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(71)	Applicant(s) Sasol Technology (Pty) Ltd			
(72)	2) Inventor(s) Van Der Walt, Janette;Koper, Edward Ludovicus;Du Plessis, Gert Hendrik;Phillips, Trevor David;Dancuart Kohler, Luis Pablo Fidel;Du Toit, Francois Jacobus			
(74)	Agent / Attorney Madderns, Level 1 64 Hindmarsh Sq, Adelaide, SA, 5000			
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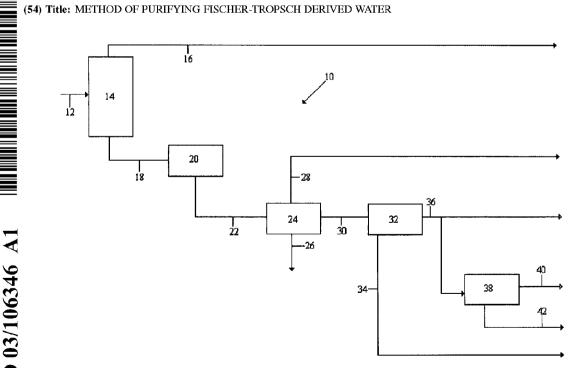
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- (71) Applicant (for all designated States except US): SASOL TECHNOLOGY (PTY) LTD [ZA/ZA]; 1 Sturdee Avenue, Rosebank, 2196 Johannesburg (ZA).
- (72) Inventors; and

(75) Inventors/Applicants (for US only): DANCUART KOHLER, Luis, Pablo, Fidel [ZA/ZA]; 20 Lombard Street, 1947 Vaalpark (ZA). DU PLESSIS, Gert, Hendrik [ZA/ZA]; 15 Vermeer Street, 2302 Secunda (ZA). DU TOIT, Francois, Jacobus [ZA/ZA]; 11 Culemborg

Street, 1947 Sasolburg (ZA). KOPER, Edward, Ludovicus [ZA/ZA]; 11 Douglas Street, 2302 Secunda (ZA). PHILLIPS, Trevor, David [ZA/ZA]; 53 Macowen Street -SE2, 1911 Vanderbijlpark (ZA). VAN DER WALT, Janette [ZA/ZA]; Gleneden 39, Constantia Avenue, 1947 Vaalpark (ZA).

- (74) Agent: HAHN & HAHN INC. LUTEREK, JF; WILLIAMS, VC; HAHN, HH, DUNLOP, AJS; MICHAEL, C; VENTER,; Hahn Forum, 222 Richard Street, 0083 Pretoria (ZA).
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(57) Abstract: A process for the production of highly purified water 40 from Fischer-Tropsch reaction water 12 includes distillation 14 as a primary treatment stage, evaporation 20 as a secondary treatment stage, aerobic treatment 24 as a tertiary treatment stage, solid-liquid separation 32 as a quartic treatment stage and membrane separation 38 as a final treatment stage.

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METHOD OF PURIFYING FISCHER-TROPSCH DERIVED WATER

Field of the Invention

5 This invention relates to the purification of water produced during Fischer-Tropsch synthesis for which synthesis a variety of carbonaceous materials are used as feedstock.

Background to the Invention

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The applicant is aware of processes for the synthesis of water from a carbonaceous feedstock, such as natural gas and coal, which processes also produce hydrocarbons.

15 One such process is the Fischer-Tropsch process of which the largest product is water and, to a lesser extent, hydrocarbons including olefins, paraffins, waxes, and oxygenates. There are numerous references to this process such as, for example on pages 265 to 278 of "Technology of the Fischer-Tropsch process" by Mark Dry, Catal. Rev. Sci.Eng., Volume 23 (1&2), 1981.

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fluids.

The products from the Fischer-Tropsch process may be processed further, for example by hydroprocessing, to produce products including synthetic crude oil, olefins, solvents, lubricating, industrial or medicinal oil, waxy hydrocarbons, nitrogen and oxygen containing compounds, motor gasoline, diesel fuel, jet fuel and kerosene. Lubricating oil includes automotive, jet, turbine and metal working oils. Industrial oil includes well drilling fluids, agricultural oils and heat transfer

In certain areas where carbonaceous feedstocks are to be found, water is in 30 short supply and a relatively costly commodity. Also, environmental concerns • prevent the disposal of polluted water derived from the Fischer-Tropsch process WO 03/106346

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into natural water ways and the sea thereby presenting a case for the production and recovery of useable water at the source of the carbonaceous feedstocks.

The carbonaceous feedstocks typically include coal and natural gas that are converted to hydrocarbons, water and carbon dioxide during Fischer-Tropsch synthesis. Naturally, other carbonaceous feedstocks such as, for example, methane hydrates found in marine deposits, can also be used.

Before the water produced during the Fischer-Tropsch process is purified in accordance with the present invention, it is typically subjected to preliminary separation aimed at isolating a water-enriched stream from the Fischer-Tropsch products.

The preliminary separation process includes condensing the gaseous product from the Fischer-Tropsch reactor and separating it in a typical three-phase separator. The three streams exiting the separator are: a tail gas, a hydrocarbon condensate including mainly hydrocarbons in the C_5 to C_{20} range and a reaction water stream containing non-acid chemicals, water, acids, and suspended hydrocarbons.

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The reaction water stream is then separated using a coalescer that separates the reaction water stream into a hydrocarbon suspension and a water-rich stream.

The coalescer is capable of removing hydrocarbons from the reaction water stream to a concentration of between 10 ppm and 1000 ppm, typically 50 ppm.

The water-enriched stream thus obtained forms the feedstock for the method according to the present invention and will be denoted in this specification by the term "Fischer-Tropsch reaction water".

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The composition of the water-enriched stream or reaction water is largely dependent on the catalyst metal used in the Fischer-Tropsch reactor and the reaction conditions (e.g. temperature, pressure) employed. The Fischer-Tropsch reaction water can contain oxygenated hydrocarbons including aliphatic, aromatic

5 and cyclic alcohols, aldehydes, ketones and acids, and to a lesser extent aliphatic, aromatic and cyclic hydrocarbons such as olefins and paraffins.

The Fischer-Tropsch reaction water may also contain small quantities of inorganic compounds including metals from the Fischer-Tropsch reactor, as well as nitrogen and sulphur containing species that originate from the feedstock.

The influence of the type of Fischer-Tropsch synthesis used on the quality of Fischer-Tropsch reaction water is illustrated in typical organic analysis (Table 1) of Fischer-Tropsch reaction water generated from three different synthesis operating modes, namely:

٠	Low Temperature Fischer-Tropsch	LTFT	Cobalt or Iron
			catalysts
٠	High Temperature Fischer-Tropsch	HTFT	Iron catalyst

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Table 1:
 Typical Organic Composition of Fischer-Tropsch reaction water

 from Different Fischer-Tropsch Synthesis Operating Modes

Component (mass %)	LTFT (Cobalt Catalyst)	LTFT (Iron Catalyst)	HTFT (Iron Catalyst)
Water	98.89	95.70	94.11
non-acid oxygenated hydrocarbons	1.00	3.57	4.47
Acidic oxygenated hydrocarbons	0.09	0.71	1.41
Other Hydrocarbons	0.02	0.02	0.02
Inorganic components	< 0.005	< 0.005	< 0.005

25 It is evident from the typical analyses of Fischer-Tropsch reaction waters of different origin (Table 1) that these waters, in particular HT Fischer-Tropsch

reaction water, contain relatively high concentrations of organic compounds, and direct application or disposal of these waters is generally not feasible without further treatment to remove undesirable constituents. The degree of treatment of the Fischer-Tropsch reaction water depends largely on the intended application,

5 and it is possible to produce a wide range of water qualities ranging from boiler feed water to partially treated water which may be suitable for discharge to the environment.

It is also possible to co-treat Fischer-Tropsch reaction water with other typical process waste water as well as rain water.

The water purification processes described in this invention may, after making minor adaptations, also be used for the processing of aqueous streams derived from generic synthesis gas conversion processes using metallic catalysts similar to the catalysts used during Fischer-Tropsch synthesis.

Summary of the Invention

According to a first aspect of the invention there is provided a process for the production of purified water from Fischer-Tropsch reaction water, which process includes at least the steps of:-

- a primary treatment stage comprising an equilibrium staged separation process having at least one stage for removing at least a fraction of nonacid oxygenated hydrocarbons from the Fischer-Tropsch reaction water to produce a primary water-enriched stream;
- 25

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 a secondary treatment stage comprising evaporation of at least a portion of the primary water-enriched stream thereby to reduce the total mass fraction of organic constituents in the unevaporated portion to produce a secondary water-enriched stream; and

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- c) a final treatment stage comprising solid-liquid separation for removing at least some solids from at least a portion of the secondary water-enriched stream.
- 5 According to a second aspect of the invention there is provided a process for the production of purified water from Fischer-Tropsch reaction water, which process includes at least the steps of:-

produce a primary water-enriched stream;

- (a) a primary treatment stage comprising an equilibrium stage separation process having at least one stage for removing at least a fraction of nonacid oxygenated hydrocarbons from the Fischer-Tropsch reaction water to
- (b) a secondary treatment stage comprising evaporation of at least a portion of the primary water-enriched stream thereby to reduce the total mass fraction of organic constituents in the unevaporated portion to produce a secondary water-enriched stream:
- (c) a tertiary treatment stage comprising biological treatment for removing at least a fraction of dissolved organic carbon from the secondary waterenriched stream to produce a tertiary water-enriched stream; and
- a quartic treatment stage comprising solid-liquid separation for removing at least some solids from at least a portion of the tertiary water-enriched stream.

The term "purified water" is to be interpreted as meaning an aqueous stream having a COD of between 20 and 500 mg/l, a pH of between 6,0 and 9,0, a suspended solids content of less than 250 mg/l and a total dissolved solids content of less than 600 mg/l.

The non-acid oxygenated hydrocarbons are typically selected from the group including: alcohols, ketones and aldehydes, and are more specifically selected 30 from the group including: acetaldehyde, propionaldehyde, butyraldehyde,

acetone, methyl-propyl-ketone, methanol, ethanol, propanol, butanol, pentanol, hexanol, and heptanol.

A number of equilibrium staged separation processes are suitable for use in the primary treatment stage. Such processes may include conventional distillation 5 processes typically used in the refining and petrochemical industry as well as solvent extraction using either conventional liquid solvents or liquefied gases.

When distillation is used as the primary treatment stage, the bulk of the non-acid oxygenated hydrocarbons contained in the Fischer-Tropsch reaction water are 10 removed, leaving predominantly mono-carboxylic acids (e.g. acetic acid, propionic acid) and optionally trace quantities of non-acid compounds. As a result of the presence of organic acids, the primary water-enriched stream is known as Fischer- Tropsch acid water.

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The overheads from distillation may be recovered and worked-up to products, or may be used for fuel or as an energy source.

The primary treatment stage may include degassing of the reaction water before further processing to remove compounds having a very low boiling point and 20 dissolved gases from the Fischer-Tropsch reaction water.

Typically, Fischer-Tropsch reaction water originating from HTFT iron catalyst processes which water has undergone primary treatment has limited application due to the relatively high concentrations (> 1% by mass) of organic acids 25 remaining in the FT acid water and further treatment of the water is required. In contrast, Fischer-Tropsch reaction water originating from cobalt-based LTFT processes which water has undergone primary treatment contains significantly lower organic acid concentrations (< 0.1 % mass %) and can therefore, following 30

neutralization, be released to the environment if sufficient dilution is available and

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discharge standards permit. The primary water-enriched stream may also have limited application as process water.

The evaporation in step b) may take place at ambient temperature and pressure 5 in a cooling tower.

During evaporation in the cooling tower, the amount of at least some of the dissolved organic constituents contained in the primary water-enriched stream are reduced through the action of micro-organisms. The constituents being removed totally or partially include acidic oxygenated hydrocarbons and methanol.

By using the primary water-enriched stream as cooling water, the abundance of oxygen caused by aeration in the cooling tower activates the growth of microorganisms that use dissolved organic constituents in the primary water-enriched

stream as food source.

The evaporation may comprise use of the primary water-enriched stream as make up water to an evaporative cooling tower. The evaporative cooling tower 20 may be selected from the group including: mechanical-draft cooling towers, natural-draft cooling towers and forced-draft cooling towers.

When using the primary water-enriched stream as cooling water, the linear flow velocity of said water through the equipment being used in the cooling must be

25 sufficiently high enough to inhibit deposition of suspended solids in said equipment.

The pH of the primary water-enriched stream should be controlled by addition of an alkali, such as caustic soda, to prevent acid corrosion of metal and/or 30 concrete surfaces that might be exposed to this water. One or more suitable additives may be added to the primary water-enriched stream before using it as cooling water to inhibit undesirable effects such as, for example, fouling, corrosion and scaling.

5 Biological treatment of the secondary water-enriched stream may include aerobic treatment.

The aerobic treatment method may be the same as is conventionally used for domestic and industrial wastewater treatment.

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The aerobic treatment may include adding nutrients in the form of nitrogen (e.g. urea, ammonia or ammonium salts) and phosphorus (e.g. phosphate salts) containing compounds to accelerate microbiological degradation of the organic carbon-containing constituents. In addition, pH control using alkali compounds such as lime, caustic and soda ash may be required to optimize the performance of the micro-organisms.

A wide range of technologies may be used in the aerobic treatment of the secondary water-enriched stream. Such technologies may be selected from a
 group including: Activated Sludge processes, High-rate Compact Reactors, Biological Aerated Filters, Trickling filters, Rotating Biological Contactors, Membrane Bioreactors and Fluidised Bed reactors. The aerobic production of Single Cell Protein (SCP) has also been successfully developed.

- 25 Apart from a water-rich or tertiary water-enriched stream, aerobic treatment typically yields carbon dioxide and sludge as byproducts. The carbon dioxide may be released to the environment. The sludge may be incinerated, used as land fill, fertilizer, soil conditioner or as a source of SCP.
- 30 The quartic treatment stage may be aimed at removing suspended solids from the tertiary water-enriched stream produced during biological treatment.

Suspended solid removal may be achieved by methods selected from the group including: sand filtration, membrane separation (e.g. micro- or ultrafiltration), flocculation sedimentation, dissolved air flotation (with or without the use of flocculants) and centrifugation.

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Local discharge standards or the intended application will dictate the required level and type of quartic treatment.

Applications for the purified water produced by the method described above may include its use as cooling water, irrigation water or general process water.

The purified water typically has the following characteristics:

Property		
Chemical Oxygen Demand (COD)	mg/l	20- 500
рН		6,0-9,0
Total Dissolved Solids (TDS)	mg/l	<600
Suspended solids (SS)	mg/l	<250

¹⁵ According to a third aspect of the invention there is provided a process for the production of highly purified water from Fischer-Tropsch reaction water, which process includes at least the steps of:-

- a primary treatment stage comprising an equilibrium staged separation process having at least one stage for removing at least a fraction of nonacid oxygenated hydrocarbons from the Fischer-Tropsch reaction water to produce a primary water-enriched stream;
- a secondary treatment stage comprising evaporation of at least a portion of the primary water-enriched stream thereby to reduce the total mass fraction of organic constituents in the unevaporated portion to produce a secondary water-enriched stream;

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- c) a tertiary treatment stage comprising solid-liquid separation for removing at least some solids from at least a portion of the secondary waterenriched stream, to produce a tertiary water-enriched stream; and
- (d) a quartic treatment stage comprising a dissolved salt and organic removal stage for removing at least some dissolved salts and organic constituents from at least a portion of the tertiary water enriched stream.

According to a fourth aspect of the invention there is provided a process for the production of highly purified water from Fischer-Tropsch reaction water, which process includes at least the steps of:-

- a primary treatment stage comprising an equilibrium stage separation process having at least one stage for removing at least a fraction of nonacid oxygenated hydrocarbons from the Fischer-Tropsch reaction water to produce a primary water-enriched stream;
- b) a secondary treatment stage comprising evaporation of at least a portion of the primary water-enriched stream thereby to reduce the total mass fraction of organic constituents in the unevaporated portion to produce a secondary water-enriched stream;
- c) a tertiary treatment stage comprising biological treatment for removing at
 20 least a fraction of dissolved organic carbon from the secondary waterenriched stream to produce a tertiary water-enriched stream;
 - a quartic treatment stage comprising solid-liquid separation for removing at least some solids from at least a portion of the tertiary waterenriched stream to produce a quartic water-enriched stream; and
- e) a final treatment stage comprising a dissolved salt and organic removal stage for removing at least some dissolved salts and organic constituents from at least a portion of the quartic water enriched stream.

The term "highly purified water" is to be interpreted as meaning an aqueous 30 stream having a COD of less than 50 mg/l, a pH of between 6,0 and 9,0, a suspended solids content of less than 50 mg/l and a total dissolved solids content of less than 100 mg/l.

The non-acid oxygenated hydrocarbons are typically selected from the group 5 including: aldehydes, ketones and alcohols, and are more specifically selected from the group including: acetaldehyde, propionaldehyde, butyraldehyde, acetone, methyl-propyl-ketone, methanol, ethanol, propanol, butanol, pentanol, hexanol, and heptanol.

- 10 A number of equilibrium staged separation processes are suitable for use in the primary treatment stage. Such processes may include conventional distillation processes typically used in the refining and petrochemical industry as well as solvent extraction using either conventional liquid solvents or liquefied gases.
- 15 When distillation is used as the primary treatment stage, the bulk of the non-acid oxygenated hydrocarbons contained in the Fischer-Tropsch reaction water are removed, leaving predominantly mono-carboxylic acids (e.g. acetic acid, propionic acid) and optionally trace quantities of non-acid compounds. As a result of the presence of organic acids, the primary water-enriched stream is known as
- 20 Fischer- Tropsch acid water.

The overheads from distillation may be recovered and worked-up to products, or may be used for fuel or as an energy source.

- 25 The primary treatment stage may include degassing of the reaction water before further processing to remove compounds having a very low boiling point and dissolved gases from the Fischer-Tropsch reaction water.
- Typically, Fischer-Tropsch reaction water originating from HTFT iron catalyst processes which water has undergone primary treatment has limited application due to the relatively high concentrations (> 1% by mass) of organic acids

remaining in the FT acid water and further treatment of the water is required. In contrast, Fischer-Tropsch reaction water originating from cobalt-based LTFT processes which water has undergone primary treatment contains significantly lower organic acid concentrations (< 0.1 % mass %) and can therefore, following

5 neutralization, be released to the environment if sufficient dilution is available and discharge standards permit. The primary water-enriched stream may also have limited application as process water.

The evaporation in step b) may take place at ambient temperature and pressure in a cooling tower.

During evaporation in the cooling tower, the amount of at least some of the dissolved organic constituents contained in the primary water-enriched stream are reduced through the action of micro-organisms. The constituents being 15 removed totally or partially include acidic oxygenated hydrocarbons and methanol.

By using the primary water-enriched stream as cooling water, the abundance of oxygen caused by aeration in the cooling tower activates the growth of microorganisms that use dissolved organic constituents in the primary water-enriched stream as food source.

The evaporation may comprise use of the primary water-enriched stream as make up water to an evaporative cooling tower. The evaporative cooling tower may be selected from the group including: mechanical-draft cooling towers, natural-draft cooling towers and forced-draft cooling towers.

When using the primary water-enriched stream as cooling water, the linear flow velocity of said water through the equipment being used in the cooling process

30 must be sufficiently high enough to inhibit deposition of suspended solids in said equipment.

The pH of the primary water-enriched stream should be controlled by addition of an alkali, such as caustic soda, to prevent acid corrosion of metal and/or concrete surfaces that might be exposed to this water.

5 One or more suitable additives may be added to the primary water-enriched stream before using it as cooling water to inhibit undesirable effects such as, for example, fouling, corrosion and scaling.

Biological treatment of the secondary water-enriched stream may include aerobic 10 treatment.

The aerobic treatment method may be the same as is conventionally used for domestic and industrial wastewater treatment.

- 15 The aerobic treatment may include adding nutrients in the form of nitrogen (e.g. urea, ammonia or ammonium salts) and phosphorus (e.g. phosphate salts) containing compounds to accelerate microbiological degradation of the organic carbon-containing constituents. In addition, pH control using alkali compounds such as lime, caustic and soda ash may be required to optimize performance of
- 20 the micro-organisms.

A wide range of technologies may be used in the aerobic treatment of the secondary water-enriched stream. Such technologies may be selected from a group including: Activated Sludge processes, High-rate Compact Reactors,

25 Biological Aerated Filters, Trickling filters, Rotating Biological Contacters, Membrane Bioreactors, and Fluidised Bed reactors. The aerobic production of Single Cell Protein (SCP) has also been successfully developed.

Apart from a water-rich or tertiary water-enriched stream, aerobic treatment 30 typically yields carbon dioxide and sludge as byproducts. The carbon dioxide

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may be released to the environment. The sludge may be incinerated, used as land fill, fertilizer, soil conditioner or as a source of SCP.

The quartic treatment stage may be aimed at removing suspended solids from the water-rich stream produced during biological treatment.

Suspended solid removal may be achieved by methods selected from the group including: sand filtration, membrane separation (e.g. micro- or ultrafiltration), flocculation sedimentation, dissolved air flotation (with or without the use of flocculants) and centrifugation.

Residual organic species not removed during biological treatment and solids removal may be removed by methods selected from the group including: chemical oxidation using agents such as ozone and hydrogen peroxide, ultraviolet light generated free radicals and adsorption/absorption processes including activated carbon treatment and organic scavenging resins.

Dissolved salts, originating from secondary treatment (i.e. pH control chemicals, nutrient addition) and/or from the co-treatment of other process effluents, can be

20 further reduced by methods from the group including: ion exchange, reverse osmosis, nano-filtration and chemical precipitation processes including hot and cold lime softening.

Applications for the highly purified water produced by the method described above may include its use as boiler feed water and drinking water.

The highly purified water typically has the following characteristics:

Property		
Chemical Oxygen Demand (COD)	mg/l	<50
рН		6,0-9,0

Total Dissolved Solids (TDS)	mg/l	< 100
Suspended Solids (SS)	mg/l	< 50

Advantages inherent in the purified and highly purified water produced according to the present invention are that the water will contain only a small amount of dissolved solids because the Fischer-Tropsch reaction water is essentially a solids-free stream. The low levels of residual salts in the purified water are a result of controlled addition of chemicals used during the purification process sequence and/or the co-treatment of other dissolved solids containing effluents. The residual salts include Ca, Mg, Na, K, Cl, SO₄, HCO₃ and CO₃ combinations. The low dissolved solids concentrations in Fischer-Tropsch reaction water can

10 simplify and reduce the cost of the purification process.

Detailed Description of the Invention

The invention will now be described by way of the following non-limiting examples with reference to the accompanying drawing.

Figure 1 shows a simplified block diagram of a method 10 according to the present invention including various treatment options.

20 Fischer-Tropsch reaction water 12 is fed to a distillation column 14 for primary treatment.

Two streams 16 and 18 exit distillation column 14. Stream 16 contains predominantly organic constituents whilst stream 18 is a primary water-enriched

25 stream.

Stream 18 is then fed to a forced draft cooling tower 20 in which the stream 18 is used as make up water for a forced-draft cooling tower. By utilizing stream 18 as cooling water, its volume is reduced by evaporation and the amount of at least WO 03/106346

some of the dissolved organic constituents in stream 18 are reduced due to the abundance of oxygen caused by aeration in the cooling tower 20 activating the growth of micro-organisms that use dissolved organic constituents in the primary water-enriched stream 18 as food source.

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Stream 22 is the blow down or unevaporated water portion (or secondary waterenriched stream) from the cooling tower 20 and is subjected to biological treatment in the form of aerobic digestion 24 which produces sludge 26, carbon dioxide 28 and a tertiary water-enriched stream 30.

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The next treatment stage involves subjecting the tertiary water-enriched stream 30 to solid-liquid separation 32 during which solids 34 in the form of sludge as well as a stream of purified water 36 are produced.

15 Some of the purified water 36 from the solid-liquid separation 32 stage is subjected to a final treatment stage in the form of membrane separation 38 to yield highly purified water 40 and concentrate 42. Stream 42 contains nonbiodegradable organic species, concentrated inorganic salts eg sulphates and chlorides of sodium and calcium as well as elevated concentrations of suspended 20 solids.

Depending on the final intended use of the purified 36 or highly purified water 40, the minimum water quality requirements are as set out in Table 2 below and the operating conditions of the equipment used in the method as well as suitable

25 treatment options can be selected accordingly.

	Process	Irrigation	Cooling	Boiler feed	Drinking
	Water	water	water	water	water
COD mg/l	0 - 75		0 - 30	0 - 10	
pН	5 - 10	6.5 - 8.4	6.5 - 8	7-8	6-9
TDS mg/l	0 - 1600	<40	0 - 450	0 - 100	0 - 450
SS mg/l	0 - 25	0 - 50	0 - 5	0 - 3	<20

 Table 2
 Water Quality – Typical Requirements

Having described the basic aspects of the invention, the following example is given to further illustrate a specific embodiment of the invention.

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Example: Treatment of Fischer-Tropsch reaction water derived from a cobalt catalyst LTFT process

After separation of by-products, a water-enriched stream from a LTFT process was degassed at atmospheric pressure. Free hydrocarbons in the water-enriched stream were reduced to 0.01 % (mass %) using a coalescer.

Primary treatment of the thus obtained Fischer Tropcsh reaction water was undertaken using distillation. Analysis of the FT acid water bottoms of the distillation column is detailed in Table 3 below. It is apparent that with the exception of trace quantities of methanol, most other non-acid oxygenated hydrocarbons were removed during primary distillation leaving an organic acid enriched or primary water-enriched stream (i.e. 0.072 mass % organic acids) with a pH value of 3.5. The measured Chemical Oxygen Demand (COD) of this

20 effluent was in the order of 800 mg $0_2/l$.

Table 3:Typical Composition of LTFT Reaction Water Feed and Acid WaterBottoms During Primary Treatment (Distillation)

Component	Reaction water feed to	Acid water – bottoms of	
Component	(mass %)	primary distillation column	
Water		(mass %)	
	97.629	99.840	
Acetaldehyde	0.019	0.000	
Propionaldehyde	0.002	0.000	
Butyraldehyde	0.001	0.000	
Acetone	0.007	0.000	
MPK	0.001	0.000	
Methanol	0.434	0.001	
Ethanol	0.369	0.000	
Propanol	0.140	0.000	
iso-Propanol	0.002	0.000	
Butanol	0.056	0.000	
Pentanol	0.047	0.000	
iso-Pentanol	0.001	0.000	
Hexanol	0.019	0.000	
lso-Hexanol	0.001	0.000	
Heptanol	0.007	0.000	
Other NAC	0.004	0.000	
Total NAC	1.106	0.001	
Formic acid	0.025	0.025	
Acetic acid	0.039	0.040	
Propionic acid	0.002	0.002	
Butyric acid	0.002	0.002	
Other acids	0.006	0.006	
Total Acids	0.070	0.072	
Other Hydrocarbons	0.011	0.011	

5 The primary water-enriched stream was fed into an open holding tank at approximately 70°C.

In order to sustain a microbiological population in the primary water-enriched stream, nitrogen in the form of urea and phosphate in the form of phosphoric acid

were added to said stream. The pH of the primary water-enriched stream was thereafter adjusted to 5 using sodium hydroxide.

The primary water-enriched stream was then fed to a forced draft cooling tower 5 as make up water. The cooling tower was operated at four cycles of concentration and a 10°C delta T.

Removal of organics (measured as COD) from the primary water-enriched stream in the tower was in the order of 45% with volatile acid removal at about

10 55%. The COD of the recirculating water in the tower was approximately 1800 mg/l and the concentration of suspended solids was approximately 200 mg/l whilst the pH of the water varied between 6,5 and 7,5.

Corrosion, fouling and scaling in the cooling tower and surrounding heat exchangers was kept within acceptable limits by applying an appropriate chemical treatment program that included a biodispersant and a scale inhibitor.

The blow down or unevaporated water portion from the cooling tower (secondary water-enriched stream) was then treated in a completely mixed activated sludge system (aerobic treatment) under the following conditions:

- pH:7,2 to 7,5
- Dissolved oxygen concentration: >2 mg/l
- Temperature: 35 °C
- 25 HRT: 30 h

- F/M ratio : 0,2 -0,4 kg COD/kg MLSS.d
- Cell retention time (sludge age) 13 days
- Feed to recycle ratio: 1 : 2
- 30 A COD removal efficiency of 91% was achieved and the thus obtained tertiary water-enriched stream had a COD of 160 mg/l. The suspended solid

concentration of the tertiary water-enriched stream was on average about 120 mg/l.

The tertiary water-enriched stream was then subjected to sand filtration to reduce its SS concentration to 25 mg/l. The thus obtained purified water has application as a both irrigation water and process cooling water. Sludge produced in the process was incinerated.

As an alternative to treatment in an activated sludge system, the blow down or 10 part of the blow down from the cooling tower was diverted to a cross-flow membrane unit fitted with a 0.2 μ m polypropylene microfiltration membrane. A permeate flux rate of 70 – 80 l/m².h was obtained during stable operation of the unit. The water recovery across the unit varied between 75 – 85 %. The resultant COD and SS concentrations in the permeate from the microfiltration unit were 15 1750 mg O₂/l and < 5 mg/l respectively.

The pH of the purified water from the microfiltration unit was adjusted to 8.5 using sodium hydroxide and the purified water was pumped to a reverse osmosis unit fitted with a high rejection polyamide sea water membrane. A permeate flux rate

- of 20 25 l/m^2 .h was obtained during stable operation of the unit. The water recovery across the reverse osmosis unit varied between 80 – 90 %. The unit yielded a highly purified water stream with COD and TDS concentrations of 45 – 50 mg O₂/l and 20 – 30 mg/l respectively.
- 25 It is to be appreciated that the invention is not limited to any specific embodiment or configuration as herein before generally described or illustrated, for example, rain water or water enriched streams from processes other than Fischer-Tropsch synthesis may be purified according to the method described above.

Claims:

- 1. A process for the production of purified water from Fischer-Tropsch reaction water wherein the purified water is an aqueous stream having a COD of between 20 and 500 mg/l, a pH of between 6,0 and 9,0, a suspended solids content of less than 250 mg/l and a total dissolved solids content of less than 600 mg/l, and wherein the process includes at least the steps of:
 - a) a primary treatment stage comprising an equilibrium staged separation process having at least one stage for removing at least a fraction of non-acid
- oxygenated hydrocarbons from the Fischer-Tropsch reaction water to produce a primary water-enriched stream;
 - b) a secondary treatment stage comprising evaporation of at least a portion of the primary water-enriched stream thereby reducing the total mass fraction of organic constituents in the unevaporated portion to produce a secondary
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- water-enriched stream; and
- c) a final treatment stage comprising solid-liquid separation for removing at least some solids from at least a portion of the secondary water-enriched stream.
- 20 2. A process for the production of purified water from Fischer-Tropsch reaction water wherein the purified water is an aqueous stream having a COD of between 20 and 500 mg/l, a pH of between 6,0 and 9,0, a suspended solids content of less than 250 mg/l and a total dissolved solids content of less than 600 mg/l, and wherein the process includes at least the steps of:-
- 25 (a) a primary treatment stage comprising an equilibrium staged separation process having at least one stage for removing at least a fraction of nonacid oxygenated hydrocarbons from the Fischer-Tropsch reaction water to produce a primary water-enriched stream;
 - (b) a secondary treatment stage comprising evaporation of at least a portion of the primary water-enriched stream thereby reducing the total mass fraction

of organic constituents in the unevaporated portion to produce a secondary water-enriched stream;

- (c) a tertiary treatment stage comprising biological treatment for removing at least a fraction of dissolved organic carbon from the secondary waterenriched stream to produce a tertiary water-enriched stream; and
- d) a quartic treatment stage comprising solid-liquid separation for removing at least some solids from at least a portion of the tertiary water-enriched stream.
- 10 3. A process as claimed in claim 1 or claim 2 wherein the non-acid oxygenated hydrocarbons are selected from the group including: acetaldehyde, propionaldehyde, butyraldehyde, acetone, methyl-propyl-ketone, methanol, ethanol, propanol, butanol, pentanol, hexanol, and heptanol.
- A process as claimed in any one of claims 1 to 3 wherein the dissolved organic carbon is selected from the group including: aldehydes, ketones, alcohols, and organic acids.
- A process as claimed in any one of the preceding claims wherein the equilibrium staged separation process for use in the primary treatment stage is selected from the group including: distillation, solvent extraction using liquid solvents, and solvent extraction using liquefied gases.
- 6. A process as claimed in any one of the preceding claims wherein the primary treatment stage includes degassing of the Fischer-Tropsch reaction water before further processing in the primary treatment stage to remove compounds having a very low boiling point and dissolved gases from the Fischer-Tropsch reaction water.

- 7. A process as claimed in any one of the preceding claims wherein the evaporation in step b) takes place at ambient temperature and pressure in a cooling tower.
- 5 8. A process as claimed in any one of the preceding claims wherein the evaporation in step b) takes place by using the primary water-enriched stream as make up water to an evaporative cooling tower.
- A process as claimed in claim 8 wherein the evaporative cooling tower is
 selected from the group including: mechanical-draft cooling towers, naturaldraft cooling towers, and forced-draft cooling towers.
 - 10. A process as claimed in any one of claims 2 to 9 wherein the biological treatment of step c) is an aerobic treatment method.
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- 11. A process as claimed in claim 10 wherein the aerobic treatment method is selected from group including: Activated Sludge processes, Biological Aerated Filters, Trickling filters, Rotating Biological Contacters, High-rate Compact Reactors, Membrane Bioreactors and Fluidised Bed reactors.

- 12. A process as claimed in any one of claims 2 to 11 wherein the quartic treatment stage removes suspended solids from the tertiary water-enriched stream produced during biological treatment.
- 13. A process as claimed in claim 12 wherein the suspended solid removal is achieved by one or more method selected from the group including: sand filtration, membrane separation, flocculation sedimentation, dissolved air flotation with the use of flocculants, dissolved air flotation without the use of flocculants, and centrifugation.

- 14. A process as claimed in claim 13 wherein the membrane separation method includes one or both of microfiltration and ultrafiltration.
- 15. A process for the production of highly purified water from Fischer-Tropsch reaction water wherein the highly purified water is an aqueous stream having a COD of less than 50 mg/l, a pH of between 6,0 and 9,0, a suspended solids content of less than 50 mg/l and a total dissolved solids content of less than 100 mg/l, and wherein the process includes at least the steps of:-
- a primary treatment stage comprising an equilibrium staged separation process having at least one stage for removing at least a fraction of nonacid oxygenated hydrocarbons from the Fischer-Tropsch reaction water to produce a primary water-enriched stream;
 - b) a secondary treatment stage comprising evaporation of at least a portion
- 15 of the primary water-enriched stream thereby to reduce the total mass fraction of organic constituents in the unevaporated portion to produce a secondary water-enriched stream;
 - a tertiary treatment stage comprising solid-liquid separation for removing at least some solids from at least a portion of the secondary waterenriched stream, to produce a tertiary water-enriched stream; and
 - (d) a final treatment stage comprising a dissolved salt and organic removal stage for removing at least some dissolved salts and organic constituents from at least a portion of the tertiary water-enriched stream.
- A process for the production of highly purified water from Fischer-Tropsch reaction water wherein the highly purified water is an aqueous stream having a COD of less than 50 mg/l, a pH of between 6,0 and 9,0, a suspended solids content of less than 50 mg/l and a total dissolved solids content of less than 100 mg/l, and wherein the process includes at least the steps of:-

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- a primary treatment stage comprising an equilibrium staged separation process having at least one stage for removing at least a fraction of nonacid oxygenated hydrocarbons from the Fischer-Tropsch reaction water to produce a primary water-enriched stream;
- b) a secondary treatment stage comprising evaporation of at least a portion of the primary water-enriched stream thereby to reduce the total mass fraction of organic constituents in the unevaporated portion to produce a secondary water-enriched stream;
 - c) a tertiary treatment stage comprising biological treatment for removing at least a fraction of dissolved organic carbon from the secondary waterenriched stream to produce a tertiary water-enriched stream;
 - d) a quartic treatment stage comprising solid-liquid separation for removing at least some solids from at least a portion of the tertiary waterenriched stream to produce a quartic water-enriched stream; and
- e) a final treatment stage comprising a dissolved salt and organic removal stage for removing at least some dissolved salts and organic constituents from at least a portion of the quartic water-enriched stream.
- A process as claimed in claim 15 or 16 wherein the non-acid oxygenated
 hydrocarbons are selected from the group including: aldehydes, ketones, alcohols, and organic acids.
 - 18. A process as claimed in any one of the claims 15 to 17 wherein the equilibrium stage separation process for use in the primary treatment stage is selected from the group including: distillation, solvent extraction using liquid solvents, and solvent extraction using liquefied gases.
 - 19. A process as claimed in any one of the claims 15 to 18 wherein the primary treatment stage includes degassing of the reaction water before further processing in the primary treatment stage to remove compounds having a

very low boiling point and dissolved gases from the Fischer-Tropsch reaction water.

- 20. A process as claimed in any one of the claims 15 to 19 wherein the
 5 evaporation in step b) takes place at ambient temperature and pressure in a cooling tower.
 - 21. A process as claimed in claim 20 wherein the evaporative cooling tower is selected from the group including: mechanical-draft cooling towers, natural-draft cooling towers, and forced-draft cooling towers.
 - 22. A process as claimed in any one of the preceding claims wherein the biological treatment of step c) is an aerobic treatment method.
- 15 23. A process as claimed in claim 22 wherein the aerobic treatment method is selected from group including: Activated Sludge processes, Biological Aerated Filters, Trickling filters, Rotating Biological Contacters, High-rate Compact Reactors, Membrane Bioreactors, and Fluidised Bed reactors.
- 20 24. A process as claimed in any one of the claims 16 to 23 wherein the quartic treatment stage removes suspended solids from the tertiary water-enriched stream produced during biological treatment.
- 25. A process as claimed in claim 24 wherein the suspended solid removal is achieved by one or more method selected from the group including: sand filtration, membrane separation, flocculation sedimentation, dissolved air flotation with the use of flocculants, dissolved air flotation without the use of flocculants, and centrifugation.
- 30 26. A process as claimed in claim 25 wherein the membrane separation method includes one or both of microfiltration and ultrafiltration.

- 27. A process as claimed in any one of claims 16 to 26 wherein residual organic species are removed in the final treatment stage by one or more methods selected from the group including: chemical oxidation, ultraviolet light generated free radicals, adsorption and/or absorption processes.
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- 28. A process as claimed in claim 27 wherein the adsorption and/ or absorption processes include one or both of activated carbon treatment and the use of organic scavenging resins.
- 10 29. A process as claimed in any one of claims 16 to 28 wherein dissolved salts originating from tertiary treatment and/or from the co-treatment of other process effluents, are reduced in the final treatment stage by one or more method selected from the group including: ion exchange, reverse osmosis, nano-filtration, and chemical precipitation processes.
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- 30. A process as claimed in claim 29 wherein the chemical precipitation processes are selected from one or both of hot lime softening and cold lime softening.
- 20 31. A process according to the invention for the production of purified water from Fischer-Tropsch reaction water substantially as hereinbefore described or exemplified.
- 32. A process of producing purified water from Fischer-Tropsch reaction water
 including any new and inventive integer or combination of integers, substantially as herein described.
 - 33. A process according to the invention for the production of highly purified water from Fischer-Tropsch reaction water substantially as hereinbefore described or exemplified.

34. A process of producing highly purified water from Fischer-Tropsch reaction water including any new and inventive integer or combination of integers, substantially as herein described.

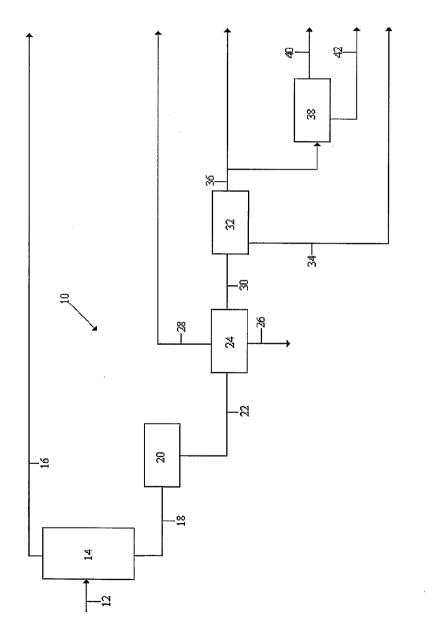


Figure 1