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Assessment of Cranial Neural Crest Proliferation Patterns between the Redeye Tetra *Moenkhausia Sanctaefilomenae* and the Zebrafish *Danio Rerio*

Lesly Ingram Illinois Wesleyan University

Brian Walter, Faculty Advisor Illinois Wesleyan University

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Assessment of cranial neural crest proliferation patterns between the redeve tetra Moenkhausia sanctaefilomenae and the zebrafish Danio rerio Lesly J. Ingram and Brian E. Walter* LINOIS WESLEYAN **Biology Department, Illinois Wesleyan University, Bloomington, IL**

Introduction and Background

fishes Teleost display remarkable amount Of morphological diversity. In this study we examined the development of two species of freshwater fishes, the redeve

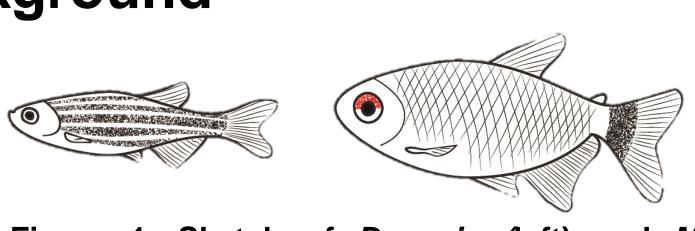


Figure 1. Sketch of *D. rerio* (left) and *M.* sanctaefilomenae.

tetra Moenkhausia sanctaefilomenae and the zebrafish Danio rerio (Figure 1). Notably, the larva of *M. sanctaefilomenae* exhibit larger jaw cartilages compared to *D. rerio* (Figure 2). Consequently, our research sought to elucidate the cellular mechanisms responsible for the morphological variations observed in the ventral arch derivatives between M. sanctaefilomenae and D. rerio.

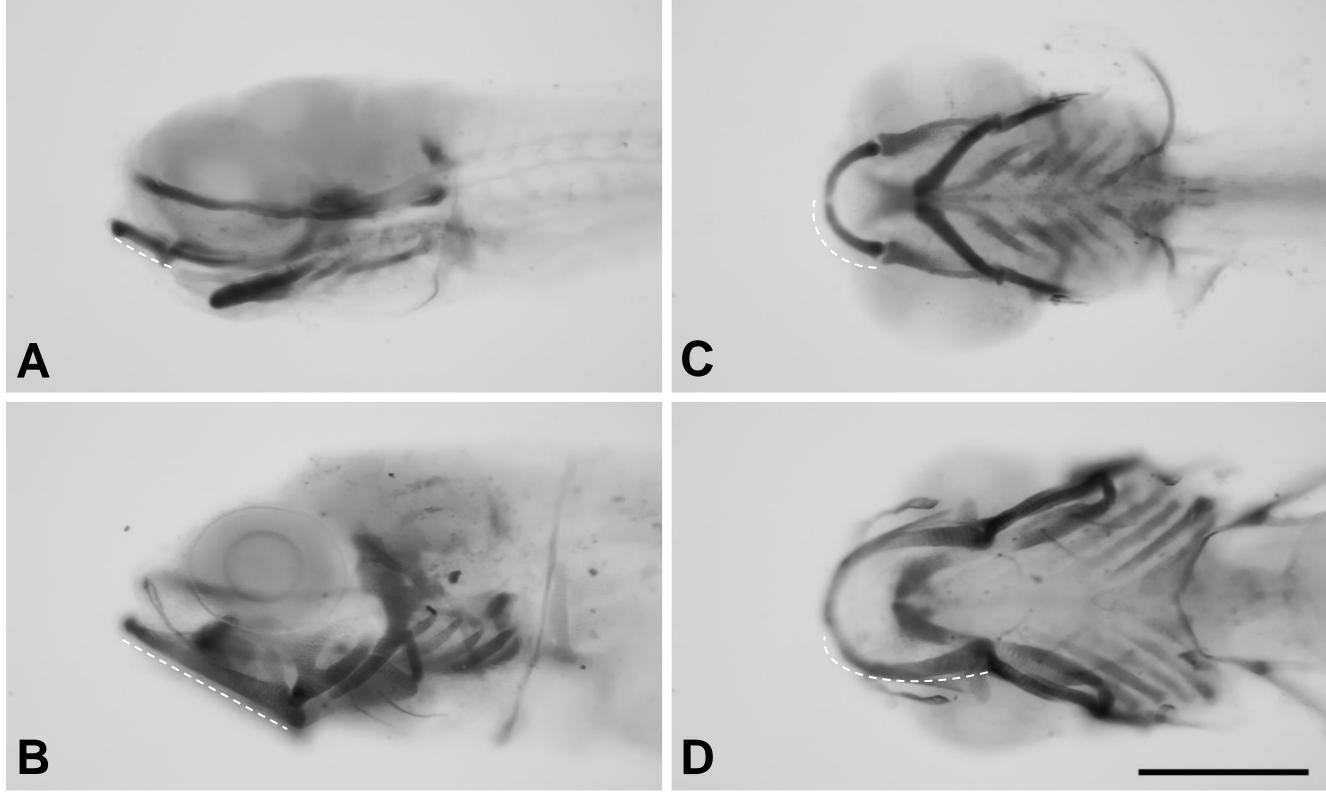


Figure 2. Developing skeletal elements of *D. rerio* (top) and *M. sanctaefilomenae* (bottom). Panels A-B depict lateral views, while panels C-D depict ventral views of the developing cartilages for each species at comparable stages. Scale bar = 0.5 mm and applies to all panels.

A specific population of cells, known as neural crest, gives rise to the jaw elements known as the Meckel's and ceratohyal cartilages. These jaw cartilages are components of the splanchnocranium and are located within the pharyngeal arches. A sequential series of developmental phenomena are known to produce these cartilages (Figure 3). Although the developmental timeframes vary for each species – and were taken into account during experimentation – the general process (proliferation, histogenesis, shaping, and growth) remains consistent.

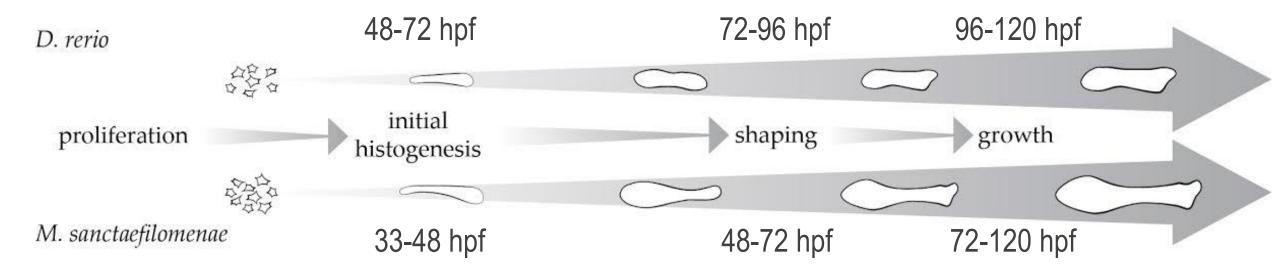


Figure 3. Developmental timeframes of skeletal element formation between two species.

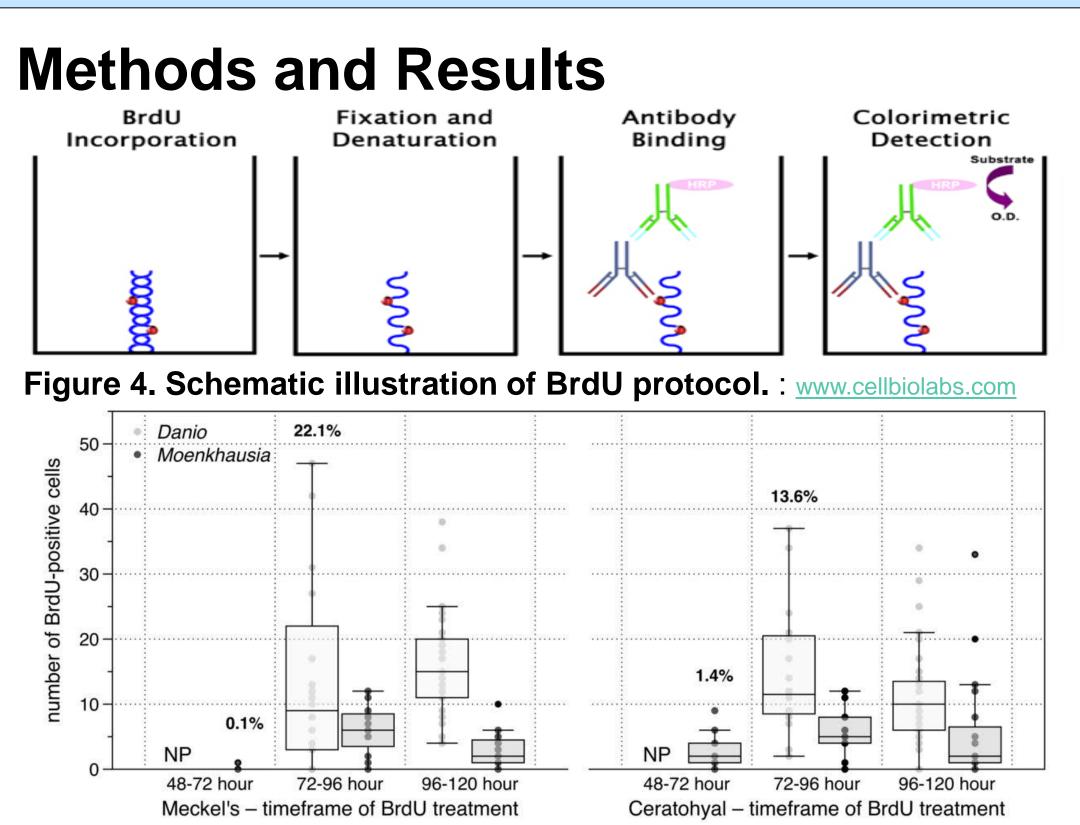


Figure 6. Numerical counts of BrdU-positive cells in the ventral arch derivatives between two species at specified time points. The percentages refer to the ratio of the average number of BrdU-positive cells to the average total number of chondrocytes within the element at the indicated timeframe.

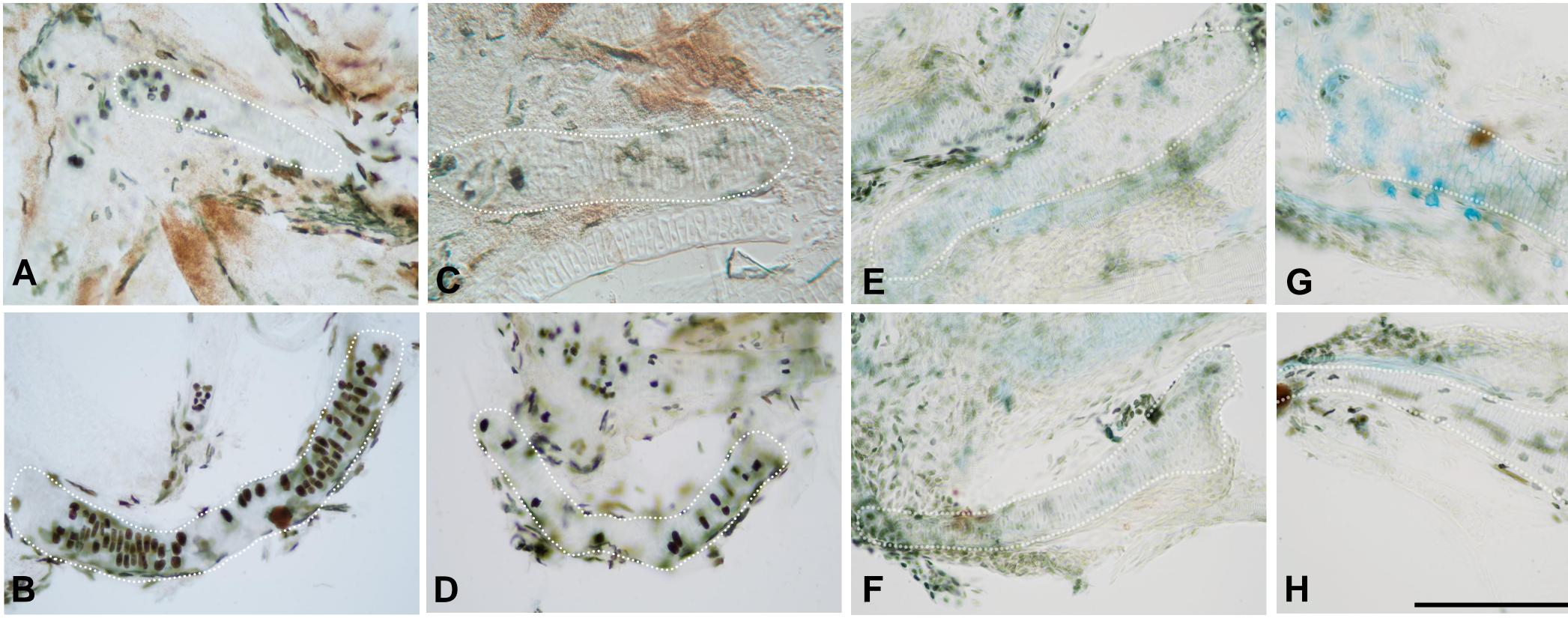


Figure 5. Comparison between BrdU-treated skeletal elements of *M. sanctaefilomenae* and *D. rerio* during various developmental timeframes. Top panels depict the ceratohyal cartilages while the bottom panels depict Meckel's cartilage Panels A – D depict elements from D. rerio. A-B correspond to the period of shaping and C-D correspond to the period of growth. Panels E-H depict elements from M. sanctaefilomenae. E-F correspond to the period of shaping and G-H correspond to the period of growth. Scale bar = 200 μ m.

Conclusions and Future work

Our research has demonstrated that there is a distinct difference in the mechanisms used for cartilage formation between the two species. Specifically, it appears that the ventral arch derivatives in *M. sanctaefilomenae* develop from an initial pool of chondroblasts that originate prior to cartilage formation, and thus exhibit limited reliance on proliferation during histogenesis, shaping, and growth. However, the ventral arch derivatives in *D. rerio* display an opposite mechanism whereby proliferation is the primary means for development during the shaping and growth phases. Further experimentation could incorporate another closely related species to determine the prevalence of these mechanisms.



To visualize proliferation patterns in the ventral arch derivatives, BrdU staining techniques were utilized at different developmental timeframes for each species. Such treatment involved BrdU (a thymidine analog) being incorporated into DNA. Figure 4 presents an overview of the detection procedure.

When comparing the specimens following BrdU treatment, it was found that both the ceratohyal and Meckel's cartilages of D. rerio displayed extensive staining during the shaping and growth periods (Figure 5, A-D). On the other hand, the ceratohyal and Meckel's cartilage of *M. sanctaefilomenae* exhibited very little to no staining during the shaping period (Figure 5, E-F) and slightly increased staining during the growth period (Figure 5, G-H). Such observations imply that proliferation is not of great significance for M. sanctaefilomenae during shaping, but does become important during the growth period. Meanwhile, it seems that proliferation is vital for D. rerio during shaping and equally important during growth. These findings are consistent with the numerical counts of BrdU-positive cells in the Meckel's and ceratohyal cartilages of both species (Figure 6).

