

YFARS

OCCASION

This publication has been made available to the public on the occasion of the $50th$ anniversary of the United Nations Industrial Development Organisation.

TOGETHER

for a sustainable future

DISCLAIMER

This document has been produced without formal United Nations editing. The designations employed and the presentation of the material in this document do not imply the expression of any opinion whatsoever on the part of the Secretariat of the United Nations Industrial Development Organization (UNIDO) concerning the legal status of any country, territory, city or area or of its authorities, or concerning the delimitation of its frontiers or boundaries, or its economic system or degree of development. Designations such as "developed", "industrialized" and "developing" are intended for statistical convenience and do not necessarily express a judgment about the stage reached by a particular country or area in the development process. Mention of firm names or commercial products does not constitute an endorsement by UNIDO.

FAIR USE POLICY

Any part of this publication may be quoted and referenced for educational and research purposes without additional permission from UNIDO. However, those who make use of quoting and referencing this publication are requested to follow the Fair Use Policy of giving due credit to UNIDO.

CONTACT

Please contact publications@unido.org for further information concerning UNIDO publications.

For more information about UNIDO, please visit us at www.unido.org

The Williams of the print of the theoretical terms

08743

Mstr. LINITED $ID/WG.293/A$ 28 March 1979

United Nations Industrial Development Organization

ENGIJ SH

Workshop on Fermentation Alcohol for Une as Fuel and Chamical Feedstock in Developing Countries

Vienna, Austria, $26 - 30$ March 1979

I IMMANOL-BASED CHENICAL INDUSTRY IN BRAZILA

۱w

F.A. Ribeiro Filho**

 \bullet The views and opinions expressed in this paper are those of the author and do not necessarily reflect the views of the secretariat of UNIDO. This document has been reproduced without formal editing.

^{**} PETROQUISA, Rio de Janeiro, Branil

with 08743

LIMITED $ID/WG.293/4$ **ABSTRACT** 5 February 1979 **ENGLISH**

Distr.

United Nations Industrial Development Organization

Workshop on Fermentation Alcohol for Use as Fuel and Chemical Feedstock in Developing Countries

Vienna, Austria, $26 - 30$ March 1979

ABSTRACT

THE ETHANOL - BASED CHENICAL INDUSTRY IN BRAZIL⁺

by

F.A. Ribeiro Filho**

The monograph presents the Brazilian background in the production of ethanol-based chemicals as well as the future outlook for the sector.

Such alcohol based industries flourished in Brazil for a pariod of time. Nevertheless several of them were shutdown due to the. use of petrochomical feedstocks for the production of those chemicals.

Following the energy crisis and the associated increase in the price of crude oil, which have particularly affected the economics of the developing countries that depend on oil imports, a new emphasis has been given in Brazil to the use of ethyl alcohol, either as motor fuel or as raw material for the chemical industry.

The views and opinions expressed in this paper are those of the author and do not necessarily reflect the views of the secretariat of UNIDO. This document has been reproduced without formal editing.

PETROQUISA, Rio de Janeiro, Brasil id.79-569

In regard to the chemical industry, a great number of old alcohol-based facilities are being reactivated and now ones are also boing planned.

A list of all ethanol-based chemical plants in Brazil is presented, with their respective capacities, process licensors, markets, the precent status and eventually their plans for expansions.

On the other hand, the prospective use of new alcohol-based chemical routes in Brazil is discussed.

The main processes ever used in Brazil for the production of ethanol-based chemicals are presented and evaluated from the economical point of view, comparing them to the conventional. pol echemical routes.

I

i

BURGARY

٦

١

Brazilian ethanol-chemical experience is analysed at different historical moments.

Initially, the author deecribes the moat important facts of the ethanol-chomical industry in Brasil, fron, installation of the first industrial unita to manufacture several products in a small scale, through 1973-74 when some large products were already during the 1973-74 when some projects were already implemented.

Next there is ^a more detailed review of present situation of the ethanol-chomical industry and its trends. After analysing tho impact of the petroleum crisis, which, through the National Alcohol Plan, led to a comeback and strengthening of this industrial activity in the country, the author discourses about the prospects of implementation, during the next years, of several ethanol-based chemical projects.

The princ .pal ethanol-ch mical processes already utilized by producing companies are described and evaluated, from a technical and economical point of view, to better judge their competitiveness as compared to alternative petrochemical routes.

The author, finally, ponders about the future of tho ethanol-chemical industry in Brasil, and tries to point out the courses of future developments.

-1-

1. DEVELOPMENT OF THE ETHANOL-CHEMICAL INDUSTRY IN BRAZIL

The ethanol-chemical industry, in the sense attributed to is in this paper, relates to the manufacture of chemicals using ethanol as raw material.

Markedly distinct phases characterize the development of this activity in Brazil until 1974, as set forth below.

1.1. Pioneer Phase (1920-57)

١

Commencement of the ethanol-chemical industry in the country goes back probably to the first years of the twenties, when PRODUTOS QUÍMICOS ELEKEIROZ, USINA COLOMBINA and CIA. QUÍMICA RHODIA BRASILEIRA, all inttalled in the State of São Paulo, obtained popularity by manufacturing perfume-squirters based on ethyl chloride, from ethanol.

Beside» ethyl chloride, RHODIA BRASILEIRA »anufactured., in its Santo André unit, acetic acid and ethyl ether in small quantities, both also obtained from ethan.1.

In 1929, the sector had a significant upeving with the installation of CIA. BRASILEIRA RHODIACETA which in subsequent years began to manufacture cellulose acetate using acetic acid and acetic anhydride, the latter also synthesized from ethanol. Later (1965) this company and RHODIA BRASILEIRA would merge to form RHODIA INDUSTRIAS QUÍMICAS E TEXTEIS (RIQT).

By 1940 the following derivatives of ethanol were already being produced in Brazil: ethyl chloride, ethyl ether (at RHODIA and at the War Ministry, in Piquete, State of Sao Paulo), acetic acid and acetic anhydride.

I

 \bullet

In the early fifties, USINA VICTOR SENCE, installed at Conceiçâo de Macabu, in Rio de Janeiro, started commercial production of butanol and acetone from residual molasses fermentation. In 1952 the company inaugurated its plant to produce acetic acid from ethanol, as well as a continuous esterification unit for the production of butyl acetate.

1.2. Expansion Phase (1958-69)

In late fifties, Brazilian ethanol-chemical industry was directed toward refinery ethylene production, then insufficient to meet requirements of the first petrochemical units using it as raw material.

Two alcohol ethylene units were installed, One by UNION CARBIDE DO BRASIL, in 1958, to supply its low density polyethylene plant, and the other by CIA. BRASILEIRA DE ESTIRENO, in 1959, to supply its styrene plant.

In 1962, INDUSTRIAS QUÍMICAS ELETRO CLORO started its ethylene production by means of ethanol denydration to meet requirements of iss high density polyethylene plant. Around the same time RHODIA's vinyl acetate unit started operations, employing acetylene and acetic acid obtained from ethanol.

Molasses availabilities then abundant which .permitted alcohol production in large scale and at Marginal costs, encouraged the implementation of more enterprising ethanol-chemical projects.

CIA. PERNAMBUCANA DE BORRACHA SINTETICA - COPERBO in 1965 started to produce butadiene from ethyl alcohol, according to a process developed by UNION CARBIDE in the States during the Second World War. Processing facilities also covered production of the intermediate aoetaldehyde.

In 1969 began production of 2-ethylhexanol, by means of an entirely eth-nol-chemical reute, by ELEKEIROZ DO NORDESTE, and butano!, acetic acid and ethyl acetato were obtained in the same unit as by-products.

The sixties determined therefore an outstanding growth of the Brazilian ethanol-chemical industry, which then was already covering a widely diversified range of products.

1.3. Decline Phase (1970-74)

In the carlies seventies, reflecting the problems which began to be felt at the end of the preceding decade, difficulties in supplying ethyl alcohol to the chemical industry worsened. Raw material was scarce and suffered successive price increases (a 67% increase during period 1967-69), in view of the favorable circumstances displayed by molasses and sugar international market.

Therefore, competitivity of ethanol-chemical products as compared to imported similar products became extremely difficult, since the latter were manufactured from a negligible price (less than \$2.00/bbl.) raw material (petroleum).

On the other hand, implementation in the country of large petrochemical projects, to take advantage of the effects of economies of scale and of the utilization of processes more advanced than those of the ethanol-chemical industry, rendered even more unfeasible use of the latter to manufacture the same products.

Such facts made the operation of some ethanol-chemical units non-economical and the ethylene units of CIA. BRASILEIRA DE ESTIRENO (1970) and UNION CARBIDE (1971) and the butadiene unit of COPERBO (1971)

 \sim).

were shut down.

r

 $-5-$

ELETRO CT.ORQ ethylene unit cvon stopped for three months, in 1973, but was started up again in view of the difficulties to supply petrochemical othyleno, and is in operation until this data.

2. COMEBACK AND STRENGTHENING OF THE FECTOR

 ~ 1000

With the supply crisis and continuous increase in petroleum prices, which affected specially the economy of developing countries depending upon imported raw material, emphasis began again to be given in Brazil to using ethyl alcohol, both as a fuel and for the manufacture of chemicals.

In this sense, the Brazilian Government, in November 1975, instituted the National Alcohol Plan (PROALCOOL), with the main purpose of rapidly expanding alcohol production in the country through modernization, expansion and implementation of independent distilleries or to sugar mills and its corresponding agricultural annexed projects.

For the purpose of defining the incentives, policies, priorities and execucion plans of the program and to review applications for acceptance of projects, the National Alcohol Commission was created, composed of representantives of six State Departments, and with the operational support of the Jugar and Alechol Institute (IAA) and the National Petroleum (cuncil (CNE).

From late 1975 through March 1979, the National Alcohol Commission had approved 218 projects for distilleries throughout the national territory, 95 of which were annexed to mills and 123 independent distilleries.

Such facilities, expected to begin normal operations until the 1932/83 crop, represent a total production capacity of 4,9 billion liters per year.

The first results of PROALCOOL are already being felt. And therefore, as evidenced in Table I, national alcohol production increased from 551,7 million liters in the 1975/76 crop to 1,47 billion liters in the 1977/78 crop.

 $-5-$

١

ROTILIAN ACHOLIC RAILIZARE

(*) Estimated

Present trend, as can be seen from said Tabl , is toward an incidase in percentage of anhydrous alcohol with a view to its progressive utilization in carburant blonds added to gasoline.

Taking as a reference year 1977, from January through December, it was found that, from 1.38 billion liters of alcohol produced, 79% were in anhydrous alcohol and the remainder in industrial alcohol. During the same year, alcohol consumption for carburant purposes was 741 million liters, whereas the portion destined to chemicals production (excluding cosmetics and pharmaceutical specialties) was of the order of 100 million liters.

The othanol-chemical industry, therefore, is shown as a secondary factor in view of use of alcohol as a fuel, to the extend that raw material availabilities, as estimated in the Fational Alcohol Plan, ensure its future expansion with no significant reflection on the volume intended for automotive uses.

 $u(t)$ is an respect of the chemical industry, the Brazilian Government decided to subsidise production from ethyl alcohol of such organic derivatives which could be manufactured alternatively by a petrochemical route. In this sense, price of a cubic meter of ethanol for such use is ensured in up to 35% of price per ton of petrochemical ethylene, and the National Petroleum Council is entrusted with establishing the quotas to be destined to the industries, whereas the Sugar and Alcohol Institute is in charge of promoting, prioritarily, its »upply.

On the other hand, scarcity, in the international technology market, of competitive cthanolchemical processes, where scale levels and economicability are concerned, as compared to processes in modern chemical industry, have led companies and research Jnsti-

*

.*.

 $-\beta$ -

tutions in the country to take up again' studies for the development of new processes and to improve those already existing.

 $-9-$

Amongst such initiatives the works already carried out by CENTRO DE PESQUISAS DA PETRONRAS - CEN-PES, should be pointed out. They comprise particularly the basic design of a 60,000 ton/year ethylene unit, from ethyl alcohol, to be constructed by SALGEMA INDUS-TRIAS QUÍMICAS S.A., intended to supply its dichlorocthanc plant.

Works in this orca, originally intended to $scale-up$ units already existing in the country (capacity limited to around 3,000 ton/year (por reactor), culminated in the development of a new procesa, which efficiency was already tested in a pilot plant.

COMPAN1ITA PEKNAMHUCANA DE BORRACHA SINTÉTICA - COPERBO, a company within PETROBRAS System, has been performing experiments, likewise successful, to obtain ethylene from alcohol at its former butadiene unit.

It should be further poin ed out that PETROBRAS QUÍMICA S.A. - PETROQUISA, through its TECH-NICAL MANAGEMENT - GETEC, is presently implementing a technological development program in the acetic acid production field. A similar program, involving technologies for acetaldchyde, acetic acid and butanol, is being carried out by OXITENO INDOSTRIA E COMERCIO S.A.

3. ALTERNATIVE RAW MATERIALS FOR ALCOHOL PRODUCTION IN BRAZIL

 $+$ Λ .

In addition to a formidable territorial extent (8,512,000 Km2), of which only a negligible fraction represents cultivated areas (about 430,000 Km2), Brazil gathers other favorable ecological conditions for the production of ethyl alcohol of vegetal origin, Buch as a tropical dina te, high insulation and quality of soil appropriate to the required cultures.

^m

Within this context, several raw materials are being considered to produce alcohol in large scale in the country, particularly saccharincous materials such as sugar cane and sorghum, and starchy materials such as manioc (cassava) and babaßau.

Table 11 gives a summary of main technological characteristics of such raw materiale, from an agricultural and industrial standpoint, with a view to their utilisation for alcohol production. Figures indicated correspond to average values presently prevailing in Brazil. In practice such values are subject to high fluctuations in accordance with the variety cultivated and the region. Therefore, in certain areas of the Center-South region of Brazil, sugar cane productivity has reached valueu as high as 65-70 t/ha.

Table III lists the projects for distilleries approved by the National Alcohol Commission until March 1979, which sum a total production capacity of the order of 4.9 Million cubic meters of cthanol per year.

3.1. Sugar Cane

Sugar cane, which for centuries has been produced and industrialized in Brazil, stands out as the most immediate use raw material for alcohol production, and shall be liable for the major part of the increase in supply of such product during the next years, specially through the so-called "annexed distilleries",

TABLE II

 \cdot

RAW MATERIALS FOR ALCOHOL PRODUCTION

As Glucose

** Estimated production considering one/two crops per year, respectively

TABLE III

 λ

PROJECTS FOR CONSTRUCTION OF ALCOHOL DISTILLERIES

APPROVED BY KATIONAL ALCOHOL COMISSION UNTIL MARCH 1979

 \bullet

which use molasses coming from sugar mills.

To lesser the effects of international sugar market fluctuations on the domestic alcohol production, the Brazilian Government decided to encourage the construction of "independent distilleries", so called those which employ cane juice directly as rnw material for alcohol production.

From sugar cane, in the preliminary phase of the industrial process, are separated the juice and the bagasse. Such two elements arc the starting point to obtain a yrcat number of industrial products, an can be seen from Figure 1.

Brazilian sugar cane production is presently approximately J00 million tons per year, with a cultivated area of the order of 2.1 billion hectares.

3.2. Sorghum

Sorghum *(corghum vulgare)*, a gramineous pTant native of Africa, was according to FAO statistics for 1974, the fifth cereal in importance around the globe in relation to cultivated area, surpassed only by wheat, rice, corn and barley.

The use of saccharineous varieties of sorghum, whose stalk is rich in sugar, to produce ethyl alcohol has been deserving a special attention in Brazil, in view of the possibility of extracting and fermenting the juice in the same equipment used for sugar cane, which therefore represents an alternative for better utilization of installed capacities of distilleries based on sugar cane use.

Other advantages noted for sorghum aro: i) short cultural cycle, which permits tuo crops per year;

-i *\~*

- li) suitability of the culture tò limited hydric resources regions;
- iii) possibility of using the grains, rich in starch, to produce alcohol, which would permit full utilization of the plant.

The commercial exploitation of sorghum in Brazil is still beginning, and the implementation of this culture in the country depends on agronomical studies to determine the cultivation characteristics and to establish the ecological regions most suitable to obtain high productivity indices.

3.3. Manioc

Production in industrial scale of manioc *{Manihot esculenta)* alcohol was previously carried out in Brazil at the Divinopolia Mill, in Minas Gerais, Which operated during ten years (1932-42), producing ^a total estimated at 5,300 liters of alcohol. Sundry factors, such as low agricultural yield, competition with low price petrol am dérivâtes, < id problems with the industrial processing for starch hydrolysis, have contributed for the halting of such production.

This activity was recently resumed, with the installation by PETROBRAS of a manioc alcohol distillery at Curvelo, Minas Gerais, with a nominal capacity of 60,000 liters/day. The process used is based on the enzymatic hydrolysis of the starch and was •ntirely developed by the NATIONAL TECHNOLOGY INSTITUTE.

As yet, the main problem with manioc utilization in manufacturing alcohol lays in its low agricultural yield which however may be explained by the fact that this is a characteristically subsistence culture, which causes this euphorbiaceous plant to be cultivated in a totally empiric manner. With the im-

!

 $-1¹$

plementation of alcohol distilleries from manioc, whose supply is to be based on an industrial culture, it is expected tnat productivity indices above 2Q *t^t* ha will be attained.

. A manioc alcohol production complex (as shown in Pigure 2) comprises an integrated agro-industrial pole, with the economic feasibility of the project depending highly on the marketing of several by-products.

As regards stilläge (a production of the order of 13 liters per liter of alcohol), with polluting characteristics, several solutions have been submitted for its utilization, such as:

- I) as a fertilizer;
- II) as a producer of a fungal biomass used as animal feed;
- III) as a methane gas generating source.

Brazilian production of manioc is the largest In the world. In 1975 it was slightly higher than 26 million tons, with a cultivated area of 2.0 million hectares.

3.1. Babassu

\

Babassu is a palm tree native of Brazil, and under this name several species of genus *Orhignya* and *Attalca* are known.

Prutification of the palm-tree commences between the seventh and the eighth year, and it produces from ³ to ⁶ clusters/year, each containing an average of 150 to 300 coconuts. The palm-tree's productive life is estimated between 10 and ³⁵ years.

The babassu coconut is constituted of three layers: epicarp (11% of the coconut, as an average), fibrous outside layer; meaocarp (231), starehy-fibroua Intermediate layer; and endocarp (591), ligneous inside

^ J:

layer, where the kernels (7%) are lodged.

The mesocarp has an average content of. 65-68% starch (15-16% of the ecconut), proving to be an important raw material for ethanol production.

By an integrated processing of the different parts of the babassu coconut, a great variety of important by-products is obtained, as shown in Figure It should be pointed out, however, that such pro- $3.$ cessing has only now become feasible, with the appearance of modern equipment to crack the coconut in an industrial scale, which used to be traditionally done in an empiric way for the sole purpose of utilizing the kernels.

In Brazil, the agricultural productivity of babassu and the palm-trac's occurrence area are data still largely debated, and it is believed, however, according to surveys made, that 2.5 ton of coco-' nut per hectare and 15.0 million hectares are, respectively, representativ values. Taking into account that only 23% of this area has a productive coverage, a productive potential of the order of 12.4 million tons of coconut per year are estimated, which could be converted into the following quantities of by-products:

The above data clearly indicate the enormous

 $-18-$

potential of babassu as a source of energetic by-produeta and ohanicals.

•V •af

**

3.5. Other Raw Materials

Within this year a pilot plant for the production of ethanol and foundry coke from vegetable residue (leaves and branches) will be in operation at Lorena, State of São Paulo, employing an acid hydrolysis process developed by the NATIONAL TECHNOLOGY INSTITUTE.

Annual availabilities of vegetable residue originating from the lumber industry are estimated at 15 million tons, which, if fully utilized, could produce approximately 4 billion liters of ethanol and 3 million tons of coke.

 $-20-$

4. PROSPECTS OF NEW ETHANOL-BASED CHEMICAL PROJECTS

Figure ⁴ shows the routeg of an ethanol chomical industry, stressing those already manufactured and those about to be utilized in Brazil, with defined operating units. The others represent projects which may be feasible, presently under consideration by different producing companies,

The already defined projects represent investments of approximately US\$ 100 million (historic value) and involve an additional alcohol consumption of the order Of 400 millions liters/year.

Prospects of Implementation in the country of some specific projects are set forth below.

4.1. Production of Ethylene and Derivatives

Brazil has at present, in operation, two large basic petrochemical complexes (PETROQUÍMICA UNIÃO S.A., in São Paulo, with a 330,00 ton/year of ethylene capacity, and PETROQUÍMICA DO NORDESTE S.A. . COPENE, in Bahia, with a 384,000 ton/year ethylene capacity), and ^a third under project to start up in 1981 (PETROQUÍMICA Do SUL - COPESUL, in Rio Grande do Sul, with a 420,000 ton/year ethylene capacity).

The production of ethylene by means of ethanol dehydration, therefore, shall probably take up, still in the future, a fundamentally supplementary character in relation to the manufacture of same by means a petrochemical route, and be directed toward strategic projects, so considered those which:

- require small quantities or raw material;
- attend to small expansions of ethylene consuming units, which would not justify expansion of naphta pyrolysis complexes;

I

— «re located far from petrochemical producing complexes ;

.
Mang-sara

- meet regional raw material supply schedules.

Within this picture is included the above mentioned project of SALCEMA ethylone unit at Macció, Alagoas, intended to produce dichloroethane locally, which shall be sent to MVC/PVC processing units in other parts of the country. This plan shall not only allow utilization of the chlorine produced there but also the regional availabilities of ethyl atcohol.

COPERSO on its turn, as anticipated, has been studying the adaptation of its butadiene unit to produce ethylene.

On the other band, the Ethylene Unit of CIA. BRASILEIRA DE ESTIRENO has just been modernized and expanded to provide the requirements of its Ethylbenzene-Styrene Unit, to supplement petrochemical ethylene feedstock.

At the same time, other opportunities are being investigated, among which the utilization of ethyl alcohol in Campos, State of Rio de Janeiro, as a possible supplement to feasible petrochemical projects.

4.2. Production of Acetaldehyne and Derivatives

Acetaldehyne and its closer derivatives (acetic acid and solvents) were always produced in Brazil by means of an exclusively ethanol-chemical route, and we can say that the technology for obtaining them from ethanol has been entirely absorbed by the companies employing same in the country.

COPERLO has a project for integrated production of MVA from ethyl alcohol and in this respect, in addition to the aforementioned plans for ethylene production, it shall be promoting the reactivation of its acctaldehyne unit and the installation of an acetic acid plant, for which the possibility of adapting

existing domentic know-how is being considered.

RHODIA, on its turn, has a project to manufacture ethyl acetate at its industrial complex, at Paulinea, São Paulo, expanding its capacity from 9,800 to 25,000 ton/year.

The technology to produce n-butanol and 2othylhoxanol from othyl alcohol may also be considered fully absorbed. Products obtained by means of this route are being placed competitively in the market, and ELEKETROZ DO EORDESTE has a project to expand fivefold its installed capacity.

Another alcohol, the local production of which is foreseen for the next years, is pentaerythritol, obtained from formuldehyde and from acetic aldehyde. Such product shall be produced at the industrial set-up which CONPANHEA PETROQUIMICA DO NORDESTE - COPENOR is. implementing at Camagari, Bahia.

Another project already defined in this area is not production of monochloroacetic acid at $Can_{(n)} = j$, which,

HONCHER DE BRESIL QUIMIEA E FARMACÉUTICA S.A. has been ut liming acctuddehyde to produce chloral and DD" at ate Sho Peulo facilities, and we are informed of a now project for the manufacture of such derivatives in Maceió, Alagoas.

Production of butudiene from sthanol process is still censidered non-economical in Brazil, and it is anticipated that there will be a growing supply of this basic item as co-product in the manufacture of petrochemical ethylone.

4.3. Production of Other Ethanol Derivatives

The cechnology to produce ethyl chloride,

 $-24-$

practiced during may years in the country and for some time suspended by producing companies, has been recently taken over by CTA. BRASTLEIRA DE ESTIRENO, which developed its own process.

 \leftarrow

Such process, already successfully tested, has been utilized in a pilot plant to produce ethyl chloride consumed by the company as a catalyst in athylbonzene manufacture.

As regards ethyl ether and glycol ethers, manufacturing processes are controlled by companies utilizing name. RHODIA intenda to expand and modernize its ethyl ether plant, while OXITENO DO NORDESTE S.A. has a project for a new glycol ethers production unit.

Table IV show: the list of ethanol-chemical units already installed and under implementation in Brazil. The list is limited to immediate ethanol derivatives, despite the fact that from the intermediates indicated therein other chemicals already produced in Brazil are obtained, such as vinyl acetate, cellulose acetate, acetic anhydride and chloral.

 \blacksquare

 \overline{a}

÷

ETHANOL-BASED CHANGEL INDUSTRY IN BRAZIL

5. MAIN ETHANOL-CUEMICAL PROCESSES UTILIZED IN BRAZIL

We give below a description of a few ethanolchemical processes already utilized in the country, as well as a preliminary economical evaluation of units employing such processes.

5.1. Description of Processes

5.1.1. Ethylene Production

The basic feature of the traditional process for the production of ethylene, as that used in Brazil by ELETRO CLORO, is the direct dehydration of ethanol using a silica-alumina catalyst:

$C_2H_5OII \longrightarrow C_2H_4 + H_2O - 10.9$ Keal/mol

The reaction is carried out at temperatures between 300 - 3600C and low pressure over a catalyst bed in a tubular reactor. It results in the formation of several byproducts, such as coke, ethyl eter, carbon monoxide and carbon dioxide. Heat for the reaction, is supplied by the circulation through the reactor's shell of a heating fluid (for example, "Dowtherm"). Ethylene is produced 97-98% pure, being necessary further distillation to give a 99 per cent pure product.

PETROBRAS RECEARCH CENTER - CENPES has developed an adiabatic variation of this process, spurred on by the high costs of the earlier inothermle process.

Feeding ethanol and steam into a row of reactors resulted in a large reduction in coke formation, longer catalyst activity, and appreciable increase in conversion and considerably less byproduct production, when tested on a pilot-

 \sim 000 000 000 0000

 \rightarrow . \rightarrow

scale. It was then necessary to test the process semi-come coially, which was done with a prototype plant at REFINARIA DUQUE DE CAXIAS' rite at Duque de Caxias, Rio de Janeiro. The capacity of the reactor was determined by the adiabatic behaviour of the reaction, wethout the need of additional heat to replace them list.

The 60 ton/yea. prototype has been in operation for a year and seems to confirm the efficiency of the adiabatic reaction. Results of running the plant have so far shown that convension in an the order of 28 per cent and there is no loss of activity or selectivity of the catalyst.

Industrial alcohed may be used with no trestment, such as purification. Also, there is a possibility of operating the reastor at tistics pressures that previously with no loss of product conversion.

The flow, he can the CENTES precess is shows in I (start botte all in acciden) and FIGURE 6 (puit cation becomes).

The process is carried out by using a contiguration of three reactors in a row, giving greater efficiency for lower investment and oper ulino costa.

Manciers are a conventional fixed bed, where the reaction is carried out. The temperature falls to cough the hed, due to the endothermic reaction.

Heat for the reaction is supplied by direct heating of the alcohol and injections of superheated steam in direct contact with the alcohol.

The configuration of three reactors in a row allows to operate each reactor with the same inlet-outlet difference of temperature and processing a greater amount óf alcohol in the second and third reactora.

Impure ethylene is produced at a ⁵ kg/om? gage pressure.

The ethylene is compressed and sent to the purification section, where it is first refrigerated and then fed into crystallizors to promote the formation of hydrate. It is then passed through a molecular sieve, to remove residual water, and a distillation column, where propylene and heavier hydrocarbon impurities are removed at the bottom.

The resultant sthylene contains a maximum of ⁵⁰ ppm propylene, 0.5 percent ethane and ita dew point is -40° C.

5.1.2. Production of Ace'aldehyde

The process utilized in Brazil comprises the catalytical dehydrogenation of ethanol, in the steam phase, according to the following reaction:

 CH_3 -CH₂OH $\frac{2800C_6}{C_1-Cr}$ CH₂ $CH₂-CHO + H₂$ ethanol acetaldahyde

Flowsheet of process ia shown in Figura 7.

I

The ethanol in the storage tank, together with alcohol recovered in the process, is vaporized and fed into a reactor (one or more converters) , wherein it goes through ^a chromecopper catalyst, over a fixed bed, at 260-330OQ temperature and low pressure (3-8 psig).

Efficiency for acetaldehyde is 85-92% for liquid by products $12-6$ %, and gaseous byproducts 3-2%. Main byproducts are ethyl acetato and acetic acid and, in smaller quantities, butyraldehyde, methyl-ethyl-ketone, ketone, butano!, methane, ethylene, carbon dioxide and monoxide and unidentified oils.

The catalyst is regenerated from time to time, which is done in two stages: withdrawal of carbon deposited by using as a burning agent air dilluted with dry steam and reduction of copper oxide into metal by passing a hydrogen stream.

Normal operation period of a converter is 700-1300 hours;, and reactivation period is 70-90 hours.

The acetaldehyde recovery system consists mainly of two condensers, one low-pressure and the other high-pressure, and a gas washing tower.

The major portion of latent heat of the converter's effluent is removed in the first condenser wherein the largest part of un-reacted acetaldehyde and alcohol is liquefied.

The remaining gas is then compressed and goes through the second condenser to remove further the heat, nnd is subsequently washed in a tray column. The hydrogen withdrawn at the top Of the gas washer leaves the unit to bo used as a fuel or sent for flare.

The washing tower bottom stream joins with the low-pressure condenser liquid effluent and, after being mixed in line with an acetaldehyde stream recovered in the process it is fed into a distillation column where high purity acetaldehyde is separated at the top and goes into storage.

 $+$

The acetaldehyde column bottom stream, containing mainly ethanol and water and a smaller quantity of light and heavy byproducts is sent to the byproducts distillation column. In this tower, compounds of a boiling point lower than ethanol arc separated at the top, whereas un reacted etlianol and higher boiling point compounds are removed at the bottom.

The overhead is sent to an oil washing aystem to recover acctaldehyde and other byproducts;. Such system comprises mainly an oil decanter, an oil extractor and an oil storage tank.

Initially, the byproducts column overhead is mixed with water, in line, and the resulting mixture is sent to the oil decanter where the major part of ethanol and acctaldehyde present, is removed, and is recycled into the acctaldehyde distillation column.

The overlayin, oil feeds the oil extractor in counter-current with a flow of process water. The bottom stream of this column, which contains a large quantity of byproducts, is fed again into the washing system or returns to the distillation system and from there to the acctaldehyde converters. The rcnidual oil leaving the top of the tower is transferred to the storage tank and used as fuel.

The bottom product of the byproducts distillation column, in the form of an ethanol water solution, is fed into ^a sot of distillation columns, where alcohol is recovered and purified and water and byproducts arc discharged into the sewage or sent to the oil recovery equipment.

The recovered alcohol is let out vaporized

and serven to feed the converters. Fresh alcohol coming from the storage section is also vaporized with the recycled material replenishing the required feed volumo.

5.1.3. Production of Acetic Acid

The acetic acid process developed hy PETROQUISA consists of an improvement of the existing domestic acetic acid technologies based on the liquid phase oxidation of acotyldohydo.

A basic feuturc of this process is the utilisation of the oxidation gas (air), prior to the reaction, to recover the acetaldehyde contents of the reactor effluent. Also, the catalyst is held in a closed loop system, so reducing louses.

The unit operates at low pressure and temperature, and a special design of the reactor, sparged, allows to operate with high gas space velocity without excessive acetaldehyde entrainment.

At present, a new catalyst system is under development toward obtaining a selectivity improvement .

Figures 8 and 9 show the reaction and purification sections of the process.

Fresh high-purity ucotaldohydc, oxidation gas, recovered acetaldehyde and catalyst are continuously fed to the reactor, where oxidation takes place.

The reaction, which can be represented by

 CII_3 CHO $+$ 1/2 O_2 \longrightarrow CH₃ COOH is highly exothermic, being developed around

.
Let il regione quote della

 $+$

70 Kcal/mol. In order to remove the heat of the reaction, the substract from the reactor is circulated through outside coolers.

The liquid effuent from the reactor contains mainly acetic acid and small amounts of by-products, such as formle acid, methyl acetate and others. This stream is first stripped of its acetaldehyde contents in the acetaldehyde stripping column. The temperature and pressure in this column are set to maintain, for safety purposer, a certain amount of acetic acid in the top stream, that goes further to the reactor.

The bottoms go to the product recovery column, where the incoming air is saturated with acetic acid. This stream is condensed and the resulting condensate, crude acetic acid, is sent to the purification section.

The gaseous effluent from the reactor contains mainly unreacted acetaldehyde, carbon dioxide, nitrogen and oxigen, of which concentration is kept low to avoid fire hazards. This stream is first washed with cooled acetic acid, in the acetaldehyde absorption column, in order to remove the entrained acetaldehyde.

The acetic acid leaving this tower in the overhead stream is recuperated in the acetic acid recovery column by washing with chilled water. The vent gas is discharged into the atmosphere and the bottoms are sent to the purification section.

The purification section receives two The first one, crude acetic acid, main streams. in the condensate from the product recovery column; the second one comes from the acetic acid recovery column.

 $-34-$

This section consists of an assotropic distillation system, where othyl acetate is employed as a solvent.

Acetic ac d coming from the reaction section is fed into the azeotropic distillation column. The solvent is fed few plates below the top plate.

Water and light byproducts are entrained by ethyl acetate, and leave the column as overhead. After condensation, this stream splits into two layers. The organic layer returns to the column.

The acqueous layer is stripped, with steam, of its ethyl acetate contents, and then is reutilized. The acetic acid product is recovered as a liquid from 3 to 6 plates above bottom and, after cooling, goes into storage. The bottoms, mainly heavy byproducts, go to a film evaporator, where they are exhausted of acetic acid contents and then sent to disposal.

5.1.4. <u>Butanol and Octanol Production</u>

In Brazil, the traditional process to obtain such products from ethanol is used.

In this route, acetaldehyde (by dehydrofenation of ethanol, such as explained in item 4.1.2. above) and crotonaldehyde (by aldolization of erotonaldehyde and subsequent dehydrogenation of aldol) are initially produced.

Next follows hydrogénation of erotonaldehyde whereby the n-butanol and butyraldehyde are obtained, the latter by uncomplete hydrogénation.

The production rates of n-butanol/butyr-

-39-

aldehyde may be controlled in order to produce a larger or smaller quantity of butyraldehyde. The reaction control is done mainly by the concontration of circulating hydrogen.

If the purpose is primarily production of n-but anol, butyraldehyde may be recycled until complete extinction.

If, on the contrary, the process is intended to obtain octanol (2-ethylhexanol), hydrogenation is controlled in such a way as to maximize butyraldehyde production, which however does not prevent formation of n-butanol as a byproduct.

In this lapt case, butyraldehyde goes through an addolization and subsequent dehydration, forming 2-ethyl hexenal. The latter is hydrogenated to 2-ethyl hexanol.

Schematically, the above route may be represented as follows:

 $-$ A()--

Acetaldehyde, in the presence of small amounts of dilluted soda (10%), is fed into a column where, by condensation, it is converted into aldol. Aldolization is carried out at a temperature of 5 to 25°C and conversion of acotaldehyde by pass rangea from ⁴⁸ to GO'S. Unconverted acetaldehyde is recovered as overhead and recycled.

Aldolization product is sent to a dehydration column system to be transformed into crotonaldehyde, which is done in the presence of monosodi, um phosphate.

Bottoms of the' first column consists mainly of water eliminated during the reaction, which is recovered and reutilized in the process.

Dehydration in this column is not complote and its ove head is sent t ^a second column to proceed with the reaction. Here, unroncted acetaldehyde is further recuperated as overhead and recycled.

Reaction mixture leaving the second column is sent to a decanter to separate the crotonaldehyde and is then fed into a third column. The overhead of the latter is recycled into the second column, whereas a new stream of process water is recovered from bottoms.

Crotonaldehyde follows toward the evaporator and then to the hydrogenation furnace, where, in the presence of a nickel-chromo catalyst, it is converted into n-butanol and butyraldchydo. Temperature of the operation is 1800C and pressure 30 psig.

i

 \mathbf{I}

Reactjon product is finally split in a two rectification towers system, and butyraldehyde is obtained from overhead of the first and n-butanol from overhead of the segond tower. When the intention is to produce only n-butanol, butyraldehyde returns to the reaction for further hydrogenation to that alcohol.

Figure 11 shows, on the other hand, the flow-sheet of 2-othyl hexanol production from butyraldehyde.

Dutyraldehyde *in* fed into the aleolic condensation reactor to form 2-cthyl hcxenal. **Small quantities of C_R** isomers are also produced.

Reaction is carried out in the presence of NaOH 10%, and fresh soda, joined with recycled soda, fed into the reactor top. This is provided with ^a serpentine for internal cooling to remove the heat of reaction.

The organic layer split at the decanter of overhead *in* cooled and sent to ^a tank where it joins the organic layer split in the bottoms decanter. The acqueous layer of the overhead constitutes the soda recycle, whereas the acqueous layer of the bottoms is sent for treatment and disposal.

The organic product then follows to be washed with water in order to remove the major part of entrained soda, whence it is transferred to the first stage of the evaporator.

Hydrogen is used to lower the steam pressure of the 2-ethyl hexenal misture, in order to Minimise the product's thermal degradation.

The liquid originated from the overhead

i

of the two stages of the evaporator, rich in 2-othyl hexenal, follows to the hydrogenation section.

The steam resulting from the first stage of the evaporator, containing mainly hydrogen, is recompresscd and reintroduced into the evaporator base.

2-ethyl hexonal is combined with hydrogen and, after pre-heating by means of exchange of heat with the effluent of the primary hydrogenation reactor, is fed into it. This equipment is provided with a fixed bed, containing a nickel catalyst. Reaction is carried out in steam phase, at a temperature of 140°C and pressure of 135 psiq, and about 99% of 2-othyl hexenal is converted at this stage. The reactor's product is cooled and the uncondensed steam is recycled for further conversion.

The liquid product then goes onto the secondary hydrogenation reactor. Reaction is carried out in the liquid phase, at about 600C and 1500 psig pressure, and through it unreacted aldehydes originated from the first hydrogenation stage arc hydrogenated. Unreacted hydrogen separate from the reactor's effluents to be used as a fuel.

Hydrogénation crude product consists mainly of 2-ethy] hoxanol, and has as impurities n-butanol, aldehydes and heavy compounds.

Recuperation and purification of 2-ethyl hexanol is carried out in three distillation
columns. The first column connects is in The first column separates in the overhead the n-butanol and other light components, which may be further distilled to recover n-butanol.

> * •»«*«•

-w W. --.««*

The second column recovers 2-ethyl hexanol

•«• i

am overhead, while the bottoms go on to the next column. In the latter, the overhead, containing 2-ethyl hexanol, is recycled to the first column, while heavier products are rejected.

5.1.5. Butadieno Production

In this process, which has been used in Brasil by COPERBO, the following chemical reaction occurs :

 CH_3CHO + CH_3CH_2OH \longrightarrow $CH_2CHCH=CH_2+ H_2O$ acetaldehyde ethanol butadiene water

It is believe, however, that such reaction takes place in two phases:

(2) $CH_3CH=CICHO + CH_3CH_2OH \longrightarrow CH_2=CHCH=CH_2 + CH_3CHO + H_2O$ crotonaldehyde et]»anol butadiene acetal- water dehyde

The process' integrated flow-sheet, from ethy alcohol, is shown in Figure 12.

The first stage of the process comprises dehydrogenation of ethanol to acetic aldehyde, and is completely identical to the process shown in item 4.1.2.

The second stage involves manufacture of butadiene proper, and is examined in more detail below.

Acetic aldehyde and ethanol vapours, from distillation columns overhead of the acetaldehyde section, are mixed at a rate of 2.5 mols ethanol for ¹ mol acetic aldehyde, pre-heated and fed into

i

DURADENE FROM ETHANCE PROCESS

the butadione converters. Here they are drained through the catalytic bed (silica gel saturated with $1.75 - 2.258$ tantalum oxide), at low pressure $(3 - 8 \text{ p} \cdot \text{sig})$ and high temperature $(320 350^{\circ}C$).

Under normal operation the conversion by pass is rather low $(8 - 12)$ butadiene, in weight), and it is necessary to recover and recycle the mejor part of the unreacted feed stream.

The effluent from converters, in addition to butadiene and steam, contains byproducts which comprise from gares, mainly ethylene, up to high molecular weight hydrocarbons, carbon deposits, and some exigenated compounds such as ethyl ether, n-butanol and acetic acid.

Total efficiency in ethanol-based butadione ethanol normally ranges between 50-63%, including losses in efficiency in acetic aldebyde production and losses relating to product recuperation and iransfer.

Converters censist of several steel tule., Hilled with catalyst, closed inside a tubular furnace. The reassion is endothermic and the hopt for the reaction is provided by a Dowtherm bath, watch surrounds the tubes.

After a 80-120 hear production period, the catalyst is reactived throughout 24-48 hours by burning carbon through a controlled passage of air at $600 - 410^{O}C$.

Vapour from the converters passes through feed pro-heaters and goes into the low pressure condenser. The condensate goes to the low

 $-AH-$

pressure condénsate tank, while uncondensed vapours flow to the top of the tank through two separate strippers, are compressed and sent to the high pressure condenser.

T

The high pressure condensate and the cooled vapour feed the bottom of the primary washer, whose gaseous overhead flow to the bottom Of the secondary washer. The gaseious overhead of the latter leaves the unit to be used as a fuel. The low pressure condensate joins with the liquid bottoms of the washers and follows toward the pre-distillaticn column.

The overhead of this column, under normal operating conditions, has 80-85% butadiene and 10-15% acetaldohyde. Bottoms are taken to the byproducts distillation column, and from there goes through other stages to recuperate other useful components.

The overhead stream of the pre-distillation column goes into the base of the washing tower where the process water is introduced at the top and absorbs the acetic aldehyde and other soluble impurities. This stream is normally carried to the oil washing systems or, alternately, sent to the feed tank of the acetaldehyde column or down the drain.

Under normal running conditions, the overhead gas of the washing tower contains about 9St (in volume) butadiene. Such stream is taken to a tower, where butadiene is absorbed selectively by Chlorex (ether di (2-ethyl chlorine)), and is separated from the major part of other component gases (methane, ethylene, propylane and butane).

In the stripping column, pressure is

 $+$

 $-49-$

reduced and the butadiene-Chlorex solution is distilled to steam, to strip all hydrocarbon present. The chilled vapour is compressed and fed to the butadiene purification columns.

 \sim C

The residual gas of the absorber is recycled to recuperate butadiene and butenes, or may be sent as fuel to the Dowtherm furnaces.

The water-Chlorex solution leaving the bottom of the stripping column is sent to a distillation column to recuperate Chlorex. The latter is reutilized in the absorption tower.

The butadiene purification systems consists basically of three distillation columns in a row.

Vaporized feed, containing 96-96.5% butadiene goes in at the first column, where the butadiene goes out as overhead, while the bottoms, containing 50-75% butadiene, passes on to the second column. Here, basically, all buteno-1 is separated in the overhead so that the critical separation effected in the second column if of butene-2 from butadiene.

In the overhead from the first and second columns butadiene has a purity of about 98%, and such streams represent the final product, which 18 sent to storage.

The third column operator with a high reflux to exhaust any butadiene still present in the butene mixture leaving the bottom of the second column. Butenes and heavy residues are collected in the circulation drum, to the extent that the butadiene content reduces evenly. At last, batch distillation of the fraction of

I
I

5.2. Technical and Economical Evaluation

Tables V through XVI present a technical and economical evaluation of the units which employ the processes described above, comprising an estimate of investments, operating costs and sales price of the products, under Brazilian conditions as of September 1978.

Tables are self-explanatory in relation to criteria utilized to calculate investments and operating costs of the respective units.

The fundamental object of this review was to evaluate the impact of the subsity instituted for alcohol over final prices of the products, as well as the competitiveness of processes under consideration in relation to those of the petrochemical industry.

In this sense, two price alternative of the raw material (ethanol) were contemplated in each case:

- industrial alcohol, at effective price in domestic market (US\$293/ton ethanol);
- alcohol subsidised for the chemical industry, at a price per cubic meter equivalent to 35% of price per ton of petrochemical ethylene (US\$170/ton ethanol).

international

Moreover, since the acetaldehyde and the acetic acid are only produced in Brazil from ethyl alcohol, in addition the review of its production by a petrochemical route was done.

Table XVI1 summarizes, for each product, the sales prices estimated for the two above alternatives, comparing same with effective prices (or which could be

TABLE - V

PROLESS ECONOMICAL EVALUATION

ESTIMATED INV. STIMENT

ESTIMATED MANUEACTURING COSTS AND TRANSEER PRICE

(*)Nonauhsidized ethanol price

1

$-53-$

$TABLF = VI$

PROGECS ECONOMICAL EVALUATION

 \mathbf{I}

ESTIMATED INVESTIMENT

ESTIMATED MANUEACTURING COSTS AND TRANSFER PRICE

When you

 $\ddot{}$

 $-94-$

 $\alpha_{\rm{eff}}=2.5$ $\mathcal{L}^{\mathcal{L}}$ **TABLE-VII**

PROCESS ECONOMICAL EVALUATION

 \mathbf{u}

Acetaldehyde **PRODUCT** PROCESS Ethanol Dely Irogenaution
CAPACITY 45,000 MTY
LOCATION Brazil LOCATION
LOCATION \sim and 1st trim. 1979...

 $\ddot{}$

 \overline{a}

ELT MATED HIVETTIMENT

ESTIMATED MANUFACTURING COSTS AND TRANSFER PRICE

الوالونهاها والعا

 $\overline{}$

(*) Fuel value

 $-5¹$

TABLE - VIII

PROCESS FCONOMICAL EVALUATION

PRODUCT Acetal delivele francess Ethanol Dilydregenation CAPACITY
LOCATION
DATE 45,000 HTY Brazil
181 trim, 1979

ESTIMATED INVESTIMENT

ESTIMATED MANUFACTURING COSTS AND TRANSITR PRICE

c available.

والموا

 α , is a con-

 $\ddot{}$

 l

 -5.1

TABLE $-1x$

PROCESS ECONOMICAL EVALUATION

 \mathbf{I}

ESTIMATED INVISTIMENT

ESTIMATED MANUFACTURING COSTS AND THANSFER PRICE

(*) Brazillan market price of Petrochemical Ethylene

 $-57-$

TABLE - X

PROCESS ECONOMICAL EVALUATION

 \sim \sim

ESTIMATED INVESTIMENT

ESTIMATED MANUFACTURING COSTS AND TRANSFER PRICE

 ~ -1

(*) Transfer Price of Acetaldehyde from non- subsidited ethanol.

TABLE $-$ XI

المتفهما والمادين المساوي

ولمحاويتهم والمراجع

PROCESS FCONOMICAL EVALUATION

PRODUCT PROCESS CAPACITY LOCATION
DATE

Acetic Acid Air Oxidation 60,000 MTY Brazil
1⁸¹ trim. 1979.

 \overline{a}

ESTIMATED INVESTIMENT

ESTIMATED MANUFACTURING COSTS AND TRANSFER PRICE

(*) Transfer Pilce of Acetaldehyde from subsidized ethanol

.
Prima provincia de la

 $-59-$ الفوات الكرامي

$TABLE - XII$

PROCESS ECONOMICAL EVALUATION

 $\mathbf{v}=\mathbf{v}$, $\mathbf{v}=\mathbf{v}$

and a straight

 $\ddot{}$

ESTIMATED INVESTIMENT

ESTIMATED MANUFACTURING COSTS AND TRANSFER PRICE.

(*) Transfer Price of Acetaldehyde from Petrochemical Ethylene

$-(\theta -$

TABLE - XIII

PROCESS ECONOMICAL EVALUATION

Service Companies Adv

 \bar{a}

 2 -ethylhexanol PRODUCT PROCESS From ethanol **CAPACITY**
LOCATION
DATE Brazil
D^{6t} trim. 1979

 $\tau \rightarrow 1$

ESTIMATED INVESTIMENT

ESTIMATED MANUFACTURING COSTS AND TRANSFER PRICE

(*) Brasilian Market Price

(**) Nöngubsidized Ethanol Price

. . . .

 $-(1-$

TABLE- XIV

PROLESS ECONOMICAL EVALUATION

2-othylexanol
From ethanol **PRODUCT PROCESS** CAPACITY
LOCATION
DATF 20,000 MTY Brazil.
1^{8t} trim. 1979

ESTIMATED INVESTIMENT

ESTIMATED MANUFACTURING COSTS AND TRANSFER PRICE

TABLE - XV

PROCESS ECONOMICAL EVALUATION

From Ethanol

ESTIMATED INVESTIMENT

ESTIMATED MANUFACTURING COSTS AND TRANSFER PRICE

 $\mathcal{L}_{\mathcal{A}}$

.
Satu mumalik

(*) Brazilian Market (**) Fuel value (***)Nonsubsidized ethanol price

Service in the

ా⊃ా استشفاء فللاعتار وراسا وراسي

TABLE - XVI

PROCESS ECONOMICAL EVALUATION

والمارا والعوالق عقران ولراميز أرز

المتصفحا وسيروز والراز

وتصامر الاستطاعين مواديسها للأرباب الرازيد

ESTIMATED IF /ESTIMENT

ESTIMATED MANUFACTURING COSTS AND TRANSFER PRICE

(*) Brazilian market price (**) Fuel Price (***)Non-subsidized ethanol price

TABI ANI

PRICES OF CHENTICALS OBTAINED FROM ETHANOL ASD PETROCHENICAL FELDSTOCK

BRAZILIAN CONDITIONS - 1st QUARTER 1979

* Brazilian market price

** Estimated transfer price according to the economical evaluation

 l

 $\begin{matrix} \bullet & \bullet \\ \bullet & \bullet \end{matrix}$

charged) in the country for the petrochemical originated product.

Such dùta indicate that, at the level of capacity taken into consideration, ethanol-chemical processes to manufacture acetaldehyde and acetic acid — widely used in the country — are competitive as compared to alternative petrochemical routes, even if considering ethanol at current market prices.

On the other hand, the saving with the production of octhanol fron ot.hyl alcohol is highly dependent on the cost of raw material and, therefore, the use of a subsidised price for the alcohol seems to be essential to ensure competitiveness of such process.

As regards ethylene, the subsidy taken as reference in the economic evaluation seems insufficient to render the price in the process from alcohol equivalent to that of the petrochemical product. By reviewin the text of the National Alcohol Plan it appears to be possible to apply for a larger subsidy, which, to reach the intended equivalence, would, in this instance, be on the basis of 25% of present petrochemical ethylene prices.

Where butadiene is concerned, results obtained indicate tnat its production by ethanol-chemical route is far from being competitive as compared to the processes to obtain this basic product in naphtha pyrolyeis complexes.

~-*6

6. CONCLUSIONS

Petroleum supply difficulties bring back the etha.iol-chemical industry as a sound technological alternative to obtain certain products, nowadays predominantly petrochemical.

Brazil's tendency toward the use of that productive route is maybe unparalleled in the world, in view of its territorial extent and propitious conditions for the culture of sugar cane, manioc and other agricultural products applicable in alcoholic fermentation.

On the other hand, the National Alcohol Plan has created favorable conditions to strengthen such activity in the country, amongst which should be pointed out the assurance of a regular supply of raw material, at a reasonable price, and encouragement to development of technology in the agricultural and industrial sectors.

Thus, the experience in manufacturing ethanol-chemical derivatives, acquired throughout many years by the producing companies, permits to consider its technology (billy absorbed in Brazil.

Such experience is now being utilized, in the scope of companies and research centers, to modernize numufactuaing processes, with a view to its prompt use in larger and more economical units.

Moreover, in view of the present stage of the art - whether agricultural, or industrial - of manufacturing ethyl alcohol, it may be anticipated that, shortly, this raw material will continue to be produced in the country at high costs. Therefore, the subsidy instituted by the Government should continue for some years, in order to ensure competitiveness of ethanol-chemical processes when compared to the petrochemical industry processes.

At medium and long term, as a result of

سرار ام

PROALCOOL efforts and the technological development programs under way, which lead to higher agricultural productivity indices and more competitive industrial processes, it is expected that ethanol-chemical industry in Brazil will at last reach its maturity, becoming self-sufficient.

 274
BIBLIOGRAPHY

- 1. ABIQUIM A INDÚSTRIA OBÍMICA BRASTLEIRA. SÃO Paulo, 1978. 255p.
- 2. CASAL1, E. O PROGRAMA NACIONAL DO ALCOOL. in Brasil Agucareiro, Rio de Janeiro, 91 (4) p. 35-42, April 1978.
- 3. DANTAS, Romen Bôto ALCOOL ETTLACO. A CRISE DE EDERGIA E A INDUSTRIA OUIMICA. Brazilian Petrochemical Congress. Rio de Jancivo, 1976. 5v. V.4, no. 309.
- 4. FRAGALE, ROGÉTIO da Silva PRODUÇÃO DE ETILEMO A PARTIR DE Brazilian Petrochemical Congrenss, Rio de Ja-ALCOOL. neiro, 1976. 5v. V.5, no. 351.
- 5. RODRIGUES, Egberto da Gama EXPERIENCIA DE UM INVESTIMEN-TO FRANCES NO BRASIL. French Brazilion Simposium on Petrochemical Industry. Salvador, Mines and Energy Sec., 1972. $425p_+$, p. $405-14$.
- 6. SANTA ROSA, Jaime da Nóbrega A THIOGETRIA QUIMICA NO ESTADO DE SÃO PAULO. Rio de Janeiro, Ed. Bornol, 1958. 182p.
- 7. SILVA FURIO, Amiliaar Petefra da O ETANOL EM SUPETITUIQAO AS MATERIAS-PRIMAS PETROQUÍBICAS. Industrial Technology Week. Petrochemical Industry. Salvador, MIC.S"L. 1976.
- 8. VILLA, Roberto DESCRET TIMESTO DE UL NOVO PROCESSO PARA A PRODUÇÃO LE ESTE O A PARTIR DE ALCOOL. 11 Latin-American Petrochemical Compensor. Cancun - México, 1978.
- Industein1 Technology Week ETANOL: COMBUSTIVEL E MATÉRIA-PRIMA. ETABOL COMO MATERIA-PRIMA NA INDUSTRIA QUÍMICA. Rio de Janeiro, MIC.STI., 1976. 469p. Pannel V, p. 261-313.
- I Simposium on Alcohol Production in Northeast Brazil. Fortaleza. Banco do Nordeste do Brasil, 1977. 452p.

 -1

30.02.04