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30 ROCKEFELLER PLAZA

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TO ALL WHOM IT MAY CONCERN:

Be it known that WE, RICHARD N. PITMAN, a citizen of the United States, residing in Sunset, County of Montague, State of Texas, whose post office address is 111 Clear Creek Dr., Sunset, Texas 76270, JOHN D. WILKINSON, JOE T. LYNCH, and HANK M. HUDSON, all citizens of the United States, all residing in Midland, County of Midland, State of Texas, whose post office addresses are 4113 Tanforan Ave., Midland, Texas 79707; 5510 Ashwood Ct., Midland, Texas 79707; and 2508 W. Sinclair Ave., Midland, Texas 79705, respectively, and TONY L. MARTINEZ, a citizen of the United States, residing in Odessa, County of Ector, State of Texas, whose post office address is 3725 W. 12th St., Odessa, Texas 79763, have invented an improvement in

HYDROCARBON GAS PROCESSING

of which the following is a

SPECIFICATION

BACKGROUND OF THE INVENTION

[0001] This invention relates to a process for the separation of a gas containing hydrocarbons. The applicants claim the benefits under Title 35, United States Code, Section 119(e) of prior U.S. Provisional Application Number 60/900,400 which was filed on February 9, 2007.

[0002] Ethylene, ethane, propylene, propane, and/or heavier hydrocarbons can be recovered from a variety of gases, such as natural gas, refinery gas, and synthetic gas streams obtained from other hydrocarbon materials such as coal, crude oil, naphtha, oil shale, tar sands, and lignite. Natural gas usually has a major proportion of methane and ethane, i.e., methane and ethane together comprise at least 50 mole percent of the gas. The gas also contains relatively lesser amounts of heavier hydrocarbons such as propane, butanes, pentanes, and the like, as well as hydrogen, nitrogen, carbon dioxide, and other gases.

[0003] The present invention is generally concerned with the recovery of ethylene, ethane, propylene, propane, and heavier hydrocarbons from such gas streams. A typical analysis of a gas stream to be processed in accordance with this invention would be, in approximate mole percent, 92.5% methane, 4.2% ethane and other C_2 components, 1.3% propane and other C_3 components, 0.4% iso-butane, 0.3% normal butane, 0.5% pentanes plus, with the balance made up of nitrogen and carbon dioxide. Sulfur containing gases are also sometimes present.

[0004] The historically cyclic fluctuations in the prices of both natural gas and its natural gas liquid (NGL) constituents have at times reduced the incremental value of ethane, ethylene, propane, propylene, and heavier components as liquid products. This has resulted in a demand for processes that can provide more efficient recoveries of these products. Available processes for separating these materials include those based upon cooling and refrigeration of gas, oil absorption, and refrigerated oil absorption. Additionally, cryogenic processes have become popular because of the availability of economical equipment that produces power while simultaneously expanding and

extracting heat from the gas being processed. Depending upon the pressure of the gas source, the richness (ethane, ethylene, and heavier hydrocarbons content) of the gas, and the desired end products, each of these processes or a combination thereof may be employed.

[0005] The cryogenic expansion process is now generally preferred for natural gas liquids recovery because it provides maximum simplicity with ease of startup, operating flexibility, good efficiency, safety, and good reliability. U.S. Patent Nos. 3,292,380; 4,061,481; 4,140,504; 4,157,904; 4,171,964; 4,185,978; 4,251,249; 4,278,457; 4,519,824; 4,617,039; 4,687,499; 4,689,063; 4,690,702; 4,854,955; 4,869,740; 4,889,545; 5,275,005; 5,555,748; 5,566,554; 5,568,737; 5,771,712; 5,799,507; 5,881,569; 5,890,378; 5,983,664; 6,182,469; 6,578,379; 6,712,880; 6,915,662; 7,191,617; 7,219,513; reissue U.S. Patent No. 33,408; and co-pending application nos. 11/430,412 and 11/839,693 describe relevant processes (although the description of the present invention in some cases is based on different processing conditions than those described in the cited U.S. Patents).

[0006] In a typical cryogenic expansion recovery process, a feed gas stream under pressure is cooled by heat exchange with other streams of the process and/or external sources of refrigeration such as a propane compression-refrigeration system. As the gas is cooled, liquids may be condensed and collected in one or more separators as high-pressure liquids containing some of the desired C_2+ or C_3+ components. Depending on the richness of the gas and the amount of liquids formed, the high-pressure liquids may be expanded to a lower pressure and fractionated. The vaporization occurring during expansion of the liquids results in further cooling of the stream. Under some

conditions, pre-cooling the high pressure liquids prior to the expansion may be desirable in order to further lower the temperature resulting from the expansion. The expanded stream, comprising a mixture of liquid and vapor, is fractionated in a distillation (demethanizer or deethanizer) column. In the column, the expansion cooled stream(s) is (are) distilled to separate residual methane, nitrogen, and other volatile gases as overhead vapor from the desired C_2 components, C_3 components, and heavier hydrocarbon components as bottom liquid product, or to separate residual methane, C_2 components, nitrogen, and other volatile gases as overhead vapor from the desired C_3 components and heavier hydrocarbon components as bottom liquid product.

[0007] If the feed gas is not totally condensed (typically it is not), the vapor remaining from the partial condensation can be split into two streams. One portion of the vapor is passed through a work expansion machine or engine, or an expansion valve, to a lower pressure at which additional liquids are condensed as a result of further cooling of the stream. The pressure after expansion is essentially the same as the pressure at which the distillation column is operated. The combined vapor-liquid phases resulting from the expansion are supplied as feed to the column.

[0008] The remaining portion of the vapor is cooled to substantial condensation by heat exchange with other process streams, e.g., the cold fractionation tower overhead. Some or all of the high-pressure liquid may be combined with this vapor portion prior to cooling. The resulting cooled stream is then expanded through an appropriate expansion device, such as an expansion valve, to the pressure at which the demethanizer is operated. During expansion, a portion of the liquid will usually vaporize, resulting in cooling of the total stream. The flash expanded stream is then supplied as top feed to the demethanizer.

Typically, the vapor portion of the flash expanded stream and the demethanizer overhead vapor combine in an upper separator section in the fractionation tower as residual methane product gas. Alternatively, the cooled and expanded stream may be supplied to a separator to provide vapor and liquid streams. The vapor is combined with the tower overhead and the liquid is supplied to the column as a top column feed.

[0009] In the ideal operation of such a separation process, the residue gas leaving the process will contain substantially all of the methane in the feed gas with essentially none of the heavier hydrocarbon components and the bottoms fraction leaving the demethanizer will contain substantially all of the heavier hydrocarbon components with essentially no methane or more volatile components. In practice, however, this ideal situation is not obtained because the conventional demethanizer is operated largely as a stripping column. The methane product of the process, therefore, typically comprises vapors leaving the top fractionation stage of the column, together with vapors not subjected to any rectification step. Considerable losses of C_2, C_3, and C_4+ components occur because the top liquid feed contains substantial quantities of these components and heavier hydrocarbon components, resulting in corresponding equilibrium quantities of C_2 components, C_3 components, C_4 components, and heavier hydrocarbon components in the vapors leaving the top fractionation stage of the demethanizer. The loss of these desirable components could be significantly reduced if the rising vapors could be brought into contact with a significant quantity of liquid (reflux) capable of absorbing the C_2 components, C_3 components, C_4 components, and heavier hydrocarbon components from the vapors.

[0010] In recent years, the preferred processes for hydrocarbon separation use an upper absorber section to provide additional rectification of the rising vapors. The source of the reflux stream for the upper rectification section is typically a recycled stream of residue gas supplied under pressure. The recycled residue gas stream is usually cooled to substantial condensation by heat exchange with other process streams, e.g., the cold fractionation tower overhead. The resulting substantially condensed stream is then expanded through an appropriate expansion device, such as an expansion valve, to the pressure at which the demethanizer is operated. During expansion, a portion of the liquid will usually vaporize, resulting in cooling of the total stream. The flash expanded stream is then supplied as top feed to the demethanizer. Typically, the vapor portion of the expanded stream and the demethanizer overhead vapor combine in an upper separator section in the fractionation tower as residual methane product gas. Alternatively, the cooled and expanded stream may be supplied to a separator to provide vapor and liquid streams, so that thereafter the vapor is combined with the tower overhead and the liquid is supplied to the column as a top column feed. Typical process schemes of this type are disclosed in U.S. Patent Nos. 4,889,545; 5,568,737; and 5,881,569, co-pending application no. 11/430,412, and in Mowrey, E. Ross, "Efficient, High Recovery of Liquids from Natural Gas Utilizing a High Pressure Absorber", Proceedings of the Eighty-First Annual Convention of the Gas Processors Association, Dallas, Texas, March 11-13, 2002.

[0011] The present invention also employs an upper rectification section (or a separate rectification column in some embodiments). However, two reflux streams are provided for this rectification section. The upper reflux stream is a recycled stream of

residue gas as described above. In addition, however, a supplemental reflux stream is provided at one or more lower feed points by using a side draw of the vapors rising in a lower portion of the tower (which may be combined with a portion of the tower overhead vapor). Because the vapor streams lower in the tower contain a modest concentration of C_2 components and heavier components, this side draw stream can be substantially condensed by moderately elevating its pressure and using only the refrigeration available in the cold vapor leaving the upper rectification section. This condensed liquid, which is predominantly liquid methane and ethane, can then be used to absorb C_2 components, C_3 components, C_4 components, and heavier hydrocarbon components from the vapors rising through the lower portion of the upper rectification section and thereby capture these valuable components in the bottom liquid product from the demethanizer. Since this lower reflux stream captures much of the C_2 components and essentially all of the C_3+ components, only a relatively small flow rate of liquid in the upper reflux stream is needed to absorb the C_2 components remaining in the rising vapors and likewise capture these C_2 components in the bottom liquid product from the demethanizer.

[0012] In accordance with the present invention, it has been found that C_2 component recoveries in excess of 97 percent can be obtained. Similarly, in those instances where recovery of C_2 components is not desired, C_3 recoveries in excess of 98% can be maintained. In addition, the present invention makes possible essentially 100 percent separation of methane (or C_2 components) and lighter components from the C_2 components (or C_3 components) and heavier components at reduced energy requirements compared to the prior art while maintaining the same recovery levels. The present invention, although applicable at lower pressures and warmer temperatures, is

particularly advantageous when processing feed gases in the range of 400 to 1500 psia [2,758 to 10,342 kPa(a)] or higher under conditions requiring NGL recovery column overhead temperatures of -50°F [-46°C] or colder.

[0013] For a better understanding of the present invention, reference is made to the following examples and drawings. Referring to the drawings:

[0014] FIG. 1 is a flow diagram of a prior art natural gas processing plant in accordance with United States Patent No. 5,568,737;

[0015] FIG. 2 is a flow diagram of an alternative prior art natural gas processing plant in accordance with co-pending application no. 11/430,412;

[0016] FIG. 3 is a flow diagram of a natural gas processing plant in accordance with the present invention; and

[0017] FIGS. 4 through 8 are flow diagrams illustrating alternative means of application of the present invention to a natural gas stream.

[0018] In the following explanation of the above figures, tables are provided summarizing flow rates calculated for representative process conditions. In the tables appearing herein, the values for flow rates (in moles per hour) have been rounded to the nearest whole number for convenience. The total stream rates shown in the tables include all non-hydrocarbon components and hence are generally larger than the sum of the stream flow rates for the hydrocarbon components. Temperatures indicated are approximate values rounded to the nearest degree. It should also be noted that the process design calculations performed for the purpose of comparing the processes depicted in the figures are based on the assumption of no heat leak from (or to) the surroundings to (or from) the process. The quality of commercially available insulating

materials makes this a very reasonable assumption and one that is typically made by those skilled in the art.

[0019] For convenience, process parameters are reported in both the traditional British units and in the units of the Système International d'Unités (SI). The molar flow rates given in the tables may be interpreted as either pound moles per hour or kilogram moles per hour. The energy consumptions reported as horsepower (HP) and/or thousand British Thermal Units per hour (MBTU/Hr) correspond to the stated molar flow rates in pound moles per hour. The energy consumptions reported as kilowatts (kW) correspond to the stated molar flow rates in kilogram moles per hour.

DESCRIPTION OF THE PRIOR ART

[0020] FIG. 1 is a process flow diagram showing the design of a processing plant to recover C_2+ components from natural gas using prior art according to assignee's U.S. Pat. No. 5,568,737. In this simulation of the process, inlet gas enters the plant at 120°F [49°C] and 1040 psia [7,171 kPa(a)] as stream **31**. If the inlet gas contains a concentration of sulfur compounds which would prevent the product streams from meeting specifications, the sulfur compounds are removed by appropriate pretreatment of the feed gas (not illustrated). In addition, the feed stream is usually dehydrated to prevent hydrate (ice) formation under cryogenic conditions. Solid desiccant has typically been used for this purpose.

[0021] The feed stream **31** is cooled in heat exchanger **10** by heat exchange with a portion (stream **46**) of cool distillation stream **39a** at -17°F [-27°C], bottom liquid product at 79°F [26°C] (stream **42a**) from the demethanizer bottoms pump, **19**, demethanizer reboiler liquids at 56°F [14°C] (stream **41**), and demethanizer side reboiler

liquids at -19°F [-28°C] (stream 40). Note that in all cases exchanger 10 is representative of either a multitude of individual heat exchangers or a single multi-pass heat exchanger, or any combination thereof. (The decision as to whether to use more than one heat exchanger for the indicated cooling services will depend on a number of factors including, but not limited to, inlet gas flow rate, heat exchanger size, stream temperatures, etc.) The cooled stream **31a** enters separator **11** at 6°F [-14°C] and 1025 psia [7,067 kPa(a)] where the vapor (stream **32**) is separated from the condensed liquid (stream **33**).

[0022] The vapor (stream **32**) from separator **11** is divided into two streams, **34** and **36**. Stream **34**, containing about 30% of the total vapor, is combined with the separator liquid (stream **33**). The combined stream **35** then passes through heat exchanger **12** in heat exchange relation with cold distillation stream **39** at -142°F [-96°C] where it is cooled to substantial condensation. The resulting substantially condensed stream **35a** at -138°F [-94°C] is then flash expanded through an appropriate expansion device, such as expansion valve **13**, to the operating pressure (approximately 423 psia [2,916 kPa(a)]) of fractionation tower **17**. The expanded stream **35b** leaving expansion valve **13** reaches a temperature of -140°F [-96°C] and is supplied to fractionation tower **17** at a mid-column feed point.

[0023] The remaining 70% of the vapor from separator **11** (stream **36**) enters a work expansion machine **14** in which mechanical energy is extracted from this portion of the high pressure feed. The machine **14** expands the vapor substantially isentropically to the tower operating pressure, with the work expansion cooling the expanded stream **36a** to a temperature of approximately -75°F [-60°C]. The typical commercially available

expanders are capable of recovering on the order of 80-88% of the work theoretically available in an ideal isentropic expansion. The work recovered is often used to drive a centrifugal compressor (such as item **15**) that can be used to re-compress the heated distillation stream (stream **39b**), for example. The partially condensed expanded stream **36a** is thereafter supplied to fractionation tower **17** at a second mid-column feed point.

[0024] The recompressed and cooled distillation stream **39e** is divided into two streams. One portion, stream **47**, is the volatile residue gas product. The other portion, recycle stream **48**, flows to heat exchanger **22** where it is cooled to -6°F [-21°C] (stream **48a**) by heat exchange with a portion (stream **45**) of cool distillation stream **39a**. The cooled recycle stream then flows to exchanger **12** where it is cooled to -138°F [-94°C] and substantially condensed by heat exchange with cold distillation stream **39** at -142°F [-96°C]. The substantially condensed stream **48b** is then expanded through an appropriate expansion device, such as expansion valve **23**, to the demethanizer operating pressure, resulting in cooling of the total stream to -144°F [-98°C]. The expanded stream **48c** is then supplied to fractionation tower **17** as the top column feed. The vapor portion (if any) of stream **48c** combines with the vapors rising from the top fractionation stage of the column to form distillation stream **39**, which is withdrawn from an upper region of the tower.

[0025] The demethanizer in tower **17** is a conventional distillation column containing a plurality of vertically spaced trays, one or more packed beds, or some combination of trays and packing. As is often the case in natural gas processing plants, the fractionation tower may consist of two sections. The upper section **17a** is a separator wherein the partially vaporized top feed is divided into its respective vapor and liquid

portions, and wherein the vapor rising from the lower distillation or demethanizing section **17b** is combined with the vapor portion (if any) of the top feed to form the cold demethanizer overhead vapor (stream **39**) which exits the top of the tower at -142°F [-96°C]. The lower, demethanizing section **17b** contains the trays and/or packing and provides the necessary contact between the liquids falling downward and the vapors rising upward. The demethanizing section **17b** also includes reboilers (such as the reboiler and side reboiler described previously) which heat and vaporize a portion of the liquids flowing down the column to provide the stripping vapors which flow up the column to strip the liquid product, stream **42**, of methane and lighter components.

[0026] Liquid product stream **42** exits the bottom of the tower at 75°F [24°C], based on a typical specification of a methane to ethane ratio of 0.025:1 on a molar basis in the bottom product. It is pumped to a pressure of approximately 650 psia [4,482 kPa(a)] in demethanizer bottoms pump **19**, and the pumped liquid product is then warmed to 116°F [47°C] as it provides cooling of stream **31** in exchanger **10** before flowing to storage.

[0027] The demethanizer overhead vapor (stream **39**) passes countercurrently to the incoming feed gas and recycle stream in heat exchanger **12** where it is heated to -17°F [-27°C] (stream **39a**), and in heat exchanger **22** and heat exchanger **10** where it is heated to 84°F [29°C] (stream **39b**). The distillation stream is then re-compressed in two stages. The first stage is compressor **15** driven by expansion machine **14**. The second stage is compressor **20** driven by a supplemental power source which compresses stream **39c** to sales line pressure (stream **39d**). After cooling to 120°F [49°C] in discharge cooler **21**, stream **39e** is split into the residue gas product (stream **47**) and the recycle stream **48** as

described earlier. Residue gas stream 47 flows to the sales gas pipeline at 1040 psia [7,171 kPa(a)], sufficient to meet line requirements (usually on the order of the inlet pressure).

[0028] A summary of stream flow rates and energy consumption for the process illustrated in FIG. 1 is set forth in the following table:

Table I

(FIG. 1)

Stream Flow Summary - Lb. Moles/Hr [kg moles/Hr]

Stream	Methane	Ethane	Propane	Butanes+	Total
31	25,384	1,161	362	332	27,451
32	25,313	1,147	349	255	27,275
33	71	14	13	77	176
34	7,594	344	105	76	8,182
35	7,665	358	118	153	8,358
36	17,719	803	244	179	19,093
39	29,957	38	0	0	30,147
48	4,601	6	0	0	4,630
47	25,356	32	0	0	25,517
42	28	1,129	362	332	1,934

Recoveries*			
Ethane	97.21%		
Propane	100.00%		
Butanes+	100.00%		
Power			
Residue Gas Compression	13,857 HP	[22,781	kW]

* (Based on un-rounded flow rates)

[0029] FIG. 2 represents an alternative prior art process in accordance with co-pending application no. 11/430,412. The process of FIG. 2 has been applied to the same feed gas composition and conditions as described above for FIG. 1. In the simulation of this process, as in the simulation for the process of FIG. 1, operating conditions were selected to minimize energy consumption for a given recovery level.

[0030] The feed stream **31** is cooled in heat exchanger **10** by heat exchange with a portion of the cool distillation column overhead stream (stream **46**) at -76°F [-60°C], demethanizer bottoms liquid (stream **42a**) at 87°F [31°C], demethanizer reboiler liquids at 62°F [17°C] (stream **41**), and demethanizer side reboiler liquids at -42°F [-41°C] (stream **40**). The cooled stream **31a** enters separator **11** at -46°F [-43°C] and 1025 psia [7,067 kPa(a)] where the vapor (stream **32**) is separated from the condensed liquid (stream **33**).

[0031] The separator vapor (stream **32**) enters a work expansion machine **14** in which mechanical energy is extracted from this portion of the high pressure feed. The machine **14** expands the vapor substantially isentropically to the tower operating pressure

of 461 psia [3,178 kPa(a)], with the work expansion cooling the expanded stream **32a** to a temperature of approximately -111°F [-79°C]. The partially condensed expanded stream **32a** is thereafter supplied to fractionation tower **17** at a mid-column feed point.

[0032] The recompressed and cooled distillation stream **39e** is divided into two streams. One portion, stream **47**, is the volatile residue gas product. The other portion, recycle stream **48**, flows to heat exchanger **22** where it is cooled to -70°F [-57°C] (stream **48a**) by heat exchange with a portion (stream **45**) of cool distillation stream **39a** at -76°F [-60°C]. The cooled recycle stream then flows to exchanger **12** where it is cooled to -133°F [-92°C] and substantially condensed by heat exchange with cold distillation column overhead stream **39**. The substantially condensed stream **48b** is then expanded through an appropriate expansion device, such as expansion valve **23**, to the demethanizer operating pressure, resulting in cooling of the total stream to -141°F [-96°C]. The expanded stream **48c** is then supplied to the fractionation tower as the top column feed. The vapor portion (if any) of stream **48c** combines with the vapors rising from the top fractionation stage of the column to form distillation stream **39**, which is withdrawn from an upper region of the tower.

[0033] A portion of the distillation vapor (stream **49**) is withdrawn from fractionation tower **17** at -119°F [-84°C] and is compressed to about 727 psia [5,015 kPa(a)] by reflux compressor **24**. The separator liquid (stream **33**) is expanded to this pressure by expansion valve **16**, and the expanded stream **33a** at -62°F [-52°C] is combined with stream **49a** at -66°F [-54°C]. The combined stream **35** is then cooled from -68°F [-56°C] to -133°F [-92°C] and condensed (stream **35a**) in heat exchanger **12** by heat exchange with the cold demethanizer overhead stream **39** exiting the top of

demethanizer **17** at -137°F [-94°C]. The resulting substantially condensed stream **35a** is then flash expanded through expansion valve **13** to the operating pressure of fractionation tower **17**, cooling stream **35b** to a temperature of -135°F [-93°C] whereupon it is supplied to fractionation tower **17** at a mid-column feed point.

[0034] The liquid product stream **42** exits the bottom of the tower at 82°F [28°C], based on a typical specification of a methane to ethane ratio of 0.025:1 on a molar basis in the bottom product. Pump **19** delivers stream **42a** to heat exchanger **10** as described previously where it is heated from 87°F [31°C] to 116°F [47°C] before flowing to storage.

[0035] The demethanizer overhead vapor stream **39** is warmed in heat exchanger **12** as it provides cooling to combined stream **35** and recycle stream **48a** as described previously, and further heated in heat exchanger **22** and heat exchanger **10**. The heated stream **39b** at 96°F [36°C] is then re-compressed in two stages, compressor **15** driven by expansion machine **14** and compressor **20** driven by a supplemental power source. After stream **39d** is cooled to 120°F [49°C] in discharge cooler **21** to form stream **39e**, recycle stream **48** is withdrawn as described earlier to form residue gas stream **47** which flows to the sales gas pipeline at 1040 psia [7,171 kPa(a)].

[0036] A summary of stream flow rates and energy consumption for the process illustrated in FIG. 2 is set forth in the following table:

Table II

(FIG. 2)

Stream Flow Summary - Lb. Moles/Hr [kg moles/Hr]

Stream	Methane	Ethane	Propane	Butanes+	Total
31	25,384	1,161	362	332	27,451
32	24,909	1,076	297	166	26,655
33	475	85	65	166	796
49	5,751	117	6	1	5,910
35	6,226	202	71	167	6,706
39	29,831	38	0	0	30,006
48	4,475	6	0	0	4,501
47	25,356	32	0	0	25,505
42	28	1,129	362	332	1,946

Recoveries*

Ethane	97.24%	
Propane	100.00%	
Butanes+	100.00%	

Power

Residue Gas Compression	12,667	HP	[20,825	kW]
Reflux Compression	664	HP	[1,092	kW]
Total Compression	13,331	HP	[21,917	kW]

* (Based on un-rounded flow rates)

[0037] Comparison of the recovery levels displayed in Tables I and II shows that the liquids recovery of the FIG. 2 process is essentially the same as that of the FIG. 1 process. However, the total power requirement for the FIG. 2 process is about 4% lower than that of the FIG. 1 process.

DESCRIPTION OF THE INVENTION

Example 1

[0038] FIG. 3 illustrates a flow diagram of a process in accordance with the present invention. The feed gas composition and conditions considered in the process presented in FIG. 3 are the same as those in FIGS. 1 and 2. Accordingly, the FIG. 3 process can be compared with that of the FIGS. 1 and 2 processes to illustrate the advantages of the present invention.

[0039] In the simulation of the FIG. 3 process, inlet gas enters the plant as stream **31** and is cooled in heat exchanger **10** by heat exchange with a portion (stream **46**) of cool distillation stream **39a** at -61°F [-52°C], the pumped demethanizer bottoms liquid (stream **42a**) at 91°F [33°C], demethanizer liquids (stream **41**) at 68°F [20°C], and demethanizer liquids (stream **40**) at -13°F [-25°C]. The cooled stream **31a** enters separator **11** at -34°F [-37°C] and 1025 psia [7,067 kPa(a)] where the vapor (stream **32**) is separated from the condensed liquid (stream **33**).

[0040] The vapor (stream **32**) from separator **11** is divided into two streams, **34** and **36**. Likewise, the liquid (stream **33**) from separator **11** is divided into two streams, **37** and **38**. Stream **34**, containing about 10% of the total vapor, is combined with stream

37, containing about 50% of the total liquid. The combined stream 35 then passes through heat exchanger 12 in heat exchange relation with cold distillation stream 39 at -137°F [-94°C] where it is cooled to substantial condensation. The resulting substantially condensed stream 35a at -133°F [-92°C] is then flash expanded through an appropriate expansion device, such as expansion valve 13, to the operating pressure (approximately 460 psia [3,172 kPa(a)]) of fractionation tower 17, cooling stream 35b to -135°F [-93°C] before it is supplied to fractionation tower 17 at a mid-column feed point.

[0041] The remaining 90% of the vapor from separator 11 (stream 36) enters a work expansion machine 14 in which mechanical energy is extracted from this portion of the high pressure feed. The machine 14 expands the vapor substantially isentropically to the tower operating pressure, with the work expansion cooling the expanded stream 36a to a temperature of approximately -103°F [-75°C]. The partially condensed expanded stream 36a is thereafter supplied as feed to fractionation tower 17 at a second mid-column feed point.

[0042] The remaining 50% of the liquid from separator 11 (stream 38) is flash expanded through an appropriate expansion device, such as expansion valve 16, to the operating pressure of fractionation tower 17. The expansion cools stream 38a to -65°F [-54°C] before it is supplied to fractionation tower 17 at a third mid-column feed point.

[0043] The recompressed and cooled distillation stream 39e is divided into two streams. One portion, stream 47, is the volatile residue gas product. The other portion, recycle stream 48, flows to heat exchanger 22 where it is cooled to -1°F [-18°C] (stream 48a) by heat exchange with a portion (stream 45) of cool distillation stream 39a. The cooled recycle stream then flows to exchanger 12 where it is cooled to -133°F [-92°C]

and substantially condensed by heat exchange with cold distillation stream **39**. The substantially condensed stream **48b** is then expanded through an appropriate expansion device, such as expansion valve **23**, to the demethanizer operating pressure, resulting in cooling of the total stream to -141°F [-96°C]. The expanded stream **48c** is then supplied to fractionation tower **17** as the top column feed. The vapor portion (if any) of stream **48c** combines with the vapors rising from the top fractionation stage of the column to form distillation stream **39**, which is withdrawn from an upper region of the tower.

[0044] A portion of the distillation vapor (stream **49**) is withdrawn from the lower region of absorbing section **17b** of fractionation tower **17** at -129°F [-90°C] and is compressed to an intermediate pressure of about 697 psia [4,804 kPa(a)] by reflux compressor **24**. The compressed stream **49a** flows to exchanger **12** where it is cooled to -133°F [-92°C] and substantially condensed by heat exchange with cold distillation column overhead stream **39**. The substantially condensed stream **49b** is then expanded through an appropriate expansion device, such as expansion valve **25**, to the demethanizer operating pressure, resulting in cooling of stream **49c** to a temperature of -137°F [-94°C], whereupon it is supplied to fractionation tower **17** at a fourth mid-column feed point.

[0045] The demethanizer in tower **17** is a conventional distillation column containing a plurality of vertically spaced trays, one or more packed beds, or some combination of trays and packing. The demethanizer tower consists of three sections: an upper separator section **17a** wherein the top feed is divided into its respective vapor and liquid portions, and wherein the vapor rising from the intermediate absorbing section **17b** is combined with the vapor portion (if any) of the top feed to form the cold demethanizer

overhead vapor (stream **39**); an intermediate absorbing (rectification) section **17b** that contains the trays and/or packing to provide the necessary contact between the vapor portion of the expanded stream **36a** rising upward and cold liquid falling downward to condense and absorb the C_2 components, C_3 components, and heavier components; and a lower, stripping (demethanizing) section **17c** that contains the trays and/or packing to provide the necessary contact between the liquids falling downward and the vapors rising upward. The demethanizing section **17c** also includes reboilers (such as the reboiler and side reboiler described previously) which heat and vaporize a portion of the liquids flowing down the column to provide the stripping vapors which flow up the column to strip the liquid product, stream **42**, of methane and lighter components.

[0046] Stream **36a** enters demethanizer **17** at a feed position located in the lower region of absorbing section **17b** of demethanizer **17**. The liquid portion of expanded stream **36a** commingles with liquids falling downward from the absorbing section **17b** and the combined liquid continues downward into the stripping section **17c** of demethanizer **17**. The vapor portion of expanded stream **36a** rises upward through absorbing section **17b** and is contacted with cold liquid falling downward to condense and absorb the C_2 components, C_3 components, and heavier components.

[0047] The expanded substantially condensed stream **49c** is supplied as cold liquid reflux to an intermediate region in absorbing section **17b** of demethanizer **17**, as is expanded substantially condensed stream **35b**. These secondary reflux streams absorb and condense most of the C_3 components and heavier components (as well as much of the C_2 components) from the vapors rising in the lower rectification region of absorbing section **17b** so that only a small amount of recycle (stream **48**) must be cooled,

condensed, subcooled, and flash expanded to produce the top reflux stream **48c** that provides the final rectification in the upper region of absorbing section **17b**. As the cold top reflux stream **48c** contacts the rising vapors in the upper region of absorbing section **17b**, it condenses and absorbs the C_2 components and any remaining C_3 components and heavier components from the vapors so that they can be captured in the bottom product (stream **42**) from demethanizer **17**.

[0048] In stripping section **17c** of demethanizer **17**, the feed streams are stripped of their methane and lighter components. The resulting liquid product (stream **42**) exits the bottom of tower **17** at 86°F [30°C], based on a typical specification of a methane to ethane ratio of 0.025:1 on a molar basis in the bottom product. Pump **19** delivers stream **42a** to heat exchanger **10** as described previously where it is heated to 116°F [47°C] (stream **42b**) before flowing to storage.

[0049] The distillation vapor stream forming the tower overhead (stream **39**) is warmed in heat exchanger **12** as it provides cooling to combined stream **35**, compressed distillation vapor stream **49a**, and recycle stream **48a** as described previously to form cool distillation stream **39a**. Distillation stream **39a** is divided into two portions (streams **45** and **46**), which are heated to 116°F [47°C] and 92°F [33°C], respectively, in heat exchanger **22** and heat exchanger **10**. Note that in all cases exchangers **10**, **22**, and **12** are representative of either a multitude of individual heat exchangers or a single multi-pass heat exchanger, or any combination thereof. (The decision as to whether to use more than one heat exchanger for the indicated heating services will depend on a number of factors including, but not limited to, inlet gas flow rate, heat exchanger size, stream temperatures, etc.) The heated streams recombine to form stream **39b** at 94°F [34°C]

which is then re-compressed in two stages, compressor **15** driven by expansion machine **14** and compressor **20** driven by a supplemental power source. After stream **39d** is cooled to 120°F [49°C] in discharge cooler **21** to form stream **39e**, recycle stream **48** is withdrawn as described earlier to form residue gas stream **47** which flows to the sales gas pipeline at 1040 psia [7,171 kPa(a)].

[0050] A summary of stream flow rates and energy consumption for the process illustrated in FIG. 3 is set forth in the following table:

Table III

(FIG. 3)

Stream Flow Summary - Lb. Moles/Hr [kg moles/Hr]

Stream	Methane	Ethane	Propane	Butanes+	Total
31	25,384	1,161	362	332	27,451
32	25,085	1,103	314	185	26,894
33	299	58	48	147	557
34	2,509	110	31	19	2,690
37	149	29	24	73	278
35	2,658	139	55	92	2,968
36	22,576	993	283	166	24,204
38	150	29	24	74	279
49	4,978	46	1	0	5,080
39	28,268	36	0	0	28,474
48	2,912	4	0	0	2,933
47	25,356	32	0	0	25,541
42	28	1,129	362	332	1,910

Recoveries*					
Ethane	97.21%				
Propane	99.99%				
Butanes+	100.00%				
Power					
Residue Gas Compression	11,841	HP	[	19,466	kW]
Reflux Compression	486	HP	[	799	kW]
Total Compression	12,327	HP	[	20,265	kW]

* (Based on un-rounded flow rates)

[0051] A comparison of Tables I, II, and III shows that, compared to the prior art processes, the present invention maintains essentially the same ethane recovery, propane recovery, and butanes+ recovery. However, comparison of Tables I, II, and III further shows that these yields were achieved with substantially lower horsepower requirements than those of the prior art processes. The total power requirement of the present invention is 11% lower than that of the FIG. 1 process and nearly 8% lower than that of the FIG. 2 process.

[0052] The key feature of the present invention is the supplemental rectification provided by reflux stream **49c** in conjunction with stream **35b**, which reduces the amount of C_2 components, C_3 components, and C_4+ components contained in the vapors rising in the upper region of absorbing section **17b**. Compare these two supplemental reflux streams in Table III with the single supplemental reflux stream, **35b**, in Table I for the FIG. 1 process. While the total supplemental reflux flow rate is about the same, the

amount of C_2+ components in these reflux streams for the FIG. 3 process is only about one-half of that of the FIG. 1 process, making these streams much more effective at rectifying the C_2+ components in the vapors rising up in the lower region of absorbing section **17b**. As a result, the methane recycle (stream **48**) that is used to create the top reflux stream for fractionation tower **17** can be significantly less for the FIG. 3 process compared to the FIG. 1 process while maintaining the desired C_2 component recovery level, reducing the horsepower requirements for residue gas compression. Also, with the supplemental reflux supplied in two separate streams, one of which (stream **49c**) has significantly lower concentrations of C_2+ components, it is possible to divide absorbing section **17b** into multiple rectification zones and thus increase its efficiency.

[0053] A further advantage provided by supplemental reflux stream **49c** is that it allows a reduction in the flow rate of supplemental reflux stream **35b**, so that there is a corresponding increase in the flow rate of stream **36** to work expansion machine **14**. This in turn provides a two-fold improvement in the process efficiency. First, with more flow to expansion machine **14**, the increase in power recovery increases the refrigeration generated by the process. Second, the greater power recovery means more power available to compressor **15**, reducing the external power consumption of compressor **20**.

[0054] Compared to the FIG. 2 process, the present invention not only provides better supplemental reflux streams, but a higher total supplemental reflux flow rate as well. Compare supplemental reflux streams **49c** and **35b** in Table III with the single supplemental reflux stream, **35b**, in Table II for the FIG. 2 process. The total supplemental reflux flow rate is about 20% higher for the present invention, and the amount of C_2+ components in these reflux streams is only about three-fourths of that of

the FIG. 2 process. As a result, the flow rate of the methane recycle (stream **48**) used as the top reflux stream for fractionation tower **17** in the FIG. 3 process is only two-thirds of that of the FIG. 2 process while maintaining the desired C_2 component recovery level, reducing the horsepower requirements for residue gas compression. Also, by supplying the supplemental reflux in two separate streams, one of which (stream **49c**) has significantly lower concentrations of C_2+ components, it is possible to divide absorbing section **17b** into multiple rectification zones and thus increase its efficiency.

[0055] Note that in the FIG. 2 process, the withdrawal location for distillation vapor stream **49** from fractionation tower **17** is below the mid-column feed point of expanded stream **32a**. For the present invention, the withdrawal location can be higher up on the column, such as above the mid-column feed point of expanded stream **36a** as in this example. As a result, distillation vapor stream **49** in the FIG. 3 process of the present invention can be subjected to more rectification, reducing the concentration of C_2+ components in the stream and improving its effectiveness as a reflux stream for absorbing section **17b**. The location for the withdrawal of distillation vapor stream **49** of the present invention must be evaluated for each application.

Example 2

[0056] FIG. 3 represents the preferred embodiment of the present invention for the temperature and pressure conditions shown because it typically requires the least equipment and the lowest capital investment. An alternative method of using the supplemental reflux streams for the column is shown in another embodiment of the present invention as illustrated in FIG. 4. The feed gas composition and conditions considered in the process presented in FIG. 4 are the same as those in FIGS. 1 through 3.

Accordingly, FIG. 4 can be compared with the FIGS. 1 and 2 processes to illustrate the advantages of the present invention, and can likewise be compared to the embodiment displayed in FIG. 3.

[0057] In the simulation of the FIG. 4 process, inlet gas enters the plant as stream **31** and is cooled in heat exchanger **10** by heat exchange with a portion (stream **46**) of cool distillation stream **39a** at -58°F [-50°C], the pumped demethanizer bottoms liquid (stream **42a**) at 93°F [34°C], demethanizer liquids (stream **41**) at 70°F [21°C], and demethanizer liquids (stream **40**) at -12°F [-24°C]. The cooled stream **31a** enters separator **11** at -31°F [-35°C] and 1025 psia [7,067 kPa(a)] where the vapor (stream **32**) is separated from the condensed liquid (stream **33**).

[0058] The vapor (stream **32**) from separator **11** is divided into two streams, **34** and **36**. Likewise, the liquid (stream **33**) from separator **11** is divided into two streams, **37** and **38**. Stream **34**, containing about 11% of the total vapor, is combined with stream **37**, containing about 50% of the total liquid. The combined stream **35** then passes through heat exchanger **12** in heat exchange relation with cold distillation stream **39** at -136°F [-94°C] where it is cooled to substantial condensation. The resulting substantially condensed stream **35a** at -132°F [-91°C] is then flash expanded through an appropriate expansion device, such as expansion valve **13**, to the operating pressure (approximately 465 psia [3,206 kPa(a)]) of fractionation tower **17**, cooling stream **35b** to -134°F [-92°C] before it is supplied to fractionation tower **17** at a mid-column feed point.

[0059] The remaining 89% of the vapor from separator **11** (stream **36**) enters a work expansion machine **14** in which mechanical energy is extracted from this portion of the high pressure feed. The machine **14** expands the vapor substantially isentropically to

the tower operating pressure, with the work expansion cooling the expanded stream **36a** to a temperature of approximately -99°F [-73°C]. The partially condensed expanded stream **36a** is thereafter supplied as feed to fractionation tower **17** at a second mid-column feed point.

[0060] The remaining 50% of the liquid from separator **11** (stream **38**) is flash expanded through an appropriate expansion device, such as expansion valve **16**, to the operating pressure of fractionation tower **17**. The expansion cools stream **38a** to -60°F [-51°C] before it is supplied to fractionation tower **17** at a third mid-column feed point.

[0061] The recompressed and cooled distillation stream **39e** is divided into two streams. One portion, stream **47**, is the volatile residue gas product. The other portion, recycle stream **48**, flows to heat exchanger **22** where it is cooled to -1°F [-18°C] (stream **48a**) by heat exchange with a portion (stream **45**) of cool distillation stream **39a**. The cooled recycle stream then flows to exchanger **12** where it is cooled to -132°F [-91°C] and substantially condensed by heat exchange with cold distillation stream **39**. The substantially condensed stream **48b** is then expanded through an appropriate expansion device, such as expansion valve **23**, to the demethanizer operating pressure, resulting in cooling of the total stream to -140°F [-96°C]. The expanded stream **48c** is then supplied to fractionation tower **17** as the top column feed. The vapor portion (if any) of stream **48c** combines with the vapors rising from the top fractionation stage of the column to form distillation stream **39**, which is withdrawn from an upper region of the tower.

[0062] A portion of the distillation vapor (stream **49**) is withdrawn from the lower region of the absorbing section of fractionation tower **17** at -129°F [-89°C] and is compressed to an intermediate pressure of about 697 psia [4,804 kPa(a)] by reflux

compressor **24**. The compressed stream **49a** flows to exchanger **12** where it is cooled to -132°F [-91°C] and substantially condensed by heat exchange with cold distillation column overhead stream **39**. The substantially condensed stream **49b** is then divided into two portions, streams **51** and **52**. The first portion, stream **51** containing about 90% of stream **49b**, is expanded through an appropriate expansion device, such as expansion valve **25**, to the demethanizer operating pressure, resulting in cooling of stream **51a** to a temperature of -136°F [-94°C], whereupon it is supplied to fractionation tower **17** at a fourth mid-column feed point as in the FIG. 3 embodiment of the present invention. The remaining portion, stream **52** containing about 10% of stream **49b**, is expanded through an appropriate expansion device, such as expansion valve **26**, to the demethanizer operating pressure, resulting in cooling of stream **52a** to a temperature of -136°F [-94°C], whereupon it is supplied to fractionation tower **17** at a fifth mid-column feed point, located below the feed point of stream **51a**.

[0063] In the stripping section of demethanizer **17**, the feed streams are stripped of their methane and lighter components. The resulting liquid product (stream **42**) exits the bottom of tower **17** at 88°F [31°C]. Pump **19** delivers stream **42a** to heat exchanger **10** as described previously where it is heated to 116°F [47°C] (stream **42b**) before flowing to storage.

[0064] The distillation vapor stream forming the tower overhead (stream **39**) is warmed in heat exchanger **12** as it provides cooling to combined stream **35**, compressed distillation vapor stream **49a**, and recycle stream **48a** as described previously to form cool distillation stream **39a**. Distillation stream **39a** is divided into two portions (streams **45** and **46**), which are heated to 116°F [47°C] and 92°F [33°C], respectively, in heat

exchanger **22** and heat exchanger **10**. The heated streams recombine to form stream **39b** at 94°F [35°C] which is then re-compressed in two stages, compressor **15** driven by expansion machine **14** and compressor **20** driven by a supplemental power source. After stream **39d** is cooled to 120°F [49°C] in discharge cooler **21** to form stream **39e**, recycle stream **48** is withdrawn as described earlier to form residue gas stream **47** which flows to the sales gas pipeline at 1040 psia [7,171 kPa(a)].

[0065] A summary of stream flow rates and energy consumption for the process illustrated in FIG. 4 is set forth in the following table:

Table IV

(FIG. 4)

Stream Flow Summary - Lb. Moles/Hr [kg moles/Hr]

Stream	Methane	Ethane	Propane	Butanes+	Total
31	25,384	1,161	362	332	27,451
32	25,118	1,109	318	190	26,943
33	266	52	44	142	508
34	2,838	125	36	21	3,045
37	133	26	22	71	254
35	2,971	151	58	92	3,299
36	22,280	984	282	169	23,898
38	133	26	22	71	254
49	4,902	50	1	0	5,000
51	4,412	45	1	0	4,500
52	490	5	0	0	500
39	28,490	36	0	0	28,702
48	3,134	4	0	0	3,157
47	25,356	32	0	0	25,545
42	28	1,129	362	332	1,906

Recoveries*					
Ethane	97.22%				
Propane	99.99%				
Butanes+	100.00%				
Power					
Residue Gas Compression	11,745	HP	[	19,309	kW]
Reflux Compression	465	HP	[	764	kW]
Total Compression	12,210	HP	[	20,073	kW]

* (Based on un-rounded flow rates)

[0066] A comparison of Tables III and IV shows that, compared to the FIG. 3 embodiment of the present invention, the FIG. 4 embodiment maintains essentially the same ethane recovery, propane recovery, and butanes+ recovery. However, comparison of Tables III and IV further shows that these yields were achieved using about 1% less horsepower than that required by the FIG. 3 embodiment. The drop in the power requirements for the FIG. 4 embodiment is mainly due to the slightly higher operating pressure of fractionation tower **17**, which is possible due to the better rectification in its absorbing section provided by introducing a portion of the supplemental reflux (stream **52a**) lower in the absorbing section. This effectively reduces the concentration of C_2+ components in the column liquids where expanded combined stream **35b** is introduced, thereby reducing the equilibrium concentrations of these heavier components in the vapors rising above this region of the absorbing section. The reduction in power requirements for this embodiment over that of the FIG. 3 embodiment must be evaluated

for each application relative to the slight increase in capital cost expected for the FIG. 4 embodiment compared to the FIG. 3 embodiment.

Example 3

[0067] An alternative method of generating the supplemental reflux streams for the column is shown in another embodiment of the present invention as illustrated in FIG. 5. The feed gas composition and conditions considered in the process presented in FIG. 5 are the same as those in FIGS. 1 through 4. Accordingly, FIG. 5 can be compared with the FIGS. 1 and 2 processes to illustrate the advantages of the present invention, and can likewise be compared to the embodiments displayed in FIGS. 3 and 4.

[0068] In the simulation of the FIG. 5 process, inlet gas enters the plant as stream **31** and is cooled in heat exchanger **10** by heat exchange with a portion (stream **46**) of cool vapor stream **43a** at -61°F [-52°C], the pumped demethanizer bottoms liquid (stream **42a**) at 92°F [33°C], demethanizer liquids (stream **41**) at 69°F [21°C], and demethanizer liquids (stream **40**) at -15°F [-26°C]. The cooled stream **31a** enters separator **11** at -35°F [-37°C] and 1025 psia [7,067 kPa(a)] where the vapor (stream **32**) is separated from the condensed liquid (stream **33**).

[0069] The vapor (stream **32**) from separator **11** is divided into two streams, **34** and **36**. Likewise, the liquid (stream **33**) from separator **11** is divided into two streams, **37** and **38**. Stream **34**, containing about 10% of the total vapor, is combined with stream **37**, containing about 50% of the total liquid. The combined stream **35** then passes through heat exchanger **12** in heat exchange relation with cold vapor stream **43** at -137°F [-94°C] where it is cooled to substantial condensation. The resulting substantially

condensed stream **35a** at -133°F [-91°C] is then flash expanded through an appropriate expansion device, such as expansion valve **13**, to the operating pressure (approximately 464 psia [3,199 kPa(a)]) of fractionation tower **17**, cooling stream **35b** to -134°F [-92°C] before it is supplied to fractionation tower **17** at a mid-column feed point.

[0070] The remaining 90% of the vapor from separator **11** (stream **36**) enters a work expansion machine **14** in which mechanical energy is extracted from this portion of the high pressure feed. The machine **14** expands the vapor substantially isentropically to the tower operating pressure, with the work expansion cooling the expanded stream **36a** to a temperature of approximately -102°F [-75°C]. The partially condensed expanded stream **36a** is thereafter supplied as feed to fractionation tower **17** at a second mid-column feed point.

[0071] The remaining 50% of the liquid from separator **11** (stream **38**) is flash expanded through an appropriate expansion device, such as expansion valve **16**, to the operating pressure of fractionation tower **17**. The expansion cools stream **38a** to -65°F [-54°C] before it is supplied to fractionation tower **17** at a third mid-column feed point.

[0072] The recompressed and cooled vapor stream **43e** is divided into two streams. One portion, stream **47**, is the volatile residue gas product. The other portion, recycle stream **48**, flows to heat exchanger **22** where it is cooled to -1°F [-18°C] (stream **48a**) by heat exchange with a portion (stream **45**) of cool vapor stream **43a**. The cooled recycle stream then flows to exchanger **12** where it is cooled to -133°F [-91°C] and substantially condensed by heat exchange with cold vapor stream **43**. The substantially condensed stream **48b** is then expanded through an appropriate expansion device, such as expansion valve **23**, to the demethanizer operating pressure, resulting in cooling of the

total stream to -140°F [-96°C]. The expanded stream **48c** is then supplied to fractionation tower **17** as the top column feed. The vapor portion (if any) of stream **48c** combines with the vapors rising from the top fractionation stage of the column to form distillation stream **39**, which is withdrawn from an upper region of the tower.

[0073] The distillation vapor stream forming the tower overhead (stream **39**) leaves fractionation tower **17** at -137°F [-94°C] and is divided into two portions, first and second vapor streams **44** and **43**, respectively. First vapor stream **44** is combined with a portion of the distillation vapor (stream **49**) withdrawn from the lower region of the absorbing section of fractionation tower **17** at -131°F [-90°C], and the combined vapor stream **50** is compressed to an intermediate pressure of about 723 psia [4,985 kPa(a)] by reflux compressor **24**. The compressed stream **50a** flows to exchanger **12** where it is cooled to -133°F [-91°C] and substantially condensed by heat exchange with the remaining portion (stream **43**) of cold distillation column overhead stream **39**. The substantially condensed stream **50b** is then expanded through an appropriate expansion device, such as expansion valve **25**, to the demethanizer operating pressure, resulting in cooling of stream **50c** to a temperature of -137°F [-94°C], whereupon it is supplied to fractionation tower **17** at a fourth mid-column feed point.

[0074] In the stripping section of demethanizer **17**, the feed streams are stripped of their methane and lighter components. The resulting liquid product (stream **42**) exits the bottom of tower **17** at 87°F [31°C]. Pump **19** delivers stream **42a** to heat exchanger **10** as described previously where it is heated to 116°F [47°C] (stream **42b**) before flowing to storage.

[0075] Second vapor stream **43** (the remaining portion of cold distillation column overhead stream **39**) is warmed in heat exchanger **12** as it provides cooling to combined stream **35**, compressed combined stream **50a**, and recycle stream **48a** as described previously to form cool second vapor stream **43a**. Second vapor stream **43a** is divided into two portions (streams **45** and **46**), which are heated to 116°F [47°C] and 94°F [34°C], respectively, in heat exchanger **22** and heat exchanger **10**. The heated streams recombine to form stream **43b** at 95°F [35°C] which is then re-compressed in two stages, compressor **15** driven by expansion machine 14 and compressor **20** driven by a supplemental power source. After stream **43d** is cooled to 120°F [49°C] in discharge cooler **21** to form stream **43e**, recycle stream **48** is withdrawn as described earlier to form residue gas stream **47** which flows to the sales gas pipeline at 1040 psia [7,171 kPa(a)].

[0076] A summary of stream flow rates and energy consumption for the process illustrated in FIG. 5 is set forth in the following table:

Table V

(FIG. 5)

Stream Flow Summary - Lb. Moles/Hr [kg moles/Hr]

Stream	Methane	Ethane	Propane	Butanes+	Total
31	25,384	1,161	362	332	27,451
32	25,079	1,102	313	184	26,886
33	305	59	49	148	565
34	2,508	110	31	19	2,689
37	152	29	24	74	282
35	2,660	139	55	93	2,971
36	22,571	992	282	165	24,197
38	153	30	25	74	283
39	28,589	36	0	0	28,800
44	572	1	0	0	576
49	4,869	35	1	0	4,950
50	5,441	36	1	0	5,526
43	28,017	35	0	0	28,224
48	2,661	3	0	0	2,681
47	25,356	32	0	0	25,543
42	28	1,129	362	332	1,908

Recoveries*					
Ethane	97.20%				
Propane	99.99%				
Butanes+	100.00%				
Power					
Residue Gas Compression	11,617	HP	[	19,098	kW]
Reflux Compression	550	HP	[	904	kW]
Total Compression	12,167	HP	[	20,002	kW]

* (Based on un-rounded flow rates)

[0077] A comparison of Tables III, IV, and V shows that, compared to the FIG. 3 and FIG. 4 embodiments of the present invention, the FIG. 5 embodiment maintains essentially the same ethane recovery, propane recovery, and butanes+ recovery. However, comparison of Tables III, IV, and V further shows that these yields were achieved using about 1% less horsepower than that required by the FIG. 3 embodiment, and slightly less horsepower than the FIG. 4 embodiment. The drop in the power requirements for the FIG. 5 embodiment is mainly due to the reduction in the flow rate of recycle stream **48**. This reduction in the flow rate of the top reflux to demethanizer **17** is possible because combining a portion (stream **44**) of the column overhead (stream **39**) with the portion of the distillation vapor (stream **49**) withdrawn from the lower region of the absorbing section of fractionation tower **17** significantly reduces the concentration of C_2+ components in reflux stream **50c**, providing better rectification in the absorbing section. This reduces the equilibrium concentrations of these heavier components in the vapors rising above this region of the absorbing section so that less rectification is

required by the top reflux stream. The reduction in power requirements for this embodiment over that of the FIG. 3 embodiment must be evaluated for each application relative to the slight increase in capital cost for the FIG. 5 embodiment compared to the FIG. 3 embodiment. The FIG. 5 embodiment may offer a slight advantage in capital cost compared to the FIG. 4 embodiment, in addition to the power reduction, but this must likewise be evaluated for each application.

Example 4

[0078] An alternative method of using the supplemental reflux streams for the column is shown in another embodiment of the present invention as illustrated in FIG. 6. The feed gas composition and conditions considered in the process presented in FIG. 6 are the same as those in FIGS. 1 through 5. Accordingly, FIG. 6 can be compared with the FIGS. 1 and 2 processes to illustrate the advantages of the present invention, and can likewise be compared to the embodiments displayed in FIGS. 3 through 5.

[0079] In the simulation of the FIG. 6 process, inlet gas enters the plant as stream **31** and is cooled in heat exchanger **10** by heat exchange with a portion (stream **46**) of cool vapor stream **43a** at -55°F [-49°C], the pumped demethanizer bottoms liquid (stream **42a**) at 93°F [34°C], demethanizer liquids (stream **41**) at 71°F [21°C], and demethanizer liquids (stream **40**) at -10°F [-24°C]. The cooled stream **31a** enters separator **11** at -31°F [-35°C] and 1025 psia [7,067 kPa(a)] where the vapor (stream **32**) is separated from the condensed liquid (stream **33**).

[0080] The vapor (stream **32**) from separator **11** is divided into two streams, **34** and **36**. Likewise, the liquid (stream **33**) from separator **11** is divided into two streams,

37 and 38. Stream 34, containing about 12% of the total vapor, is combined with stream 37, containing about 50% of the total liquid. The combined stream 35 then passes through heat exchanger 12 in heat exchange relation with cold vapor stream 43 at -136°F [-93°C] where it is cooled to substantial condensation. The resulting substantially condensed stream 35a at -132°F [-91°C] is then flash expanded through an appropriate expansion device, such as expansion valve 13, to the operating pressure (approximately 469 psia [3,234 kPa(a)]) of fractionation tower 17, cooling stream 35b to -134°F [-92°C] before it is supplied to fractionation tower 17 at a mid-column feed point.

[0081] The remaining 88% of the vapor from separator 11 (stream 36) enters a work expansion machine 14 in which mechanical energy is extracted from this portion of the high pressure feed. The machine 14 expands the vapor substantially isentropically to the tower operating pressure, with the work expansion cooling the expanded stream 36a to a temperature of approximately -99°F [-73°C]. The partially condensed expanded stream 36a is thereafter supplied as feed to fractionation tower 17 at a second mid-column feed point.

[0082] The remaining 50% of the liquid from separator 11 (stream 38) is flash expanded through an appropriate expansion device, such as expansion valve 16, to the operating pressure of fractionation tower 17. The expansion cools stream 38a to -59°F [-51°C] before it is supplied to fractionation tower 17 at a third mid-column feed point.

[0083] The recompressed and cooled vapor stream 43e is divided into two streams. One portion, stream 47, is the volatile residue gas product. The other portion, recycle stream 48, flows to heat exchanger 22 where it is cooled to -1°F [-18°C] (stream 48a) by heat exchange with a portion (stream 45) of cool vapor stream 43a. The cooled

recycle stream then flows to exchanger **12** where it is cooled to -132°F [-91°C] and substantially condensed by heat exchange with cold vapor stream **43**. The substantially condensed stream **48b** is then expanded through an appropriate expansion device, such as expansion valve **23**, to the demethanizer operating pressure, resulting in cooling of the total stream to -140°F [-95°C]. The expanded stream **48c** is then supplied to fractionation tower **17** as the top column feed. The vapor portion (if any) of stream **48c** combines with the vapors rising from the top fractionation stage of the column to form distillation stream **39**, which is withdrawn from an upper region of the tower.

[0084] The distillation vapor stream forming the tower overhead (stream **39**) leaves fractionation tower **17** at -136°F [-93°C] and is divided into two portions, first and second vapor streams **44** and **43**, respectively. First vapor stream **44** is combined with a portion of the distillation vapor (stream **49**) withdrawn from the lower region of the absorbing section of fractionation tower **17** at -128°F [-89°C], and the combined vapor stream **50** is compressed to an intermediate pressure of about 732 psia [5,047 kPa(a)] by reflux compressor **24**. The compressed stream **50a** flows to exchanger **12** where it is cooled to -132°F [-91°C] and substantially condensed by heat exchange with the remaining portion (stream **43**) of cold distillation column overhead stream **39**. The substantially condensed stream **50b** is then divided into two portions, streams **51** and **52**. The first portion, stream **51** containing about 90% of stream **50b**, is expanded through an appropriate expansion device, such as expansion valve **25**, to the demethanizer operating pressure, resulting in cooling of stream **51a** to a temperature of -136°F [-94°C], whereupon it is supplied to fractionation tower **17** at a fourth mid-column feed point as in the FIG. 5 embodiment of the present invention. The remaining portion, stream **52**

containing about 10% of stream **50b**, is expanded through an appropriate expansion device, such as expansion valve **26**, to the demethanizer operating pressure, resulting in cooling of stream **52a** to a temperature of -136°F [-94°C], whereupon it is supplied to fractionation tower **17** at a fifth mid-column feed point, located below the feed point of stream **51a**.

[0085] In the stripping section of demethanizer **17**, the feed streams are stripped of their methane and lighter components. The resulting liquid product (stream **42**) exits the bottom of tower **17** at 89°F [31°C]. Pump **19** delivers stream **42a** to heat exchanger **10** as described previously where it is heated to 116°F [47°C] (stream **42b**) before flowing to storage.

[0086] Second vapor stream **43** (the remaining portion of cold distillation column overhead stream **39**) is warmed in heat exchanger **12** as it provides cooling to combined stream **35**, compressed combined stream **50a**, and recycle stream **48a** as described previously to form cool second vapor stream **43a**. Second vapor stream **43a** is divided into two portions (streams **45** and **46**), which are heated to 116°F [47°C] and 94°F [34°C], respectively, in heat exchanger **22** and heat exchanger **10**. The heated streams recombine to form stream **43b** at 96°F [35°C] which is then re-compressed in two stages, compressor **15** driven by expansion machine **14** and compressor **20** driven by a supplemental power source. After stream **43d** is cooled to 120°F [49°C] in discharge cooler **21** to form stream **43e**, recycle stream **48** is withdrawn as described earlier to form residue gas stream **47** which flows to the sales gas pipeline at 1040 psia [7,171 kPa(a)].

[0087] A summary of stream flow rates and energy consumption for the process illustrated in FIG. 6 is set forth in the following table:

Table VI

(FIG. 6)

Stream Flow Summary - Lb. Moles/Hr [kg moles/Hr]

Stream	Methane	Ethane	Propane	Butanes+	Total
31	25,384	1,161	362	332	27,451
32	25,122	1,109	319	191	26,949
33	262	52	43	141	502
34	2,977	131	38	23	3,194
37	131	26	21	70	251
35	3,108	157	59	93	3,445
36	22,145	978	281	168	23,755
38	131	26	22	71	251
39	29,044	37	0	0	29,260
44	871	1	0	0	878
49	4,487	44	1	0	4,575
50	5,358	45	1	0	5,453
51	4,823	40	1	0	4,908
52	535	5	0	0	545
43	28,173	36	0	0	28,382
48	2,817	4	0	0	2,838
47	25,356	32	0	0	25,544
42	28	1,129	362	332	1,907

Recoveries*					
Ethane	97.22%				
Propane	99.99%				
Butanes+	100.00%				
Power					
Residue Gas Compression	11,488	HP	[	18,886	kW]
Reflux Compression	548	HP	[	901	kW]
Total Compression	12,036	HP	[	19,787	kW]

* (Based on un-rounded flow rates)

[0088] A comparison of Tables III, IV, V, and VI shows that, compared to the FIGS. 3 through 5 embodiments of the present invention, the FIG. 6 embodiment maintains essentially the same ethane recovery, propane recovery, and butanes+ recovery. However, comparison of Tables III, IV, V, and VI further shows that these yields were achieved using about 2% less horsepower than that required by the FIG. 3 embodiment, and about 1% less horsepower than the FIG. 4 and FIG. 5 embodiments. The drop in the power requirements for the FIG. 6 embodiment is mainly due to the slightly higher operating pressure of fractionation tower 17, which is possible due to the better rectification in its absorbing section provided by introducing a portion of the supplemental reflux (stream **52a**) lower in the absorbing section. This effectively reduces the concentration of C_2+ components in the column liquids where expanded combined stream **35b** is introduced, thereby reducing the equilibrium concentrations of these heavier components in the vapors rising above this region of the absorbing section. The reduction in power requirements for this embodiment over that of the FIGS. 3 through 5

embodiments must be evaluated for each application relative to the slight increase in capital cost for the FIG. 6 embodiment compared to the other embodiments.

Other Embodiments

[0089] In accordance with this invention, it is generally advantageous to design the absorbing (rectification) section of the demethanizer to contain multiple theoretical separation stages. However, the benefits of the present invention can be achieved with as few as one theoretical stage, and it is believed that even the equivalent of a fractional theoretical stage may allow achieving these benefits. For instance, all or a part of the expanded substantially condensed recycle stream **48c**, all or a part of the supplemental reflux (stream **49c** in FIG. 3, stream **50c** in FIG. 5, or streams **51a** and **52a** in FIGS. 4 and 6), all or a part of the expanded substantially condensed stream **35b**, and all or a part of the expanded stream **36a** can be combined (such as in the piping joining the expansion valve to the demethanizer) and if thoroughly intermingled, the vapors and liquids will mix together and separate in accordance with the relative volatilities of the various components of the total combined streams. Such commingling of the four or five streams shall be considered for the purposes of this invention as constituting an absorbing section. Specifically, commingling of supplemental reflux stream **52a** and expanded substantially condensed stream **35b** appears to be advantageous in many instances, as does commingling of the expanded substantially condensed recycle stream **48c** and all or a part of the supplemental reflux (stream **49c** in FIG. 3, stream **50c** in FIG. 5, or stream **51a** in FIGS. 4 and 6).

[0090] FIGS. 7 and 8 depict fractionation towers constructed in two vessels, absorber (rectifier) column 27 (a contacting and separating device) and stripper

(distillation) column **17**. In such cases, a portion of the distillation vapor (stream **49**) is withdrawn from the lower section of absorber column **27** and routed to reflux compressor **24** (optionally, as shown in FIG. 8, combined with a portion, stream **44**, of overhead distillation stream **39** from absorber column **27**) to generate supplemental reflux for absorber column **27**. The overhead vapor (stream **54**) from stripper column **17** flows to the lower section of absorber column **27** to be contacted by expanded substantially condensed recycle stream **48c**, supplemental reflux liquid (stream **51a** and optional stream **52a**), and expanded substantially condensed stream **35b**. Pump **28** is used to route the liquids (stream 55) from the bottom of absorber column **27** to the top of stripper column **17** so that the two towers effectively function as one distillation system. The decision whether to construct the fractionation tower as a single vessel (such as demethanizer **17** in FIGS. 3 through 6) or multiple vessels will depend on a number of factors such as plant size, the distance to fabrication facilities, etc.

[0091] As described in the earlier examples, the supplemental reflux (stream **49b** in FIGS. 3, 4, and 7 and stream **50b** in FIGS. 5, 6, and 8) is totally condensed and the resulting condensate used to absorb valuable C_2 components, C_3 components, and heavier components from the vapors rising through the lower region of absorbing section **17b** of demethanizer **17** (FIGS. 3 through 6) or through absorber column **27** (FIGS. 7 and 8). However, the present invention is not limited to this embodiment. It may be advantageous, for instance, to treat only a portion of these vapors in this manner, or to use only a portion of the condensate as an absorbent, in cases where other design considerations indicate portions of the vapors or the condensate should bypass absorbing section **17b** of demethanizer **17** (FIGS. 3 through 6) or absorber column **27** (FIGS. 7 and

8). Some circumstances may favor partial condensation, rather than total condensation, of the supplemental reflux stream (**49b** or **50b**) in heat exchanger **12**. Other circumstances may favor that distillation stream **49** be a total vapor side draw from fractionation column **17** (FIGS. 3 through 6) or absorber column **27** (FIGS. 7 and 8) rather than a partial vapor side draw. It should also be noted that, depending on the composition of the feed gas stream, it may be advantageous to use external refrigeration to provide some portion of the cooling of the supplemental reflux stream (**49b** or **50b**) in heat exchanger **12**.

[0092] Feed gas conditions, plant size, available equipment, or other factors may indicate that elimination of work expansion machine **14**, or replacement with an alternate expansion device (such as an expansion valve), is feasible. Although individual stream expansion is depicted in particular expansion devices, alternative expansion means may be employed where appropriate. For example, conditions may warrant work expansion of the substantially condensed recycle stream (stream **48b**), the supplemental reflux (stream **49b**, stream **50b**, or streams **51** and/or **52**), or the substantially condensed stream (stream **35a**).

[0093] When the inlet gas is leaner, separator **11** in FIGS. 3 through 8 may not be needed. Depending on the quantity of heavier hydrocarbons in the feed gas and the feed gas pressure, the cooled feed stream **31a** leaving heat exchanger **10** in FIGS. 3 through 8 may not contain any liquid (because it is above its dewpoint, or because it is above its cricondenbar), so that separator **11** shown in FIGS. 3 through 8 is not required. Additionally, even in those cases where separator **11** is required, it may not be advantageous to combine any of the resulting liquid in stream **33** with vapor stream **34**.

In such cases, all of the liquid would be directed to stream **38** and thence to expansion valve **16** and a lower mid-column feed point on demethanizer **17** (FIGS. 3 through 6) or a mid-column feed point on stripping column **17** (FIGS. 7 and 8). Other applications may favor combining all of the resulting liquid in stream **33** with vapor stream **34**. In such cases, there would be no flow in stream **38** and expansion valve **16** would not be required.

[0094] In accordance with this invention, the use of external refrigeration to supplement the cooling available to the inlet gas and/or the recycle gas from other process streams may be employed, particularly in the case of a rich inlet gas. The use and distribution of separator liquids and demethanizer side draw liquids for process heat exchange, and the particular arrangement of heat exchangers for inlet gas cooling must be evaluated for each particular application, as well as the choice of process streams for specific heat exchange services.

[0095] It will also be recognized that the relative amount of feed found in each branch of the split vapor feed and the split liquid feed will depend on several factors, including gas pressure, feed gas composition, the amount of heat which can economically be extracted from the feed, and the quantity of horsepower available. The relative locations of the mid-column feeds and the withdrawal point of distillation vapor stream **49** may vary depending on inlet composition or other factors such as desired recovery levels and amount of liquid formed during inlet gas cooling. In some circumstances, withdrawal of distillation vapor stream **49** below the feed location of expanded stream **36a** is favored. Moreover, two or more of the feed streams, or portions thereof, may be combined depending on the relative temperatures and quantities of individual streams,

and the combined stream then fed to a mid-column feed position. The intermediate pressure to which distillation stream **49** or combined vapor stream **50** is compressed must be determined for each application, as it is a function of inlet composition, the desired recovery level, the withdrawal point of distillation vapor stream **49**, and other factors.

[0096] While there have been described what are believed to be preferred embodiments of the invention, those skilled in the art will recognize that other and further modifications may be made thereto, e.g. to adapt the invention to various conditions, types of feed, or other requirements without departing from the spirit of the present invention as defined by the following claims.

WE CLAIM:

1. In a process for the separation of a gas stream containing methane, C_2 components, C_3 components, and heavier hydrocarbon components into a volatile residue gas fraction and a relatively less volatile fraction containing a major portion of said C_2 components, C_3 components, and heavier hydrocarbon components or said C_3 components and heavier hydrocarbon components, in which process

(a) said gas stream is cooled under pressure to provide a cooled stream;

(b) said cooled stream is expanded to a lower pressure whereby it is further cooled; and

(c) said further cooled stream is directed into a distillation column and fractionated at said lower pressure whereby the components of said relatively less volatile fraction are recovered;

the improvement wherein following cooling, at least a portion of said cooled stream is divided into first and second streams; and

(1) said first stream is cooled to condense substantially all of it and is thereafter expanded to said lower pressure whereby it is further cooled;

(2) said expanded cooled first stream is thereafter supplied to said distillation column at a first mid-column feed position;

(3) said second stream is expanded to said lower pressure and is supplied to said distillation column at a second mid-column feed position;

(4) a distillation vapor stream is withdrawn from a region of said distillation column below said first mid-column feed position and is compressed to an intermediate pressure;

(5) said compressed distillation vapor stream is cooled sufficiently to condense at least a part of it, thereby forming a condensed stream;

(6) at least a portion of said condensed stream is expanded to said lower pressure and is thereafter supplied to said distillation column at a third mid-column feed position located above said first mid-column feed position;

(7) an overhead vapor stream is withdrawn from an upper region of said distillation column and at least a portion of said overhead vapor stream is directed into heat exchange relation with said compressed distillation vapor stream and heated, thereby to supply at least a portion of the cooling of step (5);

(8) said heated overhead vapor stream is compressed to higher pressure and thereafter divided into said volatile residue gas fraction and a compressed recycle stream;

(9) said compressed recycle stream is cooled sufficiently to substantially condense it;

(10) said substantially condensed compressed recycle stream is expanded to said lower pressure and supplied to said distillation column at a top feed position; and

(11) the quantities and temperatures of said feed streams to said distillation column are effective to maintain the overhead temperature of said distillation column

at a temperature whereby the major portions of the components in said relatively less volatile fraction are recovered.

2. The process according to claim 1, wherein said cooled stream is cooled sufficiently to partially condense it; and

(1) said partially condensed gas stream is separated thereby to provide a vapor stream and at least one liquid stream;

(2) said vapor stream is thereafter divided into said first and second streams; and

(3) at least a portion of said at least one liquid stream is expanded to said lower pressure and is supplied to said distillation column at a fourth mid-column feed position.

3. The process according to claim 2, wherein

(1) prior to being supplied to said distillation column, said first stream is combined with at least a portion of said at least one liquid stream to form a combined stream, and said combined stream is cooled to condense substantially all of it and is thereafter expanded to said lower pressure whereby it is further cooled;

(2) said expanded cooled combined stream is thereafter supplied to said distillation column at said first mid-column feed position; and

(3) any remaining portion of said at least one liquid stream is expanded to said lower pressure and is supplied to said distillation column at said fourth mid-column feed position.

4. In a process for the separation of a gas stream containing methane, C_2 components, C_3 components, and heavier hydrocarbon components into a volatile residue gas

fraction and a relatively less volatile fraction containing a major portion of said C_2 components, C_3 components, and heavier hydrocarbon components or said C_3 components and heavier hydrocarbon components, in which process

(a) said gas stream is cooled under pressure to provide a cooled stream;

(b) said cooled stream is expanded to a lower pressure whereby it is further cooled; and

(c) said further cooled stream is directed into a distillation column and fractionated at said lower pressure whereby the components of said relatively less volatile fraction are recovered;

the improvement wherein following cooling, at least a portion of said cooled stream is divided into first and second streams; and

(1) said first stream is cooled to condense substantially all of it and is thereafter expanded to said lower pressure whereby it is further cooled;

(2) said expanded cooled first stream is thereafter supplied at a first mid-column feed position to a contacting and separating device that produces a first overhead vapor stream and a bottom liquid stream, whereupon said bottom liquid stream is supplied to said distillation column;

(3) a second overhead vapor stream is withdrawn from an upper region of said distillation column and is directed to said contacting and separating device at a first lower feed position;

(4) said second stream is expanded to said lower pressure and is supplied to said contacting and separating device at a second lower feed position;

(5) a distillation vapor stream is withdrawn from a region of said contacting and separating device below said first mid-column feed position and is compressed to an intermediate pressure;

(6) said compressed distillation vapor stream is cooled sufficiently to condense at least a part of it, thereby forming a condensed stream;

(7) at least a portion of said condensed stream is expanded to said lower pressure and is thereafter supplied to said contacting and separating device at a second mid-column feed position located above said first mid-column feed position;

(8) at least a portion of said first overhead vapor stream is directed into heat exchange relation with said compressed distillation vapor stream and heated, thereby to supply at least a portion of the cooling of step (6);

(9) said heated first overhead vapor stream is compressed to higher pressure and thereafter divided into said volatile residue gas fraction and a compressed recycle stream;

(10) said compressed recycle stream is cooled sufficiently to substantially condense it;

(11) said substantially condensed compressed recycle stream is expanded to said lower pressure and supplied to said contacting and separating device at a top feed position; and

(12) the quantities and temperatures of said feed streams to said contacting and separating device are effective to maintain the overhead temperature of said contacting and separating device at a temperature whereby the major portions of the components in said relatively less volatile fraction are recovered.

5. The process according to claim 4, wherein said cooled stream is cooled sufficiently to partially condense it; and

(1) said partially condensed gas stream is separated thereby to provide a vapor stream and at least one liquid stream;

(2) said vapor stream is thereafter divided into said first and second streams; and

(3) at least a portion of said at least one liquid stream is expanded to said lower pressure and is supplied to said distillation column at a mid-column feed position.

6. The process according to claim 5, wherein

(1) prior to being supplied to said contacting and separating device, said first stream is combined with at least a portion of said at least one liquid stream to form a combined stream, and said combined stream is cooled to condense substantially all of it and is thereafter expanded to said lower pressure whereby it is further cooled;

(2) said expanded cooled combined stream is thereafter supplied at said first mid-column feed position to said contacting and separating device; and

(3) any remaining portion of said at least one liquid stream is expanded to said lower pressure and is supplied to said distillation column at said mid-column feed position.

7. The improvement according to claim 1, 2, or 3 wherein

(1) said overhead vapor stream is divided into at least a first vapor stream and a second vapor stream;

(2) said first vapor stream is combined with said distillation vapor stream to form a combined vapor stream, whereupon said combined vapor stream is compressed to said intermediate pressure;

(3) said compressed combined vapor stream is cooled sufficiently to condense at least a part of it, thereby forming said condensed stream;

(4) said second vapor stream is directed into heat exchange relation with said compressed combined stream and heated, thereby to supply at least a portion of the cooling of step (3); and

(5) said heated second vapor stream is compressed to said higher pressure and thereafter divided into said volatile residue gas fraction and said compressed recycle stream.

8. The improvement according to claim 4, 5, or 6 wherein

(1) said first overhead vapor stream is divided into at least a first vapor stream and a second vapor stream;

(2) said first vapor stream is combined with said distillation vapor stream to form a combined vapor stream, whereupon said combined vapor stream is compressed to said intermediate pressure;

(3) said compressed combined vapor stream is cooled sufficiently to condense at least a part of it, thereby forming said condensed stream;

(4) said second vapor stream is directed into heat exchange relation with said compressed combined stream and heated, thereby to supply at least a portion of the cooling of step (3); and

(5) said heated second vapor stream is compressed to said higher pressure and thereafter divided into said volatile residue gas fraction and said compressed recycle stream.

9. The improvement according to claim 1, 2, or 3 wherein

(1) said condensed stream is divided into at least a first portion and a second portion;

(2) said first portion is expanded to said lower pressure and is thereafter supplied to said distillation column at said third mid-column feed position; and

(3) said second portion is expanded to said lower pressure and is thereafter supplied to said distillation column at a mid-column feed position below that of said expanded first portion.

10. The improvement according to claim 7 wherein

(1) said condensed stream is divided into at least a first portion and a second portion;

(2) said first portion is expanded to said lower pressure and is thereafter supplied to said distillation column at said third mid-column feed position; and

(3) said second portion is expanded to said lower pressure and is thereafter supplied to said distillation column at a mid-column feed position below that of said expanded first portion.

11. The improvement according to claim 4, 5, or 6 wherein

(1) said condensed stream is divided into at least a first portion and a second portion;

(2) said first portion is expanded to said lower pressure and is thereafter supplied to said contacting and separating device at said second mid-column feed position; and

(3) said second portion is expanded to said lower pressure and is thereafter supplied to said contacting and separating device at a mid-column feed position below that of said expanded first portion.

12. The improvement according to claim 8 wherein

(1) said condensed stream is divided into at least a first portion and a second portion;

(2) said first portion is expanded to said lower pressure and is thereafter supplied to said contacting and separating device at said second mid-column feed position; and

(3) said second portion is expanded to said lower pressure and is thereafter supplied to said contacting and separating device at a mid-column feed position below that of said expanded first portion.

13. The improvement according to claim 1, 2, or 3 wherein said at least a portion of said expanded condensed stream is combined with said expanded substantially condensed compressed recycle stream to form a combined condensed stream, whereupon said combined condensed stream is supplied to said distillation column at said top feed position.

14. The improvement according to claim 4, 5, or 6 wherein said at least a portion of said expanded condensed stream is combined with said expanded substantially condensed compressed recycle stream to form a combined condensed stream, whereupon said

combined condensed stream is supplied to said contacting and separating device at said top feed position.