Effects of water vapor on the separation of methane and carbon dioxide by gas permeation through polymeric membranes
by Gerald Thomas Paulson

A thesis submitted in partial fulfillment of the requirements for the degree of MASTER OF SCIENCE
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Abstract:

Three commercially available films and a vinylidene fluoride film modified with sulfolene were tested for the separation of carbon dioxide from carbon dioxide, methane and water vapor mixtures. Flux and separation factor values were obtained over a temperature range from 23 to 90°C and a feed gas moisture content from 0.0 to 1.32%. A feed gas containing 60% carbon dioxide and 40% methane on a dry basis was used at an operating pressure of 2068 kPa.

The commercially available films tested were cellulose acetate, polysulfone and polyethersulfone. All three films provided good separation. For example, the polysulfone film produced a permeate containing 96.0% carbon dioxide at room temperature and 0.12% water. Separation factor values for carbon dioxide ranged from 4.6 to 45.1. Flux values for the commercial films ranged from less than $10.E-05$ to 1.9×10^{-03} cu cm(STP)/sq cm(sec).

The sulfolene modified film also provided good separation. Separation factor values for carbon dioxide ranged from 9.6 to 33.6. Flux values ranged from $10.E-05$ to 5.7×10^{-05} cu cm(STP)/sq cm(sec).

Strong trends towards increasing flux and decreasing percentage of carbon dioxide in the permeate with increasing temperature were observed with all films.

A trend towards decreasing percentage of carbon dioxide in the permeate with increasing water content was observed with cellulose acetate, polysulfone and polyethersulfone films. These films demonstrated a strong trend towards increasing flux with increasing water content at low levels of water content. As the feed stream approached saturation, gas flux tended to become constant or actually decrease.

The sulfolene modified film tended towards constant or increasing percentage of carbon dioxide in the permeate with increasing water content. The flux tended to remain constant or decrease with increasing water content.

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EFFECTS OF WATER VAPOR ON THE SEPARATION OF METHANE
AND CARBON DIOXIDE BY GAS PERMEATION
THROUGH POLYMERIC MEMBRANES

by

GERALD THOMAS PAULSON

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TABLE OF CONTENTS

	<u>Page</u>
VITA	ii
ACKNOWLEDGMENTS	iii
LIST OF TABLES	vi
LIST OF FIGURES	viii
ABSTRACT	x
I. INTRODUCTION AND PURPOSE	1
II. REVIEW OF THE LITERATURE	3
A. BIOGAS PURIFICATION PROCESSES	3
B. POLYMER MEMBRANES USED FOR GASEOUS SEPARATION	4
C. LABORATORY MEMBRANE PROCESSES FOR BIOGAS PURIFICATION	5
D. COMMERCIAL PROCESSES FOR BIOGAS PURIFICATION	7
III. THEORETICAL BACKGROUND	8
A. THE NATURE OF THE TRANSPORT PROCESS	8
B. ORDERING ANALYSIS	8
C. DIFFUSION THROUGH A MEMBRANE	8
D. SIMPLIFIED MODEL OF PERMEATION OF ONE COMPOUND	9
E. TEMPERATURE EFFECTS	10
F. EFFECT OF GAS MIXTURE ON PERMEATION	10
G. EFFECT OF WATER VAPOR ON GAS PERMEATION	11

	<u>Page</u>
H. POLYMER CHEMISTRY ASPECTS	12
I. SEPARATION FACTOR DEFINITION	12
IV. EXPERIMENTAL EQUIPMENT, MATERIALS AND PROCEDURES	14
A. EXPERIMENTAL EQUIPMENT AND MATERIALS	14
1. GAS SUPPLY SYSTEM	14
2. WATER VAPOR CONTROL EQUIPMENT	14
3. PERMEATION CELL	16
4. CONSTANT TEMPERATURE ENCLOSURE FOR THE PERMEATION CELL	16
5. PERMEATE STREAM DRIER	18
6. PERMEATE RATE MEASUREMENT EQUIPMENT	19
7. GAS COMPOSITION ANALYSIS EQUIPMENT	19
B. EXPERIMENTAL PROCEDURE	20
1. GAS MIXTURE	20
2. CALIBRATION OF GAS CHROMATOGRAPH	20
3. MEMBRANE MANUFACTURE	20
4. OPERATING PROCEDURE	22
V. EXPERIMENTAL RESULTS AND DISCUSSION	24
A. MATERIALS TESTED	24
B. CONDITIONS OF THE TEST	24
C. GAS PERMEATION DATA	25
D. WATER VAPOR PERMEATION DATA	25

	<u>Page</u>
E. COMPUTER ANALYSIS OF DATA	25
F. DISCUSSION	31
1. OVERVIEW	31
2. REPRODUCIBILITY OF DATA	32
3. CELLULOSE ACETATE MEMBRANE	35
4. POLYSULFONE MEMBRANE	40
5. POLYETHERSULFONE MEMBRANE	45
6. SULFOLENE MODIFIED POLY(VINYLLIDENE FLUORIDE) MEMBRANE	50
7. WATER VAPOR FLUX	55
8. EFFECT OF WATER ON FLUX	57
VI. CONCLUSIONS AND RECOMMENDATIONS	60
A. CONCLUSIONS	60
B. RECOMMENDATIONS	60
VII. REFERENCED FOOTNOTES	62
VIII. APPENDIX	65
A. TABLE OF NOMENCLATURE	66
IX. BIBLIOGRAPHY	68

LIST OF TABLES

<u>Table</u>		<u>Page</u>
V-1	Summary of Test Results for Cellulose Acetate Film	26
V-2	Summary of Test Results for Polysulfone Film	27
V-3	Summary of Test Results for Polyethersulfone Film	28
V-4	Summary of Test Results for Modified Poly(Vinylidene Fluoride)	29
V-5	Summary of Test Results for Water Permeation	30

LIST OF FIGURES

<u>Figure</u>		<u>Page</u>
IV-1	Schematic Diagram of Permeation Equipment . . .	15
IV-2	Permeation Cell Diagram	17
IV-3	Calibration of Gas Chromatograph- Sample Volume vs. Peak Area x Attenuation	21
V-1	Reproducibility of Flux Data	33
V-2	Reproducibility of Separation Factors	34
V-3	Separation Factor vs. Percent Water in Feed Gas for Cellulose Acetate Film	36
V-4	Flux vs. Percent Water in Feed Gas for Cellulose Acetate Film	37
V-5	Plot of Separation Factor vs. Feed Stream Water Content and Temperature for Cellulose Acetate Film	38
V-6	Plot of Flux vs. Feed Stream Water Content and Temperature for Cellulose Acetate Film . .	39
V-7	Separation Factor vs. Percent Water in Feed Gas for Polysulfone Film	41
V-8	Flux vs. Percent Water in Feed Gas for Polysulfone Film	42
V-9	Plot of the Separation Factor vs. Feed Stream Water Content and Temperature for Polysulfone Film	43
V-10	Plot of Flux vs. Feed Stream Water Content and Temperature for Polysulfone Film	44
V-11	Separation Factor vs. Percent Water in Feed Gas for Polyethersulfone Film	46
V-12	Flux vs. Percent Water in Feed Gas for Polyethersulfone Film	47

<u>Figure</u>		<u>Page</u>
V-13	Plot of the Separation Factor vs. Feed Stream Water Content and Temperature for Polyethersulfone Film	48
V-14	Plot of Flux vs. Feed Stream Water Content and Temperature for Polyethersulfone	50
V-15	Separation Factor vs. Percent Water in Feed Gas for Sulfolene Modified Poly(vinylidene fluoride) Film	51
V-16	Flux vs. Percent Water in Feed Gas for Sulfolene Modified Poly(vinylidene fluoride) Film	52
V-17	Plot of Separation Factor vs. Feed Stream Water Content and Temperature for Sulfolene Modified Poly(vinylidene fluoride) Film	53
V-18	Plot of Flux vs. Feed Stream Water Content and Temperature for Sulfolene Modified Poly(vinylidene fluoride) Film	54
V-19	Water Vapor Flux vs. Feed Stream Water Content of the Films	56

ABSTRACT

Three commercially available films and a vinylidene fluoride film modified with sulfolene were tested for the separation of carbon dioxide from carbon dioxide, methane and water vapor mixtures. Flux and separation factor values were obtained over a temperature range from 23 to 90°C and a feed gas moisture content from 0.0 to 1.32%. A feed gas containing 60% carbon dioxide and 40% methane on a dry basis was used at an operating pressure of 2068 kPa.

The commercially available films tested were cellulose acetate, polysulfone and polyethersulfone. All three films provided good separation. For example, the polysulfone film produced a permeate containing 96.0% carbon dioxide at room temperature and 0.12% water. Separation factor values for carbon dioxide ranged from 4.6 to 45.1. Flux values for the commercial films ranged from less than $10.E-05$ to $1.9 \times 10E-03$ cu cm(STP)/sq cm(sec).

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INTRODUCTION AND PURPOSE

The separation of a gas mixture by a membrane process involves the selective transport of a given species through a film. This method of separation has made considerable progress in recent years in areas such as the development of more selective and permeable membranes, as well as simple and efficient equipment for large scale operations.

The separation of carbon dioxide and methane is important in a number of areas, particularly in the manufacturing of Substitute Natural Gas (SNG). A substantial amount of SNG can be produced from biomass and gaseous effluent from anaerobic sewage treatment plants, landfills, oil fields, coal mines and agricultural waste digesters(1). The upgrade of biogas to SNG requires the removal of carbon dioxide, hydrogen sulfide and water. Biogas from anaerobic digestion varies from 40 to 60% methane and 60 to 40% carbon dioxide, contains traces of hydrogen sulfide and is saturated with water vapor(2). In order to provide pipeline quality gas, the SNG must be approximately 98% methane, a few parts per million hydrogen sulfide and dried(3). Many commercial processes capable of meeting these standards are available. However, biogas membrane processes are estimated to be in the order of a third to a half the cost of traditional systems(4-5).

Membrane surface area requirements are of significant economical interest. Studies indicate that as much as 95% of the total investment costs of a gas permeation process is determined by the area(6-7). To minimize area requirements it is necessary to know the dependence of

gas permeation rates over a wide range of variables. This work was conducted in an attempt to contribute to the ultimate goal of an economical membrane separation process for biogas purification. The specific objectives of this research were:

1. to test various commercially available polymeric films to determine permeation rate and selectivity for carbon dioxide.
2. to test a plasticizer in a vinylidene fluoride film to determine permeation rates and selectivity for carbon dioxide.
3. to determine the effect of temperature on selectivity for carbon dioxide in the various membranes.
4. to determine the effect of temperature on permeation rates in the various membranes.
5. to determine the effect of water vapor on the selectivity for carbon dioxide in the various membranes.
6. to determine the effect of water vapor on permeation rates in the various membranes.
7. to determine the permeation rates of water vapor in the various membranes.

REVIEW OF THE LITERATURE

A. BIOGAS PURIFICATION PROCESSES

Carbon dioxide, hydrogen sulfide and water removal technologies include a large number of commercial processes. In addition to membrane separation systems; four major groupings can be identified in gas removal processes(8).

Physical absorption involves the dissolution of the gas in a liquid solvent. Solvent regeneration usually can be accomplished by simple flashing or by stripping with an inert gas. Historically, water scrubbing was the first method used for carbon dioxide removal.

Chemical absorption involves the formation of a reversible chemical bond between the solvent and the solute. Regeneration involves breaking those bonds. Most solvents are either aqueous solutions of amines or aqueous solutions of alkaline salts. Potassium carbonate solutions can be used to chemically absorb carbon dioxide and hydrogen sulfide.

Chemical conversion can be utilized to reduce an undesirable gas concentration. Several commercial processes are available for converting hydrogen sulfide to elemental sulfur. Carbon dioxide when mixed with hydrogen can be catalytically converted to methane and water(9).

Gas removal by condensation is achieved by cooling, compression or a combination of both. Commercial trials of biogas purification by condensation have not been successful because of high capital costs and low thermal efficiency(10).

B. POLYMERIC MEMBRANES USED FOR GASEOUS SEPARATION

Several gas permeation processes have been studied in detail. Differences in solubilities, mass transfer resistances or both account for membrane selectivity. Separation of a gas mixture is achieved by the flow of one component at a greater rate than the others.

Silicone rubber is used as a membrane for the separation of oxygen from air(11). A three-stage cascade of hollow fiber permeators results in a product stream of 38% oxygen. Oxygen could be enriched to 32.6% in a one stage process(12).

The recovery of helium from natural gas has received much attention. Union Carbide has conducted large scale tests using cellulose acetate films in a two-stage process(13). Some tests have been conducted using a Teflon FEP film(14). Teijin Limited of Japan has announced a method that employs an undisclosed hollow fiber system to recover helium(15).

A commercial installation for the separation of hydrogen from a refinery gas stream has been in operation since 1969(16). The system uses hollow fibers of dacron polyester and is marketed by the Du Pont Company.

The removal of rare gas fission products has been studied by the nuclear industry(17). Silicone rubber membranes is used in a flat plate permeator for the tests.

Polymer membrane processes have been tested for a large number of laboratory systems. Brubaker and Kammermeyer tested polyethylene, trifluoromono-chloro-ethylene and cellulose acetate over a wide variety of gas mixtures(18). Tajar and Miller reported data for the permeation of carbon dioxide, oxygen and nitrogen in a membrane consisting of polyethyleneamine, polyvinylbutyral, epoxy and water(19).

Modified poly(vinylidene fluoride) membranes have been used in four studies. Seibel and McCandless used sulfolane modified films to separate sulfur dioxide from nitrogen(20). Zavaleta studied a number of modified films for separating sulfur dioxide from nitrogen(21). Heyd studied eight modified films as well as six commercially available films for separating hydrogen sulfide from nitrogen(22). Ellig, Althouse and McCandless tested several films for separating carbon dioxide from methane(23). Their study is discussed in the following section.

C. LABORATORY MEMBRANE PROCESSES FOR BIOGAS PURIFICATION

General Electric Company has experimented with immobilized liquid membranes for carbon dioxide removal(24). The liquid membrane is immobilized by a hydrophobic porous support membrane. The liquid membrane consists of a hydrophilic porous membrane whose pores are filled with an aqueous carbonate solution. This membrane utilizes the phenomenon described as "facilitated transport" by the addition of carbon dioxide hydrolysis catalysts to the aqueous carbonate solution. Reported values

of carbon dioxide permeability vary from 0. to 160. (cc/sq cm sec) (cm/cm Hg) with a membrane thickness of 0.0015cm and a carbon dioxide feed gas with partial pressure varying from 5 to 200 cm Hg. Humidity control is critical.

The Monsanto Company has developed coated membranes(25). The process involves a multicomponent membrane consisting of a porous film and a thin(<0.005cm) coating. The film is either cellulose acetate, polycarbonate or polysulfone and the coating is either polysiloxanes or polystyrene. The combinations of films and coatings will separate carbon monoxide, nitrogen, argon, methane, ethane, carbon dioxide and sulfur hexafluoride.

Ellig, Althouse, and McCandless tested 16 membranes for the removal of carbon dioxide from a 60 percent carbon dioxide and 40 percent methane mixture(26). A moisture-free feed gas was used. Thirteen commercial membranes were tested. The best separations were obtained with polyethersulfone, polysulfone and cellulose acetate. Three modified poly(vinylidene fluoride) membranes were made by dissolving vinylidene fluoride and a modifier in dimethyl formamide. The films were cast on glass plates between thicknesses of masking tape. The solvent was evaporated by placing the plates in an electrically heated oven. The membrane modified with sulfolene demonstrated the best separation of the three. Observed separation factors ranged from less than 0.1 to 40.2

with fluxes up to 0.044 cu cm (STP)/sq cm sec . The highest fluxes were obtained from the films with the lowest separation factors.

D. COMMERCIAL PROCESSES FOR BIOGAS PURIFICATION

The General Electric Company is currently marketing a Permaselective Membrane Process(27). A 27.8 standard cubic meter per day system is operating on sewage digester gas in Southern California. A 278.0 standard cubic meter per day system is planned for the Fresh Kills Landfill located in New York City(28).

The General Electric Permaselective Membrane Process operates by a pressure differential and allows carbon dioxide to pass through the membrane surface while methane is retained in the high pressure gas stream. A second stage is used to recover methane in the carbon dioxide effluent. The system uses a silicone- polycarbonate membrane with thicknesses as thin as 200 angstroms. Water is removed from the gas prior to injection into the membrane system.

THEORETICAL BACKGROUND

A. THE NATURE OF THE TRANSPORT PROCESS

Permeation through a polymeric membrane is thought to involve the following series of steps(29):

- (i) transport from the bulk mixture to the membrane surface.
- (ii) solution of the gas into the membrane at the interface.
- (iii) transport through the membrane(diffusion).
- (iv) release of the gas from solution at the opposite interface.
- (v) transport from the membrane interface to the permeate stream.

The term permeation is used to describe the overall mass transport of gas through this sequence.

B. ORDERING ANALYSIS

Each of the five steps represents a resistance to mass transfer of various magnitude. The resistances of some steps are negligible in comparison with other steps. In gas-phase permeation, steps (i), (ii), (iv) and (v) may be negligible(30).

C. DIFFUSION THROUGH A MEMBRANE

For membrane diffusion the basic transport equations are:

$$N_A = -D \frac{\partial C_a}{\partial z} - \frac{D}{2C_m} \frac{\partial C_a^2}{\partial z} \quad (1)$$

and

$$\frac{\partial C_a}{\partial t} = \frac{\partial}{\partial z} D \frac{\partial C_a}{\partial z} + \frac{\partial}{\partial z} D \frac{C_a}{C_m} \frac{\partial C_a}{\partial z} \quad (2)$$

These equations differ from Fick's basic diffusion equations by the addition of the second terms due to the presence of the membrane.

The effect of the second term may be negligible in instances of large permeation fluxes.

D. SIMPLIFIED MODEL OF PERMEATION OF ONE COMPOUND

Assuming steady state diffusion through the membrane to be the rate controlling step and neglecting the second term of the diffusion equation (presence of the membrane), permeation reduces to Fick's basic diffusion equation.

$$N_a = -D \frac{dC_a}{dz} \quad (3)$$

Integration of this equation using the appropriate boundary conditions yields:

$$-N_a (z_1 - z_2) = -D(C_1 - C_2) \quad (4)$$

or

$$N_a = \frac{D}{L}(C_1 - C_2) \quad (5)$$

Assuming equilibrium between the gas and the interface, a Henry's Law expression, $C = (S)(p)$, is applicable. S is the solubility coefficient of the gas in the polymer. The resulting equation becomes:

$$N_a = \frac{DS}{L}(p_{a1} - p_{a2}) \quad (6)$$

The adequacy of Fick's and Henry's Law are questionable in many polymers(31). However, equation (6) does provide a basis for the study of permeation.

E. TEMPERATURE EFFECTS

For unmodified films D and S usually obey an Arrhenius type relationship:

$$D = D_0 e^{(-E/RT)} \quad (7)$$

$$S = S_0 e^{(-\Delta h/RT)} \quad (8)$$

E is the activation energy and h is the heat of solution for diffusion. Both diffusivity and gas solubility increase with increasing temperature.

For films modified with a liquid, the solubility of gases in the modifier usually decreases as temperature is increased. The flux could actually decrease with increasing temperature.

F. EFFECT OF GAS MIXTURES ON PERMEATION

For the case of a gas mixture in contact with a membrane, equation (6) becomes:

$$N_a = \frac{DS}{L}(p_{a1} - p_{a2}) \quad (9)$$

or in terms of the total pressure:

$$N_a = \frac{DS}{L}(P_1 X_{a1} - P_2 Y_{a2}) \quad (10)$$

The effect of a gas mixture on permeation is dependent on the polymer and the gaseous components. Mixtures of nitrogen and oxygen permeate in an additive fashion(32). Each gas permeates independently. Both the gases have low solubilities in most polymers.

Nitrogen and carbon dioxide mixtures, and oxygen and carbon dioxide mixtures permeate dependently(33). Oxygen has been reported to permeate at rates up to three times faster in the presence of large amounts of carbon dioxide than in pure oxygen.

G. EFFECT OF WATER VAPOR ON GAS PERMEATION

Possible interaction of water vapor with the membrane will influence the mechanism of permeation. In hydrophilic membranes a solution of water in the polymer will occur(34). This results in a softening or plasticizing effect, and may cause swelling. In hydrophobic membranes water should not interact with the polymer.

Few studies have been conducted on the effect of water vapor upon gas permeation. Simril and Hershberger reported a 50-fold increase in carbon dioxide permeability through cellophane as the relative humidity increased from 0. to 100%(35). Pilar reported a 1000-fold increase for the same conditions(36). Two methods have been adopted for the study of the effects of water on permeation. The first method preconditions the film with water and the film is then tested. The second method uses a test gas that is humidified. The results of the two methods vary(37).

In general, the permeability of gases increased as the moisture content of the film is increased. However, transmission of water vapor is pressure dependent. The solubility of water in films is reported to increase in a nonlinear fashion with pressure(38).

Some data is available on the commercial films used in this study(39). Water absorption in 24 hours for cellulose acetate, polysulfone and polyethersulfone is 8.5, 0.3 and 2.1 weight percent respectively.

H. POLYMER CHEMISTRY ASPECTS

Gases are transported through polymeric nonporous membrane by means of diffusion. The partial pressures of the gases are the driving forces and the film thickness, gas solubility and gas diffusivity control permeation rates.

Plasticizers (or modifiers) in a polymer have several effects. The addition of a plasticizer increases the diffusivity of gases in a polymer. Solubilities of gases or vapors may also increase, particularly when the gas or vapor is soluble in the modifier. The overall effect of plasticizers is generally an increase in the permeate flux.

Zavaleta studied the effect of plasticizer content and concluded that 18% sulfolene in poly(vinylidene fluoride) yielded the maximum flux(40). Films with a higher sulfone content were prone to leak.

I. SEPARATION FACTOR DEFINITION

The degree of separation of components A and B is commonly expressed in terms of a separation factor. It is defined by the concentration ratio:

$$\alpha_a = \frac{(C_a/C_b)_p}{(C_a/C_b)_s} \quad (11)$$

In terms of mole fractions the separation factor is:

$$\alpha_a = \frac{Y_a/(1-Y_a)}{X_a/(1-X_a)} \quad (12)$$

For the case of perfect mixing on the high pressure side, the overall separation factor is identical to the separation factor at any point along the membrane length.

EXPERIMENTAL EQUIPMENT, MATERIALS AND PROCEDURES

A. EXPERIMENTAL EQUIPMENT AND MATERIALS

A schematic diagram of the experimental equipment is shown in Figure IV-1. The equipment consisted of a gas supply system, water vapor control equipment, permeation cell, a constant temperature enclosure for the cell, a permeate stream drier, permeate flow measurement equipment and gas composition analysis equipment. A description of each subsystems follows.

1. GAS SUPPLY SYSTEM

The feed gas was stored in a National Cylinder Gas Division of Chemetron Corporation gas cylinder. An operating pressure of 2068 kPa was set by a Grove "Mighty Mite" back pressure regulator and was measured by a Matheson Company pressure gauge. A Matheson pressure regulator located on the outlet of the high pressure cylinder was maintained at a pressure slightly above the operating pressure. A feed flow rate (approximately 2 liters/hour) was set using a Whitey micrometering valve. The gas rate was measured periodically by a Varian Inc. bubble meter. The gas was vented through an oil seal.

2. WATER VAPOR CONTROL EQUIPMENT

The water vapor control equipment consisted of two columns in series that were submerged vertically in a Forma Scientific Masterline model 2095 company water bath(-30 to 72°C range). Feed gas enters a 2.0 cm diameter column that was filled with water and packed with 0.3 cm Fenske rings. The column was 20.3 cm in length. The wet gas continued

- (1) Feed Gas Cylinder (2) Pressure Regulator (3) Needle Valve (4) Constant Temperature Bath (5) Packed Column (6) Glass Wool Column (7) By-pass Valves (8) Feed Line Trace Heating (9) Cell Enclosure (10) Tubing Coil (11) Permeation Cell (12) Wet Permeate Sampling Septum (13) Permeate Drier (14) Vacuum Septum (15) Pressure Gauge (16) Back Pressure Regulator (17) Two Way Valve (18) Sampling Valve (19) Length Measurement (20) Oil Seal

FIGURE IV-1. SCHEMATIC DIAGRAM OF PERMEATION EQUIPMENT

through a 15.2 cm length of 1.2 cm in diameter column that was packed with glass wool. The glass wool column eliminated water droplets in the feed gas. The resulting gas was saturated at the temperature of the water bath. The bath temperature was varied to produce a feed gas of different water vapor concentrations. The gas feed line from the constant temperature bath to the permeation cell enclosure was heated by circulating water from the bath through a 0.6 cm in diameter copper line to prevent condensation. The two lines were wrapped in 2.5 cm thick fiberglass insulation. The columns were by-passed for runs requiring a moisture free feed gas.

3. PERMEATION CELL

A diagram of the permeation cell is shown in Figure IV-2. A cavity was machined into each of two stainless steel blank flanges which were 1.6cm thick and 11.4cm in diameter. A porous stainless steel disk covered with filter paper was placed in the cavity on the low pressure side to support the membrane. The membrane was sealed between two teflon gaskets and the permeation cell was assembled with eight equally spaced bolts. The gasket opening, which determine the exposed membrane surface, had an area of 20.3 sq cm. The high pressure side of the cell had a tubular thermocouple well. A sampling septum was located on the low pressure side of the cell. The septum was used to remove moist samples of the permeate.

- (1) Membrane (2) Thermistor (3) Gaskets
 (4) Filter Paper (5) Porous Stainless Steel
 Disk (6) Sampling Septum

FIGURE IV-2. PERMEATION CELL DIAGRAM

4. CONSTANT TEMPERATURE ENCLOSURE FOR THE PERMEATION CELL

The constant temperature enclosure consisted of a section of 45.5 cm diameter asbestos pipe 31.1 cm high. The wall thickness was 1.9 cm. The bottom of the enclosure was sealed with a 0.6 cm asbestos board. The bottom and sides of the enclosure were lined with 1.9 cm thick fiberglass insulation. The lid of the enclosure consisted of 1.9 cm insulation between 0.6 cm asbestos boards. To insure a tight seal the insulation and lower board extended into the interior of the enclosure. The feed gas lines and permeate line entered the enclosure through grooves cut in the top surface of the asbestos pipe section. A 0.6 cm hole was cut in the side of the enclosure for the location of a thermometer. For runs above room temperature, heat was provided by a 500 watt heater placed in the bottom of the enclosure. The heater was covered with an asbestos board to shield the permeation cell and feed lines from direct exposure to the heaters. The input current to the heater was controlled by a Yellow Springs Instrument Company Thermistemp Model 63 thermistor temperature controller. The thermistor probe was mounted in the thermowell section of a tee located at the gas inlet to the high pressure side of the permeation cell. A mercury thermometer was used to determine the temperature of the air inside the enclosure.

5. PERMEATE STREAM DRYER

A 7.6 cm length of 0.6 cm diameter polyethylene tube was placed in the permeate line as it left the permeate cell enclosure. The tube was filled with Dryrite and was supported with glass wool.

6. PERMEATE RATE MEASUREMENT EQUIPMENT

The rate of permeation was determined by timing the movement of a plug of VanWaters' and Rogers #54996 pump oil through a known distance of tubing. A 1.2m section of Chemplast Inc. Chemfluor Special FEP teflon tubing was used. The tubing was calibrated by filling a given length of tubing with oil by a 0.5 cc Precision Sampling Corporation syringe. The volume per unit length of the tubing has been determined to be 0.022 cc/cm.

7. GAS COMPOSITION ANALYSIS EQUIPMENT

The composition of samples was analyzed using a Varian Aerograph series 1400 thermal conductivity gas chromatograph and a Sargent model SR chart recorder. The recorder incorporated a disc integrator. The chromatograph column was a 1.8m section of 0.3cm stainless steel tubing packed with Walters Associated Inc. Porapak Q-S packing. Samples were introduced into the chromatograph by the use of a six port sampling valve with a 1.0 cu cm sample loop.

The following conditions were used for gas analysis:

Column Temperature- 75° C

Detector Temperature- 120° C

Carrier Gas Flow- 16.2 ml/min

Detector Current- 150 milliamperes

Carrier Gas- Hydrogen

The column temperature was increased to 170°C for the analysis of water vapor content.

B. EXPERIMENTAL PROCEDURE

1. GAS MIXTURE

A cylinder was evacuated using a vacuum pump. The cylinder was filled with laboratory grade carbon dioxide to 5727 kPa. Laboratory grade methane was then added until the final pressure of 9545 kPa was reached. The cylinder was placed on an electrical heater and was periodically heated for several days to assure a uniform composition.

2. CALIBRATION OF GAS CHROMATOGRAPH

The Porapak Q-S column was calibrated. Samples of different sizes of a sixty percent carbon dioxide and forty percent methane mixture were taken with a 1.0 cc Precision Sampling Corporation syringe through a silicon rubber septum mounted on the outlet of a low pressure regulator. Samples were injected into the gas chromatograph for analysis under the same operating conditions. The runs were repeated several times to insure reproducibility. The areas of the resulting peaks were determined from the disc integrator. These calibration data appear in Figure IV-3 as a function of the peak area multiplied by the chromatograph attenuation. The chromatograph response was shown to be linear.

3. MEMBRANE MANUFACTURE

The poly(vinylidene fluoride) film modified with sulfolene was manufactured by the following procedure. A clean pyrex beaker was

FIGURE IV-3. CALIBRATION OF GAS CHROMATOGRAPH
SAMPLE VOLUME vs. PEAK AREA $\times$ ATTENUATION

weighed on a Mettler balance (sensitivity 0.0001 gm). Twelve percent by weight of sulfolene was added to the appropriate amount of vinylidene fluoride. Dimethyl formamide was added in a ratio of 5.7 cu cm dimethyl formamide per gram of vinylidene fluoride. The mixture was stirred with a glass rod for ten minutes. The beaker was covered with polyethylene film and was placed on a hot plate at low heat until complete dissolution was achieved. The solution was degassed in a vacuum chamber at 406 mm Hg absolute for thirty minutes. The film was cast on a 12.7cm by 24.4cm glass plate between thicknesses of masking tape. The solution was distributed by drawing a glass rod over the plate with the rod resting on the masking tape. The glass plate was then placed in an electrically heated oven (125°C) for thirty minutes. The plate was cooled to room temperature before the film was stripped from the plate.

4. OPERATING PROCEDURE

A permeation experiment was accomplished by completion of the following steps:

- i) The membrane was mounted in the test cell and the system pressure was increased to the operating condition.
- ii) The temperature of the permeation cell was brought to the desired level by setting the temperature controller.

- iii) The water vapor in the feed stream was brought to the desired level by setting the temperature of the constant temperature bath.
- iv) The temperature of the permeation cell and constant temperature bath was checked by a mercury thermometer. When both were at the desired level, a vacuum was applied to the permeate line to remove residual gases.
- v) The system was allowed to run until no change was detected in the permeation rate and permeate gas composition.
- vi) The permeate flow rate and composition were measured five times.

Runs requiring a dry feed gas by-passed the water vapor equipment. The system was dried by the rapid flow of compressed air through all lines before the run. Runs requiring a moist feed gas were run in order of the least moist to saturated gas, insuring that residual moisture in the lines did not affect the results.

A 1.0cc syringe was used to extract moist permeate from the permeation cell for water vapor tests.

The above procedure was followed to achieve steady state conditions for water vapor content tests. Dry permeate flow was measured and wet permeate gas composition was determined.

EXPERIMENTAL RESULTS AND DISCUSSION

A. MATERIALS TESTED

Tests to determine permeate flux and overall separation factors for a carbon dioxide and methane mixture were conducted using the following available films:

- i) Polysulfone film: compound P-1700 (0.125mm thick) manufactured by Union Carbide Corporation.
- ii) Polyethersulfone film: compound PES 600 (0.075mm thick) manufactured by I.C.I. United States, Incorporated.
- iii) Cellulose acetate film: compound 100 CA-43 (0.025mm thick) manufactured by DuPont.
- iv) Sulfolene modified poly(vinylidene fluoride)-12% modifier produced for this experiment (0.050 mm thick approximately).

B. CONDITIONS OF THE TEST

Test runs were made with the cell temperature and feed gas moisture content as two variable parameters. The cell temperature was varied from 23 C (room temperature) to 90°C. The test conditions were chosen to be 23 ,45 ,65 and 90°C. The feed gas moisture was varied from a dry gas to a saturated gas by temperature selection of the constant temperature bath. Eight test conditions were chosen to be 0.0, 0.12, 0.24, 0.36, 0.60, 0.84, 1.08 and 1.32 mole percent water. The feed gas water content was limited by the temperature of the permeation cell. If the constant temperature bath was run at a temperature above that of the permeation cell condensate would form in the cell. Saturation was

achieved after 0.12, 0.36 and 0.84% water at 23 ,45 and 65°C respectively. Saturation at 90°C cannot be achieved because of a temperature limitation of the constant temperature bath. A maximum of 1.32% water was used at 90°C.

C. GAS PERMEATION DATA

Five values of dried permeate flow and permeate composition were measured at each test condition. The average values and standard deviations are presented in Tables V-1 through V-4. The calculated values of flux(STP) and separation factors are also listed.

D. WATER VAPOR PERMEATION DATA

Dry permeate flow and moist permeate composition were measured at two test points for each membrane. The data are presented in Table V-5. The calculated values of water vapor flux(STP) are also listed.

E. COMPUTER ANALYSIS OF DATA

Two computer programs were used to develop a three dimensional display of the data. Both programs were developed at the Harvard Center for Environmental Design Studies, Harvard University, and are available through the Honeywell CP-6 computer at Montana State University. The SYMAP program interpolated from data locations to produce a continuous surface in the form of a grid matrix. The grid matrix contained 1569 points. The program determined values by the data within a computer defined search radius. The SYMVU graphics program generated a three dimensional display of the grid matrix on a CALCOMP plotter. The plot's

TABLE V-1. SUMMARY OF TEST RESULTS FOR CELLULOSE ACETATE FILM

CELL TEMPERATURE (°C)	% H ₂ O IN FEED	FRACTION CO ₂ IN PERMEATE	STANDARD DEVIATION	SEPARATION FACTOR	FLUX(x10 ⁴) cc(STP)/cm ² sec	STANDARD DEVIATION (x10 ⁴)
23	0.0	0.965	0.002	22.8	1.7	0.1
45	0.0	0.953	0.002	16.8	3.8	0.3
65	0.0	0.931	0.005	11.2	5.6	0.1
90	0.0	0.914	0.002	8.8	7.3	0.1
23	0.12	0.875	0.013	5.8	2.0	0.1
45	0.12	0.928	0.003	10.8	9.2	0.1
65	0.12	0.913	0.002	8.7	13.2	0.1
90	0.12	0.903	0.008	7.7	19.7	0.4
45	0.24	0.923	0.003	9.9	9.9	0.3
65	0.24	0.905	0.006	7.9	9.0	0.2
90	0.24	0.887	0.007	6.5	9.1	0.3
45	0.36	0.914	0.006	8.8	3.7	0.1
65	0.36	0.920	0.005	9.5	6.4	0.3
90	0.36	0.898	0.007	7.3	10.5	0.4
65	0.60	0.913	0.002	8.7	8.0	0.2
90	0.60	0.848	0.005	4.6	10.8	0.3
65	0.84	0.905	0.005	7.9	7.8	0.2
90	0.84	0.903	0.007	7.7	7.8	0.2
90	1.08	0.886	0.002	6.4	9.0	0.1
90	1.32	0.910	0.002	8.3	9.4	0.3

TABLE V-2. SUMMARY OF TEST RESULTS FOR POLYSULFONE FILM

CELL TEMPERATURE (°C)	% H ₂ O IN FEED	FRACTION CO ₂ IN PERMEATE	STANDARD DEVIATION	SEPARATION FACTOR	FLUX(x10 ⁵) cc(STP)/cm ² sec	STANDARD DEVIATION (x10 ⁵)
23	0.0	0.971	0.003	27.7	2.0	0.5
45	0.0	0.960	0.003	19.8	2.2	0.2
65	0.0	0.942	0.001	13.4	2.0	0.4
90	0.0	0.924	0.005	10.9	3.1	0.0
23	0.12	0.962	0.003	20.9	1.6	0.1
45	0.12	0.947	0.002	14.8	4.0	0.4
65	0.12	0.948	0.001	15.1	5.6	0.2
90	0.12	0.909	0.005	8.3	5.9	0.2
45	0.24	0.934	0.002	12.3	3.4	0.5
65	0.24	0.945	0.001	14.2	4.2	0.4
90	0.24	0.927	0.005	10.5	6.2	0.1
45	0.36	0.939	0.002	12.7	2.8	0.2
65	0.36	0.933	0.005	11.5	5.3	0.1
90	0.36	0.920	0.001	9.5	5.3	0.0
65	0.60	0.937	0.003	12.3	6.1	0.2
90	0.60	0.923	0.001	9.9	7.6	0.6
65	0.84	0.922	0.005	9.8	5.6	0.1
90	0.84	0.924	0.002	10.0	5.9	0.1
90	1.08	0.921	0.004	9.6	7.5	0.1
90	1.32	0.920	0.004	9.5	4.5	0.7

TABLE V-3. SUMMARY OF TEST RESULTS FOR POLYETHERSULFONE FILM

CELL TEMPERATURE (°C)	% H ₂ O IN FEED	FRACTION CO ₂ IN PERMEATE	STANDARD DEVIATION	SEPARATION FACTOR	FLUX(x10 ⁵) cc(STP)/cm ² sec	STANDARD DEVIATION (x10 ⁵)
23	0.0	-	-	-	0.0	0.0
45	0.0	0.982	0.002	45.1	1.7	0.1
65	0.0	0.976	0.001	33.6	2.4	0.1
90	0.0	0.975	0.002	32.2	4.0	0.5
23	0.12	-	-	-	0.0	0.0
45	0.12	0.981	0.002	32.5	2.6	0.4
65	0.12	0.973	0.001	22.7	3.0	0.6
90	0.12	0.966	0.001	23.5	10.2	0.1
45	0.24	0.975	0.001	24.5	4.8	0.4
65	0.24	0.972	0.000	21.4	7.0	0.2
90	0.24	0.975	0.001	23.5	10.0	0.4
65	0.36	0.970	0.000	20.3	8.1	0.1
90	0.36	0.964	0.004	22.1	10.5	0.2
65	0.60	0.970	0.003	20.3	6.1	0.1
90	0.60	0.966	0.001	23.5	9.7	0.1
65	0.84	0.969	0.002	19.7	6.4	0.6
90	0.84	0.966	0.002	23.5	8.6	0.4
90	1.08	0.965	0.005	22.8	8.1	0.2
90	1.32	0.964	0.002	22.1	8.1	0.2

TABLE V-4. SUMMARY OF THE TEST RESULTS FOR SOLFOLENE
MODIFIED POLY (VINYLIDENE FLUORIDE) FILM

CELL TEMPERATURE (°C)	% H ₂ O IN FEED	FRACTION CO ₂ IN PERMEATE	STANDARD DEVIATION	SEPARATION FACTOR	FLUX(x10 ⁵) cc(STP)/cm ² sec	STANDARD DEVIATION (x10 ⁵)
23	0.0	0.963	0.001	21.5	1.2	0.1
45	0.0	0.958	0.006	18.9	5.2	0.0
65	0.0	0.947	0.007	14.8	16.3	0.0
90	0.0	0.921	0.003	9.6	37.6	0.2
23	0.12	0.976	0.003	33.6	1.5	0.1
45	0.12	0.964	0.003	22.1	4.2	0.0
65	0.12	0.945	0.006	14.2	17.0	0.2
90	0.12	0.925	0.003	10.2	25.7	0.0
45	0.24	0.465	0.003	22.8	5.7	0.0
65	0.24	0.939	0.002	12.7	14.5	0.1
90	0.24	0.927	0.005	10.5	28.4	0.2
65	0.36	0.931	0.002	11.2	16.7	0.2
90	0.36	0.922	0.003	9.8	32.3	0.1
65	0.60	0.922	0.004	9.8	10.0	0.3
90	0.60	0.922	0.006	9.8	30.8	0.2
65	0.84	0.946	0.002	14.5	14.2	0.0
90	0.84	0.923	0.001	9.9	32.7	0.3
90	1.08	0.922	0.001	9.8	32.7	0.0
90	1.32	0.923	0.003	9.9	22.4	0.2

TABLE V-5. SUMMARY OF TEST RESULTS FOR WATER PERMEATION

FILM	CELL TEMPERATURE (°C)	% H ₂ O IN FEED	PERMEATE DRY FLUX cc/cm ² sec	PERMEATE %H ₂ O	PERMEATE WATER FLUX cc/cm ² sec
CELLULOSE ACETATE	90.	0.25	9.0×10^{-4}	24.4	2.9×10^{-4}
CELLULOSE ACETATE	90.	0.85	7.7×10^{-4}	35.5	4.3×10^{-4}
POLYSULFONE	90.	0.36	5.3×10^{-5}	32.9	2.6×10^{-5}
POLYSULFONE	90.	0.60	7.6×10^{-5}	33.9	3.9×10^{-5}
POLYETHERSULFONE	90.	0.85	8.7×10^{-5}	13.9	1.4×10^{-5}
POLYETHERSULFONE	90.	0.96	8.3×10^{-5}	25.2	2.8×10^{-5}
SOLFOLENE MODIFIED	90.	0.36	3.23×10^{-4}	24.9	1.1×10^{-4}
SOLFOLENE MODIFIED	90.	0.50	3.12×10^{-4}	35.7	1.73×10^{-4}

base coordinates of temperature and water content were the variables of this study. The raised area represents the response surface. The stair-step shape of the response surface was produced as a result of the dependence between temperature and the feed gas water content. The area of high water content and low temperature has no physical meaning. The advantage in the use of the plots is that the effects of either variable can be seen clearly.

F. DISCUSSION

1. OVERVIEW

The use of the term "flux" is to describe the flux of the dried permeate unless otherwise noted. The discussion of film flux values does not consider the film thickness. A concentration gradient of water across the film may result in non-uniform film properties. The use of flux values adjusted to a unit thickness may be incorrect.

Flux values for the films varied from less than 0.00001 to 0.0019 cu cm(STP)/sq cm(sec) over the range of conditions investigated. Cellulose acetate gave the highest average flux followed in descending order by modified poly(vinylidene fluoride), polyethersulfone and polysulfone. Flux values for polyethersulfone and polysulfone varied little over the test conditions. Flux values for cellulose acetate and modified poly(vinylidene fluoride) varied up to ten fold.

Separation factors for the films varied from 6 to 34 over the range of conditions investigated. All the films selectively passed

carbon dioxide. Polyethersulfone gave the overall highest separation factor followed by polysulfone, modified poly(vinylidene fluoride) and cellulose acetate. The separation factor of polyethersulfone was approximately twice that of the others.

2. REPRODUCIBILITY OF DATA

Results of the Ellig, Althouse and McCandless study can be compared directly with the data from the moisture free runs of this research(41). Experimental conditions including the operating pressure and feed gas composition were identical. The flux values and separation factors of the two studies with a dry feed gas are plotted in figures V-1 and V-2 respectively.

The separation factors of sulfolene modified poly(vinylidene fluoride) film and cellulose acetate film varied by 5% and 12% between the two studies respectively. The flux values of these films varied by an average value of 18% and 19% respectively.

The polysulfone film demonstrated similar separation values (7% difference). However, flux values of this report are approximately one fourth the cited values. The thickness of the film used in this study was 2.5 times that of the cited study. After correcting for the thickness difference the flux values varied by 31%.

The separation factors and flux values of the polyethersulfone varied to a large degree between the two studies. The separation factors varied by an average of 24%, with the cited values always smaller.

FIGURE V-1. REPRODUCIBILITY OF FLUX DATA

FIGURE V-2. REPRODUCIBILITY OF SEPARATION FACTORS

The flux values at 25°C were an order of magnitude different. The difference decreased to approximately 20% at 90°C. The thickness of the film used in this study is 3.0 times that of the cited study. After correcting for the thickness difference the flux values varied by 45%. The thickness difference should have little influence on the separation factors.

Some work has been done on the permeation through membranes with various processing and treatment histories(42). In tests with polysulfone the sorption capacity for carbon dioxide varied considerably with the history of the specimen. The difference between the studies may be caused by different production methods for the polysulfone and polyethersulfone films.

3. CELLULOSE ACETATE MEMBRANE

Graphs of the separation factors versus feed stream water content and flux versus feed stream water content are shown in figures V-3 and V-4 respectively. The plot of the separation factor values versus feed stream water content and temperature is shown in Figure V-5. The plot of flux versus water content and temperature is shown in Figure V-6.

The cellulose acetate membrane behaved in the following manner:

- a) An increase in temperature decreased the separation factor.
- b) An increase in feed gas moisture decreased the separation factor.

FIGURE V-3. SEPARATION FACTOR VS. PERCENT WATER IN FEED GAS FOR CELLULOSE ACETATE

FIGURE V-4. FLUX VS. PERCENT WATER IN FEED GAS
FOR CELLULOSE ACETATE FILM

FIGURE V-5. PLOT OF SEPARATION FACTOR VS. FEED STREAM WATER CONTENT AND TEMPERATURE FOR CELLULOSE ACETATE FILM

FIGURE V-6. PLOT OF FLUX VS. FEED STREAM WATER CONTENT AND TEMPERATURE FOR CELLULOSE ACETATE FILM

- c) The separation factor varies less than the other films between temperatures as the feed gas moisture increases.
- d) An increase in water vapor from 0. to 0.12% caused the separation factor to decrease at all temperatures. The largest decrease was at room temperature by a factor of three.
- e) At 65 and 90°C separation factors were fairly constant as water content was varied.
- f) An increase in temperature increased the flux.
- g) The flux was greater with the feed gas saturated than with a dry feed gas.
- h) At 45, 65 and 90°C flux was a maximum at a feed gas moisture content between the dry and saturated condition.
- i) The flux at 0.12% water was greater than twice the dry flux at 65 and 90°C.

4. POLYSULFONE MEMBRANE

Graphs of the separation factors versus feed stream water content and flux versus feed stream water content are shown in figures V-5 and V-6 respectively. The plot of the separation factor values versus water content and temperature is shown in Figure V-9. The plot of flux versus water content and temperature is shown in Figure V-10. The polysulfone membrane behaved in the following manner:

- a) An increase in membrane temperature decreased the separation factor.

FIGURE V- 7 SEPARATION FACTOR VS. PERCENT WATER IN
FEED GAS FOR POLYSULFONE FILM

FIGURE V-8. FLUX VS. PERCENT WATER IN FEED GAS FOR POLYSULFONE FILM

FIGURE V-9. PLOT OF SEPARATION FACTOR VS. FEED STREAM WATER CONTENT AND TEMPERATURE FOR POLYSULFONE FILM

FIGURE V-10. PLOT OF FLUX VS. FEED STREAM WATER CONTENT AND TEMPERATURE FOR POLYSULFONE FILM

- b) An increase in feed gas moisture decreased the separation factor.
- c) As membrane temperature was increased, water vapor had a smaller effect on the separation factor.
- d) At 23°C and 45°C, an increase in water vapor from 0. to 0.12% caused a large (>25%) drop in the separation factor.
- e) At 65°C an increase in water vapor from 0. to 0.12% caused a 15% increase in the separation factor.
- f) At 90°C the separation factor remained approximately constant at any feed gas moisture content.
- g) An increase in temperature increased the flux.
- h) At 23°C an increase in feed gas moisture from 0. to 0.12% caused a 20% decrease in flux.
- i) At 45 ,65 and 90°C the flux was a maximum at a feed moisture content between the dry and saturated condition.
- j) At 45 ,65 and 90°C an increase in the feed gas moisture from 0. to 0.12% approximately doubled the flux. This was the largest increase in flux as moisture was varied.

5. POLYETHERSULFONE MEMBRANE

Graphs of the separation factors versus feed stream water content and flux versus feed stream water content are shown in figures V-7 and V-8 respectively. The plot of the separation factor values versus water content and temperature is shown in Figure V-13. The Plot of

FIGURE V-11. SEPARATION FACTOR VS. PERCENT WATER
IN FEED GAS FOR POLYETHERSULFONE FILM

FIGURE V-12 FLUX VS. PERCENT WATER IN FEED GAS
FOR POLYETHERSULOFNE FILM

FIGURE V- 13. PLOT OF SEPARATION FACTOR VS. FEED STREAM WATER CONTENT AND TEMPERATURE FOR POLYETHERSULFONE FILM

flux versus water content and temperature is shown in Figure V-14. The polyethersulfone membrane behaved in the following manner:

- a) An increase in the gas moisture decreased the separation factor.
- b) An increase in temperature decreased the separation factor.
- c) An increase in water vapor from 0. to 0.12% caused a large (>30%) drop in separation factor.
- d) An increase in temperature increased the flux.
- e) The flux with the feed gas saturated was greater than the flux with a dry feed gas.
- f) Flux reached a maximum at 65°C with 0.36% feed gas water content.

6. SULFOLENE MODIFIED POLY(VINYLLIDENE FLOURIDE) MEMBRANE

Graphs of the separation factors versus feed stream water content and flux versus feed stream water content are shown in figures V-9 and V-10 respectively. The plot of the separation factor values versus water content and temperature is shown in Figure V-17. The plot of flux versus water content and temperature is shown in Figure V-18. The modified poly(vinylidene fluoride) membrane behaved in the following manner:

- a) The separation factor increased as the water content of the feed gas increased at 23 and 45°C.

FIGURE V-14. PLOT OF FLUX VS. FEED STREAM WATER CONTENT AND TEMPERATURE FOR POLYETHERSULFONE FILM

FIGURE V-15. SEPARATION FACTOR VS. PERCENT WATER IN FEED GAS FOR SULFOLENE MODIFIED POLY(VINYLIDENE FLUORIDE) FILM

FIGURE V-16. FLUX VS. PERCENT WATER IN FEED GAS
FOR SULFOLENE MODIFIED POLY
(VINYLIDENE FLUORIDE) FILM

FIGURE V-17. PLOT OF SEPARATION FACTOR VS. FEED STREAM WATER CONTENT AND TEMPERATURE FOR SULFOLENE MODIFIED POLY(VINYLIDENE FLUORIDE) FILM

FIGURE V-19. PLOT OF FLUX VS. FEED STREAM WATER CONTENT AND TEMPERATURE FOR SULFOLENE MODIFIED POLY(VINYLIDENE FLUORIDE) FILM

- b) The separation factor decreased or remained approximately constant as the water content increased at 65°C.
- c) The separation factor remained approximately constant as the water content is varied at 90°C.
- d) The separation factor decreased as the temperature was increased.
- e) The flux increased as the temperature increased.
- f) The difference in flux values between 65 and 90°C did not vary as the feed gas moisture increased.
- g) The flux at 23 and 45°C generally increased as water content was increased.
- h) The flux at 23 and 45°C increased as water content was increased.
- i) The flux at 65 and 90°C generally decreased as the water content was increased.

7. WATER VAPOR FLUX

A graph of the water vapor flux versus feed stream water content is shown in Figure V-11. Cellulose acetate allowed an order of magnitude larger water vapor flux than the other films. The water vapor flux of the sulfolene modified film increased exponentially between the three conditions tested as water vapor in the feed increased.

The driving force for permeation of the water was at most a fiftieth of the driving force of the gases. Yet water permeation accounted

FIGURE V-19. WATER VAPOR FLUX VS. PERCENT WATER IN FEED STREAM FOR THE FILMS

for up to 50% of the total permeate. The films had a much higher permeability for water vapor than either carbon dioxide or methane.

8. EFFECT OF WATER ON FLUX

Polysulfone, polyethersulfone and cellulose acetate demonstrated similar behavior in terms of flux at elevated temperatures. The addition of a small amount of water (0.12 to 0.36%) increased the flux two to three fold. As the water content of the feed gas was increased further the flux became constant or decreased. The flux of the sulfone modified poly(vinylidene fluoride) film varied to a lesser degree than the other three films and had no observable peak flux.

Plasticizing effects due to the concentration of water in the membrane was apparent. The addition of a plasticizer allows for an increased mobility of polymer chain segments, as well as altering the solubilities of gases in the polymer. The net effect of a modifier in a film is an increase in gas permeation rates. Polyethersulfone, polysulfone and cellulose acetate films demonstrated an increase in gas permeation rates as the water content of the feed was altered.

The increase in polymer chain segment mobility caused by a modifier should allow a greater flux of methane and thereby decrease the separation factor. As water was introduced at isothermal conditions, the separation factor of polysulfone, polyethersulfone and cellulose acetate decreased.

Unlike modified films, additional water vapor did not increase the flux for the commercial films. As discussed earlier, the solubility of water in most films increases in a nonlinear fashion with pressure. After an initial two or three fold increase in flux for the commercial films the flux values either became fairly constant or drop. It was at these operating conditions that water vapor permeation becomes important. Flux of water vapor increased as the flux of the gases tended to become constant or decreased. Cellulose acetate flux varied the most with an initial three fold jump in flux followed by a drop to half the peak value at 90°C. Cellulose acetate also demonstrated the highest water vapor flux at these conditions.

It was evident that after an initial plasticizing effect water permeation controls gas permeation.

The discussion to this point has been limited to the three commercial films. The sulfolene modified film did not demonstrate the flux and separation factor characteristics common to the others. Flux generally decreased slightly as water was added. The separation factor actually increased at lower temperatures as water was added. The modified film was the only film tested for which water vapor flux increased in near exponential fashion with increased water in the film.

It appears that the film passed water vapor so well that there was little interaction between the polymer and water. The result of flux

and separation factor variations may be explained as small changes in the diffusivity and solubility constants in the film.

CONCLUSIONS AND RECOMMENDATIONS

A. CONCLUSIONS

1. All films tested were very effective in the separation of carbon dioxide from the methane and carbon dioxide mixture of this study. Of the four films tested, cellulose acetate has the highest average flux. The variation in the percent of carbon dioxide in the permeate was so small for the films and conditions tested that the film with the highest flux should also be considered as the best overall membrane for the separation of the dry gas mixture. On a unit thickness basis, this film would be polyethersulfone..

2. Permeate flux has a strong tendency to increase with increasing temperature, while the separation factor decreases.

3. The water flux for the films tested exhibited a nonlinear dependence on the partial pressure of water in the feed gas.

4. Water exerts a plasticizing influence on the membrane.

5. The degree of the water's plasticizing effect varies to a large degree between the films. Gas flux in the solfolene modified film appears almost independent of water permeation. The flux of cellulose acetate varies drastically as a function of feed gas water content. The ability to control feed gas water content in the 0.12% range would determine which membrane would be the best for separating a wet gas.

B. RECOMMENDATIONS

1. Direct comparison of flux values and separation factor values for the films tested is not possible because different film thicknesses

were used in the tests. It is expected that flux would be inversely proportional to the membrane thickness for a dry feed gas. However, as water is added to the feed stream a concentration gradient is established across the membrane. This gradient would no longer make flux inversely proportional to thickness. Tests should be expanded to include several thicknesses for each film.

2. Flux values for the individual components should be determined as functions of temperature and pressure. A model of a complete n-staged purification system could then be developed. Hydrogen sulfide should be included in the model.

3. A better method should be developed to measure flux and composition on a wet basis. Assuming a method could be devised, more data should be obtained at the lower temperatures.

REFERENCED FOOTNOTES

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APPENDIX

TABLE OF NOMENCLATURE

C	Concentration
D	Diffusivity
D ⁰	Temperature independent diffusivity
E	Activation energy for diffusion
h	Apparent heat of solution
H	Henry's law constant
H ⁰	Temperature independent Henry's law constant
L	Membrane thickness
N	Mass flux
P	Total pressure
p	Partial pressure
R	Gas constant
S	Solubility of gas or vapor in the polymer
S ⁰	Temperature independent solubility
T	Absolute temperature
t	Time
X	Mole fraction in the feed
Y	Mole fraction in permeate
Z	Length coordinate
α	Separation factor

SUBSCRIPTS

a	Component a
b	Component b
m	Membrane property
o	Interface
p	permeate
s	source

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