OOGENESIS IN TUBULARIA LARYNX AND TUBULARIA INDIVISA (HYDROZOA, ATHECATA)

BARRY W. SPRACKLIN

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OOGENESIS IN TUBULARIA LARYNX AND TUBULARIA INDIVISA (HYDROZOA, ATHECATA)

Abstract
In Tubularia larynx and T. indivisa, interstitial cells proliferate in the ectoderm of the gonophore stalk and peduncle of the raceme and migrate in the endoderm to the apex of the young gonophore buds where they proliferate in the endoderm to form the entocodon and germ cell mass. Eggs are produced sequentially in the gonophores of both species. Each egg is the result of fusion of 1200-2000 cells interconnected by cytoplasmic bridges into a large complex. At the center of each complex are 15-18 morphologically similar cells, all of which begin meiosis and form germinal vesicles and are considered oocytes. Toward the end of vitellogenesis, the oocytes fuse to form a single large oocyte with only one germinal vesicle surviving and completing meiosis. The remaining cells of each complex enlarge and produce yolk, but do not develop a germinal vesicle. These cells are true nurse cells, connected by cytoplasmic bridges to the oocytes. The nurse cells are engulfed by the oocytes and transformed into shrunken cells as their cytoplasm is incorporated by the oocytes. The organization of the oocyte-nurse cell complexes is variable. As many as three oocytes in each complex have four cytoplasmic bridges. Cells with four bridges may be the result of cellular fusion as both oocytes and nurse cells in many complexes contain two nuclei. An oocyte-nurse cell complex is produced from one initial oogonium by mitotic proliferation and the subsequent complexes in each gonophore are probably from other oogonia. Older polyps of T. larynx are frequently hermaphroditic while T. indivisa is strictly dioecious.

Keywords
Biology, Zoology

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OOGENESIS IN TUBULARIA LARYNX AND
TUBULARIA INDIVISA (HYDROZOA, ATHECATA)

BY

Barry W. Spracklin
B.A., Northeastern University, 1969
M.S., Northeastern University, 1974

DISSERTATION

Submitted to the University of New Hampshire
in Partial Fulfillment of
the Requirements for the Degree of

Doctor of Philosophy
in
Zoology

September, 1984
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Date 7/19/84
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# TABLE OF CONTENTS

**ACKNOWLEDGEMENTS**................................................................. iii

**LIST OF FIGURES**........................................................................... v

**ABSTRACT**.......................................................................................... vi

**SECTION**

<table>
<thead>
<tr>
<th>SECTION</th>
<th>PAGE</th>
</tr>
</thead>
<tbody>
<tr>
<td>INTRODUCTION</td>
<td>1</td>
</tr>
<tr>
<td>Oogenesis</td>
<td>1</td>
</tr>
<tr>
<td>Hydroids</td>
<td>4</td>
</tr>
<tr>
<td>Gonophores and Germ Cells</td>
<td>7</td>
</tr>
<tr>
<td>Microscopy and Autoradiography</td>
<td>15</td>
</tr>
<tr>
<td>MATERIALS AND METHODS</td>
<td>21</td>
</tr>
<tr>
<td>Light Microscopy</td>
<td>21</td>
</tr>
<tr>
<td>Autoradiography</td>
<td>22</td>
</tr>
<tr>
<td>Electron Microscopy</td>
<td>24</td>
</tr>
<tr>
<td>RESULTS</td>
<td>26</td>
</tr>
<tr>
<td>Light Microcopy</td>
<td>26</td>
</tr>
<tr>
<td>Autoradiography</td>
<td>32</td>
</tr>
<tr>
<td>Electron Microscopy</td>
<td>76</td>
</tr>
<tr>
<td>DISCUSSION</td>
<td>112</td>
</tr>
<tr>
<td>LIST OF REFERENCES</td>
<td>136</td>
</tr>
</tbody>
</table>
# LIST OF ILLUSTRATIONS

<table>
<thead>
<tr>
<th>Illustration</th>
<th>Description</th>
<th>Page</th>
</tr>
</thead>
<tbody>
<tr>
<td>1.</td>
<td>T. larvnx. Underwater photograph (20 feet)</td>
<td>34</td>
</tr>
<tr>
<td>2.</td>
<td>Clava leptostyla. Light microscopy photograph of gonophore</td>
<td>36</td>
</tr>
<tr>
<td>3.</td>
<td>T. larvnx. Light microscopy photograph of small germ cell mass</td>
<td>38</td>
</tr>
<tr>
<td>4.</td>
<td>T. larvnx. Light microscopy photograph of large germ cell mass</td>
<td>40</td>
</tr>
<tr>
<td>5.</td>
<td>T. larvnx. Light microscopy photograph of early stages of oogenesis</td>
<td>42</td>
</tr>
<tr>
<td>6.</td>
<td>T. indivisa. Light microscopy photograph of developing oocytes</td>
<td>44</td>
</tr>
<tr>
<td>7.</td>
<td>T. larvnx. Light microscopy photograph of intercellular bridges</td>
<td>46</td>
</tr>
<tr>
<td>8.</td>
<td>T. indivisa. Light microscopy photograph of intercellular bridges</td>
<td>48</td>
</tr>
<tr>
<td>9.</td>
<td>T. indivisa. Light microscopy photograph of oocyte with two nuclei</td>
<td>50</td>
</tr>
<tr>
<td>10.</td>
<td>T. indivisa. Light microscopy photograph of oocytes</td>
<td>52</td>
</tr>
<tr>
<td>11.</td>
<td>T. larvnx. Light microscopy photograph of an actinula</td>
<td>54</td>
</tr>
<tr>
<td>12.</td>
<td>T. indivisa. Light microscopy photograph of oocyte-nurse cell complex</td>
<td>56</td>
</tr>
<tr>
<td>13.</td>
<td>T. larvnx. Light microscopy photograph of mature gonophore</td>
<td>58</td>
</tr>
<tr>
<td>14.</td>
<td>T. indivisa. Light microscopy photograph of oocyte nucleus</td>
<td>60</td>
</tr>
<tr>
<td>15.</td>
<td>T. larvnx. Light microscopy photograph of spent gonophore</td>
<td>62</td>
</tr>
<tr>
<td>16.</td>
<td>T. larvnx. Light microscopy photograph of hermaphroditic polyp</td>
<td>64</td>
</tr>
<tr>
<td>17.</td>
<td>T. indivisa. Line drawing of oocyte-nurse cell complex</td>
<td>66</td>
</tr>
<tr>
<td>18.</td>
<td>T. indivisa. Line drawing of oocyte-nurse cell complex</td>
<td>68</td>
</tr>
<tr>
<td>19.</td>
<td>T. indivisa. Line drawing of oocyte-nurse cell complex</td>
<td>70</td>
</tr>
</tbody>
</table>
20. *T. indivisa*. Line drawing of oocyte-nurse cell complex...... 72  
22. *T. larynx*. Autoradiograph........................................................ 78  
23. *T. indivisa*. Autoradiograph...................................................... 80  
24. *T. indivisa*. Autoradiograph...................................................... 82  
25. *T. larynx*. Transmission electron micrograph of spadix............ 89  
27. *T. larynx*. Transmission electron micrograph of nutrient vesicles......................................................... 93  
28. *T. larynx*. Transmission electron micrograph of four intercellular bridges................................................. 95  
29. *T. larynx*. Transmission electron micrograph of intercellular bridges............................................................. 97  
31. *T. indivisa*. Plastic section of vitellogenic germ cells............. 101  
32. *T. larynx*. Transmission electron micrograph of vitellogenic germ cells......................................................... 103  
33. *T. larynx*. Transmission electron micrograph of shrunken cells................................................................. 105  
34. *T. larynx*. Transmission electron micrograph of spadix............ 107  
35. *T. larynx*. Transmission electron micrograph of vitellogenic germ cells......................................................... 109  
36. *T. larynx*. Transmission electron micrograph of yolk............. 111
ABSTRACT

OOGENESIS IN TUBULARIA LARYNX AND TUBULARIA INDIVISA (HYDROZOA, ATHECATA)

by

BARRY W. SPRACKLIN

University of New Hampshire, September, 1984

In Tubularia larynx and T. indivisa, interstitial cells proliferate in the ectoderm of the gonophore stalk and peduncle of the raceme and migrate in the endoderm to the apex of the young gonophore buds where they proliferate in the endoderm to form the entocodon and germ cell mass. Eggs are produced sequentially in the gonophores of both species. Each egg is the result of fusion of 1200-2000 cells interconnected by cytoplasmic bridges into a large complex. At the center of each complex are 15-18 morphologically similar cells, all of which begin meiosis and form germinal vesicles and are considered oocytes. Toward the end of vitellogenesis, the oocytes fuse to form a single large oocyte with only one germinal vesicle surviving and completing meiosis. The remaining cells of each complex enlarge and produce yolk, but do not develop a germinal vesicle. These cells are true nurse cells, connected by cytoplasmic bridges to the oocytes. The nurse cells are engulfed by the oocytes and transformed into shrunken cells as their cytoplasm is incorporated by the oocytes. The organization of the oocyte-nurse cell complexes is variable. As many as three oocytes in each complex have four cytoplasmic bridges. Cells
with four bridges may be the result of cellular fusion as both oocytes and nurse cells in many complexes contain two nuclei. An oocyte-nurse cell complex is produced from one initial oogonium by mitotic proliferation and the subsequent complexes in each gonophore are probably from other oogonia. Older polyps of _I. larynx_ are frequently hermaphroditic while _I. indivisa_ is strictly dioecious.
INTRODUCTION

Oogenesis

Oogenesis in long lived animals such as vertebrates is a complex process, synchronized by hormones, and involving the cooperative interaction of many organ systems. However, even with all of the complexity and capabilities of the vertebrate body, oogenesis can be a rather slow process. In certain frogs each oocyte requires three consecutive years to complete oogenesis. Human oocytes are arrested in diplotene of the first meiotic division in the fourth month of fetal life (Baker, 1963), while the first egg is not normally ready for fertilization for 12 to 14 more years. Postlethwait and Shirik (1981) observed that "people invest nearly two decades in providing sustenance and instruction to their offspring, but insects often spend only a few days in equipping their eggs with nutrient reserves and developmental instructions sufficient to sustain life and program development." In addition to insects, species in the class Hydrozoa, most of which have a brief seasonal existence and a relatively simple level of tissue organization, also manage to accomplish the entire oogenetic process in a relatively short period. *Tubularia larvax* polyps can release actinulae within 24 days of larval settlement (Pyefinch and Downing, 1949) and *T. crocea* releases actinulae only six to eight days after gonophores first appear (Mackie, 1966). Gametogenesis in *Hydra carnea* is completed in only four days (Honegger, 1981). To accomplish this rapid development hydroids employ a number of modes of germ cell formation, oocyte nutrition, and larval development. Considering the
varied developmental strategies, range of body forms, photoperiod control, local abundance, and ease of collection of many species, hydrozoans are, in many ways, excellent subjects for gametogenetic and embryogenetic studies.

The result of oogenesis is a highly specialized cell, the egg, which is developmentally totipotent and has the capacity to produce a multicellular organism formed of varied cellular types. However, with all of its latent capacity and raw materials, the haploid egg is normally incapable of further development and is merely a transitional stage between two generations. Except in parthenogenetic species, sperm penetration and pronuclear fusion are required before the egg can begin mitosis and express its developmental potential during subsequent development. Since sperm contribute only chromosomes and often a pair of centrioles, the egg is essentially a self-contained unit. Sufficient yolk, RNA, and other materials must be produced and stored during oogenesis to last until embryogenesis proceeds to the point at which endogenous production commences and a functional feeding stage can utilize an exogenous supply of food. In most species, the process of yolk production and ribosomal synthesis requires a significant expenditure of energy and nutrients by the parent organism.

Vitellogenesis, the production and storage of yolk, is thus one of the major parts of oogenesis, except in mammals and other organisms where placental nutrition sustains development from essentially yolkless eggs.

Vitellogenesis occurs in different species of animals by one of three methods. These include, autcsynthesis, in which the oocyte produces its own yolk from basic nutrients processed by the somatic cells of the organism; heterosynthesis, in which the yolk materials are
synthesized by somatic cells or other germ-line cells and then transported to the oocyte; and by a combination of the two methods referred to as heterosynthesis (Schechtman, 1955; Anderson, 1974). There is a large amount of variation in each of these three basic methods in the type and amount of yolk produced by different animals. Anderson (1974, pg. 39) observed "that nature has richly elaborated on, and in diverse ways, the mechanisms of vitellogenesis," and that there has not appeared "some common pattern that would make it an easy task to formulate a classification of vitellogenesis from an evolutionary point of view." There are examples of autosynthesis from a number of phyla including the Cnidaria, and according to Hisaw (1963) this was probably the primitive situation. However, early in the evolutionary development of many animals, the vitelлогenic requirements exceeded the capacity of a single oocyte during a breeding season. Berrill and Karp (1975, pg. 100) report that "in most animals - though many exceptions exist - the development of the oocyte takes place with the aid of another type of cell, an accessory cell."

A large number of different terms are used to refer to the various types of cells associated with oocytes during vitellogenesis; all of these are included under the term, accessory cell. This term embraces both accessory cells of somatic origin, called follicle cells, and accessory cells of germ-line origin, called nurse cells (Berrill and Karp, 1976; Browder, 1980). Following Anderson (1974) I will use the term nurse cell to include only those cells derived along with the successful oocyte from a single oogonium. All such cells are interconnected by cytoplasmic bridges that result from incomplete cytokinesis. In hydroids, germ-line accessory cells that lack
cytoplasmic bridges are termed nutritive oocytes. Examples of nutritive germ-line cells that are active during oogenesis are known from many invertebrate phyla and nurse cells, in particular, appear to have evolved independently many times. The "nurse cells" reported in sponges by Fell (1969, 1974) are not connected by cytoplasmic bridges to the oocyte and are, in different species, amoeocytes, choanocytes, or cells derived from choanocytes. With the discovery of nurse cells in Tubularia during the course of this investigation, the class Hydrozoa of the phylum Cnidaria is now the evolutionarily most primitive grade in which nurse cells have been seen and may be significant to our understanding of the evolution and function of nurse cells.

Hydroids

Within the class Hydrozoa there is a wide range of variation in the type of life cycle, the size, and the life span exhibited by different species. Some species are strictly polypoid, others strictly medusoid, and some species have both polypoid and medusoid phases in their life history. Included in the class are the interstitial actinulids, chondrophores, trachylines, and the polymorphic polypoid-medusoid siphonophores. Among the polyps are included the small hydras and the seven foot tall Brachiocerianthus imperator; the medusae include the small, short lived medusae of Millepora which survive for only a few hours after release (Hyman, 1940), and the medusae of Aequorea aquorea which may live six or seven months in the plankton and grow to 20 cm in diameter (Russell, 1953).

While hydrozoans are structurally simple, with only a tissue level of organization and two cellular layers, a wide range of diversity is seen in the development of hydrozoans and the involvement of accessory
cells supporting the growth of the oocyte during vitellogenesis. Mergner (1971) lists five different cleavage types and six methods of germ layer formation. In many hydroids, and in cnidarians in general, accessory cells are not involved during gametogenesis (Campbell, 1974) and vitellogenesis is entirely autosynthetic. At the other extreme are the species in which thousands of germ-line accessory cells support the production of a few or even a single successful or definitive oocyte.

Species in the orders Anthomedusae (Aethecata) and Leptomedusae (Thecata) are commonly referred to as hydroids (athecate and thecate) and are the most generally representative groups in the class Hydrozoa which also contains the siphonophores, trachylines, and the interstitial actinulids. Many species of athecate and thecate hydroids exhibit the classic alternation between polypoid and medusoid phases in their life history. While the thecate hydroids are all colonial and have only a planula larva, the athecte hydroids are much more variable. Both solitary and colonial species are present in the Athecata and while most species have a planula larva typical of the phylum Cnidaria, many species develop an actinula larva. There are even species in the athecate family Margelopsidae with pelagic polyps that have no attached phase in their life history.

Most studies of hydrozoan germ cell development and embryogenesis have been accomplished using species with attached gonophores. This is partly because they are usually easier to collect in a reproductive condition in the field or easier to maintain in the laboratory. However, the majority of species of hydroids have no free swimming medusoid phase with feeding tentacles, a functional mouth, and sensory organs, but instead develop attached reproductive structures called
gonophores which do not feed and have no sensory structures. The
gonophores have been considered by most investigators to represent
stages in the reduction of medusae that no longer detach (Kuhn, 1910,
1913; England, 1926; Hyman, 1940). There are, however, no basic
developmental differences between them and the species which release
free medusae.

**Tubularia**, a genus of athecate hydroids that lacks a free medusa in
its life history, is a member of the superfamily Tubularoidea (Rees,
1957) which includes **Hybocodon**, **Acaulia**, **Corymorpha**, **Euphysa**,
**Margelopsis**, **Branchiocerianthus**, and **Pelagohydra**. **Tubularia** is the best
known genus in the superfamily and *T. indivisa* Linneus, 1758 is the type
species of the family Tubulariidae Allman, 1864. **Tubularia** is one of
the most thoroughly investigated genera of marine hydroids and the fresh
water hydras may be the only hydroids to have received more attention.
Abundance and ease of collecton are among the factors that have led many
investigators to study different species of **Tubularia** and the polyps
have been described by Agassiz (1862), Allman (1871), Kingsly (1910),
Fraser (1944), and Calder (1974). Studies on various aspects of the
biology of **Tubularia** have included autotomy of hydranths (Morse, 1909),
ecology (Fenchel, 1905; Pyefinch and Downing, 1949), behavioral
physiology (Josephson, 1965; Josephson and Mackie, 1965), growth and
culture (Mackie, 1966), and regeneration (Barth, 1940; Berrill, 1948;
Davidson and Berrill, 1948; Fulton, 1959; Tardent, 1963; Rose, 1974;
Tweedell, 1974).

Species of **Tubularia** (Fig. 1) form large, colorful colonies on
floats, buoys, and boat hulls along the New England coast and in many
parts of the world they are important fouling organisms highly
resistant to copper poisons (Barnes, 1948). Polyps of T. indivisa Linnaeus, 1758, one of the larger species, may reach a height of 12 inches (Hincks, 1868) and have 40 filiform tentacles in the proximal whorl and over 40 in the distal whorl (Fraser, 1944). T. larynx Ellis and Solander, 1786 is normally 2-3 inches in height with about 20 tentacles in each whorl (Fraser, 1944). Sexually mature polyps of Tubularia bear a ring of racemes of reproductive buds or gonophores between the two whorls of tentacles. Fenchel (1905) reported a normal maximum of about 16 racemes on each polyp while Allman (1871) reported over 30 racemes on some polyps of T. larynx. Most of the polyps collected for this research had 6-8 fully developed racemes containing mature gonophores and a variable number of immature racemes. Polyps of T. larynx usually possessed 100-300 gonophores while the polyps of T. indivisa had 400-800. The oldest and most mature gonophores are located at the distal end of a raceme which elongates as new gonophores form at the base of the stalk or peduncle of the raceme. All of the polyps of T. indivisa examined were strictly dioecious while older female gonophores of T. larynx were often protogynic, which was reported in T. crocea by Liu and Berrill (1948).

Gonophores and Germ Cells

The female gonophores (Figs. 3, 4) of both T. larynx and T. indivisa are eumedusoid (Rees, 1957) which means that they develop all of the cellular layers of a free medusa (Broch, 1915; Teissier, 1926), but fail to form tentacles, sensory organs, or a functional mouth. In Tubularia the germ cells segregate at a very early stage from the cells of the entocodon (Liu and Berrill, 1948) which forms the inner ectoderm in a eumedusoid gonophore or the subumbrellar and manubrial ectodermal
layers in a free medusa (Hyman, 1940). The spadix is a finger-like projection of endodermal cells into the gonophore and the lumen of the spadix is continuous with the enteron of the raceme and polyp. In mature gonophores the germ cells are tightly packed into a horseshoe-shaped mass (Fig. 4), the germ cell mass (Liu and Berrill, 1948), that surrounds the spadix. Septal cells are scattered within the germ cell mass and interconnect the manubrial layer of the inner ectoderm with the spadix (Liu and Berrill, 1948). When the first egg is produced within a gonophore it ruptures the manubrial layer of the inner ectoderm and simultaneously moves into and creates a space between the manubrial and subumbrellar layers (Boelsterli, 1975). This space, where fertilization and embryonic development to an actinula larva occur, is the subumbrellar cavity (Boelsterli, 1975) and is equivalent to the subumbrellar space in a free medusa (Fig. 13).

An unusual aspect of gametogenesis in the female gonophores of Tubularia is the sequential production and maturation of separate clusters or generations of oocytes (Figs. 12, 13, 15, 16). Before the first egg has been produced from the germ cell mass another cluster of oocytes begins to enlarge to form a second egg (Liu and Berrill, 1948). As the eggs mature they are fertilized and develop within the gonophore which results in the release of a series of actinulae from each gonophore. The total number of eggs and actinulae normally produced by each gonophore is not known.

Many of the characteristics of Tubularia are advantageous during studies of gonophore and germ cell development. Colonies of Tubularia can be raised in the laboratory (Mackie, 1966), the polyps grow and quickly begin to produce gonophores and germ cells (Pyefinch and
Downing, 1949; Mackie, 1966), each polyp has numerous gonophores at various stages of development, one or more species are abundant at many locations, and at all times at least some of the polyps of a colony are reproductive (Pyefinch and Downing, 1949). However, other features make Tubularia a difficult research animal for developmental studies. One of these features is that it is not an easy animal to maintain in the laboratory and has been used as a water quality indicator in closed seawater systems. West and Renshaw (1970) found that T. orceae would shed its hydranths before other hydrozoans were affected by poor water quality. This has meant that most developmental studies have relied on freshly collected material, not colonies that have been under long term observation. The large number of gonophores at different developmental stages makes it difficult to follow the progress of individual gonophores and the wall of the gonophore and the large mass of germ cells in each gonophore obscures the observations of germ cells in living animals. Serial sections of fixed material also have disadvantages including the problems associated with deciphering continuous, dynamic processes such as growth, engulfment, or fusion from preparations of different gonophores and different animals. The large number of germ cells in each female gonophore, consisting of groups at different developmental stages, also add to the confusion, not to mention the problem of hermaphroditic gonophores. As a result, even though many investigators have studied the reproductive biology of species of Tubularia, much confusion and controversy remains and interpretations, even within the same species, differ at least somewhat.

Most of the work on germ cell development and gonophore formation of Tubularia has been reviewed by Lowe (1926) and Liu and Berrill (1948)
while the literature concerning larval development was summarized by Nagao (1965). There is still no complete agreement on the origin of the entocodon, the site of the initial differentiation of the germ cells, nor the cellular type from which the germ cells of Tubularia develop. This is in addition to the confusion about various aspects of the formation of a mass of about 2000 cells in each female gonophore and their subsequent fusion into a single oocyte.

Since the work by Agassiz (1860) most investigators, including Weismann (1880), Hargitt (1904), Allen (1900), and Hargitt (1909), have considered that the entocodon forms in the ectoderm of the gonophores of Tubularia by the proliferation of local ectodermal cells. They also considered the germ cells to develop from the inner layer of the entocodon without cellular migration. Benoit (1925) and Liu and Berrill (1948) also found no evidence of cellular migration, but reported the entocodon to form by the proliferation of endodermal cells at the apex of the evaginating gonophore. According to Benoit, the entire entocodon formed by the proliferation of a single interstitial cell (I-cell). There have also been advocates for a migration of the germ cells to their final position in the gonophore. Brauer (1891) and Lowe (1926) concluded that the cells of the entocodon, including the germ cells, originate from I-cells that have migrated from the ectoderm of the stalk of the gonophore. Dupont (1942) considered the germ cells and entocodon to have independent origins and according to Campbell (1974, pg. 145), the germ cells "migrate into or with the entocodon" and (pg. 148) "in well-developed medusae, gametogenesis does not occur for some time after medusa liberation." Even allowing for the probability that the development of germ cells and gonophores is not
identical in different species of *Tubularia*, the variation in the opinions of different investigators is, at the least, distracting. Liu and Berrill wrote (1948, pg. 39), "Inasmuch as the conclusions in the present paper are in marked disagreement with the more important conclusions both of Dupont and of Benoit ('25), whom Dupont also criticizes, it is optimistic to hope that this will be the last." Boelsterli (1975), the latest investigator of the development of germ cells in *Tubularia*, avoided the problem by starting with oogonia already present in the gonophore and ignoring the problem of earlier developmental stages.

While reviewing the literature on hydrozoan oogenesis it is apparent that hydroids (athecate and thecate) may be separated into two groups based on the proportion of female germ-line cells that complete development or the number of accessory cells supporting the growth of each oocyte. In the first group of hydroids there is no real development of accessory cells and most oocytes complete development, while in the second group a tremendous overproduction of germ cells occurs and these are utilized as accessory cells. Somewhat similar conclusions were obtained by Van de Vyver (1967, 1968a,b) who separated the thecate hydroids into three groups, the Filifera, corynoid Capitata, and tubularid Capitata. These groups differed in the type of larva, amount of yolk in the eggs, and the number of excess germ cells.

The first of my two groups of hydrozoans includes the majority of hydrozoan species, none of which are strictly solitary, and includes the Filifera and some of the corynoid Capitata of Van de Vyver. In this group, all, or at least a large proportion, of the female germ-line cells that differentiate, complete oogenesis and there is little or
only moderate overproduction of oocytes. According to Berrill (1961) all of the oogonia that differentiate in the gonophores of *Hydractinia echinata* and *Aselomaris mitchilli* develop into oocytes without proliferation. All of these oocytes complete oogenesis and no accessory cells are present. Vitellogenesis in these species is probably autosynthetic as the nutrients resulting from extracellular digestion of exogenous food resources are carried along the continuous enteron to the gonophores and passed across the mesoglea and the endodermal cells of the spadix to the oocytes. Josephson and Mackie (1965) and Lenhoff (1968) described the pressure changes that move food particles to the gonophores in the polyps of *Tubularia*. In species that do not release free medusae, the nutritive spadix is resorbed at the end of oogenesis while the embryo is brooded.

Species in which there is a moderate overproduction of oocytes are also included in the first group of hydroids. According to Campbell (1974), while only a few eggs develop in each gonophore, about six or seven oocytes, all of which are capable of completing development, are produced for each egg that actually matures. He reported that (pg. 161), "Most hydrozoan gonads begin with more oocytes than will eventually ripen, and frequently only a single ovum develops in each gonad. The other oocytes are expended in the process of feeding the successful oocyte." Some of these oocytes remain where they originally differentiate and later degenerate, while the rest migrate into the gonophores. In most of these hydroids there is little growth of any but the successful or definitive oocyte which Van de Vyver (1968a) calls "l'oocyte privilegie." The excess oocytes may serve as a partial source of food for the definitive oocytes (Mergner, 1957). However, except for
the degeneration and possible resorption of these relatively few excess oocytes, vitellogenesis is still basically autosynthetic. The eggs of the species, including *Clava leptostyla* (Fig. 2), which produce few excess oocytes are relatively small. The largest eggs reported in these species were produced by *Hydractinia echinata* and were 168 um in diameter.

A more complex and interesting system of oogenesis is found in the second group of hydroids which includes the tubularid Capitata and most of the corynid Capitata of Van de Vyver. In this group there is a tremendous overproduction of germ cells. The large, solitary athecate hydroids and some colonial species produce large, yolky eggs and many brood the resulting embryo to an advanced larval stage, the actinula (Fig. 11). In gonophores of these hydroids, which include *Tubularia*, *Hydra*, and *Pennaria*, hundreds to thousands of excess or nutritive oocytes are produced and utilized as food in the development of relatively few definitive oocytes. In the species that produce an actinula larva, usually only one is present in each gonophore at a time, although a sequence of actinulae may be brooded and released. Benoit (1925), working with *T. mesembryanthemum*, and Boelsterli (1975), working with *T. crocea*, have reported two or three eggs or early embryos in a single gonophore, but there are no reports of two or more actinulae simultaneously present in a single gonophore. The nutritive oocytes may be directly engulfed or may disintegrate; in any case, the resulting nutrients are absorbed by the definitive oocytes. Before being engulfed, the nutritive oocytes in *Tubularia* species, *Hydra* (Zihler, 1972), and *Pennaria* (Cowden, 1964) produce yolk and ribosomes that will be incorporated by the definitive oocytes.
There is a large amount of variation in the size of the oocytes and germinal vesicles of different hydroids and two earlier investigators, Jorgensen (1913) and Hargitt (1919), attempted to correlate the different methods of oocyte nutrition with relative nuclear size. Jorgensen proposed that oocytes that absorb their food directly from the surrounding fluid have relatively large nuclei; oocytes which are fed by special nurse cells or follicle cells or engulf other cells have relatively small nuclei. He believed that the nuclei of the accessory cells governed the growth of the oocyte and that the nuclei of the oocytes remained relatively inactive until the end of the prophase of meiosis one. An extreme example of this type of oocyte nutrition was reported by Pianka (1974) in the ctenophore, Bolinopsis microoptera, in which the oocyte nucleus does not enlarge at all during vitellogenesis.

Hargitt (1919) observed that in hydrozoans there are two different methods of oocyte nutrition which seem to overlap in certain species. In some hydroids, food is absorbed directly by oocytes from the enteron or from the endodermal cells of the spadix, while in others the oocytes absorb or engulf other cells, usually extra or nutritive oocytes. In an attempt to quantify Jorgensen's proposal, Hargitt made measurements of the oocytes of 14 hydrozoans and obtained results consistent with Jorgensen's predictions. In species that are nourished directly from the enteron, the ratio of oocyte volume to nuclear volume was found to range from 16:1 to 73:1. In species with oocytes that engulf other cells or have some form of nutritive oocytes, the ratio ranged from about 284:1 for Corymorpha pendula to about 6000:1 for T. crocea. In all of these species, the germinal vesicle of the definitive oocyte enlarges during oocyte growth. This connection between oocyte-nuclear volume and
the mode of oocyte nutrition is presumably related to the extent of participation of the oocyte nucleus in vitellogenesis.

While the entire process of oogenesis is rapid in Tubularia, Hydra, and other hydroids with nutritive oocytes, in many animals of different phyla, the meiotic prophase has a long duration. The insects which possess nurse cells (meristic oogenesis) also have a short meiotic prophase. Davidson (1968) hypothesized that there must be some rate limited synthetic activity that can be accomplished in a few days by the concerted efforts of polyploid nurse cells of insects, but which normally takes a solitary oocyte considerably longer. According to Davidson (pg. 183), "the molecules in question cannot be yolk, since yolk and other such products are typically accumulated in the course of a relatively brief period of the diplotene stage." Work on Xenopus by Perkowaka et al. (1968) revealed that the limiting factor in the rate of oogenesis is not the production of yolk, but the production of ribosomes. The nuclei of Xenopus laevis oocytes contain 1400 to 1600 nucleoli and about 5200 nucleolar DNA cores or rings while the normal number of nucleoli of somatic cells in Xenopus is two. Calculations indicate that without amplification of the ribosomal cistrons and nucleoli, about $1.7 \times 10^5$ days or about 466 years would be required for the oocytes to synthesize an equivalent number of ribosomes on their own. While there are no reports of gene amplification in the Cnidaria, the same result is achieved in the second group of hydroids in which many germ-line cells, each with genes for ribosome RNA, produce materials that are eventually incorporated by the definitive oocytes.

**Microscopy and Autoradiography**

A large number of light microscopical observations have been made
on hydrozoan oogenesis, including investigations of species that produce large numbers of excess germ cells. Van de Vyver (1968) found that oogenesis in *Arum cocksi* involves the growth of one "cellule privilegie" by the engulfment of the other oocytes in the gonophore. While this is the situation in many hydroids with nutritive oocytes, it is readily apparent that there is much variation in the oogenetic process in these hydrozoans. In addition to the absorption of the nutritive oocytes, Hargitt (1917) reported that there were indications of fusion of the oocytes in *Hybocodon prolifer*. Mangan (1909) suggested that in *Millepora* the cell membranes between the definitive oocyte and the surrounding nutritive oocytes dissolve and a plasmodium forms. In *Eudendrium hargitti*, a species with a large oocyte-nuclear volume ratio, but without nutritive oocytes, Congdon (1906) reported engulfment of somatic ectodermal cells by the oocytes. True nurse cells, with direct cytoplasmic connections to the definitive oocytes, are also present in hydroids and have been found in *Tubularia larynx* and *T. indivisa* by Spracklin (1978, 1980). It is entirely possible that the excess germ cells of other hydroids are also nurse cells, but where cytoplasmic bridges do not exist, or have not yet been found, these cells are appropriately called nutritive or abortive oocytes.

More extensive investigations of oogenesis involving hydroids with nutritive oocytes have been done on species in a number of genera including *Pennaria*, *Hydra*, and *Tubularia*. Oogenesis in *Pennaria tiarella* has been investigated by Smallwood (1899), Hargitt (1900), Berrill (1961), and Cowden (1964). In this species the primordial germ cells proliferate to produce several thousand oocytes in each gonophore. Approximately 10 oocytes in each gonophore grow at an accelerated rate
and consume the remaining nutritive oocytes. The nuclei of the consumed cells persist until the larval ectoderm develops.

Even though it is sometimes difficult to induce sexuality in *Hydra* held in the laboratory, the ease of maintaining this fresh water hydroid and its availability have led many investigators to use it extensively for investigations of oogenesis including Brauer (1891), Brien (1950a,b, 1961, 1963), Brien and Reniers-Decoen (1949, 1951), Downing (1909), Korotneff (1878), McConnell (1933), Tannreuther (1908, 1909), and Wagner (1909). Zihler (1972), who was the first to report ultrastructural observations of oogenesis in this genus, divided oogenesis in *Hydra attenuata* and *H. circumcinta* into five stages. In stage I, groups of oogonia develop from interstitial cells and proliferate in the ectoderm of the polyp. In stage II, after the oogonia have grown somewhat, one oogonia in each group develops into an oocyte. During stage III, the oocyte in each group engulfs the remaining oogonia. The remains of the engulfed oogonia are very distinctive structures called "pseudozellen" or "shrunken cells". In stage IV, the large oocytes from the original groups fuse to form a single large oocyte which matures to form the egg in stage V.

Since Agassiz (1860), oogenesis and gonophore development in the genus *Tubularia* have been investigated by numerous researchers including, most recently, Liu and Berrill (1948), Nagao (1965), Van de Vyver (1968), and Boelsterli (1965). Liu and Berrill reported that a germ cell mass of several thousand cells was formed by a rapid proliferation of the primordial germ cells and Boelsterli found that, as in *Hydra*, a small number of oocytes begin to grow and subsequently engulf the surrounding nutritive oocytes. In each gonophore, the
growing, or definitive, oocytes fuse to form a single large oocyte that completes meiosis.

The remains of the engulfed nutritive oocytes, the pseudocells, persisted in the oocytes of both Tubularia (Figs. 10, 11, 33) and Hydra. Wager (1909) realized that at least some of the pseudocells were the shrunken remains of whole cells and Zihler (1971) introduced the more descriptive term, shrunken cells. Boelsterli (1975) was the first to demonstrate that the pseudocells in Tubularia oocytes were not degenerating nuclei, but the shrunken remains of the engulfed nutritive oocytes, complete with a small amount of cytoplasm, membranes, yolk, and degenerating mitochondria. According to Nagao (1965), most of the shrunken cells persist for several days after fertilization in Tubularia and a few can still be found at the basal part of the hypostome in the settled actinula six days after fertilization.

While light microscopy has been used extensively in the investigation of germ cell development in the Hydrozoa, only a small number of papers have appeared on the ultrastructure of hydrozoan oogenesis. Very little additional work has been published since Boelsterli (1977, pg. 267) wrote, "Whereas light microscopy studies of oogenesis in hydroids and hydromedusae are quite numerous, relatively little information is available on the ultrastructural aspects of gametogenesis." The only additions I have to add to Boelsterli's list of ultrastructural papers are Van de Vyver (1964, 1967, 1968), Tucker and Wyttenback (1974), and Honegger (1981). Van de Vyver's papers were largely concerned with light microscopy and TEM of the embryological development of five athecate hydroids, but did include some information on gametogenesis. Tucker and Wyttenback included some information on
the primordial egg cells, and Honegger's work involved scanning electron microscopy (SEM) of sperm and egg release and the embryotheca, and the transmission electron microscopy (TEM) of fertilization in *Hydra carnea*. This small collection of ultrastructural papers, some of which are extremely limited in scope and contain minimal ultrastructural information, has produced only a superficial picture of hydrozoan oogenesis at the ultrastructural level. In addition, these papers consider only a few of the many genera and species with nutritive oocytes. Van de Vyver (1968a) published a single micrograph of *T. ceratogyne*; Boelsterli (1975) worked with *T. crocea*; Zihler (1972) investigated *Hydra attenuata* and *H. circumcineta*; and Honegger (1981) observed *H. carnea*.

While the ultrastructural information on hydrozoan oogenesis in the literature is sparse, autoradiographic information is non-existent. Autoradiographic techniques, which have proven valuable research tools in the study of oogenesis in other organisms, have not been used to study oogenesis in hydroids. In other animals autoradiography has been used, with great success, to follow events such as the amplification of nucleolar RNA in *Xenopus laevis* oocytes (Perkouska, Macgregor and Bernstiel, 1968) and to identify the sites of synthesis of the various yolk proteins packaged in the eggs of *Drosophila melanogaster* (Gutzeit, 1980). Isotopically labeled compounds have been especially useful in understanding the function of the nurse cells in the meroistic ovaries of insects. Bier (1963, 1965) produced striking demonstrations of the production of proteins and RNA by the nurse cells of *Drosophila* and subsequent transfer of these products to the oocytes. Autoradiographic studies with hydroids has been concerned with the incorporation of
labeled compounds (Burnett et al., 1962), hydrozoan development (Haynes, Summers, and Kessler, 1974), DNA synthesis and cell movement during regeneration (Tweedell, 1974), cellular proliferation (Campbell, 1965), the synthesis of nucleic acids and proteins (Clarkson, 1969), and the symbiotic relationship between algal and hydroid (Muscatine and Lenhoff, 1963; Sznant, 1971; Eisenstadt, 1971).

In the present study I examine the organization and ultrastructure of the oocyte-nurse cell complexes in T. larynx and T. indivisa, propose a method of formation of the complexes, and explore hydrozoan oocyte nutrition.
MATERIALS AND METHODS

Light Microscopy

Colonies of Tubularia larynx and T. indivisa were collected from floats located at the mouth of the Piscataqua River in Portsmouth, New Hampshire; from tide pools at East Point, Nahant, Massachusetts, and Eastport, Maine; and from the subtidal at the Isle of Shoals, New Hampshire. For routine light microscopical histology most specimens were fixed in Bouin's fixative, although Helly's, Zenker's, Orth's, Suse's, AFA, Sanfelice's, and 10% formalin-sea water fixatives were also used. During the water rinse racemes were removed from the fixed polyps with microdissecting scissors and separated into individual gonophores or small clusters of gonophores. To observe very young gonophores, small polyps were fixed in Bouin's fixative, the tentacles removed, and the remainder of the polyp cut into two or four pieces. The gonophores and pieces of polyps were dehydrated in a graded series of ethanol (two 10 minute rinses each of 70%, 85%, 95% and 100%) and, after an overnight infiltration at 37°C or 42°C, were embedded in polyethylene glycol 400 or 600 distearate mixed 10:1 with cetyl alcohol. Specimens were sectioned at 2, 4, 6, 8, 10, 12, 15, or 20 μm. Slides were stained with Mallory-Heidenhain Rapid Stain (Humanson, 1972), Heidenhain Iron Hematoxylin, Thionin-Methyl Green, or Azan, and mounted with Permount containing 10% terpineol to reduce crazing. Gonophores fixed in Zenker's fixative were sectioned and treated with the Feulgen and Periodic Acid-Schiff (PAS) reactions. Whole mounts were made of gonophores or racemes from polyps fixed in Bouin's or Zenker's
fixatives and were stained with Lynch’s precipitated borax carmine or Proecher’s coelestine blue B-lake (Riser, 1950). Approximately 16,000 gonophores of *T. larynx* and 4,000 gonophores of *T. indivisa* were serially sectioned and approximately 500 whole mounts were made of gonophores of each species.

Various techniques were tried to produce shrinkage or swelling to permit better observation of the intercellular bridges and the oocyte-nurse cell complexes in the gonophores of both species by spreading out the cells of the complexes. Gonophores were fixed in Bouin’s fixative lacking acetic acid, in saturated aqueous picric acid, in 70% and 90% ethyl alcohol, and in 85% ethyl alcohol mixed 5:1 with a solution of saturated aqueous picric acid, and were sectioned on a rotary microtome at 10 μm. Squash preparations with fast green were made of untreated gonophores and gonophores that were opened and then partially digested with trypsin or papain. Whole gonophores and those broken open with dissecting needles were also subjected to the maceration techniques of David (1972) and Schmid et al. (1981).

To provide specimens to compare with *Tubularia*, a small number of reproductive, female polyps of *Acaulis primaria*, *Rhizogeton fusiformis*, *Eudendrium ramosum* and *Clava leptostyla* were collected and fixed in Bouin’s fixative. The gonophores were embedded in polyethylene glycol 400 distearate, sectioned at 10 μm and stained with Mallory-Heidenhain Rapid stain.

**Autoradiography**

The volumetric capacity of the polyps was determined by injecting a solution of methylene blue sea water (0.1%) directly through the mouth of the polyp into the enteron with a number 33 needle (0.008 inch
and a 100 µl syringe. Polyps of *T. larvnx* could retain about 15 µl of stain without significant leakage while the larger polyps of *T. indivisa* could retain about 100 µl. Ten hydranths of each species were removed, blotted partially dry, and weighed to allow calculation of the amount of isotope required for a dosage of 2 µCi/gm wet weight. This amount was then doubled to allow for label lost to the stems and stolons. The variable length of the stems and the encrusting material on the stolons made it impossible to accurately weigh these parts of the colonies.

For each experiment using autoradiography, large colonies of *T. larvnx* and *T. indivisa* were collected from Portsmouth Harbor. Portions of female colonies, each consisting of about 20 polyps, were kept in finger bowls. The solutions of ^3H^-labeled uridine (5,6^-3H) and thymidine (methyl^-3H) were diluted to 0.0625 µCi/µl with sterile, Millipore filtered (0.22 µm) sea water. Under a dissecting microscope, ^3H^-labeled uridine or thymidine was injected directly into the enteron through the mouth of each polyp. Injections were initiated about one hour after collection of the colonies.

Polyps of *T. larvnx* received 4 µl (0.25 µCi) of solutions of uridine or thymidine while polyps of *T. indivisa* received 10 µl (0.625 uCi) of uridine. Uninjected control polyps from the original large colonies were placed in Bouin's fixative at the completion of the injections. Injected polyps were fixed after 2, 5, 12, 24, and 45 hours of incubation.

The first series of polyps was incubated on February 10, 1980 at 4°C. A second series of polyps, that did not include a 45 hour sample, was incubated on July 7, 1980 at 8°C (to match the warmer water
temperature at the collection site). Racemes were removed from the fixed polyps and cut into either small clusters of young gonophores or individual large gonophores. The gonophores were embedded in polyethylene glycol 400 distearate (mixed 10:1 with cetyl alcohol). Specimens were sectioned at 4 μm or 10 μm, mounted on glass slides, and were coated with Kodak NTB-3 Nuclear Track Emulsion. Resulting slides were exposed for 12 days for the first incubation and 30 days for the second incubation, developed for two minutes in Kodak D-19 at 19°C, and counterstained for two minutes with Mallory-Heidenhain Rapid stain.

**Electron Microscopy**

Small portions of colonies of *T. larynx* and *T. indivisa* were collected from floats in Portsmouth Harbor. Male and female polyps of both species were immediately placed in 2.5% phosphate buffered glutaraldehyde for two hours followed by post fixation in 2% osmium tetroxide for one hour (Cloney and Florey, 1968). Fixation was carried out at pH 7.4, 7.6, or 7.8 at room temperature (20-25°C) and at 4°C. During the first buffer wash racemes were removed and cut into small clusters of young gonophores or individual large gonophores. Specimens were embedded in Spurr's medium (Spurr, 1969; Ladd Research Industries, Inc.), Epon 812 (Ladd), and Epon 812-Araldite 506 (Mollenhauer, 1963; Polysciences). DMP-30 was the catalyst normally used with Epon 812, although DMP-10 was also used along with prolonged infiltration to improve the infiltration of yolk and shrunken cells in the large oocytes and embryos. Sectioning was carried out using glass knives on a Sorvall MT-1 or a Reichert OMI3 ultramicrotome. Some sections were mounted on formvar coated, 100 mesh, copper grids and stained 8 minutes with saturated aqueous uranyl acetate and 8 minutes with Reynold's lead
citrate (1963). Other sections were mounted on 200 and 300 mesh grids and stained 8 minutes with saturated, aqueous uranyl acetate and 10 minutes with vanadium molybdate (Callahan and Horner, 1964). All sections were examined with a Philips 200 transmission electron microscope. One micrometer thick plastic sections were stained with Richardson's stain (Richardson et al., 1960), toluidine blue (1% solution in 1% aqueous borax), or Azure II (1% aqueous). Sections were mounted in immersion oil and the cover slips rimmed with clear nail polish to yield permanent preparations.
RESULTS

Light Microscopy

Mallory-Heidenhain rapid stain and Bouin's fixative produced the best overall results and provided the most information for routine light microscopy, including observations of the initial germ cells, gonophore development, and the oocyte-nurse cell complexes. These techniques were especially useful in observations of very young gonophores. Mitotic figures were commonly found in the ectodermal I-cells in sections of both species from the base of racemes where the new gonophores form. Identical cells, some with mitotic figures, were found among the endodermal cells of the stalk and in a cluster at the apex of very young gonophore buds (Figs. 2,3). The smallest apical cluster of I-cells consisted of 17 cells, 2 of which had mitotic figures. Numerous other gonophores were found with 20-25 apical I-cells and 1 or 2 mitotic figures, but no oocytes. At this stage in gonophore development, there was no visible distinction between the distal cells of the apical cluster that would form the entocodon and inner ectoderm and the proximal cells that would develop into the germ cells. The cells of the entocodon initially form a blastula-like ball which flattens out to form the two layers of the inner ectoderm. The space or glockenohle within the entocodon is usually occluded when the entocodon flattens out. The opening reappears (Fig. 13) when the first egg forces the layers of the inner ectoderm apart creating the subumbrellar space. The smallest gonophore in which an oocyte was found contained only 26 cells in the apical mass.
In older, larger gonophore buds that contain about 200 apical cells, there are 60-75 oogonia with nuclei 7.5-8.0 μm in diameter, while the remaining apical cells, which are cells of the entocodon, have nuclei 6.0-7.5 μm in diameter. In a gonophore of this size as many as 5 oocytes (Fig. 5) have been counted. Oocytes at this time have nuclei 9.0-10.0 μm in diameter. There are also at least 2 and as many as 5 mitotic figures among the germ cells which are now organized into a relatively distinct structure, termed the germ cell mass.

The number of oocytes increases as the cells of the germ cell mass proliferate mitotically and gonophores enlarge (Figs. 6, 10). Different gonophores have been found to contain a maximum of 15-18 oocytes in the first germ cell mass. Eighteen oocytes were counted in a germ cell mass as small as 240 cells. Only the oocytes develop the chromatin network indicating the start of meiosis and develop distinct germinal vesicles. The peripheral germ cells or oogonia continue to proliferate mitotically and form a germ cell mass of 1200-2000 cells. The nuclei of the peripheral cells enlarge and appear similar to a small germinal vesicle. However, their nuclei show no signs of the chromatin organization seen in the oocytes as they begin prophase of meiosis I. These peripheral germ cells are nurse cells that can be shown with TEM to be interconnected by intercellular bridges (Figs 8, 28).

Before the first egg has been produced in a gonophore, a second group of 15-18 oocytes and their associated nurse cells begin to develop (Fig. 12) in the distal portion of the germ cell mass. The intercellular bridges in this second group or generation of germ cells are too small (0.55-0.70 μm) to be resolved with light microscopy, but
they have been observed with TEM. A single gonophore of *T. larynx* may contain up to four separate groups of germ cells or embryos at different developmental stages: proliferating oogonia, vitellogenic oocytes, early cleavage, actinula. The much larger gonophores of *T. indivisa* have been found to contain as many as five different developmental stages: proliferating oogonia, small vitellogenic oocytes, large vitellogenic oocytes, early cleavage, actinula. Numerous observations of whole mounts and living animals have revealed no gonophores of either species containing more than one actinula. Many gonophores of *T. indivisa* contain two eggs and one had three.

Nurse cell nuclei appear to have a dense layer of material (Figs. 6, 14) on the inside of the nuclear membrane when they are fixed and sectioned for light microscopy. This dense layer, which is not present in the oocytes (Figs. 9, 14) or proliferating oogonia, was thought to represent the early formation of heterochromatin which is abundant in the shrunken cells, but it did not stain with the Feulgen reaction. The dense layer was prominent in the nurse cells of both *T. larynx* and *T. indivisa* when Bouin's and Zenker's fixatives were used and was distinguishable with the other fixatives for light microscopy, except Orth's. No layer was seen on the inside of the nuclear membrane of the nurse cells that were fixed for TEM, either in the micrographs or in one micrometer thick plastic sections (Figs. 31, 32).

When the first group of oocytes complete vitellogenesis and commence to fuse into a single large oocyte, the centrally located nurse cells close to the oocytes have nuclei measuring about 15 μm in diameter. The smaller, peripheral nurse cells have nuclei 9–10 μm in diameter. Before suspension of mitosis in the second group of germ
cells in a gonophore, there is no condensed layer on the inside of the.
nuclear membranes of the peripheral nurse cells and their nuclei are
approximately 7.5 µm in diameter. These characteristics allow the
differentiation and counting of the cells of adjacent groups in the
germ cell mass of a gonophore. No special intervening layer or
cytoplasmic bridges were found between adjacent groups of germ cells at
different stages of development.

During growth of the first genetic cohort of germ cells within a
gonophore, the oocytes are usually situated adjacent to the spadix.
However, in the second and subsequent groups to develop in a gonophore,
oocytes are dispersed among their nurse cells and numerous nurse cells
are positioned between the oocytes and the spadix. As the oocytes of
each cohort of germ cells enlarge during vitellogenesis, they force
nurse cells away from and to the opposite side of the spadix. As a
result, it appears that there are fewer cells in the germ cell mass on
the side of the spadix where the oocytes develop because the oocytes are
larger than the nurse cells. The oocytes end up adjacent to the spadix
which is presumably the source of nutrients while the nurse cells are
moved away from direct contact with the spadix (Fig. 31).

Near completion of the vitellogenic phase of oogenesis several
processes occur more or less simultaneously in each oocyte-nurse cell
complex, although their initiation, duration, and conclusion are
somewhat variable. Two of these processes are the engulfment of nurse
cells and transformation of nurse cells into shrunken cells as they
discharge their cytoplasm, presumably into the oocytes. These
processes usually occur simultaneously, but in a few gonophores of
polyps of both species 20-25% of the nurse cells discharged their
cytoplasm before engulfment. This has only been observed in the peripher- al nurse cells. Engulfment of nurse cells (Fig. 10) usually commences after vitellogenesis is complete and the oocytes have started to fuse into a single oocyte. In both species, however, shrunken cells have been observed in vitellogenic oocytes before there are any indications of oocyte fusion. In gonophores of T. larvnx, virtually all of the nurse cells appear to be engulfed by the oocytes. Those not engulfed presumably do not persist long enough to be detected in sectioned gonophores. In T. indivisa most of the nurse cells are engulfed, but the excess nurse cells from several generations of oocyte-nurse cell complexes accumulate within the gonophore at the base of the spadix. The third simultaneously occurring process is the fusion of the 15-18 oocytes of the complex into one large oocyte containing a single germinal vesicle. When the oocytes begin to fuse, the intercellular bridges (Fig. 7), which reach 5-7 μm in diameter during vitellogenesis, expand and gradually disappear. When the fused oocyte of T. indivisa ruptures through the inner ectoderm into the subumbrellar space, about 100-150 shrunken nurse cells remain at the base of the spadix along with the septal cells that were intermixed with the germ cells of that complex. In some gonophores of T. indivisa there are also one or two large anucleate cellular fragments (30-55 μm diameter) that may represent portions of oocytes that did not successfully fuse.

At the base of the spadix in some gonophores of T. indivisa, there are what appear to be the remains of excess oocyte nuclei, which are surrounded by a small amount of cytoplasm. In Tubularia only one of the nuclei of the fusing oocytes survives to complete meiosis. The germinal
vesicles of the fusing oocytes seem to disappear gradually and only 5 or 6 of the original 15-18 are found in serial sections of oocytes that have almost completed fusion. No trace of abortive germinal vesicles has been found in the gonophores of T. larynx. However, up to 12 of what appear to be voided germinal vesicles have been found at the base of the spadix in gonophores of T. indivisa. These "germinal vesicles" are not found in gonophores in which the latest embryo has developed to an actinula although the discarded spetal cells and nurse cells from several oocyte-nurse cell complexes may be found at the base of the spadix.

Treatments used in attempts to increase the visibility of the intercellular bridges in the oocyte-nurse cell complexes were marginally successful in T. indivisa (Fig. 8), but failed with the gonophores of T. larynx. Rapid dehydration (directly from water to 95% ethyl alcohol) sometimes highlighted small sections of the peripheral nurse cells if the gonophores were opened before transfer to alcohol. The germ cells are tightly packed in the gonophores of T. larynx and only short sections of 3-4 oocytes could be accurately mapped, even in serial sections. The germ cells are much more loosely packed in the gonophores of T. indivisa which are much larger than those of T. larynx. The largest oocyte-nurse cell complexes mapped in T. indivisa contained 20; 22, and 45 cells although many smaller sections of other complexes were mapped (Figs. 18-21).

In most of the oocyte-nurse cell complexes each of the oocytes and central nurse cells were connected by intercellular bridges to 3 other cells. In one instance 2 of the oocytes, not connected to each other, were each connected to 4 other oocytes. In a few other instances, one
Oocyte with 4 intercellular bridges was found in each complex, but in most cases the center most and largest oocyte of a complex had only 3 intercellular bridges. In some of the complexes that were mapped, oocytes and nurse cells were found that contained 2 nuclei. Nurse cells that are directly connected to oocytes have larger nuclei than peripheral nurse cells and in some cases were difficult to distinguish from oocyte nuclei. The largest nuclei were usually found in oocytes in the center of an oocyte-nurse cell complex.

The oocyte-nurse cell complexes of both species of Tubularia contained approximately 1200-2000 cells. Since each oocyte or nurse cell normally had only one nucleolus, counts were made of the nucleoli in 10 μm serial sections of 6 vitellogenic oocyte-nurse cell complexes in each species. In T. larvnx, complexes contained 1233-1920 nucleoli (avg. 1521) and in T. indivisa, complexes contained 1293-1965 nucleoli (avg. 1570). The early embryos (20 cells or less) that were observed contained a smaller number of shrunken cells. Seven embryos of T. larvnx contained 798-2054 shrunken cells (avg. 1285) and the 6 embryos of T. indivisa contained 989-1532 shrunken cells (avg. 1287).

**Autoradiography**

The movement of labeled molecules from the enteron of injected polyps to germ cells was much slower than expected and was highly irregular in both of the experiments utilizing tritium labeled nucleotides. Two hours after injection, none of the labeled nucleotides had reached the racemes. At subsequent fixation times some racemes still showed little or no evidence of radioactive molecules while others were heavily labeled. The irregular movement of labeled molecules is partially consistent with the results from the practice
Figure 1. *T. larynx*. Underwater photograph (20 feet) of small colony. (X 1.5)
Figure 2. *Clava leptostyla*. Light microscopy photograph of a gonophore containing a single oocyte. (X 3300)
Figure 3. *T. larynx*. Light microscopy photograph of a young gonophore containing a small germ cell mass (GCM), endoderm lamella (EL), and inner ectoderm (IE) with a migrating I-cell in the endoderm. The spadix has not yet pushed into the germ cell mass. (X 600)
Figure 4. *T. larynx*. Light microscopy photograph of a gonophore with a horseshoe-shaped germ cell mass (GCM) and spadix (SP). (X 220)
Figure 5. T. larynx. Light microscopy photograph of two young gonophores; one with several oocytes in the early prophase of meiosis (arrows) and the other with a small germinal vesicle. (X 670)
Figure 6. *T. indivisa*. Light microscopy photograph of an oocyte-nurse cell complex that contains a single large oocyte (O), smaller oocytes (SO), and numerous nurse cells (NC). A dense layer (arrow) is seen on the inside of the nuclear membrane. (X 730)
Figure 7. *T. larynx*. Light microscopy photograph of an intercellular bridge between two oocytes. (X 800)
Figure 8. *T. indivisa*. Light microscopy photograph of a gonophore in which shrinkage has highlighted the intercellular bridges (arrows) between nurse cells. (X 980)
Figure 9. *T. indivisa*. Light microscopy photograph of an oocyte with two nuclei (arrow). (X 620)
Figure 10. *T. larynx*. Light microscopy photograph of a gonophore engulfment of the shrunken cells (arrow) by the oocytes has begun. (X 300)
Figure 11. *T. larvnx*. Actinula larva containing numerous shrunken cells (arrow) within the endoderm. (X 260)
Figure 12. *T. indivisa*. Light microscopy photograph of a gonophore containing a vitellogenic oocyte-nurse cell complex (ONC) and two oocytes of a young oocyte-nurse cell complex. (X 500)
Figure 13. *T. larvnx*. Light microscopy photograph of a gonophore containing an actinula (A) and egg (E) within the subumbrellar space (SC) and a developing oocyte-nurse cell complex (ONC) on the spadix. (X 166)
Figure 14. *T. indivisa*. Light microscopy photograph of a germinal vesicle (GV) at periphery of an oocyte. The layer of material on the inside of the nuclear membrane of the nurse cells (NC) is not seen in the germinal vesicles. (X 800)
Figure 15. *T. larvnx*. Light microscopy photograph of a gonophore that has ceased production of oocyte-nurse cell complexes. Septal cells and unengulfed nurse cells remain on the spadix. (X 700)
Figure 16. *T. larynx*. Light microscopy photograph of the male (M) and female gonophores of a hermaphroditic polyp. (X 130)
Figure 17. *T. indivisa.* Line drawing of 43 cells of an oocyte-nurse cell complex. Three cells contained two nuclei and three cells had four intercellular bridges.

(In each line drawing the sizes of the nuclei are in micrometers. The cells are drawn to show approximate relative cellular volume.)
Figure 18. *T. indivisa*. Line drawing of 22 cells of an oocyte-nurse cell complex.
Figure 19. *T. indivisa*. Line drawing of 20 cells of an oocyte-nurse cell complex.
Figure 20. *T. indivisa*. Line drawing of 17 cells of an oocyte-nurse cell complex.
Figure 21. *T. indivisa*. Line drawing of 13 and 11 cells of oocyte-nurse cell complexes.
injections of methylene blue-sea water during which some racemates were unstained while others were heavily stained. However, during practice injections, methylene blue usually reached some of the racemates of each polyp within 20 minutes. There were no noticeable differences in results of the two labeling experiments run at different temperatures; observations from both experiments have been combined.

Movement of the $^3$H-labeled uridine to the gonophores and germ cells was extremely slow. No labeled uridine was detected in any of the racemates 2 hours after injection. After 5 hours of incubation, most radioactivity was localized in the endoderm of the stalk and the spadix of the gonophore. Labeled material appeared to be localized in the large vesicles of food and in the cytoplasm of the endodermal cells. In a few gonophores terminated after 5 hours of incubation, radioactivity was detectable in the cytoplasm of the first generation of germ cells of younger gonophores and in the vitellogenic generation of germ cells of older gonophores. No detectable differences were noted between the gonophores terminated after 12 and 24 hours of incubation. At both sample periods some of the gonophores were labeled while others were not. A small amount of label was still present in the spadix of the labeled gonophores while most label was in the germ cells. Labeled uridine became detectable in the oocytes and nurse cells at the same time and no radioactivity was detected in the shrunken cells.

Some of the gonophores fixed after 45 hours of incubation were sectioned and stained for routine histology, but were not coated with emulsion. These polyps were fed freshly hatched brine shrimp after 24 hours of incubation, but were in poor condition after 45 hours. The gonophores that were sectioned showed signs of degeneration.
As in the case of uridine, no $^3$H-thymidine was detected in the gonophores of polyps fixed after 2 hours of incubation. A large amount of labeled nucleotide had reached the gonophore after 5 hours of incubation, but most of it was still in the spadix. After 5, 8, 12, and 24 hours of incubation, nuclei of the proliferating germ cells in young gonophores forming the first oocyte-nurse cell complex (Figs. 22, 23) were heavily labeled. In the older gonophores, only the nuclei of the proliferating, pre-vitellogenic generation were labeled. No differences were noted between the labeling of the nuclei of the nurse cells and oocytes and no labeling was detected in the shrunken cells. Label was detected in the eggs and early cleavage stages of T. indivisa, but it was not possible to determine if the label was on the surface membrane of the densely packed vesicles of yolk or in the cytoplasm surrounding the vesicles (Fig. 24).

**Transmission Electron Microscopy**

Modifications in the fixation, embedding, and staining procedures produced some variations in the ultrastructural appearance of the gonophores and germ cells. Best overall results were obtained with gonophores that were fixed at pH 7.4 or 7.6 and embedded in EPON 812 using DMP-10 as the accelerator. The cytoplasm was much less granular and better preserved at pH 7.4 or 7.6; the matrix between the cristae in mitochondria was usually better preserved at pH 7.8. Infiltration of shrunken cells and yolk was generally poor with the EPON 812-Araldite 506 mixture and the lumen of the spadix was often empty of plastic. Spurr's embedding medium, which was expected to yield the most rapid penetration of tissue as a result of its low viscosity, penetrated poorly and provided the worst support. Best infiltration and tissue
Figure 22. *T. larynx*. Autoradiograph of a young gonophore from a polyp injected with H3-thymidine. The proliferating germ cells were labeled (arrow). (X 440)
Figure 23. *T. indivisa*. Autoradiograph of a gonophore from a polyp injected with H$^3$-thymidine. The proliferating cells of a young oocyte-nurse cell complex were labeled (arrow). (X 680)
Figure 24. *T. indivisa*. Autoradiograph of an egg from a polyp labeled with H\(^3\)-thymidine. Label (arrow) is associated with yolk vesicles. (X 730)
support was obtained with EPON 812 and prolonged infiltration using DMP-10, although the shrunken cells often moved slightly during sectioning. No differences between fixation at 4°C and room temperature (20-25°C) were noted.

Membrane bound droplets 2.0-8.5 μm in diameter, assumed to be nutrient material, were found in the enteron (Fig. 25) of the stalk of the gonophore and spadix. The cells of the spadix are filled with these vesicles and many vesicles are found partially engulfed (or partially ejected). The luminal surface of the cells of the spadix have numerous microvilli (0.5-2.0 μm long) and flagella are occasionally present (Fig. 34). The luminal ends of adjacent endodermal cells are connected by a junctional complex. Discoidal coated vesicles are abundant in the cytoplasm of the endodermal cells. A homogeneous layer of cytoplasm is usually present in the base of the cells of the spadix adjacent to the secondary mesolamella that separates the spadix from the germ cell mass. The secondary mesolamella is about 0.2 μm thick and is penetrated by pores 0.22-0.28 μm in diameter. Small vesicles of homogeneous cytoplasm 0.25-0.50 μm in diameter are dispersed among the germ cells and may form a layer between them and the secondary mesolamella (Fig. 27). These vesicles appear to pass through the pores in the secondary mesolamella (Fig. 26).

During the development of the oocyte-nurse cell complexes, four categories of germ cells are distinguishable morphologically using transmission electron microscopy. These are the pre-vitelligenic germ cells, vitelligenic nurse cells with small vesicles of yolk, vitelligenic oocytes with large vesicles of yolk, and the engulfed nurse
cells or shrunken cells. Observations of pre-vitellogenic germ cells apply to the oocytes and to the nurse cells. These cells are indistinguishable until vitellogenesis begins and the difference in the size of their vesicles becomes apparent (Figs. 31, 32, 35).

The pre-vitellogenic germ cells of *Tubularia* are identical to hydrozoan I-cells. They are 8.0-8.5 μm in diameter and have nuclei 3.5-4.0 μm in diameter with 1.5-1.7 μm nucleoli. There is usually no Golgi apparatus and they have only a few mitochondria, and many ribosomes. As the pre-vitellogenic germ cells or oogonia of complex proliferate by mitosis, they do not complete cytokinesis but remain interconnected by cytoplasmic bridges (Figs. 28, 29) with a diameter of 0.55-0.70 μm. Longitudinal sections of intercellular bridges often reveal the presence of septate desmosomes connecting the cytoplasmic membranes of the two cells. The septate desmosomes between germ cells were only found in close proximity to an intercellular bridge. While some sections of bridges lacking associated desmosomes were encountered, most bridges between the pre-vitellogenic germ cells had at least one septate desmosome; three bridges were observed to have a desmosome on each side. In one thin section, a septate desmosome was encountered between a very early vitellogenic germ cell and one of the cells of the inner ectoderm. Numerous bridges were detected between spermatids in male gonophores, but no desmosomes were visible. No cross sections of intercellular bridges were encountered using TEM and the number and arrangement of the desmosomes associated with each intercellular bridge is not known. No desmosomes were detected between larger, vitellogenic germ cells (Fig. 30).

Yolk was not seen in germ cells smaller than 9.5 μm in diameter.
These cells have nuclei 4.2-4.5 μm and nucleoli 2.0-2.2 μm in diameter. It was not possible to distinguish between nurse cells and oocytes at this stage. In young nurse cells the vesicles of yolk are 0.24-0.40 μm in diameter. As the nurse cells enlarge they eventually produce vesicles that are up to 1.4 μm in diameter. In the nurse cells there are no indications of fusion of these vesicles. As vitellogenesis commences in an oocyte–nurse cell complex there is an increase in the number of Golgi apparati and mitochondria in the nurse cells. At this time, the nuclei of the nurse cells are often located near the periphery of the cell on the side facing away from the oocytes. Mitochondria are commonly found in one or two clusters and the numerous Golgi apparati are clustered in groups of five or less. Each Golgi apparatus is 0.7-1.0 μm long, consists of 5-7 lamellae about 0.2 μm thick, and produces vesicles 0.05-0.06 μm in diameter. The intercellular bridges between the nurse cells normally increase in diameter to a maximum of about 1.8-2.1 μm, although one intercellular bridge was 3.4 μm.

Using TEM, the primary oocytes and nurse cells are distinguishable when the vesicles of yolk commence to fuse in the oocytes. There are two size classes of vesicles in the oocytes once fusion starts. Smaller vesicles are 0.5-1.6 μm in diameter (Fig. 35) and are most abundant at the periphery of the oocytes near the nurse cells. In a large primary oocyte, most of the larger vesicles are 3.5-8.0 μm in diameter with a few approximately 10 μm. In the oocytes, vesicles that are intermediate between the two size classes are surrounded by a cluster of small vesicles (Fig. 36). As vitellogenesis progresses, the oocytes become densely packed with large vesicles of yolk which are eventually separated by only a very thin layer of cytoplasm. In one
micrometer thick plastic sections of the gonophores of *T. larynx* there is a distinct difference in the staining reactions of the two size classes of yolk vesicles. The smaller vesicles in both the nurse cells and oocytes stain strongly with toluidine blue, while the larger vesicles in the oocytes of the same plastic section stained lightly or not at all (Fig. 31). This differential staining was not nearly as pronounced in *T. indivisa* in which the smaller vesicles did not stain very strongly.

Engulfed nurse cells or shrunken cells (Fig. 33) were found in the large, fusing oocytes and larvae, and were enclosed by a membrane. Most of the shrunken cells viewed by TEM had been engulfed singly, but up to three were found within a single vesicle. Serial polyester sections revealed up to 12 shrunken cells in a single vesicle although most vesicles contain 1-4. The shrunken cells contained a large amount of heterochromatin in their nuclei and a small amount of cytoplasm with a few vesicles of yolk 0.5-0.7 μm in diameter.

The microvillar layer found on the surface of the embryo appears to begin to develop before the oocytes complete fusion and the single, fused oocyte ruptures through the manubrial layer of the inner ectoderm into the subumbrellar space. The microvillar layer on the surface of the embryo is 0.8-1.2 μm thick and is morphologically identical to the layer of periderm on the outside of the gonophore. The two layers of inner ectoderm have a similar microvillar layer that is 0.5-0.70 μm thick and is sometimes visible before the opening of the subumbrellar space. Both *T. larynx* and *T. indivisa* contain striated muscle fibers not found in the manubrial layer of the inner ectoderm covering the germ cells. The septal cells, which attach the manubrial layer of the
inner ectoderm to the mesolamella of the spadix, are commonly seen among the germ cells in sections of the germ cell mass. The spetal cells are much less electron dense than the germ cells (Fig. 28).
Figure 25. *T. larynx*. Electron micrograph showing engulfment of nutrients (N) by the cells of the spadix and the secondary mesolamella (ML) separating the germ cells (GC) and spadix (SP). (X 1800)
Figure 26. *T. larynx*. Micrograph showing material (arrow) passing from the spadix to the germ cell mass (GCM). (X 2900)
Figure 27. *T. larynx*. Micrograph of nutrient vesicles (NV) next to the secondary mesolamella.
Figure 28. *T. larynx*. Micrograph of intercellular bridges (arrows) between previtellogenic germ cells and a cross section of a septal cell (SC). (X 5400)
Figure 29. *T. larynx*. Micrograph of an intercellular bridge with a septate desmosome (arrow). (X 10,800)
Figure 30. *T. larynx*. Micrograph of an intercellular bridge between young vitellogenic germ cells. (X 6700)
Figure 31. *T. indivisa*. One micrometer thick plastic section of nurse cells (NC) with small vesicles of yolk and an oocyte (O) with large and small vesicles. (X 830)
Figure 32. *T. larynx*. Micrograph of a nurse cell (NC) with small vesicles of yolk and oocytes (O) with both large and small vesicles. (X 3600)
Figure 33. *T. larynx*. Micrograph of a shrunken cell with a nucleus containing heterochromatin. (X 7000)
Figure 34. *T. larynx*. Micrograph of the spadix with microvilli and discoidal coated vesicles (CV). (X 16,500)
Figure 35. T. larynx. Micrograph of early vitellogenic germ cells. (X 4300)
Figure 36. *T. larvnx*. Micrograph of an oocyte in which the small yolk vesicles appear to fuse to form the large yolk vesicles found only in the oocytes. (X 6100)
After Weismann's work (1883) it seemed definitely established that in the Hydrozoa the germ cells have an ectodermal origin and migrate to the gonophores, although the germ cells have endodermal origin in the Scyphozoa and Anthozoa according to Campbell (1974), Szmant-Froelich et al. (1980), and Larkman (1983). Other work on various hydroids, including species of *Tubularia* by Koch (1876), Hargitt (1909), Liu and Berrill (1948), and Berrill (1961), seemed to indicate that, in at least some hydroids, germ cells have an endodermal origin. Goette (1907) concluded that the spermatogonia in *Clava multicorns* had an ectodermal origin while the oogonia were derived endodermally.

My observations on the female gonophores of *T. larynx* and *T. indivisa* indicate that the oogonia are derived from undifferentiated cells (I-cells) that proliferate in the ectoderm of the stalk of the gonophore and which migrate in the endoderm to an apical position in the gonophore bud. This contrasts sharply with the work of Liu and Berrill (1948) on *T. crocea* and Benoit (1925) on *T. mesembryanthemum* who reported the direct conversion of endodermal cells into germ cells without migration. My observations, for the most part, agree with those of Lowe (1926) who reported the proliferation of I-cells in the ectoderm of the gonophore stalk and subsequent migration to the gonophore. However, she reported that the migration of I-cells occurred in the ectoderm while I found an endodermal migration. It is interesting to consider that the migration of the primordial germ cells from the endoderm at the base of the allantois and yolk sac to the germinal ridge.
in vertebrates is well established, while in the "primitive" and structurally simple hydrozoans the problem is basically unresolved.

Weismann's (1883) discovery of germ cell migration in hydroids helped to shift the emphasis from the cellular layer in which recognizable germ cells first appear, to also include the cellular type from which they originate. The origin of hydrozoan germ cells has been a controversial topic and the subject of numerous studies by many investigators. The solution to this problem has been made especially difficult since the studies have involved interpretations of a dynamic process based largely on the information based on fixed and sectioned material. Even where modern investigators have agreed that the cells that will form a gonophore bud or medusa bud are derived from I-cells, there is then the question of the origin of the I-cells. Bierbach and Hofmann (1973) argued that the localized accumulations of I-cells that produce medusa buds are derived form I-cells present in the ectoderm of the stolons. Bouillon and Werner (1965), on the other hand, considered that local somatic cells dedifferentiated to form the I-cells for medusa development.

In the Cnidaria, according to Campbell (1974, p. 152), "Germ cells and their accessory cells arise from interstitial cells." Adult hydrozoans maintain a proliferating population of I-cells in the ectoderm (Hyman, 1940; Bouillon, 1968), however, these I-cells originate from the larval endoderm during early cleavage in at least some hydroids. According to Nagao (1965) and Summers and Haynes (1969), the smaller cells produced from unequal divisions of the large endodermal cells become the I-cells and migrate to the ectoderm of the larva. While it is clear from recent work on *Hydra* that the ectodermal I-cells,
regardless of their larval origin, produce the germ cells under normal conditions (Burnett, Davis and Ruffing, 1966; Tardent, 1974; Honegger, 1981), this does not hold in the laboratory. Normandin (1960), Haynes and Burnett (1963), Davis et al. (1966), and Burnett et al. (1966) have demonstrated the differentiation of I-cells from endodermal cells under laboratory conditions. Burnett et al. (1964), after their work with \textit{Hydra viridis}, concluded that there can be no separate lines of interstitial cells to form the cnidoblasts, neurons, or germ cells since endodermal gland cells can dedifferentiate and produce a normal, sexually mature polyp.

In \textit{Tubularia}, enlargement of the oocytes that have entered the prophase of meiosis I does not begin until after the migrating I-cells reach the apex of the gonophore. This makes it difficult to separate the I-cells which develop into germ cells from the I-cells which will differentiate into nematoblasts, especially when both processes occur in the same region of the polyp. Lowe (1926) also noted this problem in \textit{T. larvnx} and in some of my sections of both \textit{T. larvnx} and \textit{T. indivisa} nematocytes were found among the germ cells in young gonophores. A potential solution to this problem may be to study hydroids in which the oocytes differentiate and begin the growth phase of meiosis I during or even before migration. In \textit{Clava leptostyla}, oocytes with a distinct germinal vesicle are first distinguishable in the endoderm of the polyp, before they reach the gonophores where they mature and are fertilized (Hargitt, 1906; Hargitt, 1916; personal observations). Wasserthal (1973) found \textit{Eudendrium armatum} an even better species to study since the oocytes begin to enlarge before migration commences. He found the young oocytes growing in the ectoderm of the hydrocaulus (stalk or stem
of the polyp) surrounded by nematocytes in various stages of nematocyst development. Later stages of oocyte growth were found in the ectoderm of the gonophore stalk, but at some location they penetrate the mesolamella of the gonophore and complete oogenesis in the endoderm of the gonophore.

David and Challoner (1974) used the thiolacetic acid-lead nitrate staining technique of Lentz and Burnett (1961) to locate the developing nematocytes in intact hydra. It might be possible to develop a similar histochemical technique to differentiate the nematoblasts and nematocytes from the early germ cells. This is especially feasible in a hydroid such as Eudendrium armatum where a morphological distinction between the two types develops quickly. This technique could then be applied to hydroids such as Tubularia in which there seem to be no morphological characters or simple staining techniques to differentiate young germ cells and nematoblasts. It might then be possible to determine young germ cells and nematoblasts. It might then be possible to determine the location of germ cell origin in living Tubularia.

Campbell (1974, p. 152) considered the problem of the germinal interstitial cells and concluded that, "it is generally impossible to assess the origin of a cell type which cannot be distinguished from other interstitial cells, at least by classical histological techniques employed."

In addition to the question of the location of origin of the germ cells, a number of other problems concerning hydrozoan reproduction have not been resolved, in spite of numerous studies. These questions include the cellular type from which germ cells originate and whether or not the germ cells actually migrate to the gonophores and are not
simply carried by tissue movements. Other questions concern the formation of the entocodon. In many species it is not known if the entocodon develops in the ectoderm or endoderm and whether the germ cells develop from the entocodon or have a separate origin. While these topics are no longer as controversial as they were in the late 1800's and early 1900's, the wide variety of results, sometimes with the same species, is troublesome. Resolution of the inconsistencies in the interpretations have been made difficult since all of the investigators have used fixed and sectioned material to interpret dynamic processes.

The present results on germ cell movement and origin are, unfortunately, also based on circumstantial evidence and interpretations of fixed and sectioned material. Examination of my specimens of *T. larynx* and *T. indivisa* has indicated that the major site of I-cell proliferation is the ectoderm at the base of the gonophores. Between the site of I-cell proliferation and the apex of the gonophore buds, single and groups of 2-4 I-cells are abundant in the endoderm. These endodermal I-cells appear to be in the process of migrating to the apex of the gonophore bud after having been produced in the ectoderm at the base of the gonophore. Endodermal I-cells were rare in other areas of the polyp including the distal portions of the racemes which contain only older gonophores. At the apex of the gonophore buds the I-cells accumulate and proliferate in the endoderm on the inside of the mesolamella that separates the endoderm and ectoderm. The entocodon and germ cell mass which form these apical I-cells are thus both endodermal structures. Brauer (1981), working with *T. mesembranaceum*, and Lowe (1926), working with *T. larynx*, also
reported that the germ cells originate in the ectoderm at the base of the gonophores. However, both of these investigators also agreed that the entocodon and the germ cells developed in the ectoderm at the apex of the gonophores. While Brauer reported that the germ cells migrated in the endoderm, Lowe found an ectodermal migration in \textit{T. larynx}, one of the species in which I am reporting an endodermal migration. Obviously, better techniques and more than circumstantial evidence are required to resolve this problem.

\textit{Tubularia larynx} and \textit{T. indivisa} produce large, yolky eggs (about 410 \(\mu\text{m}\) and 700 \(\mu\text{m}\) respectively) and brood the embryo within the gonophore to an actinula before the larva is released. The results of Hargitt (1919) and Rees (1957) indicate that the species of hydroids with large polyps produce relatively large eggs, usually greater than 200 \(\mu\text{m}\) in diameter. These species include colonial hydroids such as \textit{Pennaria}, \textit{Hybocodon} and \textit{Tubularia} which Rees considers to be essentially solitary, and the solitary hydroids including \textit{Corymorpha} and \textit{Hydra}. On the other hand, most of the colonial species, including \textit{Obelia}, \textit{Campanularia}, \textit{Clava}, \textit{Goniothyrea}, and \textit{Hydractinia}, have small polyps and produce relatively small eggs that are less than 170 \(\mu\text{m}\) in diameter. Hargitt (1919) also noted that during oogenesis in the species that produce large eggs, numerous excess oocytes, which are a type of accessory cell, are produced and consumed by the relatively few growing oocytes in each gonophore. However, the species that produced small eggs are apparently nourished directly from the enteron without any accessory cells.

A variety of accessory cells have been described and the participation of accessory cells of somatic origin, or follicle cells,
especially during vitellogenesis, is known in many animals, including vertebrates. Balinsky (1975) reported follicle cells and oocytes with interdigitating processes in all vertebrates that have been studied, with the oocyte surface typically possessing numerous microvillar projections (Anderson, 1974). These projections increase the surface area of contact between the plasma membranes of the oocyte and its follicle cells which provide an exogenous source of yolk. All of the material utilized by the oocytes are not produced by the follicle cells which may act as a barrier between the oocytes and other exogenous sources of nutrients. Bellairs (1965, p. 215) reported, "In the adult hen each oocyte is surrounded by a capsule of follicle cells and all the raw materials that enter the oocytes must pass through this capsule."

Not all of the somatic accessory cells develop interdigitations with the oocytes for the passage of nutrients. At least some lizards develop another type of follicle cell, the pyriform cell (Naeves, 1971; Taddei, 1972). Pyriform cells are unusual in that, while they are somatic cells, they develop intercellular bridges with the oocytes. These bridges were not found in small follicles and are thought to form secondarily, as the oocytes grow, and not as a result of incomplete cytokinesis. Neaves reported a system of about 10,000 pyriform cells surrounding each oocyte and suggested that the pyriform cells function in the maintenance of dormant oocytes since they disappear before the onset of yolk accumulation.

While accessory cells of germ line origin, usually termed nurse cells, have not been reported in vertebrates, numerous investigators have reported intercellular bridges between the oocytes in mammalian
ovaries. Such intercellular associations have been reported in humans (Stegner, 1967; Gondos, 1973), mice (Ruby et al., 1969), rats (Franchi and Mandl, 1962), hamsters (Weakly, 1966, 1967), and rabbits (Zamboni and Gondos 1968). Zamboni and Gondos suggest that the intercellular bridges may account for the synchronous differentiation, maturation, and degeneration of groups of germ cells and that syncytia of germ cells may form in all mammals. They also speculate that the oocytes that survive may be the ones that complete cytokinesis. Baker (1963) reported that in humans less than one oocyte in 20 survives to the age of seven years. McReynolds et al. (1984) reported the presence of mitochondria and ribosomes in the cytoplasm of the bridges and suggested that they may allow the exchange of organelles and nutrients. It is interesting to speculate that perhaps one oocyte in each cluster survives, while the remaining oocytes function as nurse cells before degenerating.

A wide variety of types of accessory cells of germ line and somatic origin are known from different groups of invertebrates, including ascidians which have been studied by Tucker (1942), Kessel and Kemp (1962), and Berrill (1975). Ascidians develop both an outer follicular epithelium and a layer of test cells during oogenesis. The primary follicle cells are derived from the germinal epithelium, from which the oocytes also differentiate. The test cells differentiate from the cells of the primary follicle and penetrate into the superficial cytoplasm of the oocyte, increasing the surface area of contact. Both the test cells, which are eventually extruded from the oocyte, and the follicle cells pass nutritive material to the oocyte. Since the unspecialized germinal epithelium produces the oocytes and
the follicle cells, Berrill (1961) considered the follicle cells to be
descendants of abortive oocytes. If this reasoning were applied to
hydrozoans, the cnidoblasts could be considered to be descended from
abortive oocytes since they also differentiate from I-cells. However,
Berrill (1975) reported that totipotent cell types exist in tunicates
and no distinct germ line had yet been demonstrated.

Projections of follicular cells penetrate into the oocytes of the
decapod cephalopods. Arnold and Williams-Arnold (1977) reported
that the cells of the follicular epithelium proliferate mitotically and
that folds of this tissue, consisting of numerous follicle cells,
penetrate deeply into the oocytes. The interdigitations tremendously
increase the surface area of contact between the follicular epithelium
and the plasma membrane of the oocyte. In addition, the follicle cells
develop multiple nucleoli and a larger nucleus to cytoplasm ratio and
eventually form a syncytium surrounding the oocyte. Autoradiography
with $^3$H-leucine was used by Selman and Wallace (1972) to demonstrate
that the follicle cells are the major site of protein synthesis in the
follicle cell-oocyte complexes in *Loligo pealei*.

In some species of turbellarians and in a few narcomedusae, the
follicular epithelium has been reduced to a single accessory cell which
surrounds the oocyte. Newton (1970), working with a freshwater
turbellarian, *Hydromedusa grisea*, found that one of the accessory
parenchymal cells completely envelops a growing oocyte and that both
nutrition by diffusion and fertilization take place through the
accessory cell. A similar development was noticed in the marine
tubellarian, *Alaurina composita* (personal observation). Berrill (1950) described work by a number of investigators including Bigelow (1909) on *Pegantha smaragdina* and Metschnikoff (1886) on *Cunina proboscidea*, both of which are narcomedusae. In both species a specialized "nurse cell", the phorocyte, whose origin is unknown, surrounds the oocyte and early embryonic stages. The nucleus of the phorocyte of *Cunina* was later incorporated into the embryo. The phorocytes of *Pegantha* become multinucleate, but their fate was not described. Davidson (1968) mentioned that certain annelids develop two "nurse cells" with polytenic nuclei along with each oocyte, but did not mention the species.

The accessory cells described above have not been demonstrated to come from the same cell line as the oocytes with which they are associated and, although they are sometimes referred to as nurse cells, are more accurately considered follicle cells (Berrill and Kapp, 1976). However, numerous examples also exist of accessory cells of germ line origin that participate in the nutrition of the oocyte or embryo.

Thorson (1950) reported examples of embryos with non-pelagic development that feed on nurse eggs in crinoids, ophiuroids, polychaets, nemerteans, and prosobranchs. In this method of embryonic nutrition, a large number of eggs are deposited together, but only a few develop into embryos. The nurse eggs are devoured by the embryos before they hatch. The extreme example was the prosobranch mollusc, *Volutopsis norvegica*, in which he reported that 50,000-100,000 nurse eggs are consumed by a single embryo.

In other invertebrates, the oocytes that do not complete
development are used as nourishment for the developing oocytes. Pollock (1975) mentions this process in tardigrads and describes the abortive oocytes functioning as "nurse cells" while Fell (1969) reports that the oocytes in the sponge, Haliclona ecbasia, engulf whole nurse cells. However, in both tardigrads and sponges the engulfed cells were not directly connected to the oocytes by intercellular bridges. If we define the term nurse cell to include only those cells of germ line origin and connected by intercellular, cytoplasmic bridges to the successful oocyte (Anderson, 1974; Berrill and Kemp, 1976; Browder, 1980), then we are left needing a term for the accessory cells of germ line origin which lack intercellular bridges to the oocytes and have not completed meiosis. Pollock (1975) used the term abortive oocytes while Raven (1961) called them nurse cells or nutrimentary eggs and considered them to be abortive germ cells. Hargitt (1919) also mentioned the problem of accessory cells which are called nurse cells, but which are "consumed instead of preparing food", but did not suggest an alternative. The term, nutritive oocytes, which has already been used (Spracklin, 1978), still appears appropriate since it implies both the function and the stage of meiosis.

The most highly organized development of accessory cells is found in the species in which true nurse cells, possessing intercellular bridges to the oocyte, are produced during oogenesis and provide the oocytes with nutrients. This type of germ cell nutrition is best known from the insects which have evolved two different types of meroistic ovaries, polytropic and telotrophic, as well as the panoistic ovaries which do not contain nurse cells (Davidson, 1968). In the telotrophic ovaries the nurse cells are connected to the oocytes by a long
nutrient cord. Woodruff and Anderson (1984) have used microinjected dyes to demonstrate the physiologic continuity between the oocytes and nurse cells that is provided by the cords, movement of dye along the cord, and the coupling of the oocytes and follicle cells, presumably by gap junctions. In the polytropic ovaries, of which Drosophila melanogaster is a typical example, a series of cell divisions, with incomplete cytokinesis, produces a complex of an oocyte and a species specific number of nurse cells. In Drosophila, a series of four divisions produces a complex consisting of a central oocyte and 15 interconnected nurse cells (Brown and King, 1964; King and Aggarwal, 1965; Koch and King, 1966). Growth of the oocyte in Drosophila is supported by the nurse cells and the cytoplasmic volume of the oocyte can increase by 90,000 times in only 3 days. After the four cycles of mitosis with incomplete cytokinesis, each of the two central cells of the oocyte-nurse cell complex have 4 intercellular bridges, a common one and 3 each to nurse cells. The other 14 cells have a maximum of 3 bridges each. The two central cells, which Koch et al. (1967) refer to as pro-oocytes, begin synopsis of meiosis I. Synapsis is later arrested in one of the two pro-oocytes which develops into another nurse cell, while the remaining pro-oocyte becomes the successful oocyte. They found that no oocytes develop in abnormal complexes in which no cell has 4 intercellular bridges, but were left with the question of what factor determines which of the pro-oocytes becomes the oocyte.

Davidson (1968) reported that chromosome diminution occurs during nurse cell formation in some beetles that also produce an oocyte-nurse cell complex of 15 nurse cells and 1 oocyte. Only the oocytes retain
the normal chromosomal complement in these species. In *Drosophila*, the nurse cells actually become highly polyploid as their DNA content reaches a maximum of 1024C and their nuclei increase in volume up to 6000-fold (Koch and King, 1966). This may actually represent amplification of only a small portion of the genome, similar to what has been reported in the oocytes of *Xenopus laevis* by Perkkowska *et al.* (1968). In *Xenopus* oocytes the number of copies of the ribosomal DNA is increased by 2500 times and 1400 to 1600 nucleoli are produced in the germinal vesicle.

As vitellogenesis progresses in *Drosophila*, the germinal vesicles enlarge and may increase to 900 times their original volume; however, the nurse cell nuclei are much larger. Raven (1961) reported that the nuclei of oocytes of animals with nurse cells are generally smaller than the oocyte nuclei of similar species that do not form nurse cells. Pianka (1974) has reported an animal in which the oocyte nuclei do not enlarge at all during oogenesis. In the ctenophore, *Bolinopsis microptera*, the oocyte is part of a complex of about 100 cells, including 99 nurse cells and 1 oocyte. In contrast to the 4 intercellular bridges of each *Drosophila* oocyte, each oocyte and the inner nurse cells of *Bolinopsis* have only 3 intercellular bridges. However, in each of the 3 clusters of nurse cells attached to each oocyte, the 7 nurse cells closest to the oocyte became highly polyploid. According to Pianka, "all major synthetic functions of oogenesis in *Bolinopsis* are apparently performed by the nurse cells, and not by the oocyte."

Hydroids display a wide range in the structural complexity of the gonophores and in the involvement of accessory cells during oogenesis.
The structurally simplest are the stylloid gonophores, whose walls are only continuations of the ectoderm and endoderm of the polyp (Kuhn, 1913; Teissier, 1926; Hyman, 1940). The gonophores of *Eudendrium armatum* are typically stylloid. Each gonophore contains only a single oocyte and no accessory cells of any kind, although the spadix is well developed and wraps about two thirds of the way around the oocyte (Wasserthal, 1973). The other types of hydrozoan gonophores, heteromedusoid, cryptomedusoid, and eumedusoid, are increasingly complex and develop additional cellular layers, an endoderm lamella and inner ectoderm, but there has been no suggestion that any of these cells function as follicle cells. However, in many hydroids, cells of germ line origin are used as food or to produce nutrients for the developing oocytes. *Pennaria tiarella* has been investigated by Smallwood (1899), Hargitt (1900), Hargitt (1909, 1919), Berrill (1952, 1961), and Cowden (1964). In the gonophores of this species, several thousand oocytes differentiate in each gonophore, but only 6-10 of the oocytes complete meiosis. The remainder of the oocytes are absorbed or engulfed and quickly digested by the growing or definitive oocytes. No trace of the engulfed oocytes remains at the completion of meiosis. Similar examples of this form of oocyte nutrition, which Cowden (1964) referred to as the "most primitive form of nutrimentary oogenesis", have been reported in other species of hydroids. Large numbers of nutritive oocytes have been reported in the gonophores of *Acaulis primaria* by Berrill (1952), *Rhizogeton fusiformis* by Spracklin (1975), and in *Millepora* by Mangan (1909). In all of these species, the excess or nutritive oocytes are engulfed and quickly digested and no trace of the engulfed oocytes remains in the eggs. Intercellular bridges have not
been reported among the oocytes in any of these species.

In other hydroids in which large numbers of nutritive oocytes are consumed by a few successful oocytes, the remains of the engulfed cells persist through embryonic development and are known as pseudocells (Brauer, 1891) or shrunken cells (Zihler, 1972). May (1903) reported that single nutritive oocytes and clusters of up to 7-8 are engulfed and enclosed in vacuoles in the eggs of _T. crocea_ while, during this research, as many as 12 shrunken cells were found in a vacuole in early cleavage stages of _T. indivisa_. The shrunken cells have been reported in all of the species of _Tubularia_ and _Hydra_ that have been investigated, most recently by Lowe (1926), Nagao (1965), and Boelsterli (1975) in _Tubularia_, and by Zihler (1972) and Tardent (1974) in _Hydra_. Zihler demonstrated conclusively that the shrunken cells were the remains of engulfed whole cells, and not simply the remains of nuclei.

During oogenesis in both _Tubularia_ and _Hydra_, a single egg is produced in each reproductive structure from the originally large number of germ cells that differentiate. There are initially over 1200 germ cells in each gonophore of _Tubularia_ and, according to Zihler's figures, at least 100 in _Hydra_. Based on the work on _Hydra circumcincta_ by Zihler and Tardent, oogenesis in each _Hydra_ ovary begins with the accumulation of a variable number (at least 3) of clusters of I-cells which differentiate into the initial germ cells, called Oogonia I. The Oogonia I develop into Oogonia II, which grow and synthesize yolk. One of the Oogonia II in each cluster becomes a primary oocyte and engulfs the remaining Oogonia II, transforming them into shrunken cells. All of the oocytes then fuse to form a single
large oocyte which completes meiosis. Only one of the oocyte nuclei survives. Tardent found intercellular bridges between the Oogonia II in a cluster, but made no mention of the number or arrangement of bridges. Burnett et al. (1966) reported the mitotic proliferation of I-cells to form syncytial nests of up to 32 cells, which differentiate into cnidoblasts or spermatogonia. The intercellular bridges in the clusters of Oogonia II suggest that each cluster is descended from a single I-cell. However, neither investigator found intercellular bridges between the primary oocytes.

There have been some previous indications of incomplete cytoplasmic divisions during oogenesis in two species of Tubularia. Lowe (1926), working with T. larynx, suspected that the cytoplasmic divisions that produced the germ cells were incomplete since the shrunken cells were engulfed in groups, but did not see the bridges. Boelsterli (1975), reporting on T. crocea, included one electron micrograph of an intercellular bridge between oogonia and in Figure 15 included drawings of clusters of young oocytes interconnected by cytoplasmic bridges. Boelsterli believed that most of the oocytes disintegrated and were incorporated, through some unknown process, by the approximately 15 growing oocytes which fused to form one large oocyte. He also briefly reported on egg fragmentation experiments with Tubularia and Hydra. His account is not extremely specific, but he confirmed that the germinal vesicles of the oocytes which fuse to form the single Hydra oocyte persist until just before the continuation of meiosis. However, he reported that similar experiments failed with Tubularia. These results seem to indicate that the oocytes which fuse in Hydra to produce the egg are essentially identical cells, while the
cells which fuse to produce an egg in *Tubularia* are not identical.

The current research on *T. larynx* and *T. indivisa* has revealed an extensive network of intercellular bridges among the germ cells in the female gonophores of both species. The cytoplasm of 1200 to 200 germ cells is incorporated into a single oocyte which completes meiosis, and it appears that all of these cells are interconnected into a single complex by intercellular bridges. It can be hypothesized that each complex, consisting of at least 1200 cells, is formed by the proliferation of one initial cell, since, except for the pyriform cells of lizards, intercellular bridges normally form only as a result of incomplete cytoplasmic division. The lizard pyriform cells also represent a much simpler system, with each pyriform cell directly connected to the oocyte. It is hard to imagine a mechanism by which secondary cell contact could result in the repeated formation of essentially identical complexes of cells, interconnected by cytoplasmic bridges in a highly organized pattern. Since they are all descended from a single cell and connected to the oocyte by intercellular bridges, the 1200 to 2000 cells in each complex that are engulfed and transformed into shrunken cells meet even the highly restricted definition of nurse cells used here.

While the majority of the cells in the germ cell complexes of *Tubularia* are nurse cells, the 15-18 centermost cells present a problem, although at least one of them has to be considered a primary oocyte. All previous investigators have considered the central cells to be oocytes; however, the existence of the intercellular bridges was previously unknown. Hargitt (1909) noted that here were two different types of oocyte nuclei in the gonophores of *T. crocea* and believed that
only the oocytes with a particular type of nucleus (the cells starting meiosis) enlarged and formed the egg. In both T. larynx and T. indivisa, the nuclei of the central cells show changes typical of meiosis I, and later all develop germinal vesicles which are identical except for slight differences in size. These germinal vesicles are distinctly different from the nuclei of the nurse cells. In addition, the oocytes produce large vesicles of yolk and fuse together at the completion of vitellogenesis, while the nurse cells contain only the smaller type of yolk vesicles and are engulfed and transformed into shrunken cells. Koch et al. (1967) reported that one of the two pro-oocytes that formed synaptinemal complexes reverted back and became a nurse cell in Drosophila. This does not appear to happen in Tubularia in which all 15-18 central cells are morphologically indistinguishable. Therefore, while only one of the germinal vesicles will persist in the fused primary oocyte, there are grounds for continuing to refer to all of the central cells as primary oocytes. An alternative would be to separate each complex into a single oocyte, about 15 large nurse cells, and about 1200 small nurse cells. Separating the cells of the complexes of Tubularia into three categories seems cumbersome and it is not entirely clear which oocyte will contribute the surviving germinal vesicle.

The oocyte-nurse cell complexes found in T. larynx and T. indivisa are similar to those reported by Pianka (1974) in the ctenophore, Bolinopsis, although there are some major differences. The complexes in Tubularia are over 10 times as large, each initially has more than 1 oocyte, and the oocytes are larger and have larger nuclei than the nurse cells. Attempts to resolve and map the interconnections in the
complexes of Tubularia have produced some confusing results. Most of the complexes contained oocytes with a maximum of 3 intercellular bridges on each cell and the largest germinal vesicle was in a cell at the center of the complex. However, some complexes were found to contain oocytes with 4 bridges and 2 germinal vesicles. These oocytes were probably produced by early fusion of the oocytes, which normally does not occur until the end of vitellogenesis when all of the oocytes fuse together. If two cells with 3 bridges apiece, including 1 common bridge, fuse together, the resulting cell will have 4 bridges and 2 nuclei. Unfortunately, some oocytes were seen that had 4 bridges and only a single nucleus, and the largest germinal vesicle was not always in a central cell of the complex. The largest complex that was mapped consisted of 43 cells and contained 3 oocytes with 4 bridges, but only 2 of these oocytes had 2 nuclei. The largest germinal vesicle in this complex was in one of the peripheral oocytes with 2 nuclei, although it was not the largest oocyte of the complex. The factors which determine the relative growth of the oocytes and their germinal vesicles and may also select the surviving germinal vesicle are not known. It is possible that all of the germinal vesicles are initially equivalent and that the one which will survive may be gradually determined during vitellogenesis.

The cooperative effort of the large number of cells of the oocyte-nurse cell complexes of Tubularia results in the production of large eggs which develop into actinulae. The actinula larva of the Tubulariidae is also found in other families of hydroids, including the Margelopsidae, Corymorphidae and Myriotheidae, as well as in other hydrozoans including the Trachymedusae, Narcomedusae, and Chondrophora
(Totton, 1954). Garstang (1946, p. 154) considers the actinula "merely a planula with precocious adult characteristics." Rees (1957) observed that an actinula is present in the more primitive families of hydroids and many authors, including Swedmark and Teissier (1966) and Rees (1966) have proposed that the ancestral hydroid was an actinula. While it is interesting to speculate about the evolutionary implications of nutritive oocytes, shrunken cells and actinulae and to make comparisons, many pieces of the puzzle are still missing. The reproductive biology of many hydroids is poorly known and few authors include sufficiently detailed information on both oogenesis and larval development.

Shrunken cells are well known in Hydra (Zihler, 1972) and have also been found in Acaulis primaris (Berrill, 1952). Hyman (1940) reported that the encysted larvae of Hydra hatch in a relatively advanced condition, with indications of tentacles, and Brinkman-Voss (1970) reported a similar process in Acaulis ilionae; however, these species do not develop an actinula. The larvae of Corymorpha palma (Torrey, 1907) and C. nutans (Rees, 1937) are also encysted and develop directly into polyps, without a free-swimming planula stage, but in these species there are no indications of shrunken cells. While all of the species whose eggs contain shrunken cells do not develop actinulae, the converse does appear to hold. According to Van De Vyver (1968a,b) the families of hydroids that develop an actinula larva have large eggs, which digest neighboring oocytes, and which accumulate shrunken cells. There are no reported exceptions to this generalization.

Hargitt (1919) found that most of the species that utilize large numbers of accessory cells are solitary species or colonial
species with large polyps, and the eggs produced are at least 200 μm in diameter. *Eudendrium ramosum* oocytes engulf large numbers of somatic accessory cells, while all of the other hydroids utilize germ line accessory cells (Hargitt, 1919). Rees (1957) reported a similar phenomenon and found that solitary species produced large eggs, while colonial species produced relatively small eggs. Other hydroids with large eggs and numerous germ line accessory cells, in addition to the species reported by Hargitt and Rees, include *Acaulis ilionae* which has 190 μm eggs (Brinkman-Voss, 1970), *Millepora* with 210-275 μm eggs (Magan, 1909), *Ectopleura dumortieri* with 350 μm eggs (Aurich and Werner, 1954), and *Acaulis primarius* with eggs 200-225 μm in diameter (Berrill, 1952). While the eggs of *Ectopleura dumortieri* are relatively large when compared to most hydroids, they are the smallest eggs of any hydroid reported to produce an actinula.

There are two major advantages to oogenesis in which many small nutritive oocytes or nurse cells support the growth of a relatively few definitive oocytes. The first is the relatively large surface area to volume ratio of many cells which increases the amount of surface area available for food absorption as compared to a single large cell. There is also the tremendous increase in the amount of DNA available to code for ribosomal RNA. Each accessory cell has at least the normal 2C DNA content and if they are germ line accessory cells that have started meiosis their DNA content will be at least 4C. The nutritive oocytes and nurse cells effectively increase the rate of synthesis of ribosomes for the definitive oocytes since each cell has at least one nucleolus where ribosomal RNA production occurs. The net result is the accelerated development of large eggs reported by Van de Vyver (1968a)
in the families of hydroids with actinulae.

Campbell (1974) reported that gametogenesis in the Cnidaria usually does not involve accessory cells. However, regardless of the type of oocyte nutrition, the eggs of hydroids with sessile gonophores are usually fertilized internally and develop in or on the gonophores. The species of hydroids with small eggs produce many of them in each gonophore and brood them to a planula before release, while the species that produce relatively few, large eggs in each gonophore brood them to more advanced stages. According to Pearse (1979) the brooding species of chitons are often relatively small compared to the non-brooders. Chia (1974) attributes brooding by small species of invertebrates to energy limitations. In general, brooding requires a greater investment in stored nutrients for each embryo than for planktotrophic species in which the egg is fertilized after being released by the parent and development is external. However, brooding increases the larval survival rate by decreasing the time a planktonic larva is exposed to filter feeders. Brooding species therefore have to release many fewer larvae than planktotrophic species to produce the same number of larvae that survive to settle and metamorphose. This may reduce the total yolk requirement during a reproductive period. The actinulae of Tubularia may have a very short planktonic existence and often settle close to the colony that released them (Pyefinch and Downing, 1949).

An alternative view of brooding is provided by Menge (1975) who suggests that internal space limitations prevent small species from producing and broadcasting the necessarily large numbers of planktotropic larvae and that they therefore concentrate on producing a
relatively small number of lecithotrophic larvae with a large yolk supply. Hydroids appear to do the opposite. All of the large solitary species produce relatively large eggs and many brood them to an actinula larva before release. The colonial species with small polyps produce relatively small eggs while *Pennaria tiarella*, a colonial species with large polyps, broods large eggs which are produced by the utilization of many nutritive ooctyes. In general, among hydroids, the fewest eggs per gonophore and the largest eggs are produced by the largest species. Rees (1957, p. 481) may have provided the answer in his suggestion that the large eggs of the solitary hydroids "may only reflect the minimum size at which such a polyp can become self supporting and at the same time be a miniature of the adult." This type of argument may also be applied to the eggs that develop into actinulae. The eggs of *Ectopleura dumortieri*, which are 360 μm in diameter, may represent the minimum size at which an egg contains enough material to form an actinula. The eggs of *Hydra* are only slightly smaller, 340 μm according to Hargitt (1919), and produce an advanced, encysted larva, but do not form an actinula.

Whatever the advantages that are involved in oogenesis when hundreds or thousands of germ line accessory cells support the production of a large egg, it is apparently a method utilized by many species of athecate hydroids. While most of these species are capitate hydroids, at least one species, *Rhizogeton fusiformis*, is in the suborder filifera. These species may be separated into two general groups depending on the presence or absence of shrunken cells in the eggs. During this investigation, intercellular bridges and nurse cells were discovered in *T. larynx*, *T. indivisa*, and *Acaulias primaria*, all
species which produce shrunken cells. The reports of bridges in T. corocea by Boelsterli (1975) and in Hydra by Tardent (1974), both species that produce shrunken cells, makes it likely that the accessory cells in these species are also nurse cells. On the basis of this information, it is also possible to make two other, more general, predictions. The first of these is that the shrunken cells of other species, including Eotopleura and Hyboecodon, will also prove to be nurse cells and that intercellular bridges will be found among the germ cells of these species. The second prediction, which is simply the converse of the first, is that no bridges will be found among the germ cells of species that lack shrunken cells and that the accessory cells of these species are nutritive oocytes.
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