

ALKYLATION PROCESS USING AN ALKYL HALIDE PROMOTED
IONIC LIQUID CATALYST

FIELD OF THE INVENTION

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The present invention relates to a process for the alkylation of light isoparaffins with olefins using a catalyst comprising an ionic liquid and an alkyl halide.

BACKGROUND OF THE INVENTION

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In general, conversion of light paraffins and light olefins to more valuable cuts is very lucrative to the refining industries. This has been accomplished by alkylation of paraffins with olefins, and by polymerization of olefins. One of the most widely used processes in this field is the alkylation of isobutane with C₃ to C₅ olefins to make gasoline cuts with high octane number using sulfuric and hydrofluoric acids. This process has been used by refining industries since the 1940's. The process was driven by the increasing demand for high quality and clean burning high-octane gasoline.

Alkylate gasoline is a high quality and efficient burning gasoline that constitutes about 14% of the gasoline pool. Alkylate gasoline is typically produced by alkylating refinery isobutane with low-end olefins (mainly butenes). Currently, alkylates are produced by using HF and H₂SO₄ as catalysts. Although these catalysts have been successfully used to economically produce the best quality alkylates, the need for safer and environmentally friendlier catalysts systems has become an issue to the industries involved.

The quest for an alternative catalytic system to replace the current environmentally unfriendly catalysts has been the subject of varying research groups in both academic and industrial institutions. Unfortunately, thus far, no viable replacement to the current processes has been put into practice at commercial refineries.

Ionic liquids are liquids that are composed entirely of ions. The so-called "low temperature" Ionic liquids are generally organic salts with melting points under 100

degrees C, often even lower than room temperature. Ionic liquids may be suitable for example for use as a catalyst and as a solvent in alkylation and polymerization reactions as well as in dimerization, oligomerization acetylation, metatheses, and copolymerization reactions.

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One class of ionic liquids is fused salt compositions, which are molten at low temperature and are useful as catalysts, solvents and electrolytes. Such compositions are mixtures of components which are liquid at temperatures below the individual melting points of the components.

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Ionic liquids can be defined as liquids whose make-up is entirely comprised of ions as a combination of cations and anions. The most common ionic liquids are those prepared from organic-based cations and inorganic or organic anions. The most common organic cations are ammonium cations, but phosphonium and sulphonium cations are also frequently used. Ionic liquids of pyridinium and imidazolium are perhaps the most commonly used cations. Anions include, but not limited to, BF_4^- , PF_6^- , haloaluminates such as Al_2Cl_7^- and Al_2Br_7^- , $[(\text{CF}_3\text{SO}_2)_2\text{N}]^-$, alkyl sulphates (RSO_3^-), carboxylates (RCO_2^-) and many other. The most catalytically interesting ionic liquids for acid catalysis are those derived from ammonium halides and Lewis acids (such as AlCl_3 , TiCl_4 , SnCl_4 , FeCl_3 ...etc). Chloroaluminate ionic liquids are perhaps the most commonly used ionic liquid catalyst systems for acid-catalyzed reactions.

Examples of such low temperature ionic liquids or molten fused salts are the chloroaluminate salts. Alkyl imidazolium or pyridinium chlorides, for example, can be mixed with aluminum trichloride (AlCl_3) to form the fused chloroaluminate salts. The use of the fused salts of 1-alkylpyridinium chloride and aluminum trichloride as electrolytes is discussed in U.S. Patent No. 4,122,245. Other patents which discuss the use of fused salts from aluminum trichloride and alkylimidazolium halides as electrolytes are U.S. Patent Nos. 4,463,071 and 4,463,072.

U.S. Patent No. 5,104,840 describes ionic liquids which comprise at least one alkylaluminum dihalide and at least one quaternary ammonium halide and/or at least

one quaternary ammonium phosphonium halide; and their uses as solvents in catalytic reactions.

U.S. Patent No. 6,096,680 describes liquid clathrate compositions useful as reusable
5 aluminum catalysts in Friedel-Crafts reactions. In one embodiment, the liquid
clathrate composition is formed from constituents comprising (i) at least one
aluminum trihalide, (ii) at least one salt selected from alkali metal halide, alkaline
earth metal halide, alkali metal pseudohalide, quaternary ammonium salt, quaternary
phosphonium salt, or ternary sulfonium salt, or a mixture of any two or more of the
10 foregoing, and (iii) at least one aromatic hydrocarbon compound.

Other examples of ionic liquids and their methods of preparation may also be found in
U.S. Patent Nos. 5,731,101; 6,797,853 and in U.S. Patent Application Publications
2004/0077914 and 2004/0133056.

15 In the last decade or so, the emergence of chloroaluminate ionic liquids sparked some
interest in $AlCl_3$ -catalyzed alkylation in ionic liquids as a possible alternative. For
example, the alkylation of isobutane with butenes and ethylene in ionic liquids has
been described in U.S. Patent Nos. 5,750,455; 6,028,024; and 6,235,959 and open
20 literature (Journal of Molecular Catalysis, 92 (1994), 155-165;
“Ionic Liquids in Synthesis”, P. Wasserscheid and T. Welton (eds.), Wiley-VCH
Verlag, 2003, pp 275).

Aluminum chloride-catalyzed alkylation and polymerization reactions in ionic liquids
25 may prove to be commercially viable processes for the refining industry for making a
wide range of products. These products range from alkylate gasoline produced from
alkylation of isobutane and isopentane with light olefins, to diesel fuel and lubricating
oil produced by alkylation and polymerization reactions.

SUMMARY OF THE INVENTION

The present invention relates to an alkylation process comprising contacting a first hydrocarbon feed comprising at least one olefin having from 2 to 6 carbon atoms and
5 a second hydrocarbon feed comprising at least one isoparaffin having from 3 to 6 carbon atoms with a catalyst under alkylation conditions, said catalyst comprising a mixture of at least one acidic ionic liquid and at least one alkyl halide, wherein the alkyl halide is produced by reacting at least a portion of the first hydrocarbon feed with a hydrogen halide under hydrohalogenation conditions to convert at least a
10 portion of the olefins contained in the first hydrocarbon feed to the alkyl halide.

DETAILED DESCRIPTION

The present invention relates to an alkylation process comprising contacting a hydrocarbon mixture comprising at least one olefin having from 2 to 6 carbon atoms and at least one isoparaffin having from 3 to 6 carbon atoms with a catalyst under alkylation conditions, said catalyst comprising a mixture of at least one acidic ionic liquid and at least one alkyl halide.

One component of a feedstock to the process of the present invention is at least one olefin having from 2 to 6 carbon atoms. This component may, for example, be any refinery hydrocarbon stream which contains olefins.

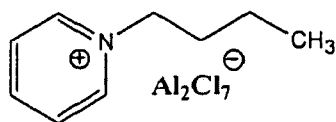
Another component of a feedstock to the process of the present invention is at least one isoparaffin having from 3 to 6 carbon atoms. This component may, for example, be any refinery hydrocarbon stream which contains isoparaffins.

The processes according to the present invention are not limited to any specific feedstocks and are generally applicable to the alkylation of C₃-C₆ isoparaffins with C₂-C₆ olefins from any source and in any combination.

In accordance with the present invention, a mixture of hydrocarbons as described above is contacted with a catalyst under alkylation conditions. A catalyst in accordance with the present invention comprises at least one acidic ionic liquid and at least one alkyl halide. The present process is being described and exemplified with reference certain specific ionic liquid catalysts, but such description is not intended to limit the scope of the invention. The processes described may be conducted using any acidic ionic liquid catalysts by those persons having ordinary skill based on the teachings, descriptions and examples included herein.

The specific examples used herein refer to alkylation processes using ionic liquid systems, which are amine-based cationic species mixed with aluminum chloride. In such systems, to obtain the appropriate acidity suitable for the alkylation chemistry, the ionic liquid catalyst is generally prepared to full acidity strength by mixing one

molar part of the appropriate ammonium chloride with two molar parts of aluminum chloride. The catalyst exemplified for the alkylation process is a 1-alkyl-pyridinium chloroaluminate, such as 1-butyl-pyridinium heptachloroaluminate.

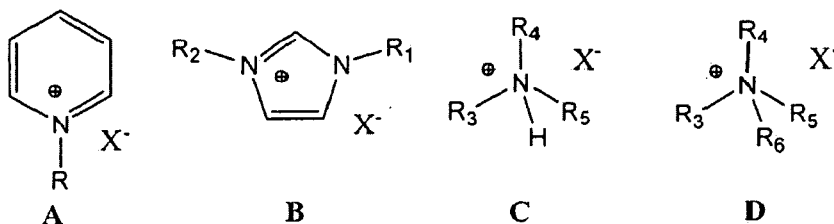


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1-Butyl-pyridinium heptachloroaluminate

As noted above, the acidic ionic liquid may be any acidic ionic liquid. In one embodiment, the acidic ionic liquid is a chloroaluminate ionic liquid prepared by mixing aluminum trichloride (AlCl_3) and a hydrocarbyl substituted pyridinium halide, a hydrocarbyl substituted imidazolium halide, trialkylammonium hydrohalide or tetraalkylammonium halide of the general formulas A, B, C and D, respectively,

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where $\text{R}=\text{H}$, methyl, ethyl, propyl, butyl, pentyl or hexyl group and X is a halide and preferably a chloride, and R_1 and $\text{R}_2=\text{H}$, methyl, ethyl, propyl, butyl, pentyl or hexyl group and where R_1 and R_2 may or may not be the same, and R_3 , R_4 , and R_5 and $\text{R}_6=\text{methyl}$, ethyl, propyl, butyl, pentyl or hexyl group and where R_3 , R_4 , R_5 and R_6 may or may not be the same. .

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The acidic ionic liquid is preferably selected from the group consisting of 1-butyl-4-methyl-pyridinium chloroaluminate, 1-butyl-pyridinium chloroaluminate, 1-butyl-3-methyl-imidazolium chloroaluminate and 1-H-pyridinium chloroaluminate .

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In a process according to the invention an alkyl halide is used as a promoter.

The alkyl halide is produced in accordance with the invention by reacting at least a portion of the olefinic feed with a hydrogen halide under hydrohalogenation conditions to convert at least a portion of the olefins to the alkyl halide. This is accomplished in accordance with the present invention by reacting at least a portion of the olefin feed stream with a hydrohalide under hydrohalogenation conditions and adding the resulting alkyl halide to the alkylation zone. In other words, the alkyl halide is generated from the olefin feed. For example, one can take a slip-stream of the olefin-containing refinery feedstock and react that with HCl under conditions which would convert the olefins in the slip-stream into alkyl halides such as sec-butyl and t-butyl chloride. This alkyl halide containing stream can be injected into the catalyst stream being injected into the alkylation reactor.

Hydrohalogenation of olefins is well known. U.S. Patent No. 5,831,137 describes a process to make t-butyl chloride by passing a mixture of anhydrous HCl and isobutene (1.2 molar ratio) in the gas phase at atmospheric pressure through concentrated aqueous HCl maintained at 60 °C. The product t-butyl chloride is condensed from the vapor phase effluent of the reactor. Other patents (e.g. FR 2,300,751) describe the reaction of anhydrous HCl and butenes to make t-butyl chloride in the presence of Lewis acid catalysts such as ZnCl₂ using t-butyl chloride as a solvent at atmospheric pressure and -25 to +50 °C. U.S. Patent No. 2,434,094 describes the reaction of anhydrous HCl and butenes in the gas phase in the presence of an inert diluent to make t-butyl chloride in the presence of Lewis acid catalysts on inert supports. Operating temperatures were below 300°F to avoid decomposition of the alkyl chloride. U.S. Patent No. 2,418,093 contains a good description of the general chemistry. Apparently isobutene reacts selectively to give t-butyl chloride in the presence of small amounts of butenes. However, some of the patents above address the selective synthesis of t-butyl chloride in the presence of excess linear butenes.

The alkyl halide acts to promote the alkylation by reacting with aluminum chloride to form the prerequisite cation ions in similar fashion to the Friedel-Crafts reactions. The alkyl halides that may be used include alkyl bromides, alkyl chlorides and alkyl iodides. Preferred are isopentyl halides, isobutyl halides, butyl halides, propyl halides

and ethyl halides. Alkyl chloride versions of these alkyl halides are preferable when chloroaluminate ionic liquids are used as the catalyst systems. Other alkyl chlorides or halides having from 1 to 8 carbon atoms may be also used. The alkyl halides may be used alone or in combination.

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For chloroaluminate ionic liquids, the alkyl halide is preferably an alkyl chloride such as ethyl chloride, tertiary butyl chloride, isopentyl chloride or butyl chlorides. The alkyl chlorides of choice are those derived from the isoparaffin and olefins used in a given alkylation reaction. For the alkylation of isobutane with butenes in chloroaluminate ionic liquids, for example, the preferable alkyl halides would be 1-butyl chloride, 2-butyl chloride or tertiary butyl chloride or a combination of these chlorides. Most preferably, the alkyl chloride is a derivative of the olefin stream to, in theory, invoke hydride transfer and the participation of the isoparaffin. The alkyl halides are used in catalytic amounts. Ideally, the amounts of the alkyl halides should be kept at low concentrations and not exceed the molar concentration of the catalyst AlCl_3 . The amounts of the alkyl halides used may range from 0.001 mol%-100 mol% of the Lewis acid AlCl_3 . Concentrations of the alkyl halides in the range of 0.001 mol%-10 mol% of the AlCl_3 are preferable in order to keep the acidity of the catalyst at the desired performing capacity. Also, the amount of the alkyl halide should be proportional to the olefin and not exceed the molar concentration of the olefin.

Without being bound to any theory, when ethyl chloride, for example is added to acidic chloroaluminate ionic liquids, ethyl chloride reacts with AlCl_3 to form tetrachloroaluminate (AlCl_4^-) and ethyl cation. Hydride shift from the isoparaffin (isopentane or isobutane) to the generated ethyl cation leads to the tertiary cation which propagates the inclusion of the isoparaffin in the reaction and, hence, the alkylation pathway.

A metal halide may be employed to modify the catalyst activity and selectivity. The metal halides most commonly used as inhibitors/modifiers in aluminum chloride-catalyzed olefin-isoparaffin alkylations include NaCl , LiCl , KCl , BeCl_2 , CaCl_2 , BaCl_2 , SrCl_2 , MgCl_2 , PbCl_2 , CuCl , ZrCl_4 and AgCl , as described by Roebuck and

Evering (Ind. Eng. Chem. Prod. Res. Develop., Vol. 9, 77, 1970). Preferred metal halides are CuCl, AgCl, PbCl₂, LiCl, and ZrCl₄.

5 HCl or any Broensted acid may be employed as co-catalyst to enhance the activity of the catalyst by boosting the overall acidity of the ionic liquid-based catalyst. The use of such co-catalysts and ionic liquid catalysts that are useful in practicing the present invention is disclosed in U.S. Published Patent Application Nos. 2003/0060359 and 2004/0077914. Other co-catalysts that may be used to enhance the activity include IVB metal compounds preferably IVB metal halides such as ZrCl₄, ZrBr₄, TiCl₄,
10 TiCl₃, TiBr₄, TiBr₃, HfCl₄, HfBr₄ as described by Hirschauer et al. in U.S. Patent No. 6,028,024.

Due to the low solubility of hydrocarbons in ionic liquids, olefins-isoparaffins alkylation, like most reactions in ionic liquids is generally biphasic and takes place at
15 the interface in the liquid state. The catalytic alkylation reaction is generally carried out in a liquid hydrocarbon phase, in a batch system, a semi-batch system or a continuous system using one reaction stage as is usual for aliphatic alkylation. The isoparaffin and olefin can be introduced separately or as a mixture. The molar ratio between the isoparaffin and the olefin is in the range 1 to 100, for example,
20 advantageously in the range 2 to 50, preferably in the range 2 to 20. In a semi-batch system the isoparaffin is introduced first then the olefin, or a mixture of isoparaffin and olefin. Catalyst volume in the reactor is in the range of 2 vol% to 70 vol%, preferably in the range of 5 vol% to 50 vol%. Vigorous stirring is desirable to ensure good contact between the reactants and the catalyst. The reaction temperature can be
25 in the range -40°C to +150°C, preferably in the range -20°C to +100°C. The pressure can be in the range from atmospheric pressure to 8000 kPa, preferably sufficient to keep the reactants in the liquid phase. Residence time of reactants in the vessel is in the range a few seconds to hours, preferably 0.5 min to 60 min. The heat generated by the reaction can be eliminated using any of the means known to the skilled person.
30 At the reactor outlet, the hydrocarbon phase is separated from the ionic phase by decanting, then the hydrocarbons are separated by distillation and the starting isoparaffin which has not been converted is recycled to the reactor.

Typical alkylation conditions may include a catalyst volume in the reactor of from 1 vol% to 50 vol%, a temperature of from -10°C to +100°C, a pressure of from 300 kPa to 2500 kPa, an isopentane to olefin molar ratio of from 2 to 8 and a residence time of 5 min to 1 hour.

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In one embodiment of a process according to the present invention, high quality gasoline blending components of low volatility are recovered from the alkylation zone. Those blending components are then preferably blended into gasoline.

- 10 The following Examples are illustrative of the present invention, but are not intended to limit the invention in any way beyond what is contained in the claims which follow.

EXAMPLES

Example 1 Preparation of Fresh 1-Butylpyridinium Chloroaluminate Ionic Liquid Catalyst A (Fresh IL A)

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1-butyl-pyridinium chloroaluminate is a room temperature ionic liquid prepared by mixing neat 1-butyl-pyridinium chloride (a solid) with neat solid aluminum trichloride in an inert atmosphere. The syntheses of butylpyridinium chloride and the corresponding 1-butyl-pyridinium chloroaluminate are described below.

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In a 2-L Teflon-lined autoclave, 400 gm (5.05 mol.) anhydrous pyridine (99.9% pure purchased from Aldrich) were mixed with 650 gm (7 mol.) 1-chlorobutane (99.5% pure purchased from Aldrich). The autoclave was sealed and the neat mixture was allowed to stir at 125°C under autogenic pressure over night. After cooling off the autoclave and venting it, the reaction mix was diluted and dissolved in chloroform and transferred to a three liter round bottom flask. Concentration of the reaction mixture at reduced pressure on a rotary evaporator (in a hot water bath) to remove excess chloride, un-reacted pyridine and the chloroform solvent gave a tan solid product. Purification of the product was done by dissolving the obtained solids in hot acetone and precipitating the pure product through cooling and addition of diethyl ether. Filtering and drying under vacuum and heat on a rotary evaporator gave 750 gm (88% yields) of the desired product as an off-white shiny solid. ¹H-NMR and ¹³C-NMR were consistent with the desired 1-butyl-pyridinium chloride and no impurities were observed.

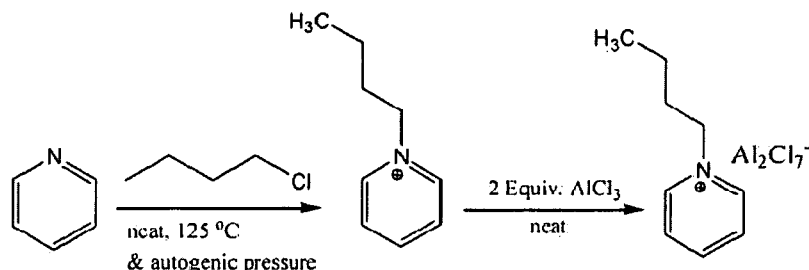
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1-butylpyridinium chloroaluminate was prepared by slowly mixing dried 1-butylpyridinium chloride and anhydrous aluminum chloride (AlCl₃) according to the following procedure. The 1-butylpyridinium chloride (prepared as described above) was dried under vacuum at 80°C for 48 hours to get rid of residual water (1-butylpyridinium chloride is hygroscopic and readily absorbs water from exposure to air). Five hundred grams (2.91 mol.) of the dried 1-butylpyridinium chloride were transferred to a 2-Liter beaker in a nitrogen atmosphere in a glove box. Then, 777.4 gm (5.83 mol.) of anhydrous powdered AlCl₃ (99.99% from Aldrich) were added in

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small portions (while stirring) to control the temperature of the highly exothermic reaction. Once all the AlCl_3 was added, the resulting amber-looking liquid was left to gently stir overnight in the glove box. The liquid was then filtered to remove any un-dissolved AlCl_3 . The resulting acidic 1-butyl-pyridinium chloroaluminate was

5 used as the catalyst for the alkylation of isopentane with ethylene.



10 Example 2 Alkylation of isoPentane with Ethylene without the presence of a promoter

A 300 cc autoclave was charged with 40 gm of ionic liquid catalyst, 100 gm anhydrous isopentane and 10 gm ethylene. The reaction was then stirred at ~1200 rpm and heated to 50°C at autogenic pressures. The starting pressure was 288 psi. The

15 reaction was allowed to run until the pressure dropped down into the single digits range (5 psi in this case after 28 minutes reaction time). In the case of slow going reaction, the reaction was allowed to run for 1hr long. At the end of the reaction, the reactor was vented out and a gas sample was checked by GC for ethylene

20 concentration. The liquid reaction mixture was allowed to settle into 2 phases. The organic phase was decanted and analyzed for product distribution by GC analysis. The reaction results are shown in Table 1.

Example 3 Alkylation of isoPentane with Ethylene in the Presence of HCl as Co-catalyst

Table 1 below shows the results of the alkylation of ethylene with isopentane in the presence of ethyl chloride and in the presence of isopentyl chloride. The alkylation of isopentane with ethylene was done according to the following procedure.

A 300 cc autoclave was charged with 40 gm of ionic liquid catalyst, 100 gm anhydrous isopentane, 10 gm ethylene, and 0.35 gm of anhydrous HCl. The reaction was then stirred at ~1200 rpm and heated to 50°C at autogenic pressures. The starting pressure was 320 psi. The reaction was allowed to run until the pressure dropped down into the single digits range (9 psi in this case after 4 minutes reaction time). In the case of slow going reaction, the reaction was allowed to run for 1hr long. At the end of the reaction, the reactor was vented out and a gas sample was checked by GC for ethylene concentration. The liquid reaction mixture was allowed to settle into 2 phases. The organic phase was decanted and analyzed for product distribution by GC analysis. The reaction results are shown in Table 1.

Example 4 Alkylation of isoPentane with Ethylene in the Presence of Chloroethane as a Promoter

The reaction described in Example 3 was repeated but chloroethane ($\text{CH}_3\text{CH}_2\text{Cl}$) was added in place of hydrochloric acid (HCl). The reaction was run on 100 gm isopentane, 10 gm ethylene and 0.9 gm chloroethane in 40 gm 1-butylpyridinium chloroaluminate ionic liquid catalyst. The use of chloroethane was as effective as using HCl in the reaction. Table 1 summarizes the reaction results and conditions.

Example 5 Alkylation of Isopentane with Ethylene in the Presence of 2-Chloro-2-methylButane (Isopentyl Chloride) as Promoter

The reaction described in Example 4 was repeated but isopentyl chloride was added in place of chloroethane. The reaction was run on 100 gm isopentane, 10 gm ethylene and 1.2 gm of isopentyl chloride in 40 gm 1-butylpyridinium chloroaluminate ionic

liquid catalyst. Isopentyl chloride appeared to be more effective than chloroethane and HCl in the alkylation of isopentane with ethylene. The reaction was extremely exothermic and there was no need to raise the reaction temperature to 50°C (reaction temperature). The pressure dropped instantaneously to the single digit mark from
 5 starting pressure of 337 psi. Table 1 summarizes the reaction results and conditions.

As shown in Table 1, when ethyl chloride or isopentyl chloride was added, the reaction time was noticeably shortened. The alkylation in the presence of isopentyl chloride was much quicker (almost instantaneous). The reactions were highly
 10 exothermic and there was no need to heat them up. The experimental results clearly indicate the very profound effect the addition of alkyl chlorides has on the progress of alkylations in ionic liquids.

	No HCl or R-Cl	W/HCl	W/Ethyl-Cl	W/Isopentyl-Cl
Start Pressure	288 psig	240 psig	331 psig	337 psig
End Pressure	5 psig	11 psig	9 psig	7
ReactionTime	28 min	4 min.	6 min.	2 min.
% Selectivity				
C3-	0.04	0	0.01	0
C4	0.86	1.88	1.93	4.97
C5	67.78	67.7	62.79	65.53
C6	1.14	2.6	2.44	5.19
C7	22.33	19.32	22.53	14.78
C8	2.94	3.25	3.68	3.27
C9	2.41	2.12	2.93	2.65
C10	1.5	1.59	1.78	1.64
C11	0.5	0.8	0.93	0.95
C12+	0.5	0.75	0.98	1.02
Total	100	100	100	100

Table 1

The alkylations in the previous Examples were done using pure isopentane feeds:

Table 2 shows a comparison between different catalyst schemes in the alkylation of refinery pentanes with ethylene using HCl, water or ethyl chloride as promoters.

Analysis of the refinery pentanes showed the feed stock to contain 86.4% iso-pentane,

5 8% n-pentane, 0.9% n-butane, 3.4% C₆s-C₉s and 0.2% olefins (C₄ and C₅ olefins).

The refinery pentane steam also contained 88ppm sulfur (as mercaptans) and 0.4ppm nitrogen. The reactions were done as described in Examples 6, 7, 8 and 9.

10 Example 6 Alkylation of Refinery Pentanes with Ethylene in 1-Butylpyridinium Chloroaluminate-No Promoters

The refinery isopentane feed with specs described earlier was dried with over a molecular sieve to remove any residual water. Then, 101 gm of the dried feed was with 10 gm of pure ethylene in 42 gm ionic liquid catalyst according to the procedure
15 described in Example 1. Table 2 summarizes the reaction and the results.

Example 7 Alkylation of Refinery Pentanes with Ethylene in 1-Butylpyridinium Chloroaluminate-with HCl as Promoter

20 Using the procedure described in Example 3, dried 101 gm of refinery isopentane feed with the specs described earlier was alkylated with 10 gm of pure ethylene in 42 gm ionic liquid catalyst in the presence of 0.6 gm HCl. Table 2 summarizes the reaction and the results.

25 Example 8 Alkylation of Refinery Pentanes with Ethylene in 1-Butylpyridinium Chloroaluminate-with H₂O as Promoter

Using the procedure described in Example 3, 101 gm of the dried refinery isopentane feed with the specs described earlier was alkylated with 10 gm of pure ethylene in 42
30 gm ionic liquid catalyst 0.1 gm water. Table 2 summarizes the reaction and the results.

Example 9 Alkylation of Refinery Pentanes with Ethylene in 1-Butylpyridinium Chloroaluminate-with Ethyl Chloride as Promoter

Using the procedure described in Example 3, 101 gm of the dried refinery isopentane
 5 feed with the specs described earlier was alkylated with 10 gm of pure ethylene in 42 gm ionic liquid catalyst 1gm of ethyl chloride. Table 2 summarizes the reaction and the results.

Table 2 summarizes the results of alkylations of refinery isopentane feed with pure
 10 ethylene described in Examples 6, 7, 8 and 9.

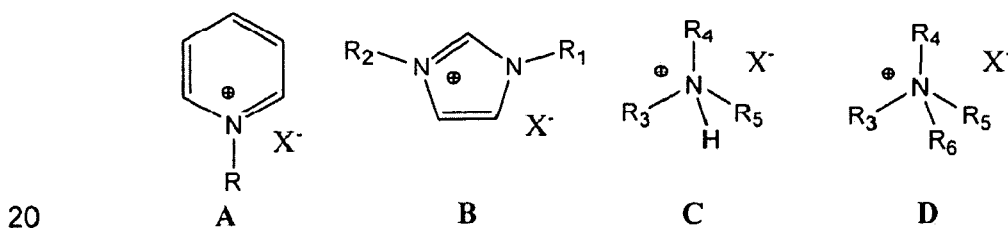
	Ref. Feed No HCl or R-Cl	Ref feed W/HCl	Ref Feed W/Ethyl-Cl	Ref Feed W/H2O
Start Pressure	226 psi	249 psi	295 psig	313 psi
End Pressure	104psi	10 psi	13 psig	15 psi
Reaction Time	64 min.	19 min.	24 min.	28 min.
%Selectivity.				
C3-	0.21	0.07	0.05	0.19
C4	0.77	1.19	1.39	1.27
C5	81.34	69.35	62.69	68.93
C6	2.82	3.06	3.87	2.97
C7	8.74	18.36	21.03	18.18
C8	3	3.8	5.05	3.93
C9	1.41	2.02	2.55	2.14
C10	0.82	1.26	1.70	1.24
C11	0.37	0.42	0.83	0.53
C12+	0.53	0.48	0.84	0.61
Total	100	100	100	100

Table 2(Alkylation of Refinery Feed in Butylpyridinium Chloride-2AlCl₃/EtCl)

There are numerous variations on the present invention which are possible in light of the teachings and supporting examples described herein. It is therefore understood that within the scope of the following claims, the invention may be practiced otherwise than as specifically described or exemplified herein.

WHAT IS CLAIMED IS:

1. An alkylation process comprising contacting a first hydrocarbon feed comprising at least one olefin having from 2 to 6 carbon atoms and a second hydrocarbon feed comprising at least one isoparaffin having from 3 to 6 carbon atoms with a catalyst under alkylation conditions, said catalyst comprising a mixture of at least one acidic ionic liquid and at least one alkyl halide, wherein the alkyl halide is produced by reacting at least a portion of the first hydrocarbon feed with a hydrogen halide under hydrohalogenation conditions to convert at least a portion of the olefins contained in the first hydrocarbon feed to the alkyl halide.
2. A process according to claim 1, where the acidic ionic liquid is a chloroaluminate ionic liquid prepared by mixing aluminum trichloride (AlCl_3) and a hydrocarbyl substituted pyridinium halide, a hydrocarbyl substituted imidazolium halide, trialkylammonium hydrohalide or tetraalkylammonium halide of the general formulas A, B, C and D, respectively,



where $R = \text{H}$, methyl, ethyl, propyl, butyl, pentyl or hexyl group and X is a halide and preferably a chloride, and R_1 and $R_2 = \text{H}$, methyl, ethyl, propyl, butyl, pentyl or hexyl group and where R_1 and R_2 may or may not be the same, and R_3 , R_4 , and R_5 and $R_6 = \text{methyl, ethyl, propyl, butyl, pentyl or hexyl group}$ and where R_3 , R_4 , R_5 and R_6 may or may not be the same.

3. A process according to claim 2, wherein the acidic ionic liquid is selected from the group consisting of 1-butyl-4-methyl-pyridinium chloroaluminate

(BMP), 1-butyl-pyridinium chloroaluminate (BP), 1-butyl-3-methyl-imidazolium chloroaluminate BMIM) and 1-H-pyridinium chloroaluminate (HP).

- 5 4. A process according to claim 1, wherein the isoparaffin is selected from the group consisting of isobutane, isopentanes and mixtures thereof.
5. A process according to claim 1, wherein the olefin is selected from the group consisting of ethylene, propylene, butylenes, pentenes and mixtures thereof.
- 10 6. A process according to claim 1, wherein the alkylation conditions include a catalyst volume in the reactor of from 1 vol% to 50 vol%, a temperature of from -10°C to 100°C, a pressure of from 300 kPa to 2500 kPa, an isopentane to olefin molar ratio of from 2 to 8 and a residence time of 1 minute to 1 hour.
- 15 7. A process according to claim 1, further comprising recovering high quality gasoline blending components of low volatility.
8. A process according to claim 1, wherein at least 50% of the olefins contained
- 20 in the first hydrocarbon feed are converted to the alkyl halide.
9. A process according to claim 1, wherein the alkyl halide has from 1 to 8 carbon atoms
- 25 10. A process according to claim 9, where the alkyl halide is selected from the group consisting of methyl halide, ethyl halide, propyl halide, 1-butyl halide, 2-butyl halide, tertiary butyl halide, pentyl halides, isopentyl halide, hexyl halides, isohexyl halides, heptyl halides, isoheptyl halides, octyl halides and isooctyl halides.
- 30 11. A process according to claim 8, where the alkyl halide is selected from the group consisting of alkyl bromides, alkyl iodides, and alkyl chlorides.

12. In an alkylation process in which at least one olefin having from 2 to 6 carbon atoms and at least one isoparaffin having from 3 to 6 carbon atoms are contacted in an alkylation zone under alkylation conditions with a catalyst comprising an acidic ionic liquid and an alkyl halide, the improvement comprising reacting a portion of said at least one olefin with a hydrohalide under hydrohalogenation conditions to convert more than 50% of the olefins contained therein to alkyl halides and adding the resulting alkyl halide rich stream into the alkylation zone.
- 10 13. A process according to claim 12, wherein the acidic ionic liquid is selected from the group consisting of 1-butyl-4-methyl-pyridinium chloroaluminate (BMP), 1-butyl-pyridinium chloroaluminate (BP), 1-butyl-3-methyl-imidazolium chloroaluminate BMIM) and 1-H-pyridinium chloroaluminate (HP).
- 15 14. A process according to claim 12, wherein the hydrohalide is anhydrous HCl.
15. A process according to claim 12, wherein the olefin is selected from the group consisting of ethylene, propylene, butylenes, pentenes and mixtures thereof.
- 20 16. A process according to claim 12, wherein the isoparaffin is selected from the group consisting of isobutane, isopentanes and mixtures thereof.