(54) Title: IMPROVED CATALYST/BIO MASS MIXING IN TRANSPORT REACTOR

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IMPROVED CATALYST/BIOMASS MIXING IN TRANSPORT REACTOR

BACKGROUND OF THE INVENTION

1. Field of the Invention

[0001] The invention relates generally to an improved reactor for pyrolysis of solid particulate biomass material, and more particularly to the mixing zones for mixing the solid biomass materials with a lifting medium and with a heat transfer medium.

2. Description of the Related Art

[0002] Entrained bed reactors, also referred to as transport reactors, are well known in the field of fluid catalytic cracking (FCC). This process is employed in crude oil refineries for cracking relatively heavy feedstocks, such as vacuum gas oil (VGO) to hydrocarbon compounds having boiling points in, for example, the gasoline and diesel ranges.

[0003] In a typical FCC reactor, hot particulate catalyst is mixed in a mixing section with a non-oxidizing lift gas. The hot catalyst particles become entrained by the flow of lift gas. The liquid feedstock is sprayed into this flow of entrained catalyst particles. Upon contact with the heat of the catalyst particles, the feedstock droplets rapidly vaporize. The formation of feedstock vapors causes a further acceleration of the catalyst particles. The hydrocarbons present in the vapors are catalytically cracked to the desired molecular weight range.

[0004] US Patent 5,961,786 to Freel et al. discloses an FCC-type reactor for carrying out a thermal or catalytic pyrolysis of a variety of feedstocks, including liquid and solid biomass materials. The reactor is an upflow, entrained bed reactor, and comprises a mixing section with a first inlet means and a secondary inlet means. Hot transfer medium particles are introduced into the mixing zone via the first inlet means. The lifting gas takes up the heat transfer particles and carries them upward. Feedstock is injected into the flow of hot heat transfer particles via the secondary inlet means.

[0005] The ’786 patent contains examples on use of the reactor in the pyrolysis of wood particles, using sand as the heat transfer medium. The stated aim is to produce wood distillation products that can be used as "liquid smoke" products for flavoring food products, such as meats. The wood distillation products are characterized by a low pH (around 3.0) and
a high oxygen content, making them unsuitable for use as liquid fuels or as a blending stock for liquid fuels.

[0006] There is a need for an entrained bed reactor for the pyrolysis of solid biomass materials to a liquid product having a low oxygen content.

[0007] Thus, there is a particular need for an entrained bed reactor for use in a catalytic pyrolysis process for solid biomass materials.

BRIEF SUMMARY OF THE INVENTION

[0008] The present invention addresses these problems by providing a reactor for fluidized cracking of solid particulate biomass material, said reactor comprising:

(i) a first mixing zone for mixing a first particulate stream comprising a particulate solid biomass material with a lifting medium and for fluidizing and transporting the first particulate stream in the lifting medium forming a fluidized first particulate stream;

(ii) a second mixing zone for injecting a second particulate stream comprising a heat carrier material, and for mixing said second particulate stream with the fluidized first particulate stream, said second mixing zone being downstream from the first mixing zone.

[0009] In a preferred embodiment the solid biomass material comprises cellulose, in particular lignocellulose.

[0010] The preferred heat carrier material is a material having catalytic activity for the cracking of the solid biomass material.

BRIEF DESCRIPTION OF THE DRAWINGS

[0011] The features and advantages of the invention will be appreciated upon reference to the following drawing, in which:

[0012] FIG. 1 is a schematic view of the mixing zones of the reactor of the present invention, showing the first and second particulate streams.
DETAILED DESCRIPTION OF THE INVENTION

[0013] In conventional FCC units, the mixing zone is generally referred to as the "wye", as in many cases it comprises a tubing assembly having a Y-shape. See, for example, US Patent 6,616,899 to Upson, in particular Figure 1 thereof, hot catalyst particles from the regenerator are metered into one branch of the wye. At the intersection the hot particles meet an upward flow of lift gas, and are entrained by it. Feedstock is injected into the flow of hot particles at a point above the intersection of the wye. In general, the injection point for the feedstock is positioned high enough above the intersection of the wye to permit thorough mixing of the catalyst particles and the lift gas before the catalyst particles are contacted with the feedstock.

[0014] US Patent 5,961,786 to Freel et al. discloses an entrained bed reactor for the thermal or catalytic pyrolysis of a variety of materials, including solid or liquid biomass materials. The reactor comprises a mixing zone having a first inlet and a secondary inlet. The mixing zone is designed as a conventional FCC wye. Specifically, the lift gas is first mixed with a hot heat carrier material. The feedstock is subsequently added to this lift gas/heat carrier flow.

[0015] The pyrolysis products reported in the '786 patent are intended to be used as "liquid smoke", a product for flavoring food products, in particular meats. These liquid smoke products are characterized by a low pH (around 3), a high oxygen content (around 26 wt%), and a propensity for developing a brown color. Although these properties may be desirable for liquid smoke products, they are highly undesirable for liquid fuels and blending stocks for liquid fuels.

[0016] In particular the propensity for developing a brown color indicates that the liquid smoke products comprise components that result from over-cracking of the biomass material. We have found that one factor causing over-cracking in the reactor of the '786 patent is the reported design of the mixing zone.

[0017] Unlike a liquid FCC feed, solid particulate biomass material cannot be sprayed into the mixing zone. As a result, as compared to a liquid FCC feed, it takes longer for a solid particulate biomass material to become fully mixed with the lift gas. In an FCC unit, the mixing action is accelerated by the rapid evaporation of the liquid feedstock. Such an effect is not present when mixing solid particulate biomass with a lift gas. The biomass particles need
to be partially pyrolyzed before any vapors are created. Thus, to the extent that evaporation aids in the mixing of the biomass particles and the lift gas, the effect is slower in onset and less pronounced than in the conventional FCC process. As a result, more time is needed to obtain good mixing of the solid particulate feedstock than if the feedstock is a liquid.

[0018] We have found it important to reverse the order of mixing, so that the solid particulate biomass material becomes mixed with the lift gas before it is contacted with a hot heat carrier material. Accordingly, the present invention relates to a reactor for fluidized cracking of solid particulate biomass material, said reactor comprising:

(i) a first mixing zone for mixing a first particulate stream comprising a particulate solid biomass material with a lifting medium and for fluidizing and transporting the first particulate stream in the lifting medium;

(ii) a second mixing zone for injecting a second particulate stream comprising a heat carrier material, and for mixing said second particulate stream with the fluidized first particulate stream, said second mixing zone being downstream from the first mixing zone.

[0019] We have also found that wall effects contribute to undesired over-cracking when solid biomass material is mixed with a lift gas flow already containing hot heat carrier material. For this reason, the advantages of the inventive mixing zones are even greater for small scale reactors, such as pilot plant scale and laboratory scale reactors.

[0020] In earlier patent applications we have reported on work done by the assignee of this patent application on developing a biomass pyrolysis process that produces a liquid pyrolysis process characterized by a pH significantly higher than 3, and an oxygen content well below 20 wt%. Both characteristics are important for materials intended to be used as liquid fuels or as blending stocks for liquid fuels, optionally after an upgrading treatment. It has been found that liquid pyrolysis products of desirable quality can be obtained by (i) using a heat carrier comprising a catalytic material; and/or (ii) giving the solid biomass material a pretreatment in order to make it more reactive.

[0021] It will be understood that both types of measures aggravate the problems encountered when the biomass material is contacted with the hot heat carrier material before it is properly mixed with the lift gas. If the heat carrier material is a catalytic material, the cracking reaction
is accelerated by it, which increases the risk of over-cracking. Likewise, the risk of over-cracking is increased if the solid biomass material has received a pretreatment making it more reactive. It will be appreciated that the design of the mixing zones of the reactor of the present invention is particularly advantageous for catalytic pyrolysis processes, and for pyrolysis processes employing activated biomass materials.

[0022] If a catalytic material is used, the material may be a solid acid, a solid base, or a solid amphoteric material.

[0023] Examples of suitable solid acid materials include zeolites, in particular zeolite Y and ZSM-5, and materials obtained by high temperature calcination of zeolites.

[0024] Examples of suitable solid bases include metal oxides, in particular MgO and CaO; mixed metal oxides; clays; hydrotalcite and hydrotalcite-like materials; layered hydroxy salts; and materials obtained by high temperature calcination of such solid base materials.

[0025] Examples of suitable amphoteric materials include aluminas, in particular a-alumina and y-alumina.

[0026] Particularly preferred are materials having a low catalytic activity, in particular materials having a specific surface area (as determined by the adsorption of liquid nitrogen, using the Brunauer-Emmett-Teller (BET) method) in the range of from 1 to 50 m²/g, preferably from 5 to 20 m²/g. Such materials may be obtained by high temperature calcination of materials having a significantly greater specific surface area. For example, a suitable catalyst may be obtained by calcining kaolin clay or a similar clay through its isotherm, i.e., to a temperature in the range of 1000 to 1100°C.

[0027] Preferably the catalyst particles are in the form of micro spheres, to facilitate their fluidization. The particle size of the catalyst particles is preferably in the range of from 20 μm to 100 μm, in order for the particles to be readily fluidizable.

[0028] It will be understood that the catalyst particles may contain catalytically inert materials, such as refractory oxides or sand. Likewise, catalyst particles may be mixed with inert particles, such as sand. In an alternate embodiment, inert materials may be used as the sole heat carrier material.
[0029] Pretreatment of the biomass material can comprise one or more of the following: impregnation of a solution of a soluble catalyst, such as KOH, NaOH, K₂CO₃, Na₂CO₃; mechanical treatment (milling, grinding, kneading, and the like), optionally in the presence of a particulate inorganic material, preferably a catalytic material; demineralization; solvent explosion; heat treatment at a temperature in the range of from 80 to 300°C; and the like.

[0030] The solid biomass material preferably comprises cellulose. Particularly preferred are materials comprising lignocellulose, because of the low cost and abundant availability of such materials.

[0031] The first mixing zone preferably has a temperature in the range of from about 0°C to about 300°C, preferably in the range of from about 125°C to about 200°C.

[0032] In one embodiment the first particulate stream further comprises a particulate inorganic material. This can be the case, for example, if the pretreatment of the solid biomass material comprises a mechanical treatment in the presence of a particulate inorganic material. In this case, the particulate inorganic material present in the first stream may be the same material as is present in the second stream, or it may be a different material. If the two materials are different it is desirable to separate the two materials after leaving the reactor, so that each can be recycled to its appropriate point in the process.

[0033] The second particulate stream comprises the hot heat carrier material. Specifically, the second particulate stream, immediately prior to entering the second mixing zone, has a temperature in the range of from about 450 °C to about 1000 °C. Preferably the second particulate stream, immediately prior to entering the second mixing zone, has a temperature in the range of from about 500°C to about 800°C.

[0034] The first particulate stream is fed into the first mixing zone by a positive displacement means, preferably selected from the group consisting of a screw feeder, a posimetric feeder, a screw feeder in combination with a vibratory feeder, and a combination of block valves and a vibratory feeder. Preferably, the first particulate stream is fed into the first mixing zone by gravity means preferably though a fluidized bed feeder and control valve.
[0035] The second particulate stream can be fed into the second mixing zone by any means known to the person skilled in the FCC art. Preferably the second particulate stream is fed into the second mixing zone by gravity means preferably though a fluidized bed feeder and control valve.

[0036] The lifting medium ("lift gas") has a velocity in the first mixing zone preferably in the range of from about 5 to about 36 feet per second (about 1.5 to about 12 m/s).

[0037] The second particulate stream can be fed to the second mixing zone at an angle which is counter-current to the flow of the fluidized first particulate stream in the reactor. The angle, as measured from the downstream length of the reactor, can be in the range of from about 30 to about 80 degrees.

DESCRIPTION OF SPECIFIC EMBODIMENTS

[0038] The following is a description of certain embodiments of the invention, given by way of example only and with reference to the drawings. Referring to FIG. 1 a), a typical biomass conversion transport reactor system is shown wherein biomass is added to a hot heat carrier material stream already fluidized by lift gas. In this operation, pre-mature heating of the biomass in the feed line was observed which, in some cases, led to plugging and over cracking of the biomass. Figures 1 b) and c) show the inventive system wherein the hot heat carrier material is added to a biomass stream already fluidized by the lift gas. In this configuration, the pre-mature heating of the biomass is avoided. Also shown in Figures 1 b) and c) is the introduction of the hot heat carrier material at an angle resulting in counter-current flow of the heat carrier material to the flow path of the fluidized biomass. This tends to increase the mixing of the hot heat carrier material and biomass, which aids in heat transfer from the hot heat carrier material to the biomass. Also, when a catalyst is the heat carrier, the counter-current flow results in better contact between biomass, thermolysis products and catalyst.
WHAT IS CLAIMED IS:

1. A reactor for fluidized cracking of solid particulate biomass material, said reactor comprising:
   (i) a first mixing zone for mixing a first particulate stream, comprising a particulate solid biomass material, with a lifting medium and for fluidizing and transporting the first particulate stream in the lifting medium thereby forming a fluidized first particulate stream;
   (ii) a second mixing zone for injecting a second particulate stream comprising a heat carrier material and for mixing said second particulate stream with said fluidized first particulate stream, said second mixing zone being downstream from said first mixing zone.

2. The reactor of claim 1 which is a riser reactor.

3. The reactor of claim 2 wherein said particulate solid biomass material comprises cellulose.

4. The reactor of claim 3 wherein said particulate solid biomass material comprises lignocellulose.

5. The reactor of any one of the preceding claims wherein said heat carrier material comprises a material selected from the group consisting of an inert particulate material, a particulate catalyst material, and combinations thereof.

6. The reactor of anyone of the preceding claims wherein said particulate solid biomass material has received a pretreatment to make it more reactive.

7. The reactor of any of the preceding claims wherein said first mixing zone has a temperature in the range of from about 0°C to about 300°C.

8. The reactor of claim 7 wherein said first mixing zone has a temperature in the range of from about 125°C to about 200°C.

9. The reactor of any one of the preceding claims wherein said first particulate stream further comprises a particulate inorganic material.

10. The reactor of claim 9 wherein said particulate inorganic material portion of said first particulate stream has the same composition as said second particulate stream.

11. The reactor of any one of the preceding claims wherein immediately prior to entering said second mixing zone said second particulate stream has a temperature in the range of from about 450°C to about 1000°C.
12. The reactor of claim 11 wherein immediately prior to entering said second mixing zone said second particulate stream has a temperature in the range of from about 500 °C to about 800 °C.

13. The reactor of any one of the preceding claims wherein said first particulate stream is fed into said first mixing zone by gravity feed, by a positive displacement means, or by fluid conveyance.

14. The reactor of claim 13 wherein said first particulate stream is fed into said first mixing zone by a positive displacement means selected from the group consisting of a screw feeder, a posimetric feeder, a screw feeder in combination with a vibratory feeder, and a combination of block valves and a vibratory feeder.

15. The reactor of claim 14 wherein said first particulate stream is fed into said first mixing zone by gravity means though a fluidized bed feeder and control valve.

16. The reactor of any one of the preceding claims wherein the lifting medium in the first mixing zone has a velocity of from about 5 to about 36 feet per second (about 1.5 to about 12 m/s).

17. The reactor of any one of the preceding claims wherein said second particulate stream is fed to said second mixing zone at an angle which is counter-current to the flow of said fluidized first particulate stream.

18. The reactor of claim 17 wherein said angle, as measured from the downstream length of said reactor, is in the range of from about 30 to about 80 degrees.
Fig 1

a) Biomass

b) Catalyst

c) Biomass