

Method for continuous producing acetic acid (variants) and method for treatment of acetic acid flow

Part of this liquid residue 27 is passed to a first evaporator 22' and then returned to the column to provide the heat required for fractionation, and the remainder is passed to a second fractionating column 24'.

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The resulting standardized solutions had the color number APHA 5, APHA 10, APHA 15, APHA 20 and APHA 25, respectively. As an example, the patent describes UD the other hexylidene of acetic acid at temperatures from about 25° With up to about 45°C.

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For practical purposes, the impeller-equipped stirring vessel type and sparger-equipped bubble column type are preferred.

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JP48030615 September, 1973 JP5564545 November, 1978 JP60222439 April, 1984 PURIFICATION OF CRUDE ACETIC ANHYDRIDE JP61002052 January, 1986 AUTOMATIC MOISTURE METER FOR GRAIN JP61056151 March, 1986 PURIFICATION OF ACETIC ACID JP01211548 August, 1989 PURIFICATION OF ACETIC ACID WITH OZONE JP02231448 September, 1990 METHOD FOR TREATING ACETIC ACID WITH HYDROGEN IN THE PRESENCE OF HYDROGENATION CATALYST JPH02231448A 1990-09-13 JPS4830615A 1973-04-23 JPH01211548A 1989-08-24 JPS5564545A 1980-05-15 JPS6156151A 1986-03-20 JPS612052A 1986-01-08 JPS60222439A 1985-11-07 What is claimed is: 1.

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