A novel high flux, cross-linked, fumed silica reinforced polyorganosiloxane thin film composite (TFC) membrane comprising a selective layer of a high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer supported by a porous support membrane formed from a glassy polymer has been developed. The novel high flux, cross-linked, fumed silica reinforced polyorganosiloxane thin film composite (TFC) membrane may be used to separate at least one component from another.
HIGH FLUX, CROSS-LINKED, FUMED SILICA REINFORCED PQLYORGANQSILQXANE MEMBRANES FOR SEPARATIONS

STATEMENT OF PRIORITY

[0001] This application claims priority to U.S. Application No. 62/423,667 which was filed November 17, 2016, the contents of which are hereby incorporated by reference in its entirety.

BACKGROUND OF THE INVENTION

[0002] Over 170 Honeywell UOP Separex™ membrane systems have been installed in the world for gas separation applications such as for the removal of acid gases from natural gas, in enhanced oil recovery, and hydrogen purification. Two new Separex™ membranes (Flux+ and Select) have been commercialized recently by Honeywell UOP, Des Plaines, IL for carbon dioxide (CO2) removal from natural gas. These Separex™ spiral wound membrane systems currently hold the membrane market leadership for natural gas upgrading. These membranes prepared from glassy polymers, however, do not have outstanding performance for organic vapor separations such as for olefin recovery, liquefied petroleum gas (LPG) recovery, fuel gas conditioning, natural gas dew point control, nitrogen removal from natural gas, etc.

[0003] Polymeric membrane materials have been found to be of use in gas separations. Numerous research articles and patents describe glassy polymeric membrane materials (e.g., polyimides, polysulfones, polycarbonates, polyamides, polycrylates, polypyrrolones) with desirable gas separation properties, particularly for use in oxygen/nitrogen separation (see, for example, U.S. 6,932,589). The polymeric membrane materials are typically used in processes in which a feed gas mixture contacts the upstream side of the membrane, resulting in a permeate mixture on the downstream side of the membrane with a greater mole fraction of one of the components than the composition of the original feed gas mixture. A pressure differential is maintained between the upstream and downstream sides, providing the driving force for permeation. The downstream side can be maintained as a vacuum, or at any pressure below the upstream pressure.
[0004] The separation of a polymeric membrane is based on a solution-diffusion mechanism. This mechanism involves molecular-scale interactions of the permeating gas with the polymer. The mechanism assumes that in a membrane having two opposing surfaces, each component is sorbed by the membrane at one surface, transported by a gas concentration gradient, and desorbed at the opposing surface. According to this solution-diffusion model, the membrane performance in separating a given pair of gases (e.g., CO2 /CH4, O2/N2, H2/CH4) is determined by two parameters: the permeability coefficient (abbreviated hereinafter as permeability or PA) and the selectivity (OCA/B). The PA is the product of the gas flux and the selective skin layer thickness of the membrane, divided by the pressure difference across the membrane. The OCA/B is the ratio of the permeability coefficients of the two gases (\( \frac{C_{A/B}}{C_{A/B}} = \frac{P_{A}}{P_{B}} \)) where P A is the permeability of the more permeable gas and P B is the permeability of the less permeable gas. Gases can have high permeability coefficients because of a high solubility coefficient, a high diffusion coefficient, or because both coefficients are high. In general, the diffusion coefficient decreases while the solubility coefficient increases with an increase in the molecular size of the gas. In high performance polymer membranes, both high permeability and selectivity are desirable because higher permeability decreases the size of the membrane area required to treat a given volume of gas, thereby decreasing capital cost of membrane units, and because higher selectivity results in a higher purity product gas.

[0005] The relative ability of a membrane to achieve the desired separation is referred to as the separation factor or selectivity for the given mixture. There are, however, several other obstacles to use a particular polymer to achieve a particular separation under any sort of large scale or commercial conditions. One such obstacle is permeation rate or flux. One of the components to be separated must have a sufficiently high permeation rate at the preferred conditions or extraordinarily large membrane surface areas are required to allow separation of large amounts of material. Therefore, commercially available glassy polymeric membranes, such as CA, polyimide, and polysulfone membranes formed by phase inversion and solvent exchange methods have an asymmetric integrally skinned membrane structure. See U.S. 3,133,132. Such membranes are characterized by a thin, dense, selectively semipermeable surface "skin" and a less dense void-containing (or porous), non-selective support region, with pore sizes ranging from large in the support region to very small proximate to the "skin". Plasticization occurs when one or more of the components of the mixture act as a
solvent in the polymer often causing it to swell and lose its membrane properties. It has been found that glassy polymers such as cellulose acetate and polyimides which have particularly good separation factors for separation of mixtures comprising carbon dioxide and methane are prone to plasticization over time thus resulting in decreasing performance of these membranes.

[0006] Natural gas often contains substantial amounts of heavy hydrocarbons and water, either as an entrained liquid, or in vapor form, which may lead to condensation within membrane modules. The gas separation capabilities of glassy polymeric membranes are affected when contacting with liquids including water and aromatic hydrocarbons such as benzene, toluene, ethylbenzene, and xylene (BTEX). The presence of more than modest levels of liquid BTEX heavy hydrocarbons is potentially damaging to traditional glassy polymeric membrane. Therefore, precautions must be taken to remove the entrained liquid water and heavy hydrocarbons upstream of the glassy polymeric membrane separation steps using expensive membrane pretreatment system. Another issue of glassy polymeric polymer membranes that still needs to be addressed for their use in gas separations in the presence of high concentration of condensable gas or vapor such as CO2 and propylene is the plasticization of the glassy polymer by these condensable gases or vapors that leads to swelling of the membrane as well as a significant increase in the permeance of all components in the feed and a decrease in the selectivity of the membranes.

[0007] Some natural gas also contains substantial amount of nitrogen (N2) in addition to the heavy hydrocarbons, water, and acid gases such as CO2 and hydrogen sulfide (H2S). Traditional glassy polymeric membranes are relatively more permeable to N2 than to methane. These membranes, however, have low N2 permeance and low N2/CH4 selectivity of <5.

[0008] For glassy polymeric gas separation membranes, permeant diffusion coefficient is more important than its solubility coefficient. Therefore, these glassy polymeric gas separation membranes preferentially permeate the smaller, less condensable gases, such as H2 and CH4 over the larger, more condensable gases, such as C3¾ and CO2. On the other hand, in rubbery polymeric membranes such as polydimethylsiloxane membrane, permeant solubility coefficients are much more important than diffusion coefficient. Thus, these rubbery polymeric membranes preferentially permeate the larger, more condensable gases...
over the smaller, less condensable gases. PDMS is the most commonly used rubbery membrane material for separation of higher hydrocarbons or methane from permanent gases such as N₂ and H₂.

[0009] Most of the polyolefn such as polypropylene (PP) and polyethylene (PE) manufacturing plants and other polymer such as polyvinyl chloride (PVC) manufacturing plants use a degassing step to remove un-reacted olefins, solvents, and other additives from the raw polyolefn. Nitrogen is normally used as the stripping gas or for the polymer transfer. Disposing of the vent stream in a flare or partial recovery of the valuable olefin or other monomers via a condensing process results in the loss of valuable monomers and undesired emissions of the highly reactive volatile monomers into the air. Typically, the vent stream of the polymer reactor is compressed and then cooled to condense the monomers such as propylene and ethylene from the PP and PE reactors. The gas leaving the condenser still contains a significant amount of the monomers. One application for rubbery polymeric membranes is to recover the valuable monomers such as propylene, ethylene, and vinyl chloride and purify nitrogen for reuse from the vent stream. For olefin splitter overhead applications, the stream leaving the column overhead is primarily olefins, mixed with light gases such as N₂ or H₂. The membrane can separate the stream into an olefin-enriched stream and a light-gas-enriched stream. The olefin-enriched stream is returned to the distillation column, where the high value olefin is recovered, and the light-gas-enriched stream is vented or flared. The condensation/membrane hybrid process will achieve significantly higher olefin recovery than condensation process alone and also allows olefin recovery at moderate temperatures and pressures than condensation process.

[0010] Ethylene recovery during the ethylene oxide (EO) production process to prevent the loss of valuable ethylene feedstock is another potential application of rubbery polymeric membranes. The rubbery polymeric membrane separates ethylene from argon purge gas by permeating ethylene at a much faster rate than argon to generate ethylene-enriched permeate that will be returned to the EO reactor and argon-enriched residue that will be flared.

[0011] The rubbery polymeric membrane can also be used for fuel gas conditioning that will reduce heavier hydrocarbons and increase CH₄ content (methane number) in the fuel gas which will be used to power upstream oil and gas operations while maintaining the pressure of the tail gas. Glassy polymeric membranes normally have very low methane permeance and also relatively low methane/heavy hydrocarbon selectivities.
SUMMARY OF THE INVENTION

[0012] This invention discloses a new high flux, cross-linked, fumed silica reinforced polyorganosiloxane thin film composite (TFC) membrane comprising a thin selective layer of a high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer on top of a porous glassy polymeric support membrane formed from a glassy polymer such as poliyethersulfone (PES), polysulfone (PSF), polyimide (PI), a blend of PES and PI, a blend of PSF and PI, and a blend of cellulose acetate (CA) and cellulose triacetate (CTA). The high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer is formed from addition cure (or hydrosilylation reaction) between a fumed silica reinforced vinyl-terminated polyorganosiloxane polymer and a methylhydrosiloxane-dimethylsiloxane cross-linking copolymer or between a mixture of a fumed silica reinforced vinyl-terminated polyorganosiloxane polymer and a vinylorganosiloxane-dimethylsiloxane copolymer and a methylhydrosiloxane-dimethylsiloxane cross-linking copolymer in the presence of a platinum complex catalyst. The present invention also discloses a method of making such a new type of high flux, cross-linked, fumed silica reinforced polyorganosiloxane TFC membrane, and the use of such a membrane for nitrogen removal from natural gas, fuel gas conditioning, olefin recovery from polyolefin production process, LPG recovery, and natural gas dew point control.

[0013] Different from glassy polymeric membranes that are highly selective to gases with smaller kinetic diameters over larger diameter gases, the new high flux, cross-linked, fumed silica reinforced polyorganosiloxane TFC membrane comprising a thin selective layer of a high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer on top of a porous glassy polymeric support membrane formed from a glassy polymer disclosed in the present invention is highly selective to olefins and heavier hydrocarbons over methane and inert gases such as N₂ and ¾ . Opposite from glassy polymeric membranes, the new high flux, cross-linked, fumed silica reinforced polyorganosiloxane TFC membrane described in the current invention has improved permeance and selectivity with the increase of operating time due to the increase of plasticization of condensable olefins on the membrane or with the decrease of operating temperature. In addition, the new high flux, cross-linked, fumed silica reinforced polyorganosiloxane TFC membrane described in the current invention has shown comparable selectivities but significantly higher permeance of C¾ for CH₄/N₂ separation, significantly higher permeances of olefins and paraffins for olefin and LPG recovery and fuel.
gas conditioning applications than those of polydimethylsiloxane rubbery polymeric membrane.

[0014] The porous glassy polymeric support membrane formed from a glassy polymer such as PES, PSF, PI, a blend of PES and PI, a blend of PSF and PI, and a blend of CA and CTA used for the preparation of the new high flux, cross-linked, fumed silica reinforced polyorganosiloxane TFC membrane disclosed in the present invention is fabricated using a phase inversion process by casting the glassy polymer solution using a casting knife. The porous glassy polymeric support membrane can be either a flat sheet support membrane or a hollow fiber support membrane. The solvents used for dissolving the glassy polymer material for the preparation of the porous glassy polymeric support membrane are chosen primarily for their ability to completely dissolve the polymers, ease of solvent removal in the membrane formation steps, and their function for the formation of pores on the skin layer of the support membrane. Other considerations in the selection of solvents include low toxicity, low corrosive activity, low environmental hazard potential, availability and cost.

Representative solvents include most amide solvents that are typically used for the formation of the porous glassy polymeric support membrane, such as N-methylpyrrolidone (NMP) and N,N-dimethyl acetamide (DMAc), methylene chloride, tetrahydrofuran (THF), acetone, methyl acetate, isopropanol, n-octane, n-hexane, n-decane, methanol, ethanol, N,N-dimethylformamide (DMF), dimethyl sulfoxide (DMSO), lactic acid, citric acid, dioxanes, 1,3-dioxolane, glycerol, mixtures thereof, others known to those skilled in the art and mixtures thereof. Preferably, the solvents used for dissolving the glassy polymer material for the preparation of the porous glassy polymeric support membrane in the current invention include NMP, 1,3-dioxolane, glycerol, and n-decane.

[0015] The thin selective layer of the high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer described in the present invention is formed on top of the porous glassy polymeric support membrane by applying a dilute hydrocarbon solution of a mixture of a fumed silica reinforced vinyl-terminated polyorganosiloxane polymer and a methylhydroxilosiloxane-dimethylsiloxane cross-linking copolymer in the presence of a platinum complex catalyst or a mixture of a fumed silica reinforced vinyl-terminated polyorganosiloxane polymer, a vinylorganosiloxane-dimethylsiloxane copolymer, and a methylhydroxilosiloxane-dimethylsiloxane cross-linking copolymer in the presence of a platinum complex catalyst to the top surface of the porous glassy polymeric support membrane by dip-
coating, spin coating, casting, soaking, spraying, painting, and other known conventional solution coating technologies. The thin selective layer of the high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer is formed by hydrosilylation reaction between the vinyl groups on the fumed silica reinforced vinyl-terminated polyorganosiloxane polymer or/and on the vinylorganosiloxane-dimethylsiloxane copolymer and the silicon hydride groups on the methylhydrosiloxane-dimethylsiloxane cross-linking copolymer after evaporating the hydrocarbon organic solvent(s) and heating at 70° to 150°C for a certain time.

[0016] Permeation experimental results demonstrate that the new high flux, cross-linked, fumed silica reinforced polyorganosiloxane TFC membrane comprising a thin selective layer of a high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer on top of a porous glassy polymeric support membrane disclosed in the present invention has higher permeance for paraffins such as ethane, propane, n-butane, and olefins such as propylene, n-butene, ethylene than inert gases such as N₂ and H₂ as well as CH₄ and has significantly higher permeances for paraffins such as ethane, propane, n-butane, and olefins such as propylene, n-butene, ethylene than those of the thermally cross-linked RTV615A/B silicone rubber membrane and UV cross-linked epoxysilicone rubbery membrane for olefin and N₂ recovery’, LPG recovery, and fuel gas conditioning applications.

[0017] This invention discloses the use of single stage or multi-stage new high flux, cross-linked, fumed silica reinforced polyorganosiloxane TFC membrane comprising a thin selective layer of a high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer on top of a porous glassy polymeric support membrane described in the current invention for olefin recovery, LPG recovery, fuel gas conditioning, natural gas dew point control, nitrogen removal from natural gas, etc.

DETAILED DESCRIPTION OF THE INVENTION

[0018] Membrane technology has been of great interest for the separation of gas, vapor, and liquid mixtures. However, despite significant research effort on separations by membrane technology, new rubbery polymeric membranes with improved performance are still needed for separations such as for olefin recovery, LPG recovery, fuel gas conditioning, natural gas dew point control, and nitrogen removal from natural gas.
This invention discloses a new type of high flux, cross-linked, fumed silica reinforced polyorganosiloxane thin film composite (TFC) membrane comprising a thin selective layer of a high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer on top of a porous glassy polymeric support membrane formed from a glassy polymer such as polyethersiloxane (PES), polysiloxane (PSF), polyimide (PI), a blend of PES and PI, a blend of PSF and PI, and a blend of cellulose acetate (CA) and cellulose triacetate (CTA). The high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer is formed from addition cure (or hydrosilylation reaction) between a fumed silica reinforced vinyl-terminated polyorganosiloxane polymer and a methylhydrosiloxane-dimethylsiloxane cross-linking copolymer or between a mixture of a fumed silica reinforced vinyl-terminated polyorganosiloxane polymer and a vinylorganosiloxane-dimethylsiloxane copolymer and a methylhydrosiloxane-dimethylsiloxane cross-linking copolymer in the presence of a platinum complex catalyst.

The present invention also discloses a method of making such a high flux, cross-linked, fumed silica reinforced polyorganosiloxane TFC membrane, and the use of such a membrane for olefin recovery from polyolefin production process, LPG recovery, fuel gas conditioning, natural gas dew point control, and nitrogen removal from natural gas.

Different from glassy polymeric membranes that are highly selective to gases with smaller kinetic diameters over larger diameter gases, the new high flux, cross-linked, fumed silica reinforced polyorganosiloxane TFC membrane comprising a thin selective layer of a high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer on top of a porous glassy polymeric support membrane formed from a glassy polymer disclosed in the present invention is highly selective to olefins and heavier hydrocarbons over methane and inert gases such as N₂ and H₂. Opposite from glassy polymeric membranes, the new high flux, cross-linked, fumed silica reinforced polyorganosiloxane TFC membrane described in the current invention has improved permeance and selectivity with the increase of operating time due to the increase of plasticization of condensable olefins on the membrane or with the decrease of operating temperature. In addition, the new high flux, cross-linked, fumed silica reinforced polyorganosiloxane TFC membrane described in the current invention has shown comparable selectivities but significantly higher permeance of C3/4 for CH₄/N₂ separation, significantly higher permeances of olefins and paraffins for olefin and LPG recovery and fuel gas conditioning applications than those of the polydimethylsiloxane rubbery membrane, the
thermally cross-linked RTV615A/B silicone rubber membrane, and UV cross-linked epoxysiicone rubbery membrane. 

[0022] The porous glassy polymeric support membrane can be formed from any glassy polymer that has good film forming properties such as PES, PSF, PI, a blend of PES and PI, a blend of PSF and PI, and a blend of CA and CTA used for the preparation of the new high flux, cross-linked, fumed silica reinforced polyorganosiloxane TFC membrane disclosed in the present invention is fabricated using a phase inversion process by casting the glassy polymer solution using a casting knife. The porous glassy polymeric support membrane described in the current invention can be either an asymmetric integrally skinned membrane or a TFC membrane with either flat sheet (spiral wound) or hollow fiber geometry.

[0023] The current invention discloses the use of a porous glassy polymeric support membrane for the preparation of the new high flux, cross-linked, fumed silica reinforced polyorganosiloxane TFC membrane by coating a thin selective layer of a high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer on top of the porous glassy polymeric support membrane. The porous glassy polymeric support membrane used for the preparation of the new high flux, cross-linked, fumed silica reinforced polyorganosiloxane TFC membrane described in the present invention has a carbon dioxide permeance of at least 100 GPU and no carbon dioxide/methane selectivity at 50 °C under 20-100 psig 10% CO₂/90% CH₄ mixed gas feed pressure.

[0024] The solvents used for dissolving the glassy polymer material for the preparation of the porous glassy polymeric support membrane are chosen primarily for their ability to completely dissolve the polymers, ease of solvent removal in the membrane formation steps, and their function for the formation of small pores on the skin layer of the support membrane. Other considerations in the selection of solvents include low toxicity, low corrosive activity, low environmental hazard potential, availability and cost. Representative solvents include most amide solvents that are typically used for the formation of the porous glassy polymeric support membrane, such as N-methylpyrrolidone (NMP) and N,N-dimethyl acetamide (DMAc), methylene chloride, tetrahydrofuran (THF), acetone, methyl acetate, isopropanol, n-octane, n-hexane, n-deene, methanol, ethanol, N,N-dimethylformamide (DMF), dimethyl sulfoxide (DMSO), lactic acid, citric acid, dioxanes, 1,3-dioxoiane, glycerol, mixtures thereof, others known to those skilled in the art and mixtures thereof. Preferably, the solvents used for dissolving the glassy polymer material for the preparation of the porous glassy
polymeric support membrane in the current invention include NMP, 1,3-dioxolane, glycerol, and n-decane.

[0025] The thin selective layer of the high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer described in the present invention is formed on top of the porous glassy polymeric support membrane by applying a dilute solution of a mixture of a fumed silica reinforced vinyl-terminated polyorganosiloxane polymer and a methylhydrosiloxane-dimethylsiloxane cross-linking copolymer in the presence of a platinum complex catalyst or a mixture of a fumed silica reinforced vinyl-terminated polyorganosiloxane polymer, a vinylorganosiloxane-dimethylsiloxane copolymer, and a methylhydrosiloxane-dimethylsiloxane cross-linking copolymer in the presence of a platinum complex catalyst to the top surface of the porous support membrane by dip-coating, spin coating, casting, soaking, spraying, painting, and other known conventional solution coating technologies. The thin selective layer of the high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer is formed by hydrosilylation reaction between the vinyl groups on the fumed silica reinforced vinyl-terminated polyorganosiloxane polymer or/and on vinylorganosiloxane-dimethylsiloxane copolymer and the silicon hydride groups on the methylhydrosiloxane-dimethylsiloxane cross-linking copolymer after evaporating the hydrocarbon organic solvent(s) and heating at 70°C to 150°C for a certain time.

[0026] The fumed silica reinforced vinyl-terminated polyorganosiloxane polymer used for the preparation of the new high flux, cross-linked, fumed silica reinforced polyorganosiloxane TFC membrane in the present invention provides the membrane with significantly improved mechanical strength under pressure for separation applications. The vinyl-terminated polyorganosiloxane polymer such as vinyl-terminated polydimethylsiloxane polymer is reinforced by fumed silica fillers such as hexamethyldisilazane treated fumed silica fillers. The vinylorganosiloxane-dimethylsiloxane copolymer used for the preparation of the new high flux, cross-linked, fumed silica reinforced polyorganosiloxane TFC membrane in the present invention can be selected from vinylmethylsiloxane-dimethylsiloxane, vinylphenylsiloxane-dimethylsiloxane, and a mixture thereof. The organic solvents that can be used for dissolving the vinyl-terminated polyorganosiloxane polymer, vinylorganosiloxane-dimethylsiloxane copolymer, and methylhydrosiloxane-dimethylsiloxane cross-linking copolymer in the present invention are essentially hydrocarbons such as n-heptane, n-hexane, n-octane, or mixtures thereof. It is preferred that
these vinyl-terminated polyorganosiloxane polymer, vinylorganosiloxane-dimethylsiloxane copolymer, and methylhydrosiloxane-dimethylsiloxane cross-linking copolymer are diluted in the hydrocarbon organic solvent or mixtures thereof in a concentration of from 1 to 20 wt% to provide a defect-free, thin, high flux, cross-linked, fumed silica reinforced polyorganosiloxane selective layer.

[0027] The platinum complex catalyst used for the preparation of the new high flux, cross-linked, fumed silica reinforced polyorganosiloxane TFC membrane in the present invention can those platinum compound catalysts that are well soluble in the reaction mixture such as platinum carbonylcyclovinylmethylsiloxane complex, platinum divinyUetramethylsiloxane complex, and platinum cyclovinylmethylsikixane complex.

[0028] The present invention also discloses a method of making the new high flux, cross-linked, fumed silica reinforced polyorganosiloxane TFC membrane comprising a thin selective layer of a high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer on top of a porous glassy polymeric support membrane comprising: a) preparation of a porous glassy polymeric support membrane from a glassy polymer such as polyethersulfone (PES), polysulfone (PSF), polyimide (PI), a blend of PES and PI, a blend of PSF and PI, and a blend of cellulose acetate (CA) and cellulose triacetate (CTA) via a phase inversion membrane fabrication process; b) coating a thin layer of a dilute solution of a mixture of a fumed silica reinforced vinyl-terminated polyorganosiloxane polymer and a methylhydrosiloxane-dimethylsiloxane cross-linking copolymer or a mixture of a fumed silica reinforced vinyl-terminated polyorganosiloxane polymer, a vinylorganosiloxane-dimethylsiloxane copolymer, and a methylhydrosiloxane-dimethylsiloxane cross-linking copolymer in the presence of a platinum complex catalyst on the top surface of the porous glassy polymeric support membrane by dip-coating, spin coating, casting, soaking, spraying, painting, and other known conventional solution coating technologies; c) evaporating the hydrocarbon organic solvents on said membrane and heating the coated membrane at 70-150 °C for a certain time to form the thin selective layer of high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer.

[0029] The new type of high flux, cross-linked, fumed silica reinforced polyorganosiloxane TFC membrane comprising a thin selective layer of a high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer on top of a porous glassy polymeric support membrane described in the present invention can be fabricated into any
convenient form suitable for a desired separation application. For example, the membranes can be in the form of hollow fibers, tubes, flat sheets, and the like. The new high flux, cross-linked, fumed silica reinforced polyorganosiloxane TFC membrane comprising a thin selective layer of a high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer on top of a porous glassy polymeric support membrane described in the present invention is in a spiral wound module that is in the form of flat sheet having a thickness from 30 to 400 µm. In another exemplary embodiment, the new high flux, cross-linked, fumed silica reinforced polyorganosiloxane TFC membrane comprising a thin selective layer of a high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer on top of a porous glassy polymeric support membrane described in the present invention is in a hollow fiber module that is in the form of thousands, hundreds of thousands, or more, of parallel, closely-packed hollow fibers or tubes. In one embodiment, each fiber has an outside diameter of from 200 micrometers (µm) to 700 millimeters (mm) and a wall thickness of from 30 to 200 µm. In operation, a feed contacts a first surface of said high flux, cross-linked, fumed silica reinforced polyorganosiloxane TFC membrane comprising a thin selective layer of a high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer on top of a porous glassy polymeric support membrane described in the present invention, a permeate permeates said membrane described in the present invention and is removed therefrom, and a retentate, not having permeated said membrane described in the present invention, also is removed therefrom. In another embodiment, the high flux, cross-linked, fumed silica reinforced polyorganosiloxane TFC membrane comprising a thin selective layer of a high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer on top of a porous glassy polymeric support membrane described in the present invention can be in the form of flat sheet having a thickness in the range of from 30 to 400 µm.
[0030] The new high flux, cross-linked, fumed silica reinforced polyorganosiloxane TFC membrane comprising a thin selective layer of a high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer on top of a porous glassy polymeric support membrane disclosed in the present invention has higher permeance for paraffins such as ethane, propane, n-butane, and olefins such as propylene, n-butene, ethylene than inert gases such as N\textsubscript{2} and H\textsubscript{2} as well as CH\textsubscript{4} and has significantly higher permeances for paraffins such as ethane, propane, n-butane, and olefins such as propylene, n-butene, ethylene than those of the thermally cross-linked RTV615A/B silicone rubber membrane for olefin and N\textsubscript{2} recovery and N\textsubscript{2} removal from natural gas applications (see Tables 1 and 2).

[0031] This invention discloses the use of single stage or multi-stage new high flux, cross-linked, fumed silica reinforced polyorganosiloxane TFC membrane comprising a thin selective layer of a high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer on top of a porous glassy polymeric support membrane described in the current invention for olefin recovery, LPG recovery, fuel gas conditioning, natural gas dew point control, nitrogen removal from natural gas, etc.

**EXAMPLES**

[0032] The following examples are provided to illustrate one or more preferred embodiments of the invention, but are not limited embodiments thereof. Numerous variations can be made to the following examples that lie within the scope of the invention.

**COMPARATIVE EXAMPLE 1**

Preparation of 5RTVS\textsubscript{i}/PES-a TFC membrane

[0033] A porous, asymmetric polyethersulfone (PES) gas separation support membrane was prepared via the phase-inversion process. A PES-a membrane casting dope comprising PES 18-25 wt\%, NMP 60-65 wt\%, 1,3-dioxolane 10-15 wt\%, glycerol 1-10 wt\% and n-decane 0.5-2 wt\% was cast on a nylon fabric then gelled by immersion in a 1 °C water bath for 10 minutes, and then annealed in a hot water bath at 85 °C for 5 minutes. The wet membrane was dried at 70 °C. The dried PES-a porous support membrane was coated with an RTVS\textsubscript{i} silicone rubber precursor polymer solution comprising RTV615A, RTV615B, and hexane (RTV615A:RTV615B=9: 1 (weight ratio), 5 wt\% of RTV615A+RTV615B in hexane)
and then thermally cross-linked at 85 °C for 1 h to form a thin, nonporous, dense RTVSi selective layer on the surface of the PES-a support membrane (abbreviated as 5RTVSi/PES-a). The 5RTVSi/PES-a TFC membrane was tested with a fuel gas mixture of 70% C1, 15% C2, 10% C3 and 5% C02 at 3549 kPa (500 psig) and 25 °C. The membrane was also tested with N2, H2, CH4, propylene, and propane single gases at 791 kPa (100 psig) and 25 °C.

EXAMPLE 1
Preparation of 5DMS-RTVB/PES-a TFC membrane

[0034] A porous, asymmetric PES gas separation support membrane was prepared via the phase-inversion process. A PES-a membrane casting dope comprising PES 18-25 wt%, NMP 60-65 wt%, 1,3-dioxolane 10-15 wt%, glycerol 1-10 wt% and n-decane 0.5-2 wt% was cast on a nylon fabric then gelled by immersion in a 1 °C water bath for 10 minutes, and then annealed in a hot water bath at 85 °C for 5 minutes. The wet membrane was dried at 70 °C. A 5 wt% DMS-RTV615B pre-cross-linked rubbery polymer solution was prepared by dissolving 6.3 g of fumed silica reinforced vinyl-terminated polydimethylsiloxane (Gelest catalog number: DMS-V31S15) and 0.7 g of RTV615B (Momentive) in 133 g of hexane at room temperature for 30 min. The dried PES-a porous support membrane was coated with the 5 wt% DMS-RTV615B pre-cross-linked polydimethylsiloxane polymer solution, dried at room temperature for 5 min, and then heated at 85 °C for 1.5-2 h to form a thin, nonporous, dense, cross-linked fumed silica reinforced DMS-RTV615B selective layer on the surface of the PES-a support membrane (abbreviated as 5DMS-RTVB/PES-a). The 5DMS-RTVZB/PES-a TFC membrane was tested with a fuel gas mixture of 70% C1, 15% C2, 10% C3 and 5% C02 at 3549 kPa (500 psig) and 25 °C. The membrane was also tested with N2, H2, CH4, propylene, and propane single gases at 791 kPa (100 psig) and 25 °C.

EXAMPLE 2
Preparation of 5DMS-RTVAB/PES-a TFC membrane

[0035] A 5DMS-RTVAB/PES-a TFC membrane was prepared using the procedure described in Example 1 except that the PES-a support membrane was coated with a 5 wt% DMS-RTVAB pre-cross-linked fumed silica reinforced polydimethylsiloxane polymer solution comprising 8.4 g of fumed silica reinforced vinyl-terminated polydimethylsiloxane (Gelest catalog number: DMS-V3 1S15), 4.2 g of RTV615A (Momentive), and 1.4 g of...
RTV615B (Momentive) in 126 g of hexane at room temperature for 30 min. The coated membrane was dried at room temperature for 5 min, and then heated at 85 °C for 1.5-2 h to form a thin, nonporous, dense, cross-linked DMS-RTV615AB selective layer on the surface of the PES-a support membrane (abbreviated as 5DMS-RTVAB/PES-a). The 5DMS-RTVAB/PES-a TFC membrane was tested with a fuel gas mixture of 70% C1, 15% C2, 10% C3 and 5% C02 at 3549 kPa (500 psig) and 25 °C. The membrane was also tested with N2, H2, CH4, propylene, and propane single gases at 791 kPa (100 psig) and 25 °C.

**TABLE 1**

Pure gas permeation results for 5RTVSi/PES-a and 5DMS-RTVB/PES-a TFC membranes for propylene recovery (propylene (C3=)/N2 separation)*

<table>
<thead>
<tr>
<th>Membrane</th>
<th>P_{C3=}/L (GPU)</th>
<th>α_{C3=}/N2</th>
</tr>
</thead>
<tbody>
<tr>
<td>5RTVSi/PES-a</td>
<td>2881</td>
<td>31.8</td>
</tr>
<tr>
<td>5DMS-RTVB/PES-a</td>
<td>4771</td>
<td>31.6</td>
</tr>
</tbody>
</table>

* Tested at room temperature and 791 kPa (100 psig); 1 GPU = 10⁻⁶ cm³(STP)/cm².sec.cmHg

**TABLE 2**

Pure gas permeation results for 5RTVSi/PES-a and 5DMS-RTVB/PES-a TFC membranes for CH4/N2 separation*

<table>
<thead>
<tr>
<th>Membrane</th>
<th>P_{CH4}/L (GPU)</th>
<th>α_{CH4}/N2</th>
</tr>
</thead>
<tbody>
<tr>
<td>5RTVSi/PES-a</td>
<td>278</td>
<td>3.05</td>
</tr>
<tr>
<td>5DMS-RTVB/PES-a</td>
<td>413</td>
<td>3.08</td>
</tr>
</tbody>
</table>

* Tested at room temperature and 791 kPa (100 psig), 1 GPU = 10⁻⁶ cm³(STP)/cm².sec.cmHg

**SPECIFIC EMBODIMENTS**

[0036] While the following is described in conjunction with specific embodiments, it will be understood that this description is intended to illustrate and not limit the scope of the preceding description and the appended claims.

[0037] A first embodiment of the invention is a high flux, cross-linked, fumed silica reinforced polyorganosiloxane thin film composite (TFC) membrane comprising a selective
layer of a high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer supported by a porous support membrane formed from a glassy polymer. An embodiment of the invention is one, any or all of prior embodiments in this paragraph up through the first embodiment in this paragraph wherein the glassy polymer is polyethersulfone (PES), polysulfone (PSF), polyimide (PI), a blend of PES and PI, a blend of PSF and PI, or a blend of cellulose acetate (CA) and cellulose triacetate (CTA). An embodiment of the invention is one, any or all of prior embodiments in this paragraph up through the first embodiment in this paragraph wherein the porous support membrane is a flat sheet support membrane or a hollow fiber support membrane. An embodiment of the invention is one, any or all of prior embodiments in this paragraph up through the first embodiment in this paragraph wherein the selective layer of a high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer is a flat sheet having a thickness from 30 nm to 40 µm. An embodiment of the invention is one, any or all of prior embodiments in this paragraph up through the first embodiment in this paragraph wherein the membrane has a higher permeance for paraffins than for inert gases. A second embodiment of the invention is one, any or all of prior embodiments in this paragraph up through the first embodiment in this paragraph wherein the membrane has a higher permeance for ethane, propane, n-butane, propylene, n-butene, and ethylene than for \( V \), \( H_2 \), and \( (1 - I) \).

[0038] A second embodiment of the invention is a method of making a high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer comprising conducting, in the presence of a platinum complex catalyst, an addition cure or hydrosilylation reaction (a) between a fumed silica reinforced vinyl-terminated polyorganosiloxane polymer and a methylhydrosiloxane-dimethylsiloxane cross-linking copolymer or (b) between a mixture of a fumed silica reinforced vinyl-terminated polyorganosiloxane polymer and a vinlyorganosiloxane-dimethylsiloxane copolymer and a methylhydrosiloxane-dimethylsiloxane cross-linking copolymer.

[0039] A third embodiment of the invention is a method of making a high flux, cross-linked, fumed silica reinforced polyorganosiloxane thin film composite (TFC) membrane comprising a selective layer of a high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer supported by a porous support membrane formed from a glassy polymer, the method comprising (a) preparing the porous support membrane using a phase inversion process by casting a glassy polymer solution using a casting knife; (b) forming the
high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer on the porous support membrane by (i) applying a dilute hydrocarbon solution of a mixture of a hydrocarbon solvent, a fumed silica reinforced vinyl-terminated polyorganosiloxane polymer and a methylhydrosiloxane-dimethylsiloxane cross-linking copolymer in the presence of a platinum complex catalyst or a mixture of a fumed silica reinforced vinyl-terminated polyorganosiloxane polymer, a vinylorganosiloxane-dimethylsiloxane copolymer, and a methylhydrosiloxane-dimethylsiloxane cross-linking copolymer in the presence of a platinum complex catalyst to the top surface of the porous support membrane; (ii) evaporating the solvent; and (iii) heating at 70° to 150°C for a period of time. An embodiment of the invention is one, any or all of prior embodiments in this paragraph up through the third embodiment in this paragraph wherein the glassy polymer solution comprises an organic solvent selected from the group consisting of N-methylpyrrolidone (NMP), N,N-dimethylacetamide (DMAc), methylene chloride, tetrahydrofuran (THF), acetone, methyl acetate, isopropanol, n-octane, n-hexane, n-decane, methanol, ethanol, N,N-dimethylformamide (DMF), dimethyl sulfoxide (DMSO), lactic acid, citric acid, dioxanes, 1,3-dioxolane, glycerol, and mixtures thereof. An embodiment of the invention is one, any or all of prior embodiments in this paragraph up through the third embodiment in this paragraph wherein the glassy polymer solution comprises NMP, 1,3-dioxolane, glycerol, and n-decane. An embodiment of the invention is one, any or all of prior embodiments in this paragraph up through the third embodiment in this paragraph wherein the applying of the dilute hydrocarbon solution to the top surface of the porous support membrane is by dip-coating, spin coating, casting, soaking, spraying, or painting. An embodiment of the invention is one, any or all of prior embodiments in this paragraph up through the third embodiment in this paragraph wherein the heating at 70° to 150°C is for 2 to 120 minutes.

A fourth embodiment of the invention is a process for removing at least one component from a stream comprising contracting the stream with a high flux, cross-linked, fumed silica reinforced polyorganosiloxane thin film composite (TFC) membrane comprising a selective layer of a high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer supported by a porous support membrane formed from a glassy polymer. An embodiment of the invention is one, any or all of prior embodiments in this paragraph up through the fourth embodiment in this paragraph wherein the at least one component is nitrogen, or hydrogen, or methane. An embodiment of the invention is one, any or all of prior
embodiments in this paragraph up through the fourth embodiment in this paragraph wherein the stream is natural gas, fuel gas, an olefin recovery stream from a polyolefin production process, LPG, and a natural gas dew point control stream. An embodiment of the invention is one, any or all of prior embodiments in this paragraph up through the fourth embodiment in this paragraph wherein the stream is natural gas, fuel gas, an olefin recovery stream from a polyolefin production process, LPG, and a natural gas dew point control stream. A embodiment of the invention is one, any or all of prior embodiments in this paragraph up through the fourth embodiment in this paragraph wherein the process is a step of an olefin recovery operation, a nitrogen recovery operation, an LPG recovery operation, a fuel gas conditioning operation, or a nitrogen removal from natural gas operation. An embodiment of the invention is one, any or all of prior embodiments in this paragraph up through the fourth embodiment in this paragraph wherein the process is a two-stage process further comprising a glassy polymeric membrane.

[0041] Without further elaboration, it is believed that using the preceding description that one skilled in the art can utilize the present invention to its fullest extent and easily ascertain the essential characteristics of this invention, without departing from the spirit and scope thereof, to make various changes and modifications of the invention and to adapt it to various usages and conditions. The preceding preferred specific embodiments are, therefore, to be construed as merely illustrative, and not limiting the remainder of the disclosure in any way whatsoever, and that it is intended to cover various modifications and equivalent arrangements included within the scope of the appended claims.

[0042] In the foregoing, all temperatures are set forth in degrees Celsius and, all parts and percentages are by weight, unless otherwise indicated.
CLAIMS:

i. A high flux, cross-linked, fumed silica reinforced polyorganosiloxane thin film composite (TFC) membrane comprising a selective layer of a high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer supported by a porous support membrane formed from a glassy polymer.

2. The high flux, cross-linked, fumed silica reinforced polyorganosiloxane thin film composite (TFC) membrane of claim 1 wherein the glassy polymer is polyethersulfone (PES), polysulfone (PSF), polyimide (PI), a blend of PES and PI, a blend of PSF and PI, or a blend of cellulose acetate (CA) and cellulose triacetate (CTA).

3. The high flux, cross-linked, fumed silica reinforced polyorganosiloxane thin film composite (TFC) membrane of claim 1 wherein the selective layer of a high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer is a flat sheet having a thickness from 30 nm to 40 \( \mu \text{m} \).

4. The high flux, cross-linked, fumed silica reinforced polyorganosiloxane thin film composite (TFC) membrane of claim 1 wherein the membrane has a higher permeance for paraffins than for inert gases or a higher permeance for ethane, propane, n-butane, propylene, n-butene, and ethylene than for \( \text{N}_2 \), \( \text{H}_2 \), and \( \text{CH}_4 \).

5. A method of making a high flux, cross-linked, fumed silica reinforced polyorganosiloxane thin film composite (TFC) membrane comprising a selective layer of a high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer supported by a porous support membrane formed from a glassy polymer, said method comprising:

(a) preparing the porous support membrane using a phase inversion process by casting a glassy polymer solution using a casting knife;

(b) forming the high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer on the porous support membrane by

(i) applying a dilute hydrocarbon solution of a mixture of a hydrocarbon solvent, a fumed silica reinforced vinyl-terminated polyorganosiloxane polymer and a methylhydrosiloxane-dimethylsiloxane cross-linking copolymer in the presence of a platinum complex catalyst or a mixture of a fumed silica reinforced vinyl-terminated polyorganosiloxane polymer, a vinylorganosiloxane-dimethylsiloxane copolymer, and a methylhydrosiloxane-dimethylsiloxane cross-linking copolymer in the
presence of a platinum complex catalyst to the top surface of the porous support membrane;

(ii) evaporating the solvent, and

(iii) heating at 70° to 150°C for a period of time.

6. The method of claim 8 wherein the glassy polymer solution comprises an organic solvent selected from the group consisting of N-methylpyrrolidone (NMP), N,N-dimethylacetamide (DMAc), methylene chloride, tetrahydrofuran (THF), acetone, methyl acetate, isopropanol, n-octane, n-hexane, n-decane, methanol, ethanol, N,N-dimethylformamide (DMF), dimethyl sulfoxide (DMSO), lactic acid, citric acid, dioxanes, 1,3-dioxolane, glycerol, and mixtures thereof.

7. A process for removing at least one component from a stream comprising contracting the stream with a high flux, cross-linked, fumed silica reinforced polyorganosiloxane thin film composite (TFC) membrane comprising a selective layer of a high flux, cross-linked, fumed silica reinforced polyorganosiloxane polymer supported by a porous support membrane formed from a glassy polymer.

8. The process of claim 13 wherein the at least one component is nitrogen, or hydrogen, or methane.

9. The process of claim 13 wherein the stream is natural gas, fuel gas, an olefin recovery stream from a polyolefin production process, LPG, and a natural gas dew point control stream.

10. The process of claim 13 wherein the process is a step of an olefin recovery operation, a nitrogen recovery operation, an LPG recovery operation, a fuel gas conditioning operation, or a nitrogen removal from natural gas operation.
**INTERNATIONAL SEARCH REPORT**

**A. CLASSIFICATION OF SUBJECT MATTER**
(see extra sheet)

According to International Patent Classification (IPC) or to both national classification and IPC

**B. FIELDS SEARCHED**

Minimum documentation searched (classification system followed by classification symbols)

B01D 71/64, 69/02, 69/06, 69/08, 53/22, 69/12, C08G 73/10, C07C 7/144

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

PatSearch (RUPTO internal), USPTO, PAJ, Esp@cenet, DWPI, EAPATIS, PATENTSCOPE, Information Retrieval System of FTPS

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

<table>
<thead>
<tr>
<th>Category</th>
<th>Citation of document, with indication, where appropriate, of the relevant passages</th>
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<td>X</td>
<td>WO 2003/039728 A2 (UOP LLC) 15.05.2003, [0001], [0017], [0025], [0028]-[0029], [0039], claims</td>
<td>1-2, 7-8</td>
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Further documents are listed in the continuation of Box C.

See patent family annex.

* Special categories of cited documents:

  "A" document defining the general state of the art which is not considered to be of particular relevance

  "E" earlier document but published on or after the international filing date

  "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

  "O" document referring to an oral disclosure, use, exhibition or other means

  "P" document published prior to the international filing date but later than the priority date claimed

  "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

  "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

  "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

  "&" document member of the same patent family

**Date of the actual completion of the international search**

15 February 2018 (15.02.2018)

**Date of mailing of the international search report**

28 February 2018 (28.02.2018)

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Telephone No. (499) 240-25-91

Form PCT/ISA/210 (second sheet) (January 2015)
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