

As can be seen from Table 1, the urine osmolality in starving leeches was very low (27 mOsm/kg H<sub>2</sub>O). The osmolality along the nephridium decreased continuously in the direction of the flow. It seemed that all parts of the nephridium contributed to the dilution of the urine. In leeches that were fed the osmotic concentration decreased only slightly along the entire nephridium and was only 27% lower in urine than in hemolymph. Thus, the resorption of solute along the nephridium appears to be regulated according to the need for electrolyte conservation.

Table 1

HEMOLYMPH, NEPHRIDIAL AND URINE OSMOLALITY mOsm/kg H<sub>2</sub>O  
IN STARVED AND FED LEECHES

	Hemolymph	I	II	III	Urine
Starved	(6) 224 ± 9	(5) 182 ± 9	(7) 150 ± 7	(2) 70 ± 5	(8) 27 ± 7
Fed	(3) 282 ± 16	(3) 262 ± 17	(3) 224 ± 8	(3) 189 ± 10	(3) 182 ± 2

Preliminary studies with <sup>14</sup>C inulin showed an inulin urine/hemolymph concentration ratio of unity. This would indicate that the urine is formed by filtration rather than secretion.

Further studies will include the determination of Na and K concentrations in the nephridial fluid.

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LIGHT AND ELECTRON MICROSCOPY OF REGENERATING Tubularia

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Complete understanding of the process of regeneration depends partly upon detailed descriptions of the morphological changes which occur. Descriptions of regeneration in the marine coelenterate Tubularia and other hydroids are of special value for several reasons. Hydroids are relatively simple in structure. Their tissue layers possess well-differentiated cell types. They undergo complex structural changes during regeneration. They are small (1 mm in diameter) and almost the entire cross section of a specimen can be seen in a low power electron micrograph. Unlike most other hydroids, which contract when they are cut, Tubularia is covered by a stiff perisarc which prevents contraction and the resulting alteration of relationships. The purpose of this work is to prepare a detailed description, at the ultrastructural level, of mouth and tentacle regeneration from isolated stems of Tubularia.

The summer was spent in preparing and embedding specimens of Tubularia in various stages of regeneration. These specimens will be sectioned for study with the electron microscope and for correlative study with the light microscope. Special attention will be given to the following stages: immediate tissue damage, healing, distal cell migration, tentacle formation and final emergence of the regenerate.