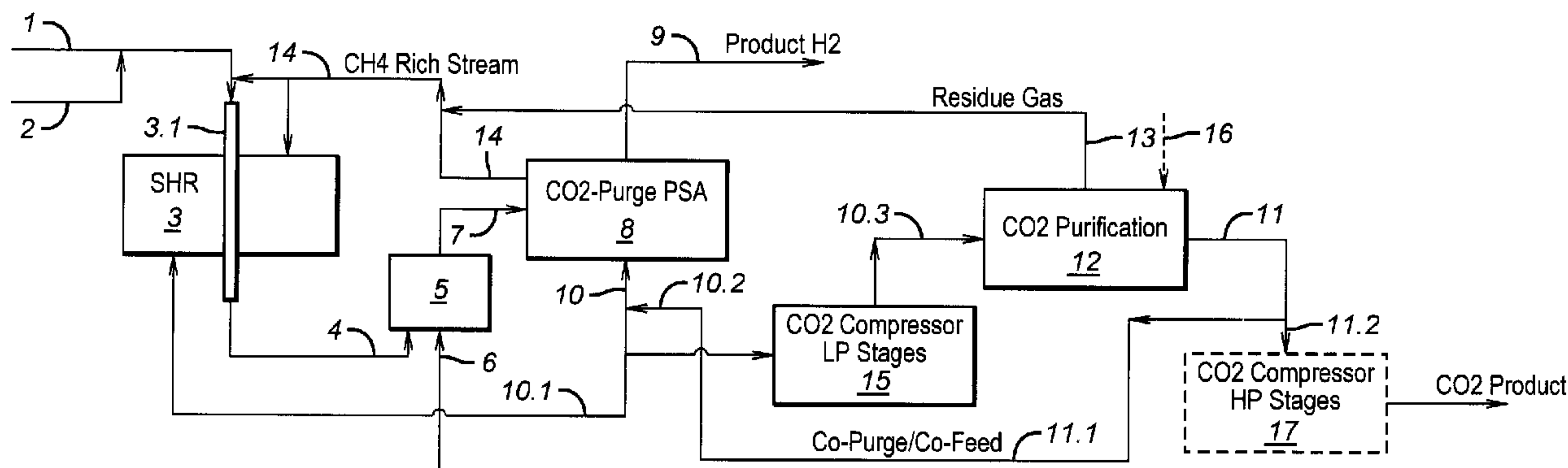




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(54) **Titre :** PROCEDE DE PRODUCTION DE DIOXYDE DE CARBONE AU MOYEN D'UNE UNITE D'ADSORPTION MODULEE EN
 PRESSION A CO-PURGE
 (54) **Title:** PROCESS FOR THE PRODUCTION OF CARBON DIOXIDE UTILIZING A CO-PURGE PRESSURE SWING ADSORPTION
 UNIT



(57) **Abrégé/Abstract:**

The present invention provides a process for recovering gaseous hydrogen and gaseous carbon dioxide from a mixture of hydrocarbons by utilizing a system that includes a reformer unit, an optional water gas shift reactor, and a pressure swing adsorption unit in conjunction with a carbon dioxide purification unit such as a cryogenic purification unit or a catalytic oxidizer. In this process, purified CO₂ from the CO₂ purification unit is used as a co-feed/co-purge in the pressure swing adsorption unit in order to produce a CO₂ tail gas that includes a higher concentration of CO₂.

Abstract

The present invention provides a process for recovering gaseous hydrogen and gaseous carbon dioxide from a mixture of hydrocarbons by utilizing a system that includes a reformer unit, an optional water gas shift reactor, and a pressure swing adsorption unit in conjunction with a carbon dioxide purification unit such as a cryogenic purification unit or a catalytic oxidizer. In this process, purified CO₂ from the CO₂ purification unit is used as a co-feed/co-purge in the pressure swing adsorption unit in order to produce a CO₂ tail gas that includes a higher concentration of CO₂.

PROCESS FOR THE PRODUCTION OF CARBON DIOXIDE UTILIZING A CO-PURGE PRESSURE SWING ADSORPTION UNIT

Field of the Invention

The present invention relates to a process for recovering gaseous hydrogen and gaseous carbon dioxide from a mixture of hydrocarbons by utilizing a system in which a purified carbon dioxide stream produced in a carbon dioxide purification unit is recycled for use as a co-feed to purge the adsorbent beds of a pressure swing adsorption unit used to treat effluent from a reformer unit/water gas shift reactor.

Background

Interest in the recovery of carbon dioxide (hereinafter "CO₂") from various CO₂ containing gas mixtures has increased due to a variety of factors including the merchant CO₂ market, enhanced oil recovery and greenhouse gas emissions reduction. The currently available systems for recovering high purity CO₂ use a variety of generic and proprietary physical and chemical solvents such as conventional pressure swing adsorption units (hereinafter "PSA units") and CO₂ recovery units downstream from the PSA unit such as amine liquid wash units, Selexol liquid wash units or methanol liquid wash units. Accordingly, the processes utilized for this recovery require a large investment due to equipment costs and also high regeneration energy requirements.

Carbon dioxide containing gas mixtures are produced as waste streams during the production of hydrogen gas from hydrocarbon streams using standard steam hydrocarbon reforming processes (hereinafter "SHR"). The most preferred of the SHR processes involves the production of hydrogen gas from hydrocarbon streams using steam methane

reforming (hereinafter “SMR”) processes since methane has a higher proportion of hydrogen than other hydrocarbons. More specifically, with regard to general SMR processes, a hydrocarbon feed gas (natural gas) is fed into a SMR device where the methane in the feed gas reacts with steam at high temperatures (from about 700 °C to about 1100 °C) in the presence of a metal-based catalyst to produce a partially reformed gas that is a mixture of carbon monoxide and hydrogen. The hydrogen yield of this mixture is increased by passing the resulting mixture through a water gas shift reactor which promotes the conversion of carbon monoxide and water into more hydrogen. Accordingly, the result is a reformed gas stream that is rich in hydrogen but also contains to a lesser degree carbon dioxide, methane, and carbon monoxide. Such units typically operate at a temperature from about 200 °C to about 500 °C. In some cases, the stream from the SHR will be at a higher temperature so optionally the stream may first be cooled with a heat exchanger before being passed through the water gas shift. In one conventional process, the reformed gas stream (the hydrogen rich stream) produced is then passed through a H₂ pressure swing adsorption unit (hereinafter “H₂ PSA unit”) in order to allow for the removal of from about 80% to about 90% or more of the hydrogen present through the use of adsorbents. The removal of the hydrogen results in a waste stream (also commonly referred to as a “PSA tail gas stream”) that is purged from the H₂ PSA unit. This PSA tail gas stream contains methane, carbon monoxide, carbon dioxide, water, and any unrecovered hydrogen. This differs from the SHR units, with the difference being that the waste stream or tail gas produced in the SHR units contains alkanes of varying size (C_nH_{2n+2}) and water. The desire has been to be able to utilize these waste streams more efficiently as in the past they have simply been recycled to be burned as make up fuel (added to the natural gas used in the SHR process or SMR process).

Recently, a CO₂ cryogenic process unit (hereinafter “CPU”) process was proposed to capture the CO₂ during steam methane reforming H₂ pressure swing adsorption off gas (by Air Liquide) in WO 2006/054008. In this process, the waste gas from the CPU plant, which normally contains significant amounts of H₂, can be recycled back to the SMR plant for additional H₂ production credit. The process requires operation at high pressure and cold temperature though. Therefore, while it may be appropriate to use the CO₂ CPU process in

a very large scale CO₂ recovery plant (>1000 TPD), when applying the CO₂ CPU process in a small size CO₂ recovery plant (typically 100 to 500 TPD merchant CO₂ plants), the energy and maintenance costs are considered to be usually high.

In an alternative conventional process, there is a CO₂ recovery unit downstream of the H₂ PSA. This CO₂ recovery unit can be a liquid wash unit such as an amine liquid wash unit, a Selexol liquid wash unit or a methanol liquid wash unit or a cryogenic unit. In this schematic, the H₂ PSA tail gas is optionally compressed upstream of the CO₂ recovery unit. The tail gas after CO₂ recovery is then recycled to the PSA unit, recycled as the SMR feed or used as fuel in the SMR furnace. By recycling the tail gas, H₂ recovery is increased.

Even with the above conventional methods, there exists a need to provide a process that allows for a more economical recovery of highly concentrated CO₂ from a pressure swing adsorption process without effecting hydrogen recovery.

Summary of the Invention

The present invention relates to a process for recovering gaseous hydrogen and gaseous carbon dioxide from a mixture of hydrocarbons by utilizing a system that comprises a reformer unit, an optional water gas shift reactor, and a pressure swing adsorption unit in conjunction with a carbon dioxide purification unit (such as a cryogenic purification unit or a catalytic oxidizer unit) in which purified carbon dioxide from the carbon dioxide purification unit is used as a co-feed to purge the adsorbent beds in the pressure swing adsorption unit.

Brief Description of the Figures

Figure 1 provides a schematic of the present invention which includes a SHR unit.

Figure 2 provides an alternative schematic of the present invention which includes an ATR unit.

Figure 3 provides a schematic of two adsorption vessels from a hydrogen pressure swing adsorption unit with each adsorption vessel having two adsorbent beds within one

interior area, wherein within each of the adsorption vessels, the adsorbent beds are separated by an intermediate draw area.

Figure 4 provides a schematic of two parallel adsorption vessels from a hydrogen pressure swing adsorption unit, each of the adsorption vessels having two separated sections separated by a pipe with each section having two adsorbent beds.

Detailed Description of the Invention

By using the purified CO₂ from a purification unit such as a CPU or a catalytic oxidizer unit (hereinafter "CatOx unit") to serve as a co-purge (co-feed) for the adsorbent beds in a H₂ PSA unit after the adsorption step of the H₂ PSA cycle, it is possible to increase the concentration of the CO₂ in the CO₂ rich tail gas coming from the H₂ PSA unit thereby providing a process and system which not only results in the production of a high purity hydrogen (hereinafter "H₂") gas but also a highly concentrated CO₂ gas while at the same time reducing the costs for the production of the same due to less downstream treatment (purification) being required.

In the process of the present invention, it is possible to recover gaseous H₂ and gaseous CO₂ from a mixture of hydrocarbons utilizing a reformer unit in conjunction with an optional water gas shift reactor (hereinafter "WGS reactor"), a co-purge H₂ PSA unit and CO₂ purification unit (such as a CPU or CatOx unit). This is achieved by utilizing the purified CO₂ product from the CO₂ purification unit as a co-feed or co-purge in the H₂ PSA process. The proposed processes of the present invention include a variety of embodiments for achieving this result, some of which include: 1) the use of a SHR unit, a WGS reactor, a co-purge H₂ PSA unit and a CPU or 2) the use of a oxygen fed autothermal reformer (hereinafter "ATR") unit, a WGS reactor, a co-purge H₂ PSA unit and a CPU or 3) the use of a SHR unit, a WGS reactor, a co-purge H₂ PSA unit and a CatOx unit or 4) the use of an ATR unit, a WGS unit, a co-purge H₂ PSA unit and a CatOx unit, in order to increase the concentration of the CO₂ in the tail gas stream produced from the H₂ PSA unit and accordingly provide for additional use of the CO₂ that would normally be used for other purposes such as makeup fuel in the reformer unit.

The overall processes of the present invention involve recovering high purity gaseous hydrogen and highly concentrated gaseous CO₂. As used herein, the phrase “highly concentrated gaseous CO₂” refers to the tail gas stream that is the product of the process of the present invention in which a CO₂ stream from the CO₂ purification unit is used as a co-feed to purge the adsorbents beds of a H₂ PSA unit, said highly concentrated gaseous CO₂ having a CO₂ content that is greater than about 75 mol% carbon dioxide, preferably from about 85 mol% carbon dioxide to about 99 mol % carbon dioxide.

The first stages of this process, as shown in Figures 1 and 2, involve the generation of a hydrogen rich effluent stream by the treatment of one or more hydrocarbon streams (also referred to as one or more hydrocarbon feed gases) provided via line **1** in a reformer unit **3** (**3A** in Figure 2). As used herein, the phrase “reformer unit” is meant to encompass those processes which are utilized to produce a synthesis gas. In the preferred embodiments of the present invention, the reformer unit **3** will be selected from a SHR unit **3** as depicted in Figure 1 and an oxygen fed ATR unit **3A** as depicted in Figure 2.

In the first embodiment as shown in Figure 1, the SHR unit **3** is used. The present invention is not meant to be limited by the SHR unit **3** or the process for carrying out the reaction in the SHR unit **3**. Accordingly, any SHR unit **3** known in the art may be used in the process of the present invention. By way of general description, such SHR units **3** typically contain tubes **3.1** (only one shown in Figure 1) packed with a SHR catalyst (typically a nickel catalyst) through which the steam **2** /hydrocarbon stream **1** mixture passes. As used throughout with regard to the present invention, the phrase “steam hydrocarbon reformer unit” or “SHR unit” refers not only to the actual reformer units, but also to all of the additional components (not shown) that typically are considered to make up a steam hydrocarbon reformer, including, but not limited to, one or more components selected from heat exchangers, pre-reformer units, the reformer, tubes with one or more types of catalyst, etc. The reaction product from the steam hydrocarbon reformer unit (hereinafter “SHR product stream”) is principally a hydrogen rich effluent that contains H₂, carbon monoxide (hereinafter “CO”), methane (hereinafter “CH₄”), water vapor and

CO₂ in proportions close to equilibrium amounts at the elevated temperature and pressure. When the SHR unit is utilized, it is preferably a SMR unit.

With regard to this first embodiment, in certain situations, especially where there is no need to have CO as a product, the scheme may optionally contain a WGS reactor **5** which functions to react CO and water to form H₂ and CO₂ and obtain a WGS effluent. In the preferred embodiment where the WGS reactor **5** is included, the SHR product stream is introduced via line **4** into a WGS reactor **5** (which can contain a variety of stages or one stage; stages not shown) along with steam which is introduced via line **6** to form additional H₂ and CO₂. The combination of a SHR unit **3** and a WGS reactor **5** is well known to those skilled in the art.

In an alternative embodiment of the present invention as shown in Figure 2, the reformer unit will be an oxygen fed ATR unit **3A**. With regard to the ATR unit **3A**, combustion of the one or more hydrocarbon streams provided via line **1** is carried out with steam provided via line **2** and substoichiometric amounts of oxygen provided via line **2A** by flame reactions in a burner combustion zone and, subsequently, autothermal reforming of the partially combusted feedstock. The present invention is not meant to be limited by the ATR unit **3A** or the process for carrying out the reaction in the ATR unit **3A**. Accordingly, any ATR unit **3A** known in the art may be used in the process of the present invention. In addition, as used herein, the phrase “autothermal reformer unit” or “ATR unit” refers not only to the actual reformer units, but also to all of the additional components (not shown) that typically are considered to make up an ATR unit, including, but not limited to, one or more components selected from heat exchangers, pre-reformer units, the reformer, beds with one or more types of catalyst, etc.

As with the SHR unit, in certain situations when there is no need to have CO as a product, the scheme may optionally include a WGS reactor **5** which functions to react CO and water to form additional H₂ and CO₂ by further reacting or treating the hydrogen rich effluent provided to the WGS reactor **5** via line **4** from the ATR unit **3A** with steam

supplied via line **6** in order to obtain a WGS effluent. The combination of an ATR unit **3A** and a WGS reactor **5** is known to those skilled in the art.

With regard to each of the embodiments discussed above, a H₂ rich effluent is produced in the corresponding reformer unit **3**, **3A** that contains in addition to H₂, other components such as CO, CO₂, CH₄ and water vapor. Preferably, this H₂ rich effluent is optionally further treated in the WGS reactor **5** in order to further enrich the H₂ content of the H₂ rich effluent and to also increase the CO₂ content in the H₂ rich effluent by oxidizing a portion of the CO present in the effluent to CO₂ thereby obtaining a WGS effluent. For purposes of the present discussion, reference will be made to those embodiments which include the WGS reactor **5**. However, those skilled in the art will recognize that the effluent to be introduced into the H₂ PSA unit **8** may simply be taken from the reformer unit without passing through a WGS reactor **5**. Both embodiments are considered to be within the scope of the present invention.

With regard to each of these embodiments, once the WGS effluent is obtained, this effluent is introduced into a co-purge H₂ PSA unit **8** via line **7** in order to produce high purity H₂. Prior to introduction into the co-purge H₂ PSA unit **8**, if necessary, the WGS effluent will typically be cooled down to less than about 50 °C and to a pressure that allows for the adsorption step of the H₂ PSA cycle to be run at a pressure from about 250 psig to about 700 psig. The cooling down step is typically accomplished via a heat exchanger (not shown). Typically the H₂ rich effluent and the WGS effluent utilized are at pressure or may be pressurized via a compressor (not shown). As used herein, the phrase “co-purge H₂ PSA unit” or “co-feed H₂ PSA unit” refers to a conventional H₂ PSA unit which allows for the introduction of an additional gas stream (co-purge or co-feed stream) during the purge step of the H₂ PSA cycle. The H₂ PSA unit **8** utilized can be any H₂ PSA unit known in the art that comprises two or more adsorption vessels **18** as shown in Figures 3 to 4, each of the adsorption vessels **18** independently having anywhere from two to six adsorbent beds **19** although more than six adsorbent beds may be utilized. Typically, the H₂ PSA unit **8** will have anywhere from two to sixteen, preferably from four to sixteen, adsorption vessels **18** that are parallel with regard to one

another. The actual number of adsorption vessels **18** present will typically depend upon the size of the facility where the H₂ PSA unit **8** is located and the quantity of gas to be treated. Each of the adsorption vessels **18** will typically contain multiple adsorbent beds **19** which are also parallel to one another and in certain embodiments; two or more of the multiple adsorbent beds **19** may be adjacent to one another. A variety of configurations with regard to the adsorption vessels **18** and the adsorbent beds **19** as shown in Figures 3 to 4 may be utilized in the present invention.

During the process of H₂ production, each of the adsorption vessels **18** will individually undergo a cycle that includes an adsorption step, a co-feed step, a depressurization step, a regeneration step, and a re-pressurization step. With regard to the adsorption vessels **18** of the H₂ PSA units **8** that are in use at the same time, the cycles in these various adsorption vessels **18** will run consecutively. However, while the cycles run consecutively, they do not all run such that each of the adsorption vessels **18** is within the same cycle step at the same time. Instead, at least one of the adsorption vessels **18** of the total number of adsorption vessels **18** being utilized within the co-purge H₂ PSA unit **8** will be in a different step of the cycle when compared to the step of at least one of the remaining adsorption vessels **18** that is being utilized (the cycles are staggered with regard to at least two of the adsorption vessels **18**).

As noted above, the actual adsorption vessels **18** and adsorbent beds **19** can take on a variety of different configurations as shown in Figures 3 to 4. In one such configuration as shown in Figure 3, the co-purge H₂ PSA unit **8** has two or more adsorption vessels **18** that are parallel to one another (note only two adsorption vessels are depicted in Figure 3). Each adsorption vessel **18** comprises one internal space **20** into which multiple adsorbent beds **19** are placed (in this schematic two adsorbent beds **19**).

During the co-purge step of the H₂ PSA cycle, a purified CO₂ stream is fed into the adsorption vessel **18** and consequently the adsorption beds **19** via line **11.1**. As the purified CO₂ stream enters the adsorbent beds **19** of the adsorption vessel **18**, the CO₂ from the purified CO₂ stream begins to displace the components (such as CH₄ and CO)

that adhered to the adsorbents of the adsorbent beds **19** during the adsorption step of the H₂ PSA cycle (when the WGS effluent stream was brought into contact with the adsorbent beds **19**). As a result, within this particular step of the H₂ PSA cycle (the co-purge step), a CH₄ rich stream is formed from the components that are displaced by the CO₂. As a result, this CH₄ rich stream is withdrawn from the intermediate draw area **24** with the aid of a valve **25** that is associated with line **14**. This allows for the withdrawal of a stream from the first adsorbent bed **19.1** during the co-purge step of the H₂ PSA cycle before the stream goes through the second adsorbent bed **19.2**. Furthermore, with regard to this particular alternative, the adsorbent beds **19** may still be considered to be “stacked” in that one of the adsorbent beds **19.2** is considered in an upper position in relation to the other adsorbent bed **19.1** which would then be considered to be in a lower position in relation to the first adsorbent bed **19.2**. Accordingly, in this particular alternative, in the H₂ PSA cycle, the adsorbent beds are in full communication with one another. Note that the CH₄ rich stream that is withdrawn can be recycled to the hydrocarbon feed stream **1** to supplement the hydrocarbon feed or can be directly injected into the reformer unit (see Figures 1 and 2).

As noted in Figure 3, the adsorbent beds **19** are arranged within an adsorption vessel **18** such that they are separated from one another by the intermediate draw area **24**. Accordingly, as shown in Figure 3, in the co-purge H₂ PSA unit **8**, an actual space (intermediate draw area **24**) in which there is no adsorbent beds **19** present extends between the two adsorbent beds **19** that are present in that particular adsorption vessel **18**. In this alternative, only two adsorbent beds **19** are shown within each adsorption vessel **18**. While this embodiment is referenced with regard to two adsorbent beds **19**, those skilled in the art will recognize that this embodiment may be carried out with regard to any number of adsorbent beds **19**. Accordingly, with regard to an adsorption vessel **18** having greater than two adsorbent beds **19**, it is still possible to withdraw a stream from the area between any two adsorbent beds **19** using a valve **25** located along line **14** which extends from the intermediate draw area **24**. More specifically, in the case where there are more than two adsorbent beds **19** present, there may be an intermediate draw area **24** present between each pair of adsorbent beds **19**; between only one pair of adsorbent beds

19 with the remaining adsorbent beds **19** being adjacent to one another, or multiple intermediate draw areas **24** dispersed among the multiple adsorbent beds **19** with there being some adsorbent beds **19** that are adjacent to one another and others that are separated by an intermediate draw area **24**.

With regard to the adsorbent beds **19** of Figure 3, it would be possible to have different adsorbents in the one or more adsorbent beds **19**. For example, the first adsorbent bed **19.1** (or adsorbent beds **19** when more than one adsorbent bed **19** is present on one side of the intermediate draw area **24**) can include one or more adsorbents that are selective for the removal of CH₄, CO, and nitrogen (hereinafter "N₂") while the second adsorbent bed **19.2** (or adsorbent beds **19** when more than one adsorbent bed **19** is present on the other side of the intermediate draw area **24**) can contain one or more adsorbent beds **19** that each contain one or more adsorbents that allow for the removal of CO₂ (one or more adsorbents that are selective for this component).

In another configuration as shown in Figure 4, the H₂ PSA unit **8** has two or more adsorption vessels **18** that are parallel to one another with each of these adsorption vessels **18** containing two or more adsorbent beds **19**. Unlike the previous embodiment set forth in Figure 3 where the adsorbent beds **19** are stacked in relation to one another, and are separated by an intermediate draw area **24** but are still all contained within the same interior space **20** of the adsorption vessel **18** of the H₂ PSA unit **8**, in this alternative embodiment, the adsorption vessel **18** comprises a split adsorption vessel **18** (two or more separated sections in series **18a** and **18b**) wherein each section **18a**, **18b** of the adsorption vessel **18** includes one or more adsorbent beds **19**. The manner in which the adsorption vessel **18** is split is to allow the sections **18a**, **18b** to be partitioned from one another to optionally limit the flow of any gas introduced into the one or more adsorbent beds **19** of one section **18b** of the adsorption vessel **18** with regard to the one or more of the adsorbent beds **19** of the other section **18a** of the adsorption vessel **18**. In other words, the adsorbent beds **19** of the various sections in essence function as separate entities with regard to one another by residing in separated sections **18a**, **18b** within the same general adsorption vessel **18**. Accordingly, as shown in Figure 4, in the co-purge H₂ PSA unit **8**,

a pipe **21** is present that extends between the two adsorption sections **18a, 18b** and serves to “connect” the adsorption sections **18a, 18b** for the purpose of flow between the two separate individually contained adsorption sections **18a, 18b**. This pipe **21** allows for the isolation or separation of the adsorption sections **18a, 18b** with regard to one another. This pipe **21**, through the use of a withdrawal valve **22** located on line **14**, in turn also allows for the withdrawal of a CH₄ rich stream between the two adsorption vessel sections **18a, 18b** during the co-purge step of the H₂ PSA cycle. In essence, this pipe **21** functions much in much the same manner as the intermediate draw area of Figure 3 but with more control over the flow between the two adsorption sections **18a, 18b** (two adsorbent beds **19** in Figure 3). Furthermore, with regard to this particular alternative embodiment, the adsorbent beds **19** of the adsorption vessel sections **18a, 18b** may still be considered to be “stacked” in that one of the adsorption vessel sections **18b** is considered in an upper position in relation to the other adsorption vessel section **18a** which would then be considered to be in a lower position in relation to the other adsorption vessel section **18b**. With this configuration of an upper and lower position, the adsorbent beds **19** are also considered to be in a stacked position with regard to one another as the one or more adsorbent beds **19** of the upper adsorption vessel section **18b** are now considered to be in the upper position with regard to the one or more adsorbent beds **19** of the lower adsorption vessel section **18a**.

While in some PSA cycle steps it may be desirable to run the H₂ PSA cycle with the pipe **21** fully communicating between the two adsorption vessel sections **18a, 18b**, in other instances, it would be desirable to have a means for isolating or separating **23** (closing off the communication) one adsorption vessel section **18a** with regard to the other adsorbent bed section **18b**. In such an instance, it would be possible to have different adsorbents in the one or more adsorbent beds **19** of the different adsorption vessel sections **18a, 18b**. For example, the lower adsorption vessel section **18a** can contain one or more adsorbent beds **19** that each include one or more adsorbents that are selective for CO₂ while the upper adsorption vessel section **18b** can contain one or more adsorbent beds **19** that each contain one or more adsorbents that allow for the removal of CH₄, CO, and nitrogen (hereinafter “N₂”) (one or more adsorbents that are selective for

these components). Accordingly, with such a pipe **21** in place that connects the two adsorption vessel sections **18a, 18b** and a means for isolating or separating **23** the two adsorption vessel sections **18a, 18b** from one another during the depressurization step of the H₂ PSA cycle, it would be possible to prevent the CH₄, CO and N₂ adsorbed to the adsorbent in the upper adsorption vessel section **18b** from mixing with the CO₂ adsorbed to the adsorbent of the lower adsorption vessel section **18a** or the CO₂ from flowing up into the upper adsorption vessel section **18b**. In a still further alternative, each of the adsorption vessel sections **18a, 18b** that are separated from one another may have greater than one adsorbent bed **19** within that particular section **18a, 18b** with each of the adsorbent beds **19** containing a different adsorbent from the other adsorbent bed **19** contained in that particular section **18a, 18b**. The means for isolating or separating **23** the two adsorption vessel sections **18a, 18b** will typically comprise one or more valves **23** located along the pipe **21** that connects the two adsorption vessel sections **18a, 18b**. The CH₄ stream withdrawn during the co-purge step can be withdrawn via line **14** through the use of valve **22** and recycled to line **1** for use to supplement the hydrocarbon feed stream.

An even further alternative to Figure 4 is possible in that the two or more beds in the various individual sections **18a** or **18b** of the adsorption vessel **18** may be separated from one another by an intermediate draw area **24** as shown within Figure 3.

With regard to the various embodiments of the present invention, preferably at least two types of adsorbents are used in the adsorbent beds **19** of the co-purge H₂ PSA unit **8**. These adsorbents include, but are not limited to, activated alumina, silica gel, activated carbon, zeolites, and combinations thereof. In one preferred embodiment of the present invention, the co-purge H₂ PSA unit **8** comprises multiple adsorbent beds **19** (most preferably two to four) that are stacked in relation to one another and in which different adsorbents are used. Accordingly, for example as shown in Figure 3, in the embodiment where there are two adsorbent beds **19**, one adsorbent bed **19** may be referred to as the first adsorbent bed **19.1** and contain one or more adsorbents that are selective for CH₄ and CO, while the other adsorbent bed **19** may be referred to as the

second adsorbent bed **19.2** and contain one or more adsorbents that are selective for CO₂ and water vapor. In a still further embodiment, the adsorption vessel **18** can comprise a split adsorption vessel as shown in Figure 4, for example, a split adsorption vessel having a first section **18a** and a second section **18b**, wherein the first section **18a** contains two or more adsorbent beds **19**, each bed **19** containing one or more adsorbents that are selective for CH₄ and CO, and the second section **18b** containing two or more adsorbent beds **19**, each adsorbents selective for CO₂ and water vapor.

As a result of the co-purge H₂ PSA cycle within the adsorption vessel **18**, as shown in Figures 1 and 2, three separate gas streams are obtained as a result of the co-purge H₂ PSA process. More specifically, in the embodiments depicted in Figures 3 and 4 where the adsorbent beds **19** are separated due to the adsorbent beds **19** being divided into sections **18a**, **18b** or due to an intermediate draw area **24** being positioned between the adsorbent beds **19**, three separate gas streams are obtained: one that is a gaseous high purity H₂ stream that is withdrawn via line **9**, a PSA tail gas which is rich in CO₂ and is withdrawn via line **10** after desorption of the two or more adsorbent beds **19** in the various adsorption vessels **18** and a CH₄ rich stream that is withdrawn via line **14**.

With regard to the PSA tail gas, a portion of this gas stream may be passed back to the CO₂ purification unit **12** via line **10.2** and line **10.3** or it may be passed back to the SHR unit **3** as shown in Figure 1 (or to the ATR unit **3A** as shown in Figure 2) via line **10.1**. The CH₄ rich stream **14** is recycled back to the SHR or the ATR. In the cycle, the WGS effluent is introduced into the H₂ PSA vessel via line **7** where it then flows over the adsorbent beds **19** (not shown). The H₂ contained in the WGS effluent (which is also a H₂ rich effluent) passes over the adsorbents and is capable of being withdrawn via line **9** during the adsorption step of the H₂ PSA cycle where it is passed on for further use as H₂ product. The other components from the WGS effluent make up the CH₄ rich stream and PSA tail gas and are adsorbed to the adsorbents of the adsorbent beds **19** during the adsorption step of the H₂ PSA cycle. Accordingly, these components are still within the adsorbent beds **19** of the H₂ PSA adsorption vessel **18** once the introduction of the WGS effluent flow is stopped. The H₂ PSA cycle is carried out using conventional parameters.

With regard to the present embodiments, the pressure during the adsorption steps will generally range from about 250 psig to about 700 psig.

As noted, the H₂ PSA cycle of the present process also includes a co-purge (co-feed) step in which a CO₂ stream is used to aid in purging the adsorbent beds **19**. The purpose of this co-purge step is to provide an economically efficient manner to increase the concentration of the CO₂ in the H₂ PSA tail gas. According to the present processes as noted above, after the adsorption step, the injection of the WSG effluent is stopped and a purified CO₂ stream that is obtained from the CO₂ purification unit **12** via line **11** downstream from the H₂ PSA unit **8** is then injected via lines **11.1** and **10** into the adsorption vessel **18** of the H₂ PSA unit **8** which is in the co-purge step of the H₂ PSA cycle and allowed to pass over the adsorbent beds **19**. In the preferred embodiment, the purified CO₂ stream is introduced into the co-purge step of the H₂ PSA cycle at a pressure that is higher than the pressure during the adsorption step of the H₂ PSA cycle. Preferably, the pressure during the co-purge step of the H₂ PSA cycle is in the range of from about 300 psig to about 800 psig. An optional CO₂ compressor **15** having multiple compression stages may be used to achieve this range if necessary.

As noted, the purified CO₂ stream is obtained from a CO₂ purification unit **12** via line **11**. In one embodiment of the present invention, the CO₂ purification unit **12** is a CPU unit (specifics not shown). In an alternative embodiment, the CO₂ purification unit is a CatOx unit (specifics not shown). Due to the degree of affinity of the various components in the WGS effluent for the adsorbents in the adsorbent beds **19** of the H₂ PSA unit **8**, the CO₂ that is injected from the purified CO₂ stream (the co-purge) begins to displace the CH₄ and CO that is adsorbed on the adsorbents in the adsorbent beds **19** during the adsorption step of the H₂ PSA cycle. The result of this is the displacement of these other components (CH₄ and CO) by CO₂ on the adsorbents. Consequently, there is a larger concentration of CO₂ that is adsorbed on the adsorbents. CH₄ and CO displaced from the adsorbent and void spaces in the adsorbent is removed as a gas stream that is CH₄ rich but also contains amounts of CO and H₂. As the purified CO₂ stream is injected into the H₂ PSA vessel **8**, it is also possible to remove the resulting CH₄ rich stream from

the H₂ PSA vessel via line 14. The CH₄ rich stream can be recycled to the reformer feed 1, injected directly into the SHR unit 3 (or in the second embodiment the ATR unit 3A), or used in any other manner known in the art for the use of such streams.

With further reference to various Figures, the PSA tail gas is withdrawn from the adsorbent beds 19 of the adsorption vessels 18 of the H₂ PSA unit 8 via line 10 during the depressurization step of the H₂ PSA cycle. By depressurizing the adsorbent beds 19 of the at least one H₂ PSA adsorption vessel 18, it is possible to release the CO₂, and CO, CH₄, water vapor and remaining H₂ adsorbed to or on the adsorbents, or held in the void spaces in the adsorbent, and produce a CO₂ rich pressure swing adsorption tail gas during the depressurization step of the H₂ PSA cycle. The CO₂ recovered during the depressurization step can be collected at two or more different pressure levels. In one embodiment, the suction pressure of the various stages of the CO₂ compressors 15 are the same as the pressure levels in the depressurization step of the H₂ PSA cycle. Once the adsorbent beds 19 are depressurized and the CO₂ rich PSA tail gas is withdrawn via line 10, the adsorbent beds 19 of the adsorption vessel 18 may be further regenerated and repressurized utilizing procedures that are readily known in the art.

Once the CO₂ rich H₂ PSA tail gas is withdrawn from the H₂ PSA unit 8 via line 10, the CO₂ rich PSA tail gas can optionally be compressed in a compressor unit 15 having various pressure stages to a pressure that ranges from about 250 psig to about 1000 psig in order to obtain a compressed CO₂ rich H₂ PSA tail gas. Compressor units 15 such as the one utilized in the present invention are known by those skilled in the art and include compressor units 15 that have a variety of compression stages thereby allowing for staged compression. After the CO₂ rich H₂ PSA tail gas is compressed to the desired level, the compressed CO₂ rich H₂ PSA tail gas is then introduced into a CO₂ purification unit via line 10.3 that is either a CPU (a cryogenic purification unit) such as those known in the art or a CatOx unit such as those known in the art.

In the embodiment that utilizes the CPU, the CO₂ in the compressed CO₂ rich H₂ PSA tail gas is condensed at a temperature between ambient temperature and -56 °C to

provide a liquid stream and a gas stream. The condensation may be in one stage or in various stages. Such cryogenic purification units and associated processes are readily known to those skilled in the art. As a result of this condensation, the CO₂ liquefies and the remaining components (the incondensibles) remain in a gaseous state. The incondensibles stream typically contains CH₄, CO, and H₂. Following the condensing step, the liquid CO₂ stream is separated from the incondensibles stream. The liquid CO₂ is vaporized to recover refrigeration, and optionally compressed further to 2000 psig in CO₂ Compressor HP Stages 17. The incondensibles stream is then passed along for further use according to the prior art.

As noted above, at least a portion of the CO₂ produced will be recycled via line 11.1 for use in the H₂ PSA unit 8 as a co-purge (co-feed) during the co-purge step of the H₂ PSA cycle. The remaining portion of the CO₂ produced may be withdrawn via line 11.2 and used according to the various uses known in the art. When the purified CO₂ stream is recycled to the H₂ PSA unit 8, the purified CO₂ stream will be introduced into the H₂ PSA adsorption vessel and allowed to pass over the various adsorbent beds during the co-purge step in order to increase the concentration of CO₂ in the H₂ PSA tail gas.

In the alternative embodiment, the CO₂ purification unit 12 is a catalytic oxidation (CatOx) unit which functions to burn off the light ends (the CH₄ and CO) thereby leaving a stream that contains basically CO₂ and water. In this embodiment, after the CO₂ rich H₂ PSA tail gas is compressed in the compressor unit 15, the compressed CO₂ rich H₂ PSA tail gas is introduced into a CatOx unit such as those that are known in the art. Following the introduction of the CO₂ rich H₂ PSA tail gas, gaseous oxygen (hereinafter "O₂") is then introduced into the CatOx unit via line 16. With the aid of a combustion catalyst, the CO, CH₄, and H₂ that is present in the CO₂ rich H₂ PSA tail gas is combusted to produce an oxidized stream that contains CO₂ and water. The amount of O₂ utilized in the process will typically be slightly below the stoichiometric requirement. Once the combustion takes place, the resulting oxidized stream of CO₂ and water is condensed in order to allow for the removal of water by cooling the oxidized stream to ambient temperature to produce a purified CO₂ stream. This purified CO₂ stream is withdrawn

via line **11** and at least a portion of the purified stream is recycled to the H₂ PSA unit **8** via lines **11.1** and **10** where the stream is introduced into the adsorbent beds **19** of the adsorption vessels **18** during the co-purge step in order to increase the concentration of CO₂ in the H₂ PSA tail gas. A portion **11.2** of purified CO₂ stream **11** is optionally compressed to 2000 psig in CO₂ Compressor HP Stages **17** and passed on as CO₂ product.

By utilizing a purified CO₂ stream as a co-purge during the co-purge step of the PSA process cycle, it is possible to obtain a H₂ PSA tail gas that contains a considerably higher concentration of CO₂.

Preferred processes and apparatus for practicing the present invention have been described. It will be understood and readily apparent to the skilled artisan that many changes and modifications may be made to the above-described embodiments without departing from the spirit and the scope of the present invention. The foregoing is illustrative only and that other embodiments of the integrated processes and apparatus may be employed without departing from the true scope of the invention defined in the following claims.

Elements of the Figures:

- 1 - line to provide hydrocarbon stream to reformer unit (reformer feed)
- 2 - line that provides steam
- 2A – line that provides oxygen
- 3 - steam hydrocarbon reformer unit
- 3.1 - catalyst holding tubes for reformer unit
- 3A - autothermal reformer unit
- 4 - line that provides reformer product stream to the water gas shift reactor
- 5 - water gas shift reactor
- 6 - line that introduces steam into the water gas shift reactor
- 7 - line that introduces water gas shift effluent into the co-purge hydrogen pressure swing adsorption unit

- 8 - co-purge hydrogen swing adsorption unit
- 9 - line by which high purity hydrogen is withdrawn from the co-purge hydrogen pressure swing adsorption unit
- 10 - line through which the pressure swing adsorption tail gas is withdrawn
- 10.1 - line by which a portion of the hydrogen pressure swing adsorption tail gas is passed to the reformer unit or ATR
- 10.2 - line by which a portion of the hydrogen pressure swing adsorption tail gas is passed to a carbon dioxide compressor (LP stages)
- 10.3 - line by which the compressed carbon dioxide rich hydrogen pressure swing adsorption tail gas is introduced into the carbon dioxide purification unit
- 11 - line by which a purified stream of carbon dioxide is withdrawn from the carbon dioxide purification unit
- 11.1 - line by which the purified stream of carbon dioxide is injected into the hydrogen pressure swing adsorption unit vessel as a co-purge stream
- 11.2 - line by which carbon dioxide product is withdrawn from the from the carbon dioxide purification unit and passed to an optional carbon dioxide compressor (HP stages)
- 12 - carbon dioxide purification unit
- 13 - line from which residue gas is withdrawn from the carbon dioxide purification unit
- 14 - line by which a methane rich stream is withdrawn from the hydrogen pressure swing adsorption unit
- 15 - carbon dioxide compressor (LP stages)
- 16 - line by which gaseous oxygen is introduced into the CO₂ purification unit (CatOx unit)
- 17 - Optional CO₂ compressor (HP stages)
- 18 - adsorption vessel of the H₂ PSA unit
- 18a - first section of the adsorption vessel (also referred to as lower)
- 18b - second section of the adsorption vessel (also referred to as upper)
- 19 - adsorbent bed
- 19.1 – first adsorbent bed
- 19.2 – second adsorbent bed

20 - internal space of adsorption vessel

21 - pipe

22 - withdrawal valve

23 - means of isolating and separating the adsorbent beds in the upper and lower section of the adsorption vessel

24 – intermediate draw area

What is claimed is:

1. A process for recovering gaseous hydrogen and gaseous carbon dioxide from a mixture of hydrocarbons utilizing a reformer unit in conjunction with a pressure swing adsorption unit and a cryogenic purification unit, the process comprising the steps of:

a) generating a hydrogen rich effluent that also contains at least carbon monoxide, carbon dioxide, methane and water vapor from one or more hydrocarbon feed streams in a reformer unit;

b) optionally treating the hydrogen rich effluent in a water gas shift reactor in order to further enrich the hydrogen content of the hydrogen rich effluent and to also increase the carbon dioxide content in the hydrogen rich effluent by oxidizing a portion of the carbon monoxide present in the effluent to carbon dioxide thereby obtaining a water shift effluent;

c) providing a pressure swing adsorption unit having at least two adsorption vessels that each contain two or more adsorbent beds, the at least two adsorption vessels being parallel to one another and being independently operated on a cycle that includes an adsorption step, a co-feed step, a depressurization step, a regeneration step and a re-pressurization step, with the proviso that the cycles in the at least two adsorption vessels run consecutively with at least one of the adsorption vessels being in a different step of the cycle compared to at least one of the remaining adsorption vessels and the two or more adsorbent beds having an intermediate draw area positioned between at least two of the adsorbent beds thereby allowing for the withdrawal of a methane rich gas stream from the intermediate draw area during the co-feed step of the hydrogen pressure swing adsorption cycle;

d) introducing the hydrogen rich effluent or water gas shift effluent into at least one of the adsorption vessels of the pressure swing adsorption unit during the adsorption step of the hydrogen pressure swing adsorption cycle in order to allow for the recovery of a high purity gaseous hydrogen stream from the hydrogen rich effluent or the water gas shift effluent and the adsorption of the carbon monoxide, carbon dioxide, methane, water vapor and remaining hydrogen onto the adsorbents in the

adsorbent beds of the pressure swing adsorption unit, the adsorption step being carried out at a pressure from about 250 psig to about 700 psig;

e) withdrawing the high purity gaseous hydrogen stream from the pressure swing adsorption unit during the adsorption step and passing this high purity gaseous hydrogen stream on for further use;

f) allowing the at least one adsorption vessel to enter into the co-feed step while introducing a purified carbon dioxide stream obtained from a downstream cryogenic purification unit in order to allow the carbon dioxide to displace methane and carbon monoxide adsorbed onto the adsorbents in the adsorbent beds thereby producing a methane rich stream;

g) withdrawing the methane rich stream from the intermediate draw area positioned between at least two of the adsorbent beds and recycling this methane rich stream to the reformer unit;

h) allowing the at least one adsorption vessel to enter the depressurization step by depressurizing the adsorbent beds of the at least one adsorption vessel in order to release the carbon dioxide, carbon monoxide, methane, water vapor and hydrogen adsorbed to the adsorbents in the adsorbent beds thereby producing a carbon dioxide rich pressure swing adsorption tail gas having an increased concentration of carbon dioxide;

i) withdrawing the carbon dioxide rich pressure swing adsorption tail gas from the pressure swing adsorption unit;

j) regenerating and re-pressurizing the adsorbent beds for further use;

k) compressing the carbon dioxide rich pressure swing adsorption tail gas in a compressor unit to a pressure that ranges from about 250 psig to about 1000 psig in order to obtain a compressed carbon dioxide rich pressure swing adsorption tail gas;

l) introducing the compressed carbon dioxide rich pressure swing adsorption tail gas into the cryogenic purification unit in order to produce the carbon dioxide purified stream and an incondensibles stream that contains methane, carbon monoxide and hydrogen; and

m) withdrawing the carbon dioxide purified stream from the cryogenic purification unit and recycling at least a portion of the carbon dioxide purified stream to the pressure

swing adsorption unit where the carbon dioxide purified stream is introduced into the adsorption vessels during the co-feed step of the cycle in order to increase the concentration of carbon dioxide in the hydrogen pressure swing adsorption tail gas.

2. The process of claim 1, wherein the reformer unit is selected from a steam hydrocarbon reformer unit and an oxygen fed autothermal reformer unit.

3. The process of claim 2, wherein the adsorption vessel contains four adsorbent beds with two of the adsorbent beds being adjacent to one another on one side of the intermediate draw area and the remaining two adsorbent beds being positioned on the other side of the intermediate draw area.

4. The process of claim 2, wherein the adsorption vessel comprises a first adsorbent bed that is separated from a second adsorbent bed by the intermediate draw area.

5. The process of claim 4, wherein the first adsorbent bed and the second adsorbent bed each independently comprise one or more adsorbents selected from activated alumina, silica gel, activated carbon, zeolites, and combinations thereof.

6. The process of claim 5, wherein the one or more adsorbents are in layers within the individual adsorbent beds.

7. The process of claim 4, wherein the first adsorbent bed contains one or more adsorbents that are selective for methane and carbon monoxide and the second adsorbent bed contains one or more adsorbents that are selective for carbon dioxide and water vapor.

8. The process of claim 7, wherein the one or more adsorbents are in layers within the individual beds.

9. The process of claim 7, wherein when the hydrogen rich effluent or the water gas shift effluent is introduced into the adsorption vessel during the adsorption step of the hydrogen pressure swing adsorption cycle, the hydrogen rich effluent or the water gas shift effluent comes into contact with the first adsorbent bed before it comes into contact with the second adsorbent bed.

10. The process of claim 2, wherein the carbon dioxide purified stream is introduced into the co-feed step of the pressure swing adsorption cycle at a pressure that is higher than the pressure during the adsorption step of the pressure swing adsorption cycle.

11. The process of claim 10, wherein the pressure during the co-feed step of the pressure swing adsorption cycle is from about 300 psig to about 750 psig.

12. The process of claim 2, wherein the CO₂ recovered during the depressurization step of the pressure swing adsorption cycle is collected at two or more different pressure levels utilizing a compressor.

13. The process of claim 12, wherein the pressure levels of the various stages of the carbon dioxide compressor are the same as the pressure levels in the depressurization step of the pressure swing adsorption cycle.

14. The process of claim 1, wherein the portion of the carbon dioxide purified stream obtained from the cryogenic purification unit that is not utilized as a co-feed in the co-feed step of the pressure swing adsorption cycle is further compressed for additional use.

15. A process for recovering gaseous hydrogen and gaseous carbon dioxide from a mixture of hydrocarbons utilizing a reformer unit in conjunction with a pressure swing adsorption unit and a cryogenic purification unit, the process comprising the steps of:

a) generating a hydrogen rich effluent that also contains at least carbon monoxide, carbon dioxide, methane and water vapor from one or more hydrocarbon feed streams in a reformer unit;

b) optionally treating the hydrogen rich effluent in a water gas shift reactor in order to further enrich the hydrogen content of the hydrogen rich effluent and to also increase the carbon dioxide content in the hydrogen rich effluent by oxidizing a portion of the carbon monoxide present in the effluent to carbon dioxide thereby obtaining a water shift effluent;

c) providing a pressure swing adsorption unit having at least two adsorption vessels, each adsorption vessel comprising two sections that are separate from one another and connected by a pipe that allows for the withdrawal of a methane rich gas stream from the pipe during a co-feed step of a hydrogen pressure swing adsorption cycle, each section of the adsorption vessels independently containing two or more adsorbent beds, the at least two adsorption vessels being parallel to one another and being independently operated on a process cycle that includes an adsorption step, the co-feed step, a depressurization step, a regeneration step and a re-pressurization step, with the proviso that the cycles in the at least two adsorption vessels run consecutively with at least one of the adsorption vessels being in a different step of the cycle compared to at least one of the remaining adsorption vessels;

d) introducing the hydrogen rich effluent or water gas shift effluent into at least one of the adsorption vessels of the pressure swing adsorption unit during the adsorption step of the hydrogen pressure swing adsorption cycle in order to allow for the recovery of a high purity gaseous hydrogen stream from the hydrogen rich effluent or the water gas shift effluent and the adsorption of the carbon monoxide, carbon dioxide, methane, water vapor and remaining hydrogen onto the adsorbents in the adsorbent beds of the pressure swing adsorption unit, the adsorption step being carried out at a pressure from about 250 psig to about 700 psig;

e) withdrawing the high purity gaseous hydrogen stream from the pressure swing adsorption unit during the adsorption step and passing this high purity gaseous hydrogen stream on for further use;

f) allowing the at least one adsorption vessel to enter into the co-feed step while introducing a purified carbon dioxide stream obtained from a downstream cryogenic purification unit into a lower (second) section, in order to allow the carbon dioxide to displace methane and carbon monoxide adsorbed onto the adsorbents in the adsorbent beds thereby producing a methane rich stream; an upper (first) section is isolated from the lower section;

g) withdrawing the methane rich stream from the pipe positioned between the two sections of the adsorption vessel and recycling this methane rich stream to the reformer unit;

h) allowing the at least one adsorption vessel to enter the depressurization step by depressurizing the adsorbent beds of the at least one adsorption vessel in order to release the carbon dioxide, carbon monoxide, methane, water vapor and hydrogen adsorbed to the adsorbents in the adsorbent beds thereby producing a carbon dioxide rich pressure swing adsorption tail gas having an increased concentration of carbon dioxide;

i) withdrawing the carbon dioxide rich pressure swing adsorption tail gas from the pressure swing adsorption unit;

j) regenerating and re-pressurizing the adsorbent beds for further use;

k) compressing the carbon dioxide rich pressure swing adsorption tail gas in a compressor unit to a pressure that ranges from about 250 psig to about 1000 psig in order to obtain a compressed carbon dioxide rich pressure swing adsorption tail gas;

l) introducing the compressed carbon dioxide rich pressure swing adsorption tail gas into the cryogenic purification unit in order to produce the carbon dioxide purified stream and an incondensibles stream that contains methane, carbon monoxide and hydrogen; and

m) withdrawing the carbon dioxide purified stream from the cryogenic purification unit and recycling at least a portion of the carbon dioxide purified stream to the pressure swing adsorption unit where the carbon dioxide purified stream is introduced into the adsorption vessels during the co-feed step of the cycle in order to increase the concentration of carbon dioxide in the pressure swing adsorption tail gas.

16. The process of claim 15, wherein the reformer unit is selected from a steam hydrocarbon reformer unit and an oxygen fed autothermal reformer unit.

17. The process of claim 16, wherein each section of the adsorption vessel comprises two or more adsorbent beds that are adjacent to one another within each section of the adsorption vessel.

18. The process of claim 16, wherein the pipe further contains a first valve which allows for the withdrawing of the methane rich stream between the two sections of the adsorption vessel during the co-feed step of the pressure swing adsorption cycle .

19. The process of claim 18, wherein the pipe further contains a means of isolating the first section of the adsorption vessel from the second section of the adsorption vessel during the withdrawal of the methane rich stream from the pipe through the first valve.

20. The process of claim 19, wherein the means of isolating the first section of the adsorption vessel from the second section of the adsorption vessel comprises a second valve which remains closed during the withdrawal of the methane rich stream from the pipe.

21. The process of claim 15, wherein the first section of the adsorption vessel contains adsorbent beds that include an adsorbent that is selective for methane, carbon monoxide, and nitrogen and the second section of the adsorption vessel contains adsorbent beds that include an adsorbent that allows for the removal of carbon dioxide.

22. The process of claim 21, wherein the pipe that connects the two sections of the adsorption vessel includes a means for isolating the two adsorption vessel sections from one another during the depressurization step of the pressure swing adsorption cycle, thus preventing the methane, carbon monoxide and nitrogen adsorbed to the adsorbent in the first section of the adsorption vessel from mixing with the carbon

dioxide adsorbed to the adsorbent of the second section of the adsorption vessel or the carbon dioxide from flowing back into the first section of the adsorption vessel.

23. The process of claim 15, wherein the carbon dioxide purified stream is introduced into the co-feed step of the pressure swing adsorption cycle at a pressure that is higher than the pressure during the adsorption step of the pressure swing adsorption cycle.

24. The process of claim 23, wherein the pressure during the co-feed step of the pressure swing adsorption cycle is from about 300 psig to about 750 psig.

25. The process of claim 15, wherein the carbon dioxide recovered during the depressurization step of the pressure swing adsorption cycle is collected at two or more different pressure levels utilizing a compressor.

26. The process of claim 25, wherein the pressure levels of the various stages of the carbon dioxide compressor are the same as the pressure levels in the depressurization step of the pressure swing adsorption cycle.

27. A process for recovering gaseous hydrogen and gaseous carbon dioxide from a mixture of hydrocarbons utilizing a reformer unit in conjunction with a pressure swing adsorption unit and a catalytic oxidizer unit, the process comprising the steps of:

a) generating a hydrogen rich effluent that also contains at least carbon monoxide, carbon dioxide, methane and water vapor from one or more hydrocarbon feed streams in a reformer unit;

b) optionally treating the hydrogen rich effluent in a water gas shift reactor in order to further enrich the hydrogen content of the hydrogen rich effluent and to also increase the carbon dioxide content in the hydrogen rich effluent by oxidizing a portion of the carbon monoxide present in the effluent to carbon dioxide thereby obtaining a water shift effluent;

c) providing a pressure swing adsorption unit having at least two adsorption vessels that each contain two or more adsorbent beds, the at least two adsorption vessels being parallel to one another and being independently operated on a cycle that includes an adsorption step, a co-feed step, a depressurization step, a regeneration step and a re-pressurization step, with the proviso that the cycles in the at least two adsorption vessels run consecutively with at least one of the adsorption vessels being in a different step of the cycle compared to at least one of the remaining adsorption vessels and the two or more adsorbent beds having an intermediate draw area positioned between at least two of the adsorbent beds thereby allowing for the withdrawal of a methane rich gas stream from the intermediate draw area during the co-feed step of the hydrogen pressure swing adsorption cycle;

d) introducing the hydrogen rich effluent or water gas shift effluent into at least one of the adsorption vessels of the pressure swing adsorption unit during the adsorption step of the hydrogen pressure swing adsorption cycle in order to allow for the recovery of a high purity gaseous hydrogen stream from the hydrogen rich effluent or the water gas shift effluent and the adsorption of the carbon monoxide, carbon dioxide, methane, water vapor and remaining hydrogen onto the adsorbents in the adsorbent beds of the pressure swing adsorption unit, the adsorption step being carried out at a pressure from about 250 psig to about 700 psig;

e) withdrawing the high purity gaseous hydrogen stream from the pressure swing adsorption unit during the adsorption step and passing this high purity gaseous hydrogen stream on for further use;

f) allowing the at least one adsorption vessel to enter into the co-feed step while introducing a purified carbon dioxide stream obtained from a downstream catalytic oxidizer unit in order to allow the carbon dioxide to displace methane and carbon monoxide adsorbed onto the adsorbents in the adsorbent beds thereby producing a methane rich stream;

g) withdrawing the methane rich stream from the intermediate draw area positioned between at least two of the adsorbent beds and recycling this methane rich stream to the reformer unit;

h) allowing the at least one adsorption vessel to enter the depressurization step by depressurizing the adsorbent beds of the at least one adsorption vessel in order to release the carbon dioxide, carbon monoxide, methane, water vapor and hydrogen adsorbed to the adsorbents in the adsorbent beds thereby producing a carbon dioxide rich pressure swing adsorption tail gas having an increased concentration of carbon dioxide;

i) withdrawing the carbon dioxide rich pressure swing adsorption tail gas from the pressure swing adsorption unit;

j) regenerating and re-pressurizing the adsorbent beds for further use;

k) compressing the carbon dioxide rich pressure swing adsorption tail gas in a compressor unit to a pressure that ranges from about 250 psig to about 1000 psig in order to obtain a compressed carbon dioxide rich pressure swing adsorption tail gas;

l) introducing the compressed carbon dioxide rich pressure swing adsorption tail gas into the catalytic oxidizer unit followed by introducing gaseous oxygen thereby allowing for the carbon monoxide, methane, and hydrogen to be combusted to produce an oxidized stream that contains carbon dioxide and water, the amount of oxygen being introduced being slightly below the stoichiometric requirement;

m) condensing the oxidized stream to allow for the removal of water by cooling the oxidized stream to ambient temperature to produce the carbon dioxide purified stream; and

n) withdrawing the carbon dioxide purified stream from the catalytic oxidizer unit and recycling at least a portion of the carbon dioxide purified stream to the pressure swing adsorption unit where the carbon dioxide purified stream is introduced into the adsorption vessels during the co-feed step of the cycle in order to increase the concentration of carbon dioxide in the hydrogen pressure swing adsorption tail gas.

28. The process of claim 27, wherein the reformer unit is selected from a steam hydrocarbon reformer unit and an oxygen fed autothermal reformer unit.

29. The process of claim 28, wherein the adsorption vessel contains four adsorbent beds with two of the adsorbent beds being adjacent to one another on one

side of the intermediate draw area and the remaining two adsorbent beds being positioned on the other side of the intermediate draw area.

30. The process of claim 28, wherein the adsorption vessel comprises a first adsorbent bed that is separated from a second adsorbent bed by the intermediate draw area.

31. The process of claim 30, wherein the first adsorbent bed and the second adsorbent bed each independently comprises one or more adsorbents selected from activated alumina, silica gel, activated carbon, zeolites, and combinations thereof.

32. The process of claim 31, wherein the one or more adsorbents are in layers within the individual adsorbent beds.

33. The process of claim 30, wherein the first adsorbent bed contains one or more adsorbents that are selective for methane and carbon monoxide and the second adsorbent bed contains one or more adsorbents that are selective for carbon dioxide and water vapor.

34. The process of claim 33, wherein the one or more adsorbents are in layers within the individual beds.

35. The process of claim 33, wherein when the hydrogen rich effluent or the water gas shift effluent is introduced into the adsorption vessel during the adsorption step of the hydrogen pressure swing adsorption cycle, the hydrogen rich effluent or the water gas shift effluent comes into contact with the first adsorbent bed before it comes into contact with the second adsorbent bed.

36. The process of claim 28, wherein the carbon dioxide purified stream is introduced into the co-feed step of the pressure swing adsorption cycle at a pressure

that is higher than the pressure during the adsorption step of the pressure swing adsorption cycle.

37. The process of claim 36, wherein the pressure during the co-feed step of the pressure swing adsorption cycle is from about 300 psig to about 750 psig.

38. The process of claim 28, wherein the CO₂ recovered during the depressurization step of the pressure swing adsorption cycle is collected at two or more different pressure levels utilizing a compressor.

39. The process of claim 38, wherein the pressure levels of the various stages of the carbon dioxide compressor are the same as the pressure levels in the depressurization step of the pressure swing adsorption cycle.

40. The process of claim 27, wherein the portion of the carbon dioxide purified stream obtained from the cryogenic purification unit that is not utilized as a co-feed in the co-feed step of the pressure swing adsorption cycle is further compressed for additional use.

41. A process for recovering gaseous hydrogen and gaseous carbon dioxide from a mixture of hydrocarbons utilizing a reformer unit in conjunction with a pressure swing adsorption unit and a catalytic oxidizer unit, the process comprising the steps of:

a) generating a hydrogen rich effluent that also contains at least carbon monoxide, carbon dioxide, methane and water vapor from one or more hydrocarbon feed streams in a reformer unit;

b) optionally treating the hydrogen rich effluent in a water gas shift reactor in order to further enrich the hydrogen content of the hydrogen rich effluent and to also increase the carbon dioxide content in the hydrogen rich effluent by oxidizing a portion of the carbon monoxide present in the effluent to carbon dioxide thereby obtaining a water shift effluent;

c) providing a pressure swing adsorption unit having at least two adsorption vessels, each adsorption vessel comprising two sections that are separate from one another and connected by a pipe that allows for the withdrawal of a methane rich gas stream from the pipe during a co-feed step of a hydrogen pressure swing adsorption cycle, each section of the adsorption vessels independently containing two or more adsorbent beds, the at least two adsorption vessels being parallel to one another and being independently operated on a process cycle that includes an adsorption step, the co-feed step, a depressurization step, a regeneration step and a re-pressurization step, with the proviso that the cycles in the at least two adsorption vessels run consecutively with at least one of the adsorption vessels being in a different step of the cycle compared to at least one of the remaining adsorption vessels;

d) introducing the hydrogen rich effluent or water gas shift effluent into at least one of the adsorption vessels of the pressure swing adsorption unit during the adsorption step of the hydrogen pressure swing adsorption cycle in order to allow for the recovery of a high purity gaseous hydrogen stream from the hydrogen rich effluent or the water gas shift effluent and the adsorption of the carbon monoxide, carbon dioxide, methane, water vapor and remaining hydrogen onto the adsorbents in the adsorbent beds of the pressure swing adsorption unit, the adsorption step being carried out at a pressure from about 250 psig to about 700 psig;

e) withdrawing the high purity gaseous hydrogen stream from the pressure swing adsorption unit during the adsorption step and passing this high purity gaseous hydrogen stream on for further use;

f) allowing the at least one adsorption vessel to enter into the co-feed step while introducing a purified carbon dioxide stream obtained from a downstream cryogenic purification unit into a lower (second) section, in order to allow the carbon dioxide to displace methane and carbon monoxide adsorbed onto the adsorbents in the adsorbent beds thereby producing a methane rich stream;

g) withdrawing the methane rich stream from the pipe positioned between the two sections of the adsorption vessel and recycling this methane rich stream to the reformer unit;

h) allowing the at least one adsorption vessel to enter the depressurization step by depressurizing the adsorbent beds of the at least one adsorption vessel in order to release the carbon dioxide, carbon monoxide, methane, water vapor and hydrogen adsorbed to the adsorbents in the adsorbent beds thereby producing a carbon dioxide rich pressure swing adsorption tail gas having an increased concentration of carbon dioxide;

i) withdrawing the carbon dioxide rich pressure swing adsorption tail gas from the pressure swing adsorption unit;

j) regenerating and re-pressurizing the adsorbent beds for further use;

k) compressing the carbon dioxide rich pressure swing adsorption tail gas in a compressor unit to a pressure that ranges from about 250 psig to about 1000 psig in order to obtain a compressed carbon dioxide rich pressure swing adsorption tail gas;

l) introducing the compressed carbon dioxide rich pressure swing adsorption tail gas into a catalytic oxidizer unit followed by introducing gaseous oxygen thereby allowing for the carbon monoxide, methane, and hydrogen to be combusted to produce an oxidized stream that contains carbon dioxide and water, the amount of oxygen being introduced being slightly below the stoichiometric requirement;

m) condensing the oxidized stream to allow for the removal of water by cooling the oxidized stream to ambient temperature to produce the carbon dioxide purified stream; and

n) withdrawing the carbon dioxide purified stream from the catalytic oxidizer unit and recycling at least a portion of the carbon dioxide purified stream to the pressure swing adsorption unit where the carbon dioxide purified stream is introduced into the adsorption vessels during the co-feed step of the cycle in order to increase the concentration of carbon dioxide in the pressure swing adsorption tail gas.

42. The process of claim 41, wherein the reformer unit is selected from a steam hydrocarbon reformer unit and an oxygen fed autothermal reformer unit.

43. The process of claim 42, wherein each section of the adsorption vessel comprises two or more adsorbent beds that are adjacent to one another within each section of the adsorption vessel.

44. The process of claim 42, wherein the pipe further contains a first valve which allows for the withdrawing of the methane rich stream between the two sections of the adsorption vessel during the co-feed step of the pressure swing adsorption cycle .

45. The process of claim 44, wherein the pipe further contains a means of isolating the first section of the adsorption vessel from the second section of the adsorption vessel during the withdrawal of the methane rich stream from the pipe through the first valve.

46. The process of claim 45, wherein the means of isolating the first section of the adsorption vessel from the second section of the adsorption vessel comprises a second valve which remains closed during the withdrawal of the methane rich stream from the pipe.

47. The process of claim 41, wherein the first section of the adsorption vessel contains adsorbent beds that include an adsorbent that is selective for methane, carbon monoxide, and nitrogen and the second section of the adsorption vessel contains adsorbent beds that include an adsorbent that allows for the removal of carbon dioxide.

48. The process of claim 47, wherein the pipe that connects the two sections of the adsorption vessel includes a means for isolating the two adsorption vessel sections from one another during the depressurization step of the pressure swing adsorption cycle, thus preventing the methane, carbon monoxide and nitrogen adsorbed to the adsorbent in the first section of the adsorption vessel from mixing with the carbon dioxide adsorbed to the adsorbent of the second section of the adsorption vessel or the carbon dioxide from flowing back into the first section of the adsorption vessel.

49. The process of claim 41, wherein the carbon dioxide purified stream is introduced into the co-feed step of the pressure swing adsorption cycle at a pressure that is higher than the pressure during the adsorption step of the pressure swing adsorption cycle.

50. The process of claim 49, wherein the pressure during the co-feed step of the pressure swing adsorption cycle is from about 300 psig to about 750 psig.

51. The process of claim 41, wherein the carbon dioxide recovered during the depressurization step of the pressure swing adsorption cycle is collected at two or more different pressure levels utilizing a compressor.

52. The process of claim 51, wherein the pressure levels of the various stages of the carbon dioxide compressor are the same as the pressure levels in the depressurization step of the pressure swing adsorption cycle.

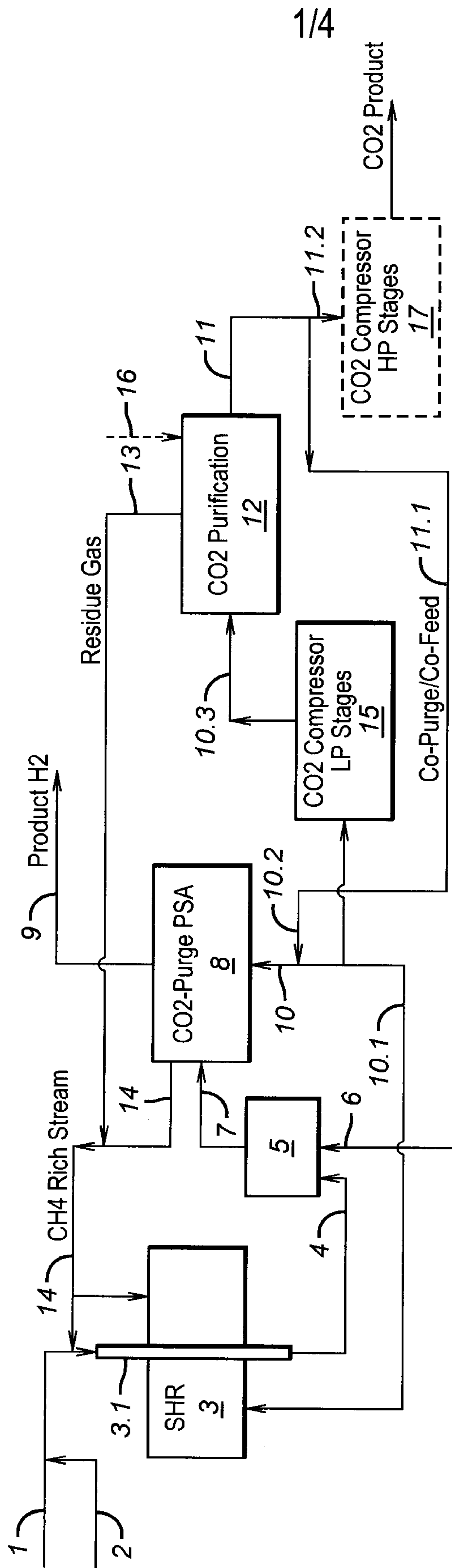


FIG. 1

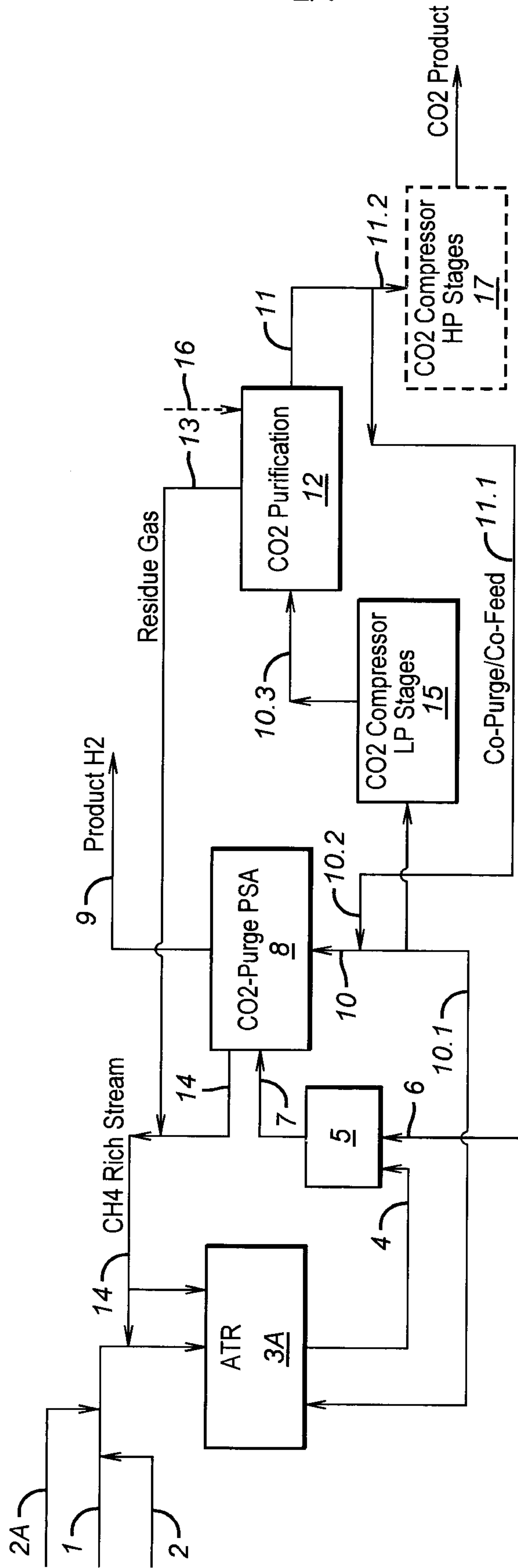


FIG. 2

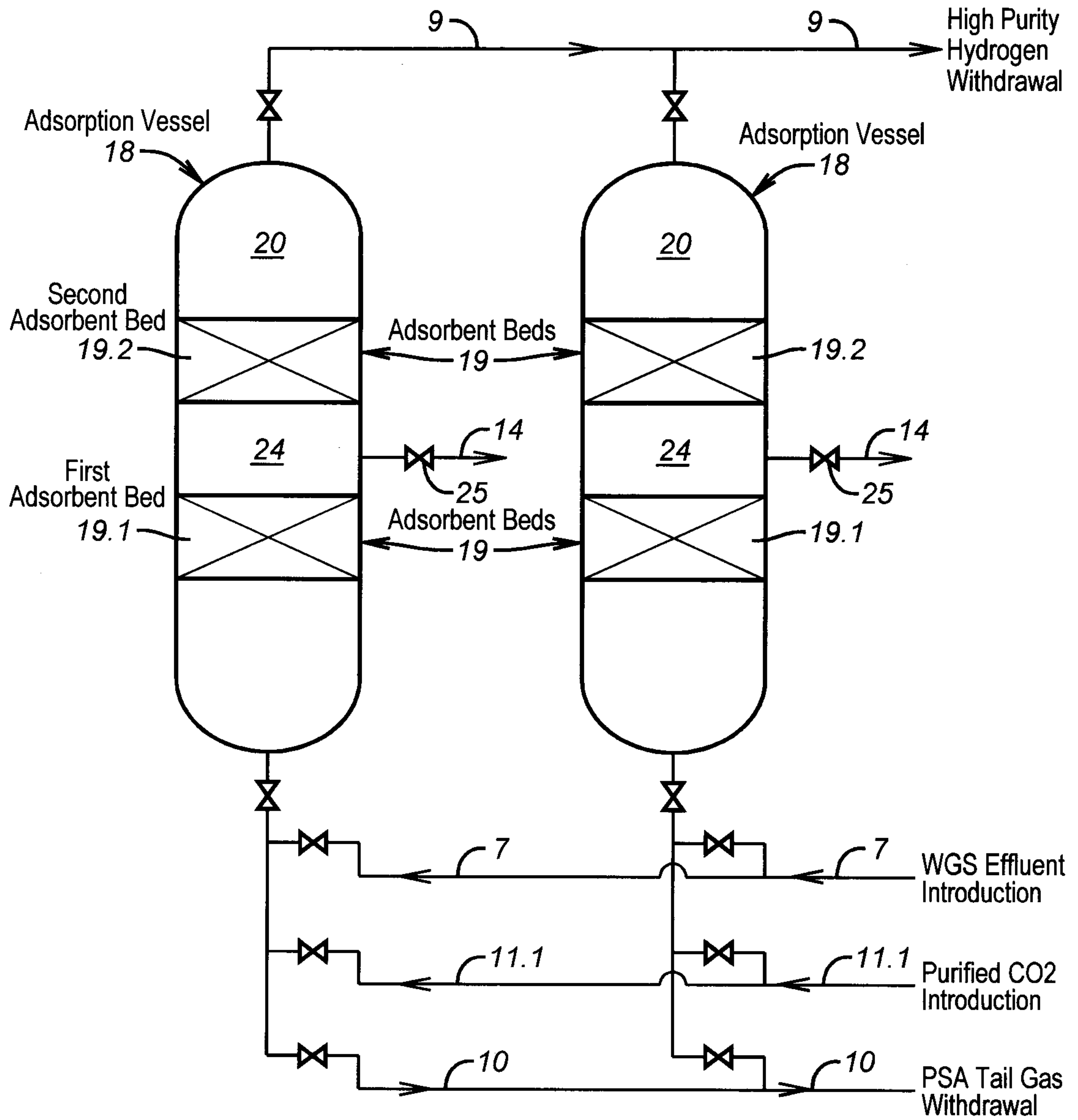


FIG. 3

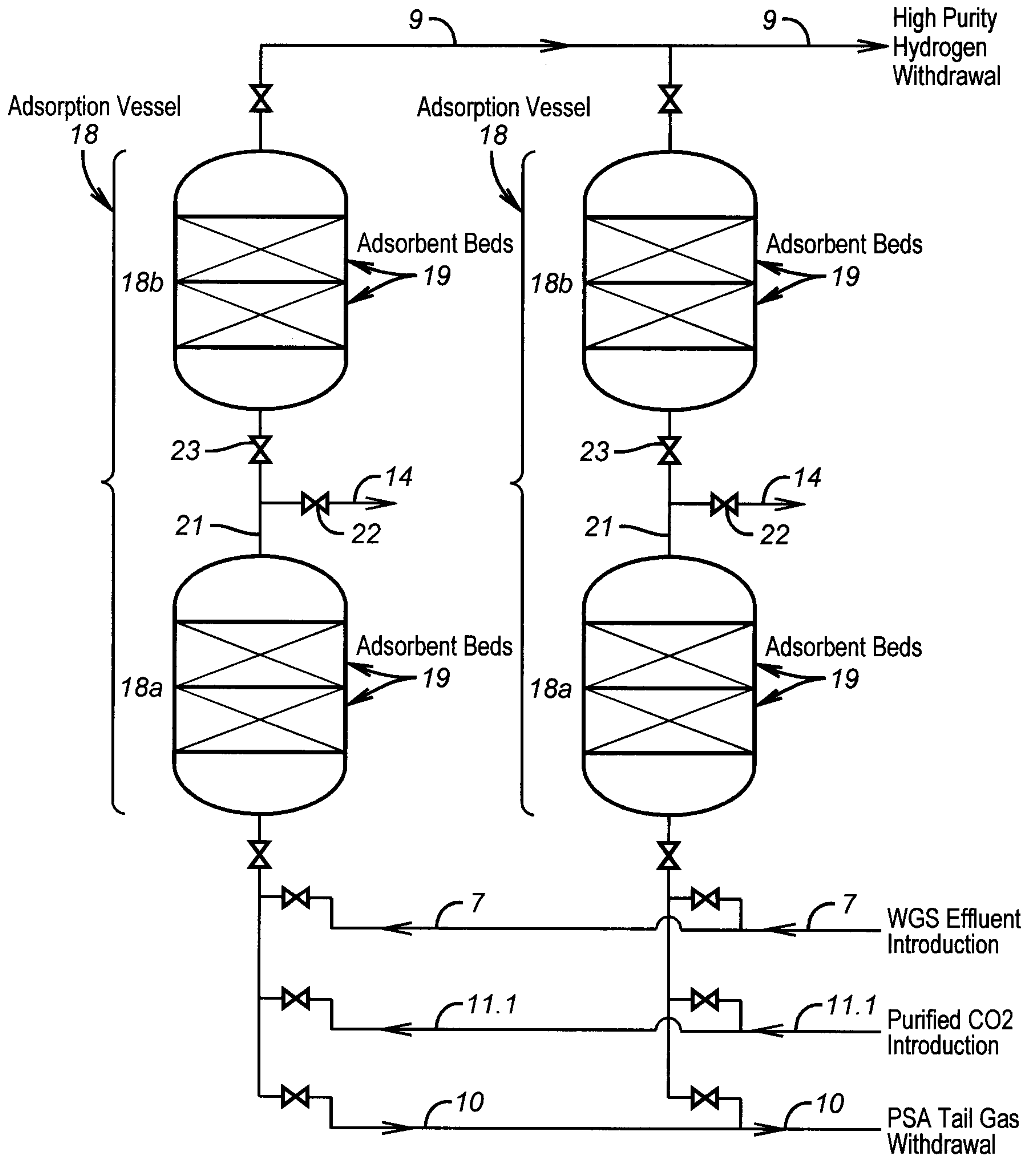


FIG. 4

