ITEM I - ANALYTICAL TETHODS

SUMMARY REPORT

This item is apparently taken from the files of Gelsenberg Benzin, Inc., covering the period from late 1938 to about January, 1940.

Sub-items 1, 2, 4 through 39 and 44 deal with the analyses common to the production of steel. Included are methods for the analysis of various types of finished steels and pig iron, of slag (Bessemer ?) obtained from the production of steel, of refractory material used for the linings of steel melting furnaces, materials used in the production of slag, of coal for the production of coke, and, of coke used in the production of steel. They have all been classified as having "no value" to the petroleum industry and there is considerable question in the reviewers' minds if anything would be gained by passing this film on to steel manufacturing interests.

Sub-items 10, 41, and 42 deal with analysis of plant and waste vaters by more or less obvious means. Sub-item 45 through 64a, 77, 85, 89, 91, 103, and 104 deal with analysis of coke oven gases and their treating solutions. The remainder deal with analysis of light oil fractions (benzine, or gasoline) from coal distillation. A number of these have been abstracted, particularly those dealing with the analysis of materials having compositions similar to petroleum material, if a published source of the material could not be readily identified.

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ABSTRACTED AND TRANSLATED SUB-ITEMS

45) Procedure for the Determination of Liouefiable Gases (C2, C3, C4 vithout C5 Fraction) in Strippers and Benzine. (Abstract)

This method describes the separation of C_{1} and lighter (no C_{5} present) hydrocarbons as a single fraction from stripper liquids and gasoline. The separation is made in a large vacuum jacketed column having a metal spiral wire packing. The dephlegmator consists of a bulb condenser maintained at a constant temperature of -15°C. The distillation is reputed to be rapid but the apparatus seems far below current U.S. design quality.

46) Calculation of Alkali Content in Phenol Extractor and Similar Materials. (Abstract)

The method assumes that the extract contains free NaOH, Na₂S, and sodium salts of phenol mixtures (30% phenols and 70% cresols). The total alkali is determined by titration with 1 N acid, the Na₂S by precipitation with acidified cadmium acetate and the phenols sprung with acid and measured. Alternatively the total alkali is determined by a double titration (with indicators not described) to obtain the Na₂S content by difference. The ratio of phenol to cresols is determined from the density of the spring phenols. Caustic is determined by difference.

50) Description and Manipulation of Apparatus for the Detection of Water in Liouefied Gages. (Abstract)

Liquefied sample is transferred to a transparent Dewar flask and inspected for turbidity or large separated ice crystals. Results are qualitative as only the appearance of the liquid is reported.

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53) Analysis of "Alkazid" Solutions. (Abstract)

Describes the analysis of saturated and regenerated "alkazid" solution (apparently an alkaline-solution for the removal of H₂S from 'gases). In the former the specific gravity and total gas (00₂ plus H₂S) are determined. In the latter the specific gravity, total gas, H₂S (alone) and thiosulfate are determined. All determinations are made according to common practices.

62) Determination of Dissolved Iron in Alkazid Solution. (Abstract)

Describes a procedure for the gravimetric determination of iron (as Fe(OH)₃). If SiO₂ or aluminum present, iron is determined colorimetrically as thiocyanate.

64) Directions for Determination of Arsenic in "Alkazid" Solution. (Abstract)

Arsenic is removed from an acidified solution with H₂S. The sulfice is oxidized to arsenate with KClO₃ and in ammonical solution is precipitated as magnesium arsenate with magnesia-mixture. The precipitate is ignited to Mg2As₂O₇ and weighed.

65) C3 and C4 Determination in Benzine (Gasoline). (Abstract)

All the C_3 and C_{ij} is removed from the gasoline by distilling over the first 25% in an ASTM appearatus catching the overhead in a cold trap. The distillate is distilled through a vacuum jacketed column packed with small aluminum spirals. The dephlegmator bulb condenser is held at $-35^{\circ}C$ until no more C_3 is collected. By slowly varning to $-8^{\circ}C$, an isobutane rich fraction is obtained, to $-3^{\circ}C$ a butylene rich fraction is obtained; the remaining C_{ij} is taken over at $+3^{\circ}C$. The analysis is reputed to be rapid.

66) C3, C4 and C5 in Gases. (Abstract)

This is an extension of the procedure described in No. 65 differing principally in that the amounts of different components collected in the traps are recorded together with the temperature of the dephlegmator head which is allowed to increase slowly

68) Temo Concerning Tank Gas Shipment Samples. (Abstract)

This is a certified report of properties of tank car shipment of gas oil from Teerver-waltung D-Meiderich to Gelsenberg as determined by a referee laboratory.

69) Description of "Sarmiza" Process for Production of High Octane Fuel (Non-Catalytic). (Abstract)

This is a brief abstract of a lecture given by one H. Weiss describing the Rumanian "Sarmiza" process for the production of aromatic rich aviation fuels. Paraffin and gas oils are vaporized (300°C),

ITEM I - ANALYTICAL METHODS (cont.)

cracked (500°C) polymerized at higher temperatures (600°C) and dehydrogenated (720°C) all being accomplished without pressure or catalyst. Some data on properties of finished product given. No details or references given.

82) Oxidation of NO to NO2. (Translation)

"With kieselgur alone or after impregnation with NiO, CoO, MnO2 according to Burdick (JACS 111, 2114 (1922)) gives only a slight increase in the oxidation velocity. Silica gel is capable of increasing the exidation of NO considerably. The velocity constant on the surface of the silica gel is about 20 times greater than in the gas phase. With decreasing bulk density in general the catalytic activity of the silica gel increases. But since the total surface of the gel decreases per volume unit with decreasing bulk density the use of highly voluminous silica gel is not advisable."

90) Preparation of Non-foaming 50% KOH for Microazotometer. (Translation)

"Some pulverized activated charcoal (T-Kohle) is triturated with water to a thin paste until all the charcoal is well wetted. Suspension is drawn through a glass filter funnel of medium porosity (with vacuum) until a layer of 1 cm. is formed. The filtrate should be absolutely clear; otherwise wash the mat with water until a clear filtrate is obtained. After the charcoal has been dried as much as possible with suction, the hot 50% KOH is filtered under continuous suction. The filtrate is clear and does not foam."