SHORT COMMUNICATION

SARCOMERE LENGTH CHANGES DURING FISH SWIMMING

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The first noninvasive measurements of muscle sarcomere length were made by
laser diffraction during swimming in the glass catfish (Kryptopterus bicirri). Sarcomere length changes were recorded at a relatively constant swimming speed (approximately 3.2 body lengths s\(^{-1}\), tailbeat frequency 5–7 beats s\(^{-1}\)) just dorsal to the vertebral column. 18 s of continuous diffraction data were obtained and yielded cyclic sarcomere length changes of only 65±10 nm. Based on published fish morphology, architecture and behaviour, we suggest that these values represent white muscle cyclic sarcomere shortening and elongation during swimming powered by red muscle.

The interaction between actin and myosin represents the fundamental force-producing mechanism in all skeletal muscles. Interaction between analogous proteins also occurs within cells, resulting in movement of subcellular organelles. Although considerable detail is known about the muscle sarcomere length-tension relationship and how isometric force potential is a function of myofilament overlap (Gordon et al. 1966), there are no direct measurements of myofilament movement during locomotion. We have taken advantage of the transparent morphology of the glass catfish (Kryptopterus bicirri, Fig. 1) to investigate noninvasively sarcomere length changes which occur during free swimming.

An acrylic ‘fish mill’ was designed allowing fish to swim against a regulated water flow (average speed 3.2 body lengths s\(^{-1}\)) while remaining stationary relative to a laser beam (Fig. 2). (At higher flow rates, fish maintained their position by abrupt changes in swimming speed, suggesting the recruitment of white muscle. Thus, for the purposes of this investigation, we assume that the locomotion was powered by the red muscle.) The sides of the chamber were constructed of flat glass that permitted transillumination of the fish during swimming by a helium–neon laser beam. The resultant diffraction pattern was imaged onto a translucent screen and video taped at 60 Hz. Off-line, the position of the first-order diffraction line was measured and sarcomere length calculated according to the grating equation:

Key words: sarcomere, locomotion, actin, myosin, myofilament, Kryptopterus bicirri.
Fig. 2. Experimental apparatus used in this study. The fish is placed in the acrylic chamber and transilluminated by a HeNe laser. The resultant diffraction pattern is composed of different orders (0, ±1, ±2, ...). The angular position of a diffracted order is recorded and sarcomere length is calculated according to the grating equation: 

\[ n\lambda = dsin\theta, \]

where \( n \) is the diffraction order number (1 in this experiment), \( \lambda \) is the laser wavelength (0.632 \( \mu \)m), \( \theta \) is the diffraction angle and \( d \) is the sarcomere length (Cleworth and Edman, 1972; Kawai and Kunz, 1973; Yeh et al. 1980). Measurement resolution was approximately 5nm, while accuracy was 18nm determined using diffraction gratings of known spacing placed into the mill. Even though fish often moved slightly relative to the laser beam during swimming, it was possible to obtain uninterrupted sarcomere length records lasting several seconds from a single region of the fish. Generally, the fish were transilluminated near the axis of the vertebral column. From nine separate records obtained from eight different fish, 18s (range 1.5–5.2s) of sarcomere length data from single regions were recorded successfully. In addition to sarcomere lengths from live fish, sarcomere length and fibre pennation angle distribution were measured from muscles embedded in plastic by scanning plastic sections with a laser beam.
Fig. 1. Photograph of the glass catfish (*Kryptopterus bicirri*) illuminated by white light. The rainbow pattern on the fish represents diffraction of the white light by the sarcomeres within the fish musculature. Scale bar, 1 cm.
Sarcomere changes during swimming

Fig. 3. Sarcomere length during 4 s of uninterrupted measurement. Each data point represents a single diffraction pattern analysis. Sarcomere length change frequency was 4.19±0.04 Hz, while tailbeat frequency was 4.20±0.03 Hz. Average sarcomere excursion was 76±4 nm.

Sarcomere length excursion during swimming was always less than 0.1 μm (Fig. 3). Assuming that the actin and myosin filament lengths were identical to those found by Sosnicki et al. (1991) in carp (0.99 μm and 1.54 μm respectively) and thus that the maximum sarcomere length was 3.52 μm while minimum sarcomere length was 1.54 μm, sarcomeres shortened by only 3.3% of their potential range. Average sarcomere excursion was 65±10 nm (mean±s.d., N=9). Average minimum sarcomere length (1.829±0.021 μm; mean±s.e., N=9) was significantly less (P<0.001) than average maximum sarcomere length (1.894±0.023 μm). In three fish, sarcomere length change was measured along with tailbeat frequency. These tailbeat frequencies were not significantly different from sarcomere length change frequencies (range 4–10 Hz, mean difference=0.03±0.06 Hz, P>0.8).

The relatively small sarcomere length change reported here is consistent with that provided by Rome and Sosnicki (1991) if one assumes that measurements were obtained from the white muscle during swimming powered by red muscle. Since sarcomere length changes for red muscle are four times those of white muscle for a given spinal curvature, much larger sarcomere length changes would have been expected if records had been obtained from active red muscle (Alexander, 1969; Rome et al. 1988).

We must be cautious in interpreting these data for two reasons. First, the
muscles from which sarcomere length was measured have not been definitely identified. The argument presented here is based on a synthesis of available information from the literature. It is possible that active white muscle was being recorded. Future experiments will be required to elucidate this phenomenon further. Second, there is some uncertainty as to the nature of laser light diffraction from the fish itself. The laser traverses two halves of the fish. It is, therefore, not clear precisely why the diffraction pattern obtained appears to represent only one side of the fish. Our working hypothesis is that the diffraction pattern created by the muscle closest to the laser beam is then diffracted again by the second half of the fish at relatively high angles. Empirical support for this hypothesis was obtained by diffracting the laser beam using two blocks of fish muscle embedded in plastic, one of which was fixed at a long sarcomere length (2.1 \( \mu \)m) and one at a short sarcomere length (1.9 \( \mu \)m). When the position of these blocks was interchanged, the sarcomere length measured was that of the muscle block closest to the screen, including its fine structure. Average sarcomere length measured with the 'shorter' block closest to the screen was 1.93±0.04 \( \mu \)m, which was significantly less than that measured with the 'longer' block closest to the screen (2.15±0.05 \( \mu \)m, \( P<0.01 \)). A second argument against multiple diffraction is that the relative ratios of the zero: first: second order were found to be 1.0:0.05:0.01, which is similar to what would be expected from a thin grating. If significant multiple diffraction effects occurred, we would expect to see intensity 'dumping' to the higher orders, as has been demonstrated experimentally by Magnusson and Gaylord (1977).

References


