Galactose Induces Formation of Chains of the Retinal Adhesion Protein, 
Retinoschisin

J. Bernard Heymann¹, Camasamudram Vijayasarathy², Rick Huang³, Altaira Dearborn¹, Paul A. 
Sieving²,⁴ and Alasdair C. Steven¹

¹ Laboratory for Structural Biology Research, NIAMS, NIH, Bethesda, MD.
² Section on Translational Research for Retinal and Macular Degeneration, NIDCD, NIH, Bethesda, 
MD.
³ Cryo-Electron Microscopy Facility, Janelia Research Campus, HHMI, Ashburn, VA.
⁴ NEI, NIH, Bethesda, MD.

Retinoschisin (RS1) is involved in cell-cell junctions in the retina, but is unique among known cell-
adhesion proteins in that it is a soluble secreted protein. Loss-of-function mutations in RS1 lead to early 
vision impairment in young males, called X-linked retinoschisis (XLRS). This name refers to the 
separation of the inner retinal layers, disrupting synaptic signaling. We previously reported the structure 
to 4.1 Å, revealing double octamer rings [1]. Each 24 kDa subunits has a small N-
terminal RS1 domain 
in the middle of the ring, and a larger C-terminal discoidin domain at the periphery. As a soluble protein, 
RS1 is unlikely to interact directly with the hydrophobic parts of the cell membrane. Importantly, RS1 
binds several sugar moieties, of which the best characterized is galactose [2]. It may tie retinal membranes 
together through interaction with the abundant glycans found on lipid-embedded and extracellular matrix 
proteins.

We imaged RS1 (purified as in [1]) in the presence of galactose by cryo-electron microscopy. Surprisingly, 
the RS1 double octamer rings formed strings of molecules (Figure 1A) not seen without galactose. They 
aligned at the air-water interface to show mostly one or a small number of side views. We processed these 
(using the Bsoft package [3]) to find classes for averaging and interpretation: Figure 1B shows the six best 
class averages. The repeat along the chain is 124 Å, somewhat less than the maximum diameter of 140 Å 
(measured between the subunit spike tips). Curiously, the central planes through the rings from one 
molecule to the next do not line up, but are offset by ~13 Å. Along a chain of molecules, the offset switches 
direction at every interface, giving a slightly zig-zag appearance. This indicates that the interaction 
propagates along the chain, with some communication of the conformation through the rings. We 
determined the orientation of each double ring to within a small uncertainty (±3°), allowing us to place 
3D maps in the correct relationship with each other (Figure 2A). The interaction between the double rings 
is through the spikes, with contributions from four subunits of each double ring (Figure 2B). The interface 
between neighboring double rings appears to be extensive and well-formed (Figure 2C). However, the 
two rings must break their symmetry to produce the alternating offset, likely with a different interface 
from that portrayed in Figure 2C. Future studies will focus on obtaining better 3D maps of the chains, 
locating galactose in the map, investigating the binding of other carbohydrates and assessing the 
implications of these interactions in the retina [4].

References:

This work was supported by the Intramural Research Programs of NEI, NIDCD and NIAMS.

Figure 1. (A) Cryo-electron micrograph of retinoschisin (~0.14 mg/ml) with galactose (10 mM) taken on a FEI Krios with a Gatan K2 camera in super-resolution mode (dose of 4 e⁻/Å² each 0.25s frame for 20 frames = 80 e⁻/Å²). Scale bar: 300 Å. (B) Class averages of chains of RS1 double rings (with the number of contributing images), showing a lateral offset propagated in alternating fashion along the chain. Scale bar: 100 Å.

Figure 2. Assembly of 3D maps and models consistent with the major view of the double rings in the chains. (A) Side view as seen in the micrographs. (B) Top view, showing the interaction between the subunits of the double rings. (C) The interface between the double rings shows good connectivity and little overlap, despite being refined on isolated double rings in a previous study [1]. Two different interface types connect the two pairs of rings. The important R141 is buried in the subunit core and may not directly contribute to the interface. A few other residues that lead to disease when mutated are noted. Map source: EMDB 6425; Atomic structure source: PDB 3JD6.