FOREIGN TECHNOLOGY DIVISION



A METHOD FOR OBTAINING NITRIC ACID

by

G. A. Skvortsov, M. M. Karavayev, et al.



Distribution of this document is unlimited. It may be released to the Clearinghouse, Department of Commerce, for sale to the general public.

Reproduced by the CLEARINGHOUSE for Federal Scientific & Technical Information Springfield Va. 27151

This translation was made to provide the users with the basic essentials of the original document in the shortest possible time. It has not been edited to refine or improve the grammatical accuracy, syntax or technical terminology.

UNEDITED ROUGH DRAFT TRANSLATION

A METHOD FOR OBTAINING NITRIC ACID

By: G. A. Skvortsov, M. M. Karavayev, et al.

English pages: 2

SOURCE: Patent No. 183194 (Appl. No. 893995/23-26, April 13, 1964), 2 pages.

Translated by: J. Anderson/TDBXT

TA7000887

THIS TRANSLATION IS A RENDITION OF THE ORIGINAL POREIGN TEXT WITHOUT ANY ANALYTICAL OR EDITORIAL COMMENT. STATEMENTS OR THEORIES ABYOCATED OR IMPLIED ARE THOSE OF THE SOURCE AND BO NOT NECESSARILY REPLECT THE POSITION OR OPINION OF THE FOREIGN TECHNOLOGY DIVISION.

PREPARED BY

TRANSLATION DIVISION POREIGN TECHNOLOGY DIVISION WP-APB, ONIO.

FTD-HT - 23-953-67

Date 21 July 19 67

ITIS INDEX CONTROL FORM

97	TA7000887		FTD-HT	68 Translation Nr FTD-HT-23-953-67 as 64 Control Marking		Ref Acc N 025584		76 Reel/Frame Nr 1654 0707		
02	UNCL		ICL, O	0	ings			40 Ctry Info		
-	UR	0000	04 Yr 00	05 Vol	06 Iss		9. 45 E. Pg.	10 Date		
	Transli	erated T			000	0001	0002	13APR64		
09	English	Title		OB POLUCHEN			Υ			
43	Source	PATEN	<u>A METH</u> T 183194 (<u>OD FOR OBTA</u> 893995/23-2	INING NITE	CIC ACID	^			
42	Author SKVORT	SOV, G.			Document Lo		0			
16	16 Co-Author				47 Subject Codes					
16	KARAVAYEV, M. M. 16 Co-Author				07, 11					
16	KIRILLOV, I. P. 16 Co-Author				39 Topic Tags: nitric acid, nitrogen compound nitric oxide, temperature					
16	FERD, N	<u> </u>								
	ALEKSEYENKO, D. A.									

ABSTRACT: This Author Certificate presents a method for obtaining nitric acid under the pressure of 5--10 atm, out of nitrogen oxides in the system of condensation of water vapors. To increase the concentration of nitric acid, the unreacted nitrogen oxides are absorbed by the produced acid at a temperature of 25--45C in the absorption part of the bleaching column. English translation: 2 pages.

U. S. BOARD ON GEOGRAPHIC NAMES TRANSLITERATION SYSTEM

Block A a 6 B F F A B F F A B F F A B F F F F F F F	Italic A a B b B c C d E w C w C w C w C w C w C w C w C w C w C	Transliteration A, a B, b V, v G, g D, d Ye, ye; E, e* Zh, zh Z, z I, i Y, y K, k L, l M, m N, n O, o	Block P C T y 中 x u u u u u b u b b b b b b b b b b b b	Italic P P C T Y P X Y Y W W W W W W W W W W W W W W W W W	Transliteration R, r S, s T, t U, u F, f Kh, kh Ts, ts Ch, ch Sh, sh Shch, shch " Y, y E, e Yu, yu
Ооп	0 o	0, o	a Q	O O	Yu, yu
	17 n	P, p	R R	A A	Ya, ya

^{*} ye initially, after vowels, and after ъ, ь; e elsewhere. When written as ë in Russian, transliterate as yë or ë. The use of diacritical marks is preferred, but such marks may be omitted when expediency dictates.

The state of the s

A METHOD FOR OBTAINING NITRIC ACID

G. A. Skvortsov, M. M. Karavayev, I. P. Kirillov, M. L. Ferd, D. A. Alekseyenko, and I. M. Kaganskiy

This invention concerns the industrial production of nitric acid of increased concentration.

The process of obtaining nitric acid from nitrous gases containing nitric oxide under conditions of condensation of water vapors and subsequent concentration of nitric acid by processing it with gases containing nitric oxide is well known.

By using the method offered here it is possible to increase the concentration of acid to 70-74% without using additional nitric oxides, resulting in conversion of a maximum amount of the oxide of nitrous gases to acid. In this method the process of acid formation under conditions of condensation of water vapors takes place in a refrigerator-condenser at a pressure of 5 atm.

Unreacted nitric oxides are absorbed by nitric acid at temperatures no higher than -5°C in absorption column, with subsequent bleaching.

The obtained 58-62-% nitric acid is strengthened to 70-74% in the absorption part of the bleaching column at a temperature of 25-45°C by concentrated gases in the lower part of the bleaching column.

Example: Nitrous gases from conversion of ammonia, containing vol. % 6.56 NC; 4.37 NO₂; 68.76 N₂; 1.91 O₂, and 18.4 H₂O, after preliminary cooling to 120-150°C at a pressure of 5-10 atm, are passed through cooler—ondensers, where at a temperature of 20-25°C the process of acid formation takes place. Unreacted gases, containing 4.77% O₂; 0.2-0.4% O₂; and 91.02% N₂, after cooling to -5°C, are fed to the absorption column, where at this temperature the nitric oxides are absorbed by the 72.0% HNO₃: The acid which has absorbed the nitric oxides from the absorption column is fed to the center of the bleaching column, in the lower part of which in air at 45-55°C the oxides are desorbed. To the upper, strengthening, part of the bleaching column is fed condensate from the cooler-condensers, containing 60.1% HNO₃ and 39.9% H₂O. The strengthening of acid to 72% is due to the desorbed nitric oxides in the bleaching part. Air, together with unreacted nitric oxides in the bleaching column, combines with nitrous gases of conversion and goes to the cooler-condensers. The degree of conversion of nitric oxides is 98.5-99.5%.

Object of invention

A method of obtaining nitric acid under pressure of 5-10 atm from nitric oxides under conditions of condensation of water vapors, <u>distinguished</u> by the fact that for the purpose of increasing the concentration of nitric acid the unreacted nitric oxides are absorbed by the produced acid at a temperature of not more than -5°C, with subsequent bleaching, these being used to strengthen the acid at a temperature of 25-45°C in the absorption part of the bleaching column.