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- (54) Title: SYNTHESIS AND USE OF MOLECULAR SIEVE ITQ-32

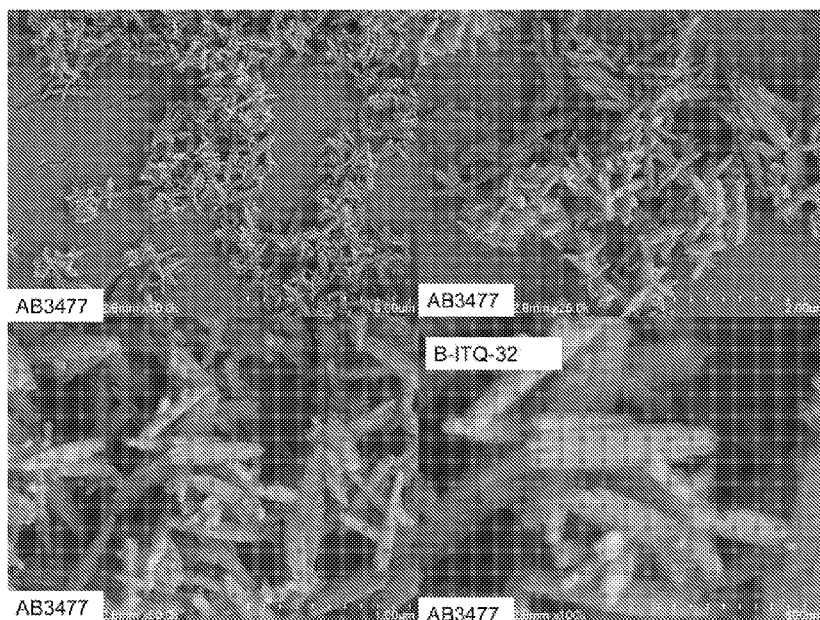


FIG. 2

(57) Abstract: The present invention relates to molecular sieves having the structure of ITQ-32 is synthesized from a reaction mixture substantially free of fluoride ions and comprising 4,4- dimethyl, 1-cyclohexyl-piperazinium cations in its pore structure, as well as methods of making such molecular sieves and methods of using them.



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SYNTHESIS AND USE OF MOLECULAR SIEVE ITQ-32

FIELD OF THE INVENTION

[0001] This invention relates to the synthesis of ITQ-32 and use of the resultant molecular sieve as an adsorbent and a catalyst for organic conversion reactions.

BACKGROUND OF THE INVENTION

[0002] ITQ-32 is a molecular sieve having a two-dimensional pore system comprising cages linked through windows of 8-membered rings of tetrahedrally coordinated atoms. Although the widest cross section of the large cages is circumscribed by 12-rings, these 12-rings can only be accessed by diffusion through the 8-ring channels. ITQ-32 has been assigned the framework type IHW by the Structure Commission of the International Zeolite Association.

[0003] ITQ-32 is of interest for the separation of small molecular species like carbon dioxide and methane because of the small dimensions of its pores (3.5Å x 4.3Å). This and other separations using ITQ-32 are described in, for example, U.S. Patent Application Publication No. 2009/0202416.

[0004] The characteristic X-ray diffraction pattern of ITQ-32 and its synthesis in the presence of N,N,N',N'-tetramethyldecahydrocyclobuta[1,2-*c*;3,4-*c'*]dipyrrolidinium and 4,4-dimethyl,1-cyclohexylpiperazinium cations as organic structure directing agents are disclosed in U.S. Patent No. 7,582,278. However, the synthesis mixtures described in U.S. Patent No. 7,582,278 require the presence of fluoride ions, which in commercial scale operations is disadvantageous in that they may lead to extra safety and cost considerations. There is, therefore, interest in synthesizing ITQ-32 in the absence of fluoride ions.

[0005] According to the present invention, it has now been found that ITQ-32, particularly in a borosilicate form, can be prepared in the absence of fluoride. The fluoride-free synthesis can be effected using 4,4-dimethyl,1-cyclohexylpiperazinium as the organic structure directing agent. The synthesis can be accomplished in the absence of alkali metal cations thereby obviating the need for ion-exchange of the product after calcination to remove the occluded organic structure directing agent.

SUMMARY OF THE INVENTION

[0006] In one aspect, the invention resides in a molecular sieve having the structure of ITQ-32 and, in its as-synthesized form, being substantially free of fluoride ions.

[0007] In one embodiment, the molecular sieve comprises 4,4-dimethyl,1-cyclohexylpiperazinium cations in its pores.

[0008] In one embodiment, the as-synthesized molecular sieve has a composition comprising the molar relationship $qQ^+:xX_2O_3:SiO_2$ (normalized to ~1.0 silica content) wherein $0 < q \leq 0.06$, $0 \leq x \leq 0.05$, Q^+ comprises a 4,4-dimethyl,1-cyclohexylpiperazinium cation, and X comprises a trivalent element, such as at least one of aluminum, boron, chromium, gallium, and iron, desirably including or being boron.

[0009] In a further aspect, the invention resides in a process for producing a molecular sieve having the structure of ITQ-32, the process comprising: (i) preparing a synthesis mixture capable of forming said molecular sieve and comprising water, a source of silica, a source of a trivalent element (X), and a source of 4,4-dimethyl,1-cyclohexylpiperazinium cations (Q^+), said synthesis mixture being substantially free of fluoride ions and having a composition, in terms of molar ratios, within the following amounts: SiO_2/X_2O_3 from about 2 to about 100; Q^+/SiO_2 from about 0.02 to about 1.0; and H_2O/YO_2 from about 10 to about 60; (ii) heating said mixture under crystallization conditions comprising a temperature of from about 100 °C to about 200 °C and a time from about 1 day to about 28 days until crystals of said molecular sieve are formed; and (iii) recovering said molecular sieve from step (ii).

[0010] In one embodiment, the synthesis mixture comprises a source of chloride ions, desirably such that the Cl^-/SiO_2 molar ratio of said synthesis mixture is (non-zero and) up to about 0.3.

BRIEF DESCRIPTION OF THE DRAWINGS

[0011] FIG. 1 shows the powder XRD patterns of the as-made and calcined forms of the borosilicate ITQ-32 produced in Example 2.

[0012] FIG. 2 provides SEM images of the borosilicate ITQ-32 produced in Example 2. Magnification increases from left to right and top to bottom within the four images.

DETAILED DESCRIPTION OF THE EMBODIMENTS

[0013] Described herein is a process for the synthesis of the molecular sieve ITQ-32 from a synthesis mixture which is substantially free of fluoride ions and optionally may contain other halide ions, particularly chloride ions. As used herein, the term “substantially

free of fluoride ions” means that the synthesis mixture contains less than 100 wppm, *e.g.*, less than 50 wppm, less than 25 wppm, less than 10 wppm, or no measurable quantity, of fluoride ions.

[0014] The present synthesis employs 4,4-dimethyl,1-cyclohexylpiperazinium cations as a structure directing agent, and the processes described herein can produce ITQ-32 material having, in its as-synthesized form, a composition comprising the molar relationship: $qQ^+:xX_2O_3:SiO_2$, wherein $0 < q \leq 0.06$, $0 \leq x \leq 0.05$, Q^+ comprises a 4,4-dimethyl,1-cyclohexylpiperazinium cation, and X comprises a trivalent element, such as at least one of aluminum, boron, chromium, gallium, and iron. In one embodiment, boron is present in the as-synthesized molecular sieve, either alone or in combination with one or more other trivalent elements, such as aluminum. It should be understood that the molar relationship above was normalized to the content of the silica component, such that the (absent) coefficient for silica is meant to be ~ 1.0 .

[0015] In its as-synthesized form, the ITQ-32 produced by the present process can exhibit an X-ray diffraction (XRD) spectrum including the characteristic peak maxima listed in Table 1 below.

Table 1

Two-theta (degrees) ± 0.3	Relative Intensity (100xI/I _o)
7.3	m-s
8.8	vs
9.7	w
16.4	w-m
19.6	s
20.2	s-vs
21.0	s-vs
21.9	vs
24.2	m
26.0	m
27.0	m-s
27.5	m

[0016] After calcination to remove the organic material Q^+ occluded in its pores, the ITQ-32 produced by the present process can have an XRD pattern including the characteristic peak maxima listed in Table 2 below.

Table 2

Two-theta (degrees) ± 0.3	Relative Intensity (100xI/I _o)
7.4	s-vs
8.9	vs
9.7	w
12.9	w
19.3	w
20.3	w-m
20.9	m
22.0	m
24.3	w
26.0	w-m
27.1	m
27.6	w

[0017] The X-ray diffraction data reported herein were collected with a Panalytical X'Pert Pro diffraction system with an Xcelerator multichannel detector, equipped with a germanium solid state detector, using copper K-alpha radiation. The diffraction data were recorded by step-scanning at 0.02 degrees of two-theta, where theta is the Bragg angle, and using an effective counting time of 2 seconds for each step. The interplanar spacings, d-spacings, were calculated in Angstrom units, and the relative intensities of the lines, I/I_o is the ratio of the peak intensity to that of the intensity of the strongest line, above background. The intensities are uncorrected for Lorentz and polarization effects. The relative intensities are given in terms of the symbols "vs" = very strong (60-100%), "s" = strong (40-60%), "m" = medium (20-40%), and "w" = weak (0-20%). In certain cases, some (select) or all weak peaks listed may have non-zero intensities.

[0018] The ITQ-32 described herein can be produced from a synthesis mixture comprising water, a source of silica, a source of a trivalent element (X), and a source of 4,4-dimethyl,1-cyclohexylpiperazinium cations (Q⁺), wherein the synthesis mixture can be substantially free of fluoride ions and can have a composition, in terms of molar ratios, within the following amounts:

<u>Reactants</u>	<u>Useful</u>	<u>Preferred</u>
SiO ₂ /X ₂ O ₃	2-100	8-30
Q ⁺ /SiO ₂	0.02-1.0	0.025-0.6; and
H ₂ O/YO ₂	10-60	15-40.

[0019] Suitable sources of silica in the above mixture can include, but are not necessarily limited to, colloidal suspensions of silica, precipitated silicas, alkali metal silicates, tetraalkyl-orthosilicates, silicon hydroxides, silicon oxy-hydroxides, and the like, and combinations thereof. Suitable sources of the trivalent element X can depend on the relevant trivalent element(s) employed but, in the case of boron, can include but are not necessarily limited to boric acid, sodium borate, potassium borate, boron oxide, and combinations thereof. Suitable sources of aluminum, when present, can include but are not necessarily limited to hydrated alumina (also called aluminum hydroxides), aluminum oxide, aluminum oxy-hydroxides, water-soluble aluminum salts such as aluminum nitrate, and combinations thereof.

[0020] Suitable sources of Q^+ can include the hydroxides, halides (*e.g.*, chloride, bromide, and/or iodide), and/or other salts of the relevant quaternary ammonium compound.

[0021] In some embodiments, the synthesis mixture can comprise a source of chloride ions, for example such that the Cl^-/SiO_2 molar ratio of said synthesis mixture can be (non-zero and) up to about 0.3.

[0022] Though the pH of the synthesis mixture may not necessarily be critical, it can be highly dependent on the combination of various different components in the synthesis mixture. While pH values can extend from 7.5 to 14, it can sometimes be difficult to attain relatively high yield and/or relatively high purity at lower pH values in the synthesis mixture. Nevertheless, in certain embodiments, the pH of the synthesis mixture can be relatively low, *e.g.*, from 8 to 10, while still allowing substantially fluoride-free ITQ-32 product to be effectively made.

[0023] In some embodiments, the synthesis mixture may also include seeds of a molecular sieve material, such as ITQ-32 from a previous synthesis, *e.g.*, in an amount from about 0.1 wt% to about 10 wt% or from about 0.5 wt% to about 5 wt% of the synthesis mixture. Additionally or alternately, when seeds such as ITQ-32 seeds are present in the synthesis mixture, they can be present in an amount from about 0.1 wt% to about 10 wt% or from about 0.5 wt% to about 5 wt% of the silica component of the synthesis mixture.

[0024] Crystallization of ITQ-32 from the synthesis mixture described herein can be carried out at static or stirred conditions in a suitable reactor vessel (*e.g.*, polypropylene jars or TeflonTM-lined or stainless steel autoclaves) at a temperature from about 100°C to about 200°C, such as from about 140°C to about 180°C, for a time sufficient for crystallization to occur at the temperature used (*e.g.*, from about 12 hours to about 100 days, from about 1 day to about 7 days, or from about 2 days to about 20 days). Thereafter, the crystals can be separated from the liquid and recovered. It should be appreciated that not all of the trivalent

element present in the synthesis mixture may be transferred to the framework of the molecular sieve, such that the $\text{SiO}_2/\text{X}_2\text{O}_3$ molar ratio of the ITQ-32 product may indeed be higher than that of the synthesis mixture. The same can be true with one or more other components of the synthesis mixture.

[0025] Using the synthesis mixture and processes described herein, it is possible to synthesize ITQ-32 that may not only be substantially free of fluoride ions in its as-synthesized form, but that can also contain less than 25 wt% (*e.g.*, contain less than 20 wt%, contain less than 15 wt%, contain less than 11 wt%, or be substantially free) of large pore molecular sieve/zeolite materials (such as ZSM-12), *e.g.*, as measured by XRD techniques. Large pore molecular sieve/zeolite materials, as used herein, have a microporous structure with at least one pore opening cross-sectional dimension being greater than 6\AA . As used herein, the term “substantially free of large pore” materials means that the product contains less than 10 wt%, *e.g.*, less than 8 wt%, less than 6 wt%, less than 5 wt%, less than 4 wt%, less than 3 wt%, or no detectable quantity by XRD techniques), of large pore materials, which can include, but are not necessarily limited to, ZSM-12 and the like. Being a relatively large pore molecular sieve, ZSM-12 can be a relatively undesirable impurity when ITQ-32 is used for adsorption applications. Indeed, it is additionally or alternately possible to synthesize ITQ-32 product using the synthesis mixture and/or processes described herein so as to attain less than 25 wt% (*e.g.*, less than 20 wt%, less than 15 wt%, less than 11 wt%, less than 10 wt%, less than 8 wt%, less than 6 wt%, less than 5 wt%, less than 4 wt%, less than 3 wt%, or no detectable quantity) of impurity (non-ITQ-32) phases, *e.g.*, by XRD techniques.

[0026] In one embodiment, the ITQ-32 product synthesized by the present process can comprise silicon and boron in an atomic ratio (Si/B) of greater than 7, *e.g.*, greater than 8.

[0027] The as-synthesized ITQ-32 product may also be subjected to treatment to remove and/or decompose all or part of the organic directing agent Q used in its synthesis. This can be conveniently accomplished by thermal treatment, in which the as-synthesized material can be heated to a temperature of at least about 370°C for at least 1 minute and generally not longer than 20 hours. While subatmospheric pressure can be employed for the thermal treatment, atmospheric pressure can be desired for reasons of convenience. The thermal treatment can be performed at a convenient and effective temperature, *e.g.*, up to about 925°C . The thermally treated product, especially in its metal, hydrogen, and ammonium forms, can be particularly useful as an adsorbent and/or in the catalysis of certain organic (*e.g.*, hydrocarbon) conversion reactions.

[0028] The present synthesis of ITQ-32 can be accomplished in the absence of alkali metal cations, thereby obviating the need for ion-exchange of the product after thermal treatment to remove any occluded structure directing agent. However, depending on the X_2O_3/SiO_2 molar ratio of the material, any cations in the as-synthesized ITQ-32 can be replaced in accordance with techniques well known in the art, *e.g.*, by ion exchange with other cations. Preferred replacing cations can include, but are not necessarily limited to, metal ions, hydrogen ions, hydrogen precursor (*e.g.*, ammonium) ions, and mixtures thereof. Particularly preferred cations can include those that can tailor the catalytic activity for adsorption and/or for certain hydrocarbon conversion reactions (*e.g.*, hydrogen, rare earth metals, and/or one or more metals of Groups 2-15 of the Periodic Table of the Elements). As used herein, the numbering scheme for the Periodic Table Groups is as disclosed in Chemical and Engineering News, 63(5), 27 (1985).

[0029] ITQ-32, as produced herein, may be intimately combined with a hydrogenating component, such as molybdenum, rhenium, nickel, cobalt, chromium, manganese, and/or noble metal (such as platinum and/or palladium), where a hydrogenation-dehydrogenation function may be desirable to perform. Such component can be in the composition by way of co-crystallization, exchanged into the composition (to the extent a trivalent element, *e.g.*, boron, is present in the structure), impregnated therein, and/or intimately physically admixed therewith. Such component can be impregnated in/on to the composition, *e.g.*, in the case of platinum, by treating the silicate with a solution containing a platinum metal-containing ion. Thus, suitable platinum compounds for such purpose can include chloroplatinic acid, platinumous chloride, and/or various compounds containing the platinum amine complex.

[0030] The present molecular sieve, when employed as an adsorbent and/or as a catalyst, can generally be at least partially dehydrated. This can be done, *e.g.*, by heating to a temperature from about 120°C to about 400°C (*e.g.*, from about 200°C to about 370°C) in an atmosphere such as air, nitrogen, etc., and at atmospheric, subatmospheric, or superatmospheric pressures for an appropriate time (*e.g.*, from about 15 minutes to about 48 hours). Dehydration can alternately be performed at room temperature (~20-25°C) merely by placing the ITQ-32 in a vacuum, but a longer time may be required to sufficiently dehydrate.

[0031] The ITQ-32 described herein can be particularly useful as an adsorbent in separating a first component, such as carbon dioxide, from a gaseous mixture comprising the first component and an additional second component, such as methane.

[0032] Additionally or alternately, the present invention can include one or more of the following embodiments.

[0033] Embodiment 1. A molecular sieve having the structure of ITQ-32 and, in its as-synthesized form, being substantially free of fluoride ions.

[0034] Embodiment 2. The molecular sieve of embodiment 1, which comprises silicon and boron in an atomic ratio (Si/B) greater than 8:1.

[0035] Embodiment 3. The molecular sieve of embodiment 1 or embodiment 2, wherein the molecular sieve is substantially free of large pore molecular sieve materials, such as ZSM-12.

[0036] Embodiment 4. The molecular sieve of any one of the previous embodiments, further comprising 4,4-dimethyl,1-cyclohexylpiperazinium cations in its pore structure.

[0037] Embodiment 5. The molecular sieve of embodiment 4, which has a composition comprising the molar relationship: $qQ^+:xX_2O_3:SiO_2$, wherein $0 < q \leq 0.06$, $0 \leq x \leq 0.05$, Q^+ comprises a 4,4-dimethyl,1-cyclohexylpiperazinium cation, and X comprises a trivalent element (*e.g.*, comprising at least one of aluminum, boron, chromium, gallium, and iron, such as comprising at least boron and/or aluminum).

[0038] Embodiment 6. A process for producing a molecular sieve having the structure of ITQ-32 and/or a molecular sieve according to any one of the previous embodiments, the process comprising: (i) preparing a synthesis mixture capable of forming the molecular sieve and comprising water, a source of silica, a source of a trivalent element (X), and a source of 4,4-dimethyl,1-cyclohexylpiperazinium cations (Q^+), the synthesis mixture being substantially free of fluoride ions and having a composition wherein: an SiO_2/X_2O_3 ratio is from about 2 to about 100; a Q^+/SiO_2 ratio is from about 0.02 to about 1.0; and an H_2O/YO_2 ratio is from about 10 to about 60; (ii) heating the mixture under crystallization conditions comprising a temperature of from about 100°C to about 200°C and a time from about 1 day to about 28 days until crystals of the molecular sieve are formed; and (iii) recovering the molecular sieve from step (ii).

[0039] Embodiment 7. The process of embodiment 6, wherein the synthesis mixture comprises a source of chloride ions.

[0040] Embodiment 8. The process of embodiment 7, wherein the synthesis mixture exhibits a (non-zero) Cl/SiO_2 molar ratio up to about 0.3.

[0041] Embodiment 9. The process of any one of embodiments 6-8, wherein the synthesis mixture also contains seeds, *e.g.*, from about 0.01 wppm to about 10000 wppm seeds or from about 100 wppm to about 5000 wppm seeds.

[0042] Embodiment 10. The process of embodiment 9, wherein the seeds comprise a crystalline material having the structure of ITQ-32.

[0043] Embodiment 11. The process of any one of embodiments 6-10, further comprising removing and/or decomposing at least part of the organic material from the molecular sieve crystals recovered in step (iii).

[0044] Embodiment 12. A molecular sieve produced by the process of any one of embodiments 6-11.

[0045] Embodiment 13. A process for separating a first component from a gaseous mixture comprising said first component and a second component, the process comprising contacting the gaseous mixture with the molecular sieve of any one of embodiments 1-4 or 12.

[0046] Embodiment 14. The process of embodiment 13, wherein the first component comprises carbon dioxide and the second component comprises methane.

[0047] The invention will now be more particularly described with reference to the following Examples and the accompanying drawings.

EXAMPLES

[0048] In the Examples, the alpha test is a measure of molecular sieve acidic functionality and is described together with details of its measurement in U.S. Patent No. 4,016,218, and in J. Catalysis, Vol. VI, pp. 278-287 (1966).

Example 1

[0049] A reaction mixture was prepared in a sealed 1-mL stainless steel reactor vessel with the following mole ratios using boric acid as the boron source, Ludox™ AS-40 as the silica source, and substantially no fluoride ions: SDA/Si \approx 0.2; Si/B \approx 5; HCl/Si \approx 0.10; and H₂O/Si \approx 35. 4,4-dimethyl,1-cyclohexylpiperazinium hydroxide was used as the structure directing agent (SDA).

[0050] The reaction mixture was heated in a convection oven under tumbling conditions (\sim 30 rpm) for about 28 days at \sim 160°C. The sample was worked up by a series of three centrifugations and washings with deionized water. Powder XRD of the product appeared to indicate substantially pure phase ITQ-32.

Example 2

[0051] First, \sim 5.1 grams deionized water were added to \sim 2.20 grams of 1,1-dimethyl-4-cyclohexylpiperazinium hydroxide (29.2%) inside a Teflon™ liner for a \sim 23-mL steel Parr autoclave. Next, \sim 0.19 grams boric acid was added to the solution and mixed manually with a spatula until near dissolution. In the next step, \sim 1.50 grams of 1 N HCl and then \sim 2.25 grams of Ludox™ AS-40 were added to the solution and mixed to create a relatively uniform

suspension having the same molar ratios as the reaction mixture of Example 1. About 0.04 grams of seeds from the Example 1 product were added. The liner was then capped, sealed inside the ~23-mL autoclave, and heated to ~160°C under tumbling conditions (~50 rpm). After about 14 days of heating, the reaction was quenched, and the solids were isolated by filtering through a Buchner funnel, washing with deionized water, and drying in a vacuum oven at ~60°C. The yield of solid product was ~1.02 grams. Powder XRD of the product appeared to indicate substantially pure phase ITQ-32. Figure 1 shows the powder XRD patterns of the as-made and calcined forms of the product of Example 2. Figure 2 shows SEM images of the product of Example 2.

[0052] The calcined form of the product of Example 2 was heated inside a muffle furnace from ambient temperature (~20-25°C) to about 400°C at a heating rate of ~4°C/min under a nitrogen atmosphere, heated to about 600°C at a heating rate of ~4°C/min in air, and then maintained at about 600°C in air for about another 2 hours. The calcined product was then measured with nitrogen physisorption and the data were analyzed using the t-plot method. The micropore volume was determined to be ~0.16 cm³/g, and the external (mesopore) surface area was determined to be about 52 m²/g.

Example 3

[0053] Example 2 was repeated, except that ~0.023 grams of aluminum hydroxide and ~0.06 g of the products seeds from Example 2 were included in the synthesis mixture. The yield of solid product after about 14 days crystallization was ~1.05 grams, and powder XRD of the product appeared to indicate substantially pure phase ITQ-32. This sample was calcined and then alpha tested for n-hexane cracking activity. The alpha value of the sample was determined to be ~6.

Example 4

[0054] Example 2 was repeated, except that no 1 N HCl was added. Additional deionized water was added to maintain the same H₂O/Si level. After ~8 days of heating, the product appeared (by XRD) to be substantially pure phase ITQ-32.

Example 5

[0055] Example 4 was repeated, except that ~0.018 grams of sodium chloride was included in the synthesis mixture. After ~7 days of heating, the product appeared (by XRD) to be substantially pure phase ITQ-32.

Example 6

[0056] Example 3 was repeated, except that ~0.033 grams of aluminum hydroxide was included in the synthesis mixture to give an initial Si/Al ratio of ~35 and an initial Si/B ratio

of ~5. After ~9 days of heating, the product appeared (by XRD) to comprise predominantly ITQ-32 with a minor gibbsite impurity.

Example 7

[0057] Example 2 was repeated, but without boron and with sufficient aluminum hydroxide included in the synthesis mixture to give an initial Si/Al ratio of ~20. The product appeared (by XRD) to be amorphous after heating for ~17 days.

[0058] Although the present invention has been described in terms of specific embodiments, it is not so limited. Suitable alterations/modifications for operation under specific conditions should be apparent to those skilled in the art. It is therefore intended that the following claims be interpreted as covering all such alterations/modifications as fall within the true spirit/scope of the invention.

CLAIMS

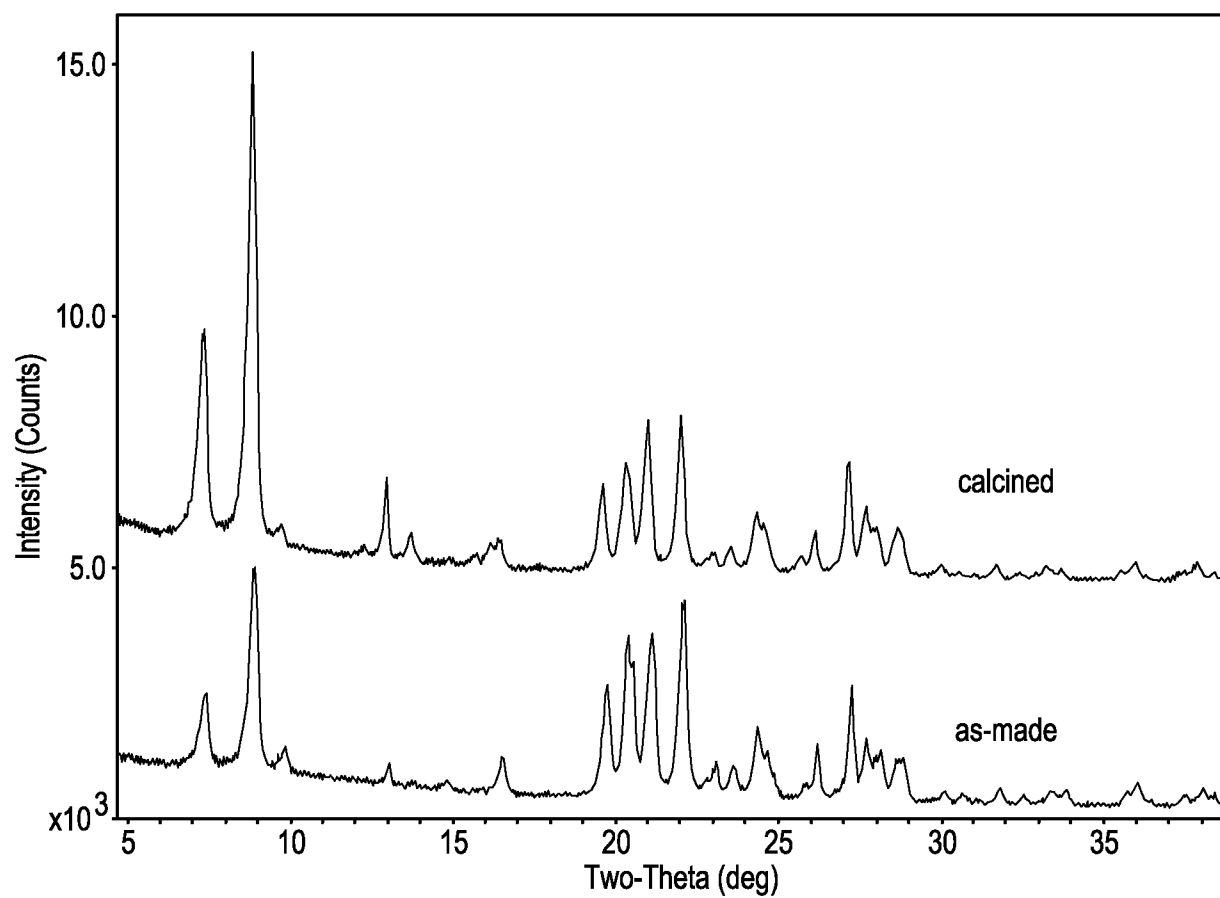
WHAT IS CLAIMED IS:

1. A molecular sieve having the structure of ITQ-32 and, in its as-synthesized form, being substantially free of fluoride ions.
2. The molecular sieve of claim 1, which comprises silicon and boron in an atomic ratio (Si/B) greater than 8:1.
3. The molecular sieve of claim 1 or claim 2, wherein the molecular sieve is substantially free of large pore molecular sieve materials, such as ZSM-12.
4. The molecular sieve of any one of the previous claims, further comprising 4,4-dimethyl,1-cyclohexylpiperazinium cations in its pore structure.
5. The molecular sieve of claim 4, which has a composition comprising the molar relationship: $qQ^+:xX_2O_3:SiO_2$, wherein $0 < q \leq 0.06$, $0 \leq x \leq 0.05$, Q^+ comprises a 4,4-dimethyl,1-cyclohexylpiperazinium cation, and X comprises a trivalent element (*e.g.*, comprising at least one of aluminum, boron, chromium, gallium, and iron, such as comprising at least boron and/or aluminum).
6. A process for producing a molecular sieve having the structure of ITQ-32 and/or a molecular sieve according to any one of the previous claims, the process comprising:
 - (i) preparing a synthesis mixture capable of forming the molecular sieve and comprising water, a source of silica, a source of a trivalent element (X), and a source of 4,4-dimethyl,1-cyclohexylpiperazinium cations (Q^+), the synthesis mixture being substantially free of fluoride ions and having a composition wherein: an SiO_2/X_2O_3 ratio is from about 2 to about 100; a Q^+/SiO_2 ratio is from about 0.02 to about 1.0; and an H_2O/YO_2 ratio is from about 10 to about 60;
 - (ii) heating the mixture under crystallization conditions comprising a temperature of from about 100°C to about 200°C and a time from about 1 day to about 28 days until crystals of the molecular sieve are formed; and
 - (iii) recovering the molecular sieve from step (ii).
7. The process of claim 6, wherein the synthesis mixture comprises a source of chloride ions.
8. The process of claim 7, wherein the synthesis mixture exhibits a (non-zero) Cl^-/SiO_2 molar ratio up to about 0.3.

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9. The process of any one of claims 6-8, wherein the synthesis mixture also contains seeds, *e.g.*, from about 0.01 wppm to about 10000 wppm seeds or from about 100 wppm to about 5000 wppm seeds.
10. The process of claim 9, wherein the seeds comprise a crystalline material having the structure of ITQ-32.
11. The process of any one of claims 6-10, further comprising removing and/or decomposing at least part of the organic material from the molecular sieve crystals recovered in step (iii).
12. A molecular sieve produced by the process of any one of claims 6-11.
13. A process for separating a first component from a gaseous mixture comprising said first component and a second component, the process comprising contacting the gaseous mixture with the molecular sieve of any one of claims 1-4 or 12.
14. The process of claim 13, wherein the first component comprises carbon dioxide and the second component comprises methane.

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**FIG. 1**

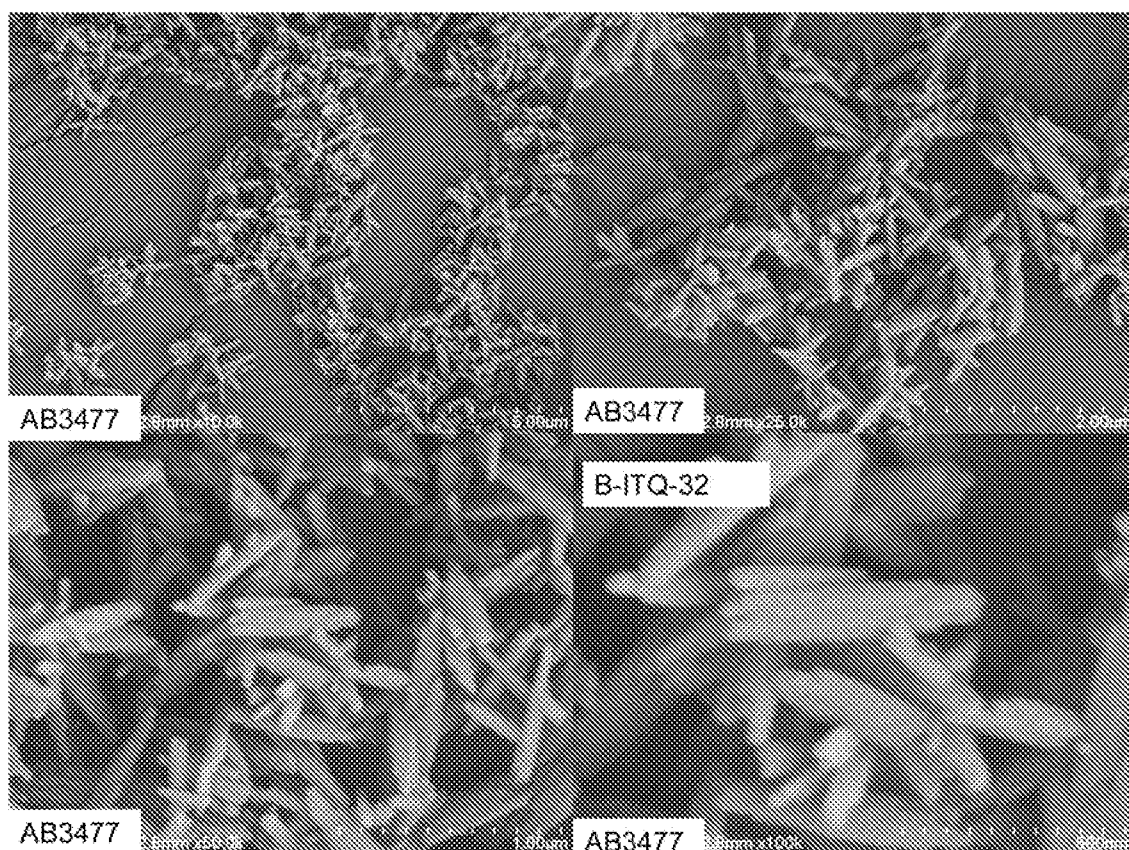


FIG. 2

INTERNATIONAL SEARCH REPORT

International application No

PCT/US2013/069577

A. CLASSIFICATION OF SUBJECT MATTER INV. C01B39/48 B01D53/04 ADD.				
According to International Patent Classification (IPC) or to both national classification and IPC				
B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) C01B B01D				
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched				
Electronic data base consulted during the international search (name of data base and, where practicable, search terms used) EPO-Internal, COMPENDEX, INSPEC, WPI Data				
C. DOCUMENTS CONSIDERED TO BE RELEVANT				
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.		
X	US 7 582 278 B2 (CORMA CANOS AVELINO [ES] ET AL) 1 September 2009 (2009-09-01) cited in the application	1-5, 12-14		
A	column 2, line 3 - column 3, line 35 column 9, lines 1-45 column 10, line 54 - column 11, line 5; examples 2-9	6-11		
X	US 2009/202416 A1 (CORMA CANOS AVELINO [ES] ET AL) 13 August 2009 (2009-08-13) cited in the application	1-5, 12-14		
A	paragraphs [0009], [0012] - [0025]; examples 1,2	6-11		
A	US 2009/318696 A1 (STROHMAIER KARL G [US] ET AL) 24 December 2009 (2009-12-24) paragraphs [0007] - [0022]; examples 5,7-8	1-14		
<input type="checkbox"/> Further documents are listed in the continuation of Box C. <input checked="" type="checkbox"/> See patent family annex.				
* Special categories of cited documents : <table border="0"> <tr> <td> "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier application or patent but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed </td> <td> "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "&" document member of the same patent family </td> </tr> </table>			"A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier application or patent but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "&" document member of the same patent family
"A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier application or patent but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed	"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art "&" document member of the same patent family			
Date of the actual completion of the international search 6 May 2014		Date of mailing of the international search report 12/05/2014		
Name and mailing address of the ISA/ European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Fax: (+31-70) 340-3016		Authorized officer Follens, Lana		

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

PCT/US2013/069577

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 7582278	B2	01-09-2009	EP 1852394 A1 07-11-2007
			ES 2263369 A1 01-12-2006
			JP 4964150 B2 27-06-2012
			JP 2008528435 A 31-07-2008
			US 2009068092 A1 12-03-2009
			WO 2007036589 A1 05-04-2007

US 2009202416	A1	13-08-2009	AU 2006273997 A1 01-02-2007
			BR PI0614173 A2 15-03-2011
			CA 2616705 A1 01-02-2007
			CN 101351257 A 21-01-2009
			EP 1918012 A1 07-05-2008
			ES 2275422 A1 01-06-2007
			JP 2009502467 A 29-01-2009
			KR 20080071548 A 04-08-2008
			US 2009202416 A1 13-08-2009
			WO 2007012690 A1 01-02-2007
			ZA 200801734 A 27-05-2009

US 2009318696	A1	24-12-2009	CA 2726816 A1 23-12-2009
			CN 102066259 A 18-05-2011
			EP 2300370 A2 30-03-2011
			JP 5377632 B2 25-12-2013
			JP 2011524848 A 08-09-2011
			US 2009318696 A1 24-12-2009
			WO 2009154783 A2 23-12-2009
