

SCM-10 MOLECULAR SIEVE, A PROCESS FOR PRODUCING SAME AND USE THEREOF

Technical Field

The present invention relates to an SCM-10 molecular sieve, a process for producing same and use thereof.

Background Art

In industry, porous inorganic materials have been widely used as catalysts and catalyst carriers. These porous materials generally include amorphous porous materials, crystalline molecular sieves and modified layered materials. Minute difference in the structure between any two materials may indicate significant difference in properties like catalytic performance and adsorbing ability therebetween, and further difference in available parameters used to characterize same, such as morphology, specific surface area or pore size.

The structure of a molecular sieve is specifically confirmed by the X-ray diffraction pattern (XRD), while the X-ray diffraction pattern (XRD) is determined by X-ray powder diffraction with a Cu-K α -ray source and a Ni filter. Different molecular sieves have different characterizing XRD patterns. Known molecular sieves, like A-Type Zeolite, Y-Type Zeolite, MCM-22 molecular sieve and so on, have their characterizing XRD patterns respectively.

At the same time, two molecular sieves, if sharing the same characterizing XRD pattern but comprising different combination of skeleton elements, will be identified as different molecular sieves. For example, TS-1 molecular sieve (US4410501) and ZSM-5 molecular sieve (US3702886), share the same characterizing XRD pattern but comprise different combination of skeleton elements. Specifically, TS-1 molecular sieve comprises Si and Ti as the skeleton elements, exhibiting a catalytic oxidation ability, while ZSM-5 molecular sieve comprises Si and Al as the skeleton elements, exhibiting an acidic catalytic ability.

Further, two molecular sieves, if sharing the same characterizing XRD pattern and the same combination of skeleton elements but with different relative amounts of the skeleton elements, will be identified as different molecular sieves as well. For example, Zeolite X (US2882244) and Zeolite Y (US3130007), share the same characterizing XRD pattern and the same combination of skeleton elements (Si and Al), but with different relative amounts of Si and Al. Specifically, Zeolite X has a Si/Al molar ratio of less than 1.5, while Zeolite Y has a Si/Al molar ratio of greater than 1.5.

Invention Summary

The present inventors, on the basis of the prior art, found a novel molecular sieve having the SFE structure, and further identified beneficial properties for same.

Specifically, this invention relates to the following aspects.

1. An SCM-10 molecular sieve, having an empirical chemical composition as illustrated by the formula "the first oxide · the second oxide", wherein the ratio by molar of the first oxide to the second oxide is less than 40, preferably in the range of from 3 to less than 40, more preferably 5-30, the first oxide is at least one selected from the group consisting of silica and germanium dioxide, preferably silica, the second oxide is at least one selected from the group consisting of

alumina, boron oxide, iron oxide, gallium oxide, titanium oxide, rare earth oxides, indium oxide and vanadium oxide, preferably boron oxide or a combination of boron oxide and at least one selected from the group consisting of alumina, iron oxide, gallium oxide, titanium oxide, rare earth oxides, indium oxide and vanadium oxide, more preferably boron oxide or a combination of boron oxide and alumina, more preferably boron oxide, and the molecular sieve in the calcined form has X ray diffraction pattern as substantially illustrated in the following table,

2 θ (°) ^(a)	d-spacing (Å) ^(b)	Relative intensity (I/I ₀ ×100)
6.50	13.59	w-s
7.98	11.07	s-vs
9.36	9.45	m
11.27	7.85	w-m
20.02	4.43	s
22.65	3.92	vs
24.13	3.69	vs
26.45	3.37	w-m
27.92	3.19	w-m
35.95	2.50	m

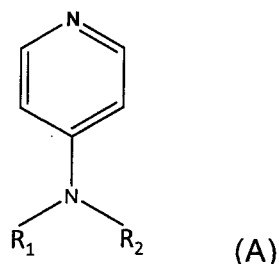
a: $\pm 0.3^\circ$, b: changed with 2 θ .

2. The SCM-10 molecular sieve according to anyone of the preceding aspects, wherein the X-ray diffraction pattern further includes X-ray diffraction peaks as substantially illustrated in the following table,

2 θ (°) ^(a)	d-spacing (Å) ^(b)	Relative intensity (I/I ₀ ×100)
14.12	6.27	w
16.12	5.49	w
18.82	4.71	w
37.52	2.40	w

a: $\pm 0.3^\circ$, b: changed with 2 θ .

3. A process for producing an SCM-10 molecular sieve, including a step of crystallizing a mixture comprising a first oxide source, a second oxide source, an organic template and water to obtain the molecular sieve, and optionally, a step of calcining the obtained molecular sieve, wherein the organic template is selected from a compound represented by the following formula (A), a quaternary ammonium salt thereof and a quaternary ammonium hydroxide thereof, preferably 4-dimethylamino pyridine,



wherein R_1 and R_2 may be identical to or different from each other, each independently representing a C_{1-8} alkyl, preferably a C_{1-4} alkyl, more preferably a C_{1-2} alkyl.

4. The process according to anyone of the preceding aspects, wherein the mixture does not contain an alkaline source.

5. The process according to anyone of the preceding aspects, wherein the mixture does not contain a fluorine source.

6. The process according to anyone of the preceding aspects, wherein the mixture has a $pH=6-14$, preferably $pH=7-14$, more preferably $8-14$, more preferably $8.5-13.5$, more preferably $9-12$, more preferably $9-11$.

7. The process according to anyone of the preceding aspects, wherein the first oxide source is at least one selected from the group consisting of a silicon source and a germanium source, preferably a silicon source, the second oxide source is at least one selected from the group consisting of an aluminum source, a boron source, an iron source, a gallium source, a titanium source, a rare earth source, an indium source and a vanadium source, preferably a boron source or a combination of a boron source and at least one selected from the group consisting of an aluminum source, an iron source, a gallium source, a titanium source, a rare earth source, an indium source and a vanadium source, more preferably a boron source or a combination of a boron source and an aluminum source, more preferably a boron source, the ratio by molar between the first oxide source (as the first oxide), the second oxide source (as the second oxide), the organic template and water is $1:(0.025-1/3):(0.01-1.0):(4-50)$, preferably $1:(1/30-1/3):(0.02-0.9):(4-40)$, more preferably $1:(1/30-1/5):(0.04-0.8):(4-30)$, with the proviso that the ratio by molar of the first oxide source (as the first oxide) to the second oxide source (as the second oxide) is less than 40, preferably less than 30.

8. An SCM-10 molecular sieve composition, comprising the SCM-10 molecular sieve according to anyone of the preceding aspects or an SCM-10 molecular sieve produced in line with the process according to anyone of the preceding aspects, and a binder.

9. Use of the SCM-10 molecular sieve according to anyone of the preceding aspects, an SCM-10 molecular sieve produced in line with the process according to anyone of the preceding aspects or the SCM-10 molecular sieve composition according to anyone of the preceding aspects as an adsorbent or a catalyst for converting an organic compound.

10. Use according to anyone of the preceding aspects, wherein the catalyst for converting an organic compound is at least one selected from the group consisting of an alkane isomerization catalyst, a catalyst for the alkylation between olefins and aromatics, an olefin isomerization catalyst, a naphtha cracking catalyst, a catalyst for the alkylation between alcohols and aromatics, an olefin hydration catalyst and an aromatic disproportionation catalyst.

Technical effects

According to the present invention, the SCM-10 molecular sieve has the SFE structure, but with a chemical composition that has never been obtained in this field before.

Figure Description

Fig.1 illustrates the X-ray diffraction pattern (XRD) of the molecular sieve produced in Example 1.

Specific Mode to Carry Out This Invention

This invention will be described in details hereinafter with reference to the following specific embodiments. However, it should be noted that the protection scope of this invention should not be construed as limited to these specific embodiments, but rather determined by the attached claims.

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Further, to the extent that any meaning or definition of a term in this document conflicts with any meaning or definition of the same term in a document incorporated by reference, the meaning or definition assigned to that term in this document shall govern.

In the context of this specification, a molecular sieve, before any other material (for example, organic templates) than water and metal ions that has been filled into its pores during the production thereof is removed from the pores, is referred to as "precursor".

In the context of this specification, in the XRD data of the molecular sieve, w, m, s, vs indicate the intensity of a diffraction peak, with w referring to weak, m to medium, s to strong, vs to very strong, which has been well known in this field. In general, w represents a value of less than 20; m represents a value of 20-40; s represents a value of 40-70; vs represents a value of greater than 70.

In the context of this specification, the molecular sieve structure is confirmed by the X-ray diffraction pattern (XRD), while the X-ray diffraction pattern (XRD) is determined by X-ray powder diffraction with a Cu-K α -ray source and a Ni filter. Before determination, the crystalline state of the test sample is observed under scanning electron microscope (SEM), to confirm that there presents only one type of crystal therein, which indicates that the molecular sieve as the test sample presents as a pure phase, and then the XRD determination is conducted thereon, in order to ensure that there is no interfering peak of other crystal in the XRD pattern.

In the context of this specification, by specific surface area, it refers to the total area per unit of mass of a sample, including the internal surface area and the external surface area. A non-porous material has only external surface area, like Portland cement or some clay mineral powder, while a porous material has an external surface area and an internal surface area, like asbestos fiber, diatomite or molecular sieves. In a porous material, the surface area of pores having a

diameter of less than 2nm is referred to as internal surface area, while the surface area obtained by subtracting the internal surface area from the total surface area is referred to as external surface area. The external surface area per unit of mass of a sample is referred to as external specific surface area.

In the context of this specification, by pore volume, it refers to the volume of pores per unit of mass of a porous material (e.g. a molecular sieve). By total pore volume, it refers to the volume of all pores (generally involving only pores having a pore diameter of less than 50nm) per unit of mass of a molecular sieve. By micropore volume, it refers to the volume of all micropores (generally referred to pores having a pore diameter of less than 2nm) per unit of mass of a molecular sieve.

The present invention relates to an SCM-10 molecular sieve. The SCM-10 molecular sieve has the SFE structure, but with a chemical composition that has never been obtained in this field before.

According to the present invention, the SCM-10 molecular sieve has an empirical chemical composition as illustrated by the formula " the first oxide · the second oxide ". It is known that, a molecular sieve will sometimes (especially immediately after the production thereof) contain a certain amount of water, however, this invention does not specify or identify as to how much this amount may be, since the presence or absence of water will not substantially change the XRD pattern of the present molecular sieve. In this context, the empirical chemical composition actually represents an anhydrous chemical composition of this molecular sieve. Further, it is obvious that the empirical chemical composition represents the framework chemical composition of the molecular sieve.

According to the present invention, in the SCM-10 molecular sieve, the ratio by molar of the first oxide to the second oxide is generally less than 40, preferably in the range of from 3 to less than 40, more preferably 5-30.

According to the present invention, in the SCM-10 molecular sieve, the first oxide is at least one selected from the group consisting of silica and germanium dioxide, preferably silica.

According to the present invention, in the SCM-10 molecular sieve, the second oxide is at least one selected from the group consisting of alumina, boron oxide, iron oxide, gallium oxide, titanium oxide, rare earth oxides, indium oxide and vanadium oxide, preferably boron oxide or a combination of boron oxide and at least one selected from the group consisting of alumina, iron oxide, gallium oxide, titanium oxide, rare earth oxides, indium oxide and vanadium oxide, more preferably boron oxide or a combination of boron oxide and alumina, more preferably boron oxide.

According to one embodiment of the present invention, the first oxide is silica, and the second oxide is boron oxide.

According to another embodiment of the present invention, the first oxide is at least one selected from the group consisting of silica and germanium dioxide, and the second oxide is boron oxide.

According to another embodiment of the present invention, the first oxide is silica, and the second oxide is at least one selected from the group consisting of boron oxide and alumina.

According to one embodiment of the present invention, if multiple oxides are used

in combination, the ratio by molar between each two oxides is generally 1-99.6:99-0.4, preferably 33-99.5:67-0.5, more preferably 50-99:50-1, more preferably 60-99:40-1, more preferably 66-98:34-2, more preferably 66-97:34-3.

According to the present invention, the molecular sieve in the calcined form has X ray diffraction pattern as substantially illustrated in the following table.

2 θ (°) ^(a)	d-spacing (Å)	Relative intensity (I/I ₀ × 100)
6.50	13.62±0.63	w-s
7.98	11.09±0.42	s-vs
9.36	9.45±0.30	m
11.27	7.85±0.21	w-m
20.02	4.43±0.07	s
22.65	3.92±0.05	vs
24.13	3.69±0.05	vs
26.45	3.37±0.04	w-m
27.92	3.19±0.03	w-m
35.95	2.50±0.02	m

a:±0.3°.

Further, the X-ray diffraction pattern further includes X-ray diffraction peaks as substantially illustrated in the following table,

2 θ (°) ^(a)	d-spacing (Å)	Relative intensity (I/I ₀ × 100)
14.12	6.27±0.13	w
16.12	5.50±0.10	w
18.82	4.71±0.07	w
37.52	2.40±0.02	w

a:±0.3°.

According to the present invention, the SCM-10 molecular sieve has a specific surface area (by the BET method) of 250-600m²/g, preferably 280-450m²/g.

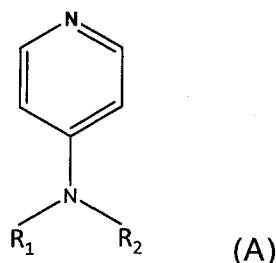
According to the present invention, the SCM-10 molecular sieve has a micropore volume (by the t-plot method) of 0.05-0.25 cm³/g, preferably 0.08-0.18 cm³/g.

According to the present invention, the SCM-10 molecular sieve has a pore size (by the Argon adsorption method) of 0.6-0.73nm, preferably 0.62-0.68nm.

According to the present invention, the SCM-10 molecular sieve can be produced in line with the following process. In view of this, the present invention further relates to a process for producing an SCM-10 molecular sieve, including a step of crystallizing a mixture comprising a first oxide source, a second oxide source, an organic template and water (hereinafter referred to as the mixture) under crystallization conditions to obtain the molecular sieve (hereinafter referred to as the crystallization step).

According to the present invention, in the process for producing the molecular sieve,

the organic template may be a compound represented by the following formula (A), a quaternary ammonium salt thereof or a quaternary ammonium hydroxide thereof, preferably 4-dimethylamino pyridine.



According to the present invention, in the formula (A), R_1 and R_2 may be identical to or different from each other, each independently representing a C_{1-8} alkyl, preferably a C_{1-4} alkyl, more preferably a C_{1-2} alkyl, most preferably both methyl.

According to the present invention, as the quaternary ammonium salt of the compound represented by the formula (A), there may be exemplified a quaternary nitrogen (N^+) structure obtained by additionally bonding a C_{1-8} alkyl (preferably a C_{1-4} alkyl, more preferably a C_{1-2} alkyl or methyl) to the N atom in addition to the groups R_1 and R_2 . As the counterion of the quaternary nitrogen, there may be exemplified a halo ion like Br^- , but not limiting thereto.

According to the present invention, as the quaternary ammonium hydroxide of the compound represented by the formula (A), there may be exemplified a quaternary nitrogen (N^+) structure obtained by additionally bonding a C_{1-8} alkyl (preferably a C_{1-4} alkyl, more preferably a C_{1-2} alkyl or methyl) to the N atom in addition to the groups R_1 and R_2 . As the counterion of the quaternary nitrogen, a hydroxyl ion (OH^-) is needed.

According to the present invention, in the process for producing the molecular sieve, the crystallization step may be conducted in any way known in this field, there may be exemplified a way wherein the first oxide source, the second oxide source, the organic template and water are mixed in predetermined ratios, and then the obtained mixture is allowed to hydrothermally crystallize under crystallization conditions.

According to the present invention, in the process for producing the molecular sieve, the crystallization conditions include: a crystallization temperature of 140-210 degrees Celsius, preferably 150-190 degrees Celsius, more preferably 160-180 degrees Celsius, a crystallization duration of 10 hrs to 10 days, preferably 1-7 days, more preferably 1-5 days, more preferably 1-3 days.

According to the present invention, in the process for producing the molecular sieve, the first oxide source is at least one selected from the group consisting of a silicon source and a germanium source, preferably a silicon source.

According to the present invention, in the process for producing the molecular sieve, the second oxide source is at least one selected from the group consisting of an aluminum source, a boron source, an iron source, a gallium source, a titanium source, a rare earth source, an indium source and a vanadium source, preferably a boron source or a combination of a boron source and at least one selected from the group consisting of an aluminum source, an iron source, a gallium source, a titanium source, a rare earth source, an indium source and a vanadium source, more preferably a boron source or a combination of a boron

source and an aluminum source, more preferably a boron source.

According to the present invention, in the process for producing the molecular sieve, as the first oxide source, any corresponding oxide source known in this field for this purpose can be used. For example, if the first oxide is silica, as the first oxide source (silicon source), there may be exemplified silicic acid, silica gel, silica sol, water glass or tetralkoxysilane. If the first oxide is germanium dioxide, as the first oxide source (germanium source), there may be exemplified tetralkoxy germanium, germanium dioxide, germanium nitrate. As the first oxide source, one kind or a mixture of two or more kinds at any ratio therebetween could be used.

According to the present invention, in the process for producing the molecular sieve, as the second oxide source, any corresponding oxide source known in this field for this purpose can be used, including but not limiting to the oxides, alkoxides, oxometallates, acetates, oxalates, ammonium salts, sulfates and nitrates of the corresponding metal in the second oxide. For example, if the second oxide is alumina, as the second oxide source (aluminum source), there may be exemplified aluminum hydroxide, sodium aluminate, aluminum salts, aluminum alkoxides, kaolin or montmorillonite. As aluminum salts, there may be exemplified aluminum sulfate, aluminum nitrate, aluminum carbonate, aluminum phosphate, aluminum chloride or alum. As aluminum alkoxides, there may be exemplified aluminum isopropoxide, aluminum ethoxide, aluminum butoxide. If the second oxide is boron oxide, as the second oxide source (boron source), there may be exemplified boric acid, borate salt, borax, diboron trioxide. If the second oxide is iron oxide, as the second oxide source (iron source), there may be exemplified ferric nitrate, ferric chloride, iron oxide. If the second oxide is gallium oxide, as the second oxide source (gallium source), there may be exemplified gallium nitrate, gallium sulfate, gallium oxide. If the second oxide is titanium oxide, as the second oxide source (titanium source), there may be exemplified titanium tetralkoxide, titania, titanium nitrate. If the second oxide is rare earth oxides, as the second oxide source (rare earth source), there may be exemplified lanthanum oxide, neodymium oxide, yttrium oxide, cerium oxide, lanthanum nitrate, neodymium nitrate, yttrium nitrate, ceric ammonium sulfate. If the second oxide is indium oxide, as the second oxide source (indium source), there may be exemplified indium chloride, indium nitrate, indium oxide. If the second oxide is vanadium oxide, as the second oxide source (vanadium source), there may be exemplified vanadium chloride, ammonium metavanadate, sodium vanadate, vanadium dioxide, vanadyl sulfate. As the second oxide source, one kind or a mixture of two or more kinds at any ratio therebetween could be used.

According to one embodiment of the present invention, for the first oxide source, if multiple oxide sources are used in combination, the ratio by molar between each two oxide sources is generally 1-99.6:99-0.4, preferably 33-99.5:67-0.5, more preferably 50-99:50-1, more preferably 60-99:40-1, more preferably 66-98:34-2, more preferably 66-97:34-3.

According to one embodiment of the present invention, for the second oxide source, if multiple oxide sources are used in combination, the ratio by molar between each two oxide sources is generally 1-99.6:99-0.4, preferably 33-99.5:67-0.5, more preferably 50-99:50-1, more preferably 60-99:40-1, more preferably 66-98:34-2, more preferably 66-97:34-3.

According to one embodiment of the present invention, in the process for

producing the molecular sieve, from the point of facilitating obtaining the molecular sieve of the present invention, the mixture does not contain an alkaline source. As the alkaline source, there may be exemplified alkaline substances excepting the first oxide source, the second oxide source and the organic template, specifically there may be exemplified any alkaline source conventionally used in this field for alkalizing the reaction system, more specifically there may be exemplified inorganic alkali containing alkali metal or alkaline earth metal as the cation, especially NaOH and KOH. Herein, by "does not contain an alkaline source", it means not intentionally or on purpose introducing into the mixture an alkaline source.

According to one embodiment of the present invention, in the process for producing the molecular sieve, from the point of facilitating obtaining the molecular sieve of the present invention, the mixture does not contain a fluorine source. As the fluorine source, there may be exemplified fluoride or an aqueous solution thereof, especially HF. Herein, by "does not contain a fluorine source", it means not intentionally or on purpose introducing into the mixture a fluorine source.

According to one embodiment of the present invention, in the process for producing the molecular sieve, from the point of facilitating obtaining the molecular sieve of the present invention, at least at the beginning of the crystallization step, preferably throughout the crystallization step, the mixture is controlled at pH=6-14, preferably pH=7-14, more preferably 8-14, more preferably 8.5-13.5, more preferably 9-12.

According to the present invention, in the process for producing the molecular sieve, the ratio by molar between the first oxide source (as the first oxide), the second oxide source (as the second oxide), the template and water is 1:(0.025-1/3):(0.01-1.0):(4-50), preferably 1:(1/30-1/3):(0.02-0.9):(4-40), more preferably 1:(1/30-1/5):(0.04-0.8):(4-30), with the proviso that the ratio by molar of the first oxide source (as the first oxide) to the second oxide source (as the second oxide) is less than 40, preferably less than 30.

According to this invention, in the process, upon completion of the crystallization step, any separation method conventionally known in this field may be used to isolate a molecular sieve from the obtained reaction mixture as the final product, whereby obtaining the molecular sieve of the present invention. As the separation method, there may be exemplified a method wherein the obtained reaction mixture is filtered, washed and dried. Herein, filtering, washing and drying can be conducted in any manner conventionally known in this field. Specifically, as the filtration, there may be exemplified a method of simply suction filtering the obtained reaction mixture. As the washing, there may be exemplified a method of washing with deionized water. As the drying temperature, there may be exemplified a temperature of 40-250 degrees Celsius, preferably a temperature of 60-150 degrees Celsius, as the drying duration, there may be exemplified a duration of 8-30h, preferably a duration of 10-20h. The drying could be conducted under the normal pressure or a reduced pressure.

According to this invention, in the process, if needed, the obtained molecular sieve could be calcinated so as to remove the organic template and any water therefrom, whereby obtaining a calcinated molecular sieve (i.e. the molecular sieve in the calcined form), which corresponds to the molecular sieve of the present invention as well. The calcination could be conducted in any manner conventionally known

in this field, for example, the calcination temperature is generally 300-800 degrees Celsius, preferably 400-650 degrees Celsius, while the calcination duration is generally 1-10h, preferably 3-6h. Further, the calcination is generally conducted under an oxygen containing atmosphere, for example, under the air atmosphere or under oxygen atmosphere

According to the present invention, the obtained molecular sieves may be used in any physical form, for example, powder, particulate or a molded product (for example, strip, clover). These physical forms can be obtained in any manner conventionally known in this field, without any specific limitation thereto.

The molecular sieve according to this invention may be combined with other material, whereby obtaining a molecular sieve composition. As these other materials, there may be exemplified an active material and a non-active material. As the active material, there may be exemplified synthesized zeolites and natural zeolites, as the non-active material (generally referred to as binder), there may be exemplified clay, white earth, silica gel and alumina. As these other materials, one kind or a mixture of two or more kinds at any ratio therebetween could be used. As the amount of these other materials, any amount conventional used in this field could be used, without any specific limitation thereto.

The molecular sieve or the molecular sieve composition of the present invention can be used as an adsorbent, for example, that to be used in a gaseous or liquid phase to isolate at least one component from a mixture made of multiple components. In this way, a part of or substantially all of the at least one component can be isolated from the mixture. Specifically there may be exemplified a way wherein the molecular sieve or the molecular sieve composition is made to contact the mixture, whereby selectively adsorbing this component.

The molecular sieve or the molecular sieve composition of the present invention may be directly or after treated or converted (for example after ion exchanged) in a way conventionally used in this field regarding a molecular sieve used as a catalyst for converting an organic compound (or as a catalytic active component thereof). Specifically, according to the present invention, for example, reactants can be made to conduct a predetermined reaction in the presence of the catalyst for converting an organic compound to obtain the aimed product. As the predetermined reaction, there may be exemplified isomerization of normal paraffins, liquid phase alkylation between benzene and ethylene to produce ethyl benzene, liquid phase alkylation between benzene and propene to produce iso-propyl benzene, butene isomerization, naphtha cracking reaction, alkylation of benzene with ethanol, cyclohexene hydration, toluene disproportionation to produce p-xylene, alkylation of toluene with methanol to produce p-xylene or disproportionation of iso-propyl naphthalene to produce 2,6-di(iso-propyl) naphthalene. In view of this, as the catalyst for converting an organic compound, there may be exemplified an alkane isomerization catalyst, a catalyst for the alkylation between olefins and aromatics, an olefin isomerization catalyst, a naphtha cracking catalyst, a catalyst for the alkylation between alcohols and aromatics, an olefin hydration catalyst or an aromatic disproportionation catalyst.

EXAMPLE

The following examples illustrate rather than limit this invention.

Example 1

10.995g of the organic template 4-dimethylamino pyridine, 54.0g water, 1.879g boric

acid, 22.5g silica sol (containing SiO₂ 40wt%) were mixed till homogeneous, to obtain a mixture with a ratio (ratio by molar) of:

SiO₂/B₂O₃=10

4-dimethylamino pyridine /SiO₂=0.6

H₂O/SiO₂=25

and then charged into a stainless steel reactor, under stirring at 175 degrees Celsius crystallized for 3 days, upon completion of the crystallization, filtered, washed, dried to obtain a molecular sieve precursor, and then the precursor was at 650 degrees Celsius in air calcined for 6 hours to obtain a molecular sieve.

The XRD data of the resultant molecular sieve were listed in Table 1, and the XRD pattern was as illustrated in Fig.1.

The resultant molecular sieve has a specific surface area of 297m²/g, a micropore volume of 0.11 cm³/g.

If determined by inductively coupled plasma-atomic emission spectroscopy (ICP), the sample after calcined has SiO₂/B₂O₃=16.2.

Table 1

2θ / °	d / Å	I/I ₀ × 100
6.555	13.472	36.2
8.053	10.9694	85
9.376	9.4243	29.9
11.35	7.7899	10
14.145	6.2559	7.4
16.278	5.4409	14.5
16.831	5.2633	13.6
18.624	4.7604	16.3
18.956	4.6778	13.1
20.221	4.3879	65
22.825	3.8928	81.2
24.225	3.6709	100
26.749	3.33	20.4
28.146	3.1678	20.1
32.118	2.7846	3.2
32.683	2.7377	3.7
33.976	2.6364	2.8
36.156	2.4823	24.8
37.71	2.3835	3.7

Example 2

36.651g of the organic template 4-dimethylamino pyridine, 45g water, 3.488g germanium dioxide, 3.34g boric acid, 75g silica sol (containing SiO₂ 40wt%) were mixed till homogeneous, to obtain a mixture with a ratio (ratio by molar) of:

$$(\text{SiO}_2 + \text{GeO}_2) / \text{B}_2\text{O}_3 = 20$$

$$\text{4-dimethylamino pyridine} / \text{SiO}_2 = 0.6$$

$$\text{H}_2\text{O} / \text{SiO}_2 = 10$$

and then charged into a stainless steel reactor, under stirring at 180 degrees Celsius crystallized for 2 days, upon completion of the crystallization, filtered, washed, dried, to obtain a molecular sieve in the synthesized form.

The XRD data of the resultant molecular sieve were listed in Table 2 while the XRD pattern is similar to Fig.1.

The resultant molecular sieve has a specific surface area of 342m²/g, a micropore volume of 0.12 cm³/g.

If determined by inductively coupled plasma-atomic emission spectroscopy (ICP), the sample after calcined has SiO₂/B₂O₃=25.2.

Table 2

2θ / °	d / Å	I/I ₀ × 100
6.517	13.5515	19.2
7.993	11.0521	71.8
9.336	9.465	21.7
11.331	7.8028	23.7
14.13	6.2626	6.7
16.218	5.4609	12.2
16.671	5.3134	12.4
18.506	4.7904	14.4
18.801	4.716	10.7
20.123	4.4091	60.8
22.765	3.9029	71.7
24.146	3.6828	100
26.473	3.3641	23.4
27.952	3.1894	20
29.25	3.0507	2.9
31.955	2.7984	3.2
32.668	2.7389	5.1
33.057	2.7076	3.7
34.678	2.5846	4

36.038	2.4901	26.2
36.924	2.4324	4.2
37.517	2.3953	4.3
39.017	2.3066	3.2

Example 3

109.95g of the organic template 4-dimethylamino pyridine, 540g water, 9.394g boric acid, 3.939g aluminum hydroxide, 225.0g silica sol (containing SiO₂ 40wt%) were mixed till homogeneous, to obtain a mixture with a ratio (ratio by molar) of:

$$\text{SiO}_2/(\text{B}_2\text{O}_3+\text{Al}_2\text{O}_3)=15$$

$$\text{4-dimethylamino pyridine /SiO}_2=0.6$$

$$\text{H}_2\text{O/SiO}_2=25$$

and then charged into a stainless steel reactor, under stirring at 170 degrees Celsius crystallized for 3 days, upon completion of the crystallization, filtered, washed, dried, and then the precursor was at 650 degrees Celsius in air calcined for 6 hours to obtain a molecular sieve.

The XRD data of the resultant molecular sieve were listed in Table 3 while the XRD pattern is similar to Fig.1.

The resultant molecular sieve has a specific surface area of 321m²/g, a micropore volume of 0.13 cm³/g.

If determined by inductively coupled plasma-atomic emission spectroscopy (ICP), the sample after calcined has SiO₂/B₂O₃=25.2, SiO₂/Al₂O₃=63.0.

Table 3

2θ / °	d / Å	I/I ₀ × 100
6.612	13.3574	39.2
8.034	10.9964	95.6
9.395	9.4055	31.6
11.329	7.8042	10.7
14.145	6.2561	8.4
16.201	5.4666	15.9
16.73	5.2947	15.9
18.882	4.696	11.7
20.182	4.3964	63.4
22.805	3.8963	74.4
24.186	3.6768	100
26.453	3.3666	25.4
28.011	3.1828	20.5

31.936	2.8	2.9
32.961	2.7152	4
34.496	2.5978	3.1
36.058	2.4888	20.8
37.729	2.3823	5.2

Example 4

Similar to Example 2, except that $\text{SiO}_2/\text{GeO}_2=19.2$, $(\text{SiO}_2+\text{GeO}_2)/\text{B}_2\text{O}_3=35.2$, 4-dimethylamino pyridine / $\text{SiO}_2=0.8$, $\text{H}_2\text{O}/\text{SiO}_2=25$, at 170 degrees Celsius crystallized for 70 hours.

The XRD data of the resultant molecular sieve were listed in Table 4 while the XRD pattern is similar to Fig.1.

The resultant molecular sieve has a specific surface area of $297\text{m}^2/\text{g}$, a micropore volume of $0.11\text{ cm}^3/\text{g}$.

If determined by inductively coupled plasma-atomic emission spectroscopy (ICP), the sample after calcined has $\text{SiO}_2/\text{B}_2\text{O}_3=35.2$.

Table 4

$2\theta / ^\circ$	$d / \text{\AA}$	$I/I_0 \times 100$
6.575	13.4316	21.3
8.034	10.9958	79.5
9.377	9.4242	25.5
11.425	7.7385	28.7
13.387	6.6087	3
14.148	6.2549	6.8
16.24	5.4536	12.8
16.75	5.2885	14.8
18.546	4.7802	14.2
18.921	4.6864	11.4
20.22	4.388	63.2
21.533	4.1233	2.4
22.844	3.8896	74.9
24.225	3.6709	100
26.572	3.3518	23.9
28.07	3.1762	21
29.271	3.0486	3.5
32.012	2.7935	2.7

33.02	2.7105	4.1
34.516	2.5964	4.7
36.136	2.4836	25.7
36.95	2.4308	2.9
37.637	2.3879	5
39.155	2.2988	3.3

Example 5

Similar to Example 1, except that $\text{SiO}_2/\text{B}_2\text{O}_3=27$, 4-dimethylamino pyridine / $\text{SiO}_2=0.3$, $\text{H}_2\text{O}/\text{SiO}_2=25$, at 170 degrees Celsius crystallized for 3 days.

The XRD data of the resultant molecular sieve were listed in Table 5 while the XRD pattern is similar to Fig.1.

The resultant molecular sieve has a specific surface area of $343\text{m}^2/\text{g}$, a micropore volume of $0.13\text{ cm}^3/\text{g}$.

If determined by inductively coupled plasma-atomic emission spectroscopy (ICP), the sample after calcined has $\text{SiO}_2/\text{B}_2\text{O}_3=31.5$.

Table 5

$2\theta / ^\circ$	$d / \text{\AA}$	$I/I_0 \times 100$
6.575	13.4314	32.8
8.053	10.9703	81.4
9.452	9.3493	29.7
11.309	7.818	10.8
14.167	6.2465	8.8
16.277	5.4412	12.4
16.751	5.2882	15.4
18.587	4.7698	16.4
18.919	4.6869	12.7
20.203	4.3917	64.1
22.864	3.8863	84.4
24.226	3.6708	100
26.711	3.3347	22.1
28.072	3.176	24.5
29.409	3.0346	4
32.08	2.7878	2.3
33.018	2.7107	3.4
34.541	2.5945	3.3

36.154	2.4824	25.3
37.072	2.423	2.3
37.715	2.3831	4

Example 6

Similar to Example 1, except that $\text{SiO}_2/\text{B}_2\text{O}_3=20$, 4-dimethylamino pyridine / $\text{SiO}_2=0.2$, $\text{H}_2\text{O}/\text{SiO}_2=30$, at 170 degrees Celsius crystallized for 80 hours.

The XRD data of the resultant molecular sieve were listed in Table 6 while the XRD pattern is similar to Fig.1.

The resultant molecular sieve has a specific surface area of $347\text{m}^2/\text{g}$, a micropore volume of $0.12\text{ cm}^3/\text{g}$.

If determined by inductively coupled plasma-atomic emission spectroscopy (ICP), the sample after calcined has $\text{SiO}_2/\text{B}_2\text{O}_3=28.0$.

Table 6

$2\theta / ^\circ$	$d / \text{\AA}$	$I/I_0 \times 100$
6.556	13.4705	31.5
8.053	10.9705	69.9
9.377	9.4239	24.2
11.366	7.7789	10.3
14.224	6.2214	8.2
16.296	5.4349	12.9
16.809	5.27	14.7
18.568	4.7747	15.9
18.94	4.6818	13.3
20.222	4.3878	63.1
22.864	3.8863	83.3
24.245	3.668	100
26.672	3.3395	24.6
28.05	3.1784	21.6
29.47	3.0284	3.6
31.208	2.8637	1.9
32.132	2.7834	3.6
32.887	2.7212	3.7
33.941	2.639	2.6
34.674	2.5849	2.5
36.155	2.4824	26.8

36.855	2.4368	2.3
37.677	2.3855	4.2
39.184	2.2971	2.1

Example 7

Similar to Example 1, except that $\text{SiO}_2/\text{B}_2\text{O}_3=10$, 4-dimethylamino pyridine / $\text{SiO}_2=0.4$, $\text{H}_2\text{O}/\text{SiO}_2=17.5$, at 170 degrees Celsius crystallized for 2 days.

The XRD data of the resultant molecular sieve were listed in Table 7 while the XRD pattern is similar to Fig.1.

The resultant molecular sieve has a specific surface area of $357\text{m}^2/\text{g}$, a micropore volume of $0.13\text{ cm}^3/\text{g}$.

If determined by inductively coupled plasma-atomic emission spectroscopy (ICP), the sample after calcined has $\text{SiO}_2/\text{B}_2\text{O}_3=14.2$.

Table 7

$2\theta / ^\circ$	$d / \text{\AA}$	$I/I_0 \times 100$
6.595	13.3912	30.4
8.054	10.9683	80.6
9.416	9.3843	26.7
11.347	7.7917	9.4
14.169	6.2453	7.9
16.3	5.4336	13.8
16.79	5.2761	11.9
18.605	4.7651	15.6
18.923	4.6858	13.5
20.223	4.3875	63.6
22.865	3.8861	82.1
24.264	3.6651	100
26.75	3.3299	22.5
28.091	3.1739	20.4
29.356	3.0399	3.5
32.095	2.7865	4.2
32.924	2.7182	3.7
34.6	2.5903	2.5
36.174	2.4811	26.3
36.967	2.4297	2.5
37.754	2.3808	4.6

Example 8

Similar to Example 1, except that $\text{SiO}_2/\text{B}_2\text{O}_3=5$, 4-dimethylamino pyridine / $\text{SiO}_2=0.5$, $\text{H}_2\text{O}/\text{SiO}_2=25$, at 170 degrees Celsius crystallized for 66 hours.

The XRD data of the resultant molecular sieve were listed in Table 8 while the XRD pattern is similar to Fig.1.

The resultant molecular sieve has a specific surface area of $342\text{m}^2/\text{g}$, a micropore volume of $0.12\text{ cm}^3/\text{g}$.

If determined by inductively coupled plasma-atomic emission spectroscopy (ICP), the sample after calcined has $\text{SiO}_2/\text{B}_2\text{O}_3=12.1$.

Table 8

$2\theta / ^\circ$	$d / \text{\AA}$	$I/I_0 \times 100$
6.594	13.3928	42.3
8.053	10.9692	90.8
9.415	9.386	32.5
11.366	7.7787	12.1
14.168	6.2461	10.4
16.258	5.4473	13.3
16.75	5.2884	16.8
18.567	4.7748	18
18.901	4.6912	14.6
20.203	4.3917	67.1
22.845	3.8895	83
24.263	3.6652	100
26.691	3.3371	25.3
28.032	3.1805	25.1
29.256	3.0501	3.1
32.919	2.7186	4.3
33.933	2.6396	2.7
34.494	2.598	3.2
36.136	2.4836	24
36.914	2.433	3.5
37.715	2.3832	4.7

Example 9

Similar to Example 1, except that $\text{SiO}_2/\text{B}_2\text{O}_3=35$, 4-dimethylamino pyridine / $\text{SiO}_2=0.36$,

H₂O/SiO₂=25, at 170 degrees Celsius crystallized for 3 days.

The XRD data of the resultant molecular sieve were listed in Table 9 while the XRD pattern is similar to Fig.1.

The resultant molecular sieve has a specific surface area of 331m²/g, a micropore volume of 0.12 cm³/g.

If determined by inductively coupled plasma-atomic emission spectroscopy (ICP), the sample after calcined has SiO₂/B₂O₃=39.5.

Table 9

2θ / °	d / Å	I/I ₀ × 100
6.556	13.4713	41
8.055	10.9668	88.3
9.415	9.3855	27.7
11.366	7.7787	9.7
14.169	6.2457	9.4
16.335	5.4221	14.9
16.752	5.2879	15.4
18.566	4.775	16.1
18.915	4.6877	11.8
20.222	4.3877	62.3
22.845	3.8895	78.1
24.245	3.6679	100
26.612	3.3468	25.5
28.05	3.1784	21.5
29.414	3.0341	4
32.092	2.7868	2.9
32.847	2.7244	3.6
34.49	2.5983	3.1
36.155	2.4824	23.6
37.657	2.3867	4.7

Example 10

Similar to Example 3, except that aluminum sulfate was used as the aluminum source, SiO₂/(Al₂O₃+B₂O₃)=20, 4-dimethylamino pyridine /SiO₂=0.4, H₂O/SiO₂=25, at 170 degrees Celsius crystallized for 3 days.

The XRD data of the resultant molecular sieve were listed in Table 10 while the XRD pattern is similar to Fig.1.

The resultant molecular sieve has a specific surface area of 288m²/g, a micropore

volume of 0.09 cm³/g.

If determined by inductively coupled plasma-atomic emission spectroscopy (ICP), the sample after calcined has SiO₂/B₂O₃=35.2, SiO₂/Al₂O₃=63.3.

Table 10

2θ / °	d / Å	I/I ₀ × 100
6.478	13.6332	29.6
7.995	11.0488	100
9.375	9.4258	31.8
11.268	7.8459	9
14.147	6.2551	8.4
16.238	5.4542	13.1
16.714	5.2997	14.1
18.268	4.8524	20.8
18.858	4.7018	12.3
20.123	4.4091	64.3
22.746	3.9063	76
24.166	3.6798	99.9
26.414	3.3714	25.4
27.913	3.1937	19.7
31.974	2.7968	4.3
32.665	2.7392	3.5
34.328	2.6101	3.3
36.074	2.4877	23.5
37.573	2.3918	5.1

Example 11

Similar to Example 1, except that HF was added as the fluorine source, SiO₂/B₂O₃=15, 4-dimethylamino pyridine /SiO₂=0.5, F/SiO₂=0.4, H₂O/SiO₂=25, at 170 degrees Celsius crystallized for 60 hours.

The XRD data of the resultant molecular sieve were listed in Table 11 while the XRD pattern is similar to Fig.1.

The resultant molecular sieve has a specific surface area of 318m²/g, a micropore volume of 0.10 cm³/g.

If determined by inductively coupled plasma-atomic emission spectroscopy (ICP), the sample after calcined has SiO₂/B₂O₃=21.2.

Table 11

$2\theta / ^\circ$	$d / \text{\AA}$	$I/I_0 \times 100$
6.595	13.3909	30.3
8.053	10.9699	100
9.432	9.3693	39.2
11.31	7.817	12.4
14.166	6.2466	8.7
16.258	5.4473	12.8
16.729	5.2952	14.2
18.528	4.7848	15.8
18.883	4.6958	12.9
20.145	4.4042	31.4
22.784	3.8997	48.1
24.168	3.6795	55.9
26.629	3.3448	20.4
27.971	3.1872	20.6
28.507	3.1285	4.8
29.154	3.0606	4.3
32.028	2.7922	1.5
32.783	2.7295	3.7
34.42	2.6034	3.4
36.057	2.4889	10.8
37.598	2.3903	4.2
38.997	2.3077	1.6

Example 12

Similar to Example 1, except that tetra-n-butyl titanate was added as the titanium source, $\text{SiO}_2/\text{B}_2\text{O}_3=10$, $\text{SiO}_2/\text{TiO}_2=30$, 4-dimethylamino pyridine / $\text{SiO}_2=0.8$, $\text{H}_2\text{O}/\text{SiO}_2=25$, at 170 degrees Celsius crystallized for 3 days.

The XRD data of the resultant molecular sieve were listed in Table 12 while the XRD pattern is similar to Fig.1.

The resultant molecular sieve has a specific surface area of $335\text{m}^2/\text{g}$, a micropore volume of $0.10\text{ cm}^3/\text{g}$.

If determined by inductively coupled plasma-atomic emission spectroscopy (ICP), the sample after calcined has $\text{SiO}_2/\text{B}_2\text{O}_3=16.2$, $\text{SiO}_2/\text{TiO}_2=39.1$.

Table 12

$2\theta / ^\circ$	$d / \text{\AA}$	$I/I_0 \times 100$
--------------------	------------------	--------------------

6.457	13.6776	34.2
7.993	11.0518	95
9.337	9.4641	27.1
11.29	7.8307	8.2
14.104	6.274	7
16.199	5.4672	14.5
16.632	5.3259	13.6
18.526	4.7853	14.9
20.122	4.4093	58.9
22.668	3.9195	71.9
24.088	3.6915	100
26.396	3.3738	23.1
27.991	3.185	20.4
31.951	2.7987	3.1
32.594	2.745	4.8
35.959	2.4954	24.3
37.495	2.3966	5.7

Example 13

30g of the molecular sieve produced in Example 3 in the form of powder was ion-exchanged by an aqueous ammonium nitrate solution (with a concentration of 1mol/L) for 4 times, filtered, and dried at 110 degrees Celsius, calcined at 500 degrees Celsius in air for 6 hours. Then, 1.5g of the calcined molecular sieve was charged into a 100ml stainless steel reactor, further introducing therein 35g iso-propyl naphthalene, and closed the reactor. At 250 °C, under 200 rpm stirring, the reaction was conducted for 48 hours. Upon completion of the reaction, the system was cooled to the room temperature, after centrifugally isolating the molecular sieve powder therefrom, the reaction product was analysed on an Agilent 19091N-236 gas chromatograph, indicating an iso-propyl naphthalene conversion of 26.82%, and a total selectivity to the aimed product 2,6-di(iso-propyl) naphthalene and 2,7-di(iso-propyl) naphthalene of 74.88%.

We claim

1. An SCM-10 molecular sieve, having an empirical chemical composition as illustrated by the formula " the first oxide · the second oxide ", wherein the ratio by molar of the first oxide to the second oxide is less than 40, preferably in the range of from 3 to less than 40, more preferably 5-30, the first oxide is at least one selected from the group consisting of silica and germanium dioxide, preferably silica, the second oxide is at least one selected from the group consisting of alumina, boron oxide, iron oxide, gallium oxide, titanium oxide, rare earth oxides, indium oxide and vanadium oxide, preferably boron oxide or a combination of boron oxide and at least one selected from the group consisting of alumina, iron oxide, gallium oxide, titanium oxide, rare earth oxides, indium oxide and vanadium oxide, more preferably boron oxide or a combination of boron oxide and alumina, more preferably boron oxide, and the molecular sieve in the calcined form has X ray diffraction pattern as substantially illustrated in the following table,

2 θ (°) ^(a)	d-spacing (Å) ^(b)	Relative intensity (I/I ₀ × 100)
6.50	13.59	w-s
7.98	11.07	s-vs
9.36	9.45	m
11.27	7.85	w-m
20.02	4.43	s
22.65	3.92	vs
24.13	3.69	vs
26.45	3.37	w-m
27.92	3.19	w-m
35.95	2.50	m

a: ±0.3°, b: changed with 2 θ .

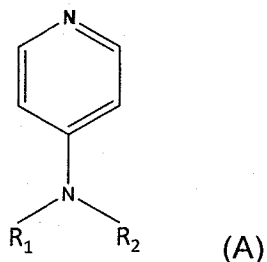
2. The SCM-10 molecular sieve according to Claim 1, wherein the X-ray diffraction pattern further includes X-ray diffraction peaks as substantially illustrated in the following table,

2 θ (°) ^(a)	d-spacing (Å) ^(b)	Relative intensity (I/I ₀ × 100)
14.12	6.27	w
16.12	5.49	w
18.82	4.71	w
37.52	2.40	w

a: ±0.3°, b: changed with 2 θ .

3. A process for producing an SCM-10 molecular sieve, including a step of crystallizing a mixture comprising a first oxide source, a second oxide source, an organic template and water to obtain the molecular sieve, and optionally, a step of calcining the obtained molecular sieve, wherein the organic template is selected from a compound represented by the following formula (A), a quaternary ammonium

salt thereof and a quaternary ammonium hydroxide thereof, preferably 4-dimethylamino pyridine,



wherein R_1 and R_2 may be identical to or different from each other, each independently representing a C_{1-8} alkyl, preferably a C_{1-4} alkyl, more preferably a C_{1-2} alkyl.

4. The process according to Claim 3, wherein the mixture does not contain an alkaline source.

5. The process according to Claim 3, wherein the mixture does not contain a fluorine source.

6. The process according to Claim 3, wherein the mixture has a pH=6-14, preferably pH=7-14, more preferably 8-14, more preferably 8.5-13.5, more preferably 9-12, more preferably 9-11.

7. The process according to Claim 3, wherein the first oxide source is at least one selected from the group consisting of a silicon source and a germanium source, preferably a silicon source, the second oxide source is at least one selected from the group consisting of an aluminum source, a boron source, an iron source, a gallium source, a titanium source, a rare earth source, an indium source and a vanadium source, preferably a boron source or a combination of a boron source and at least one selected from the group consisting of an aluminum source, an iron source, a gallium source, a titanium source, a rare earth source, an indium source and a vanadium source, more preferably a boron source or a combination of a boron source and an aluminum source, more preferably a boron source, the ratio by molar between the first oxide source (as the first oxide), the second oxide source (as the second oxide), the organic template and water is $1:(0.025-1/3):(0.01-1.0):(4-50)$, preferably $1:(1/30-1/3):(0.02-0.9):(4-40)$, more preferably $1:(1/30-1/5):(0.04-0.8):(4-30)$, with the proviso that the ratio by molar of the first oxide source (as the first oxide) to the second oxide source (as the second oxide) is less than 40, preferably less than 30.

8. An SCM-10 molecular sieve composition, comprising the SCM-10 molecular sieve according to Claim 1 or an SCM-10 molecular sieve produced in line with the process according to Claim 3, and a binder.

9. Use of the SCM-10 molecular sieve according to Claim 1, an SCM-10 molecular sieve produced in line with the process according to Claim 3 or the SCM-10 molecular sieve composition according to Claim 8 as an adsorbent or a catalyst for converting an organic compound.

10. Use according to Claim 9, wherein the catalyst for converting an organic compound is at least one selected from the group consisting of an alkane isomerization catalyst, a catalyst for the alkylation between olefins and aromatics, an olefin isomerization catalyst, a naphtha cracking catalyst, a catalyst for the alkylation between alcohols and aromatics, an olefin hydration catalyst and an aromatic disproportionation catalyst.