

CO-CURRENT GAS ABSORPTION METHOD AND SYSTEM WITH AN ENHANCED ABSORBENT

CROSS REFERENCE TO RELATED APPLICATION

[0001] This application claims the priority benefit of United States Provisional
5 Patent Application No. 62/769144 filed November 19, 2018, entitled ENHANCED
ACID GAS REMOVAL WITHIN A GAS PROCESSING SYSTEM.

FIELD

[0002] The present techniques provide for the enhanced removal of acid gas from
a gas stream using treated solvent within a gas processing system including a co-
10 current flow scheme. More specifically, the present techniques provide for the use of
a solvent treater in connection with the final stage of the gas processing system to
enhance the solvent's ability to selectively remove acid gas from the gas stream.

BACKGROUND

[0003] This section is intended to introduce various aspects of the art, which may
15 be associated with exemplary embodiments of the present techniques. This
discussion is believed to assist in providing a framework to facilitate a better
understanding of particular aspects of the present techniques. Accordingly, it should
be understood that this section should be read in this light, and not necessarily as
admissions of prior art.

20 [0004] The production of hydrocarbons from a reservoir oftentimes carries with it
the incidental production of non-hydrocarbon gases. Such gases include
contaminants such as hydrogen sulfide (H₂S) and carbon dioxide (CO₂). When H₂S
and CO₂ are produced as part of a hydrocarbon gas stream, the raw gas stream is
sometimes referred to as "sour gas." The H₂S and CO₂ are often referred to together
25 as "acid gases."

[0005] In addition to hydrocarbon production streams, acid gases may be
associated with synthesis gas streams, or with refinery gas streams. Acid gases may
also be present within so-called flash-gas streams in gas processing facilities. Further,
acid gases may be generated by the combustion of coal, natural gas, or other
30 carbonaceous fuels.

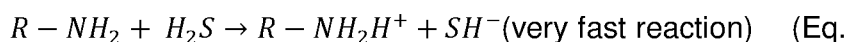
[0006] Natural gas streams may contain not only H₂S and CO₂, but may also contain other “acidic” impurities. These include mercaptans and other trace sulfur compounds (e.g., COS). In addition, natural gas streams may contain water. Such impurities are often removed prior to industrial or residential use. For example, natural gas streams are typically purified to concentrations of less than 4 parts per million (ppm) H₂S and less than 2-3 volume percent (vol. %) CO₂ prior to sale. The extent to which such impurities must be removed is dictated by pipeline regulations, which help to ensure public safety and maintain the integrity of the pipeline by reducing corrosion.

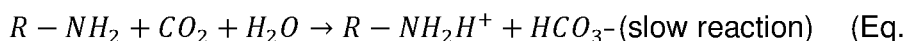
[0007] Acid gas removal is an expensive and equipment-intensive process. The removal of H₂S from natural gas streams is especially complicated due to the safety, health, and environmental considerations when working with toxic H₂S and the processing of sulfur by-products into solid sulfur, or the injection of H₂S-rich gas through acid gas injection methods.

[0008] Various processes have been devised to remove acid gases from a raw natural gas stream. For example, the raw natural gas stream may be treated with a solvent. Solvents may include chemical solvents such as amines. Examples of amines used in sour gas treatment include monoethanol amine (MEA), diethanol amine (DEA), and methyl diethanol amine (MDEA).

[0009] Physical solvents are sometimes used in lieu of chemical solvents. Examples include SELEXOL™ (available from Dow Chemical Company) and RECTISOL® (available from The Linde Group). However, chemical solvents are generally more effective than physical solvents, particularly at feed gas pressures below about 300 psia (2.07 MPa). In some instances, hybrid solvents, meaning mixtures of physical and chemical solvents, have been used. An example is Sulfinol®.

[0010] Chemical solvents, such as amine-based solvents, rely on a chemical reaction between the solvent and the acid gases within the natural gas stream. The reaction process is sometimes referred to as “gas sweetening.” As an example, the reactions of acid gases with tertiary amines ($R_1R_2R_3 - N$) are shown below in Eq. 1 and 2.





2)

As shown in Eq. 1, the reaction of H₂S with the amine is inherently very fast and is often considered instantaneous with respect to diffusion and other kinetic limitations.

5 However, as shown in Eq. 2, the reaction of CO₂ is somewhat slower. The difference in these reaction rates can be utilized to selectively remove one impurity over another within a gas processing system. Note that primary and secondary amines offer a faster reaction route with CO₂ to form carbamates. Consequently, those amines normally cannot be used for selective H₂S removal. Exceptions to this include “sterically-
10 hindered” amines, which prevent the CO₂ from reacting with the amino hydrogen to form carbamates.

[0011] Shale gas often requires H₂S removal with little to no CO₂ removal. Therefore, selective H₂S removal is becoming a central part of the processing facility for natural gas assets. To accomplish this, a solvent with a high selectivity for H₂S
15 may be used. The “H₂S selectivity” of the solvent is defined as the ratio of H₂S removal to CO₂ removal, which is a function of the respective reaction rates. A high H₂S selectivity may be obtained by using solvents that have a slower reaction rate with CO₂. Similarly, the contact time of the gas and liquid phases can be minimized to enhance H₂S uptake over CO₂.

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SUMMARY

[0012] An exemplary embodiment provides a gas processing system. The gas processing system includes a number of co-current contacting systems configured to contact a sour feed gas stream including acid gas with a solvent stream to produce a partially-sweetened gas stream and a rich solvent stream including absorbed acid gas.
25 At least one of the co-current contacting systems is configured to send the rich solvent stream to a regenerator. The regenerator is configured to remove the absorbed acid gas from the rich solvent stream to produce a lean solvent stream. The gas processing system also includes a solvent treater configured to treat at least a portion of the lean solvent stream to produce an enhanced solvent stream, and a final co-current
30 contacting system configured to contact the partially-sweetened gas stream with the enhanced solvent stream to produce a partially-loaded solvent stream and a final gas stream.

[0013] Another exemplary embodiment provides a method for enhanced acid gas removal within a gas processing system. The method includes contacting a sour feed gas stream including acid gas with a solvent stream within a number of co-current contacting systems to produce a partially-sweetened gas stream and a rich solvent stream including absorbed acid gas. The method also includes removing the absorbed acid gas from the rich solvent stream within a regenerator to produce a lean solvent stream, and treating at least a portion of the lean solvent stream within a solvent treater to produce an enhanced solvent stream. The method further includes contacting the partially-sweetened gas stream with the enhanced solvent stream within a final co-current contacting system to produce a partially-loaded solvent stream and a final gas stream.

[0014] Another exemplary embodiment provides a gas processing system. The gas processing system includes a number of co-current contacting systems configured to contact a sour feed gas stream including acid gas with a solvent stream to produce a partially-sweetened gas stream and a rich solvent stream including a first portion of the acid gas. The gas processing system includes a first regenerator configured to remove the first portion of the acid gas from the rich solvent stream to regenerate the solvent stream, and recirculate the solvent stream into at least one of the co-current contacting systems. The gas processing system also includes a final co-current contacting system configured to contact the partially-sweetened gas stream with an enhanced solvent stream to produce a final gas stream and a partially-loaded solvent stream including a second portion of the acid gas. The gas processing system further includes a second regenerator configured to remove the second portion of the acid gas from the partially-loaded solvent stream to produce a lean solvent stream, and a solvent treater configured to treat at least a portion of the lean solvent stream to produce the enhanced solvent stream that is contacted with the partially-sweetened gas stream within the final co-current contacting system.

DESCRIPTION OF THE DRAWINGS

[0015] The advantages of the present techniques are better understood by referring to the following detailed description and the attached drawings, in which:

[0016] **FIG. 1A** is a process flow diagram of a gas processing system that includes a co-current flow scheme and is configured for enhanced acid gas removal;

[0017] **FIG. 1B** is a process flow diagram of another gas processing system that includes a co-current flow scheme and is configured for enhanced acid gas removal;

[0018] **FIG. 2** is a process flow diagram of a separation system that includes a number of co-current contacting systems and a chiller for enhanced acid gas removal;

5 [0019] **FIG. 3** is a process flow diagram of a separation system that includes a number of co-current contacting systems and an anion exchange bed for enhanced acid gas removal;

[0020] **FIG. 4** is a process flow diagram of a first separation system including a number of co-current contacting systems and a second separation including a final co-current contacting system and a corresponding solvent treater;

[0021] **FIG. 5** is a schematic of a co-current contacting system; and

[0022] **FIG. 6** is a process flow diagram showing a method for enhanced acid gas removal within a gas processing system.

DETAILED DESCRIPTION

15 [0023] In the following detailed description section, specific embodiments of the present techniques are described. However, to the extent that the following description is specific to a particular embodiment or a particular use of the present techniques, this is intended to be for exemplary purposes only and simply provides a description of the exemplary embodiments. Accordingly, the techniques are not

20 limited to the specific embodiments described below, but rather, include all alternatives, modifications, and equivalents falling within the true spirit and scope of the appended claims.

[0024] At the outset, for ease of reference, certain terms used in this application and their meanings as used in this context are set forth. To the extent a term used

25 herein is not defined below, it should be given the broadest definition persons in the pertinent art have given that term as reflected in at least one printed publication or issued patent. Further, the present techniques are not limited by the usage of the terms shown below, as all equivalents, synonyms, new developments, and terms or techniques that serve the same or a similar purpose are considered to be within the

30 scope of the present claims.

[0025] “Acid gas” refers to any gas that dissolves in water, producing an acidic solution. Non-limiting examples of acid gases include hydrogen sulfide (H₂S), carbon dioxide (CO₂), sulfur dioxide (SO₂), carbon disulfide (CS₂), carbonyl sulfide (COS), mercaptans, or mixtures thereof.

- 5 [0026] “Co-current contactor” refers to a vessel that receives a stream of gas and a separate stream of solvent in such a manner that the gas stream and the solvent stream contact one another while flowing in generally the same directions within the contactor.

- 10 [0027] The term “co-currently” refers to the internal arrangement of process streams within a unit operation that can be divided into several sub-sections by which the process streams flow in the same direction.

- 15 [0028] The term “dehydrated natural gas stream” refers to a natural gas stream that has undergone a dehydration process. Typically, the dehydrated natural gas stream has a water content of less than 7 lb H₂O/million standard cubic feet for US pipeline applications, or less than 0.1 ppm for LNG applications. Any suitable process for dehydrating the natural gas stream can be used. Typical examples of suitable dehydration processes include, but are not limited to, treatment of the natural gas stream with molecular sieves (for LNG specifications) or dehydration using glycol or methanol (for U.S. pipeline specifications). Alternatively, the natural gas stream can be dehydrated by formation of methane hydrates; for example, using a dehydration process as described in WO 2004/070297.

- 20 [0029] As used herein, the term “dehydration” refers to the pre-treatment of a raw feed gas stream to obtain a dehydrated natural gas stream by partially or completely removing water and, optionally, some heavy hydrocarbons from the feed gas stream.
- 25 This can be accomplished by means of a pre-cooling cycle, against an external cooling loop or a cold internal process stream, for example. Water may also be removed by means of pre-treatment with molecular sieves, e.g. zeolites, or silica gel or alumina oxide or other drying agents. Water may also be removed by means of washing with glycol, monoethylene glycol (MEG), diethylene glycol (DEG) or triethylene glycol
- 30 (TEG), or glycerol. The amount of water in the gas feed stream is suitably less than 1 volume percent (vol. %), preferably less than 0.1 vol. %, more preferably less than 0.01 vol. %.

[0030] As used herein, the term “fluid” refers to gases, liquids, and combinations of gases and liquids, as well as to combinations of gases and solids, and combinations of liquids and solids.

[0031] The term “flue gas” refers to any gas stream generated as a by-product of hydrocarbon combustion.

[0032] The term “gas” is used interchangeably with “vapor,” and is defined as a substance or mixture of substances in the gaseous state as distinguished from the liquid or solid state. Likewise, the term “liquid” means a substance or mixture of substances in the liquid state as distinguished from the gas or solid state.

[0033] A “hydrocarbon” is an organic compound that primarily includes the elements hydrogen and carbon, although nitrogen, sulfur, oxygen, metals, or any number of other elements may be present in small amounts. As used herein, the term “hydrocarbon” generally refers to components found in natural gas, oil, or chemical processing facilities. Moreover, the term “hydrocarbon” may refer to components found in raw natural gas, such as CH₄, C₂H₆, C₃ isomers, C₄ isomers, benzene, and the like.

[0034] With respect to fluid processing equipment, the term “in series” means that two or more devices are placed along a flow line such that a fluid stream undergoing fluid separation moves from one item of equipment to the next while maintaining flow in a substantially constant downstream direction. Similarly, the term “in line” means that two or more components of a fluid mixing and separating device are connected sequentially or, more preferably, are integrated into a single tubular device.

[0035] The term “industrial plant” refers to any plant that generates a gas stream containing at least one hydrocarbon or an acid gas. One non-limiting example is a coal-powered electrical generation plant. Another example is a cement plant that emits CO₂ at low pressures.

[0036] The term “liquid solvent” refers to a fluid in substantially liquid phase that preferentially absorbs one component over another. For example, a liquid solvent may preferentially absorb an acid gas, thereby removing or “scrubbing” at least a portion of the acid gas component from a gas stream. Moreover, a liquid solvent may preferentially absorb one acid gas over another.

[0037] “Natural gas” refers to a multi-component gas obtained from a crude oil well or from a subterranean gas-bearing formation. The composition and pressure of natural gas can vary significantly. A typical natural gas stream contains methane (CH₄) as a major component, i.e., greater than 50 mole percent (mol. %) of the natural gas stream. The natural gas stream can also contain ethane (C₂H₆), higher molecular weight hydrocarbons (e.g., C₃-C₂₀ hydrocarbons), acid gases (e.g., carbon dioxide and hydrogen sulfide), or any combinations thereof. The natural gas can also contain minor amounts of contaminants such as water, nitrogen, iron sulfide, wax, crude oil, or any combinations thereof. The natural gas stream may be substantially purified prior to use in embodiments described herein, so as to remove compounds that may act as poisons.

[0038] “Non-absorbing gas” means a gas that is not significantly absorbed by a solvent during a gas treating or conditioning process.

[0039] As used herein, “purification” includes separation processes by which impurities that may cause problems to downstream processes are removed.

[0040] “Solvent” refers to a substance capable at least in part of dissolving or dispersing other substances, such as to provide or form a solution. The solvent may be polar, nonpolar, neutral, protic, aprotic, or the like. The solvent may include any suitable element, molecule, or compound, such as methanol, ethanol, propanol, glycols, ethers, ketones, other alcohols, amines, salt solutions, or the like. The solvent may include physical solvents, chemical solvents, or the like. The solvent may operate by any suitable mechanism, such as physical absorption, chemical absorption, chemisorption, physisorption, adsorption, pressure swing adsorption, temperature swing adsorption, or the like. Specific solvents that are useful for acid gas absorption include, but are not limited to, monoethanolamine (MEA), 2(2-aminoethoxy) ethanol [Diglycolamine® (DGA)], diethanolamine (DEA), diisopropanolamine (DIPA), methyldiethanolamine (MDEA), triethyleneamine, FLEXSORB® SE, 2-amino-2-methyl-1-propanol (AMP), or formulated amines such as FLEXSORB® SE PLUS, the UCARSOL™ family of products, or formulated MDEA solutions.

[0041] “Substantial” when used in reference to a quantity or amount of a material, or a specific characteristic thereof, refers to an amount that is sufficient to provide an effect that the material or characteristic was intended to provide. The exact degree of

deviation allowable may depend, in some cases, on the specific context.

[0042] The term “sweetened natural gas stream” refers to a natural gas stream that has had at least a portion of acid gas components removed.

Overview

5 [0043] The present techniques provide for the enhanced removal of acid gas from a sour gas stream using treated solvent within a gas processing system including a co-current flow scheme. In various embodiments, the present techniques are used to selectively remove H₂S from a sour natural gas stream. Specifically, the sour natural gas stream may be purified to a concentration of less than 4 ppm H₂S to meet pipeline
10 regulations.

[0044] The co-current flow scheme may utilize any number of co-current contacting systems connected in series within a pipe. A natural gas stream and a liquid solvent stream move together, i.e., co-currently, within the co-current contacting systems. In some embodiments, the natural gas stream and the liquid solvent stream move
15 together generally along the longitudinal axis of the co-current contacting system.

[0045] Each co-current contacting system within the gas processing system may include a co-current contactor that facilitates the absorption of acid gas, such as H₂S and CO₂, into the solvent stream. In addition, each co-current contacting system may include a separator that is capable of separating the natural gas stream from the solvent stream with the absorbed acid gas, producing a sweetened, liquid-free natural
20 gas stream.

[0046] In some embodiments, the solvent stream is an amine-based solvent that is capable of absorbing acid gases, such as H₂S and CO₂, within the natural gas stream. For example, the solvent may include, but is not limited to, monoethanolamine (MEA),
25 2(2-aminoethoxy) ethanol [Diglycolamine® (DGA)], diethanolamine (DEA), diisopropanolamine (DIPA), methyldiethanolamine (MDEA), triethylethanolamine, FLEXSORB® SE, 2-amino-2-methyl-1-propanol (AMP), or formulated amines such as FLEXSORB® SE PLUS, the UCARSOL™ family of products, or formulated MDEA solutions.

30 [0047] According to current techniques, as the solvent flows through the series of co-current contacting systems, the solvent becomes loaded with contaminants, such

as H₂S and CO₂. For rapidly-reacting components like H₂S, this can result in a pinch condition near the outlet of the co-current contacting systems as the CO₂, which is thermodynamically favored for absorption, begins to displace the H₂S. For a typical shale gas application, in which it is desirable to remove up to 1,000 ppm H₂S, three or four co-current contacting systems may be used. However, it can be particularly challenging to reach a 4 ppm H₂S specification in the final stage because the solvent will load with H₂S and CO₂ as it flows through the device, thus reducing its capacity to further reduce the H₂S concentration of the natural gas stream.

[0048] Therefore, according to embodiments described herein, enhanced H₂S removal is achieved through the use of a solvent treater upstream of the final co-current contacting system. The solvent treater is configured to modify the solvent stream for increased H₂S absorption in the final co-current contacting system, as described further with respect to FIGS. 1A, 1B, 2, 3, 4, and 6.

Gas Processing System

[0049] FIG. 1A is a process flow diagram of a gas processing system **100** that includes a co-current flow scheme and is configured for enhanced acid gas removal. The gas processing system **100** may be used for the removal of acid gas components, such as H₂S and CO₂, from a feed gas stream **102**. In addition, the gas processing system **100** may be used for the removal of water or other impurities from the feed gas stream **102**. The gas processing system **100** may employ a number of co-current contacting systems **104A-F**. Each co-current contacting system **104A-F** may include a co-current contactor and a separator, for example, as discussed further with respect to FIG. 5.

[0050] The feed gas stream **102** may be, for example, a natural gas stream from a hydrocarbon production operation, a flue gas stream from a power plant, or a synthesis gas (syn-gas) stream. If the feed gas stream **102** is a syn-gas stream, the feed gas stream **102** may be cooled and filtered before being introduced into the gas processing system **100**. The feed gas stream **102** may also be a flash gas stream taken from a flash drum in a gas processing facility itself. In addition, the feed gas stream **102** may be a tail gas stream from a Claus sulfur recovery process or an impurities stream from a solvent regenerator. Furthermore, the feed gas stream **102** may be an exhaust emission from a cement plant or other industrial plant. In this instance, CO₂ may be

absorbed from excess air or from a nitrogen-containing flue gas.

[0051] The feed gas stream **102** may include a non-absorbing gas, such as methane, and one or more impurities, such as CO₂ and H₂S. In some embodiments, the feed gas stream **102** includes a large amount of H₂S, such as, for example, on the order of 1,000 ppm H₂S. The gas processing system **100** may convert the feed gas stream **102** into a sweetened gas stream **106** by removing the CO₂ and H₂S. In various embodiments, the sweetened gas stream **106** contains concentrations of less than 4 ppm H₂S and less than 2-3 vol. % CO₂.

[0052] In operation, the feed gas stream **102** may be flowed into a first co-current contacting system **104A**, where it is mixed with a solvent stream **108**. In various embodiments, the solvent stream **108** includes an amine solution, such as monoethanol amine (MEA), diethanol amine (DEA), or methyldiethanol amine (MDEA). The solvent stream **108** may include a lean solvent that has undergone a desorption process for the removal of acid gas impurities. For example, in the gas processing system **100** shown in FIG. 1A, the solvent stream **108** introduced into the first co-current contacting system **104A** includes a semi-lean solvent that is taken from a central portion of a regenerator **110**. A lean solvent stream **112** taken from the regenerator **110** may also be directed into the fifth and final co-current contacting systems **104E** and **104F**.

[0053] In various embodiments, the gas processing system **100** employs a series of co-current contacting systems **104A-F**. Each co-current contacting system **104A-F** removes a portion of the acid gas content from the feed gas stream **102**, thereby releasing a progressively sweetened gas stream in a downstream direction. The final co-current contacting system **104F** provides the final sweetened gas stream **106**.

[0054] Before entering the first co-current contacting system **104A**, the feed gas stream **102** may pass through an inlet separator **114**. The inlet separator **114** may be used to clean the feed gas stream **102** by filtering out impurities, such as brine and drilling fluids. Some particle filtration may also take place. The cleaning of the feed gas stream **102** can prevent foaming of solvent during the acid gas treatment process.

[0055] In some embodiments, the feed gas stream **102** may also be pretreated upstream of the inlet separator **114** or the first co-current contacting system **104A**. For

example, the feed gas stream **102** may undergo a water wash to remove glycol or other chemical additives. This may be accomplished via a separate processing loop (not shown) wherein water is introduced to the gas, such as via an additional co-current contacting system. Water has an affinity for glycol and will pull the glycol out of the feed gas stream **102**. This, in turn, will help control foaming within the co-current contacting systems **104A-F**. In the case of flue gas applications, corrosion inhibitors may be added to the solvent to retard the reaction of O₂ with the steel in the processes.

[0056] As shown in FIG. 1A, the solvent stream **108** is flowed into the first co-current contacting system **104A**. Movement of the semi-lean solvent stream **108** into the first co-current contacting system **104A** may be aided by pumps **116A** and **116B** and a cooler **117**. The cooler **117** may cause the solvent stream **108** to flow into the first co-current contacting system **104A** at a suitable temperature, while the pumps **116A** and **116B** may cause the solvent stream **108** to flow into the first co-current contacting system **104A** at a suitable pressure of, for example, about 15 psia to about 1,500 psig.

[0057] Once inside the first co-current contacting system **104A**, the feed gas stream **102** and the solvent stream **108** move along the longitudinal axis of the first co-current contacting system **104A**. As they travel, the liquid amine (or other treating solution) interacts with the H₂S and CO₂ in the feed gas stream **102**, causing the H₂S and CO₂ to chemically attach to or be absorbed by the amine molecules. A first partially-loaded, or “rich,” solvent stream **118A** may be flowed out of a bottom portion of the first co-current contacting system **104A**. In addition, a first partially-sweetened gas stream **120A** may be flowed out of a top portion of the first co-current contacting system **104A** and into a second co-current contacting system **104B**.

[0058] As shown in the example illustrated in FIG. 1A, a third co-current contacting system **104C** may be provided after the second co-current contacting system **104B**, and a fourth co-current contacting system **104D** may be provided after the third co-current contacting system **104C**. In addition, a fifth co-current contacting system **104E** may be provided after the fourth co-current contacting system **104D**, and a final co-current contacting system **104F** may be provided after the fifth co-current contacting system **104E**. Each of the second, third, fourth, and fifth co-current contacting systems **104B**, **104C**, **104D**, and **104E** may generate a respective partially-sweetened gas

stream **120B**, **120C**, **120D**, and **120E**. In addition, each of the second, third, fourth, fifth, and final co-current contacting systems **104B**, **104C**, **104D**, **104E**, and **104F** may generate a respective partially-loaded solvent stream **118B**, **118C**, **118D**, **118E**, and **118F**. If an amine is used as the solvent stream **108**, the partially-loaded solvent streams **118A-F** may include rich amine solutions. In the gas processing system **100**, the second partially-loaded solvent stream **118B** merges with the first partially-loaded solvent stream **118A** and goes through a regeneration process in the regenerator **110**.

[0059] As the progressively-sweetened gas streams **120A-E** are generated, the gas pressure in the gas processing system **100** will progressively decrease. As this occurs, the liquid pressure of the progressively-rich solvent streams **118A-F** may be correspondingly increased. This may be accomplished by placing one or more booster pumps (not shown) between each co-current contacting system **104A-F** to boost liquid pressure in the gas processing system **100**.

[0060] In the gas processing system **100**, solvent streams may be regenerated by flowing the partially-loaded solvent streams **118A** and **118B** through a flash drum **122**. Absorbed natural gas **124** may be flashed from the partially-loaded solvent streams **118A** and **118B** within the flash drum **122**, and may be flowed out of the flash drum **122** via an overhead line **126**.

[0061] The resulting rich solvent stream **128** may be flowed from the flash drum **122** to the regenerator **110**. The rich solvent stream **128** may be introduced into the regenerator **110** for desorption. The regenerator **110** may include a stripper portion **130** including trays or other internals (not shown). The stripper portion **130** may be located directly above a reboiler portion **132**. A heat source **134** may be provided with the reboiler portion **132** to generate heat. The regenerator **110** produces the regenerated, lean solvent stream **112** that is recycled for re-use in the fifth and final co-current contacting systems **104E** and **104F**. Stripped overhead gas from the regenerator **110**, which may include concentrated H_2S and CO_2 , may be flowed out of the regenerator **110** as an overhead impurities stream **136**.

[0062] The overhead impurities stream **136** may be flowed into a condenser **138**, which may cool the overhead impurities stream **136**. The resulting cooled impurities stream **140** may be flowed through a reflux accumulator **142**. The reflux accumulator **142** may separate any remaining liquid, such as condensed water, from the impurities

stream **140**. This may result in the generation of a substantially pure acid gas stream **144**, which may be flowed out of the reflux accumulator **142** via an overhead line **146**.

[0063] In some embodiments, the H₂S in the acid gas stream **144** is then converted into elemental sulfur using a sulfur recovery unit (not shown). The sulfur recovery unit
5 may be a so-called Claus unit. Those of ordinary skill in the art will understand that a “Claus process” is a process that is sometimes used by the natural gas and refinery industries to recover elemental sulfur from H₂S-containing gas streams.

[0064] In practice, the “tail gas” from the Claus process, which may include H₂S, SO₂, CO₂, N₂, and water vapor, can be reacted to convert the SO₂ to H₂S via
10 hydrogenation. The hydrogenated tail gas stream has a high partial pressure, a large amount of CO₂, e.g., more than 50%, and a small amount of H₂S, e.g., a few percent or less. This type of gas stream, which is typically near atmospheric pressure, is amenable to selective H₂S removal. The recovered H₂S may be recycled to the front of the Claus unit, or may be sequestered downstream. Alternatively, a direct oxidation
15 of the H₂S to elemental sulfur may be performed using various processes known in the field of gas separation.

[0065] Because the H₂S reaction is instantaneous relative to the CO₂ reactions, lowering the residence time, i.e., the contact time between the vapor and liquid phases, will result in less CO₂ being absorbed into the solvent. The design of the co-current
20 contacting systems **104A-F** enhances selective H₂S removal due to the short contact time inherent in the equipment design.

[0066] As shown in FIG. 1A, a residual liquid stream **148** may be flowed out of the bottom of the reflux accumulator **142**. The residual liquid stream **148** may be flowed through a reflux pump **150**, which may boost the pressure of the residual liquid stream
25 **148** and pump the residual liquid stream **148** into the regenerator **110**. The residual liquid stream **148** may be flowed out of the regenerator **110**, for example, from the bottom of the reboiler portion **132** as part of the lean solvent stream **112**. Some water may be added to the lean solvent stream **112** to balance the loss of water vapor to the sweetened gas stream **106** and the acid gas stream **144**. This water may be added
30 at an intake or suction of the reflux pump **150**.

[0067] The lean solvent stream **112** may be at a low pressure. Accordingly, the

lean solvent stream **112** may be passed through a pressure boosting pump **152**. From the pressure boosting pump **152**, the lean solvent stream **112** may be flowed through a cooler **154**. The cooler **154** may cool the lean solvent stream **112** back to near ambient temperatures after it has been heated by the regenerator **110**.

5 [0068] In some embodiments, the lean solvent stream **112** may then be flowed into a solvent tank **156**. In other embodiments, the solvent tank **156** is off-line and provides a reservoir for the lean solvent stream **112**.

10 [0069] Movement of the lean solvent stream **112** towards the fifth and final co-current contacting systems **104E** and **104F** may be aided by a pump **158**. The pump **158** may cause the lean solvent stream **112** to flow at a suitable pressure, for example, of about 15 psia to about 1,500 psig.

15 [0070] A first portion **160** of the lean solvent stream **112** may be joined with the partially-loaded solvent stream **118F** and flowed into the fifth co-current contacting system **104E**. A second portion **162** of the lean solvent stream **112** may be flowed into a solvent treater **164**, which is configured to treat the lean solvent stream **112** to produce an enhanced solvent stream **166**. According to embodiments described herein, the enhanced solvent stream **166** is a treated solvent stream that is capable of absorbing a higher concentration of acid gas than the lean solvent stream **112**. The enhanced solvent stream **166** may be a highly H₂S-selective solvent stream that is capable of selectively absorbing a higher concentration of H₂S as opposed to CO₂. In various embodiments, an H₂S concentration of less than 4 ppm within the final sweetened gas stream **106** is achieved using the enhanced solvent stream **166**.

20 [0071] In various embodiments, the solvent treater **164** is a chiller that is configured to produce the enhanced solvent stream **166** by cooling the lean solvent stream **112** to at least about ambient, such as about 20 °C to 25 °C, or to at least about 5 °C below ambient, or to at least about 10 °C below ambient, or to at least about 20 °C below ambient, or to a temperature that is the same as, or slightly lower than, that of the partially-sweetened gas stream **120E** entering the final co-current contacting system **104F**. For example, the solvent treater **164** may be an ammonia chiller, a cold water flow from a cooling water tower, or any other suitable type of chiller.

30 [0072] In some embodiments, an analyzer (not shown) and a controller (not shown)

may be coupled to the solvent treater **164**, as described in further detail with respect to FIG. 2. The analyzer may be configured to determine the acid gas concentration of the final sweetened gas stream **106** exiting the final co-current contacting system **104F**. The controller may then adjust the temperature of the solvent treater **164** based on the analysis of the sweetened gas stream **106**.

[0073] In some embodiments, the solvent treater **164** is an anion exchange bed that produces the enhanced solvent stream **166** by removing residual HS⁻ and HCO₃⁻ from the lean solvent stream **112**. In other embodiments, the solvent treater **164** is an electrodialysis unit that is configured to produce the enhanced solvent stream **166** by reducing the lean load of the lean solvent stream **112**. In these embodiments, the solvent treater **164** may also be used to remove heat stable salt contaminants from the lean solvent stream **112**.

[0074] In other embodiments, the solvent treater **164** produces the enhanced solvent stream **166** by injecting an enhancement fluid into the lean solvent stream **112** that enhances the solvent's ability to selectively absorb acid gas. For example, a liquid H₂S scavenger may be added to the lean solvent stream **112** to increase the solvent's ability to remove residual H₂S within the partially-sweetened gas stream **120E**. In some embodiments, a controlled bypass (not shown) may be used to ensure that the final sweetened gas stream **106** has a concentration of 3-4 ppm H₂S, not 0 ppm, which would be a waste of scavenger.

[0075] The process flow diagram of FIG. 1A is not intended to indicate that the gas processing system **100** is to include all of the components shown in FIG. 1A. Further, any number of additional components may be included within the gas processing system **100**, depending on the details of the specific implementation. For example, the gas processing system **100** may include any suitable types of heaters, chillers, condensers, liquid pumps, gas compressors, blowers, bypass lines, other types of separation and/or fractionation equipment, valves, switches, controllers, and pressure-measuring devices, temperature-measuring devices, level-measuring devices, or flow-measuring devices, among others. Moreover, the gas processing system **100** may include any number of additional co-current contacting systems.

[0076] In some embodiments, the lean solvent stream **112** taken from the regenerator **110** is only directed into the final co-current contacting system **104F**, not

the fifth co-current contacting system **104E**. In those embodiments, the entire lean solvent stream **112** is sent through the solvent treater **164** to produce the enhanced solvent stream **166**.

[0077] In some embodiments, a portion of the rich solvent stream **128** is acidified
5 before the rich solvent stream **128** enters the regenerator **110**. This may be accomplished by adding 1-2 weight percent (wt. %) phosphoric acid to the rich solvent stream **128**. Acidifying the rich solvent stream **128** may allow the rich solvent stream **128** to release more acid gas during the regeneration process.

[0078] FIG. 1B is a process flow diagram of another gas processing system **168**
10 that includes a co-current flow scheme and is configured for enhanced acid gas removal. Like numbered items are as described with respect to FIG. 1A. Operation of the gas processing system **168** of FIG. 1B is similar to that of the gas processing system **100** of FIG. 1A. However, in the gas processing system **168** of FIG. 1B, the first co-current contacting system **104A** receives the partially-loaded solvent stream
15 **118B** from the second co-current contacting system **104B**. Therefore, the gas processing system **168** does not include the semi-lean solvent stream **108**.

[0079] Because the partially-loaded solvent stream **118B** received by the first co-current contacting system **104A** in FIG. 1B has already been processed through the second co-current contacting system **104B**, the partially-loaded solvent stream **118B**
20 received by the first co-current contacting system **104A** may be very rich. For this reason, it may be desirable to provide some level of intermediate processing of the partially-loaded solvent stream **118B**.

[0080] Alternatively, a semi-lean solvent stream could be taken from other sweetening operations in the gas processing system **168** and used, at least in part, as
25 an amine solution for the first or second co-current contacting system **104A** or **104B**. In this respect, there are situations in which a single type of solvent is used for more than one service in the gas processing system **168**. This is referred to as integrated gas treatment. For example, MDEA may be used both for high-pressure, H₂S-selective acid gas removal, as well as in a Claus tail gas treating (TGT) process. The
30 rich amine stream from the TGT process is not heavily loaded with H₂S and CO₂, owing to the low pressure of the process. Thus, in some embodiments, the rich amine stream from the TGT process is used as a semi-lean stream for the first or second co-

current contacting system **104A** or **104B**. The semi-lean stream (not shown) may be pumped to a suitable pressure and injected into the first or second co-current contacting system **104A** or **104B**, possibly along with the partially-loaded solvent stream from the succeeding co-current contacting system.

5 [0081] Further, in the gas processing system **168** of FIG. 1B, the first partially-loaded solvent solution **118A** is flowed through a heat exchanger **170** after being flowed through the flash drum **122**. Within the heat exchanger **170**, the temperature of the first partially-loaded solvent solution **118A** is increased via heat exchange with the lean solvent stream **112** taken from the regenerator **110**. This serves to heat the
10 first partially-loaded solvent solution **118A** before introduction into the regenerator **110**, while cooling the lean solvent stream **112**.

[0082] The process flow diagram of FIG. 1B is not intended to indicate that the gas processing system **168** is to include all of the components shown in FIG. 1B. Further, any number of additional components may be included within the gas processing
15 system **168**, depending on the details of the specific implementation. For example, the gas processing system **168** may include any suitable types of heaters, chillers, condensers, liquid pumps, gas compressors, blowers, bypass lines, other types of separation and/or fractionation equipment, valves, switches, controllers, and pressure-measuring devices, temperature-measuring devices, level-measuring devices, or flow-
20 measuring devices, among others. Moreover, the gas processing system **168** may include any number of additional co-current contacting systems.

[0083] According to the embodiments described in FIGS. 1A and 1B, the gas processing systems **100** and **168** include a countercurrent arrangement of co-current contactors. However, it is to be understood that the embodiments described herein
25 also apply to a parallel configuration of co-current contactors where each stage is fed with a freshly-generated solvent stream. Moreover, the individual co-current contactors may be arranged in a myriad of different configurations, including both horizontal and vertical sections, stages with or without in-line separation immediately following contacting, and with dehydration, H₂S removal, and CO₂ removal occurring
30 in subsequent portions of a single in-line device. In addition, all of the co-current contacting systems may be bundled into a single pressure vessel oriented vertically or horizontally.

Separation System

[0084] FIG. 2 is a process flow diagram of a separation system **200** that includes a number of co-current contacting systems **202A-C** and a chiller **204** for enhanced acid gas removal. The separation system **200** may be implemented as part of a gas processing system, such as the gas processing system **100** or **168** discussed with respect to FIG. 1A or 1B. The gas processing system may utilize a number of co-current contacting systems **202A-C** connected in series, such as the co-current contacting systems **104A-F** discussed with respect to FIGS. 1A and 1B. In the illustrative arrangement shown in FIG. 2, a first co-current contacting system **202A**, a second co-current contacting system **202B**, and a third co-current contacting system **202C** are provided.

[0085] A sour feed gas stream **206** may be flowed into the first co-current contacting system **202A**. The first co-current contacting system **202A** may generate a first partially-sweetened gas stream **208A**, which may be flowed from the first co-current contacting system **202A** to the second co-current contacting system **202B**. The second co-current contacting system **202B** may then generate a second partially-sweetened gas stream **208B**, which may be flowed from the second co-current contacting system **202B** to the third co-current contacting system **202C**. In some embodiments, the third co-current contacting system **202C** generates a final sweetened gas stream **210**.

[0086] Each of the first, second, and third co-current contacting systems **202A-C** also generates a respective rich solvent stream **212A-C**. The third rich solvent stream **212C** may be directed back to the second co-current contacting system **202B**, and the second rich solvent stream **212B** may be directed back to the first co-current contacting system **202A**. In addition, the first rich solvent stream **212A** may be returned to a regenerator **214**. In some embodiments, the regenerator **214** is the same as, or similar to, the regenerator **110** discussed with respect to FIGS. 1A and 1B.

[0087] The regenerator **214** may remove absorbed acid gases and other impurities from the first rich solvent stream **212A**, producing a lean solvent stream **216**. The lean solvent stream **216** may then be sent through the chiller **204**, which may reduce the temperature of the lean solvent stream **216** to produce an enhanced solvent stream **218**. In various embodiments, the chiller **204** corresponds to the solvent treater **164**

described with respect to FIGS. 1A and 1B. Further, in various embodiments, the enhanced solvent stream **218** is a solvent stream that is capable of absorbing a higher concentration of acid gas from the second partially-sweetened gas stream **208B**. For example, the enhanced solvent stream **218** may be a highly H₂S-selective solvent stream that is capable of absorbing a higher concentration of H₂S as opposed to CO₂.

[0088] The chiller **204** may be any suitable type of chiller that is capable of lowering the temperature of the lean solvent stream **216** to at least 10 °F below ambient, or to a temperature that is the same as, or slightly lower than, that of the second partially-sweetened gas stream **208B** entering the third co-current contacting system **202C**. For example, the chiller **204** may be an ammonia chiller or a cold water flow from a water tower. Moreover, in some embodiments, the chiller **204** includes a small addition system for injecting enhancement fluid into the lean solvent stream **216**.

[0089] From the chiller **204**, the enhanced solvent stream **218** may be flowed into the third co-current contacting system **202C**. Within the third co-current contacting system **202C**, the enhanced solvent stream **218** contacts the second partially-sweetened gas stream **208B** and absorbs an increased amount of acid gas, such as H₂S, from the second partially-sweetened gas stream **208B**. The resulting sweetened gas stream **210** may include a low concentration of acid gas, such as, for example, less than 4 ppm H₂S.

[0090] The separation system **200** may also include an analyzer **220** and a controller **222**. The analyzer **220** may be configured to conduct an external analysis of the final sweetened gas stream **210** exiting the third co-current contacting system **202C** to determine the acid gas concentration, such as the H₂S and CO₂ concentrations, of the final sweetened gas stream **210**. The controller **222** may then adjust the temperature of the chiller **204** based on the analysis of the sweetened gas stream **210**. In some embodiments, using the analyzer **220** and the controller **222** within the separation system **200** results in energy savings because the chiller **204** may be operated at different temperatures depending on the acid gas concentration of the final sweetened gas stream **210**.

[0091] The process flow diagram of FIG. 2 is not intended to indicate that the separation system **200** is to include all of the components shown in FIG. 2. Further, any number of additional components may be included within the separation system

200, depending on the details of the specific implementation. For example, any number of additional co-current contacting systems may be included within the separation system **200**.

[0092] FIG. 3 is a process flow diagram of a separation system **300** that includes a number of co-current contacting systems **302A-C** and an anion exchange bed **304** for enhanced acid gas removal. The separation system **300** may be implemented as part of a gas processing system, such as the gas processing system **100** or **168** discussed with respect to FIG. 1A or 1B. The gas processing system may utilize a number of co-current contacting systems **302A-C** connected in series, such as the co-current contacting systems **104A-F** discussed with respect to FIGS. 1A and 1B. In the illustrative arrangement shown in FIG. 3, a first co-current contacting system **302A**, a second co-current contacting system **302B**, and a third co-current contacting system **302C** are provided.

[0093] A sour feed gas stream **306** may be flowed into the first co-current contacting system **302A**. The first co-current contacting system **302A** may generate a first partially-sweetened gas stream **308A**, which may be flowed from the first co-current contacting system **302A** to the second co-current contacting system **302B**. The second co-current contacting system **302B** may then generate a second partially-sweetened gas stream **308B**, which may be flowed from the second co-current contacting system **302B** to the third co-current contacting system **302C**. In some embodiments, the third co-current contacting system **302C** generates a final sweetened gas stream **310**.

[0094] Each of the first, second, and third co-current contacting systems **302A-C** also generates a respective rich solvent stream **312A-C**. The third rich solvent stream **312C** may be directed back to the second co-current contacting system **302B**, and the second rich solvent stream **312B** may be directed back to the first co-current contacting system **302A**. In addition, the first rich solvent stream **312A** may be returned to a regenerator **314**. In some embodiments, the regenerator **314** is the same as, or similar to, the regenerator **110** discussed with respect to FIGS. 1A and 1B.

[0095] The regenerator **314** may remove absorbed acid gases and other impurities from the first rich solvent stream **312A**, producing a lean solvent stream **316**. The lean solvent stream **316** may then be sent through the anion exchange bed **304**. In various

embodiments, the anion exchange bed **304** corresponds to the solvent treater **164** described with respect to FIGS. 1A and 1B.

[0096] The anion exchange bed **304** may produce an enhanced solvent stream **318** by removing residual acid gases from the lean solvent stream **316**. In addition, the
5 anion exchange bed **304** may remove heat stable salt contaminants from the lean solvent stream **316**.

[0097] According to embodiments described herein, the enhanced solvent stream **318** is a solvent stream that is capable of absorbing a higher concentration of acid gas from the second partially-sweetened gas stream **308B**. For example, in some
10 embodiments, the enhanced solvent stream **318** may be a highly H₂S-selective solvent stream that is capable of absorbing H₂S more quickly than CO₂. In those embodiments, the anion exchange bed **304** produces the enhanced solvent stream **318** by removing residual HS⁻ and HCO₃⁻ from the lean solvent stream **316**.

[0098] The process flow diagram of FIG. 3 is not intended to indicate that the
15 separation system **300** is to include all of the components shown in FIG. 3. Further, any number of additional components may be included within the separation system **300**, depending on the details of the specific implementation. For example, any number of additional co-current contacting systems may be included within the separation system **300**.

[0099] FIG. 4 is a process flow diagram of a first separation system **400** including
20 a number of co-current contacting systems **402A-C** and a second separation system **404** including a final co-current contacting system **406** and a corresponding solvent treater **408**. The first and second separation systems **400** and **404** may be implemented as part of a gas processing system, such as a gas processing system
25 that is similar to the gas processing system **100** or **168** discussed with respect to FIG. 1A or 1B. The gas processing system may utilize a number of co-current contacting systems **402A-C** and **406** connected in series, such as the co-current contacting systems **104A-F** discussed with respect to FIGS. 1A and 1B. In the illustrative arrangement shown in FIG. 4, the first separation system **400** includes a first co-
30 current contacting system **402A**, a second co-current contacting system **402B**, and a third co-current contacting system **402C**, and the second separation system **404** includes the final co-current contacting system **406**.

[0100] Within the first separation system **400**, a sour feed gas stream **410** may be flowed into the first co-current contacting system **402A**. The first co-current contacting system **402A** may generate a first partially-sweetened gas stream **412A**, which may be flowed from the first co-current contacting system **402A** to the second co-current contacting system **402B**. The second co-current contacting system **402B** may then generate a second partially-sweetened gas stream **412B**, which may be flowed from the second co-current contacting system **402B** to the third co-current contacting system **402C**. The third co-current contacting system **402C** may then generate a third partially-sweetened gas stream **412C**, which may be flowed from the second co-current contacting system **402B** to the final co-current contacting system **406** within the second separation system **404**.

[0101] Each of the first, second, and third co-current contacting systems **402A-C** also generates a respective rich solvent stream **414A-C**. The third rich solvent stream **414C** may be directed back to the second co-current contacting system **402B**, and the second rich solvent stream **414B** may be directed back to the first co-current contacting system **402A**. In addition, the first rich solvent stream **414A** may be returned to a first regenerator **416**. In some embodiments, the first regenerator **416** is the same as, or similar to, the regenerator **110** discussed with respect to FIGS. 1A and 1B.

[0102] The first regenerator **416** may remove absorbed acid gases and other impurities from the first rich solvent stream **414A**, producing a first lean solvent stream **418**. The first lean solvent stream **418** may then be recirculated into the third co-current contacting system **402C**.

[0103] Within the final co-current contacting system **406**, the third partially-sweetened gas stream **412C** is contacted with an enhanced solvent stream **420**, producing a final sweetened gas stream **422**. According to embodiments described herein, the enhanced solvent stream **420** is a treated solvent stream that is capable of absorbing a high concentration of acid gas. Therefore, the final sweetened gas stream **422** may include a low concentration of acid gas, such as, for example, less than 4 ppm H₂S.

[0104] The final co-current contacting system **406** also generates a partially-loaded solvent stream **424**. The partially-loaded solvent stream **424** may be sent to a second

regenerator **426**. In some embodiments, the second regenerator **426** is the same as, or similar to, the regenerator **110** discussed with respect to FIGS. 1A and 1B. The second regenerator **426** may remove absorbed acid gases and other impurities from the partially-loaded solvent stream **424**, producing a second lean solvent stream **428**.

5 [0105] According to embodiments described herein, the second lean solvent stream **428** is then sent through the solvent treater **408**. The solvent treater **408** is configured to treat the second lean solvent stream **428** to produce the enhanced solvent stream **420**. In various embodiments, the solvent treater **408** is the same as, or similar to, the solvent treater **164** described with respect to FIGS. 1A and 1B. The
10 enhanced solvent stream **420** may then be sent through a pressure boosting pump **430** before being directed back into the final co-current contacting system **406**.

[0106] The process flow diagram of FIG. 4 is not intended to indicate that the first and second separation systems **400** and **404** are to include all of the components shown in FIG. 4. Further, any number of additional components may be included
15 within the first and second separation systems **400** and **404**, depending on the details of the specific implementation. For example, any number of additional co-current contacting systems may be included within the first separation system **400** or the second separation system **404**.

Co-Current Contacting System

20 [0107] FIG. 5 is a schematic of a co-current contacting system **500**. The co-current contacting system **500** may provide for the separation of components within a gas stream. In addition, the co-current contacting system **500** may aid in the implementation of various gas processing systems, such as the gas processing systems **100** and **168** of FIGS. 1A and 1B, where the rapid separation of components
25 is desired. In some embodiments, the co-current contacting system **500** is one of the co-current contacting systems **104A-F**, **202A-C**, **302A-C**, **402A-C**, and **406** discussed with respect to FIGS. 1A, 1B, 2, 3, and 4.

[0108] The co-current contacting system **500** may include a co-current contactor **502** that is positioned in-line within a pipe **504**. The co-current contactor **502** may
30 include a number of components that provide for the efficient contacting of a liquid solvent stream with a flowing gas stream **506**. The liquid solvent stream can be used

for the separation of impurities, such as H₂S and CO₂, from a gas stream **506**.

[0109] In various embodiments, the co-current contactor **502** includes a mixer **508** and a mass transfer section **510**. As shown in FIG. 5, the gas stream **506** may be flowed through the pipe **504** and into the mixer **508**. A liquid solvent stream **512** may also be flowed into the mixer **508**, for example, through a hollow space **514** coupled to flow channels **516** in the mixer **508**.

[0110] From the flow channels **516**, the liquid solvent stream **512** is released into the gas stream **506** as fine droplets through injection orifices **518**, and is then flowed into the mass transfer section **510**. This may result in the generation of a treated gas stream **520** within the mass transfer section **510**. The treated gas stream **520** may include small liquid droplets dispersed in a gas phase. The liquid droplets may include impurities from the gas stream **506** that were absorbed or dissolved into the liquid solvent stream **512**.

[0111] The treated gas stream **520** may be flowed from the mass transfer section **510** to a separator **522**, such as a cyclonic separator, a mesh screen, or a settling vessel. The separator **522** removes the liquid droplets from the gas phase. The liquid droplets may include the original liquid solvent stream with the absorbed impurities **524**, and the gas phase may include a purified gas stream **526**. In various embodiments, the purified gas stream **526** is a gas stream that has been purified via the removal of H₂S and CO₂.

Method for Enhanced Acid Gas Removal within a Gas Processing System

[0112] FIG. 6 is a process flow diagram showing a method **600** for enhanced acid gas removal within a gas processing system. The method **600** is implemented by a gas processing system, such as the gas processing systems **100** and **168** discussed with respect to FIGS. 1A and 1B. In addition, the gas processing system may include one or more separation systems, such as the separation systems **200**, **300**, **400**, and **404** discussed with respect to FIGS. 2, 3, and 4. Further, the gas processing system may include a number of co-current contacting systems. In various embodiments, the co-current contacting systems correspond to the co-current contacting system **500** described with respect to FIG. 5.

[0113] The method begins at block **602**, at which a sour feed gas stream including

an acid gas is contacted with a solvent stream within a number of co-current contacting systems to produce a partially-sweetened gas stream and a rich solvent stream including an absorbed acid gas. At least one of the co-current contacting systems is configured to send the rich solvent stream to a regenerator. In addition, in some
5 embodiments, each of the co-current contacting systems is configured to recirculate a corresponding solvent stream to a preceding one of the co-current contacting systems.

[0114] In various embodiments, the sour feed gas stream is a sour natural gas stream. Moreover, in various embodiments, the sour feed gas stream includes at least 1,000 ppm H₂S. In those embodiments, the absorbed acid gas is primarily H₂S, and
10 the partially-sweetened gas stream is a gas stream that has had a portion of the H₂S removed.

[0115] At block **604**, the absorbed acid gas is removed from the rich solvent stream within the regenerator to produce a lean solvent stream. In some embodiments, the regenerator may correspond to the regenerator **110** described with respect to FIGS.
15 1A and 1B.

[0116] At block **606**, at least a portion of the lean solvent stream is treated within a solvent treater to produce an enhanced solvent stream. The solvent treater may correspond to the solvent treater **164** described with respect to FIGS. 1A and 1B. In some embodiments, the solvent treater is a chiller that produces the enhanced solvent
20 stream by reducing the temperature of the lean solvent stream. In other embodiments, the solvent treater is an anion exchange bed that produces the enhanced solvent stream by removing residual HS⁻, HCO₃⁻, and heat stable salt contaminants from the lean solvent stream. In other embodiments, the solvent treater is an electrodialysis unit that produces the enhanced solvent stream by reducing the lean load of the lean
25 solvent stream and removing heat stable salt contaminants from the lean solvent stream. Further, in other embodiments, the solvent treater produces the enhanced solvent stream by injecting an enhancement fluid into the lean solvent stream that increases its ability to absorb acid gas.

[0117] At block **608**, the partially-sweetened gas stream is contacted with the enhanced solvent stream within a final co-current contacting system to produce a partially-loaded solvent stream and a final gas stream. In various embodiments, the
30 final gas stream includes concentrations of less than 4 ppm H₂S and less than 2-3 vol.

% CO₂.

[0118] The process flow diagram of FIG. 6 is not intended to indicate that the steps of the method **600** are to be executed in any particular order, or that all of the steps of the method **600** are to be included in every case. Further, any number of additional
5 steps not shown in FIG. 6 may be included within the method **600**, depending on the details of the specific implementation. For example, in embodiments in which the solvent treater is a chiller, the method **600** may also include determining an acid gas concentration of the final gas stream using an analyzer, and then increasing or decreasing the temperature of the chiller based on the acid gas concentration of the
10 final gas stream using a controller.

[0119] In some embodiments, the method **600** is implemented using two separate separation systems, such as the first and second separation systems **400** and **404** described with respect to FIG. 4. In those embodiments, the final co-current contacting system may employ intra-stage recycling of its own solvent stream using a separate
15 regenerator. Specifically, the partially-loaded solvent stream exiting the final co-current contacting system may be sent to the regenerator, which may produce a lean solvent stream. The lean solvent stream may then be sent through the solvent treater, which may produce the enhanced solvent stream. The enhanced solvent stream may then be recirculated through the final co-current contacting system to produce the final
20 gas stream.

[0120] While the present techniques may be susceptible to various modifications and alternative forms, the embodiments discussed above have been shown only by way of example. However, it should again be understood that the techniques are not intended to be limited to the particular embodiments disclosed herein. Indeed, the
25 present techniques include all alternatives, modifications, and equivalents falling within the true spirit and scope of the appended claims.

CLAIMS

1. A method for enhanced acid gas removal within a gas processing system, comprising:
contacting a sour feed gas stream comprising an acid gas with a solvent stream within a plurality of co-current contacting systems to produce a partially-sweetened gas stream and a rich solvent stream comprising an absorbed acid gas;
removing the absorbed acid gas from the rich solvent stream within a regenerator to produce a lean solvent stream;
dividing the lean solvent stream into a first portion and a second portion;
combining the first portion of the lean solvent stream with a partially-loaded solvent stream from a final co-current contacting system;
treating the second portion of the lean solvent stream within a solvent treater to produce an enhanced solvent stream; and
contacting the partially-sweetened gas stream with the enhanced solvent stream within the final co-current contacting system to produce the partially-loaded solvent stream and a final gas stream.
2. The method of claim 1, comprising:
recirculating a corresponding solvent stream from each of the plurality of co-current contacting systems to a preceding one of the plurality of co-current contacting systems; and
recirculating the partially-loaded solvent stream from the final co-current contacting system and the first portion of the lean solvent stream to a preceding one of the plurality of co-current contacting systems.
3. The method of claim 1, wherein the solvent treater comprises a chiller configured to reduce a temperature of the second portion of the lean solvent stream to at least about 5°C. below ambient temperature to produce the enhanced solvent stream.
4. The method of claim 3, comprising:
analyzing the final gas stream to determine an acid gas concentration of the final gas stream; and
increasing or decreasing a temperature of the chiller based on the acid gas concentration of the final gas stream.
5. The method of claim 1, wherein contacting the sour feed gas stream with the solvent stream within the plurality of co-current contacting systems comprises:
flowing the solvent stream into a co-current contactor via a mixer through flow channels;
flowing the sour feed gas stream into the co-current contactor via a mixer;
contacting the sour feed gas stream with the solvent stream to provide for incorporation of liquid droplets formed from the solvent stream into the sour feed gas stream; and
separating the liquid droplets from the sour feed gas stream within a separator.
6. The method of claim 1, wherein treating the second portion of the lean solvent stream within the solvent treater comprises injecting an enhancement fluid into the second portion of the lean solvent stream to produce the enhanced solvent stream, wherein the enhancement fluid is different than the solvent and capable of enhancing the ability of the solvent to selectively absorb acid gas.
7. The method of claim 6, wherein the enhancement fluid is a liquid H₂S scavenger.