

AUTHORS: Vorozhtsov N.N., Corresponding Member ^{SOV/68-58-11-14/25} of the Academy of Science of the USSR, Doctor of Chemical Science, Lisitsyn V.N., Candidate of Chemical Science, Agafonov A.V. and Krasivichev V.V., Candidates of Technical Science, and Abayeva B.T., Candidate of Chemical Science

TITLE: Transformation of Higher Homologues of Phenol into Lower Ones (Prevrashcheniye vysshikh gomologov fenola v nizshiye)

PERIODICAL: Koks i Khimiya, 1958, Nr 11, pp 42-47 (USSR)

ABSTRACT: The results of an investigation on the dealkylation of technical xylenol with simultaneous alkylation of benzole in a pilot plant of the All-Union Scientific Research Institute of the Petroleum Industry in which bead aluminosilicate was used are described. This was a continuation of the previously published work (Ref 1) on the transformation of xylenols (on interaction with benzole) into phenols and cresols on cracking under mild conditions on an aluminosilicate catalyst. The experimental plant used (Fig 1) is outlined. It was established that, on passing xylenol in mixture with benzole

Card 1/3

SOV/68-58-11-14/25

Transformation of Higher Homologues of Phenol into Lower Ones

x (1 : 3.65 by weight) over aluminosilicate catalyst at temperatures in the range 300-400°C and volume velocities of 0.42-1.47hr⁻¹, up to 60% (on weight of starting xylenol) of phenolic compounds (phenol, o-, m- and p-cresols, xylenols) including 20-22% of phenolic-cresolic fraction, are obtained. Simultaneously 11-19% of benzene homologues with a boiling temperature of 100-185°C and 13-18% of neutral compounds with boiling temperatures above 185°C are formed. 8-25% of coke is deposited on the catalyst. The influence of the temperature of the reaction, the volume velocity of reactants (Table 1), additions of water vapour and various proportions of benzole (Table 2) on the transformation of xylenol and changes in the activity of the catalyst with time of operation (Table 3) were established. It was found that at temperatures 300-320°C and volume velocities 0.92-1.47hr⁻¹ more phenolic-cresolic fraction and less of neutral compounds and coke on the catalyst is obtained (taking into consideration the transformation of xylenol). At 300°C and a volume velocity 0.92hr⁻¹ 330kg of

Card 2/3

30V/68-58-11-14/25

Transformation of Higher Homologues of Phenol into Lower Ones

phenolic-cresolic fraction and about 200kg of benzene homologues with a boiling temperature 100-185°C can be obtained from 1 ton of xylenol.

There are 3 tables, 3 figures and 6 references (4 Soviet, 1 English and 1 German)

ASSOCIATION: MGNTI im. D.I. Mendeleeva, VNII NP

Card 3/3

AGAFONOV, A. V., ZHERDEVA, L. G., KARASEVA, A. A., VOZNESENSKAYA, E. V.,
ALTSHULER, A. E., KROL, B. B., OROCHKO, D. I., AKIMOV, V. S., MIKHAYLOV, B. B.,
DRUZHININA, A. V.

"Production of Lubricating Oils and Paraffin from Sulfurous Oils
in the USSR."

Report submitted at the Fifth World Petroleum Congress, 30 May -
5 June 1959. New York City.

AGAFONOV, A. V., TOPCHIYEVA, K. V., KALIKO, M. A., FIGUZOVA, L. I.
AGAFONOV, A. V., PANCHENKOV, G. M., KAMAKIN, N. M., MIRSKIY, Y. S.

"Studying the Nature of Activity of Aluminosilicate Catalysts."

Report submitted at the Fifth World Petroleum Congress, 30 May -
5 June 1959. New York.

SOV/65-59-4-4/14

AUTHORS: Agafonov, A.V., Abayeva, B.T., Andreyeva, A.S.,
Eygenon, A.S., Kantor, I.I. and Ivchenko, Ye.G.

TITLE: Catalytic Cracking of Crude and Hydro-Purified Vacuum Gas-Oil from Arlan. Petroleum (Kataliticheskiy kreking iskhodnogo i gidroochishchennogo vakuumnogo gazoylya arlanskoy nefti)

PERIODICAL: Khimiya i tekhnologiya topliv i masel, 1959, Nr 4, pp 18-24 (USSR)

ABSTRACT: Vacuum gas-oil from Arlan. petroleum contains 3.2% sulphur compounds, 0.11% nitrogen compounds and 24% tarry substances; these quantities are larger than the corresponding quantities in heavy gas-oil from Tatariya and Bashkiriya petroleums. These components block the active surface of the catalyst during cracking, prevent the access of hydro-carbon molecules and therefore decrease the degree of conversion of the crude material. Considerable amounts of coke are deposited on the catalyst which inhibits secondary reactions and leads to decreased yields and inferior quality end-products. Hydro-purification was carried out on a continuous apparatus in the VNII NP by

Card 1/5

SOV/65-59-4-4/14

Catalytic Cracking of Crude and Hydro-Purified Vacuum Gas-Oil from
Arlan. Petroleum

N.A.Chepurov and R.N.Yudinon; a stationary aluminium-cobalt-molybdenum catalyst was used at 380°C, a pressure of 50 atm and space velocity of the supplied crude material of 0.7 hour⁻¹. The properties of the starting material and of the hydro-purified vacuum gas-oil are tabulated (table 1). The octane number of the end product was appreciably higher than when using fractional distillation (58.5 as compared to 41.0) and contained considerably less sulphur (0.013 as against 0.17%). The properties of the gas-oil fractions are listed in table 2. Cracking experiments of both the crude and hydro-purified vacuum gas-oil were carried out on a pilot plant with a synthetic bead catalyst at temperatures within the limits of 430 to 520°C, atmospheric pressure and a space velocity of 0.65 to 1.5, calculated on the volume of the catalyst per hour. The ratio of the catalyst to the crude material was constant in all experiments and equalled 5:1 (table 3). Optimum

Card 2/5

SOV/65-59-4-4/14

Catalytic Cracking of Crude and Hydro-Purified Vacuum Gas-Oil from
Arlan Petroleum

yields of petrol were obtained at temperatures between 450 and 475°C when the optimum space velocity of the supplied raw material was within the limits of 1.0 to 0.65 hours⁻¹. The hydro-purified vacuum gas-oil could more easily be processed; an optimum yield of light components at the same space velocities was achieved at 50°C. The authors concluded that the presence of a considerable quantity of light fractions boiling up to 350°C (37.6 as against 19.4%) influences the yield of the light components. The optimum yield at this temperature reached 66 to 67% by weight as against 58 to 59%. Results of the cracking experiments indicate (Fig 1) that the hydro-purification of the crude (by separating the tarry substances, metals, sulphur and nitrogen) improves the process conditions and also the yields and properties of the cracking products (compare table 4). The gasoline obtained by this process is less unsaturated, contains more aromatic compounds and has higher octane numbers (80 to 81.5 as compared to

Card 3/5

SOV/65-59-4-4/14

Catalytic Cracking of Crude and Hydro-Purified Vacuum Gas-Oil from Arlan Petroleum

77.7 to 80.7) (Fig 2). A lower content of unsaturated compounds renders the gasoline more stable. Its induction period exceeds 600 minutes. The light catalytic gas-oils, obtained during the cracking of hydro-purified crudes, show improved properties. Their cetane number is 34 to 38 (as against 30 to 33) and they contain 0.21 to 0.38% sulphur (as against 2.6 to 3.3%) (Fig 3). These light gas-oils can be used directly as components of diesel fuels. The heavy catalytic gas-oils (fractions boiling above 350°C) can be used for the production of lubricating oils or re-used as recycles. In both cases 2 to 3% of the tarry (tail) fractions have to be separated. The gaseous hydrocarbons produced by this process are of interest as starting materials for petro-chemical syntheses. The influence of the temperature on the ratio of unsaturated and saturated hydrocarbons in gaseous reaction products, and on the

Card 4/5

SOV/65-59-4-4/14

Catalytic Cracking of Crude and Hydro-Purified Vacuum Gas-Oil from
Arlan: Petroleum

content of unsaturated hydrocarbons in the gas, is
shown in a graph (Fig 4). There are 4 figures,
4 tables and 2 English references.

Card 5/5

SOV/65-59-4-5/14

AUTHORS: Agafonov, A.V., Basov, A.N., Manakov, N.Kh. and
Manshillin, V.V.

TITLE: Combined Plant for Fractional Distillation of Petroleum
and of Catalytic Cracking Residues on a Microspherical
Natural Catalyst (Kombinirovannaya ustanovka pryamoy
peregonki nefti i kataliticheskogo krekinga ostatochnogo
syr'ya na mikrosfericheskom prirodnom katalizatore)

PERIODICAL: Khimiya i tekhnologiya topliv i masel, 1959, Nr 4,
pp 25-31 (USSR)

ABSTRACT: Petroleum refineries have to process asphalt-tar
substances of petroleum which can be extremely difficult.
Processing methods hitherto applied use high temperatures
(above 450°C) at high or low pressures. A high yield of
tarry residues and poor quality gasoline or distillate
fractions and petrols of low quality and also hard
residues in the form of petroleum coke are obtained by
thermo-cracking. The temperature is an important factor
during thermal destructive processes. It has been found
that temperatures should be selected to give fractions
with octane numbers exceeding 70 and that the cetane

Card 1/4

SOV/65-59-4-5/14

Combined Plant for Fractional Distillation of Petroleum and of
Catalytic Cracking Residues on a Microspherical Natural Catalyst

number of the diesel fuel fraction should not exceed 42 to 43. The VNII NP have developed an economical catalytic destructive process for the treatment of residual petroleum crudes which makes it possible to obtain high grade gasoline and diesel fuels in industrial quantities. The process was tested under laboratory, pilot plant and industrial conditions. The VNII NP is, in collaboration with the Giproneftezavod Institute, at present designing two plants where the simultaneous fractional distillation and catalytic cracking of the petroleum crude can be carried out, one with an annual capacity of 2 million tons and a second of 3 million tons. The lay-out of both factories will be the same as is shown in Fig 1. The asphalt-tar substances will be subjected to the direct action of aluminium silicate catalysts which will be sufficiently active to ensure decomposition of the high molecular petroleum fractions (boiling above 530 to 550°C). The light gas-oil fractions of the petroleum will not be decomposed and the cetane number of the diesel fuel

Card 2/4

SOV/65-59-4-5/14

Combined Plant for Fractional Distillation of Petroleum and of
Catalytic Cracking Residues on a Microspherical Natural Catalyst

fraction, obtained during the process, should be 42 to 43 or higher. The newly-formed fraction of the gasoline should have an octane number of 76 to 78 and above. The crude petroleum or fuel oil can be directly supplied into the reactor. Various further improvements in the process are described. The percentage composition of the end product obtained on a natural microspherical catalyst in an industrial plant is given, as well as experimental data, obtained by VNII NP during 1958, on fuel oil subjected to catalytic cracking on a pilot plant. The coke deposited on the catalyst can be separated by roasting at a temperature of about 600°C; the importance of the catalyst is discussed. By using pneumatic transport for the catalyst in a highly concentrated current it is possible to decrease the height of the plant and, therefore, to lower construction costs. The regeneration of the catalyst is intensified. The considerable enlargement of the desorption zone in the reactor, and also the creation of a counter-current

Card 3/4

SOV/65-59-4-5/14

Combined Plant for Fractional Distillation of Petroleum and of Catalytic Cracking Residues on a Microspherical Natural Catalyst

desorption zone in the regenerator for degasification and activation of the regenerated catalyst, decreases coke-formation and the yield of methane, gives higher grade gasoline and simplifies the further separation of cracking gases. Practically all the heat, generated by burning the coke and other component gases, is utilised. These vapours are used as power and also for desorption or for heating. The plant is also equipped for utilising the effluents. Comparative technical and economical characteristics are listed in a table. The authors also refer to a relevant article by Sherwood which was published in "Petroleum", 1959, Nr 2. There are 2 figures, 1 table and 1 English reference.

Card 4/4

SOV/65-59-4-7/14

AUTHORS: Agafonov, A.V., Soskind, D.M. and Abayeva, B.T.

TITLE: The Operation and Methods of Reconstruction of Catalytic Cracking Plants Where Bead Catalysts are Used (Opyt ekspluatatsii i puti rekonstruktsii ustanovok kataliticheskogo krekinga s sharikovym katalizatorom)

PERIODICAL: Khimiya i tekhnologiya topliv i masel, 1959, Nr 4, pp 34-44 (USSR)

ABSTRACT: Investigations were carried out in the VNII NP which made it possible to work out conditions and make recommendations for the manufacture of high quality petrols. Heavy distillates, boiling at temperatures between 300 and 500°C, can be used. The process is carried out in one stage and therefore, the efficiency of the plant increased by 30 to 35%. The properties of the gas-oil fractions of Romashkaya petroleum are given in table 1. It can be seen that the heavy crudes differ from the kerosine-gas-oil fractions by their high boiling and solidification points, by their high content of tar, sulphur and aromatic compounds as well as by appreciable content of polycyclic aromatic compounds,

Card 1/3

SOV/65-59-4-7/14

The Operation and Methods of Reconstruction of Catalytic Cracking
Plants Where Bead Catalysts are Used

asphaltenes and metal salts. Various investigations carried out in the Novoufimka plant during 1954 to 1955, and modifications of the plant carried out at the time, are discussed in detail. The reconstructions, carried out at present, aim to increase the efficiency of the plant 1.5-fold (first modification) and 1.7-fold (second modification) without altering the principal layout of the plant. A further reconstruction is to achieve a considerable improvement in the conversion process which will increase the efficiency of the plant by 100% (third modification). The first modification is based on recommendations made by the authors, the Novoufimka factory Giproneftemash and Giproneftezavod. This type of reconstruction was carried out on one plant of the NUNPZ and three plants of the Salavatskiy factory. The various modifications are listed in a table on page 40 and the most important of these discussed in detail. Table 2 shows the improvements achieved during 1956 to 1957 and the first nine months of 1958 in various plants where the recommended

Card 2/3

SOV/65-59-4-7/14

The Operation and Methods of Reconstruction of Catalytic Cracking
Plants Where Bead Catalysts are Used

reconstructions have been carried out. The second modification was recommended by GrozNII and Giproneftezavod and the third by VNII NP and Giproneftemash. The lay-out of the last plant is given in Fig 3 and the authors suggest that this last modification should only be incorporated in newly-erected plants. There are, however, various drawbacks e.g. the circulation time of the catalyst is rather low, the generator is not completely efficient, the cooling pipes of the regenerator are unsatisfactory and this leads to an increased catalyst consumption. There are 3 figures and 2 tables.

Card 3/3

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S/081/63/000/002/069/088
B160/B144

11.0140
AUTHORS: Osipov, L. N., Gol'dshteyn, D. .., Agafonov, A. V.

TITLE: Hydrofining of diesel fuels

PERIODICAL: Referativnyy zhurnal. Khimiya, no. 2, 1963, 461, abstract
2P128 (Tr. Vses. n.-i. in-t po pererabotke nefti i gaza i
polucheniya iskusstv. zhidk. topliva, no. 8, 1959, 54 - 73)

TEXT: The process of hydrofining sulfurous straight-run distillates and secondary distillates was studied in laboratory high-pressure circulation equipment with an industrial Al-Co-Mo catalyst. The rate of hydrating the S-compounds and unsaturated hydrocarbons at the given partial H₂ pressure is shown to increase as the temperature rises to 420°C; at a higher temperature of the order of 460°C the rate of hydration decreases. The optimum partial H₂ pressure in the hydrofining of diesel-fuel distillates depends on the chemical composition of the crude. Hydrofining of low-aromatic distillates can be carried out at a comparatively low partial H₂ pressure (15 - 20 atm.) and hydrofining of aromatized distillates (e.g. catalytic-
Card 1/2

Hydrofining of diesel fuels

S/081/63/000/002/069/088
B160/B144

cracking distillate) at a high partial H_2 pressure. In the process of hydrofining low-aromatic distillates at a comparatively low partial H_2 pressure, the amount of H_2 consumed in the reaction decreases as the temperature of the processes rises ($> 380^\circ C$) as a result of dehydration of the naphthene hydrocarbons contained in the crude. Certain gases containing H_2 , which are produced in a number of processes (e.g. the circulating waste gas of the hydroforming process) can be used for hydrofining straight-run and other low-aromatic distillates. 16 references. [Abstracter's note: Complete translation.]

Card 2/2

AGAFONOV, A-V

PHASE I BOOK EXPLOITATION

SOV/4659

Osnovy tekhnologii neftekhimicheskogo sinteza (Fundamentals of Synthesis Technology in Petroleum Chemistry) Moscow, Gostoptekhizdat, 1960. 852 p. 3,800 copies printed.

Eds.: Dintsés, Arkadiy Il'ich, Professor, and Lev Aleksandróvich Potolovskiy, Professor; Executive Ed.: L.A. L'vova; Tech. Ed.: E.A. Mukhina.

PURPOSE: This book is intended for engineers and chemists of petroleum refineries and chemical plants, for councils of the national economy, planning organizations and scientific research institutes engaged in chemical processing and large-scale utilization of petroleum stock for the production of synthetic products.

COVERAGE: The book describes important commercial methods of producing hydrocarbon petroleum and gas stock and coal stock for the manufacture of alcohols, aldehydes, ketones, acids, detergents, synthetic fibers, and synthetic rubber. Flow sheets are included, and the basic equipment of the petrochemical industry is described. The physicochemical properties and use of intermediate and end synthetic products are also described. The state of the petrochemical industry outside the USSR and prospects for its development are covered. No personalities are mentioned. References follow each chapter.

Card 1721

Fundamentals of Synthesis Technology (Cont.)

SOV/4659

TABLE OF CONTENTS:

| | |
|---|----|
| Foreword | 3 |
| Ch. I. Development of Petroleum Refining and Petrochemical Processing | 7 |
| I. From straight-run distillation of petroleum to the petrochemical industry (raw materials for the manufacture of petrochemicals [A.I. Dintses]) | 7 |
| II. Petrochemical processing industry abroad [L.A. Potolovskiy] | 15 |
| Ch. II. Processes of Raw Material Production for the Petrochemical Industry [B.T. Abayeva and A.V. Agafonov] | 40 |
| I. Destructive distillation of petroleum, the main source of raw materials for petrochemical processing | 40 |
| II. Thermal cracking | 41 |
| III. Pyrolysis of hydrocarbon gases and liquid petroleum products | 47 |
| 1. Chemical nature of the pyrolysis process | 48 |
| 2. Pyrolysis of liquid raw materials | 51 |
| 3. Pyrolysis of gaseous raw materials | 60 |
| 4. Tube heaters for the pyrolysis process | 69 |
| IV. Cooking of petroleum residue | 71 |
| V. Catalytic cracking of petroleum products | 76 |

Card 2/21

AGAFONOV, A.V.; ABAYEVA, B.T.; OKINSHEVICH, N.A.

Distribution of sulfur in the cracking products of heavy charge
stocks. Khim.sera-i azotorg.soed.sod.v نفت. i نفتprod. 3:183-192
'60. (MIRA 14:6)

1. Vsesoyuznyy nauchno-issledovatel'skiy institut po pererabotke
nefti i gaza i pplucheniyu iskusstvennogo zhidkogo topliva.
(Petroleum products) (Sulfur organic compounds)

GOL'DSHTEYN, D.L.; OSIPOV, L.N.; AGAFONOV, A.V.

Selective hydrofining of catalytically cracked gasolines. *Khim. sera i azotorg. soed. v neft. i nefteprod.* 3:389-395 '60. (MIRA 14:6)

1. Vsesoyuznyy nauchno-issledovatel'skiy institut po pererabotke nefti i gaza i polucheniyu iskusstvennogo zhidkogo topliva.
(Gasoline) (Cracking process)

S/065/60/000/006/006/008/XX
E194/E484

AUTHORS: Agafonov, A.V., Gel'ms, I.E. and Rabikovich, E.I.
TITLE: The Selection of Catalyst for Cracking Residual
Petroleum Fractions and Study of its Poisoning During
the Process

PERIODICAL: Khimiya i tekhnologiya topliv i masel, 1960, No.6,
pp.6-12

TEXT: A special feature of the operating conditions of catalyst when cracking residual feed is the high rate of poisoning of the catalyst by the combined influence of temperature, steam, sulphur compounds and resinous substances containing metallo-organic compounds. The poisoning is specially marked with catalyst in powder form. The high molecular hydro-carbons of the heavy part of the feed are the main source of products that are of low stability at temperatures used for cracking so that there is no need for a great reduction in the energy of activation. Moreover, the use for this purpose of high activity catalysts is accompanied by considerable increase in gas and coke formation. The measure of the necessary activity of catalyst used in cracking residual feed should be the production of gasoline of good engine properties combined with
Card 1/4

S/065/60/000/006/006/008/XX
E194/E484

The Selection of Catalyst for Cracking Residual Petroleum Fractions and Study of its Poisoning During the Process

favourable gas composition and a high rate of conversion of the residual fractions. It is desirable to use large plants for cracking residual fractions which requires large quantities of catalyst of appropriate quality. It is found that the requisite conditions are satisfied by fire-resistant high-alumina kaolin clays, many kinds of which after simple heat treatment have sufficiently good and stable catalytic properties. Table 1 gives the characteristics of semi-industrial quantities of natural catalysts obtained from various natural clays. In this table the activity of the catalysts is characterized by the cracking of light feed as this gives the more sensitive index of performance. Catalyst poisoning is then considered and Table 2 gives experimental data about the deactivation of a simple, natural, microspherical catalyst. Characteristics of the cracker feed-stocks are given in Table 3. The tests, results of which are given in Table 2, were continued for 20 days and the observed changes in catalyst properties may be considered as the results of poisoning by metals and sulphur combined with poisoning due to reduction in the surface and porosity.

Card 2/4

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S/065/60/000/006/006/008/XX
E194/E484

The Selection of Catalyst for Cracking Residual Petroleum Fractions and Study of its Poisoning During the Process

caused by heat and steam. The selectivity of the catalyst was much reduced. Comparison between the test results and those of laboratory tests with artificial poisoning of the catalyst by metal showed that in the tests most of the poisoning was due to metals and not sulphur, see Table 4. The influence of metals deposited on the catalysts on certain characteristics of cracking of distillate and crude oil feeds are given in Table 5. Note should be made of the much smaller degree of poisoning of the catalyst by an equal quantity of deposited metal when cracking residual feed rather than distillate. The influence of steam was studied in the laboratory and the results are given in Table 6; it will be seen that treatment of poisoned catalyst by steam promotes recovery of activity and improves the selectiveness. It is concluded from the work that catalytic cracking of residual feed stock on micro-spherical natural catalyst is practical and a stable process can be achieved. The capital cost of constructing a catalyst manufacturing works should not be above 500 roubles per ton of catalyst produced per year. Investment in quarries and other
Card 3/4

S/065/60/000/006/006/008/XX
E194/E484

The Selection of Catalyst for Cracking Residual Petroleum Fractions
and Study of its Poisoning During the Process

workings is from 100 to 200 roubles per ton per year. The cost of one ton of finished catalyst is about 400 roubles. It is calculated that the use of natural catalyst rather than synthetic economizes both capital investment in the production of catalyst and the cost of the first charge by not less than 15 roubles for each ton per year of feed stock delivered for catalytic crackings. Accordingly, the economy that results from the use of natural rather than synthetic catalyst for cracking residual feed stock is considerable. There are 6 tables and 8 references: 2 Soviet and 6 English.

ASSOCIATION: VNII NP

Card 4/4

AG-AFONOV, A. V.

B/065/61/000/004/003/011
E194/EEB4

BoGov, B. P., Danilevich, A. F., Gol'dsheyn, D. L.,
Kysakov, N. V. and Agafonov, A. V.

Hydrofining of Lubricating Oils

PERIODICAL: Khimiya i tekhnologiya topliv i masei, 1961, No. 4,
pp. 25-27

TEXT: Hydrofining is under consideration as a replacement for earth treating in finishing of solvent raffinate. This article describes tests on the hydrofining of distillate (spindle oil) and machine oil Type AC-5 (AS-5) and residual de-waxed phenol fractions of the Kovokhyshevsk Niz. The hydrofining was carried out on a large laboratory pilot plant with gas circulation. This work was carried out in a laboratory with steam stripping. It was concluded that the pressure of 40 atmospheres, the highest tried, was the best in respect of improving the viscosity index, reducing the coke number and sulphur content and improving the colour of the finished oils. The ratio of volumes of oil per hour to volume of catalyst ranged from 1 to 4. The influence of treatment temperature was then studied using

Card 1/5

on the one hand an aluminium-cobalt-molybdenum catalyst and on the other an aluminium-molybdenum catalyst. These tests were made with machine oil Type AC-5 at a total pressure of 40 atm and a delivery rate by volume relative to catalyst of 3 l/hours and a circulation rate of 275, 300, 325 and 350°C. It was shown that increasing the temperature has such the same effect as decreasing the feed rate. As a rule increasing the temperature somewhat increase the pour point which rose from -18°C with a treatment of hydrofined spindle oil to -15°C with a treatment of hydrofined spindle oil. Tables are then given of the characteristics of hydrofined spindle oils under optimal conditions. Table 3 was obtained with an aluminium-molybdenum catalyst and Table 4 with aluminium-cobalt-molybdenum catalyst.

B/065/61/000/004/003/011
E194/EEB4

Hydrofining of Lubricating Oils

Table 3

| | Treated Oil | |
|---|-------------|-------|
| | 200° | 225° |
| Viscosity centistokes: | 19.03 | 18.25 |
| at 100°C | 4.87 | 4.80 |
| at 150°C | 92.3 | 95.7 |
| Viscosity Index | -14 | 112 |
| Pour point °C | 190 | 198 |
| Flash point °C | 2.5 | 1.5 |
| Colour, HFA | 0.96 | 0.86 |
| Sulphur content % weight | 0.03 | 0.01 |
| Coke % weight | 0.02 | 0.01 |
| Conductivity Pinkovich gms/m ² | 6.65 | 2.13 |
| Yield % weight | 100.0 | 99.1 |

Card 3/5

8/065/61/000/004/003/011
E194/0228A

Hydrofining of Lubricating Oils

Table A

| | Feed | Treated Oil |
|--------------------------|--------|-------------|
| Viscosity centistokes: | | |
| at 50°C | 159.35 | 155.87 |
| at 100°C | 20.98 | 20.80 |
| Viscosity index | 85.1 | 88.4 |
| Pour point °C | -10 | -8 |
| Flash point °C | 246 | 270 |
| Colour NEA | 6.5 | 5 |
| Sulphur content % weight | 1.03 | 0.27 |
| Coke No. % weight | 0.38 | 0.27 |
| Yield % weight | 100 | 99.1 |

The hydrogen consumption in treating the distillate oil was 0.13% weight and in treating the residual oil 0.15% weight. The results of hydrofining and earth finishing are then compared and it is Card 4/5

shown that hydrofinishing gave the greater yield, about 2% on distillates and 4% on residual lubricants. The hydrofined oils have lower coke number but there is some loss in the viscosity and a slight increase in the pour point. Hydrofining has little influence on the chemical composition of the lubricants. The increase in viscosity index on hydrofining mainly results from newly formed paraffinic, naphthenic and light aromatic hydrocarbons. Preliminary technical and economic calculations show that hydrofinishing of lubricants is promising as a replacement for earth treatment. There is not much to choose between the performance of the two catalysts tested but the aluminum-molybdenum catalyst is cheaper. Full scale tests carried out at the Novokubyshevsk NPZ confirmed the laboratory test results of the VNIIP. There are 6 tables and 2 non-Soviet references.

ASSOCIATION: VNIIP

Card 5/5

AGAFANOV, A.V., RYSAKOV, M.V., GOLDSHTEYN, D.L., GUSENKOVA, YE.A.,
ALFIMOVA, YE.A., POSHITNOV, V.N.,

Gewinnung von Motorölen aus schwefelhaltigen Rohölen durch
Hydrierung.

Report to be submitted for the Symposium Lubricants and
Lubrication, Dresden, 27-30 June 1961

S/064/61/000/007/003/005
B124/B206

AUTHORS: Agafonov, A. V., Dubinin, M. M., Onusaytis, B. A.,
Tororeshnikov, N. S.

TITLE: Studies on production and application of new selective
adsorbents - molecular sieves - in the USSR

PERIODICAL: Khimicheskaya promyshlennost', no. 7, 1961, 26 - 30

TEXT: The authors give a short summary of the main results of studies in the field of synthetic zeolites conducted in various scientific institutes in 1960 on the basis of the coordination plan of the Komissiya po tseolitam (Zeolite Commission). The Zeolite Commission under the chairmanship of Academician M. M. Dubinin was established at the Otdeleniye khimicheskikh nauk AN SSSR (Department of Chemical Sciences, AS USSR) in 1959, in order to coordinate studies in the field of synthesis and application of synthetic zeolites. Its activities comprised: 1) development of synthesis- and technological processes for synthetic zeolites; 2) investigation of structural properties and adsorption of synthetic and natural zeolites, and 3) study of the application of synthetic zeolites for the drying and separation of gases. Crystallization of zeolites and their ion exchange prop-
Card 1/5

Studies on production...

S/064/61/000/007/003/005
B124/B206

erties were investigated at the Institut fizicheskoy khimii AN USSR (Institute of Physical Chemistry, AS USSR) under the direction of I. Ye. Neymark, zeolites of the types CaA, KA, LiA, BeA etc having been produced (the authors use the designations NaA, CaA, NaX and CaX, approved by the above-mentioned Commission, instead of the customary designations 4A, 5A, 10X and 13X). One of the institutes of the chemical industry under the direction of G. I. Mikulin and V. Ya. Nikolenko investigated the technological conditions for the synthesis of zeolites, and one of the institutes of the petroleum industry under the direction of Ya. V. Mirskiy the conditions for the production of crystalline zeolites of the type NaA and CaA in the laboratory and pilot plant. Optimum conditions for the synthesis of zeolites of the types NaA and NaX, as well as the ion exchange for the production of the CaA and CaX zeolites were studied in the laboratory under the direction of M. S. Misin and L. M. Maksimova. The conditions for the synthesis of zeolites of the types A and X were studied at the institut neftyanoy promyshlennosti (Institute of the Petroleum Industry) under the direction of A. V. Agafonov, L. I. Piguzova and B. A. Lipkind, applying the process used by N. S. Kurnakov (Ref. 3: Izv. AN SSSR, 6, 1381, (1937)) for the production of Permutit. The use of aluminum sulfate and aluminum oxy-

Card 2/5

Studies on production...

S/064/61/000/007/003/005
B124/B206

chloride in the synthesis of zeolites was studied in a laboratory of the chemical industry under the direction of V. S. Vinogradova and L. S. Kofman. The institut khimii AN Gruz. SSR (Institute of Chemistry of the Georgian SSR) under the direction of G. V. Tsitsishvili dealt with the kinetics of the crystallization of the NaA zeolites, and the Institut khimii silikatov AN SSSR (Institute of Silicate Chemistry, AS USSR) under the direction of S. P. Zhdanov with the optimum conditions for the production of Na zeolites by hydrothermal synthesis in the temperature range of from 70 to 200°C from strongly basic aluminum silica gels with a base excess of 300 - 500%. The studies by the laboratoriya GEOKhI AN SSSR (Laboratory of the GEOKhI, AS USSR) under the direction of N. I. Khitarov dealt with the drying of gases by means of the natural zeolites natrolite, desmine, thomsonite and limonite, while the use of the chemical-catalytical method for the production of natrolite granules was tried out at the IGI AN SSSR (IGI, AS USSR) under the direction of B. A. Onusavtis. D. P. Dobychin elaborated a process for the production of porous glasses of the molecular sieve type yielding a molecular sieve with a porosity close to that of the CaA zeolite from the $H_2O-7/23$ (Na-7/23) glass, and one with a porosity similar to that of the NaX zeolite from the $H_2O-10/30$

Card 3/5

S/064/61/000/007/003/005
B124/B206

Studies on production...

(Na-10/30) glass. A number of investigations of the structure and adsorption of synthetic and natural zeolites was conducted at the Institute of Physical Chemistry, AS USSR under the direction of M. M. Dubinin. The distribution curves of the zeolite crystals were determined by the electron microscope investigation conducted by V. M. Luk'yanovich.

D. P. Timofeyev studied the kinetics of steam adsorption, A. V. Kiselev the adsorption of nitrogen, benzene vapors and hexane on the molecular sieves NaA and CaX as well as the adsorption of benzene and n-hexane and their mixtures on the molecular sieve CaA. X-ray photographic investigations were made under the direction of N. A. Shishakov. Studies conducted under the direction of I. Ye. Neymark at the Institute of Physical Chemistry, AS USSR showed that the equilibrium adsorption on zeolites is well described by the potential theory, and that the thermal stability of zeolites drops in the sequence $CaA > KA > NaA > NH_4A$. The properties of Soviet

and American molecular sieves during drying of gases were compared at the Leningradskiy tekhnologicheskii institut im. Lensovet (Leningrad Technological Institute imeni Lensovet) under the direction of T. G. Plachenov and G. M. Belotserkovskiy. Studies on the drying and purification of gases by means of molecular sieves were conducted at the Moskovskiy khimiko-

Card 4/5

S/064/61/000/007/003/005
B124/B206

Studies on production...

tekhnologicheskii institut im. D. I. Mendeleeva (Moscow Institute of Chemical Technology imeni D. I. Mendeleev) under the direction of N. S. Torocheshnikov and N. V. Kel'tsev, and by V. S. Vinogradova, L. S. Kofman and Ya. V. Mirskiy. In 1960 the Zeolite Commission held three meetings (in Moscow, Leningrad, and Grozny) in the form of scientific colloquia with 120 - 150 participants. There are 4 references: 2 Soviet-bloc and 2 non-Soviet-bloc. The two references to English-language publications read as follows: R. M. Barrer, Brit. Chem. Eng., No. 5, 1 (1959) and US Patents 2882243, 2882244, 1959.

Card 5/5

11.0140

26520
S/065/61/000/008/003/009
E030/E135

AUTHORS: Rogov, S.P., Gol'dshteyn, D.L., Osipov, L.N., and Agafonov, A.V.

TITLE: Hydrofining the high-sulphur kerosine-gas oil fraction of Arlan crude

PERIODICAL: Khimiya i tekhnologiya topliv i masel, 1961⁶ No.8, pp. 13-19

TEXT: The preparation of satisfactory diesel fuels from Arlan crudes has been investigated by VNII NP. In the laboratory straight fractions were hydrofined; this process lowers the flash point and it was found necessary to remove subsequently the fractions boiling up to 180 °C to keep the flash point in the 60-65 °C region. However, the diesel fuel then fails specification ГОСТ 4749-49 (GOST 4749-49) and 305-58, on pour point (-9 °C instead of -10 °C). However, hydrofining cat. cracked products gives satisfactory diesel fuels, and it is recommended that these be blended with the straight run components. In order to increase the output of the benzine fractions, without raising the diesel pour point, hydrofining experiments were then conducted on a
Card 1/2

Hydrofining the high-sulphur

26520
S/065/61/000/008/003/009
E030/E135

tungsten-nickel oxide catalyst at 300 atmospheres pressure. One can then obtain about 20% benzine and 80% diesel fuel, the latter being of satisfactory quality. However, the benzine octane number is only about 50 and it should be used either for motor spirit blending or for further platinum reforming. There are 6 tables, and 3 Soviet references.

ASSOCIATION: VNII NP

Card 2/2

GLIKMAN, L.S.; BOCHAROV, I.V.; VIKHMAN, G.L.; ABROSIMOV, B.Z.; KIRILOV,
Ye.A.; MEL'NIKOV, S.M.; AGAFONOV, A.V.; SOSKIND, D.M.

Rebuilding catalytic cracking units with a combined reactor-regenerator.
Khim. i tekhn. topl. i masel 6 no.11:6-10 N '61. (MIRA 14:12)

1. Gosudarstvennyy nauchno-issledovatel'skiy i proyektnyy institut
neftyanogo mashinostroyeniya.
(Cracking process)

128

PHASE I BOOK EXPLOITATION

SOV/6246

Soveshchaniye po tseolitam. 1st, Leningrad, 1961.

Sinteticheskiye tseolity; polucheniye, issledovaniye i primeneniye
(Synthetic Zeolites: Production, Investigation, and Use). Mos-
cow, Izd-vo AN SSSR, 1962. 286 p. (Series: Its: Doklady)
Errata slip inserted. 2500 copies printed.

Sponsoring Agency: Akademiya nauk SSSR. Otdeleniye khimicheskikh
nauk. Komisiya po tseolitam.

Resp. Eds.: M. M. Dubinin, Academician and V. V. Serpinskiy, Doctor
of Chemical-Sciences; Ed.: Ye. G. Zhukovskaya; Tech. Ed.: S. P.
Golub'.

PURPOSE: This book is intended for scientists and engineers engaged
in the production of synthetic zeolites (molecular sieves), and
for chemists in general.

Card 1/3 3

Synthetic Zeolites: (Cont.)

SOV/6246

COVERAGE: The book is a collection of reports presented at the First Conference on Zeolites, held in Leningrad 16 through 19 March 1961 at the Leningrad Technological Institute imeni Lensovet, and is purportedly the first monograph on this subject. The reports are grouped into 3 subject areas: 1) theoretical problems of adsorption on various types of zeolites and methods for their investigation, 2) the production of zeolites, and 3) application of zeolites. No personalities are mentioned. References follow individual articles.

TABLE OF CONTENTS:

| | |
|------------------------------|---|
| Foreword | 3 |
| Dubinina, M. M. Introduction | 5 |

Card 2/12

Synthetic Zeolites: (Cont.)

14
SOV/6246

Misin, M. S., L. M. Maksimova, V. A. Litvinova, and L. B. Khandros. Production and Adsorption Properties of NaA, NaP, CaA and CaP Zeolites

135

Misin, M. S., L. M. Maksimova, V. A. Litvinova, L. B. Khandros, G. A. Polyakova, and L. S. Urin. Production and Adsorption Properties of NaX, CaX, and AgX Zeolites

143

Figuzova, L. I., A. V. Agafonov, A. S. Vitukhina, V. F. Dmitriyeva, A. T. Slepneva, V. A. Burylov, and N. A. Ghepurov. Synthesis Conditions and Thermal Stability of Type X Zeolites

152

Mirskiy, Ya. V., M. G. Mitrofanov, and T. N. Bredikhina. Ion Exchange of Na for Ca in Type A Synthetic Zeolite

167

Mirskiy, Ya. V., M. G. Mitrofanov, B. M. Popkov, L. T. Bolotov, and A. I. Mezhlumova. Production of Synthetic Zeolites Under Industrial Conditions

169

Card 7/15 5/3

S/065/62/000/006/004/007
E194/E436

AUTHORS: Manshilin, V.V., Manakov, N.Kh., Agafonov, A.V.,
Vasilenko, V.P., Maslov, I.Ya., Knyazev, V.S.

TITLE: Testing of engineering development of a new system
for fluid catalytic cracking

PERIODICAL: Khimiya i tekhnologiya topliv i masel, no.6, 1962, 41-50

TEXT: To prevent spontaneous afterburning of carbon monoxide and other combustible gases which can occur in the regenerators of fluid catalytic crackers, damaging the cyclone and causing other faults, the regenerator temperature is kept below 600°C, though in many respects it would be advantageous to raise it to 650°C. To achieve this the free oxygen content of the gas in the regenerator must be reduced by raising the level of coking of the catalyst, by greatly improving the contact between air and catalyst or by a combination of these two methods. A regenerator which achieves this combined effect is the main feature of the system here described. The construction of a pilot plant reactor unit which includes the reactor, a turbulent scrubber, a regenerator and two pneumatic catalyst transport lines is
Card 1/3

Testing of engineering ...

S/065/62/000/006/004/007
E194/E436

described. The regenerator is a vertical cylinder with fireproof lining of 1400 mm internal diameter; it has a three stage cyclone in the upper part. Within the zone of the fluid bed is an inner hollow steel cylinder 600 mm diameter containing cooling coils with air distribution arrangements. The spent catalyst is delivered to the annular zone of the regenerator and, under conditions close to those of ideal mixing, sufficient coke is burned to maintain the temperature in this zone at about 600°C. Because of the intensive mixing there is little local overheating. Combustion of the coke is completed in the control zone and the temperature of the catalyst leaving the lower part of the zone for the reactor can be controlled by the cooling coil. The regeneration process is split into these two stages to improve combustion of the coke. Most of the coke is removed in the first zone, where the mean content of coke on the catalyst is high, the combustion being intensified by the counter current conditions and most of the oxygen used up. Operating conditions are given for the various parts of the unit and the results obtained provide all the necessary data for designing full-scale industrial plant with Card 2/3

Testing of engineering ...

S/065/62/000/006/004/007
E194/E436

reactor and regenerator at the same high level using
pneumatic transport of (PVK). The two-stage and three-stage
cyclones in the reactor and regenerator respectively gave
satisfactory retention of catalyst dust and returned it to the
fluid bed. There are 5 figures and 5 tables.

ASSOCIATION: VNII NP

Card 3/3

S/282/63/000/002/003/005
A059/A126

AUTHORS: Kurganov, V. M., Gonsales, M. A., Agafonov, A. V.

TITLE: Methods of supplying stocks to a reactor of catalytic cracking

PERIODICAL: Referativnyy zhurnal, otdel'nyy vypusk, 47. Khimicheskoye i kholodil'noye mashinostroyeniye, no. 2, 1963, 33, abstract 2.47.186
(Novosti neft. u gaz. tekhn. Neftepererabotka i neftekhimiya, no. 8, 1962, 15 - 21)

TEXT: Stock feeding to the reactor by single vapor-liquid flow has considerable advantages over the separate feeding of the liquid and vapor phases to the reactor, greatly simplifies the operation and reduces the operating expenses of stock preparation. The contacting method based on spraying of the liquid phase over the surface of the catalyst layer is the most unsuitable of all known methods, since it does not exclude coking of the internal surfaces and conglomerate formation. The utilization of any cross section of dropping catalyst film for contacting with the stock creates a uniform distribution of the liquid residue on the greater part of the catalyst, but does not exclude coking of the

Card 1/2

Methods of supplying stocks to a...

S/282/63/000/002/003/005
A059/A126

reactor. The most advantageous of the alternatives considered is the setup based on the method of phase contacting under restricted conditions on moving in the suspended state below the distributing plate (model ВНИИП - K-18 (VNIINP-K-18)). Fitting out the reactors of catalytic-cracking devices with an inlet assembly for the stock according to the model VNIINP-K-18 permits: to process heavy petroleum stocks without coking of the reactor and conglomerate formation; to increase the yield of light petroleum products by 3 to 5%, to reduce catalyst consumption by 0.5 to 1.5 kg/t of the stock; to reduce the temperature of the stock on discharge from the furnace from 480 - 490°C to 420 - 450°C; to prolong the time of passage through the setups and to stabilize their capacity during the whole cycle; to eliminate laborious and dangerous work involving the removal of coke from the internal surface of the reactor. There are 4 figures and 8 references.

[Abstracter's note: Complete translation]

Card 2/2

L 12295-63

S/031/63/000/005/052/075

AUTHOR: Kozlov, G. N. and Agafonov, A. V. 44

TITLE: A study of the regeneration process of alumo-silicate cracking catalyst

PERIODICAL: Referativnyy zhurnal, Khimiya, no. 5, 1963, 500, abstract 5P156, (Tr. Vses. n.-i in-t, po pererabotke neti i gaza i polucheniya isskust-b. zhidk. topliva, no. 8, 39-53) 1962

TEXT: Investigation of the regeneration process of alumosilicate catalysts by air in the temperature range of 500-680°C was conducted on laboratory apparatus with 15 g capacity of the reaction tube for catalyst. Upon introduction of air at a rate of 70 l/hr the regeneration gases were purified from moisture, the CO was oxidized to CO₂ and the latter was absorbed by ascarite. The results of this operation may be judged from the rate of formation of CO and CO₂. The investigation was conducted under kinetic as well as diffusion areas of reaction path, temperature limits of which were previously determined. Catalysts with wide spaced pores and large internal surfaces are easier to regenerate. The rate of regeneration depends on the nature, the chemical composition and the structure of the catalysts. In this process the effect of structural characteristics of the catalyst are

Card 1/2

L 12295-63

S/081/63/000/005/052/075

A study of the regeneration

manifested mainly in the diffusion area. A catalyst which is coked in cracking of petroleum is easier to regenerate than one used in cracking light or heavy sulfur-containing raw materials. The use of this method is proposed for standard determination of the regenerative characteristics of granular alumo-silicate catalysis, with measurement of speed of the process at two temperatures, corresponding to the kinetic and diffusive areas of the reaction path. The article contains a 25-item bibliography. V. Kel'tsev.

Abstractor's note: Complete translation

Card 2/2

AGAFONOV, A. V.

"Production of high-grade petroleum oils and waxes by hydrogenation"

report to be submitted for the Sixth World Petroleum Congress,
Frankfurt am Main, W. Germany, 19-26 Jun 63.

S/065/63/000/001/004/005
E075/E436

AUTHORS: Morozov, V.I., Agafonov, A.V., Abayeva, B.T.,
Ryabov, V.A., Karpenko, L.P., Gilyazetdinov, L.P.

TITLE: The preparation of feedstock carbon black in thermal
cracking units

PERIODICAL: Khimiya i tekhnologiya topliv i masel, no.1, 1963,
39-42

TEXT: A threefold increase in the production of carbon black is scheduled in the current 7-year plan. New feedstocks suitable for conversion into carbon black are therefore required to supplement green and anthracene oils used at present. Catalytic gas oils and lubricating oil extracts (phenol extracts) were subjected to thermal cracking to produce oils suitable for the production of carbon black. The cracked oils (43.5, 36.0 and 54.4% yields of the feedstock for light gas oil, heavy gas oil and phenol extract respectively) contained from 70 to 80% of aromatic hydrocarbons, of which at least 50% were heavy aromatics. The cost of these oils was about half that of green oil and a quarter of anthracene oil. The yields of carbon black from the oils ranged from 47 to Card 1/2

The preparation of feedstock ...

S/065/63/000/001/004/005
E075/E436

56.7%, which compares well with the yields from green oils. The carbon blacks satisfy the ГОСТ 7885-56 (GOST 7885-56) specification. There are 1 figure and 4 tables. ✓

ASSOCIATION: Omskiy Neftepererabatyvayushchiy zavod VNII NP
(Omsk Refinery VNII NP)

Card 2/2

S/065/63/000/003/001/006

E075/E436

AUTHORS: Rysakov, M.V., Agafonov, A.V., Gol'dshteyn, D.L.,
Osipov, L.N., Rogov, S.P., Khavkin, V.A.

TITLE: Hydrofining of diesel fuels with a considerable
reduction of hydrogen consumption

PERIODICAL: Khimiya i tekhnologiya topliv i masel, no.3, 1963, 7-11

TEXT: In an attempt to refine sulphurous diesel fuels with a reduced quantity of hydrogen, a method was developed with the use of internal H₂ (autofining) as well as external H₂. It was applied to a 1:1 mixture of diesel fuel fractions from Arlan crude and catalytic gas oil from Romashkino crude. The method gave the optimum results at 30 kg/cm² and 400°C. Lowering the pressure to 22 kg/cm² does not affect the H₂ consumption. Increase of temperature to 420 - 440°C, although decreasing the H₂ consumption, may shorten the catalyst life (alumino-cobaltomolybdate). At 400°C and 30 kg/cm² the content of aromatics decreases to 16.3% from 21.6% with a simultaneous increase in the amount of naphthene-paraffins. The catalyst was used without losing its activity for 400 hours at a space velocity of 2.0 h⁻¹, temperature 400°C, pressure 30 kg/cm² and H₂ circulation of 300 m³/m³. The
Card 1/2

Hydrofining of diesel ...

S/065/63/000/003/001/006
E075/E436

consumption of H₂ was 0.2 to 0.3 wt.% of the diesel fuel.
The refined fuel contained 0.12 to 0.13% S (originally 1.62%).
There are 4 tables.

ASSOCIATION: VNII NP

Card 2/2

MOROZOV, V.I.; AGAFONOV, A.V.; ABAYEVA, B.T.; KARPENKO, L.P.

Results of the industrial adoption of the production of crude
for carbon black in thermal cracking devices. Neftoper. i
neftekhim. no.4:18-21 '63 (MIRA 17:7)

1. Omskiy neftepererabatyvayushchiy zavod i Vsesoyuznyy nauch-
no-issledovatel'skiy institut po pererabotke nefli i gaza i
polucheniyu iskusstvennogo shidkogo topliva.

GONSALES, A.A.; KURGANOV, V.M.; AGAFONOV, A.V.; ABAYEVA, B.T.;
POLETAYEV, V.B.; VIV'YER, A.S.; RUDOVICH, M.A.; BELYAYEVA, Z.G.;
RUTMAN, G.I.

Results of redesigning an industrial catalytic-cracking device.
Nefteper. i neftekhim. no.9:6-10 '63. (MIRA 17:8)

1. Salavatskiy kombinat i Vsesoyuznyy nauchno-issledovatel'skiy
institut po pererabotke nefli.

AGAFONOV, A.V.; ABAYEVA, B.T.; OKINZHEVICH, N.A.

Catalytic cracking products for petrochemical syntheses.
Trudy VNII NP no. 9:27-51 '63. (MIRA 17:6)

VASIL'YEV, S.F.; AGAFONOV, A.V.

Obtaining high-octane gasolines. Nefteper. i neftekhim. no.
11:3-6 '63. (MIRA 17:5)

1. Institut goryuchikh iskopayemykh AN SSSR i Vsesoyuznyy
nauchno-issledovatel'skiy institut po pererabotke nefi i
gaza i polucheniyu iskusstvennogo zhidkogo topliva.

ABAYEVA, B.T.; AGAFONOV, A.V.; GILYAZETDINOV, L.P.; GYUL'MISARYAN, T.G.;
ZUYEV, V.P.; MOROZOV, V.I.

Testing thermocatalytic gas oil in the production of furnace black.
Nefteper. i neftekhim. no.12:17-19 '63. (MIRA 17:4)

1. Vsesoyuznyy nauchno-issledovatel'skiy institut po pererabotke
nefti i Nauchno-issledovatel'skiy institut shinnoy promyshlennosti.

MANSHILIN, V.V.; AGAFONOV, A.V.; MANAKOV, N.Kh.; VASILENKO, V.P.;
MASLOV, I.Ya.; KNYAZEV, V.S.; STEPANENKO, I.A.; Primali
uchastiye: VAYL', Yu.K.; NEMETS, L.L.; BELOUSOVA, I.V.;
STOLYARENKO, Ye.G.; YEMEL'YANOV, A.A.; RYABOV, V.M.;
BEREZOVSKIY, V.D.; ZEFIROVA, Ye.G.; CHELOGUZOVA, Ye.F.;
SOLOTSINSKIY, S.Ye.; BOL'SHAKOVA, K.A.; KHRAMOV, A.Ye.

Catalytic cracking of raw heavy distillates on a microspheric
catalyst of Troshkovskiy clay. Khim. i tekh. topl. i masel. 8
no.3:1-6 Mr '63. (MIRA 16:4)

1. Vsesoyuznyy nauchno-issledovatel'skiy institut po pererabotke
nefti i gazov i polucheniyu iskusstvennogo zhidkogo topliva.
(Cracking process) (Catalysts)

AGAFONOV, A.V.; ABAYEVA, B.T.; OKINSHEVICH, N.A.; ANDREYEVA, A.S.;
MOROZOV, V.I.

Developing extraction methods for obtaining carbon black
stock from catalytically cracked gas oils. Khim. i tekhn.
topl. i masel 9 no.5:13-16 5 My'64 (MIRA 17:7)

1. Vsesoyuznyy nauchno-issledovatel'skiy institut po pererabotke
nefti i gaza i polucheniya iskusstvennogo zhidkogo topliva.

AGAPONOV, A.V.; ZHAYDA, B.T.; ORISHENICH, N.A.; SEKUYAN, F.N.; FIBELONOV, V.P.; LYSAROV, G.A.; TRAPUNOVICH, L.B.; FEDIN, S.A.; KHLEVSKIY, I.S.

Obtaining raw stock for the production of active carbon black by extraction with the selective solvents of the gas oils of catalytic cracking. Khim. i tekhn. topl. i masel 9 no.7:36-39 JI '64.

(DATA 17:22)

1. Vsesoyuznyy nauchno-issledovatel'skiy tsentr po obrabotke nefli i gaza i polucheniya iskusstvennogo zhidkogo topliva.

OSIPOV, L.N.; AGAFONOV, A.V.; KHAVKIN, V.A.; ROGOV, S.P.

Effect of nitrogen compounds on the hydrocracking of heavy distillates.
Khim. i tekhn. topl. i masel 10 no.8:1-4 Ag '65. (MIRA 18:9)

1. Vsesoyuznyy nauchno-issledovatel'skiy institut po pererabotke nefi
i gazov i polucheniya iskusstvennogo zhidkogo topliva.

L 49791-65 EWT(d)/EWT(m)/EWP(w)/EWA(d)/EWP(v)/EWP(k)/EWA(h) Pf-4/Pg-4/
Feb IJP(c) Ww/EM

ACCESSION NR: AP5010189

UR/0373/65/000/001/0148/0150

AUTHOR: Agafonov, A. V. (Leningrad)

TITLE: Action of suddenly applied concentrated force on a cylindrical shell

SOURCE: AN SSSR. Izvestiya. Mekhanika, no. 1, 1968, 118-120

TOPIC TAGS: shell theory, partial differential equation, Fourier transform, integral transform

ABSTRACT: A simple asymptotic solution was obtained for small times t describing the deflection of an infinite cylindrical shell under a suddenly applied concentrated load. The governing differential equation is given by

$$\frac{Eh^3}{12(1-\nu^2)} \Delta^2 \left(\Delta + \frac{1}{a^2} \right) v + \frac{Eh}{a^2} \frac{\partial^2 v}{\partial t^2} + \rho h \frac{\partial^2 \Delta v}{\partial t^2} = \Delta \left(\Delta + \frac{\partial^2}{\partial t^2} + \frac{\partial^2}{\partial a^2} \right) v$$

A double Fourier integral transform is used, in x and in a , to solve the above equation with zero initial condition and a zero boundary condition at infinity. The solution for the shell deflection is then given by

Card 1/2

L 49791-65

ACCESSION NR: AP5010189

$$w_0 = \frac{\pi}{2} \int_0^{2\pi} d\varphi \int_0^2 J_0(\cos^2 \varphi \tau) d\tau -$$

$$- \int_0^{2\pi} d\varphi \int_0^2 d\tau \int_0^2 J_0(\cos^2 \varphi \sqrt{\tau^2 - s^2}) \frac{\sin \gamma s}{2s} ds$$

where the second term represents the correction to the deflection itself, as is obtained in the theory of thin shells. The result shows that, at least up to $\gamma = 0.1 \gamma^{-1}$, this correction is of the order X and hence is negligible. In the above result

$$\tau = \frac{\sqrt{E} \rho}{a}$$

Orig. art. has: 14 equations and 1 figure.

ASSOCIATION: none

SUBMITTED: 27 May 64

ENCL: 00

SUB CODE: M3

NO REF SOV: 003

OTHER: 003

Card 2/2

L 53663-65 EWT(m)/EPF(c)/T Pr-4 DJ

ACCESSION NR: AP5013515

UR/0313/65/COC/005/0006/0008
665.534.4

AUTHORS: Dets, M. M.; Agafonov, A. V.

TITLE: Obtaining oil fractions with a high viscosity index by deep hydrogenation of crude oil

SOURCE: Neftepererabotka i neftekhimiya, no. 5, 1965, 6-8

TOPIC TAGS: lubricating oil, lubrication, lubricant viscosity, hydrogenation, oil, solvent action / 3076 sulfide nickel tungsten catalyzer, TsIATIM 58 hydrogenation apparatus

ABSTRACT: The effect of deep hydrogenation on two types of crude (the straight distillation fraction and the gas oil fraction of the catalytic cracking) was studied in an effort to obtain lubricants with the viscosity of 4 and 8 centistoke at 50C. A commercial sulfide-nickel-tungsten catalyzer (3076) was used in all the experiments (pressure of 300 atm and hydrogen cycling 1200-1500 nm³/nm³ of crude). High aromatic gas oil fractions were hydrogenated in two stages to lower the reaction heat. Conditions for the hydrogenation of primary and secondary crudes are tabulated. Paraffin was removed from the fractions obtained in the TsIATIM-58

Card 1/2

L 53663-65

ACCESSION NR: AP5013515

hydrogenation device by selective solvents. Optimal deparaffinization conditions were as follows: solvent—methyl-ketone and toluene (60:40% by vol); the solvent-crude ratio—5:1 by weight; cooling temperature—-20 to -40°C. The output of the lube oils at +50°C and -40°C from the straight distillation crude was higher than that from gas oil. The hydrogenation of narrow fractions resulted in an increased production of oils of similar qualities. Best quality lube oils with viscosity 4 centistoke at 50°C were obtained by hydrogenation of crudes with boiling points 300–350°C at negative temperatures and under less stable conditions, while the processing of crudes with higher boiling points required more stable hydrogenation conditions for the same quality of lube oils. See Tables.

ASSOCIATION: VNIINP

SUBMITTED: 00

ENCL: 00

SUB CODE: FP

NO REF SOV: 000

OTHER: 000

482
Card 2/2

L 58285-65 EWT(m)/EPF(c)/I Pr-4 WE

ACCESSION NR: AP5016198

UR/0318/65/000/006/0029/0032
543.2.665.4

22
20
B

AUTHORS: Dets, M. M.; Agafonov, A. V.

TITLE: Viscosity-temperature characteristics of low solidifying lube oil fractions obtained by hydrogenation

SOURCE: Neftepерerabotka i neftekhimiya, no. 6, 1965, 29-32

TOPIC TAGS: petroleum, hydrogenation, paraffin hydrocarbon, naphthenic ring, temperature, viscosity

ABSTRACT: The effect of hydrogenation intensity on viscosity and temperature of lube oil fractions with 3,0 centistoke at 50C and low viscosity at negative temperatures was studied on three fractions of the Romashkinskaya oil: Nos. 1 and 2 obtained at 350-400C and No. 2 at 300-400C. Two catalysts were used: (8376)—NiO-Al₂O₃-SiO₂ and (8377)—with the same composition of aluminum oxide. The fractions with boiling points at 100-150C were separated from the original mixture and separated. The fractions with boiling points at 150-200C were obtained by the adsorption separation of the 300-400C fraction on silica gel in an effort to clarify the destructive hydrogenation mechanism. The

Card 1/3

L 58285-65
ACCESSION NR: AP5016198

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characteristics of the crudes, the hydrogenates, the paraffin-naphthenic hydrocarbons and the experimental conditions are presented. It was noted that transformations of the paraffin-naphthene hydrocarbons in the hydrogenation process were very important for the end results (their viscosities being equal at 50C differed at negative temperatures). Hydrocarbons obtained at a higher reaction temperature had better qualities than those obtained at a lower temperature. The hydrocarbons obtained at a higher temperature had a separation of stages, increasing, as lower temperatures were used, the concentration of hydrocarbons concentration in the initial fraction. At higher reaction temperatures, the paraffin-naphthenic hydrocarbons were separated into two fractions, one containing the paraffin-naphthenic hydrocarbons and the other containing the naphthenic hydrocarbons. The hydrocarbons obtained at a higher temperature had a smaller number of condensed hydrocarbons.

Card 2/3

L 58285-65

ACCESSION NR: AP5016198

2

This transformation scheme was sustained by the mass-spectral analysis of the paraffin-naphthene hydrocarbons made according to the n-d-m method. The analysis was carried out in the laboratory under the leadership of K. I. Zimina. Orig. art. has: 3 tables.

ASSOCIATION: VNIINP

SUBMITTED: 00

ENCL: 00

SUB CODE: FP

NO REF SOV: 003

OTHER: 001

Card ^{LR} 3/3

L 27931-66 EWT(m)/T/EWP(t)/ETI IJP(c) JD/WE
ACC NR: AP6017743 SOURCE CODE: UR/0065/65/000/008/0001/0004

AUTHOR: Osipov, L. N.; Agafonov, A. V.; Khavkin, V. A.; Rogov, S. P. 37
B

ORG: VNII NP

TITLE: Effect of nitrogen compounds on hydrocracking of heavy distillates

SOURCE: Khimiya i tekhnologiya topliv i masel, no. 8, 1965, 1-4

TOPIC TAGS: organic nitrogen compound, catalytic cracking, hydrogenation, gasoline, petrochemistry

ABSTRACT: The main results of studies on the effect of organic nitrogen compounds on the yield and quality of two-stage hydrocracking products are presented. A bifunctional catalyst on a carrier with increased acidity served as the catalyst of the second stage. The experiments were carried out on laboratory circulating high pressure equipment with a 160 ml catalyst charge. The hydrogen content in the circulating gas was 90-95%, by volume. The original crude for the second hydrocracking stage consisted of products of the hydrogenation of vacuum gas oil of a mixture of eastern sulfur-containing petroleum stocks on an alumina-cobalt-molybdenum catalyst at 425°C, the space velocity of the crude was 1 hour⁻¹, and the hydrogen pressures were 50, 150 and 250 atmospheres, which made it possible to produce 3 hydro-
genates with different nitrogen contents. The experiments on hydrocracking of crude containing 0.06, 0.01, and less than 0.01% nitrogen on a bifunctional

Card 1/2

UDC: 665.554:661.5

L 27931-66

ACC NR: AP6017743

0

catalyst showed that nitrogen has a substantial effect on the activity and stability of the second stage catalyst of the process. The deactivating effect of nitrogen when its content in the crude was 0.01% and less can be eliminated by increasing the total pressure to 150 atmospheres; the duration of the reaction cycle here was not less than 1400 hours. Two-stage hydro-cracking makes it possible to obtain gasoline with an octane number of about 76 and diesel fuel with a cetane number of 50-55. Orig. art. has: 3 figures and 2 tables. [JPRS]

SUB CODE: 11, 07 / SUBM DATE: none / ORIG REF: 002 / OTH REF: 019

Card 2/2 BLG

L 46019-66 ENT(m)/T WE

ACC NR: AP6021342

(A)

SOURCE CODE: UR/0318/66/000/002/0008/0010

AUTHOR: Osipov, L. N.; Agafonov, A. V.; Rogov, S. P.

31
B

ORG: VNIINP^W

TITLE: Production of motor fuels // by a two-stage hydrocracking of the vacuum distillate

SOURCE: Neftepererabotka i neftekhimiya, no. 2, 1966, 8-10

TOPIC TAGS: catalytic cracking, motor vehicle gasoline, diesel fuel

ABSTRACT: The article describes the results of a two-stage hydrocracking of the vacuum distillate of eastern sulfur-bearing crudes which boils in the 350-500° range, with the use of an aluminum-cobalt-molybdenum catalyst in the first stage and of a special bifunctional catalyst in the second stage. Experiments performed on laboratory units produced good results with a satisfactory removal of nitrogen, sulfur, and other undesirable components in the first stage and high yields of motor fuels in the second stage. A distinctive feature of the second stage is the possibility of controlling the selectivity of the hydrocracking (production of either mostly gasoline or mostly diesel fuel) by varying the temperature and the volume feed rate of the stock over relatively narrow limits. All the gasoline fractions obtained had a low content of unsaturated and aromatic hydrocarbons and consisted mainly of naphthenes, isoparaffins, and n-paraffins. The diesel fuel obtained had a cetane rating of 50 and met

Card 1/2

UDC: 665.644.092.57:662.753

L 46019-66

ACC NR: AP6021342

all the GOST requirements for a low-sulfur summer diesel fuel. Orig. art. has: 1 figure and 3 tables.

SUB CODE: 21/ SUBM DATE: none

Card 2/2

L 45674-66 ENT(m)/T WE
ACC NR: AP6023622

SOURCE CODE: UR/0318/66/000/004/0012/0015

AUTHOR: Agafonov, A. V.; Osipov, L. N.; Rogov, S. P.; Uzunkoyan, P. N.; Finelonov, V. P.; Zhandanovskiy, N. B.; Perezhigina, I. Ya.; Kel'man, I. V.; Pisarchik, A. N.; Afanas'yev, V. I.; Khavkin, V. A.; Laz'yan, N. G. 47 46

ORG: All-Union Scientific Research Institute of Petroleum Refining (Vsesoyuznyy nauchno-issledovatel'skiy institut po pererabotke nefli); Novokuybyshev Petroleum Refinery (Novokuybyshevskiy neftepererabatyvayushchiy zavod) 5

TITLE: Experience with catalytic hydrocracking of vacuum distillate on the hydrofining assembly of the Novokuybyshev Petroleum Refinery

SOURCE: Neftopererabotka i neftekhimiya, no. 4, 1966, 12-15

TOPIC TAGS: catalytic cracking, petroleum product, gas oil fraction, diesel fuel, gasoline

ABSTRACT: The VNIINP has developed a variant of the process for producing diesel fuel involving one-step hydrocracking of sulfur-containing vacuum distillates on an aluminum-cobalt-molybdenum catalyst. The results of laboratory experiments with this variant were successfully applied at the experimental industrial hydrofining assembly of the Novokuybyshev Petroleum Refinery. The operation of the hydrocracking assembly is described. The feed stock for the plant hydrocracking was vacuum gas oil obtained from distillation of sulfur feed stock. Distillation of the hydrogenate produced:

Card 1/2

UDC: 665.644.2.048.5:665.658.2

L 45674-66

ACC NR: AP6023622

diesel oil which met all the requirements of GOST 4749-49 for DL grade; a gasoline fraction characterized by a low sulfur content (0.002-0.03), a relatively heavy fractional composition (melting range 120-180°), and a low octane number (42), and is recommended as feed stock for catalytic reforming; the gaseous products methane (49.2 wt. %), ethane (29.4%), propane (17.8%) and butanes (3.65). The residue of the distillation of fuel fractions is recommended as feed stock for catalytic cracking. It is concluded that the hydrocracking of vacuum gas oil on the hydrofining assembly of NKNPZ confirmed the results of work carried out by the VNIINP on pilot plants for the purpose of designing high-capacity units. Orig. art. has: 1 figure and 2 tables.

SUB CODE: 11/ SUBM DATE: none/ ORIG REF: 001/ OTH REF: 003

Card 2/2 fv

ACC NR: AP6032842

SOURCE CODE: UR/0000/00/

(A, N)
AUTHOR: Pereshigina, I. Ya.; Agafonov, A. V.; Rysakov, M. V.; Osipov, L. N.; Rogov, S. P.

ORG: VNIINP

TITLE: Study of the fundamentals of hydrocracking of a heavy distillate with high sulfur content

SOURCE: Khimiya i tekhnologiya topliv, i masel, no. 10, 1966, 15-18

TOPIC TAGS: petroleum refinery product, petroleum refining gasoline, liquid fuel, diesel oil, desulfurization

ABSTRACT: A study of hydrocracking of high-sulfur vacuum distillate (2.16 wt % S, 0.1 wt % N, 0.9163 specific gravity, and containing 50% aromatics and 50% paraffins and naphthenes) over Co-Mo/alumina catalyst at 50-250 atm, 380-425°C, 0.5-6.0 hourly volume space velocity, and a hydrogen to feed volume ratio of 300-1500 was made. The object of the work was to define the optimal process condition for the greatest yield of low-sulfur diesel oil fraction. It was found that in the 600-1500 range of H₂:feed ratio, the H₂:feed ratio did not affect the hydrocracking process. It was also found that the optimal conditions leading to 30-45% yield of low-sulfur diesel oil and very low yields of gas and gasoline fraction are: 50 atm, 400-425°C, and 1-2 hourly volume space ve-

UDC: 665.534:665.521.4

Card 1/2

ACC NR: AP6032842

locity. Under these process conditions, the life of the catalyst was found to be at least three months. Orig. art. has: 3 figures, 3 tables.

SUB CODE: 07,21/

SUBM DATE: none

Card 2/2

DRUYANOV, Lev Aleksandrovich; PANTSKHAV, I.D., prof., red.;
AGAFONOV, A.Ye., red.; ZYKINA, T.N., tekhn. red.

[What matter is] Chto takoe materia. Pod red. I.D.
Pantskhava. Moskva, Uchpedgiz, 1961. 80 p.
(MIRA 15:10)

(Matter--Constitution)

RAYSKIY, Boris Fedorovich; AGAFONOV, A.Ye., red.; KREYS, I.G.,
tekhn. red.

[Technical and work training of the senior grade students]
Politekhnikheskaia i trudovaia podgotovka uhashchikhsia
starshikh klassov. Moskva, Uchpedgiz, 1963. 166 p.
(MIRA 17:3)

SOLOV'YEV, Aleksandr Fedorovich; AGAFONOV, A.Ye., red.; KREYS, I.G.,
tekhn. red.

[Work training of grade 5-8 students; from the rural
school practice of Ul'yanovsk Province] Trudovoe vospi-
tanie uchashchikhsia V - VIII klassov; iz opyta raboty
sel'skikh shkol Ul'ianovskoi oblasti. Moskva, Uchpedgiz,
1963. 165 p. (MIRA 17:1)

AGAFONOV, B.; BABANSKIY, I.

Organization of work and students' wages in school brigades.
Politekh. obuch. no.7:89 JI '59. (MIRA 12:9)

1.Kirovogradskiy oblastnoy institut usovershenstvovaniya uchiteley.
(Kirovograd Province--Agriculture--Study and teaching)

21. 4500

31455
S/626/60/000/012/010/010
D298/D303

AUTHORS: Agafonov, B. M., Dolgikh, T. I., Savchenko, M. I., and Timofeyev-Resovskiy, N. V.

TITLE: Distribution of dispersed elements among the components of reservoirs. IV. Experiments on the distribution of strontium, ruthenium, cesium, cerium and an unseparated solution of uranium fragments in series of tanks

PERIODICAL: Akademiya nauk SSSR. Ural'skiy filial. Institut biofiziki. Trudy. no. 12. Moscow, 1960. Sbornik rabot Laboratorii biofiziki. no. 2: Problemy biofiziki, 238-271 ✓

TEXT: The article describes the results of experiments to study the biological purification of water from a weak solution of an unseparated mixture of uranium fragments, and biological purification from the four main components of this mixture: Strontium-90, ruthenium-106, cesium-137 and cerium-144. The aim of the work was to discover possible differences in the degree of deactivation of the
Card 1/4

Distribution of dispersed ...

31155
S/626/60/000/012/010/010
D298/D303

water from the above-mentioned chemical elements under the same experimental conditions and also to determine similarities and differences in the distribution of these elements among the main components of the reservoirs. The method consisted in passing solutions of the isotopes through a series of small aquarium tanks containing earth, water plants and appropriate microplankton and periphyton. Experiments with strontium showed that with a daily flow of 6 liters of a solution with a concentration of 10 $\mu\text{c}/\text{l}$, the concentration of strontium at the end of the tank series is 4.5 - 10.8% of the original concentration. By decreasing the daily flow of solution to 3 liters and by increasing the size of the first tank the water is deactivated of strontium. Strontium was found to be distributed evenly among the components of the tanks. With a flow of 6 liters/day the concentration of ruthenium at the end of the tank series is 1.5 - 2.5% of the original concentration. Study of its distribution among the components of the tanks showed that it was absorbed mainly by the biomass. In the experiments with cesium, complete deactivation of the water was achieved. Cesium was mainly

Card 2/4

Distribution of dispersed ...

31455
S/626/60/000/012/010/010
D298/D303

absorbed by the ground material in the tanks. With a flow of 6 liters/day complete deactivation of cerium was also attained. Decrease in the daily flow of solution and an increase in the size of the first tank made no essential difference to the results of the experiments. Cerium, like ruthenium, was mainly absorbed by the biomass. A high degree of deactivation was achieved in experiments with an unseparated solution of uranium fragments. The longer the experiment continued, the less was the deactivation of the water. By reducing the flow of the solution to 3 liters/day and by increasing the volume of the first and last tanks, a high degree of deactivation was achieved, even in protracted experiments (more than 6 months). The coefficient of accumulation was highest in the periphyton and detritus, lower in the higher plants and lowest in the ground material. Of the elements studied the highest coefficient of accumulation in the biomass was given by cerium, and the lowest by strontium. In the ground material the highest coefficient of accumulation was given by cesium and the lowest by ruthenium. On the basis of the results the authors divide the radioactive elements studied into three main groups according to their

Card 3/4

Distribution of dispersed ...

31455
S/626/60/000/012/010/010
D298/D303

distribution among the various components of reservoirs: Evenly distributed (strontium), mainly sorbed by the ground material (cesium), absorbed mainly by the biomass (ruthenium, cerium and an unseparated solution of uranium fragments). There are 12 figures, 41 tables and 3 Soviet-bloc references. X

Card 4/4

AGAFONOV, B.M.; DOLGIKH, T.I.; SAVCHENKO, M.I.; TIMOFEYEV-RESOVSKIY, N.V.

Distribution of scattered elements in different components of water reservoirs. Report No.4: Experiments on the distribution of strontium, ruthenium, cesium, cerium, and the unseparated solution of uranium fission fragments in a series of tanks. Trudy Inst. biol. UFAN
SSSR no.12:238-277 '60. (MIRA 14:1)
(Radioactive substances) (Water--Pollution)

21.4500

29421

S/081/61,000/017/062/166
B110/B138

AUTHORS: Timofeyeva-Resovskaya, Ye. A., Agafonov, B. M.,
Timofeyev-Resovskiy, N. V.

TITLE: Biological soil deactivation of water

PERIODICAL: Referativnyy zhurnal. Khimiya, no. 17, 1961, 302, abstract
174321 (Tr. In-ta biol. Ural'skiy fil. AN SSSR, no. 13,
1960, 35-48)

TEXT: The investigations were carried out on laboratory filters, in
pools and in aquariums with weak current. On an average, the following
was retained in filters (filtering rate 0.6 m/hr) filled with mud, clay,
activated carbon etc. (in %): Cs = 100; Sr and Y = 99; a mixture of Nb, Zr,
Ce and U fragments = 80 - 90; Ru = 60 - 70. Optimum deactivation was
observed with natural mud (no active solution passed when some hundreds of
volumes of a solution with a concentration of 10 - 20 $\mu\text{Cu}/\text{l}$ were filtered).
The accumulation coefficients were calculated. For higher aquatic plants
they are $10^2 - 10^3$, for large water invertebrates $10^2 - 10^3$, for small ones

Card 1/2

29421

S/081/61, J00/017/062/166

B110/B138

Biological soil deactivation of water

(zoo- and phytoplankton and periphyton) 10^3 - 10^4 . In experiments made in pools a 90 % water deactivation was attained. When a uranium solution with a concentration of $10 \mu\text{Cu}/\text{l}$ was passed through three successive pools, 99 % of the initial activity were kept back. The activity in the concentration of 10^{-6} - $10^{-4} \mu\text{Cu}$ stimulates the development of fresh water biocoenosis and of the microflora of the soil and the water.
[Abstracter's note: Complete translation.]

X

Card 2/2

TIMOFYEVA-RESOVSKAYA, Ye.A.; AGAFONOV, B.M.; TIMOFYEV-RESOVSKIY, N.V.

Fate of radioisotopes in the bodies of water. Trudy Inst. biol.
UFAN SSSR. no. 22:49-67 '62. (MIRA 16:3)
(RADIOISOTOPES) (WATER--POLLUTION)

AGRE, A.L.; AGAFONOV, B.M.

Some data on the migration of cesium and strontium radioisotopes
from reservoirs with slow circulation. Report No.1. Biul.MOIP.
Otd.biol. 67 no.3:154-155 My-Je '62. (MIRA 15:11)
(Radioisotopes) (Water—Purification)

AGAFONOV, B.N.

"Some Experiments in Biological Deactivation of Water
Laboratory Experiments in a Series of Tanks." p. 54

Trudy Vsesoyuznoy konferentsii po Meditsinskoj Radiologii
(Voprosy Gigieny i Dozimetrii) Medgiz, 1957, Moscow Russian, OK.

Proceedings of the All-Union Conference on Medical Radiology
(Hygienic and Dosimetric Problems).

AGAFONOV, B.S.

Load characteristics of modern oscillator tubes. Radiotekhnika 8
no.4:13-19 J1-Ag '53. (MIRA 11:6)
(Electron tubes) (Oscillators, Electron-tube) (Radio, Shortwave)

PHASE I BOOK EXPLOITATION

452

Agafonov, B.S.

Teoriya i raschet radiotelegrafnykh rezhimov generatornykh lamp
(Theory and Calculation of Radiotelegraph Operating Conditions
for Oscillator Tubes) Moscow, Izd-vo "Sovetskoye Radio", 1954.
534 p. Number of copies printed not given.

Eds.: Shamshur, V.I. and Dikareva, A.I.; Tech. Ed.: Koruzev, N.N.

PURPOSE: The monograph is intended for radio engineers and radio technicians as a manual on oscillator tube characteristics and operating conditions.

COVERAGE: The book attempts to create methods of calculating radiotelegraph operating conditions for newly-developed oscillators. Only such oscillators are described, the design of which has recently been greatly improved. Tubes with a screen grid (pentode or beam tetrode) and an oxide-coated cathode or with a cathode of thoriated carbide tungsten are the most widely used.

Card 1/18

Theory and Calculation (Cont.)

452

TABLE OF
CONTENTS:

| | |
|---|----|
| Foreword | 3 |
| Ch. 1. Special Features of Oscillator Cathodes | 7 |
| 1. General characteristics of oscillator cathodes | 7 |
| 2. Concepts of saturation and space charge operating conditions | 7 |
| 3. Developing the concept of cathode current | 9 |
| 4. Cathode parameters | 11 |
| 5. Dynamic efficiency of the cathode according to the feed current | 13 |
| 6. Dynamic efficiency of the cathode according to the plate current of various types of tubes | 14 |
| 7. Dynamic efficiency of the cathode according to the direct component of the cathode current | 16 |
| 8. Cathode peak efficiency under telegraph operating conditions | 16 |
| 9. Developing the concept of cathode peak efficiency | 17 |

Card 3/18

| | |
|--|-----|
| Theory and Calculation (Cont.) | 452 |
| 10. Tungsten cathode | 18 |
| 11. Operating conditions of a power tube heater circuit during frequent interruptions in operation | 20 |
| 12. Thoriated carbide-tungsten cathode | 20 |
| 13. Oxide-coated cathode | 23 |
| 14. Special performance features of the oxide-coated cathode | 23 |
| 15. Special features of the indirectly-heated cathode | 25 |
| 16. Design of the tube base | 26 |
| Ch. II. Oscillator-tube Plates | 32 |
| 1. Power dissipation by the plate | 32 |
| 2. Materials used for manufacturing plates | 34 |
| 3. Self-cooled oscillator plates | 35 |
| 4. Grid operation and construction | 36 |
| 5. Water-cooled tubes | 37 |
| 6. Forced-air cooled tubes | 41 |
| Ch. III. Characteristics and Parameters of Modulator and Transmitting Triodes | 43 |
| 1. Characteristics and parameters of diodes | 43 |
| 2. Characteristics and parameters of modulator triodes | 45 |

* Card 4/18

| | |
|---|-----|
| Theory and Calculation (Cont.) | 452 |
| 5. Developing the concept of the boundary condition line for tetrodes and pentodes | 79 |
| 6. Fan-shaped plate-grid characteristics of tetrodes and pentodes | 83 |
| 7. Necessity of selecting correct transconductance values in technical calculations of operating conditions for oscillators | 85 |
| 8. Secondary importance of μ and R_1 parameters for screen-grid tubes | 86 |
| 9. Tetrode and pentode plate-grid characteristics for various screen grid voltages | 87 |
| 10. Pentode plate characteristics for various E_{g2} and E_{g3} voltages | 89 |
| 11. The controlling voltage in a tetrode | 92 |
| Ch. V. Physical Processes Occurring in Oscillators with Separate Excitation | 94 |
| 1. Oscillator with separate excitation | 94 |
| 2. Plate current pulse components | 99 |
| 3. Graphoanalytical study of oscillator performance | 101 |

Card 6/ 18

| | |
|---|-----|
| Theory and Calculation (Cont.) | 452 |
| 10. Developing the concept of a-c power existing in the plate circuit | 145 |
| 11. Parallel-fed oscillator circuit | 147 |
| 12. Oscillator equivalent circuits | 149 |
| Ch. VII. Special Features in Calculating Telegraph Operating Conditions for Transmitting Triodes | 152 |
| 1. Methods of calculating the plate circuit and the control-grid circuit for tubes of various design | 152 |
| 2. Shape of plate current pulses in transmitting triodes | 153 |
| 3. Calculating telegraph operating conditions of transmitting triodes having a stepped-up μ | 155 |
| 4. Oscillators working on plate current with square-topped pulses | 158 |
| 5. Calculating telegraph operating conditions of triodes having a medium μ | 161 |
| 6. Calculating amplitude of excitation voltage for triodes having a medium μ and operating on plate current with cosinoidal pulses | 178 |
| 7. Calculating bias voltage E_g and amplitude of excitation voltage U_{pg} for triodes having a low μ and operating on plate current with cosinoidal pulses | 183 |
| Card 8/18 | |

| | |
|---|-----|
| Theory and Calculation (Cont.) | 452 |
| 9. Evaluating various tubes as power amplifiers | 218 |
| Ch. IX. Design of Screen Grid Circuit | 219 |
| 1. Static characteristics of the screen grid current of the type: $i_{g2} = f(e_g)$ | 219 |
| 2. Splitting of screen grid current pulse | 221 |
| 3. Duration of screen grid current pulse. Calculation of current I_{g2} and power P_{g2} | 222 |
| 4. Screen grid current behavior under conditions of undervoltage and overvoltage | 224 |
| 5. Approximate relationships between voltages E_a and E_{g2} , and currents I_{a0} and I_{g2} for various tubes | 225 |
| 6. Methods of screen grid feeding | 227 |
| 7. Danger of secondary emission from the screen grid of ordinary tetrodes | 228 |
| 8. Resultant efficiency of screen-grid oscillator | 231 |
| 9. Current in a suppressor grid circuit | 233 |
| 10. Methods of creating grid bias by using the cathode current | 234 |
| 11. Comparing autonomous and automatic biasing methods | 236 |

Card 10/18

| | |
|--|-----|
| Theory and Calculation (Cont.) | 452 |
| 12. Accuracy of calculations for currents I_{g2} and I_g and excitation voltage amplitude U_{mg} | 238 |
| Ch. X. Load Characteristics of Modern Oscillator Tubes | 239 |
| 1. Concept of load characteristics | 239 |
| 2. Load characteristics of ultrashort wave oscillators | 245 |
| 3. Calculation of load characteristics | 247 |
| 4. Developing the concept of optimum operating conditions for oscillators | 253 |
| 5. Load characteristics of triodes with a low Mu | 257 |
| Ch. XI. Standard Operating Conditions of Oscillators | 260 |
| 1. Possible damages to oscillators | 260 |
| 2. Some operational requirements | 261 |
| 3. Vacuum degree of a tube | 262 |
| 4. Standard operating conditions | 263 |
| 5. Limit electrical values determining the correct utilization of oscillators | 265 |
| 6. Limit electrical values ensuring normal operating conditions for various grids | 273 |

Card 11/18

| | | |
|--|-----|-----|
| Theory and Calculation (Cont.) | 452 | |
| 7. Examining the factors determining the rated plate power dissipation | | 275 |
| 8. Relationships between the rated values of plate power dissipation P_{a_n} and the useful output P_n in tubes designed exclusively for telegraph use | | 276 |
| 9. Relationship between P_{a_n} and P_n for various groups of tubes | | |
| 10. Necessity of reducing plate voltage in oscillators operating on higher than maximum permissible frequencies | | 278 |
| 11. Selecting standard operating conditions for various types of oscillators | | 283 |
| Ch. XII. Possible Methods of Calculating Telegraph Operating Conditions for Oscillators | | 293 |
| 1. Changes in methods of calculation in accordance with changes in oscillator design | | 293 |
| 2. Possible methods of calculating telegraph operating conditions for oscillators | | 296 |
| 3. Calculating telegraph operating conditions for power tubes with tungsten cathodes | | 298 |
| 4. Calculating conditions for screen grid tubes with tungsten cathodes | | 304 |
| Card 12/18 | | |