

2

SEC

AD-A211 420

PORT DOCUMENTATION PAGE

FILE COPY

|                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        |                                              |                                                                                             |                           |
|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|----------------------------------------------|---------------------------------------------------------------------------------------------|---------------------------|
| 1a. <b>AD-A211 420</b>                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 |                                              | 1b. RESTRICTIVE MARKINGS<br>NA                                                              |                           |
| 2a. NA                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                 |                                              | 3. DISTRIBUTION / AVAILABILITY OF REPORT<br>Distribution unlimited                          |                           |
| 2b. DECLASSIFICATION / DOWNGRADING SCHEDULE<br>NA                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                      |                                              | 5. MONITORING ORGANIZATION REPORT NUMBER(S)<br>NA                                           |                           |
| 4. PERFORMING ORGANIZATION REPORT NUMBER(S)<br>University of California                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                |                                              | 7a. NAME OF MONITORING ORGANIZATION<br>Office of Naval Research                             |                           |
| 6a. NAME OF PERFORMING ORGANIZATION<br>University of California                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        | 6b. OFFICE SYMBOL (if applicable)<br>NA      | 7b. ADDRESS (City, State, and ZIP Code)<br>800 N. Quincy Street<br>Arlington, VA 22217-5000 |                           |
| 6c. ADDRESS (City, State, and ZIP Code)<br>Department of Zoology<br>University of California<br>Davis, California 95616                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                |                                              | 9. PROCUREMENT INSTRUMENT IDENTIFICATION NUMBER<br>N00014-85-K-0242                         |                           |
| 8a. NAME OF FUNDING / SPONSORING ORGANIZATION<br>Office of Naval Research                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                              | 8b. OFFICE SYMBOL (if applicable)<br>ONR     | 10. SOURCE OF FUNDING NUMBERS                                                               |                           |
| 8c. ADDRESS (City, State, and ZIP Code)<br>800 N. Quincy Street<br>Arlington, VA 22217-5000                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                            |                                              | PROGRAM ELEMENT NO.<br>61153N                                                               | PROJECT NO.<br>RR04108    |
|                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        |                                              | TASK NO.<br>441K712                                                                         | WORK UNIT ACCESSION NO.   |
| 11. TITLE (Include Security Classification)<br>Role of water in proton-hydroxide conductance across model and biological membranes                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                     |                                              |                                                                                             |                           |
| 12. PERSONAL AUTHOR(S)<br>Deamer, David W.                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             |                                              |                                                                                             |                           |
| 13a. TYPE OF REPORT<br>Annual                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                          | 13b. TIME COVERED<br>FROM 10-1-88 TO 9-30-89 | 14. DATE OF REPORT (Year, Month, Day)                                                       | 15. PAGE COUNT            |
| 16. SUPPLEMENTARY NOTATION                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                             |                                              |                                                                                             |                           |
| 17. COSATI CODES                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                       |                                              | 18. SUBJECT TERMS (Continue on reverse if necessary and identify by block number)           |                           |
| FIELD                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  | GROUP                                        | SUB-GROUP                                                                                   |                           |
|                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        |                                              |                                                                                             |                           |
|                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                        |                                              |                                                                                             |                           |
| 19. ABSTRACT (Continue on reverse if necessary and identify by block number)<br><p>The goal of this research is to understand the mechanism of proton translocation in model and biological membranes. The significance is that electrochemical proton gradients and proton transport are central to many bioenergetic functions of cells. It is essential to address both the barrier function of membranes, and the process by which protons are directed to specific reaction sites within the membrane. The primary theme concerns the role of water in proton translocation, and particularly the possibility that hydrogen-bonded water structures contribute to the transport process. In previous studies, we have shown that proton permeability of lipid bilayers is orders of magnitude greater than that of other cations. To explain this observation, we suggested that transient hydrated defects are continuously being produced by thermal fluctuations in the lipid. If the water in the defects were associated by hydrogen bonding, protons could cross the defect by hydrogen bond exchange along the associated water molecules, accounting for the high relative permeability of protons. This concept can be extended to other membrane systems. Gramicidin A contains a 0.4 nm channel which conducts protons through hydrogen bonded water, thereby providing an important model for investigating such processes. The Fo subunit of coupling membranes has the function of directing protons to the site of ATP synthesis in the F1 ATPase. In the work planned for next year, biophysical aspects of proton transport will be studied in lipid bilayers, gramicidin channels and the Fo subunit of coupling membranes.</p> |                                              |                                                                                             |                           |
| 20. DISTRIBUTION / AVAILABILITY OF ABSTRACT<br><input checked="" type="checkbox"/> UNCLASSIFIED/UNLIMITED <input type="checkbox"/> SAME AS RPT. <input type="checkbox"/> DTIC USERS                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                    |                                              | 21. ABSTRACT SECURITY CLASSIFICATION<br>(U)                                                 |                           |
| 22a. NAME OF RESPONSIBLE INDIVIDUAL<br>Dr. A.J. Majde                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                                  |                                              | 22b. TELEPHONE (Include Area Code)<br>(916) 752-2175                                        | 22c. OFFICE SYMBOL<br>ONR |

89 8 16 035

## Introduction

The overall thrust of the work supported by the Office of Naval Research in our laboratory has been to better understand the role of water in ion transport. We have particularly focused on proton flux mechanisms, building on our observation that protons diffuse across lipid bilayer membranes by a process quite different from that of other cations (Nichols and Deamer, 1980; Deamer and Nichols, 1983). As a working hypothesis, we proposed that proton equivalents move along hydrogen bonded chains of water molecules which occur in transient defects in the bilayer. We have extended this concept to the action of certain membrane perturbants on the bilayer. We are also working with a known hydrogen bonded chain of water - the gramicidin channel - and have proposed that proton conductance along similar water structures may play a role in biological membranes as well (Deamer and Nichols, 1989).

The results provide insight into the nature of the lipid bilayer, and the manner in which hydrated defects contribute to ion permeation across the bilayer barrier. It has also permitted us to better understand the effects of anesthetic molecules on the ability of synaptic vesicles to maintain proton gradients necessary for neurotransmitter uptake. Finally, recent results suggest that hydrogen bonded chains of water may be involved in conducting proton equivalents through the Fo subunit of coupling membranes, and we are now in a position to test this exciting possibility in reconstituted planar membranes.

## Progress Report.

Publications related to ONR contract NO0014-85-K:

1. Barchfeld, G.L. and Deamer, D.W. (1985) Effect of general anesthetics on proton and potassium permeability of liposomes. *Biochim. Biophys. Acta* 189:161-169.
2. Deamer, D.W. and Gutknecht, J. (1986) *Methods Enzymol.* 127:471-480.
3. Deamer, D.W. and Bramhall, J. (1986) Permeability of lipid bilayers to water and ionic solutes. *Chem. Phys. Lipids* 40:167-181.
4. Deamer, D.W. (1987) Proton permeability of lipid bilayers. *J. Bioenerg. Biomembr.* 19:457-479.
5. Barchfeld, G.L. and Deamer, D.W. (1988) Effect of alcohols on proton permeability: relation to general anesthetics. *Biochim. Biophys. Acta* 944:40-48.
6. Deamer, D.W. and Nichols, J.W. (1989) Proton flux in model and biological membranes. *J. Membr. Biol.* 107:91-103.
7. Akeson, M. and Deamer, D.W. (1989) A test of the pump-leak hypothesis for general anesthesia. *Biochemistry* 28:5120-27.
8. Deamer, D.W. and Akeson, M. (1989) General anesthetics and membranes: a critical review. *Advances in Membrane Fluidity*. (In press).
9. Deamer, D.W. Annual reports to the Office of Naval Research in 1986, 1987 and 1988, and distribution to other Principal Investigators.

Our original goals over the past contract period can be summarized as follows:

1. To understand the role of proton flux in the action of general anesthetics.



|     |  |
|-----|--|
| 1   |  |
| 2   |  |
| 3   |  |
| 4   |  |
| 5   |  |
| 6   |  |
| 7   |  |
| 8   |  |
| 9   |  |
| 10  |  |
| 11  |  |
| 12  |  |
| 13  |  |
| 14  |  |
| 15  |  |
| 16  |  |
| 17  |  |
| 18  |  |
| 19  |  |
| 20  |  |
| 21  |  |
| 22  |  |
| 23  |  |
| 24  |  |
| 25  |  |
| 26  |  |
| 27  |  |
| 28  |  |
| 29  |  |
| 30  |  |
| 31  |  |
| 32  |  |
| 33  |  |
| 34  |  |
| 35  |  |
| 36  |  |
| 37  |  |
| 38  |  |
| 39  |  |
| 40  |  |
| 41  |  |
| 42  |  |
| 43  |  |
| 44  |  |
| 45  |  |
| 46  |  |
| 47  |  |
| 48  |  |
| 49  |  |
| 50  |  |
| 51  |  |
| 52  |  |
| 53  |  |
| 54  |  |
| 55  |  |
| 56  |  |
| 57  |  |
| 58  |  |
| 59  |  |
| 60  |  |
| 61  |  |
| 62  |  |
| 63  |  |
| 64  |  |
| 65  |  |
| 66  |  |
| 67  |  |
| 68  |  |
| 69  |  |
| 70  |  |
| 71  |  |
| 72  |  |
| 73  |  |
| 74  |  |
| 75  |  |
| 76  |  |
| 77  |  |
| 78  |  |
| 79  |  |
| 80  |  |
| 81  |  |
| 82  |  |
| 83  |  |
| 84  |  |
| 85  |  |
| 86  |  |
| 87  |  |
| 88  |  |
| 89  |  |
| 90  |  |
| 91  |  |
| 92  |  |
| 93  |  |
| 94  |  |
| 95  |  |
| 96  |  |
| 97  |  |
| 98  |  |
| 99  |  |
| 100 |  |

dist  
A-1

Codes

d/or  
Special

2. To characterize the effects of transmembrane peptides on proton permeation of bilayers, and to relate this to bioenergetic functions of coupling membranes.

3. To establish a model system for measuring proton flux along hydrogen bonded chains of water in hydrophobic phases.

We have completed our research on the first aim, with three published papers which will be described below. A review is also in press in *Advances in Membrane Fluidity*. We have made substantial progress on the second aim, with two published papers/reviews, and a manuscript in preparation. The results will guide our present aims and will be described below. We decided not to initiate research on the third aim, because a more powerful approach - microbilayers on a modified patch clamp device - is now established in our laboratory. We intend to use this in future research.

### **General anesthetics and proton permeability.**

A variety of compounds perturb lipid bilayers in such a way that the barrier to ion flux is reduced. This can lead to inhibition of certain membrane functions that depend on ionic gradients. The classic example is the effect of uncoupling agents on mitochondrial and thylakoid membranes. Uncouplers introduce "leaks" that allow gradients of protons or other ions to decay, with the result that chemiosmotic phosphorylation is uncoupled from electron transport.

Physical perturbation of membranes has also been used to account for the action of general anesthetics (for review, see Akesson and Deamer, 1989). The perturbing effect on physical properties of lipid bilayers - fluidity, volume, phase transitions - does not appear to be sufficient to explain anesthetic effects (Franks and Lieb, 1978; 1982; 1984; 1986). However, Bangham and co-workers have suggested that increased permeability to ions may in fact offer a unitary explanation for the action of general anesthetics. These considerations led to the pump-leak hypothesis (Bangham and Mason, 1980; Bangham and Hill, 1986) which integrates several aspects of anesthetic effects on the bilayer phase of nerve cell membranes. First, it focuses on synaptic transmission, which is the most plausible cellular site of anesthetic action. Second, it incorporates the effect of general anesthetics on lipid bilayer ionic permeability in a way that accounts for other anesthetic effects, particularly the manner in which cold, heat and hypoxia might produce anesthesia. It therefore offers a general and broadly based explanation of anesthetic action on any organism sufficiently complex to have a nervous system.

The pump-leak hypothesis notes that synaptic vesicles have an ATP-dependent proton transport enzyme (the pump) which works against a continuing proton leak across the vesicle membrane. The pump is required to maintain an electrochemical proton gradient equivalent to perhaps 2 pH units, which in turn is used as an energy source to concentrate neurotransmitters such as catecholamines and indolamines (Figure 1). The hypothesis proposes that anesthetics increase membrane permeability to protons, leading to a collapse of the gradient. As a result, neurotransmitters are lost from the vesicles, synaptic transmission is inhibited, and if this occurs in sensitive portions of the CNS, anesthesia ensues.

The pump leak hypothesis is valuable because it makes clear predictions that can be tested experimentally. Results of such tests have provided information about quantitative aspects of anesthetic effects on lipid bilayers. For instance, one would expect anesthetics to cause marked increments in

bilayer permeability to protons in model systems. Second, anesthetics should release neurotransmitters such as catecholamines from vesicles in which they have accumulated in response to a proton gradient. Finally, these effects must be consistent with the principles stated in the introduction. That is, they will be measureable at ED<sub>50</sub> concentrations of anesthetics, and the kinetics of the process should match the known time factors with which anesthetics produce the anesthetic state.

Bangham and Mason (1980) measured effects of benzyl alcohol and other anesthetics (halothane, chloroform, butanol) on permeability of synaptic vesicles isolated from rat brain. The vesicles were shown to accumulate labeled dopamine in an ATP-dependent process corresponding to their function in the synapse. It was demonstrated that benzyl alcohol did in fact increase proton permeability in a dose-dependent manner, with the result that the dopamine was released. It was also shown that liposomes with pH gradients accumulate dopamine, which was released at a more rapid rate in the presence of benzyl alcohol, butanol or halothane. Barchfeld and Deamer (1985, 1988) in research supported by the Office of Naval Research, confirmed these results in liposome systems, and extended the observations to comparative proton and potassium permeabilities. Several general anesthetics produced similar increments in both proton and potassium permeabilities, demonstrating that the leak was due to a general defect in the bilayer permeability barrier, and not specific for protons.

Despite the early positive results, significant questions remained. At ED<sub>50</sub> concentrations, anesthetic effects on permeability were minimal. For instance, if we assume that a significant permeability increment would be a doubling of the proton permeability coefficient, concentrations of anesthetics several times ED<sub>50</sub> are required. A second concern is that anesthetic-induced decay of proton gradients had half-times in the range of 15 - 30 minutes, much longer than required for the onset of anesthesia in organisms.

To define the pump-leak mechanism more precisely, Akeson and Deamer (1989) initiated direct measurements of catecholamine loss from chromaffin granules. This system was chosen because it has a robust proton ATPase activity which produces large proton gradients. Furthermore, the electrochemical proton gradient and membrane potential are responsible for the accumulation and maintenance of internal catecholamines to concentrations approaching 0.4 M (Johnson, 1988). These factors permitted direct measurements of ATPase activity, proton permeability and catecholamine flux in the presence and absence of anesthetics.

Some results are shown in Figure 2 with corresponding values from synaptic vesicles plotted on the same time scale for comparison. It is clear that there is a measureable loss of catecholamine from the chromaffin granules at ED<sub>50</sub> concentrations of anesthetics. This correlates closely with an increased proton permeability which was also measured, and a corresponding decrease in the magnitude of the pH gradient. Moreover, the kinetics of catecholamine efflux are similar in the chromaffin granules and synaptic vesicles.

These positive results qualitatively agree with the pump-leak hypothesis. However, analysis at a more quantitative level is less favorable. First, the leak is measureable only because of the sensitivity of the method. In fact, the pH gradient in chromaffin granules decays just 0.05 pH units in 20 minutes, and perhaps 5% of the catecholamine is lost. Clearly, the release is slow. Using the principle that an anesthetic effect must correlate with onset times in organisms, the amount of catecholamine lost in the first minute following anesthetic addition is miniscule. Even when the pH gradient was

completely released by addition of uncouplers or ammonium chloride, loss of catecholamine had half times of 30 minutes.

In summary, we have concluded that anesthetic effects on ionic permeability of membranes, while measurable, is not sufficient to account for the effects of general anesthetics on the nervous system. Our publication in *Biochemistry* summarizes these results, and we will not continue further investigative effort on this anesthetic mechanism. Instead, there is good evidence that anesthetics such as the benzodiazepines, barbiturates, steroids and alcohols all act on the GABA receptor, and we have initiated collaborative research with Dr. Mark McNamee (Biochemistry, UC Davis) which will focus on this receptor, particularly the effects of steroids.

### **The nature of the proton-conducting bilayer defect.**

#### Plots of proton flux and driving force

Nagle (1987) proposed three mathematical treatments of proton flux through transient hydrogen-bonded defects. These depend on the lifetime of the defect and the mechanism of proton transfer. Each model produces a characteristic curve when driving force ( $\Delta\text{pH}$  or voltage) is plotted against flux. Results from earlier investigations show a surprising lack of consensus for such curves. In an attempt to clarify such results for lipid bilayers, we have made a series of careful measurements for liposome systems in which  $\Delta\text{pH}$  was the driving force. These are shown in Figure 3, together with similar results from other laboratories. There is general agreement that proton flux is linear with  $\Delta\text{pH}$ .

The conclusion that the relationship between proton flux and  $\Delta\text{pH}$  is linear suggests a transport mechanism in which the effect of the rate limiting step increases exponentially with driving force. Our next goal is to determine the nature of the rate limitation by direct comparisons of voltage-driven proton currents in liposomes and PLM systems. This will also test the generality of the result, assuring that it is not related to arbitrary choices of lipid, driving force and buffers.

#### Chain length effects lipid bilayer permeability to protons

One of the most convincing demonstrations that transient hydrated defects are involved in ion permeation of lipid bilayers is illustrated in Figure 4 (unpublished results). In this experiment, we measured proton permeability of homologous phospholipids containing hydrocarbon chains ranging from 14 to 22 carbons. The remarkable observation is that permeability increases approximately an order of magnitude between C14 and C18, even though this represents only a fractional decrease in the thickness of the bilayer.

Our interpretation of these results is that the hydrophobic effect maintaining bilayer stability becomes increasingly weaker with shorter chain lengths, so that thermal fluctuations become more numerous or longer lived. Finally, at ten carbons, thermal fluctuations overwhelm stabilizing forces and the phospholipid becomes micellar. One of our goals in future work is to extend these observations to C10 and C12 lipid membranes, using both voltage and concentration gradients to drive protonic current. One significant aspect of this work is that it may explain why membranes maintain a given chain length. For instance, if mitochondrial lipids were on average four carbons shorter, they would be ten times more permeable to protons, and

the electron transport coupled to proton pumping would be unable to keep up with the leak.

### Proton flux in the gramicidin channel

One of the most striking characteristics of proton flux in bilayers is the independence of flux and hydrogen ion concentration. If the gramicidin channel is a useful model of the defect occurring in bilayers, one would expect a similar independence for proton flux through the channel. However, Gutknecht (1987) has reported that conductance through the channel is linearly related to hydrogen ion concentration (Figure 5).

We have carried out preliminary comparisons of such data in the liposome systems, and observed independence of flux and pH, also illustrated in Figure 5. The results are necessarily over a restricted range, due to limitations of the pyranine dye system, but are convincingly different from the PLM results. Because the proton current/voltage relationship in gramicidin should be characteristic of conductance along hydrogen bonded water chains, we wish to compare gramicidin and the Fo subunit in this regard. One of our goals is therefore to resolve this apparent discrepancy by direct comparisons of liposomes and PLM systems.

In other preliminary work with gramicidin, we have established that proton flux in D<sub>2</sub>O is approximately half that in H<sub>2</sub>O under identical experimental conditions, while potassium flux is unaffected. Similar results have been reported by Lear et al. (1988) and by Lill et al. (1987) in other proton channels. This is a significant result, in that an isotope effect would provide a useful probe for characterizing proton conductance along tHBC in biological membranes. In the research proposed here, we intend to confirm and extend this observation.

### REFERENCES

- Akeson M and Deamer DW (1989): A test of the pump-leak hypothesis of general anesthesia. *Biochemistry* 28:5120-27.
- Akeson M and Deamer DW (1989): General anesthetics and membranes: a critical review. *Adv. Membr. Fluidity*. (In press)
- Bangham AD, and Mason WT (1980): *Anesthesiol* 53: 135-141.
- Bangham AD, Hill MW, Mason WT (1980): *Molecular Mechanisms of Anesthesia*, Vol 2, Fink BR (ed) Raven Press, NY pp 69-77
- Bangham AD, Hill MW (1986): *Chem Phys Lipids* 40: 189-205.
- Barchfeld, G and Deamer DW (1985) *Biochim Biophys Acta* 819:161-167.
- Barchfeld, G and Deamer DW (1988) *Biochim Biophys Acta* 944:40-48.
- Franks NP, Lieb WR (1978) *J Mol Biol* 133: 469-500.
- Franks NP, Lieb WR (1982): *Nature* 300: 487-493.
- Franks NP, Lieb WR (1984): *Nature* 310:599-601.

- Franks NP, Lieb WR (1986) Proc Natl Acad Sci USA 83: 5116-5120.
- Gutknecht, J. (1987) J. Bioenerg. Biomembr. 19: 427-442.
- Johnson, R. (1988) Physiol. Rev. 68: 232-307.
- Lear, J.D., Wasserman, Z.R., and DeGrado, W.F. (1988) Science 240:1177-1181.
- Nagle, J. (1987) J. Bioenerg. Biomembr. 19: 413-426.

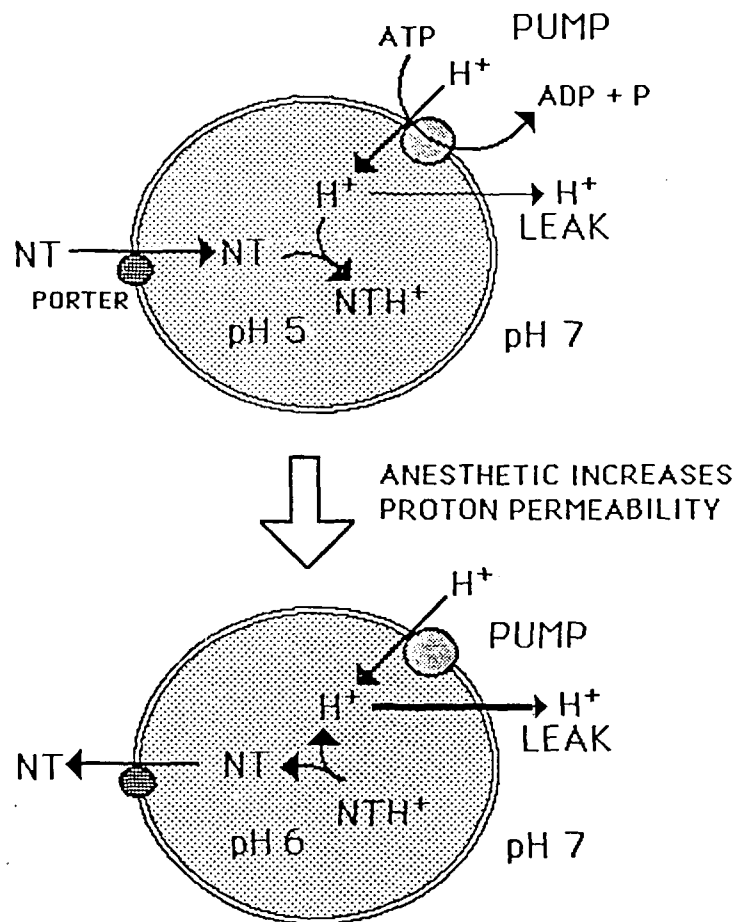


Figure 1. The pump-leak hypothesis. Protons are pumped into the interior of a synaptic vesicle, as shown here, and the interior pH decreases to near 5. The proton gradient can provide an energy source for the movement of weak bases such as catecholamines. If the membrane is made more permeable to protons, for instance by anesthetic molecules partitioning into the membrane, the pH gradient decays and neurotransmitter is released, thereby disabling synaptic transmission. For purposes of illustration, the sketch exaggerates the magnitude of the pH shift, which is actually closer to 0.05 pH units. It is also simplified, because the pH gradient alone represents only part of the energy available for catecholamine uptake (Johnson, 1988).



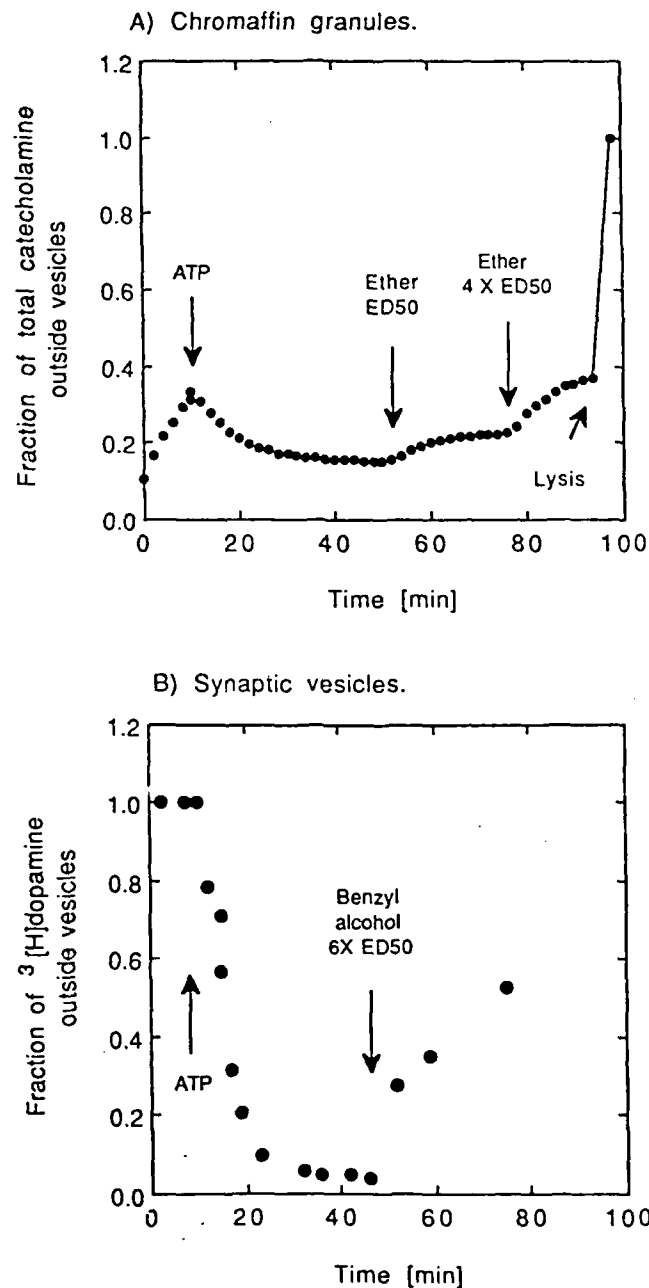


Figure 2: Comparison of volatile anesthetic effects on catecholamine loss from chromaffin granules (top - Akeson and Deamer, 1989) and synaptic vesicles (bottom - Bangham and Mason, 1980). The results for chromaffin granules measured loss from the total pool of internal catecholamines, while the data for synaptic vesicles measured only the loss of labeled dopamine which had been previously accumulated. See text for details.

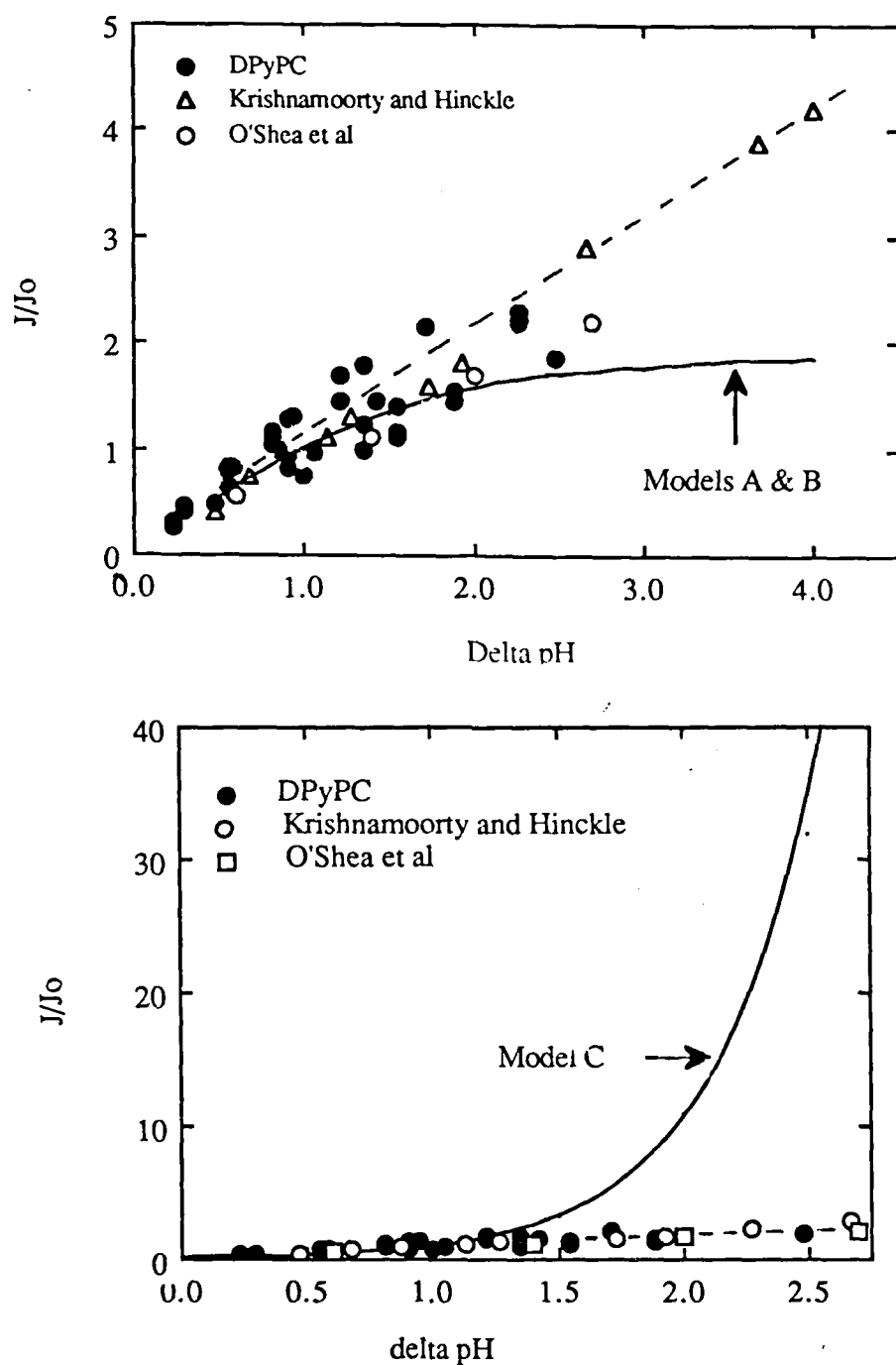


Figure 3. Plots of proton flux (normalized) as a function of driving force, in this case pH gradients across liposome membranes. Our data for diphytenoyl phosphatidylcholine (unpublished) is shown as closed circles, and compared with lines calculated from models A, B and C of Nagle (1987). Normalized values are also calculated from data of other investigators. The results best fit a linear relationship (dashed line) rather than sub- or superlinear.

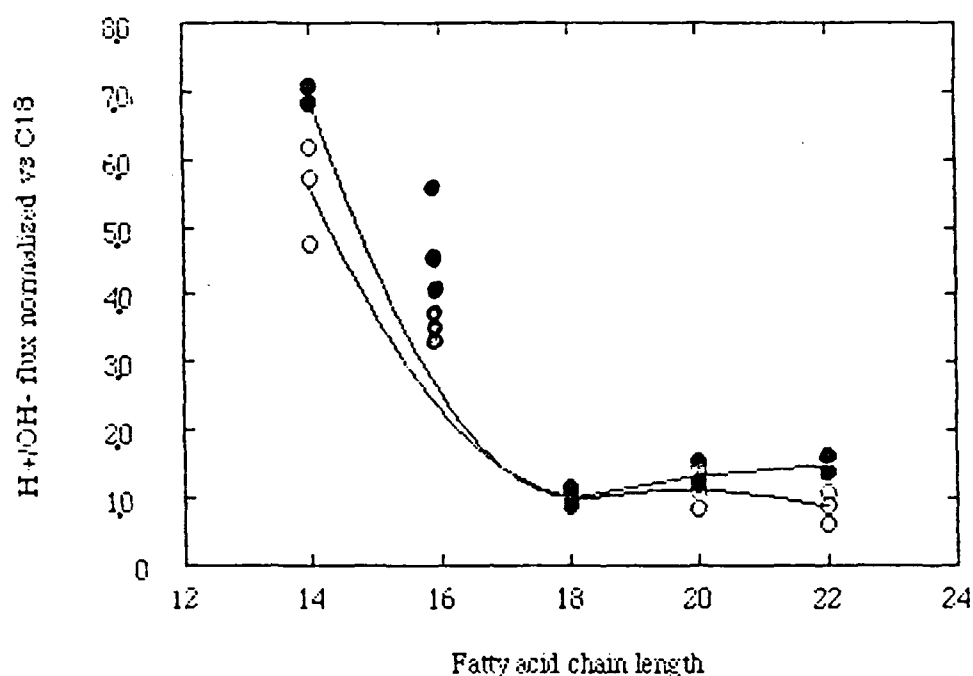


Figure 4. Effect of chain length on proton flux. Unpublished results are shown from six different experiments in which proton flux under standardized conditions was driven by 1 pH unit gradient. As expected, decreasing chain length in lipid bilayers led to marked proton permeability increments.

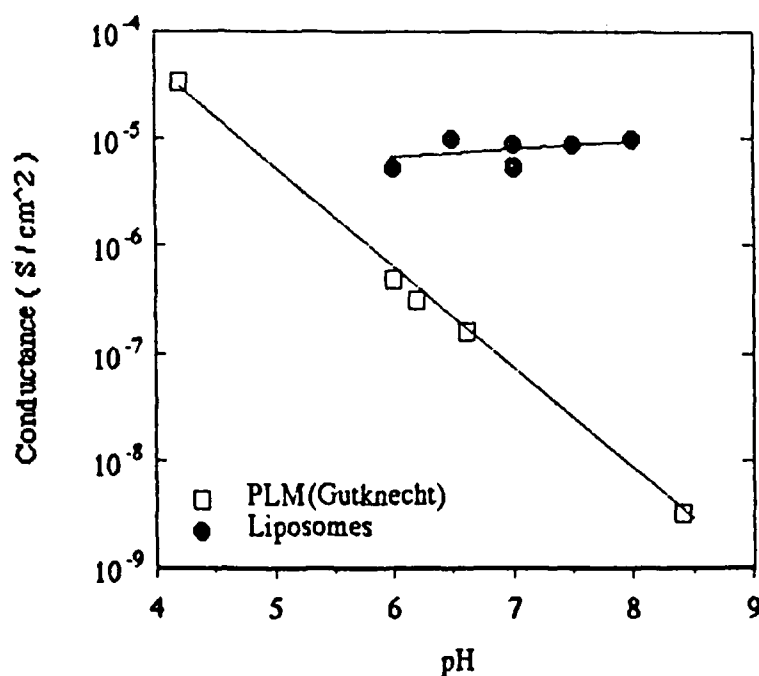


Figure 5. Comparison of pH effects on proton flux through gramicidin embedded in PLM (Gutknecht, 1987) and liposomes (Akeson and Deamer, unpublished). Proton current in PLM was driven by voltage, while the flux in liposomes was driven by pH gradients. (See text for details.)