



(51) International Patent Classification:

B01J 21/04 (2006.01) **C07C 41/01** (2006.01)
B01J 23/80 (2006.01) **C07C 43/04** (2006.01)
B01J 29/06 (2006.01)

(21) International Application Number:

PCT/EP2013/052959

(22) International Filing Date:

14 February 2013 (14.02.2013)

(25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data:

61/598,931 15 February 2012 (15.02.2012) US

(71) Applicant: **BASF SE** [DE/DE]; 67056 Ludwigshafen (DE).

(71) Applicant (for CY only): **BASF SCHWEIZ AG** [CH/CH]; Klybeckstrasse 141, CH-4057 Basel (CH).

(72) Inventors: **SCHAEFER, Alexander**; Maxburgstrasse 20, 67117 Limburgerhof (DE). **VON FEHREN, Thorsten**; Gunterstrasse 16, 68642 Buerstadt (DE). **MADON, Rostam**; 17, Hendrick Road, Flemington, New Jersey 08822 (US).

(74) Agent: **FÉAUX DE LACROIX, Stefan**; Isenbruck Bösl Hörschler LLP, Patentanwälte, Eastsite One, Seckenheimer Landstrasse 4, 68163 Mannheim (DE).

(81) Designated States (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, BN, BR, BW, BY, BZ, CA, CH, CL, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PA, PE, PG, PH, PL, PT, QA, RO, RS, RU, RW, SC, SD, SE, SG, SK, SL, SM, ST, SV, SY, TH, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW.

(84) Designated States (unless otherwise indicated, for every kind of regional protection available): ARIPO (BW, GH, GM, KE, LR, LS, MW, MZ, NA, RW, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, RU, TJ, TM), European (AL, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MK, MT, NL, NO, PL, PT, RO, RS, SE, SI, SK, SM, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

Published:

- with international search report (Art. 21(3))
- before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments (Rule 48.2(h))



WO 2013/120945 A1

(54) Title: CATALYTICALLY ACTIVE BODY FOR THE SYNTHESIS OF DIMETHYL ETHER FROM SYNTHESIS GAS

(57) Abstract: The invention relates to a catalytically active body for the synthesis of dimethyl ether from synthesis gas. In particular, the invention relates to an improved catalytically active body for the synthesis of dimethyl ether, whereby the components of the active body comprise a defined particle size distribution. Furthermore, the present invention concerns a method for the preparation of a catalytically active body, the use of the catalytically active body and a method for preparation of dimethyl ether from synthesis gas.

Catalytically active body for the synthesis of dimethyl ether from synthesis gas

Field of the invention

5 The invention relates to a catalytically active body for the synthesis of dimethyl ether from synthesis gas. In particular, the invention relates to an improved catalytically active body for the synthesis of dimethyl ether, whereby the components of the active body comprise a defined particle size distribution. Furthermore, the present invention concerns a method for the preparation of a catalytically active body, the use of the catalytically active body and a method for preparation of dimethyl ether from synthesis gas.

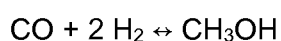
Background of the invention

15 Hydrocarbons are essential in modern life and used as fuel and raw materials, including the chemical, petrochemical, plastics, and rubber industry. Fossil fuels such as oil and natural gas are composed of hydrocarbons with a specific ratio of carbon to hydrogen. In spite their wide application and high demand, fossil fuels also have limitations and disadvantages in the view of being a finite resource and their contribution to global warming if they are burned.

20 Research on alternative fuels was mainly started due to ecological and economical considerations. Among the alternative fuels, dimethyl ether (DME), which is recently discovered as a clean fuel, can be synthesized from syngas that was generated from different primary sources. These primary sources can be natural gas, coal, heavy oil and also biomass. Up to now, only two DME synthesis procedures from synthesis gas have been claimed, whereby one is the traditional methanol synthesis, followed by a dehydration step and the other is a direct conversion of synthesis gas to DME in one single step.

30 Recently, attention has been directed towards the direct synthesis of dimethyl ether from synthesis gas, using a catalytic system that combines a methanol synthesis catalyst and a catalyst for dehydration of said alcohol. It was confirmed on the basis of experimental studies that both, the stage of methanol synthesis and the stage of methanol dehydration, could be conducted simultaneously on one appropriate catalytic system. Depending upon the applied synthesis gas the catalyst might additionally show water gas shift activity.

35 Most known methods of producing methanol involve synthesis gas. Synthesis gas is a mixture of mainly hydrogen, carbon monoxide and carbon dioxide, whereby methanol is produced out of it over a catalyst.

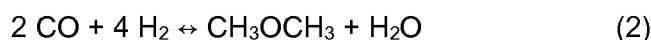
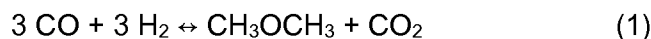


40 In a following step Methanol can be converted into DME by dehydration over an acidic catalyst.



In the direct DME production there are mainly two overall reactions that occur from synthesis gas. These reactions, reaction (1) and reaction (2), are listed below.

5



10 Reaction (1) occurs with the combination of three reactions, which are methanol synthesis reaction, methanol dehydration reaction, and water gas shift reaction:



20 The reaction (1) has a stoichiometric ratio H_2/CO of 1:1 and has some advantages over reaction (2). For example reaction (1) generally allows higher single pass conversions and less energy-consuming in comparison to the removal of water from the system in reaction (2).

25 Methods for the preparation of dimethyl ether are well-known from prior art. Several methods are described in the literature where DME is produced directly in combination with methanol by the use of a catalyst active body in both the synthesis of methanol from synthesis gas and methanol dehydration. Suitable catalysts for the use in the synthesis gas conversion stage include conventionally employed methanol catalyst such as copper and/or zinc and/or chromium-based catalyst and methanol dehydration catalyst.

30 The document US 6,608,114 B1 describes a process for producing DME by dehydrating the effluent stream from the methanol reactor, where the methanol reactor is a slurry bubble column reactor (SBCR), containing a methanol synthesis catalyst that converts a synthesis gas stream comprising hydrogen and carbon monoxide into an effluent stream comprising methanol.

35 Document WO 2008/157682 A1 provides a method of forming dimethyl ether by bimolecular dehydration of methanol produced from a mixture of hydrogen and carbon dioxide, obtained by reforming methane, water, and carbon dioxide in a ratio of about 3 to 2 to 1. Subsequent use of water produced in the dehydration of methanol in the bi-reforming process leads to an overall ratio of carbon dioxide to methane of about 1:3 to produce dimethyl ether.

40

Document WO 2009/007113 A1 describes a process for the preparation of dimethyl ether by catalytic conversion of synthesis gas to dimethyl ether comprising contacting a stream of synthesis gas, comprising carbon dioxide with one or more catalysts active in the formation of methanol and the dehydration of methanol to dimethyl ether, to form a product mixture comprising the components dimethyl ether, carbon dioxide and unconverted synthesis gas, washing the product mixture comprising carbon dioxide and unconverted synthesis gas in a first scrubbing zone with a first solvent rich in dimethyl ether and subsequently washing the effluent from the first scrubbing zone in a second scrubbing zone with a second solvent rich in methanol to form a vapor stream comprising unconverted synthesis gas stream with reduced content of carbon dioxide transferring the vapor stream comprising unconverted synthesis gas stream with reduced carbon dioxide content for the further processing to dimethyl ether.

Document WO 2007/005126 A2 describes a process for the production of synthesis gas blends, which are suitable for conversion either into oxygenates such as methanol or into Fischer-Tropsch-liquids.

The US 6,191,175 B1 describes an improved process for the production of methanol and dimethyl ether mixture rich in DME from essentially stoichiometrically balance synthesis gas by a novel combination of synthesis steps.

In document US 2008/125311 A1 is a catalyst used for producing dimethyl ether, a method of producing the same, and a method of producing dimethyl ether using the same. More particularly, the present invention relates to a catalyst used for producing dimethyl ether comprising a methanol synthesis catalyst produced by adding one or more promoters to a main catalyst comprised of a Cu-Zn-Al metal component and a dehydration catalyst formed by mixing Aluminium Phosphate (AlPO_4) with gamma alumina, a method of producing the same, and a method of producing dimethyl ether using the same, wherein a ratio of the main catalyst to the promoter in the methanol synthesis catalyst in a range of 99/1 to 95/5, and a mixing ratio of the methanol synthesis catalyst to the dehydration catalyst is in a range of 60/40 to 70/30.

The processes for the preparation of dimethyl ether according to the prior art bear the drawbacks that different steps have to be undergone to get an efficient DME production. Besides this, the catalyst used in the method known in prior art does not achieve the thermodynamic possibilities. Therefore it is still desirable to increase the yield of DME in the synthesis gas conversion.

Summary of the invention

The object of the present invention is to provide a catalytically active body that shows the ability to convert CO-rich synthesis gas selectively into dimethyl ether and CO_2 , whereby ideally the yield of the DME is increased in comparison to the state of the art. If the conversion is incom-

plete, the resulting off-gas comprises hydrogen and carbon monoxide preferably in the ratio $H_2/CO \sim 1$. Thus the off-gas can be recycled directly after the separation of the product DME and CO_2 . In addition, it is an object of the present invention to provide a method for the preparation of a catalytically active body and a method for the preparation of dimethyl ether from synthesis gas, comprising the inventive catalytically active body and also the use of the catalytically active body.

These objects are achieved by a catalytically active body for the synthesis of dimethyl ether from synthesis gas, comprising a mixture of:

- (A) 70-90 % by weight of a methanol-active component, selected from the group consisting of copper oxide, aluminium oxide, zinc oxide, amorphous aluminium oxide, ternary oxide or mixtures thereof,
- (B) 10-30 % by weight of an acid component, selected from the group consisting of aluminosilicate, γ -alumina and zeolite or mixtures thereof,
- (C) 0-10 Gew.-% by weight of at least one additive, whereby the sum of the components (A), (B) and (C) is in total 100 % by weight.

Preferably, the ternary oxide is a zinc-aluminium-spinel.

In a preferred embodiment of the catalytically active body the mixture comprises:

- (A) 70-90 % by weight of a methanol-active component, selected from the group consisting of copper oxide, aluminium oxide, zinc oxide, amorphous aluminium oxide, ternary oxide or mixtures thereof, whereby the component (A) has a particle size distribution characterized by a D-10 value of 5-140 μm , a D-50 value of 40-300 μm , and a D-90 value of 180-800 μm ,
- (B) 10-30 % by weight of an acid component, selected from the group consisting of aluminosilicate, γ -alumina and zeolite or mixtures thereof, whereby the component (B) has a particle size distribution characterized by a D-10 value of 5-140 μm , a D-50 value of 40-300 μm , and a D-90 value of 180-800 μm ,
- (C) 0-10 Gew.-% by weight of a at least one additive, whereby the sum of the components (A), (B), and (C) is in total 100 % by weight and the particle size of components (A) and (B) is maintained in the catalytically active body.

This particle size distribution can be determined via state of the art analysis techniques, e.g. via analysis apparatus like Mastersizer 2000 or 3000 by Malvern Instruments GmbH. The particle size distribution in the sense of the invention is characterized by the D10-, D50-, and D-90 value. The definition of D10 is: that equivalent diameter where 10 mass % (of the particles) of the sample has a smaller diameter and hence the remaining 90% is coarser. The definition of D50 and D90 can be derived similarly (see: HORIBA Scientific, A Guidebook to Particle Size Analysis" page 6)

The inventive catalytically active body is characterized by a high turnover of carbon monoxide, preferably at 180°C to 350°C and particular at 200°C to 300°C. For example, a suitable pressure for the synthesis of DME is preferably in the range from 20 to 80 bar and in particular from 30 to 50 bar.

Preferably, the components (A) or (B) have a particle size distribution characterized by a D-10, D-50, and D-90 value of 5-80 µm, 40-270 µm, and 250-800 µm respectively. In a further embodiment the particle size distribution from component (A) can be different from component (B) and (C), in particular, the components (A) or (B) have a particle size distribution characterized by a D-10, D-50, and D-90 value of 5-50 µm, 40-220 µm, and 350-800 µm respectively. In a further embodiment the particle size distribution from component (A) can be different from component (B) and (C).

In the sense of the present invention a catalytically active body can be a body known in the art that contains pores or channels or other features for enlargement of surface, which will help to bring the educts in contact that they can react to the desired product. A catalytically active body in the sense of the present invention can be understood as a physical mixture, whereby the components (A) and (B) contact each other and presenting channels and/or pores between their contact surfaces. Preferably, the components (A) and (B) are not melted or sintered at their contact surfaces.

A methanol-active component in the sense of the present invention is a component which leads to the formation of methanol, starting from hydrogen, carbon monoxide or carbon dioxide or mixtures thereof. Preferably, the methanol-active compound is a mixture of copper oxide, aluminium oxide and zinc oxide, whereby copper oxide can consist of all kinds of oxides of copper. In particular, copper has the oxidation state (I) or (II) in the oxide. The aluminium oxide according to the present invention can also be referred to γ-alumina or corundum, whereby zinc in zinc oxide in the sense of the present invention preferably has the oxidation state (II).

In a preferred embodiment of the catalytically active body, the component (A) comprises 50-80 % by weight of copper oxide, 15-35 % by weight of ternary oxide and 15-35 % by weight of zinc oxide and the sum of which is in total 100 % by weight. In particular the component (A) com-

prises 65-75 % by weight of copper oxide, 20-30 % by weight of ternary oxide and 20-30 % by weight of zinc oxide and the sum of which is in total 100 % by weight.

5 In a preferred embodiment of the catalytically active body, the component (A) comprises 50-80 % by weight of copper oxide, 2-8 % by weight of boehmite and 15-35 % by weight of zinc oxide and the sum of which is in total 100 % by weight. In particular the component (A) comprises 65-75 % by weight of copper oxide, 3-6 % by weight of boehmite and 20-30 % by weight of zinc oxide and the sum of which is in total 100 % by weight.

10 In a preferred embodiment of the catalytically active body, the component (A) comprises 50-80 % by weight of copper oxide, 2-8 % by weight of amorphous aluminium oxide and 15-35 % by weight of zinc oxide and the sum of which is in total 100 % by weight. In particular the component (A) comprises 65-75 % by weight of copper oxide, 3-6 % by weight of amorphous aluminium oxide and 20-30 % by weight of zinc oxide and the sum of which is in total 100 % by weight.

15 In a preferred embodiment of the catalytically active body, the component (A) comprises 50-80 % by weight of copper oxide, 2-8 % by weight of aluminium oxide and 15-35 % by weight of zinc oxide and the sum of which is in total 100 % by weight. In particular the component (A) comprises 65-75 % by weight of copper oxide, 3-6 % by weight of aluminium oxide and 20-30 % by weight of zinc oxide and the sum of which is in total 100 % by weight.

25 In the sense of the present invention an acid component (B) is selected from the group consisting of aluminosilicate, γ -alumina and zeolite or mixtures thereof. Aluminosilicate comprise minerals composed of aluminium, silicon and oxygen. They are the major components of kaolin and other clay minerals like for example Halloysite, Kaolinite, Illite, Montmorillonite, Vermiculite, Talc, Palygorskite, Pyrophyllite, which are also suitable as component (B).

30 In a preferred embodiment, the zeolite comprises 35-55 % by weight of silicon, 0.15-4 % by weight of aluminium, 45-65 % by weight of oxygen and where appropriate 0-0.3 by weight of sodium and the sum of which is in total 100 % by weight. Preferably the zeolite comprises 40-52 % by weight of silicon, 0.8-3.5 % by weight of aluminium, 50-60 % by weight of oxygen and where appropriate 0-0.3 % by weight of sodium and the sum of which is in total 100 % by weight. In a preferred embodiment zeolite can be Zeolithe A, Zeolithe X, Zeolithe Y, Zeolithe L, Mordenit, ZSM-5, ZSM-11.

35 In the sense of the present invention an additive (C) can be a structure-promoter like but not limited a thermally decomposable compound like polymers, wood dust, flour, graphite, film material, a painting, straw, stearic acid, palmitic acid, celluloses or a combination thereof. For example, the structure-promotor can help to build up pores or channels.

40

In a preferred embodiment the catalytically active body consists of (A) 70-90 % by weight of a methanol-active component and 10-30 % by weight of a zeolite (B) and the sum of (A) and (B) being in total 100 % by weight. Preferably the catalytically active body consists of (A) 75-85 % by weight of a methanol-active component and 15-25 % by weight of a zeolite (B) and the sum of (A) and (B) being in total 100 % by weight. One advantage of this composition is that the turnover of the reaction of the methanol-active compound (A) and the acid compound (B) is favored, because the highly integrated catalyst system combines the methanol synthesis, water gas shift activity, and methanol dehydration catalyst in a close proximity. Therefore an optimum efficiency can be obtained.

In a preferred embodiment the catalytically active body is a pellet with a size in the range from 1 x 1 mm to 10 x 10 mm, preferably in the range from 2 x 2 mm to 7 x 7 mm. The pellet is obtained by pressing the mixture of the components (A), (B) and (C) to a pellet. In the sense of the present invention a pellet can be obtained by pressing the components (A), (B) and optionally (C) under force to the pellet, whereby the shape of the pellet can be ring-shaped, star-shaped or spherical-shaped. Furthermore the pellet can be hollow strings, trilobes, multihole pellets, extrudates and alike.

The present invention further relates to a method for the preparation of a catalytically active body, comprising the step:

c) preparation a physical mixture comprising:

- (A) 70-90 % by weight of a methanol-active component, selected from the group consisting of copper oxide, aluminium oxide, zinc oxide, amorphous aluminium oxide, ternary oxide or mixtures thereof,
- (B) 10-30 % by weight of an acid component, selected from the group consisting of aluminosilicate, γ -alumina and zeolite or mixtures thereof,
- (C) 0-10 Gew.-% by weight of a at least one additive, whereby the sum of the components (A), (B) and (C) is in total 100 % by weight.

In this context, the meanings of the features are the same as for the catalytically active body already mentioned.

In the sense of the present invention preparing a physical mixture means that the different compounds (A), (B) and (C) are brought in contact without further chemical modification.

In a preferred embodiment of the method, the component (A) has a particle size distribution characterized by a D-10 value of 5-80 μm , a D-50 value of 40-270 μm , and a D-90 value of 250-800 μm , whereby the component (B) has a particle size distribution characterized by a D-10 value of 5-80 μm , a D-50 value of 40-270 μm , and a D-90 value of 250-800 μm and the particle size distribution of components (A) and (B) is maintained in the catalytically active body. In a particular embodiment of the method, the component (A) has a particle size distribution characterized by a D-10 value of 5-50 μm , a D-50 value of 40-220 μm , and a D-90 value of 350-800 μm , whereby the component (B) has a particle size distribution characterized by a D-10 value of 5-50 μm , a D-50 value of 40-220 μm , and a D-90 value of 350-800 μm and the particle size distribution of components (A) and (B) is maintained in the catalytically active body.

In a preferred embodiment the method comprising further the steps:

- a) precipitation a copper-, zinc-, or aluminiumsalt or a mixture thereof,
- b) calcination of the product obtained in step a).

Preferably, the steps a) and b) are carried out before the step c). Preferably, the obtained product consists after step c) of 70-90 % by weight of a methanol-active component (A), selected from the group consisting of copper oxide, aluminium oxide and zinc oxide or mixtures thereof, 10-30 % by weight of an acid component (B), selected from the group consisting of aluminosilicate, γ -alumina and zeolite or mixtures thereof. Preferably, after step c) the component (A) has a particle size distribution characterized by a D-10 value of 5-140 μm , a D-50 value of 40-300 μm , and a D-90 value of 180-800 μm and the component (B) has a particle size distribution characterized by a D-10 value of 5-140 μm , a D-50 value of 40-300 μm , and a D-90 value of 180-800 μm .

Preferably, the method comprises at least spray drying, filtration, grinding, sieving or further steps, known in the art to create a catalytically active body, or combinations thereof.

In the sense of the present invention precipitation is a method for the formation of a solid in a solution or inside another solid during a chemical reaction or by diffusion in a solid. The precipitation techniques are known in the art, see also Ertl, Gerhard, Knözinger, Helmut, Schüth, Ferdi, Weitkamp, Jens (Hrsg.) "Handbook of Heterogeneous Catalysis" 2nd edition 2008, Wiley VCH Weinheim, Vol. 1, chapter 2. For example salts of copper, zinc or aluminium are dissolved in a solvent, in particular water. At least two of the salts of either copper, zinc, or aluminium can be heated and a basic solution can be prepared and added. Both solutions can be added in parallel to the template, till the salt-solution is consumed. After this the suspension is vacuumed, dried, and calcinated under air flow.

Preferred anions in the salts for copper, zinc, or aluminium are selected from the group consisting of nitrate, acetate, halide, carbonate, nitrite, sulfate, sulfite, sulfide, phosphate ion or silicate. In particular, salts of copper, zinc or aluminium formed with the above mentioned anions can be converted into oxides of copper, zinc or aluminium applying a calcination step.

5

Calcination in the sense of the present invention can be understood as a thermal treatment process applied to ores and other solid materials to bring about a thermal decomposition, phase transition, or removal of a volatile fraction. The calcination process normally takes place at temperatures below the melting point of the product materials. Mostly it is done under oxygen-containing atmosphere. In some cases the calcination can be performed under inert atmosphere (e.g. nitrogen). Calcination is to be distinguished from roasting, in which more complex gas-solid reactions take place between the furnace atmosphere and the solids.

10

In particular the components (A), (B) and (C) can be compacted in a presser, a squeezer, a crusher or a squeezing machine, preferably after step a), b) or c). Compacting in the sense of the present invention can mean that particles of a defined particle size distribution are pressed to bodies, which have a diameter in the range of 1 to 10 mm and a height of 1 to 10 mm. Preferably the particle size distribution is still left after the compacting.

15

In a preferred embodiment of the method a pellet is formed, preferably with a size in the range from 1 x 1 mm to 10 x 10 mm, especially in the range from 2 x 2 mm to 7 x 7 mm.

20

In a preferred embodiment of the method, the components (A) and (B) are independently pressed through at least one sieve, whereby the sieve exhibits a mesh size from 0.005 to 5 mm in order to obtain a particle size distribution characterized by a D-10 value of 5-140 μm , a D-50 value of 40-300 μm , and a D-90 value of 180-800 μm . Preferably the sieve exhibits a mesh size from 0.005 to 1.50 mm and in particular a mesh size from 0.005 to 0.80 mm. In particular the particles can also exhibit particle size distribution characterized by a D-10, D-50, and D-90 value of 5-140 μm , 40-300 μm , and 180-800 μm respectively. Thereby the components (A) and (B) can be obtained as particles with a defined particle size distribution, also referred in the sense of the present invention as a split-fraction. Because of this split-fraction the CO-conversion increases when synthesis gas contacts the split-fraction. Furthermore the yield of the DME increases, when synthesis gas is converted to DME by the catalytically active body. Preferably, this step is included in step c).

25

30

35

In a further embodiment component (C) is admixed to the components (A) and (B) before sieving.

40

In a preferred embodiment of the preparation of a catalytically active body at least three different sieves are used, whereby the components (A) and (B) are pressed in direction from the sieve with the biggest mesh size to the sieve with the smallest mesh size. By using three sieves

with different mesh sizes the components (A) and (B) are initially pressed into the sieve with the biggest mesh size, which results in particles with the maximal size of the mesh size of this sieve. Preferably, the particle size distribution of the components (A) and (B) is characterized by a D-10 value of 5-140 μm , a D-50 value of 40-300 μm , and a D-90 value of 180-800 μm . These particles can also be broken during the first sieving, so that smaller particles are obtained, which can go through the second sieve, which exhibits a smaller mesh size. Therefore a first fraction with a specific particle size distribution can be obtained before the second sieve. This fraction can also be used as a catalytically active body. Besides this, the particles which go through the second sieve with a mesh size smaller than the first sieve, but bigger than the third sieve, can be obtained behind the second sieve and before the smallest sieve with the smallest mesh size. Also here the particles obtained after the second (middle) sieve can be used as a catalytically active body. In addition to this, the particles obtained after the sieve with the biggest mesh size could be pressed through the second sieve in order to reduce the particle size.

In a preferred embodiment of the method according to the present invention in step a) a part of the component (A) is prepared by precipitation reaction and/or calcination. In the sense of the present invention precursors of the component (A) in form of a salt in a solution can be heated and adjusted to a defined pH-value. After this, a calcination step can be carried out, whereby calcination is known from prior art. These steps can lead to the desired component (A).

In a preferred embodiment of the inventive method at least one part of component (A) is precipitated and whereby at least another part of component (A), which is not subjected to the first precipitation, is added to the precipitate. Preferably, it is added by spray drying or precipitation.

In a preferred embodiment of the inventive method, the method further comprises the step d) adding a mixture of hydrogen and nitrogen to component (A) and/or (B). Preferably the content of the volume of the hydrogen is less than 5% in the mixture.

The present invention further relates to a method for the preparation of dimethyl ether from synthesis gas comprising at least the steps:

- e) reducing the catalytically active body
- f) contacting the catalytically active body in a reduced form with hydrogen and at least one of carbon monoxide or carbon dioxide.

In a further embodiment the method comprising the steps:

- g) providing the inventive catalytically active body, in particular in form of pellets
- h) disposing the catalytically active body in a reactor,
- i) reducing the catalytically active body at a temperature between 140 $^{\circ}\text{C}$ and 240 $^{\circ}\text{C}$ with at least a nitrogen and hydrogen mixture.

The present invention further relates to the use of a catalytically active body according to the present invention for the preparation of dimethyl ether. Preferred admixtures and preferred methods for the preparation are mentioned above and also included in the use.

5

The present invention is further illustrated by the following examples:

A) Synthesis of the methanol-active compounds:

10 1. Example

Two solutions are prepared for the precipitation of the components:

Solution 1: A solution of 1.33 kg copper nitrate, 2.1 kg zinc nitrate and 0.278 kg aluminium nitrate are solved in 15 L water.

15

Solution 2: 2.344 kg sodium bicarbonate is dissolved in 15 L water.

Precipitation:

Both solutions are separately heated to 90°C, followed by the fast addition of solution 1 to solution 2 within 1-2 minutes under stirring. Afterwards 15 min is stirred and the precipitation is filtered and washed with water till it is free of nitrates. The filter cake is dried at 110°C and is calcinated for 4h at 270°C under nitrogen atmosphere. The metal content of the catalyst is in atom%: Cu 38.8; Zn 48.8 and Al 12.9.

25 2. Example

Two solutions are prepared for the precipitation of the components:

Solution 1: A solution of 2.66 kg copper nitrate, 1.05 kg zinc nitrate and 0.278 kg aluminium nitrate are solved in 15 L water.

30

Solution 2: 2.344 kg sodium bicarbonate is dissolved in 15 L water.

Precipitation:

The same procedure as described in the 1. Example, whereby the metal content of the catalyst is in atom%: Cu 61.6; Zn 28.1 and Al 10.9.

35

3. Example - Preparation of Me₃O:

i. Precipitation:

40

A sodium bicarbonate solution (20%) is prepared, whereby 11 kg sodium bicarbonate is dissolved in 44 kg demineralised water. Also a Zn/Al-solution is prepared consisting of 6.88 kg zinc nitrate and 5.67 kg aluminium nitrate and 23.04 kg water. Both solutions are heated to 70°C. A template filled with 12.1 L demineralised water is also heated to 70°C. Both solutions are added in parallel to the template at a pH=7, till the Zn/Al-solution is consumed. Afterwards 15 h is stirred at a pH=7. After this the suspension is vacuumed and washed to a content of sodium oxide <0.10% and the water is free of nitrate. The product is dried for 24h at 120°C and calcinated for 1h at 350°C under air flow.

10 ii. Precipitation:

A sodium bicarbonate solution (20%) is prepared, whereby 25 kg sodium bicarbonate is dissolved in 100 kg demineralised water. Also a Cu/Zn-nitrate solution is prepared consisting of 26.87 kg copper nitrate and 5.43 kg zinc nitrate and 39 kg water. Both solutions are heated to 70°C. After the Cu/Zn-nitrate solution has reached a temperature of 70°C, the product of the 1.precipitation is added slowly and the pH-value is adjusted to pH=2. Also a solution of nitric acid (65%) is provided (650g conc. HNO₃ and 350 g demineralised water). A template filled with 40.8 L demineralised water is also heated to 70°C. Both solutions (sodium bicarbonate and Cu/Zn-nitrate solution) are added in parallel to the template at a pH=6.7, till the Cu/Zn-nitrate solution is consumed. Afterwards 10h is stirred whereby the pH-value is adjusted to pH=6.7 with the nitric acid (65%). After this the suspension is vacuumed and washed to a content of sodium oxide <0.10% and the water is free of nitrate. The product is dried for 72h at 120°C and calcinated for 3h at 300°C under air flow. After cooling to room temperature the material is ready for use.

25

B) Preparation of the final catalytically active body:

The methanol-active compound and the acid compound are compacted separately in a tablet press and/or pelletizing machine. The obtained molding (diameter = ca, 25mm, height = ca, 2mm), is squeezed through sieves with an appropriate mesh size, so that the desired split fraction is obtained. From both fractions the proper quantity is weight in (9/1, 8/2, or 7/3 methanol-active/acidic compound) and mixed with the other compound in a mixing machine (Heidolph Reax 2 or Reax 20/12).

35 C) Testing conditions for non-pelletized mixtures:

The catalytically active body (5 cm³ by volume) is incorporated in a tubular reactor (inner diameter 0.4 cm, bedded in a metal heating body) on a catalyst bed support consisting of alumina powder as layer of inert material and is pressure-less reduced with a mixture of 1 Vol.-% H₂ and 99 Vol.-% N₂. The temperature is increased in intervals of 8 h from 150°C to 170°C and from 170°C to 190°C and finally to 230°C. At a temperature of 230°C the synthesis gas is introduced

40

and heated within 2h up to 250°C. The synthesis gas consists of 45% H₂ and 45% CO and 10% inert gas (argon). The catalytically active body is run at an input temperature of 250°C, GHSV of 2400h⁻¹ and a pressure of 50 bar.

5 D) Testing conditions for pelletized mixtures:

Tests for pelletized materials are conducted in a similar test rick compared to the setup described above for non-pelletized materials using the same routine. Only no tubular reactor with an inner diameter of 0.4 cm is used but a tubular reactor having an inner diameter of 3 cm. Tests for pelletized materials are done with a catalyst volume of 100 cm³.

10

Results:

According to table 1 the different mixtures are listed.

Methanol-active component:

15

Me30: Consists of 70% by weight of CuO, 5.5% by weight Al₂O₃ and 24.5% by weight of ZnO.

Acid Component:

20 The applied acid components have the following composition:

ZSM5-400H

Al 0.23 g/100g

Na 0.09 g/100g

25 Si 45.5 g/100g

ZSM5-100H

Al 0.84 g/100g

Na 0.02 g/100g

30 Si 44 g/100g

ZSM5-80H

Al 0.99 g/100g

Na < 0.01g/100g

35 Si 44 g/100g

ZSM5-50H

Al 1.7 g/100g

Na 0.02 g/100g

40 Si 43 g/100g

ZSM5-25H

Al	2.7 g/100g
Na	0.16 g/100g
Si	41 g/100g

5

In the following table 1 the results are presented. Me30 and ZSM5 (corresponds to ZSM-5) with different ratios of Al, Na and Si are used. The different mixtures of split-fractions (the corresponding D-10, D-50, and D-90 values of Me30 and ZSM5-100H are presented in table 2) show different CO-conversions. The comparison experiments C1 to C7 showing a lower turnover, whereby the inventive experiments E1 and E2 showing an increased value. Surprisingly the mixtures of inventive materials showing a particle size distribution that is characterized by a D-10, D-50, and D-90 value of 5-140 μm , 40-300 μm , and 180-800 μm , respectively, show a significantly increased CO-conversions compared to the comparison experiments C1 to C7. With respect to the selectivity patterns it is worth to mention that within the DME forming samples an equal selectivity of DME and CO₂ can be observed. This shows that all catalysts have a sufficient water gas shift activity that is needed to convert the water generated in the Methanol dehydration step with CO into CO₂. Furthermore all catalysts show an adequate MeOH dehydration capability apart from C4. This can be seen in the MeOH contents in the product streams in table 1.

15

Inventive experiment E3 shows that the superior performance of E1 is maintained if this mixture is properly transferred into a pellet.

Table 1:

25

Experiment	Mixture (A):(B)	Split-fraction	CO conversion [%]	S(MeOH)	S(DME)	S(CO ₂)	S(Other s)
C1	Me30 : ZSM5-25H 8 : 2	0.05-0.1	38.5	2.46	48.45	48.7	0.39
C2	Me30 : ZSM5-50H 8 : 2	0.05-0.1	70.59	1.89	48.39	49.37	0.35
C3	Me30 : ZSM5-100H 8 : 2	0.05-0.1	73.46	1.06	49.09	49.23	0.63
C4	Me30 : ZSM5-400H 8 : 2	0.05-0.1	15.92	96.2	0.44	1.33	2.03
C5	Me30 : ZSM5-100H 8 : 2	0.05-0.1	73.46	1.06	49.09	49.23	0.63
C6	Me30 : ZSM5-100H 8 : 2	0.1-0.15	65.86	2.86	50.95	46.12	0.07

E1	Me30 : ZSM5-100H 8 : 2	0.15-0.2	81.43	2.91	50.22	46.79	0.08
E2	Me30 : ZSM5-100H 8 : 2	0.2-0.5	79.43	1.91	51.88	48.17	0.08
C7	Me30 : ZSM5-100H 8 : 2	0.5-0.7	61.88	3.76	48.69	47.67	0.07
E3	Pellet from E1	3x3 mm	80.78	1.73	49.17	48.89	0.21

All gaseous streams were analyzed via online-GC. Argon was used as internal standard to correlate in and off gas streams.

CO conversion is given as follows: $(CO_{in} - (CO_{out} * Argon_{in} / Argon_{out})) / CO_{in} * 100\%$

5 $S(\text{MeOH}) = \text{Volume (MeOH) in product stream} / \text{Volume (MeOH+DME+CO}_2\text{+Others without hydrogen and CO) in product stream} * 100\%$

$S(\text{DME}) = \text{Volume (DME) in product stream} / \text{Volume (MeOH+DME+CO}_2\text{+Others without hydrogen and CO) in product stream} * 100\%$

$S(\text{CO}_2) = \text{Volume (CO}_2\text{) in product stream} / \text{Volume (MeOH+DME+CO}_2\text{+Others without hydrogen and CO) in product stream} * 100\%$

10 $S(\text{Others}) = \text{Volume (Others) in product stream} / \text{Volume (MeOH+DME+CO}_2\text{+Others without hydrogen and CO) in product stream} * 100\%$

“Others” are compounds that are formed out of H₂ and CO in the reactor that are not MeOH, DME, or CO₂.

15 Table 2:

	D-10 [μm]	D-50 [μm]	D-90 [μm]
Me30 (0.05-0.1)	2.42	46.57	89.14
Me30 (0.1-0.15)	5.06	129.53	143.06
Me30 (0.15-0.2)	6.33	131.69	189.23
Me30 (0.2-0.5)	20.71	275.6	396.86
ZSM5-100H (0.05-0.1)	2.87	56.38	82.17
ZSM5-100H (0.1-0.15)	5.47	100.92	184.78
ZSM5-100H (0.15-0.2)	5.27	163.57	196.22
ZSM5-100H (0.2-0.5)	5.15	373.09	489.57

Claims

1. Catalytically active body for the synthesis of dimethyl ether from synthesis gas, comprising a mixture of:
- 5
- (A) 70-90 % by weight of a methanol-active component, selected from the group consisting of copper oxide, aluminium oxide, zinc oxide, amorphous aluminium oxide, ternary oxide or mixtures thereof,
- 10
- (B) 10-30 % by weight of an acid component, selected from the group consisting of aluminosilicate, γ -alumina and zeolite or mixtures thereof,
- (C) 0-10 Gew.-% by weight of at least one additive, whereby the sum of the components (A), (B) and (C) is in total 100 % by weight.
- 15
2. Catalytically active body according to claim 1, whereby the component (A) has a particle size distribution characterized by a D-10 value of 5-140 μm , a D-50 value of 40-300 μm , and a D-90 value of 180-800 μm , whereby the component (B) has a particle size distribution characterized by a D-10 value of 5-140 μm , a D-50 value of 40-300 μm , and a D-90 value of 180-800 μm and the particle size distribution of components (A) and (B) is maintained in the catalytically active body.
- 20
3. Catalytically active body according to claim 1 or claim 2, characterized in that component (A) comprises 50-80 % by weight of copper oxide, 15-35 % by weight of ternary oxide and 15-35 % by weight of zinc oxide and the sum of which is in total 100 % by weight.
- 25
4. Catalytically active body according to claim 1 or claim 2, characterized in that component (A) comprises 50-80 % by weight of copper oxide, 2-8 % by weight of boehmite and 15-35 % by weight of zinc oxide and the sum of which is in total 100 % by weight.
- 30
5. Catalytically active body according to claim 1 or claim 2, characterized in that component (A) comprises 50-80 % by weight of copper oxide, 2-8 % by weight of amorphous aluminium oxide and 15-35 % by weight of zinc oxide and the sum of which is in total 100 % by weight.
- 35
6. Catalytically active body according to claim 1 or claim 2, characterized in that component (A) comprises 50-80 % by weight of copper oxide, 2-8 % by weight of aluminium oxide and 15-35 % by weight of zinc oxide and the sum of which is in total 100 % by weight.
- 40
7. Catalytically active body according to any of claims 1 to 6, wherein the zeolite comprises 35-55 % by weight of silicon, 0.15-4 % by weight of aluminium, 45-65 % by weight of oxy-

gen and where appropriate 0-0.3 by weight of sodium and the sum of which is in total 100 % by weight.

- 5 8. Catalytically active body according to any of claims 1 to 7, wherein the catalytically active body consists of (A) 70-90 % by weight of a methanol-active component and 10-30 % by weight of a zeolite (B) and the sum of (A) and (B) being in total 100 % by weight.
9. Catalytically active body according to any of claims 1 to 8, wherein the catalytically active body is a pellet with a size in the range from 1 x 1 mm to 10 x 10 mm.
- 10 10. Method for the preparation of a catalytically active body, comprising the step:
- c) preparation of a physical mixture comprising:
- 15 (A) 70-90 % by weight of a methanol-active component, selected from the group consisting of copper oxide, aluminium oxide, zinc oxide, amorphous aluminium oxide, ternary oxide or mixtures thereof,
- (B) 10-30 % by weight of an acid component, selected from the group consisting of aluminosilicate, γ -alumina and zeolite or mixtures thereof,
- 20 (C) 0-10 Gew.-% by weight of a at least one additive, whereby the sum of the components (A), (B) and (C) is in total 100 % by weight.
- 25 11. Method for the preparation of a catalytically active body according to claim 10, whereby the component (A) has a particle size distribution characterized by a D-10 value of 5-140 μm , a D-50 value of 40-300 μm , and a D-90 value of 180-800 μm , whereby the component (B) has a particle size distribution characterized by a D-10 value of 5-140 μm , a D-50 value of 40-300 μm , and a D-90 value of 180-800 μm and the particle size distribution of
- 30 components (A) and (B) is maintained in the catalytically active body.
12. Method for the preparation of a catalytically active body according to claim 10 or 11, comprising further the steps:
- 35 a) precipitation a copper-, zinc-, or aluminium salt or a mixture thereof,
b) calcination of the product obtained in step a).
13. Method for the preparation of a catalytically active body according to any of the claims 10 to 12, wherein a pellet is formed.
- 40

14. Method for the preparation of a catalytically active body according to any of the claims 10 to 13, wherein the components (A) and (B) are independently pressed through at least one sieve, whereby the sieve exhibits a mesh size from 0.005 to 5 mm in order to obtain a particle size distribution characterized by a D-10 value of 5-140 μm , a D-50 value of 40-300 μm , and a D-90 value of 180-800 μm .
15. Method for the preparation of a catalytically active body according to any of the claims 10 to 14, wherein at least three different sieves are used, whereby the components (A) and (B) are pressed in direction from the sieve with the biggest mesh size to the sieve with the smallest mesh size.
16. Method for the preparation of a catalytically active body according to any of the claims 10 to 15, wherein in step a) at least a part of the component (A) is prepared by precipitation reaction and/or calcination.
17. Method for the preparation of a catalytically active body according to any of the claims 10 to 16, whereby at least one part of component (A) is precipitated and whereby at least another part of component (A), which is not subjected to the first precipitation, is added to the precipitate.
18. Method for the preparation of a catalytically active body according to any of the claims 10 to 17, wherein the method further comprises the step d) adding a mixture of hydrogen and nitrogen to component (A) and/or (B).
19. Method for the preparation of dimethyl ether from synthesis gas comprising at least the steps:
- e) reducing the catalytically active body as defined in any of the claims 1 to 9,
 - f) contacting the catalytically active body in a reduced form with hydrogen and at least one of carbon monoxide or carbon dioxide.
20. Use of a catalytically active body according to any of the preceding claims 1 to 9 or obtained by a method according to the claims 10 to 19 for the preparation of dimethyl ether.

INTERNATIONAL SEARCH REPORT

International application No
PCT/EP2013/052959

A. CLASSIFICATION OF SUBJECT MATTER
 INV. B01J21/04 B01J23/80 B01J29/06 C07C41/01 C07C43/04
 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)
 B01J C07C

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, 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 4 423 155 A (BELL WELDON K [US] ET AL) 27 December 1983 (1983-12-27) claims; examples	1-20
X	US 2008/125311 A1 (BAEK YOUNG SOON [KR] ET AL) 29 May 2008 (2008-05-29) cited in the application claims; examples	1-20
X	US 2011/105306 A1 (CHIEN CHUN-CHING [TW] ET AL) 5 May 2011 (2011-05-05) claims; examples	1-20
	----- -/--	

Further documents are listed in the continuation of Box C.

See patent family annex.

* Special categories of cited documents :

<p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier application or patent but published on or after the international filing date</p> <p>"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)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p>	<p>"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</p> <p>"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</p> <p>"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</p> <p>"&" document member of the same patent family</p>
---	---

Date of the actual completion of the international search 24 July 2013	Date of mailing of the international search report 01/08/2013
---	--

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 Schoofs, Bart
--	---

INTERNATIONAL SEARCH REPORT

International application No
PCT/EP2013/052959

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	<p>SOFIANOS A C ET AL: "CONVERSION OF SYNTHESIS GAS TO DIMETHYL ETHER OVER BIFUNCTIONAL CATALYTIC SYSTEMS", INDUSTRIAL & ENGINEERING CHEMISTRY RESEARCH, AMERICAN CHEMICAL SOCIETY, US, vol. 30, no. 11, 1 November 1991 (1991-11-01), pages 2372-2378, XP000264760, ISSN: 0888-5885, DOI: 10.1021/IE00059A002 the whole document</p> <p>-----</p>	1-20

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No PCT/EP2013/052959

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 4423155	A	27-12-1983	CS 234042 B2 14-03-1985
			JP S57156041 A 27-09-1982
			US 4423155 A 27-12-1983
			ZA 8108124 A 27-07-1983

US 2008125311	A1	29-05-2008	CN 101190415 A 04-06-2008
			JP 2008132467 A 12-06-2008
			KR 100812099 B1 12-03-2008
			US 2008125311 A1 29-05-2008

US 2011105306	A1	05-05-2011	TW 201114491 A 01-05-2011
			US 2011105306 A1 05-05-2011
