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Nanodomain organization of rhodopsin in native human and murine rod outer segment disc membranes

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Abstract

Biological membranes display distinct domains that organize membrane proteins and signaling molecules to facilitate efficient and reliable signaling. The organization of rhodopsin, a G protein-coupled receptor, in native rod outer segment disc membranes was investigated by atomic force microscopy. Atomic force microscopy revealed that rhodopsin is arranged into domains of variable size, which we refer to herein as nanodomains, in native membranes. Quantitative analysis of 150 disc membranes revealed that the physical properties of nanodomains are conserved in humans and mice and that the properties of individual disc membranes can be variable. Examining the variable properties of disc membranes revealed some of the factors contributing to the size of rod outer segment discs and the formation of nanodomains in the membrane. The diameter of rod outer segment discs was dependent on the number of rhodopsin molecules incorporated into the membrane but independent of the spatial density of rhodopsin. The number of nanodomains present in a single disc was also dependent on the number of rhodopsin molecules incorporated into the membrane. The size of the nanodomains was largely independent of the number or spatial density of rhodopsin in the membrane.

Keywords

Atomic force microscopy; G protein-coupled receptor; lipid rafts; membrane structure; photoreceptor; phototransduction

1. Introduction

Rhodopsin is a prototypical G protein-coupled receptor (GPCR)¹ with 7 transmembrane domains (Fig. 1A). This light receptor is located in the outer segment of rod photoreceptor cells and initiates vision upon photon capture. The rod outer segment (ROS) is comprised of discs that are stacked one on top of another in a highly ordered manner [1, 2] (Fig. 2A).

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¹Abbreviations: AFM, atomic force microscopy; GPCR, G protein-coupled receptor; ROS, rod outer segment(s).

There are 500-2000 discs in a single vertebrate ROS [3], depending on the species, encased by a plasma membrane. Rhodopsin is primarily found in the disc membranes of ROS.

Rhodopsin is densely packed and found in high concentrations in ROS disc membranes [1, 4]. It is estimated that rhodopsin represents about 70-90% of the protein content in ROS and greater than 90% of the protein content in the disc membranes [5-9]. The high purity and concentration of rhodopsin in the ROS has greatly facilitated the structural, biophysical, and biochemical characterization of the native receptor, making it the most thoroughly studied GPCR in native form. The high concentration of rhodopsin in retinal membranes has been advantageous for experiments, yet this high concentration of molecules creates a crowded disc membrane environment in which signaling must occur with high efficiency, sensitivity, and reliability [10-13].

A crowded membrane environment is not unique to ROS disc membranes. All biological membranes have crowded environments with most being occupied by a heterogeneous complement of membrane proteins. Crowded environments are not ideal for efficient signaling if the signaling cascade proceeds in a random manner via freely diffusing proteins, a condition originally envisioned for biological membranes [14]. More recent evidence suggests biological membranes are more organized than initially recognized forming domains such as lipid rafts [15-18], which can facilitate the efficiency, sensitivity, and reliability that is required for signaling in the ROS [19-21]. The sizes of these domains are usually less than 200 nm [22] and are, therefore, difficult to visualize and study by conventional microscopy methods.

Atomic force microscopy (AFM) is uniquely suited to visualize the nanoscale organization of proteins and domains in biological membranes [23-27]. Imaging can be performed under physiological conditions and requires minimal processing of samples compared to other structural methods. AFM has revealed that ROS disc membranes like other types of biological membranes exhibit order. Two types of packing arrangements for rhodopsin in ROS disc membranes have been observed by AFM: a densely packed paracrystalline lattice arrangement and a less densely packed nanodomain organization. In both cases, rhodopsin forms an oligomeric complex composed of rows of dimeric receptor [28](Fig. 1B).

The first AFM image of an intact single bilayer disc membrane displayed a densely packed paracrystalline lattice of rhodopsin molecules [29]. This image has received the most attention to date and has provided the highest resolution of rhodopsin in native membranes, thereby allowing the generation of an oligomeric model of rhodopsin using the geometric constraints revealed in the image [28, 30] (Fig. 1B). Subsequent AFM images have consistently revealed a nanodomain organization of rhodopsin [28, 30-32], similar to that detected in the current study. Rhodopsin appears to form similar oligomeric complexes within these nanodomains to those present in the densely packed paracrystalline lattice arrangement [28]. The nanodomain organization has received less attention than the densely packed paracrystalline lattice organization. The nanodomain organization, however, may represent the native arrangement since it is the most consistently observed arrangement in AFM studies and is also consistent with cryoelectron tomograms of disc membranes in preserved intact ROS [1].

Dimerization/oligomerization appears to be a common trait among GPCRs [33-35]. Rhodopsin dimers appear to be the basic unit of oligomeric complexes in either densely packed paracrystalline lattices or in nanodomains [28]. The role of dimeric interactions in rhodopsin is beginning to be revealed. The arrangement of rhodopsin into dimers provides a platform for binding signaling partners such as transducin and arrestin, where the stoichiometry can be 2:1 between rhodopsin and the signaling partner [36-38]. Structural and functional asymmetry exists between rhodopsin molecules in a dimeric unit when bound to either transducin or arrestin [37, 39, 40]. This asymmetry may contribute to signaling efficiency and may play a protective role in photoreceptor cells under intense lighting conditions [37, 39, 40].

In contrast to dimeric interactions in rhodopsin, less is known about the role and properties of rhodopsin nanodomains. One limitation has been that the nanodomain organization of rhodopsin has only been qualitatively characterized. Therefore, basic knowledge about the nanodomains, including their size and factors contributing to their formation, has yet to be determined but is required to better understand their role in phototransduction and photoreceptor biology. To gain some of these important insights, AFM was utilized to quantitatively assess the nanodomain organization of rhodopsin in native ROS disc membranes.

Previous AFM studies have focused on samples obtained from murine and bovine retinas. Murine and bovine rhodopsin have served as models to understand the effect of the over 100 mutations detected in the rhodopsin gene that cause retinitis pigmentosa or congenital night blindness in human patients [41]. While murine and bovine rhodopsin are predicted to be structurally indistinguishable [42], it is unclear how structurally similar they are with human rhodopsin. There are 18 amino acid residue differences between human and murine rhodopsin (Fig. 1A). A comparison of rhodopsin nanodomains in ROS disc membranes from human and murine retinas was conducted to determine whether the amino acid residue differences alters the organization of the receptor in the membrane.

2. Materials and Methods

2.1. ROS disc membrane preparation

All experimental procedures were conducted under dim red light. Murine ROS disc membranes were prepared from the retina of 10-15 C57BL/6J mice (The Jackson Laboratory, Bar Harbor, ME) that were 4-6 weeks old. Eight preparations were used to collect data reported in the study. Mice were dark-adapted overnight prior to being sacrificed. Human ROS disc membranes were prepared from the retina of a single whole donor globe (Saving Sight, Columbia, MO). Five preparations were used to collect data reported in the study. The sex, age, and cause of death of donors are as follows: female, 45, septic shock; female, 63, esophageal cancer; female, 70, cerebral vascular accident; female, 75, lung cancer; male, 51, liver failure. Donors had no reported ocular history known to impact rod photoreceptor cells. Donor cadavers were cooled within 2 hours and eyes procured between 3.5-18 hours after death of the donor. Eyes were placed in Life4°C preservation media (Numedis Incorporated, Isanti, MN) and shipped on ice in a light-tight container. Eyes were received in the laboratory within 24-36 hours of donor death and

processed immediately. ROS and ROS disc membranes were obtained from murine and human retina using procedures reported previously [1, 43]. ROS disc membranes were resuspended in 50 μ L of Ringer's buffer (10 mM HEPES, 130 mM NaCl, 3.6 mM KCl, 2.4 mM $MgCl_2$, 1.2 mM $CaCl_2$, 0.02 mM EDTA, pH 7.4).

2.2. AFM imaging

All AFM procedures were conducted at ambient temperatures under dim red light. Samples were prepared for AFM by adding 40 μ L of ROS disc membranes (5-10 μ g/mL) onto freshly cleaved mica and incubating for 10 min. The mica was washed 5 times with 40 μ L of Ringer's buffer to remove unadsorbed material. ROS disc membranes adsorbed on mica were imaged by AFM in imaging buffer (20 mM Tris, 150 mM KCl, 25 mM $MgCl_2$, pH 7.8). AFM was performed using a Multimode II atomic force microscope equipped with an E scanner (13 μ m scan size) and silicon nitride cantilevers with a nominal spring constant of 0.06 N/m (NP-S, Bruker Corporation, Santa Barbara, CA). Samples were imaged using contact mode at a scan rate of 5.09 Hz to acquire images at a resolution of 512 lines per frame. Minimal force (< 100 pN) was applied to samples during imaging. Height and deflection images were collected and analyzed [44, 45].

2.3. Analysis of AFM images

The dimension of ROS disc membranes and nanodomains were measured using Nanoscope 5.3 software (Bruker Corporation, Santa Barbara, CA). The height of ROS disc membrane features were measured from height images that were flattened to the first order (Fig. 2E). The average height of the rim region, nanodomains, and protein-free lipid bilayer were obtained from peaks in height distribution histograms of individual images. These values were confirmed by line scan analysis. The lateral dimensions of ROS disc membranes and nanodomains were measured from deflection images. Accuracy of lateral dimensions is limited by tip convolution effects of the AFM probe [46, 47]. ROS disc membranes were presumed to be circular. The diameter of the entire disc membrane and the disc membrane excluding the rim region were measured. The latter measurement was used to compute the inner disc area. Nanodomains were presumed to be elliptical (Fig. 2G). The diameters of the ellipse were measured to compute the surface area of an individual nanodomain. To calculate the number of rhodopsin molecules in a nanodomain, rhodopsin was presumed to form an oligomer with a surface area of 84 nm² for six rhodopsin molecules (Fig. 1B). Statistical analysis was performed using Prism 6 (GraphPad Software Incorporated, La Jolla, CA). Mean values are reported with the associated standard deviation.

2.4. SDS-PAGE

ROS disc membranes were resuspended in LDS sample buffer containing 50 mM dithiothreitol (Expedeon Incorporated, San Diego, CA). Solubilized samples (500 ng of protein) were loaded onto a 4-12% Tris-Glycine precast gel (Life Technologies, Grand Island, NY) and electrophoresis was conducted. Gels were silver-stained to detect proteins extracted from ROS disc membranes.

3. Results

3.1. AFM of native ROS disc membranes

ROS were purified from human and murine retinas and disc membranes were released from the ROS (Figs. 2A-C). The purity of ROS disc membranes was assessed by SDS-PAGE (Fig. 2D). SDS-PAGE of ROS disc membranes isolated from both human and murine retinas displayed a single major band corresponding to rhodopsin [43]. Thus, preparations of ROS disc membranes were pure and contained predominantly rhodopsin.

ROS disc membranes were adsorbed on mica and imaged by AFM (Fig. 2C). Height images contain information about the height of material adsorbed onto mica, and were used to measure vertical features in ROS disc membranes (Fig. 2E). Deflection images accentuate the edges of sample features, and were used to measure lateral features of ROS disc membranes (Figs. 2F and 2G). Measurements made from AFM images are summarized in Table 1.

ROS disc membranes displayed a distinct topography in AFM images (Fig. 2E). Discs contain double lamellar membranes that are circumscribed by a hairpin loop forming a rim (Fig. 2C). A majority of adsorbed discs displayed only a single lamellar membrane with a rim region. Incisures were not regularly observed in AFM images, presumably because they are disrupted during adsorption of ROS discs to the mica support. Single-bilayer disc membranes predominantly adsorb on mica exposing the extracellular surface [32, 48]. Thus, AFM images of ROS disc membranes represent the topography of the extracellular surface in most instances. Individual rhodopsin molecules cannot be resolved by AFM in these instances since covalently linked sugar groups at the amino terminal region of rhodopsin interfere with the AFM probe [28].

3.2. Features of ROS disc membranes in AFM images

The average diameter of ROS disc membranes of both human and murine retinas was 1.2 μm (Table 1). The highest features in height profiles corresponded to the rim region (Fig. 2E). The heights of the rim region were variable and corresponded to intact and partially disrupted rims. The lamellar region of disc membranes exhibited two major classes of heights (Fig. 2E). One class had a height of about 4 nm (Table 1), which corresponds to the thickness of a lipid bilayer without embedded proteins. The protein-free lipid bilayer often formed a pool, presumably at the location where the disc breaks open, which may form from lipids in the rim region. The other class of heights from the lamellar region had a height of about 8 nm (Table 1), which corresponds to the height of rhodopsin [49]. The areas in the lamellar region with this height have been shown by AFM and single-molecule force spectroscopy to consist of high concentrations of rhodopsin [28, 32, 48].

Rhodopsin predominantly resides in the lamellar region of the ROS disc and is excluded from the rim region where structural proteins such as peripherin and Rom-1 are localized [50]. In AFM images of human and murine ROS discs, the lamellar region of the disc membrane is organized into clusters of distinct domains that are surrounded by protein-free lipid bilayer (Fig. 2F). The size of the domains is heterogeneous, but almost all have dimensions of less than 100 nm; therefore, these domains are referred to as nanodomains.

Both human and murine ROS discs displayed nanodomains in the lamellar region (Fig. 3). Since rhodopsin is the predominant protein species in the lamellar region of discs, rhodopsin molecules must largely form the observed nanodomains. It is possible that rhodopsin coexists with a minor population of additional proteins in these nanodomains.

3.3. Size of nanodomains and density of rhodopsin in disc membranes

Nanodomains have been observed in previous AFM studies on murine and bovine samples but have not been characterized quantitatively. The dimensions of the nanodomains were manually measured to determine their surface area (Fig. 2G). The surface areas of nanodomains from a single human or murine ROS disc generally displayed a skewed distribution that could be fit by a Log Gaussian function (Figs. 3A and 3B). Histograms of nanodomain surface areas obtained from all ROS disc membranes analyzed were also generated. These histograms also displayed a skewed distribution that could be fit by a Log Gaussian function (Figs. 3C and 3D).

The average size of nanodomains in human and murine ROS disc membranes was 1,109 nm² and 1,245 nm², respectively (Table 1). Human nanodomains occupied 22 % of the lamellar area of ROS disc membranes whereas murine nanodomains occupied 24 % of the same area. These values are consistent with previous estimates of 25 % of the ROS disc membrane area being occupied by rhodopsin [4].

The number of rhodopsin molecules present in a nanodomain was previously estimated assuming a monomeric arrangement of rhodopsin [32]. Thus, each rhodopsin molecule was estimated to occupy 15 nm² of space within the membrane [49]. High-resolution AFM images of nanodomains, however, display oligomeric complexes of rhodopsin organized as rows of dimers [28]. The surface area of oligomeric rhodopsin was therefore used to estimate the number of rhodopsin molecules present in a nanodomain of a given size. Using the oligomeric model derived from AFM data [28, 30] (Fig. 1B), six rhodopsin molecules are estimated to occupy 84 nm² of space on the extracellular surface. Using this value, the number and spatial density of rhodopsin in ROS disc membranes were computed (Table 1).

The average spatial density of rhodopsin in a human ROS disc membrane is 15,865/μm² (Table 1). This value represents the density of rhodopsin if it were homogeneously distributed within the membrane. The spatial density of rhodopsin in murine ROS disc membranes is 17,486/μm². The rhodopsin density values derived from AFM images of nanodomains are much lower than that estimated from a densely packed paracrystalline lattice arrangement, where the density was estimated to be 48,300/μm² [29]. Rhodopsin density values derived from a nanodomain arrangement are more consistent with previously reported estimates of about 25,000/μm² [1, 4], albeit a little lower. The discrepancies are small and may be due to a variety of factors. Errors associated with different methods may account for the small differences in the estimate of rhodopsin density. In the current AFM study, rhodopsin monomers may coexist with rhodopsin oligomers that form nanodomains. Diffusely distributed monomers would not be distinguishable in areas that have been classified as protein-free lipid bilayers (Fig. 2E). The presence of monomers would increase rhodopsin density estimates. In addition, computed density values may be underestimated

because the estimate for the inner disc area included a proposed girdle zone separating the rim region from the area containing rhodopsin [32].

3.4. Correlation of parameters computed from AFM images

In addition to revealing the average properties of ROS discs, the analysis of 150 individual AFM images revealed the variability that exists among discs. Quantification of parameters describing features of ROS disc membranes provided insights about the properties of nanodomains and their influence on the ROS disc. The variability and correlation of parameters were similar between human and murine samples (Fig. 4).

Although the average diameter of ROS disc membranes is consistent with previous estimates [1], variability was observed in the diameter of individual disc membranes. The diameter of ROS disc membranes ranged from 0.7 – 1.9 μm . No correlation was observed between the diameter of ROS disc membranes and the spatial density of rhodopsin in the disc membrane (Fig. 4A). In contrast, a strong positive correlation was observed between the diameter of ROS disc membranes and the number of rhodopsin molecules present in the membrane (Fig. 4B).

The number of nanodomains found in a ROS disc membrane is strongly correlated with the number of rhodopsin molecules and is more weakly correlated with the spatial density of rhodopsin (Figs. 4C and 4D). In contrast, the average size of nanodomains did not show a correlation, or only a weak correlation, with either the number or spatial density of rhodopsin in the ROS disc membrane (Figs. 4E and 4F).

4. Discussion

4.1. Nanodomain organization of rhodopsin

In the current study, investigation of ROS disc membranes from human and murine retinas by AFM revealed that rhodopsin is organized into nanodomains in both species. Nanodomains of rhodopsin have also been observed in ROS disc membranes from bovine retina [32]. The nanodomain organization of rhodopsin is the most consistently observed arrangement in ROS disc membranes by AFM and is also consistent with the heterogeneous densities observed within disc membranes in cryoelectron tomograms of preserved intact murine ROS [1, 28, 30-32]. The observed nanodomain organization differs from the densely packed paracrystalline lattice arrangement reported in the first AFM study of ROS disc membranes [29]. ROS disc membranes displaying a densely packed paracrystalline lattice were not observed in the current study or in a previous study [32]. Thus, the nanodomain organization of rhodopsin likely represents the most common or physiologically relevant arrangement in the hundreds of stacked discs of the intact ROS. The formation of rhodopsin nanodomains is conserved among humans, mice, and cows, and likely other vertebrate species as well.

4.2. Factors that impact the size of ROS discs

The analysis of individual ROS disc membranes allows for the detection of variability that exists among ROS discs. Variability is present in both the size of ROS discs and the number

of rhodopsin molecules incorporated into the disc (Fig. 4). The number of rhodopsin molecules incorporated into ROS disc membranes has a strong influence on the size of ROS discs (Fig. 4B). The correlation between disc diameter and the number of rhodopsin molecules is consistent with observations where either overexpression or underexpression of rhodopsin in genetically modified mice resulted in larger or smaller diameters of ROS, respectively, which in turn is accompanied by altered electrophysiological responses [31, 51, 52].

The variability in the number of rhodopsin molecules incorporated into a ROS disc likely does not arise from differences at the level of transcription or rhodopsin synthesis, but instead, may be due, in part, to the variability in the rate of ROS disc formation induced by light [53, 54]. Constant rhodopsin synthesis and variable rates of ROS disc formation would result in different amounts of rhodopsin being incorporated into the discs. In contrast to the number of rhodopsin molecules incorporated into a ROS disc, the density of rhodopsin in the disc membrane is not a factor in determining the size of ROS discs (Fig. 4A). Thus, the size of ROS discs may adjust to maintain a constant range or average density of rhodopsin in disc membranes [55], perhaps in order to maintain signaling efficiency or sensitivity.

4.3. Factors that impact the number and size of nanodomains

Understanding the physical properties of nanodomains will be required to better understand their functional role. The incorporation of more rhodopsin molecules into disc membranes results in the formation of a greater number of nanodomains (Figs. 4C and 4D). In contrast, the size of nanodomains is largely unaffected by either the number or spatial density of rhodopsin in the membrane (Figs. 4E and 4F).

Rhodopsin likely arranges in nanodomains as an oligomeric complex consisting of rows of dimeric receptor [28]. The factors contributing to the oligomerization of rhodopsin, or other GPCRs, are poorly understood. The hydrophobic mismatch between monomeric GPCRs and the lipid bilayer appears to play a significant role in the spontaneous aggregation of GPCRs [56, 57]. Amino acid residue differences can result in different levels of hydrophobic mismatch, thereby resulting in different sizes of oligomers [56]. Since the sizes of nanodomains in human and murine ROS disc membranes are similar (Table 1), the 18 amino acid residue differences in human and murine rhodopsin do not appear to play a role in determining the size of nanodomains. This is consistent with the putative oligomeric model of rhodopsin where the amino acid residue differences occur in regions of the receptor that are not directly involved in the putative oligomeric interfaces (Fig. 1B), although some are in close proximity to these proposed interfaces.

The skewed distributions displayed in histograms of nanodomain size (Fig. 3) indicate that the variability in the size of nanodomains is random and arises from the product of multiple independent factors. The number or spatial density of rhodopsin in the ROS disc membrane is not one of the factors contributing to this variability (Figs. 4E and 4F). The variability in the average size of nanodomains observed among different ROS discs must then arise from extrinsic factors. One factor may be the lipid bilayer in which rhodopsin is embedded. Properties of the lipid bilayer can have significant influence on the structure and function of rhodopsin [58-63]. The lipid composition is heterogeneous in discs at different axial

positions in ROS [64-67], and lipids can impact the oligomerization of rhodopsin [57, 62]. Thus, ROS discs with different complements of lipid in the membrane may result in variable sizes of nanodomains, which in turn may impact phototransduction.

4.4. Nanodomains and rhodopsin diffusion

Classical studies demonstrating the apparent rapid lateral and rotational diffusion of rhodopsin in the disc membranes of amphibian ROS led, in part, to the notion that rhodopsin freely diffuses within the membrane as a monomer [68-71]. These studies are inconsistent with a densely packed paracrystalline lattice arrangement of rhodopsin [72], where the receptor would be expected to be relatively immobile. Determining the size of rhodopsin nanodomains in the current AFM study allows for the assessment of whether this type of organization is consistent with reported lateral diffusion coefficients.

Since the size of a protein embedded in membranes only weakly impacts the lateral diffusion rate [73], it is difficult to differentiate between monomeric and oligomeric rhodopsin by monitoring diffusion alone. A range of lateral diffusion coefficients has been reported for rhodopsin in amphibian ROS: 0.1-0.6 $\mu\text{m}^2/\text{s}$ [68, 70, 71, 74-79]. Multiple factors can lead to incorrect estimates of rhodopsin lateral diffusion (e.g., [78, 79]), and improvements in experimental procedures indicate that the lower end of the reported range better describes the lateral diffusion of rhodopsin [79].

Lipid rafts in the plasma membrane of various types of cells are predicted to be heterogeneous with sizes in the range of 10-200 nm [22]. Rhodopsin nanodomains fall within the lower half of this size range. It is unclear whether rhodopsin nanodomains represent a typical lipid raft found in plasma membranes since the majority of rhodopsin molecules in ROS disc membranes are not found in the detergent-insoluble fractions typically associated with lipid rafts [80]. However, the similarity in size between rhodopsin nanodomains and lipid rafts suggest that some parallels may exist. Lipid rafts are mobile and those with sizes on the lower end of the spectrum are predicted to exhibit lateral diffusion coefficients of about 0.1 $\mu\text{m}^2/\text{s}$ [81]. If rhodopsin nanodomains exhibit similar properties as smaller lipid rafts, then rhodopsin nanodomains would be expected to be mobile within the ROS disc membrane and exhibit a lateral diffusion coefficient consistent with the lower end of reported values obtained from amphibian ROS.

4.5. Nanodomains and phototransduction

A functional role for nanodomains may be to provide order to the ROS disc membranes so that efficient and sensitive signaling in ROS can be achieved despite a crowded environment [13, 82]. Theoretical considerations suggest that the nanodomain organization of rhodopsin can facilitate signaling that is quantitatively consistent with classical kinetic data on the initial steps of phototransduction in ROS [83, 84]. Rhodopsin must be arranged as oligomeric complexes consisting of rows of dimeric receptor within nanodomains to achieve consistency with classical kinetic data [84, 85]. Consistency with classical kinetic data can be achieved even in the absence of rhodopsin diffusion within the membrane. Oligomeric complexes of rhodopsin in nanodomains can provide a platform for transient interactions between rhodopsin and transducin in the dark state, which may contribute to the efficiency

observed in the phototransduction cascade initiated upon light activation of rhodopsin [86]. Nanodomains are consistent with classical diffusion and kinetic data and therefore may represent the organizing principle utilized by ROS disc membranes to achieve signaling efficiency and sensitivity.

5. Conclusions

AFM has provided an avenue to directly visualize the organization of rhodopsin into nanodomains in native ROS disc membranes. The analysis of single ROS disc membranes has revealed the variability in the size of nanodomains and the number and spatial density of rhodopsin in the ROS disc membranes. This quantification has provided insights about some of the basic rules that define the properties of ROS discs and the organization of rhodopsin within the membrane under normal conditions. The number of rhodopsin molecules is a determinant for the size of ROS discs and the number of nanodomains formed. In contrast, the concentration of rhodopsin in the membrane is not a factor in determining the size of ROS discs. The size of nanodomains is largely independent of the number or density of rhodopsin in the disc membranes, and instead, is presumably affected by extrinsic factors. Despite 18 amino acid residue differences (Fig. 1), both human and murine rhodopsin form nanodomains with similar physical properties.

The results from this study lay the groundwork to begin dissecting out the specific factors governing nanodomain formation. The similarities in the properties of human and murine nanodomains indicate that factors contributing to the formation of nanodomains are conserved and that murine ROS disc membranes can serve as a good model for understanding the organization of human rhodopsin in ROS disc membranes. Additionally, the availability of different animal models will allow the testing of whether pathological conditions can alter the normal organization of rhodopsin in the ROS disc membrane and provide insights into the detrimental effects that may follow (e.g., [87, 88]). Although rhodopsin is unique among GPCRs in that it is expressed at high concentrations and purity in ROS disc membranes, the organization of receptors into nanodomains may be a common feature among GPCRs [89].

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Highlights

- The organization of rhodopsin in native membranes was visualized by AFM
- Human and murine rhodopsin form nanodomains with similar physical properties
- The size of nanodomains is heterogeneous within a single rod outer segment disc
- Heterogeneity was observed among individual rod outer segment disc membranes
- Factors contributing to nanodomain and disc membrane properties were determined

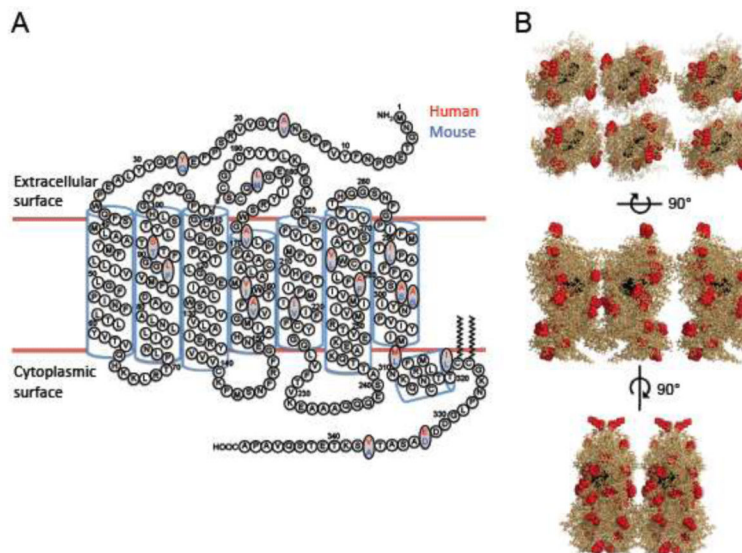


FIGURE 1. Structure of rhodopsin

(A) The secondary structure of rhodopsin is shown. The 18 amino acid residue differences in the sequences of human (red) and mouse (blue) rhodopsin are highlighted on the secondary structure. (B) The oligomeric model of rhodopsin (PDB: 1N3M) derived from AFM studies is shown in stick representation. The location of the 18 amino acid residue differences in the sequences of human and mouse rhodopsin are highlighted as red spheres. The chromophore 11-*cis* retinal is shown as black spheres. The first structure is a top view of the extracellular surface. The other structures are side views with the extracellular surface on top and the cytoplasmic surface on the bottom.

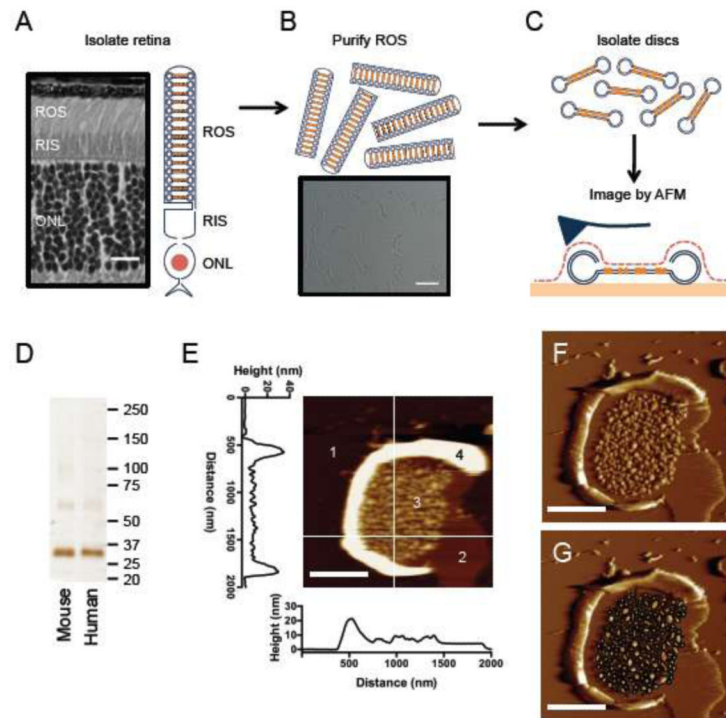


FIGURE 2. Preparation and AFM imaging of ROS disc membranes

(A) A cross-section of a mouse retina and a cartoon of a rod photoreceptor cell are shown. The rod outer segment (ROS), rod inner segment (RIS) and outer nuclear layer (ONL) are highlighted. Scale bar, 15 μm . (B) Purified ROS are shown as a cartoon and in a light micrograph. Scale bar, 15 μm . (C) ROS discs are isolated from purified ROS by osmotic bursting and washing steps. ROS disc membranes are adsorbed onto mica for AFM imaging. In AFM, a sharp probe is raster-scanned over the sample surface to generate topographical images. (D) SDS-PAGE on ROS disc membrane preparations from mouse and human samples reveal that rhodopsin is the predominant protein species. The sizes of protein standards are indicated in kDa. (E) A height image of a ROS disc membrane. Four different surfaces are revealed: 1, mica; 2, protein-free lipid bilayer; 3, rhodopsin nanodomains; 4, rim region. The height profiles of the highlighted line scans are shown. Scale bar, 500 nm. (F) A deflection image of the same ROS disc membrane. Scale bar, 500 nm. G, The deflection image with nanodomains outlined by black ellipses. The diameters of the ellipses were measured to determine the surface area of the nanodomains.

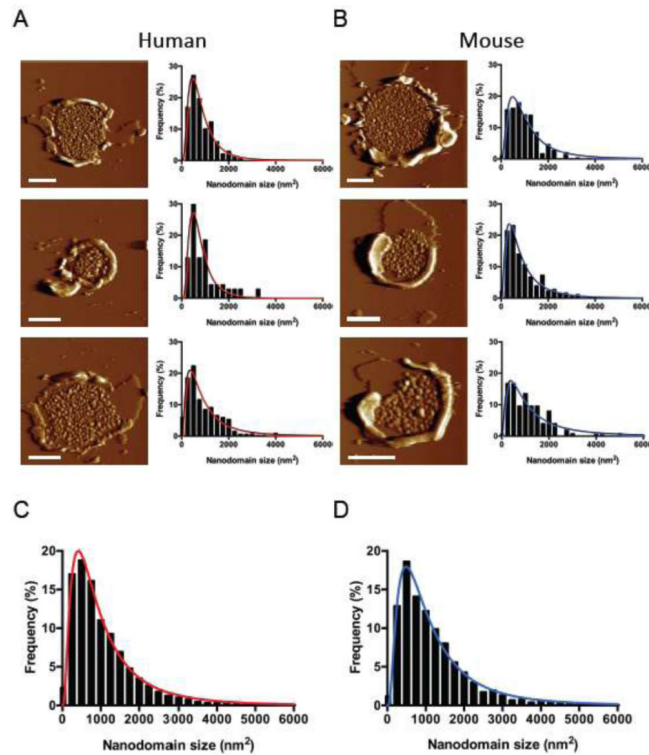


FIGURE 3. Deflection images of human and murine ROS disc membranes
 (A, B) A sampling of deflection images of human and murine ROS disc membranes is shown. A histogram of rhodopsin nanodomain surface areas measured in each ROS disc membrane is shown along with the fit of the data to a Log Gaussian function. (C) A histogram of nanodomain surface areas measured from 50 images of human ROS disc membranes is shown along with the fit of the data to a Log Gaussian function ($n = 7,793$). (D) A histogram of nanodomain surface areas measured from 100 images of murine ROS disc membranes is shown along with the fit of the data to a Log Gaussian function ($n = 15,608$).

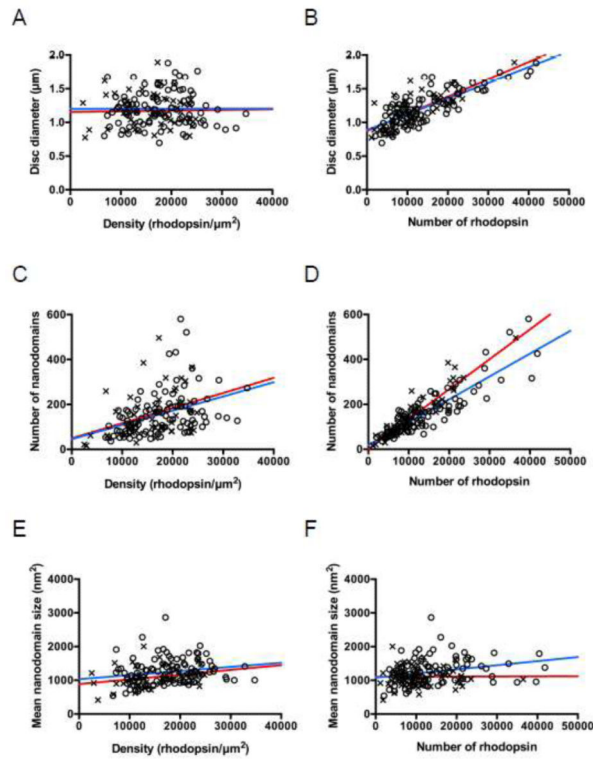


FIGURE 4. Correlation between different parameters computed from AFM images of ROS disc membranes

Human (red, \times , $n = 50$) and mouse (blue, o , $n = 100$) data were plotted and fit by linear regression. A correlation analysis was conducted to compute the Pearson coefficient (r) and level of significance (p). (A) Disc diameter vs. density of rhodopsin. Human, $r = 0.02$, $p = 0.87$. Mouse, $r = 0.002$, $p = 0.98$. (B) Disc diameter vs. number of rhodopsin molecules. Human, $r = 0.75$, $p < 0.0001$. Mouse, $r = 0.79$, $p < 0.0001$. (C) Number of nanodomains vs. density of rhodopsin. Human, $r = 0.38$, $p = 0.006$. Mouse, $r = 0.40$, $p < 0.0001$. (D) Number of nanodomains vs. number of rhodopsin molecules. Human, $r = 0.92$, $p < 0.0001$. Mouse, $r = 0.88$, $p < 0.0001$. (E) Mean nanodomain size vs. density of rhodopsin. Human, $r = 0.27$, $p = 0.06$. Mouse, $r = 0.20$, $p = 0.05$. (F) Mean nanodomain size vs. number of rhodopsin molecules. Human, $r = 0.008$, $p = 0.95$. Mouse, $r = 0.28$, $p = 0.006$.

Table 1

Summary of quantitative analysis of AFM images

Properties of a ROS disc	Parameter value	
	Human	Mouse
Disc diameter	1.17 ± 0.24 μm	1.20 ± 0.25 μm
Inner disc area	0.74 ± 0.37 μm ²	0.77 ± 0.39 μm ²
Nanodomain height	8.11 ± 0.21 nm	8.10 ± 0.44 nm
Lipid bilayer height	3.68 ± 0.31 nm	3.69 ± 0.28 nm
Number of nanodomains	156 ± 103	156 ± 99
Nanodomain density	208 ± 79 μ ⁻²	207 ± 79 μ ⁻²
Mean nanodomain size	1,109 ± 311 nm ²	1,245 ± 379 nm ²
Disc area occupied by nanodomains	22 ± 8 %	24 ± 9 %
Number of rhodopsin molecules	11,642 ± 7,108	13,513 ± 8,522
Rhodopsin spatial density	15,865 ± 5,894 μm ⁻²	17,486 ± 6,170 μm ⁻²

Parameter values were determined from individual ROS disc membranes. The mean values are reported with the associated standard deviation. The mean was determined from the analysis of 50 human ROS disc membranes or 100 murine ROS disc membranes. Significant differences between human and mouse data was assessed by a t-test. No significant differences between parameters were observed ($p > 0.05$) except for the mean nanodomain size in a ROS disc membrane, where there is a small but significant difference ($p = 0.03$).