Actin in the Photoreceptor Connecting Cilium: Immunocytochemical Localization to the Site of Outer Segment Disk Formation

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ABSTRACT  Actin has been localized in Rana pipiens retinas that were fixed and embedded in aldehyde cross-linked BSA. Thin sections were reacted sequentially with (a) affinity-purified antiactin antibodies induced in rabbits; (b) biotinyl-sheep anti-rabbit antibodies; and (c) avidin-ferritin conjugates. As expected, antiactin labeling density was high in the apical pigment epithelial cell processes and in the calycal processes of photoreceptors. Actin was also localized in a new site. The connecting cilium that joins the inner and outer segments of both rods and cones was heavily labeled by antiactin at its outer segment (OS), or distal, end. In this region of the cilium, the plasma membrane evaginates to form new OS disks and these basal disks were labeled in some instances. Below the new disks in rods, the cytoplasm of liplike expansions of the distal cilium was also heavily labeled. The plasma membrane and interior of the connecting cilium and the remainder of the OS were unlabeled. These findings suggest that actin may participate in the vectorial transport of opsin and other intrinsic membrane proteins that are incorporated into newly forming OS disks. The results also implicate actin in the membrane expansion involved with OS disk formation.

Retinal rod and cone photoreceptors are highly polarized cells in which the light sensitive outer segment (OS) is connected to the remainder of the cell by a nonmotile cilium (Fig. 1). The OS of rods is comprised of a plasma membrane enclosed stack of hundreds of membranous disks that contain the visual pigment rhodopsin as an integral membrane protein. Opsin, the apoprotein of rhodopsin, is synthesized in the inner segment of the cell (1–4) and is membrane bound in vesicles or cisternae as it is vectorially transported to the base of the connecting cilium (CC; 3, 5). It has been suggested that these vesicles fuse with the apical plasma membrane of the rod inner segment (5, 6), possibly in a recently described surface domain known as the periciliary ridge complex (PRC; 7) (Fig. 2). Opsin may migrate from the PRC along the CC plasma membrane to its site of incorporation into OS disks. A similar pathway is postulated for a high molecular weight intrinsic membrane protein recently localized to the margins and incisures of OS disks (8–10).

The OS undergoes a unique form of membrane renewal. At the base of the OS the CC plasma membrane evaginates to form flattened lamellae that expand in size to the full diameter of the OS (7, 11) (Fig. 2). In cones these lamellae remain open to the extracellular space, whereas in rods they separate from the OS plasma membrane and become isolated disks. In rods, as newer disks are formed the older ones are gradually displaced toward the apex of the OS where they are intermittently shed in packets that are ingested by adjacent pigment epithelial (PE) cells (12, 13) (Fig. 1). Rods shed the major portion of their OS disks following light onset (14, 15). Cones also shed their OS material but the timing of this event varies in different species (16, 17).

The PE cells extend long, thin, cytoplasmic processes that interdigitate with and ensheath the OS (Fig. 1). These processes play a role in the phagocytosis of shed OS material and contain parallel arrays or bundles of axially oriented actin filaments (18–20). Actin filament bundles are also observed in photoreceptor calycal processes (21). These are microvillus-like projections that arise from the inner segment and extend alongside the basal OS (Figs. 1 and 2).

A relatively constant OS length is maintained by a balance
between the processes of disk formation and disk shedding. Although much information has been gathered about the effects of light and temperature on these two processes (for review see reference 22), the molecular events that underlie OS renewal are still not understood. A contractile mechanism may mediate the vectorial transport of opsins to the OS, as well as the membrane expansion involved in OS disk formation. To examine this hypothesis, we conducted an immunocytochemical study in which thin sections of albumin embedded retinas were labeled with antiactin antibodies. As expected, heavy labeling of actin occurred in the PE processes and calycal processes. The presence of actin was also demonstrated in a new site. A dense, localized concentration of actin was observed in the distal portion of the CC at the site of OS disk formation in both rod and cone photoreceptors.

**MATERIALS AND METHODS**

**Animals:** Frogs (*Rana pipiens* adults) were kept on a 12-h light/12-h dark cycle and sacrificed at various times of the day and night. For all samples taken at night, the frogs were killed and the eyes dissected under dim red light. One of the frogs was exposed to the natural midautumn diurnal cycle and was killed 1 h after the onset of light in the morning.

**Tissue Processing:** Animals were decapitated, the eyes removed, and the eyecups fixed at room temperature either in 2% paraformaldehyde, 2.5% glutaraldehyde, 0.1 M sodium cacodylate, pH 7.4, for 2 or 3.5 h, or in 1.2% glutaraldehyde, 0.15 M sodium phosphate, pH 7.0 for 2 h. The eyecups were rinsed in cacodylate or phosphate buffer, cut into strips and embedded in formaldehyde-or glutaraldehyde-crosslinked BSA (Armour Reheis, fraction V) as described previously (4, 23-25).

**Preparation of Antibodies:** Antiactin antibodies were induced in rabbits using actin purified from chicken gizzards (26). The immunization schedule followed was that described by Jockusch et al. (27) and the antiactin was purified from immune serum using affinity chromatography. For this study the actin used to prepare an actin-Sepharose column was purified as follows. An acetone powder was prepared from fresh chicken gizzards as outlined by Herman and Pollard (28). Actin was extracted following the method of Spudich and Watt (29), with modifications (26). The depolymerized actin was passed over a Sephadex G-150 column equilibrated in the actin depolymerizing buffer. The peak actin containing fractions were pooled, concentrated, and bound to CNBr-activated Sepharose 4B (Pharmacia Fine Chemicals, Inc.) as previously described (26, 27).

Immune serum was mixed with the actin-Sepharose beads. The bound antiactin antibodies were eluted with 4 M M9C12 and dialyzed against borate buffer, pH 8.0 containing 0.02% sodium azide. The antiactinantibodies were concentrated to 0.15 mg/ml and BSA (CalBiochem-Behring Corp., San Diego, CA, fraction V) was added to 1%. For immunocytochemistry, antiactin was used at this concentration or diluted to a concentration of 0.10 or 0.05 mg/ml.

Biotinyl IgG or Fab(Fab')2 of sheep anti-rabbit Fab(Fab')2 was prepared by reaction with biotin N-hydroxysuccinimide ester (30). Six times crystallized horse spleen ferritin (Miles Laboratories, Inc., Elkhart, IN), with no further purification, was conjugated to avidin (Sigma Chemical Co., St. Louis, MO, # A-9390) following the glutaraldehyde conjugation method (31).

Retinal thin sections were collected on carbon and formvar coated copper grids and were stored dry until use. At room temperature, grids were transferred sequentially to drops of 0.1 M Tris-Cl, pH 7.4 (5 min); 4% BSA in Tris-Cl (10
Diagram showing the connecting cilium and junctional region between the inner and outer segments of a rod. Outer segment disks develop from evaginations of the cilium plasma membrane. In rods these lamellae expand to the full diameter of the outer segment and then separate from the plasma membrane to form isolated disks. During this renewal process, what had been the space between two expanding lamellae becomes the intradiscal space of an isolated disk. A liplike structure protrudes from the distal cilium just below the developing disks. Note the difference between a calycal process and the lateral inner segment process, which is a ridge of the periciliary ridge complex.

In control experiments nonimmune rabbit IgG was substituted for the antiactin in the labeling procedure. All tissue sections were examined in a Philips 300 electron microscope operated at 60 KeV.

Morphometric Analysis:  For the quantitation of antiactin antibody labeling, micrographs were taken at a magnification of x17,000 of different areas of photoreceptor and PE cells. Contact positives on the same film were made and viewed on a Bellco Plaque Viewer (Bellco Glass, Inc., Vineland, NJ). A template consisting of a double lattice grid system was placed on the screen. Ferritin grains per square were counted over areas of interest and the mean grain counts per square micron (± SE) were calculated as previously described (25).

RESULTS
The affinity-purified antiactin antibodies had previously been used in an immunofluorescence study with cultured rat PE cells (26). The antiactin antibodies labeled stress fibers in these cultured cells, as well as actin containing feltworks associated with newly phagocytized rod outer segment (ROS) fragments. Similar patterns of actin distribution have been reported for other types of cultured cells. In the present study...
we used these antiactin antibodies to label actin in thin sections of albumin-embedded frog retinas. The antiactin heavily labeled PE processes that surround ROS and cone outer segments (COS) (Fig. 3). Label was also concentrated in the apical region of the PE adjacent to the ROS tips (Fig. 4). Newly formed phagosomes that contained shed ROS material were surrounded by actin in the apical PE, whereas phagosomes that had moved deeper into the PE cell body were no longer surrounded by actin (Fig. 4). Antiactin did not specifically label either ROS or COS (Figs. 3 and 4, Table 1).

Antiactin also heavily labeled photoreceptor calycal processes which are long tapered projections arising from the inner segments and extending alongside the basal portions of the OS (Fig. 5a). The label extended from the calycal processes deep into the inner segment, subjacent to the plasma membrane. In cross-sectioned rods, labeled calycal processes were observed to lie along the indentations of the ROS (Fig. 5b). Labeled actin bundles of the calycal processes continued into the ellipsoid (mitochondria rich region) of the inner segment (Fig. 5c). In the lower ellipsoid, near the junction with the myoid, these bundles separated into a more uniform distribution of actin lying just beneath the plasma membrane (Fig. 5d). The inner segment of frog photoreceptors was not specifically labeled aside from the actin bundles which extended from the calycal processes (Fig. 5, Table I).

Actin was also localized to a new site in rods and cones. Antiactin label was heavily concentrated in the distal portion of the rod CC and in the interior of a lip-like structure that protrudes from the rod CC just below the basal ROS disks (Fig. 6a). In this region the CC plasma membrane evaginates to form new ROS disks. The plasma membrane of the lip was unlabeled. When the plane of section was rotated such that the CC appeared to be in the middle of the cell, more than one labeled lip could be viewed (Fig. 6, b and c). In cones, antiactin was also concentrated in the distal portion of the CC (Fig. 6d) at the site where new COS disks are formed, however lip-like expansions of the CC were not observed. In cross-section, the proximal portion of the CC below the actin-rich domain was observed to be free of label (Fig. 7). Antiactin label occasionally extended from the distal portion of the CC along the length of the basal OS disks in both rods and cones (Fig. 8). The OS itself was otherwise unlabeled. In both rods and cones, antiactin did not specifically label the interior of the CC or its plasma membrane, nor did it label the basal body or the periciliary ridge complex (Fig. 6–8, Table I).

As a control, thin sections of albumin embedded retinas were labeled with nonimmune rabbit IgG. No specific labeling of the PE processes, the calycal processes, the CC, or the photoreceptor inner and outer segments occurred (Fig. 9).

**DISCUSSION**

Actin filaments have previously been identified in the apical PE and PE processes by using heavy meromyosin or myosin subfragment-1 binding to filaments exposed in glycerinated retinas (18–20, 33). In one immunofluorescence study (34) the PE processes in frozen sections of retina became labeled with antiactin antibodies. These actin filaments may mediate the movement of melanin granules in response to changes in the light-dark cycle (18, 20, 33) and they may participate in the phagocytosis of shed packets of OS disks (20). In the present study, when thin sections of albumin embedded retinas were labeled with antiactin antibodies, the apical PE and PE processes were densely labeled (Figs. 3, and 4). These results confirm those of the previous studies, and serve as a positive control for both the specificity and reactivity of the antiactin antibodies used in these experiments. The reactive
FIGURE 5 Antiactin antibody labeling of actin bundles within calycal processes of rod photoreceptors. (a) Photoreceptor showing the basal portion of a rod outer segment (ROS) and the ellipsoid (mitochondria rich) region of the rod inner segment (RIS). Antiactin densely labels the actin bundles within a calycal process (arrows). This label continues along the bundles as they extend into the RIS, subjacent to the plasma membrane (double arrows). ROS and RIS lack specific antiactin label. The arrowhead indicates the level of the ellipsoid-myoid junction in the RIS. (Inset) Higher magnification of the apical end of the calycal process (CP). (b) Cross-sectioned ROS showing antiactin labeled calycal processes that lie along indentations of the ROS plasma membrane. (c) Cross-sectioned upper ellipsoid region of the RIS showing discrete labeled actin bundles that extend from the calycal processes. (d) Cross-sectioned lower ellipsoid, near junction with the myoid region of the RIS. Antiactin labels actin which is now more circumferentially distributed beneath the lateral plasma membrane. Bars, 0.5 μm (a); 0.25 μm (b–d, and inset); × 18,480 (a); × 42,500 (inset); × 49,300 (b); × 60,420 (c); × 56,525 (d).
FIGURE 6 Longitudinal sections of photoreceptor connecting cilia (CC) labeled with antiactin antibodies. (a) This rod CC is heavily labeled by antiactin in the distal portion of the CC and in the interior of a lip-like structure (L) that protrudes from the CC just below the basal rod outer segment (ROS) disks. Actin in a calycal process (CP) of a neighboring photoreceptor is also labeled. A lateral inner segment process adjacent to the CC in this plane of section (arrow) is always observed to be free of antiactin label. This process is a ridge of the periciliary ridge complex (PRC). BB, basal body; RIS, rod inner segment. (b) The plane of section in this image appears to be rotated around the CC so that more than one lip (L) is observed to be labeled by antiactin. A calycal process (CP) is also labeled. (c) When the plane of section is rotated approximately 90° from that in Fig. 6a, the CC appears to lie in the middle of the cell and two antiactin labeled lips (L) are observed. Since this section is slightly oblique to the long axis of the CC, a gap appears between the lips and the ROS. (d) Longitudinal section through a cone CC. Antiactin antibodies heavily label the distal portion of the CC (arrow), and in this cone the label extends into the adjacent portions of the most basal outer segment lamellae (double arrows). CIS, cone inner segment; COS, cone outer segment. Bars, 0.1 μm. × 68,000 (a); × 95,040 (b); × 84,480 (c); × 105,600 (d).
sites in PE cells also indicate antigen preservation and exposure in these tissue sections.

Adjacent to ROS tips, recently formed phagosomes were situated within antiactin-labeled regions of the apical PE (Fig. 4). As phagosomes moved deeper into the PE cell body the surrounding actin was lost. In recent studies of the phagocytosis of ROS fragments by cultured rat PE cells (26), ROS that attached to the PE cells were shown to be ingested by actin-containing pseudopods that extended from the PE cell surface to surround the ROS. Actin feltworks surrounded newly formed phagosomes, but the feltwork was lost as a phagosome moved deeper into the PE cell body. Thus our in vivo study in frog retinas and the in vitro studies of cultured rat PE cells indicate that a similar sequence of phagocytic events occurs. This result is important for evaluating the use of PE cell cultures as models of PE cell function.

 Bundles of actin filaments are also found in the calycal processes of photoreceptors. In teleost cones these filament bundles extend deep into the inner segment, subjacent to the plasma membrane (21). They are believed to take part in a light-induced actomyosin mediated contraction of teleost cones (35). Retinomotor movement also occurs in frog photoreceptors. Our results indicate that actin in the calycal processes of Rana pipiens photoreceptors is highly reactive with antiactin antibodies and that the label extends into the depths of the inner segment (Fig. 5) in a distribution similar to that observed in teleost cones. These actin filaments might also function in rod and cone contraction in the frog.

Antiactin did not specifically label photoreceptor outer segments in the present study. However, recent studies of freeze-fractured and deep-etched, rapidly frozen ROS demonstrated filament-like structures on the rims of ROS disks. These filaments connected the rims of adjacent disks and also spanned the deep incisures of amphibian ROS to connect adjacent rims from the same disk (36, 37). Other filaments connected the disk rims to the ROS plasma membrane. The absence of specific antiactin labeling in ROS or COS suggests that these filaments do not contain actin.

At the base of both ROS and COS, in the distal portion of the CC, actin was detected with a high labeling density (Fig. 6, Table I). New disk formation is initiated in this region by an evagination of the CC plasma membrane (11). Actin was also highly concentrated in the cytoplasm of lip-like structures that protrude from the CC just below the basal disks of rods. The plasma membrane of these lips was unlabeled by antiactin antibodies. More than one lip had previously been observed in frog rods using high resolution scanning electron microscopy (7). Lips appear not to be primordial disks (7) and at the present time their function is unknown. Thus actin could be part of a cytoskeleton that maintains lip structure. The concentration of actin in such a small domain in the distal CC and lips indicates that actin may be involved in the final stages of the transport and processing of opsin and other intrinsic membrane proteins that are destined for the newly forming OS disks. As pointed out by Peters et al. (7), the simple lateral diffusion of opsin from the periciliary ridge complex along the CC plasma membrane to the ROS would require a downhill gradient. However, the ROS plasma membrane has a high concentration of opsin. Thus actin in the

| Tissue Site               | Ferritin grains/square micron ± SE | Tissue site               | Ferritin grains/square micron ± SE |
|---------------------------|-----------------------------------|---------------------------|-----------------------------------|
| Rod lip (11)              | 1,324 ± 70                        | Rod outer segment (21)    | 20 ± 3                            |
| Distal rod cilium (5)     | 1,349 ± 216                       | Distal inner segment (13) | 25 ± 2                            |
| Distal cone cilium (5)    | 1,434 ± 134                       | Cone outer segment (5)    | 12 ± 5                            |
| Rod calycal process (9)   | 1,825 ± 141                       | Rod inner segment (5)     | 16 ± 7                            |
| Cone calycal process (3)  | 1,778 ± 314                       | Rod basal body (9)        | 40 ± 13                           |
| Pigment epithelial process (3) | 3,012 ± 386                         | Cone basal body (5)       | 38 ± 26                           |

Values in parentheses indicate the number of cells counted for each tissue site. Ferritin grain counts for the lips, distal cilia, and inner and outer segments were taken from longitudinally or obliquely cut photoreceptors. Counts for the calycal processes and pigment epithelial processes were taken from longitudinally cut cells only. Counts for the basal bodies were taken from those that were transversely cut as shown in Fig. 6.
determined how a local concentration of actin is restricted to receptor CC, and its possible roles in the vectorial transport communication, Dr. Joseph C. Besharse, Dept. of Anatomy, Emory University, Atlanta, Georgia). In control retinas, such open basal disks have been used to assay new disk assembly (38).

The localization of actin to the distal portion of the photoreceptor CC, and its possible roles in the vectorial transport of OS membrane proteins and in OS disk formation suggest yet another example of cell membrane motility involving actin. The presence of myosin and an actomyosin contractile mechanism are currently being investigated. It is yet to be determined how a local concentration of actin is restricted to the distal CC and whether or not this actin is maintained in the polymerized or unpolymerized states. Also of interest is how this actin may be associated with the CC plasma membrane. Since photoreceptor OS are renewed throughout life it is possible that a defect in actin-mediated OS disk formation could lead to photoreceptor cell death. Additional studies are in progress to further evaluate the role of actin in OS renewal.

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