1381±5°; cooling curves of their mixtures indicate transformations at 1191° and 1080°. $Mg_2O.B_2O_3$ melts at 1400±5° and forms eutectics at 1300±5° (B_2O_3 , 43% wt.-%) and 1340±



Replacement of sodium in crystals of " β -alumina" with calcium, strontium and barium. N. A. Toropov and M. M. Stukalova. *Compt. rend. acad. sci. U.R.S.S.* S. 27, 974-7(1940)(in English); cf. *C. A.* 33, 44331--"beta-Alumina" serves to denote a rather extensive group of aluminates characterized by relatively large Al_2O_5 contents. One g. of Na β -aluminate powder (200-mesh) was fused with a 6-fold quantity of alk. earth chloride for 1 hr. The resulting crystals were not homogeneous, but after a 2nd 1-hr. fusion period the crystals became optically homogeneous. The prep'd. $\text{CaO}\cdot\text{Al}_2\text{O}_5$ possessed a d. of 3.28, s. of 1.702, and a birefringence of 0.035 with a neg. optic sign. The BaO - and $\text{SrO}\cdot\beta$ -aluminates were similar in properties to the CaO compd. These crystals are of the holohedral class of the hexagonal system and

have parameters in A. units for the CaO-, SrO- and BaO-aluminates, resp., of $a = 5.536$, 5.887 and 5.877 , and $c = 21.925$, 21.945 and 22.67 . H. E. Meissner

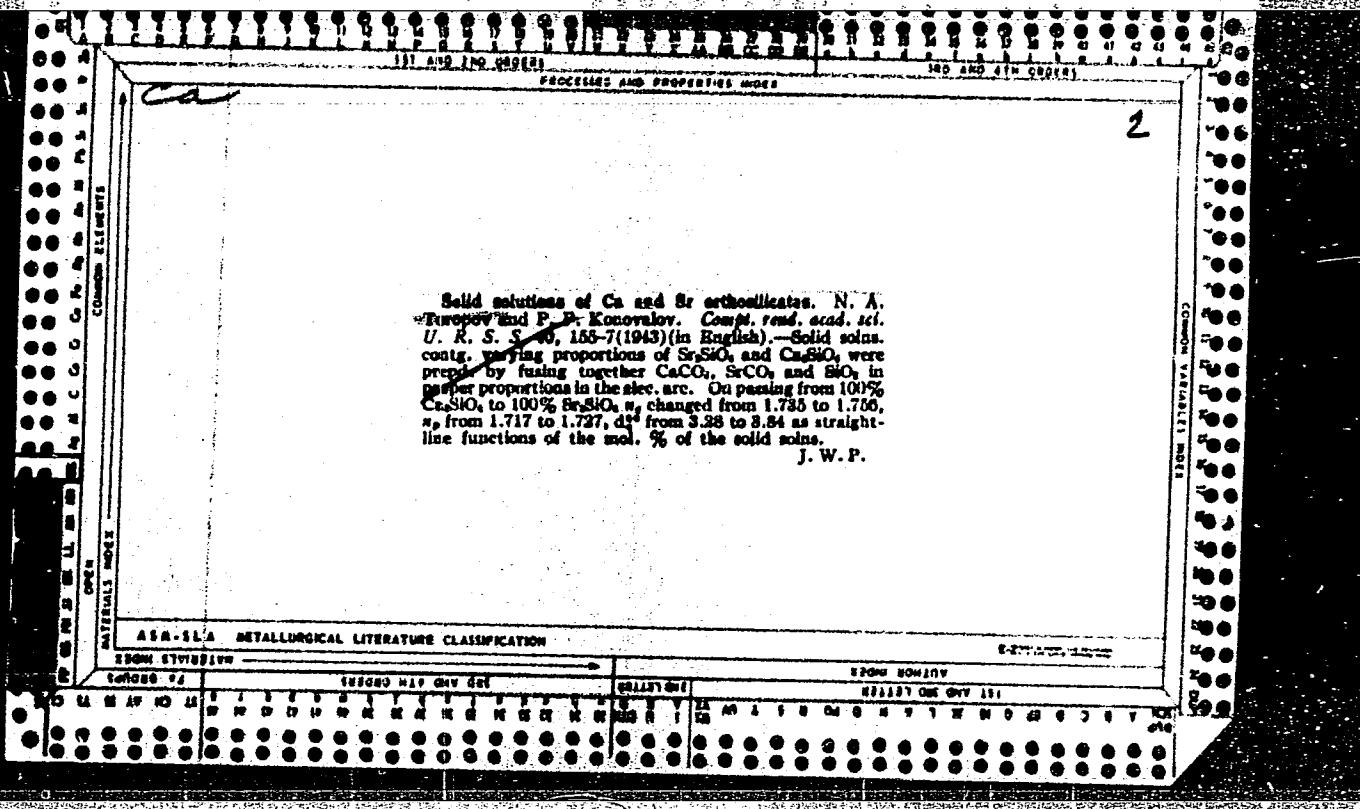
H. E. Measmore

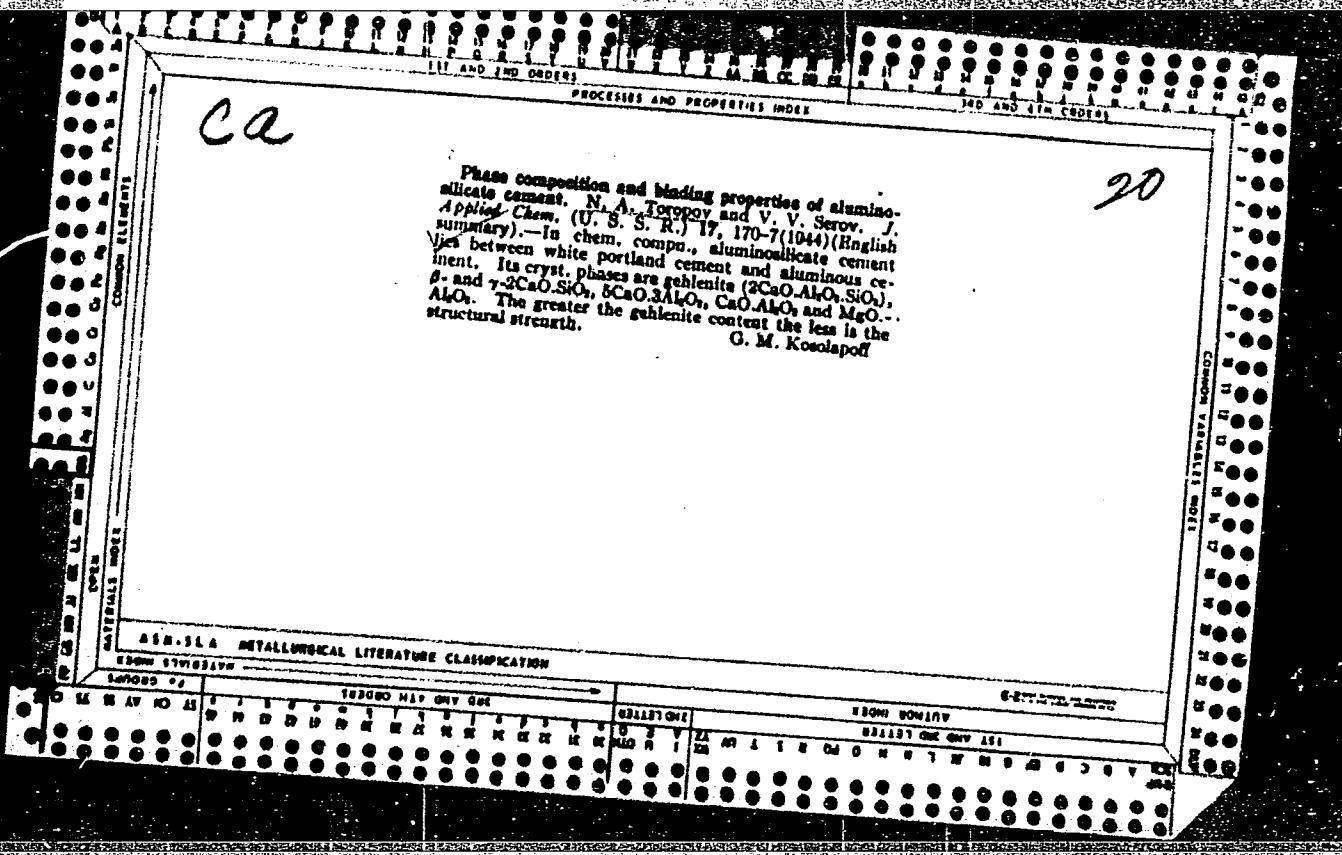
ABSTRACTS OF METALLURGICAL LITERATURE CLASSIFIED BY SUBJECT

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APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001756330003-3"





TOPOROV, N. A.

C
Review of research on the forming and constitution of
clinker and of the structures of hydrated cements. N. A.
TOPOROV. Trudy Vsesoyuz. Soveschaniya Zavod. Lab.
Iz cement. Prom., 3, 45-61 (1915). The following topics
are reviewed: (1) liquid phase of Portland cement clinker,
(2) constitution of clinker, and (3) microstructure of hy-
drated cements. No references. B.Z.K.

TOROPOV, N. A.

Toropov, N. A. - "A survey of the newest achievements in the area of the chemistry and petrography of cements," Trudy 4-go Vsesoyuz. soveshchaniya zavodskikh laboratori tsement, prom-stil, Leningrad, 1948, p. 43-69.

SO: U-3850, 16 June 53, (Letopis 'Zhurnal 'nykh Statey, No. 5, 1949).

"APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001756330003-3

TOROPOV, N.A., doktor tekhnicheskikh nauk, professor.

Academician D.S. Beliakin's work. TSegment 14 no.6:3-4 N-D '48.
(Building materials) (MLRA 9:5)

APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001756330003-3"

TOROPOV, N. A.

PA 35/49T66

USSR/Metals

Nickel Oxide

Ferric Oxide

Dec 48

"Solid Solutions in the System NiO-Fe₂O₃," N. A. Toropov, A. I. Borisenko, All-Union Sci Res Inst Giprostremen, 4 pp

"Dok Ak Nauk SSSR" Vol LXIII, No 6 1970-8

Conducted chemical, crystal-optic, and X-ray analysis of mixtures of NiO and Fe₂O₃ for various temperatures to determine temperature interval in which ferrites form. Table shows content of FeO (in %) during the heating of various mixtures in a platinum furnace for 3½ hours, and relative

35/49T66

USSR/Metals (Contd)

Dec 48

intensity (angstroms) of Debye crystallogram lines for samples, obtained by clinkering at 1,100°. Another table shows composition of the mixture, content of NiO and Fe₂O₃ in weight, and characteristics of the mixture. Submitted by Acad D. S. Belyankin 27 Oct 48.

35/49T66

CA

//Belyakin, D. S., Terekov, N. A., and Lepin, V. V.:
Fiziko-khimicheskie sistemy silikatnoj tekhnologii (Physico-
chemical Systems of Silicate Technology). Moscow:
Promtrotizdat. 1949. 251 pp. R. IR.76. Reviewed
in J. Am. Ceram. Soc. 34, No. 4, Ceram. Abstracts 7A(1951).

1957

TOROPOV, N. A.

SOLID SOLUTIONS IN THE SYSTEM $\text{CoO-Fe}_2\text{O}_3$. N. A. TOROPOV, E. A. PORAI-KOSHITS, AND A. I. BORISENKO. Doklady Akad. Nauk S.S.R. v. 66, 905-8 (1949); cf. C. A. 43, 4552b. In an equimol. mixt. of the pptd. hydroxides, ferritization is complete on 40 hrs. heating at 1100° of pellets dried at 120° and pressed under 3000 kg. /sq. cm. The thermal dissociation $6\text{Fe}_2\text{O}_3 \rightarrow 4\text{Fe}_3\text{O}_4 + \text{O}_2$, in mixts. with CoO , is less intense than in mixts. with NiO . In the presence of excess CoO , the product obtained is porous, and contains, on microscopic exam., 2 phases, one dark-gray, the other dark-yellow. The amt. of the former decreases with decreasing CoO . With increasing Fe_2O_3 content, the color of the crystals seen in specimens etched with HNO_3 changes from dark-yellow in the equimol. compon., to yellow for $2\text{CoO} + 5\text{Fe}_2\text{O}_3$. With further increasing excess of Fe_2O_3 , a new light phase appears along with the yellow crystals, attaining 45-50% in $\text{CoO} + 6\text{Fe}_2\text{O}_3$. By x-ray examn., free Fe_2O_3 is

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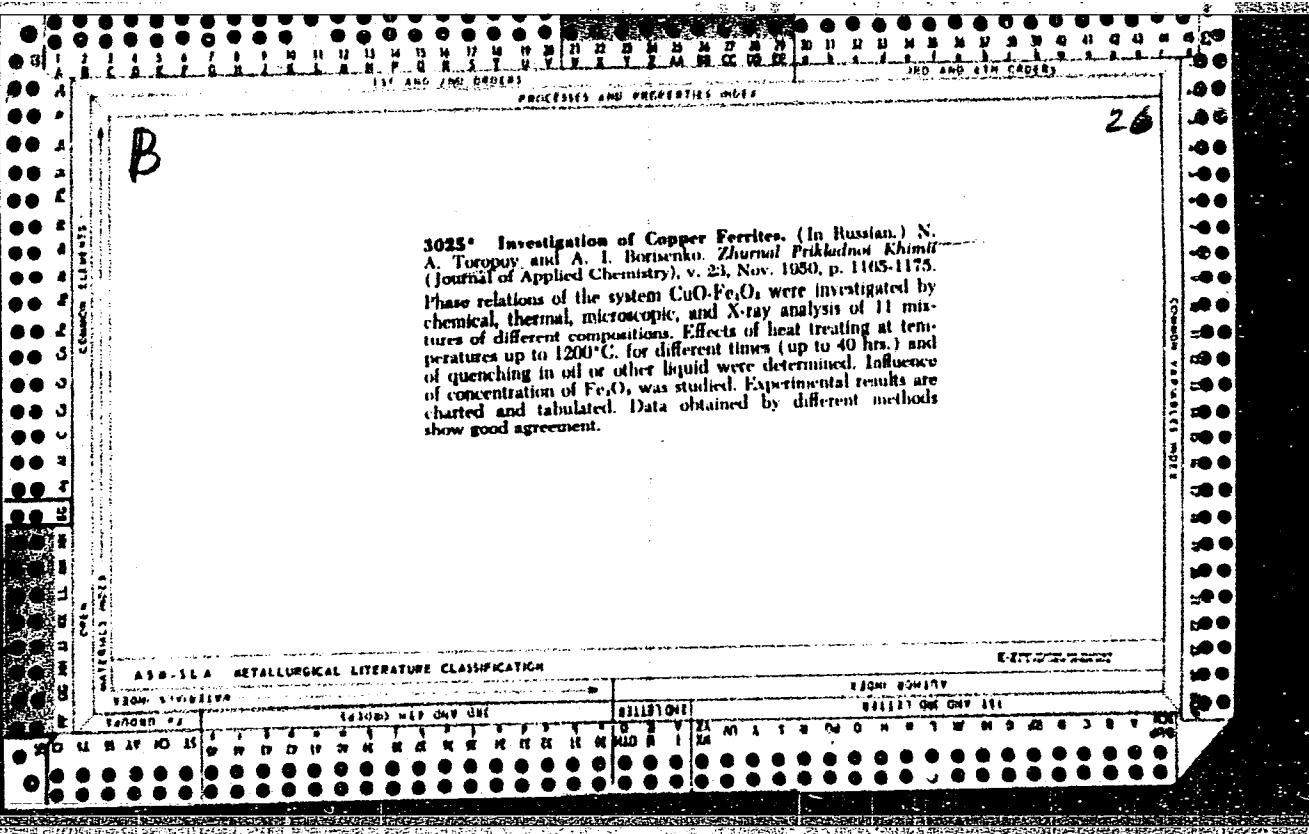
Zinc borates. N. A. Toropov and P. F. Konovalov.
Doklady Akad. Nauk S.S.R. '66, 1105-8(1969); cf.
C.A. 68, 22961. —The system $ZnO-B_2O_3$ was investigated
by means of heating curves, and the solid phases were
identified by x-ray diffraction studies. Three compounds were
found: $3ZnO.B_2O_3$, m. 1120° ; $ZnO.B_2O_3$, m. 1000° .
 $ZnO.3B_2O_3$, decomp. at 900° into $ZnO.B_2O_3$ and a liquid
contg. 2% ZnO . The system is characterized by having
a 2-liquid-phase region extending from 8 to 48% ZnO and
with a lower limit of 990° . The upper limit was not
detd. but was above 2000° . There are 3 eutectics:
 990° , 48% ZnO ; 1010° , 64% ZnO ; 1080° , 82% ZnO .
 m_1 and m_2 are, resp.: $ZnO.B_2O_3$, 1.643, 1.676; $3ZnO.B_2O_3$,
1.669, 1.720; $ZnO.3B_2O_3$, 2.004, 2.020. Arild J. Miller

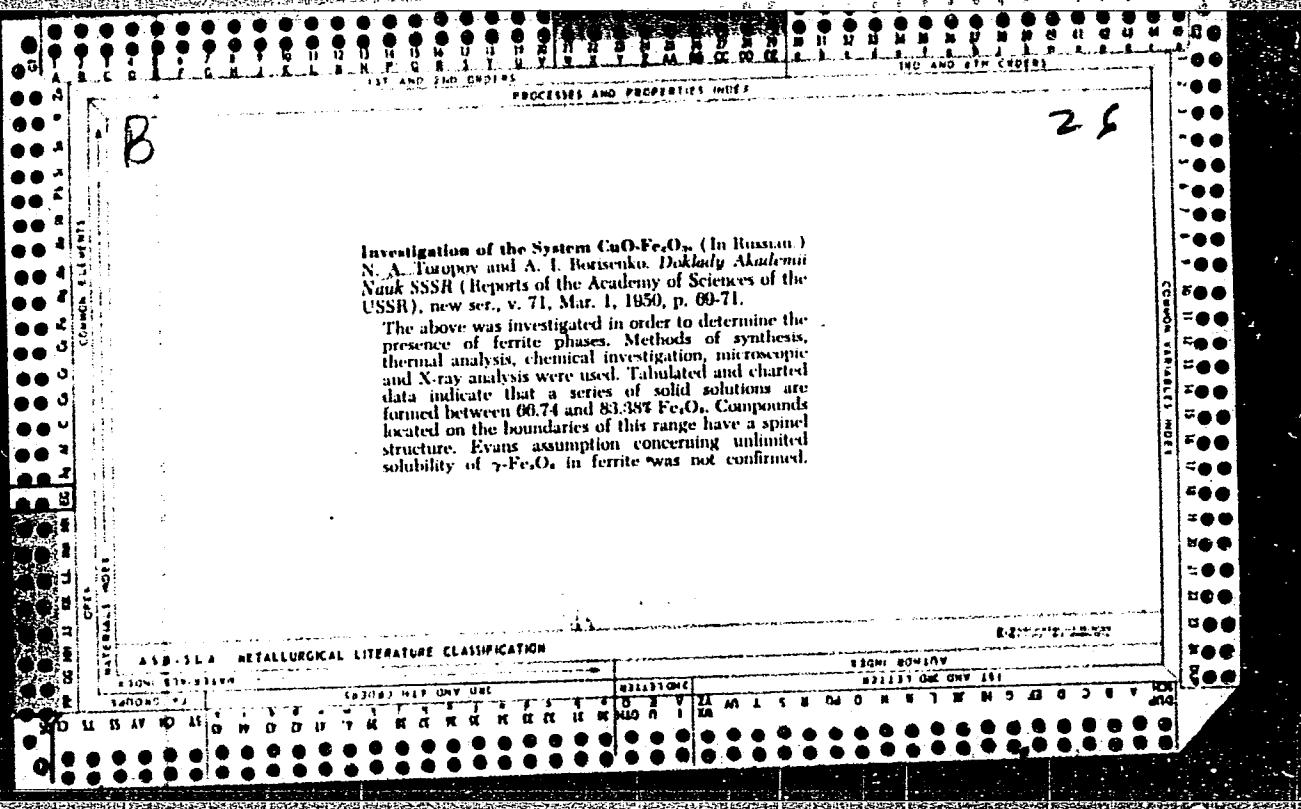
BIR

26

11010 *Khimika Kremala i Fizicheskaya Khimika Silikatov, (The Chemistry of Silicon and the Physical Chemistry of Silicates.)* (Russian.) K. S. Evstrop'ev and N. A. Toropov. 304 pages. 1950. State Publishing House of Constructional Material Literature, Moscow, U.S.S.R. (QD181.56 Ev79k)

A textbook. Subjects covered include the structure of silicates in crystalline, vitreous and fused states; bases of the theory of constructing constitution diagrams for silicate systems and their application; and description of high-Si compounds.





TOROPOV, N. A.

PA 160T80

USSR/Minerals - Bauxite
Silicates

11 May 50

"New Orthosilicates of Potassium and Sodium," N. A.
Toropov, O. I. Arakelyan, All-Union Aluminum-Magnesium
Inst, 4 pp

"Dok Ak Nauk SSSR" Vol LXXII, No 2 - p. 365

During systematic investigations of reactions in
process of roasting bauxite with limestone and soda,
authors observed two new crystal phases in binary
system $2\text{CaO}\cdot\text{SiO}_2\text{-Na}_2\text{O}\cdot\text{CaO}\cdot\text{SiO}_2$. Gives characteris-
tics of these phases, and formulas of formation of
new silicates and X-ray measurements of interplanar
distances.

160T80

178T85

TOROPOV, N. A.

USSR/Metals - Oxides, Structure

1 Jan 51

"Solid Solutions in the Systems NiO-ZnO-Fe₂O₃ and CuO-ZnO-Fe₂O₃," N. A. Toropov, A. I. Borisenko, Leningrad Tech Inst imeni Lensoviet'

"Dok Ak Nauk SSSR" Vol LXXVI, No 1, pp 85-88

Studies 2 ternary systems. Concludes solid soln of limited concn are being realized in these systems. Mixed ferrites and ferric oxides are components of solid soln in one part of systems, and mixed ferrites and oxides of bivalent metals in the other part. Presents and discusses diagrams of both systems. Submitted 27 Oct 50 by Acad D. S. Belyankin.

178T85

"APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001756330003-3

APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001756330003-3"

TOROPOV, N. A., GALAKHOV, F. YA.

Mulite

"New data on the system $Al_2O_3-SiO_2$." N. A. Toropov, F. Ya. Galakhov. Reviewed by Prof. S. V. Glebov, Ogneupory, 17, No. 7, 1952.

9. Monthly List of Russian Accessions, Library of Congress, October 1952 ~~1953~~, Uncl.

CP

State diagram of the system barium oxide-alumina.
N.A. Tsyppov and V.Ya. Galakhov. *Doklady Akad. Nauk S.S.R.* 82, 69-70 (1952).—The complete m. diagram was
drawn with samples contg. no BaCO₃. The system has 4
eutectic points (compns. in wt. %): 100% BaO-Ba₂Al₅O₁₂
15%; 1710°, 77.5, 22.5; 1700°, 48, 35; 1890°, 17, 83.
There are 3 maxima, corresponding to the compds.: 3BaO-Al₂O₃ m. 1750°; BaO-Al₂O₃ m. 1850°; BaO-Ba₂Al₅O₁₂ m.
1900°.

TOROFOV, N. A.

USSR/Chemistry - Abrasives, Oxide Magnets 1 Feb 52

"Solid Solutions in the System ZnO - Fe₂O₃." N. A. Toropov, A. I. Borisikov

"Dok Ak Nauk SSSR" Vol LXXXII, No 4, pp 607-609

The cryst phases in the system ZnO - Fe₂O₃ are studied microscopically and by X-ray diffraction. The compounds studied were ZnO-Fe₂O₃, ZnO-Fe₂O₃, and others. Comparison of results of X-ray study of the synthesized zinc ferrite with those of New Jersey franklinite shows the 2 substances to be analogous. Compns contg 66.24 - 83.07% Fe₂O₃ by wt form homogeneous solid solns on being heated to

213TR19

1,100°. Compns lying within the limits of a homogeneous ferrite phase exhibit spinel structure, ZnFe₂O₄ (zinc ferrite) is an important ingredient of oxide magnets.

213TR19

CR

Kinetics of formation of dicalcium silicate. N. A. Toto
rov, A. M. Glinting, and L. G. Luginina. *Doklady Akad. Nauk S.S.R.* **84**, 203-5 (1952). — A mixt. (~ 0.9 g.) of 2
moles CaCO_3 + 1 mole SiO_2 , grain size of both reagent:
 $70-80 \mu$, pressed under $250 \text{ kg}/\text{sq. cm}$, was heated at
 $1,350^\circ$, and free CaO and unbound SiO_2 were detd. every
20 min., up to 320 min. The results were evaluated with
the aid of the equation of diffusional kinetics in spherical
particles, $I = 1 - e^{-t/\tau} = (1 - G)^{-1} = Kt$, where G =
fraction reacted, and t = time in min. From $G \geq 0.30$ up,
 K has a practically const. value of $\sim 1.3 \times 10^{-3}$; I in-
creases linearly with t from 140 to 320 min. The silicate
formation rate is detd. by the diffusion of CaO to the SiO_2
across a layer of the product. Jander's equation (C.A. 21,
3738) is inapplicable to these exptl. results, as J 's const.
varies with the degree of conversion. N. Thor

Independent reactions. A. Skralch (Osterr. Akad. Wissenschafl., Graz, Austria). *Monatsh.* **83**, 530-40 (1952). — The concept of independent reaction, i.e. reactions between
which no math. relations exist, is explained and illustrated
in terms of the system of 9 possible reactions among the
compds. A_2O , R_2O , AOH , ROH , AOR , and H_2O , where A
is a univalent acyl and R a univalent alkyl group. Both
the math. method, which involves setting up a matrix of
reactions, and the "chem." method which makes a more
direct use of the chem. equations, indicate that in this sys-
tem the no. of independent reactions is 3 and the no. of inde-
pendent relations among the reactions is 6. In a given sys-
tem there are in general several possible sets of independent
reactions. The no. of independent reactions corresponds
to the no. of independent analyses necessary to det. the
progress of the reactions at a given time and also to the max
no. of stages in which the system may react. E. R. F.

TOROPOV, N.A.; BORISENKO, A.I.

Physicochemical study of solid solutions formed by orthosilicates
of calcium and barium. (In: Soveshchanie po eksperimental'noi
mineralogii i petrografii. 4th, Moscow, 1952. Trudy, Moskva, 1953.
No.2, p.214-229).
(MLRA 7:3)

1. Fiziko-khimicheskaya laboratoriya Instituta khimii silikatov
Akademii nauk SSSR. (Silicates) (Systems (Chemistry))
(Solutions, Solid)

TOROPOV, Nikita Aleksandrovich, 1908-

BULAK, L.N.; CHETVERIKOV, S.D., redaktor.

[Course in mineralogy and petrography and the principles of geology]
Kurs mineralogii i petrografii s osnovami geologii. Pod red. S.D.Chetverikova. Moskva, Gos. izd-vo lit-ry po stroit. materialam, 1953. 486 p.

(MLRA 7:1)

(Mineralogy) (Petrology) (Geology, Stratigraphic)

"APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001756330003-3

APPROVED FOR RELEASE: 08/31/2001

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180°-185° until clear, cooling, diluting with water and warming to 100°, re-dissolution, followed by drying at 100° for 4 hr. Chemical analysis and X-ray data fully support its purity. The reaction proceeds through the formation of a glycerate which reacts with the amorphous SiO_2 . This product is hydrated on filtration and washing with water. When the effluent is heated for 2 hr. at 900° it forms fine grained aggregates with refractive index 1.714 and

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CIA-RDP86-00513R001756330003-3"

USSR

The similar problem. N. A. Toropov and V. Yu. Galaktionov. *Voprosy Parozidimostnosti Akad. Nauk S.S.R.* 2, 245-254 (1971); cf. Belyankin and Lapin. *GA* 45, 68976. A vacuum microfurnace is described, for temps up to 2000°, with a W coil as the heating element, and for cylindrical samples of about 2-4 mm. in diam. and height. The temps were measured by an optical pyrometer, and the vacuum was brought to 10⁻⁴ mm. Hg. The formation of thin layers of W metal deposited on the surface of the samples was important because it limited the volatilization

PM

discussed on the basis of the results of the investigation of the low-volatilizing nature of this system.

Facts were reproduced: BULK VOLATILIZATION WAS OBSERVED IN THE PRESENCE OF AN OXIDIZING AGENT.

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CIA-RDP86-00513R001756330003-3

measuring, electrocast multite glass-tint blocks. The
dimensions of the

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CIA-RDP86-00513R001756330003-3"

TOROPOV, N.A., professor; VOL'FSOY, S.L., dotsent.

Intensification process of clinker firing. TSement no. 4:12-16 Jl-Ag '53.
(MIRA 6:8)
(Cement kilns)

TOROPOV, N.A.

Chemical Abstracts
May 25, 1954
Cement, Concrete and
other Building Materials

(5) 7
The binding of calcium oxide in burning portland cement raw mixes, as a function of the size of granules. (4) A. V. Toropov and I. G. Likhachev, *Silikattech*, 4, 470-1 (1953); cf. C.A. 48, 972h. For a special study of the optimum conditions for accelerated burning of portland cement, the authors detd. the functional relations between the size of granules from the raw mix and the temp. distribution in a kiln, the time of exposure to the firing temps. in the sintering zone, finally the effects of addns. of NaF or CaF₂. The degree of reaction of free CaO with the ingredients of the raw mix is measured by analytical methods and controlled under the microscope. The firing process is distinctly accelerated by reducing the diam. of the raw mix granules from 5 to 1.25 mm. The mineralizing effects of the fluorides are better for relatively lower firing temps., while at higher temps. their volatilization is marked. The time for a complete clinkering is for granules of 1.25 mm. diam. at 1600° only 3.5 min., and in the presence of fluorides (for a clinker rich in 2CaO·SiO₂) even only 10-15 sec. Industrial kilns, with granules of more than 5 mm. in diam., require in general a sintering time at 1500° of 6 to 7 min., and if CaF₂ was added, only of 1.5 to 3 min. In the last sintering period, the rate of CaO binding in the clinker minerals is distinctly decreased. It is important to know that granules of less than 5 mm. in diam. in most industrial kilns are unnecessarily retained in the sintering zone.
W. Hitzel

TOROPOV, N. A., Prof.; LUGINAN, I. G.

Cement

Effect of quick heating on the formation of cement clinker. TSement 19, No. 1, 1953.

9. Monthly List of Russian Accessions, Library of Congress, June 1953. Unclassified.

TOROPOV, N.A.

Glory
B

✓ Effect of the grain size of raw materials on assimilation of CaO during burning of portland cement. N. A. Toropov and I. G. Lukinina. *Cement* 19, No. 2, 17-22(1959).—

With decreasing grain size, assimilation of CaO increases considerably. Mineralizers (NaF and CaF₂) exert an accelerating influence; however, the effect is less the higher the temp. and the smaller the grain size. By decreasing grain size to 1.25 mm., burning at 1600° is reduced to 3.5 min. for av. clinker and to 10-15 sec. for belite clinker with addn. of mineralizers. However, rise in temp. to 1600° is not always possible. The grain size of the product from the rotary kiln is diverse. Of the total clinker, 52% by wt. were grains < 5.00 mm. In 50% of the mixt., clinker

formation is completed in 5-7 min. after entrance into the burning zone. However, the presence of large grains favors the increase of CaO in the total mass of clinker. When burning av. clinker with CaF₂, assimilation of CaO in small grains is completed in 1.5-3.0 min. in the burning zone; for large grains, the burning time is increased 4-5 times.

B. Z. Kamich

TOROPOV, N. A.

(2)

Chemical Abstracts
Vol. 48 No. 5
Mar. 10, 1954
Cement, Concrete, and Other Building
Materials

Intensifying the firing of clinker. N. A. Toropov and
G. L. Vol'son. *Cement* 19, No. 4, 12-16(1953). Two
portland-cement mixes, differing only in the content of
di- and tri-Ca silicates were fired at 1200 and 1300° with
0.06, 0.025, and 0.012 g.-equivs. of fluorides and fluosili-
cates (superphosphate by-products) per 100 g. of the ce-
ment mix. The fluosilicates and fluorides proved more
beneficial than fluor spar. The fluosilicates were, in turn,
more effective than the fluorides of the corresponding cations.
Optimum dosage of fluosilicate was 0.012 g.-equiv. Strength
of the cement specimens was not lowered by these admixts.
B. Z. Kamich

"APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001756330003-3

APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001756330003-3"

TOROPOV, N.A.

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26/3/

(4) 5656* Phase Composition and Certain Ferromagnetic Properties of Manganese-Zinc Ferrites. (Russian.) N. A. Toropov, L. I. Kubkin, E. Zh. Freidenfeld's, and D. Sh. Epstein. Zhurnal Prikladnoi Khimii, v. 26, no. 9, Sept. 1953, p. 682-690. Describes chemical, microscopic, X-ray, and magnetic studies of synthesized ferrites. Tables, diagrams, graphs, 9 ref.

11/19/34

1. TOROPOV, N. A., GALAKHOV, F. YA., BONDAR, Y. A.
 2. USSR (600)
 4. Aluminum Silicates
 7. Structural diagram of the ternary system BaO-Al₂O₃- SiO₂. Dokl. AN SSSR 89, no. 1, 1953.
9. Monthly List of Russian Accessions, Library of Congress, May 1953, Uncl.

TOROPOV, N. A.

USSR

/ Investigation of the system tricalcium silicate-tricalcium phosphate. N. A. TOROPOV, A. I. BORISENKO, AND P. V. SUTOKOVA. Zhurn. Akad. Nauk S.S.R., 92 [5] 1015-18

(1973).—Disks made from mixtures of $3\text{CaO}\cdot\text{SiO}_2$ (I) and $3\text{CaO}\cdot\text{P}_2\text{O}_5$ (II) were heated at 1450°C , and every 10 min. a different disk was removed and subjected to microscopic and chemical analysis. Disks prepared of 98 to 50 mole % I and 2 to 50 mole % II were found to consist of tricalcium silicate, free CaO , solid solutions of silicophosphates, and tetracalcium phosphate (III), in amounts depending on the original proportions of I and II. Free CaO increased with time, reaching maximum in 80 min. At this point, tricalcium silicate, dicalcium silicate, CaO , tricalcium phosphate, and traces of tetracalcium phosphate were observed. After 70 min. at 1450°C , free CaO dropped to a minimum. Free CaO is formed in accordance with the reaction $3\text{CaO}\cdot\text{P}_2\text{O}_5 + \text{CaO} \rightarrow 4\text{CaO}\cdot\text{P}_2\text{O}_7$. Detection of III is indicative of the reaction $3\text{CaO}\cdot\text{P}_2\text{O}_5 + \text{CaO} \rightarrow 4\text{CaO}\cdot\text{P}_2\text{O}_7$.

After 80 min., silicocarmelite was also detected and there was a further increase in the amount of free CaO , reaching a constant limit at 100 to 140 min. The final content of free CaO depended on the composition of the disks, increasing to 11.47 wt. % for 0 to 20 mole % II and dropping to 0 for 50 mole % and more II. In all cases, free CaO was less than that expected from decomposed I. This was traced to solution of CaO in silicocarmelite and magelschmidtite with the formation of more basic silicophosphates. D.Z.K.

TOROPOV, N. A.

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Fiziko-khimicheskiye sistemy silikatnoy tekhnologii (Physico-chemical systems of silicate technology, by) D. S. Belyankin, V. V. Lapin, N. A. Toropov. Moskva, Promstroyizdat, 1954.
370 p. diagrs., graphs, tables.
Includes bibliographies.

TOROPOV, N. A.

Diagram of state for the ternary system $\text{BaO}-\text{Al}_2\text{O}_3-\text{SiO}_2$.
N. A. Toropov, B. Ya. Galakhov, and I. A. Bondar. *Bull. Acad. Sci. U.S.S.R., Div. Chem. Sci.* 1954, 647-65 (Engl. translation).—See *C.A.* 49, 6711h.
B.M.B. (2)

TOROPOV, N.A.

TOROPOV, N.A.; GALAKHOV, F.Ya.; BONDAR', I.A.

Equilibrium diagram of the ternary system: BaO - Al₂O₃ - SiO₂.
Izv. AN SSSR Otd. khim. nauk no. 5:753-764 S-0 '54.
(MLRA 8:3)

1. Institut khimii silikatov Akademii nauk SSSR.
(Phase rule and equilibrium)(Aluminum silicates)
(Barium salts)

TOROPOV, N. N.

Investigation of cements with ionization x-ray analysis.
N. A. Toropov, P. F. Korovalov, A. I. Efremov, and G. V.
Afanasyev. Tsvetnoye 20, No. 3, 17-20(1954).—Processes of
dissoc. of limestone, transformations of kaolinite, and for-
mation of dicalcium silicate were investigated with an ion-
ization x-ray installation comprising source of x-rays (x-ray
tube), goniometer, high-temp. furnace, receiver of x-ray
radiation (gas amplifier), electrometric amplifier of d.e., and
recorder of ionization curves. Dissoc. of carbonate oc-
curs somewhat above 500° and is complete at 700°. At
900°, dicalcium silicate changes from the γ -modification into
a new α' -modification. At 500°, kaolinite undergoes struc-
tural changes and becomes amorphous; it remains amor-
phous to 900-1000°, after which mullite starts to form slowly
and then speeds up at 1200°. At 1200°, α -cristobalite forms
from excess amorphous silica. B. Z. Kamich

(3)

TOROPOV N A

Interaction of calcium phosphate with clinker materials
N. A. TOROPOV AND A. I. BORISENKO, *Cement*, 20 [8] 10-14
The study dealt with the high-temperature reactions of tricalcium phosphate with the synthetic clinker materials $2\text{CaO} \cdot \text{SiO}_3$, $3\text{CaO} \cdot \text{SiO}_3$, $1\text{CaO} \cdot \text{Al}_2\text{O}_3 \cdot \text{Fe}_2\text{O}_3$, $1\text{CaO} \cdot 2\text{Al}_2\text{O}_3 \cdot \text{Fe}_2\text{O}_3$, $3\text{CaO} \cdot \text{SiO}_3 + 2\text{CaO} \cdot \text{SiO}_3$, and $2\text{CaO} \cdot \text{SiO}_3 + 3\text{CaO} \cdot \text{SiO}_3 + 4\text{CaO} \cdot \text{Al}_2\text{O}_3 \cdot \text{Fe}_2\text{O}_3$. Incomplete linking of CaO during burning of raw cement mixtures containing P_2O_5 is due to thermal decomposition of tricalcium silicate in the presence of P_2O_5 , with the formation of dicalcium silicate and free CaO . The influence of tricalcium phosphate on a mixture of dicalcium and tricalcium silicates during burning is determined by the proportion of these silicates in the mixture. The permissible content of P_2O_5 increases with decreasing content of tricalcium silicate in the clinker. The content of Fe_2O_3 should be established on the basis of physical tests. The behavior of mixtures of tricalcium silicate with tetracalcium aluminoferrite in the presence of additions of tricalcium phosphate is analogous to the behavior of tricalcium silicate. Tricalcium phosphate does not react with Ca aluminoferrite at 1500°C when it dissolves therein during melting. The P_2O_5 is in the clinker as a component of belite, which dissolves Ca silrophosphates, and in the ferruginous phase, which dissolves up to 28% $3\text{CaO} \cdot \text{P}_2\text{O}_5$ at 1500°C . Change in the appearance of alite due to the presence of P_2O_5 in mixtures is determined by nonuniformity of crystals, indicating the start of decomposition of $3\text{CaO} \cdot \text{SiO}_3$. Use of carbonates containing P_2O_5 should be limited to the production of belite cements.

B.Z.K. ①

TOROPOV, N.A.; ARAKELYAN, O.I.

Crystallization of γ -alumina from melts of the system: NaF -
 AlF_3 - Al_2O_3 . TSvet. met. 27 no.1:57-58 Ja-F '54. (MLRA 10:9)
(Alumina) (Crystallization)

TOROPOV, N. A.

TOROPOV, N.A.; KONOVALOV, P.F.; YEFREMOV, A.I.; ANAN'YEVA, G.V.

Use of the high-temperature X-ray ionization method for studying
processes that take place in alumina production. TSvet.met. 27
no.2:37-42 Mr-Ap '54. (MIRA 10:10)

1. Giprotsement.
(Alumina) (X rays)

"APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001756330003-3

APPROVED FOR RELEASE: 08/31/2001

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APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001756330003-3"

Toropov, N. A.

Investigation of Cr_2O_3 -colored crystals of synthetic ruby.
N. A. Toropov and I. F. Andreev. *Trudy Leningrad. Tekhnol. Inst. im. Lensovetka* 1954, No. 29, 90-5. The absorption spectrum of the ruby crystals is used for a photometric detn. of the amt. of the staining oxide. The transmittance (T) curves which are related to the extinction curves are defined by the relation $D = \log T$ (D is the optical d.), following the Lambert-Beer law. The measurements were made with crystal plates oriented parallel and perpendicular to the optical axis. The characteristic absorption peaks are at 410, and 550 to 560 m μ parallel to the optical axis, and 420 and 500 to 570 perpendicular to the optical axis. The optical d. and the extinction coeffs. of samples perpendicular to the optical axis are considerably higher than those for the orientation parallel to the optical axis. They are linear functions of the concen. in Cr_2O_3 . Also a_0 and c_0 of the unit cell increase proportionally to the Cr_2O_3 content. X-ray powder data are given for leucosapphire and a ruby with 3.2% Cr_2O_3 . W. Eitel

Toropov N.A.

Determination of the content of the staining oxide (Cr_2O_3)
in the crystals of synthetic ruby. N.A. Toropov and I. F.
Serev. "Trudy Leningrad. Tekhn. Inst. im. Lensovet'a"
1954, No. 20, 95-8. The n_{D} and n_{A} are determined by the prism
method for three rubies with 0.13, 0.33, and 2.02% Cr_2O_3 .
The indexes are functions of the concn. of Cr_2O_3 in the syn-
thetic rubies. They are tabulated for six wave lengths of
the visible spectrum. W. Eitel

Toro PoV, N. A.

USSR/Chemistry - Silicates

Card 1/1 Pub. 22 - 24/48

Authors : Toropov, N. A., and Skue, E. R.

Title : Effect of fluoride compounds on solid calcium alumo-ferrite solutions

Periodical : Dok. AN SSSR 98/3, 415-418, Sep 21, 1954

Abstract : The effect of certain fluoride compounds on the stability of solid solutions of $2\text{CaO} \cdot \text{Fe}_2\text{O}_3$, $4\text{CaO} \cdot \text{Al}_2\text{O}_3 \cdot \text{Fe}_2\text{O}_3$ and $6\text{CaO} \cdot 2\text{Al}_2\text{O}_3 \cdot \text{Fe}_2\text{O}_3$, was investigated at high temperatures. It was established that the crystals of the investigated solid calcium alumo-ferrite solutions form a so-called "celite" of Portland cement clinker and also make it possible to determine certain technical properties of the cement. Six references: 2-USA; 2-German; 1-Italian and USSR (1928-1951). Tables; illustrations.

Institution : Academy of Sciences USSR, Institute of Chemistry of Silicates

Presented by: Academician S. I. Vol'fkovich, April 28, 1954

BOTVINKIN, O.K.; YEVSTROP'YEV, K.S., doktor khimicheskikh nauk, professor, retsenzent; TOROPOV, N.A., doktor tekhn.nauk, professor, retsenzent; MAZURIN, O.V., kandidat khim. nauk, retsenzent; KUKOLEV, G.V., doktor tekhnicheskikh nauk, professor, retsenzent; ALKIND, I.Ya., kandidat tekhnicheskikh nauk, redaktor; DEMINA, G.A., redaktor; LYUDKOVSKAYA, N.I., tekhnicheskiy redaktor.

[Physical chemistry of silicates] Fizicheskaya khimiia silikatov. Izd. 2-oe, perer. i dop. Moskva, Gos.izd-vo lit-ry po stroit. materialam, 1955. 285 p.
(MLRA (9:5))

1. Kafedra obshchey tekhnologii silikatov Leningradskogo ordena Trudovo-go Krasnogo Znameni Tekhnologicheskogo instituta imeni Lensoveta (for Yevstrop'yev, Toropov, Mazurin).
(Silicates)

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M. A. KOUTZ

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✓ Structure of Glass—Report of a Symposium on the Structure of Glass, Leningrad, November 23 to 27, 1955. Edited by A. A.

Lebedev, N. A. Torgov, V. P. Barzakovskii, and A. A.

Afipen Akademii Nauk S.S.R., Leningrad-Moscow, 1955.

308 pp.—Upon the invitation of the Institute of Silicate Chemistry of the Academy of Sciences U.S.S.R., the State Optical

Institute, and the Leningrad Section, All-Union Society of Silicate

Technological Research, a symposium on the structure of glass was organized, which was attended by more than 500 representa-

tives from 96 institutions and 28 cities of the U.S.S.R. The introductory address by A. A. Lebedev emphasizes the importance of structural conversions in glass as the basis of many phenomena which cannot be explained otherwise, e.g., the annealing range of optical glass, the luminescence, and the diffraction of X rays, electrons, and neutrons. The crystallites, as assumed in some theories, in the order of magnitude of 10 to 15 a.u. should be detectable through more accomplished electron microscopic methods in the future. The chain structures assumed by others are still hypothetical; their confirmation would be an important approach toward the manufacture of unbreakable glass. The titles of the papers read in the symposium are as follows: K. S. Evstrop'ev: "Crystallite theory of glass structure" (pp. 9-18). P. P. Kobeko: "Structure and properties of organic glasses" (pp. 19-28). O. K. Botvinkin: "Glass structure" (pp. 26-29). E. A. Perel-Koshtik: "Possibilities and results of X-ray methods in the investigation of glassy materials" (pp. 30-43). O. A. Esim and P. V. Gel'd: "Structural

H.H. LEBEDEV

nature of glassy and liquid silicates" (pp. 44-55). E. F. Gross and V. A. Kolesova: "Combination scattering of light and structure of glassy materials" (pp. 50-61). V. V. Taratov: "Quantum theory of heat conductance and structure of silicate glasses" (pp. 62-69). V. A. Florinskaya and R. S. Fechenkina: "Spectra of simple glasses in the infrared and their relations to the structure of glass" (pp. 70-85). A. A. Appen: "Coordination principle in the distribution of ions in silicate glasses" (pp. 96-106). L. I. Demkina: "Ideas on the fine structure of silicate glasses resulting from investigations on the properties of glasses in simple systems" (pp. 107-19). A. I. Stožarov: "Measurement of the thermal expansion of glass as a method for the investigation of its structure" (pp. 120-25). L. G. Mel'nicenko: "Theoretical opinions of D. I. Mendeleev on the structure of silicates and glasses and their importance for the actual science" (pp. 126-35). V. P. Barzakovskii: "Ideas of D. I. Mendeleev on the chemical character of silicates" (pp. 136-40). O. S. Molchanova: "Properties of glasses in the ternary system $\text{Na}_2\text{O}-\text{B}_2\text{O}_3-\text{SiO}_2$ " (pp. 141-44). E. A. Porai-Koshits: "Structure of Na borosilicate glasses" (pp. 145-61). S. P. Zhdanov: "Structure of glass as seen from the results of the investigation of the structure of porous glasses and films" (pp. 162-75). D. P. Dobychin: "State of SiO_4 in microporous glass" (pp. 170-80). S. I. Dubrovu: "Corrosion of glassy silicates and of Na aluminosilicates by aqueous solutions in its relation to the state of SiO_4 in glass" (pp. 181-84). A. F. Zuk: "Existence of distinct chemical compounds in the glass structure" (pp. 185-80). Yu. A. Gastev: "Chemical stability of glass" (pp. 187-89). N. A. Tudorovskaya: "Structural variabilities of the light refractive index of glass at temperatures below 300°C." (pp. 190-97). D. I. Levin: "Rayleigh scattering in glasses and the glass structure" (pp. 198-201). M. M. Gurevich: "Spectral relation of

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light scattering in Na borosilicate glasses" (pp. 202-206). A. N. Sevchenko: "Application of the luminescence method for the investigation of the glassy state" (pp. 207-15). G. O. Bagdyk'yants: "The problem of an oriented structure of glass" (pp. 216-18). V. I. Shelyubskii: "Application of the electron microscope to the investigation of glass" (pp. 219-23). L. A. Afanas'ev: "Experiments on the electronographic study of industrial glasses" (pp. 224-26). A. I. Avgustinik: "Some properties of highly aluminous glasses" (pp. 227-29). N. V. Solomin: "Chemical compounds in borate glasses" (pp. 230-31). G. A. Kolykov: "Selective volatility of components of the system $\text{Na}_2\text{O}-\text{B}_2\text{O}_3-\text{SiO}_2$, a method for the investigation of the nature of the glassy state" (pp. 234-44). A. G. Bergman: "Visual-polythermic method for the investigation of crystallization in glasses and silicate systems" (pp. 245-47). V. A. Kozhevnikov: "Phenomenon of limited miscibility in binary silicate melts" (pp. 248-50). V. T. Slavyanskii: "Temperature function of viscosity and structure in some glassy and liquid materials" (pp. 251-55). M. M. Skornyakov: "Viscosities of glasses above and below the liquidus temperature" (pp. 256-57). V. A. Ioffe: "Dielectric losses in silicate glasses" (pp. 258-63). B. I. Markin: "Electric conductance of simple borate systems in the glassy state" (pp. 264-66). V. A. Presnov: "I, Electric

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conductance of glasses in strong electric fields; II. The wetting of metals by glass" (pp. 207-60). V. P. Pryanishnikov: "Electric conductance of silica glass" (pp. 270-72). E. P. Azarov: "Structure of enamels and their properties" (pp. 273-75). A. G. Repa: "Oxygen potential of glass" (pp. 276-79). L. V. Sergeyev: "The glassy state of organic polymers" (pp. 280-82). Yu. N. Andreyev: "Problems of the methodological basis of the actual ideas on the structure of glass" (pp. 283-89). The ample discussions (pp. 293 to 302) include the following main items: general remarks on the nature of glass; physical chemistry of polycomponent systems and the glass structure; optical properties and glass structure; caloric and electrical properties and the structure of glass; crystallochemistry and glass structure; and problems of further development of glass science. The concluding address of A. A. Lebedev (pp. 360-62), and the official resolution of the Symposium Meeting (pp. 303-05) announce plans for another Symposium on the Structure of Liquids and, in 1956, the third Symposium on the Structure of Glass, under the auspices of the Academy of Sciences U.S.S.R. The present volume is excellently printed and illustrated; it is a real milestone in the evolution of modern investigations on glass structure.

W. EITZ

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"APPROVED FOR RELEASE: 08/31/2001

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CIA-RDP86-00513R001756330003-3"

Toropov, N. A.

USSR/ Chemistry - Silicates

Card 1/1 Pub. 40 - 1/27

Authors : Toropov, N. A.; Galekhov, F. Ya.; and Bondar', I. A.

Title : Solid solutions formed by celsian, dibaum trisilicate and barium disilicate (Sanbornite)

Periodical : Izv. AN SSSR. Otd. khim. nauk 1, 3-8, Jan-Feb 1955

Abstract : Experiments were conducted to establish the zone, boundaries and liquidus of a ternary solid solution formed by barium disilicate, dibaum trisilicate and celsian. It was found that the refraction index for this zone depends largely upon the barium disilicate and aluminum oxide contents of the solution. The refraction index decreases with the increase of barium disilicate and Al_2O_3 . The equilibrium ratio of the investigated solution was established on the basis of several polythermal samples with constant Al_2O_3 contents. Two USA references (1922 and 1950). Graphs; table; illustrations.

Institution : Acad. of Sc., USSR, Institute of Chem. of Silicates

Submitted : January 28, 1954

TOROPOV, N.A., professor; AVGUSTINIK, A.I., professor; BARZAKOVSKIY, V.P.,
doktor khimicheskikh nauk

Scientific research conducted in Czechoslovakia on the technology
of silicates. Stek. i ker. 12 no.10:12-13 O '55. (MLRA 9:1)
(Silicates) (Czechoslovakia--Research)

Toropov N.A.

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Influence of fluorine salts on tricalcium aluminate at high temperatures. N. A. TOROPOV, B. V. VOLKONSKIY, AND V. I. SAPKOV. *Tsvetnaya Metallurgiya*, 12-13 (1955).—Roentgen-ionization analysis was used to determine the effect of 5% K, Na, and Ca fluoride on tricalcium aluminate at temperatures up to 1600°C. Sodium and K fluorides begin to exert their influence and cause decomposition of tricalcium aluminate at 800° into pentacalcium aluminate and free CaO. Above 1300°, the influence of these salts ceases. The action of CaF₂ is similar to that of NaF and KF, but tricalcium aluminate decomposes at $t \approx 1000^\circ$.
B.Z.K.

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R.M. ✓

Subject : USSR/Chemistry AID P - 1371
Card 1/1 Pub. 119 - 4/6
Authors : Toropov, N. A. and Bondar', I. A., (Leningrad)
Title : Fluoberyllates and other crystallochemical analogs of silicates and like substances
Periodical : Usp. khim., 23, no. 1, 52-68, 1955
Abstract : A survey of the literature on fluoberyllates is given; most of the references are to non-Russian sources. A high degree of analogy is found between BeF₂ and SiO₂. Many binary systems are reviewed. Twenty diagrams, 2 tables, 37 references (7 Russian: 1939-53).
Institution : None
Submitted : No date

"APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001756330003-3

APPROVED FOR RELEASE: 08/31/2001

CIA-RDP86-00513R001756330003-3"

TOKO PUV, N. H.

AID P - 2290

Subject : USSR/Chemistry

Card 1/1 Pub. 152 - 15/21

Authors : Toropov, N. A. and M. M. Sychev

Title : Study of the temperature resistance of mineral wool fibers

Periodical: Zhur. prikl. khim., 28, no.3, 322-325, 1955

Abstract : A method for determination of devitrification with the use of Kurnakov's differential pyrometer is given. Increase in the Fe₂O₃-content lowers and increase in Al₂O₃-content increases the resistance of mineral wool fibers to high temperature. Two tables, 1 diagram, no references.

Institution: None

Submitted : S 6, 1953

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TOROPOV, N.A.

BOYKOVA, A.I. [translator]; BONDAR', A.I. [translator]; VOANO, V.G. [translator]; YEGOROVA, Ye.N. [translator]; MIKOGOSYAN, Kh.S. [translator]; TOROPOV, N.A., professor, redaktor; ZAKHAR'YEVSKIY, V.A., redaktor; OGANDZHANOVA, N.A., redaktor; DUMBRE, I.Ya., tekhnicheskiy redaktor

[Physical chemistry of silicates; a collection of articles. Translated from the English and German] Fizicheskaya khimiia silikatov; sbornik statei. Perevod s angliiskogo i nemetskogo A.I.Boikovo i dr. Pod red. N.A.Toropova. Moskva, Izd-vo inostrannoi lit-ry, 1956. 302 p. (MIRA 9:7) (Silicates)

YEVSTROP'YEV, Konstantin Sergeyevich, professor, doktor khimicheskikh nauk;
TOROPOV, Nikita Aleksandrovich, professor, doktor tekhnicheskikh
nauk; GURBICH, E.A., redaktor; GIADKIKH, N.N., tekhnicheskiy
redaktor

[The chemistry of silicon and the physical chemistry of silicates]
Khimiia kremniia i fizicheskaya khimiia silikator. Izd. 2-oe.
Moskva, Gos. izd-vo lit-ry po stroit. materialam, 1956. 339 p.
(Silicon) (Silicates) (MLRA 10:3)

TOROPOV, N. A.

Dr. Tech. Sci.

"Latest Data on the Phase Diagram of Aluminum Oxide -- Silicon Dioxide and on
the Behavior of Aluminous Refractories in Glass-Making Furnaces," a paper given at the
4th International Congress on Glass, Paris, 2-7 Jul 1956

Sinn. 1274

15-57-7-9444

Translation from: Referativnyy zhurnal, Geologiya, 1957, Nr 7,
pp 103-104 (USSR)

AUTHORS: Toropov, N. A., Bondar', I. A.

TITLE: Synthesis of a Fluoberyllate Type of the Double Calcium
and Barium Metasilicate (Sintez ftoroberillatnoy modeli
dvoynogo metasilikata kal'tsiya i bariya)

PERIODICAL: Sb. nauch. rabot po khimi i tekhnol. silikatov, Moscow,
Promstroyizdat, 1956, pp 20-23.

ABSTRACT: The compound $2\text{NaF}\cdot\text{KF}\cdot3\text{BeF}_2$, the fluoberyllate analogue
of the double calcium and barium metasilicate $2\text{CaO}\cdot\text{BaO}\cdot$
 3SiO_2 , has been synthesized. The following were used
in the synthesis: 1) sodium fluoride (98 percent NaF),
2) potassium fluoride in the form of $\text{KF}\cdot\text{H}_2\text{O}$ (77 percent
KF), and 3) beryllium fluoride, obtained by treating
beryllium oxide with hydrofluoric acid (97.5 percent
 BeF_3). The fusion was made in a covered platinum cruci-
ble in a crucible furnace and then quenched (the
melt poured out into a pan). To compare the compound

Card 1/4