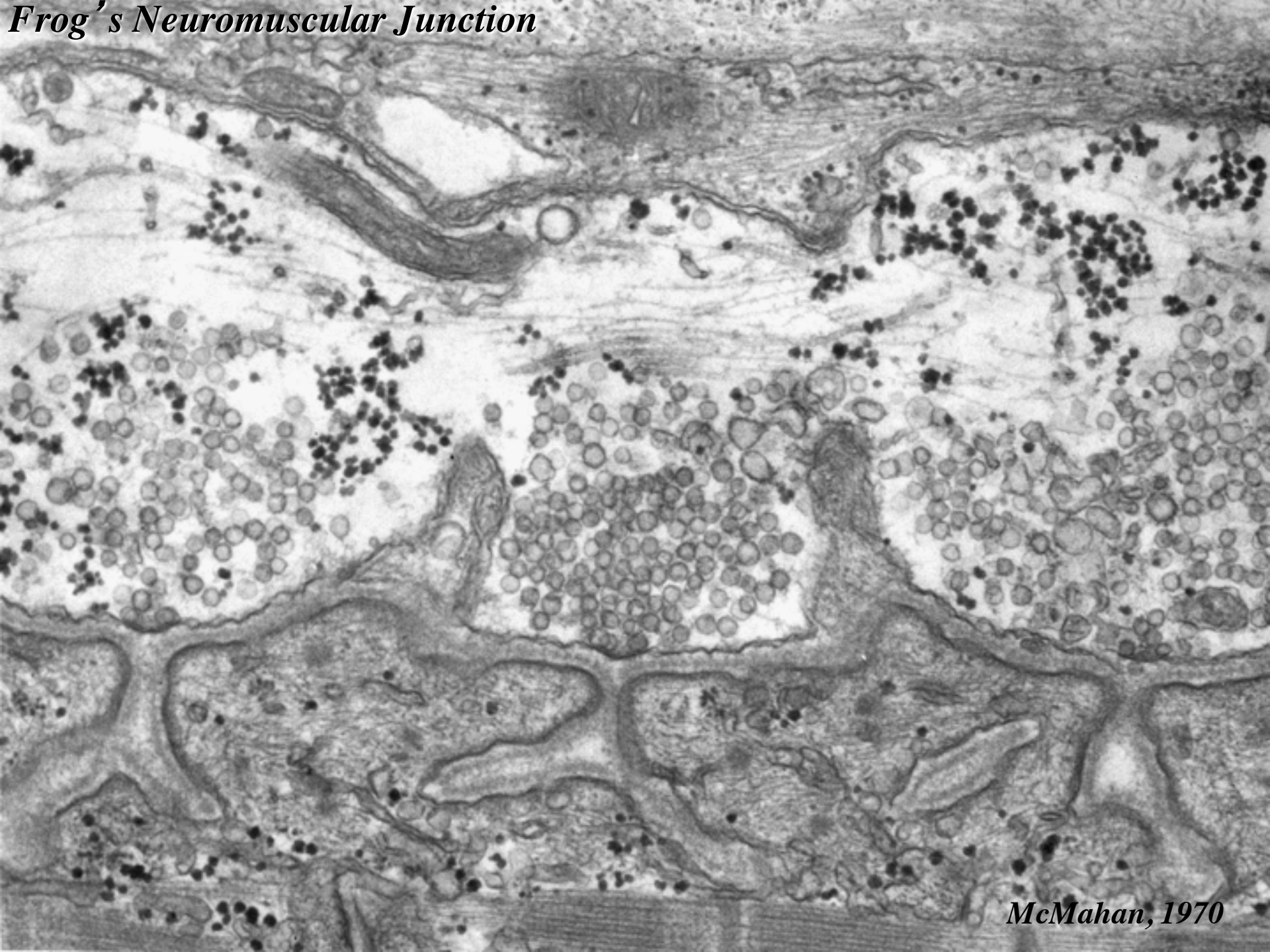
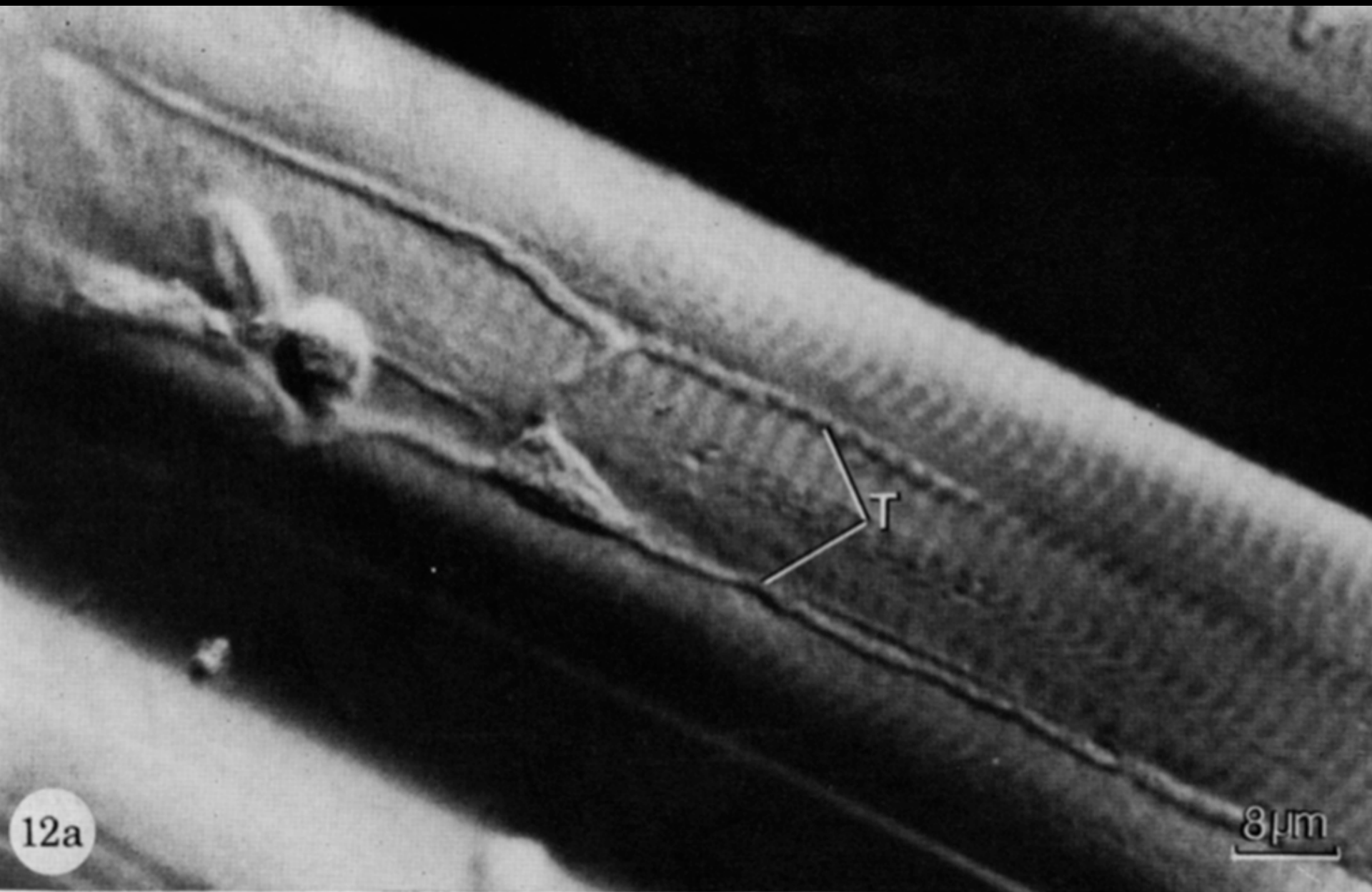
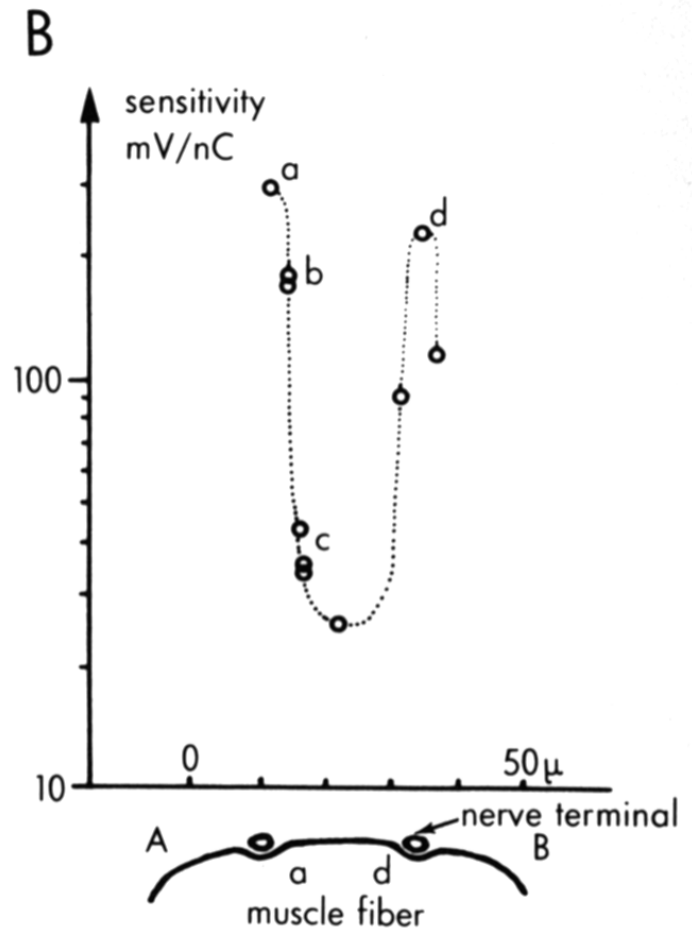
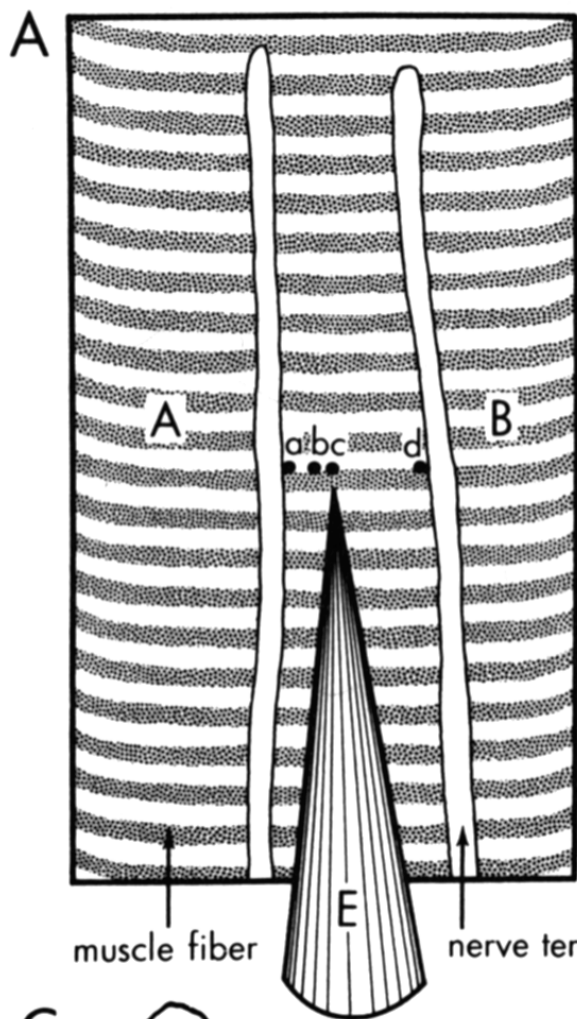


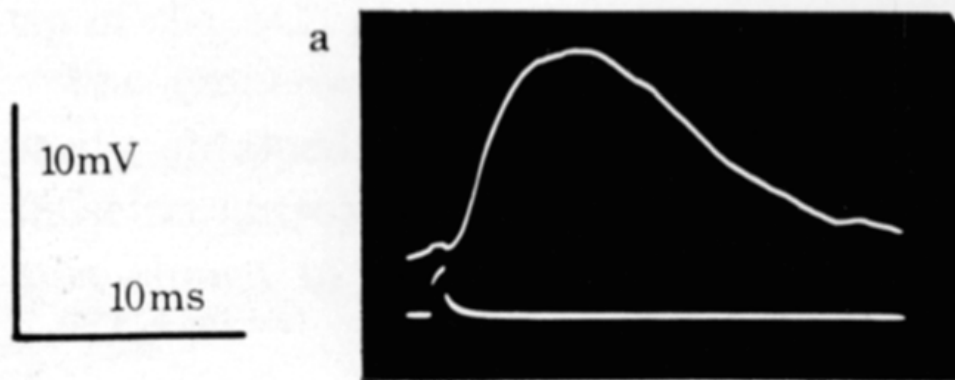
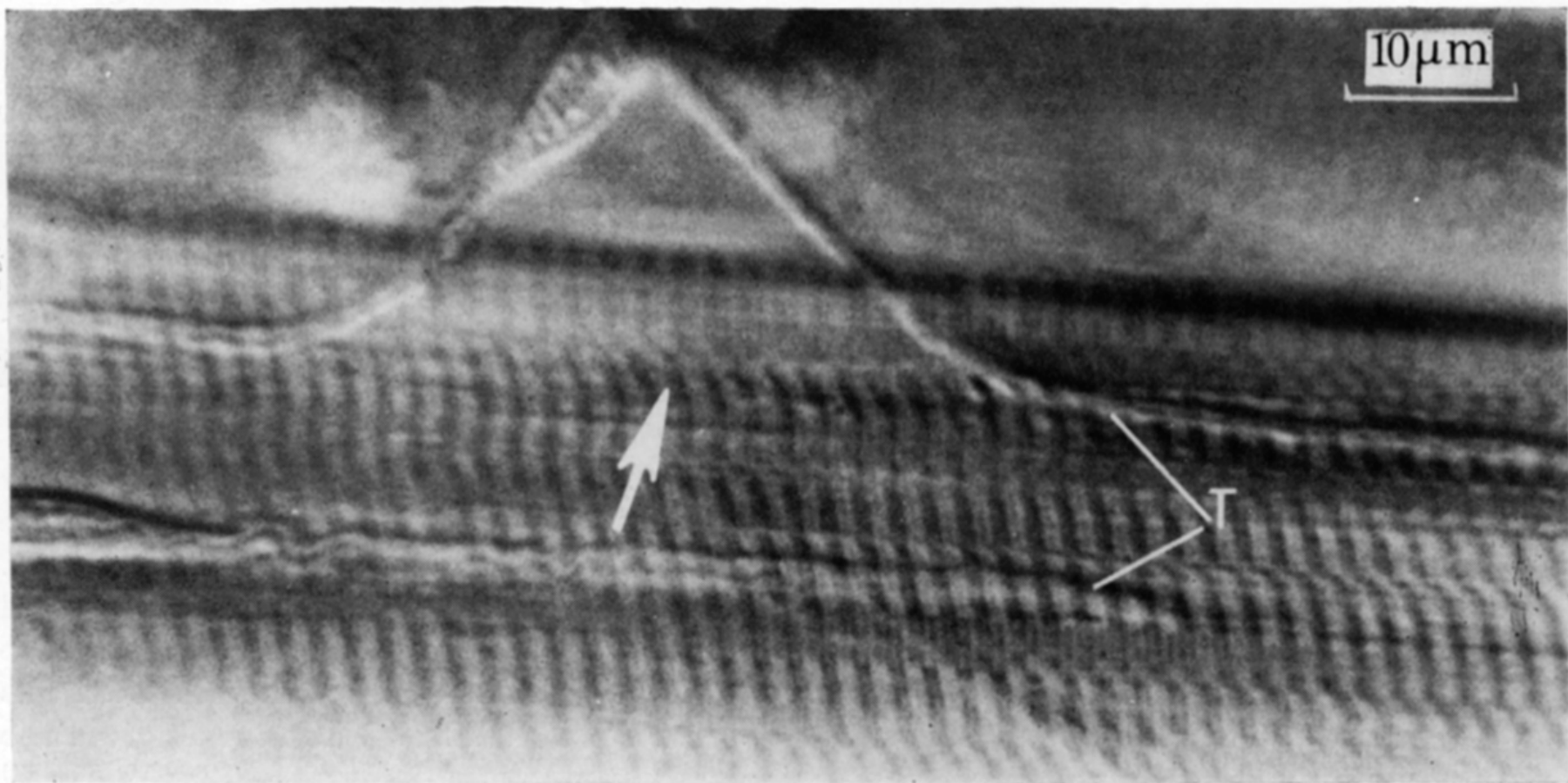
Frog's Neuromuscular Junction



McMahan, 1970







Frog NMJ/ HRP- α -BTX



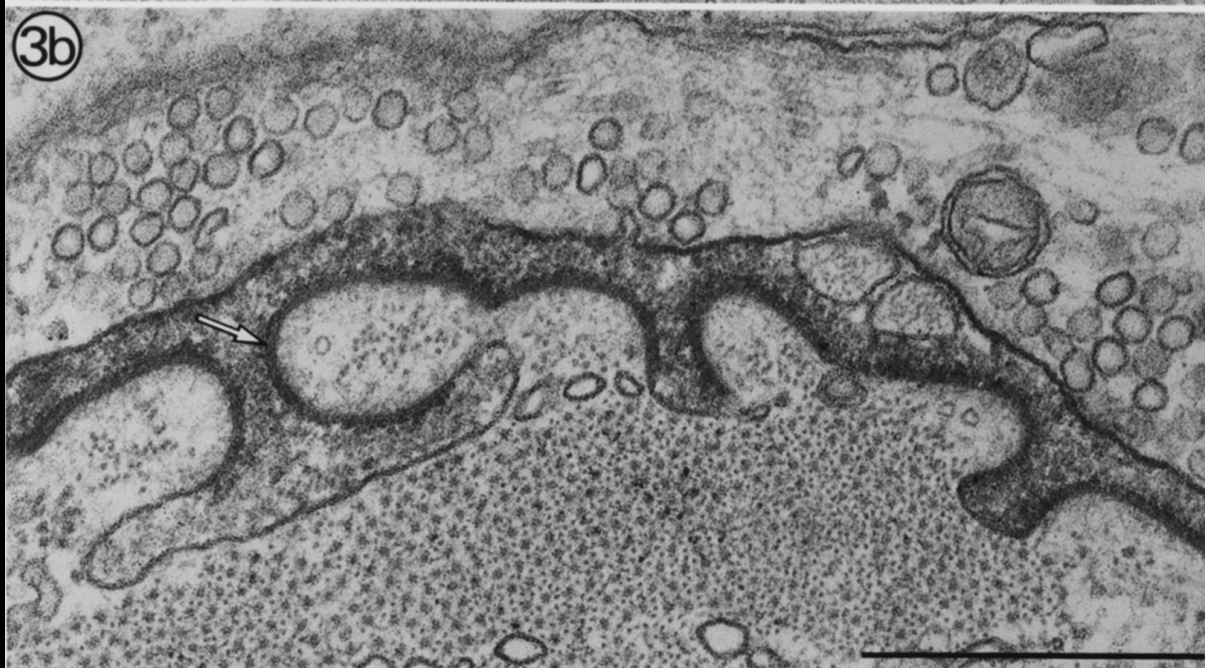
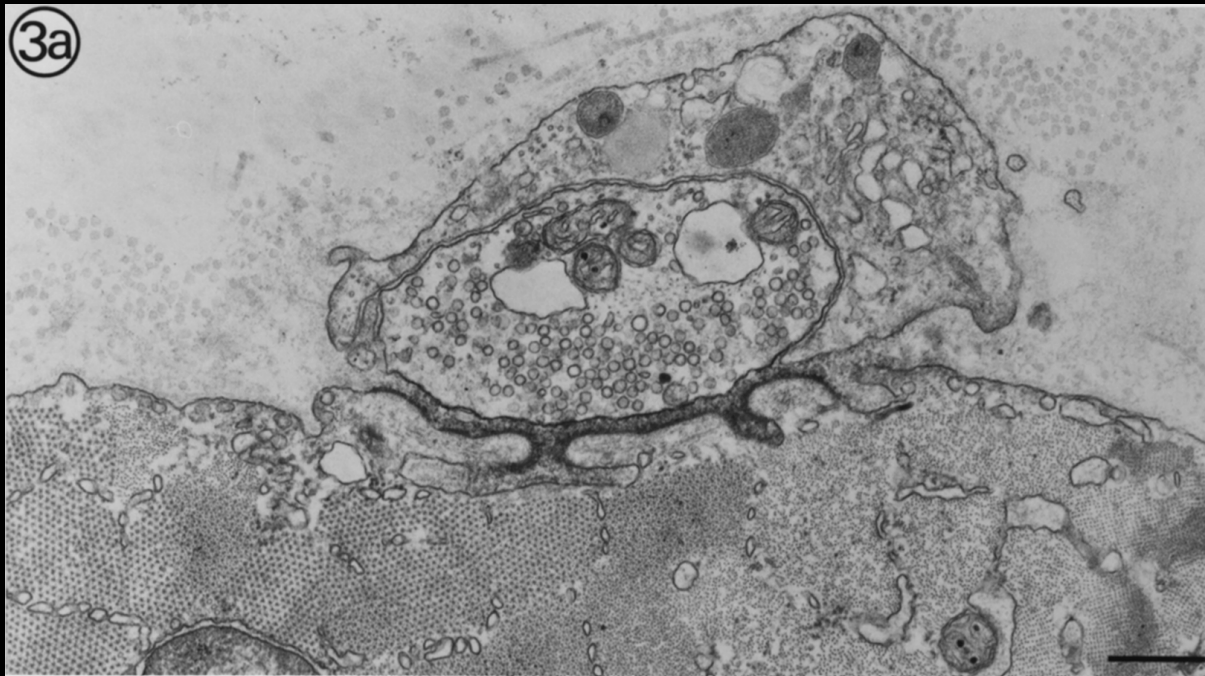
20 μ m

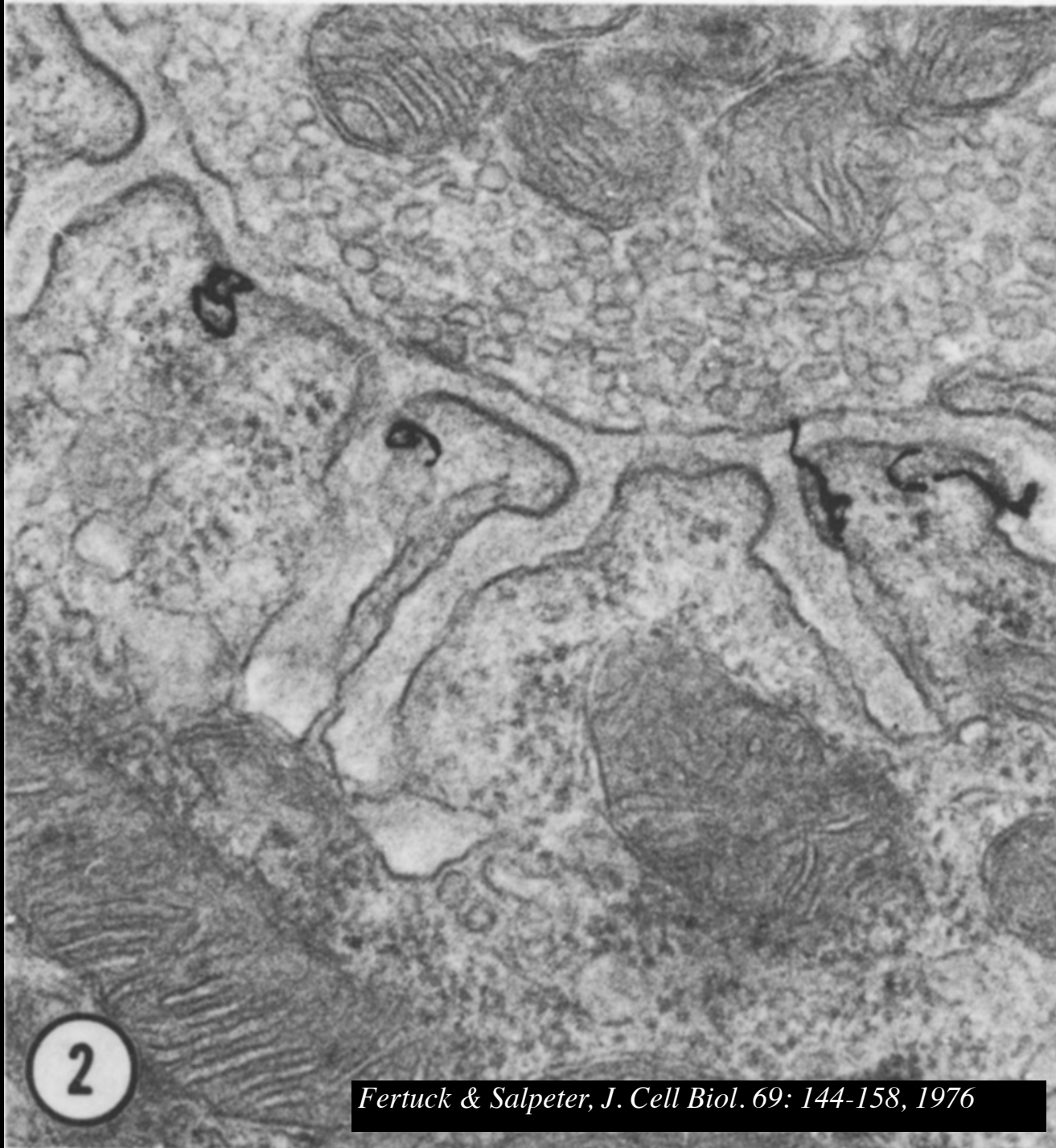
Rat NMJ/ HRP- α -BTX



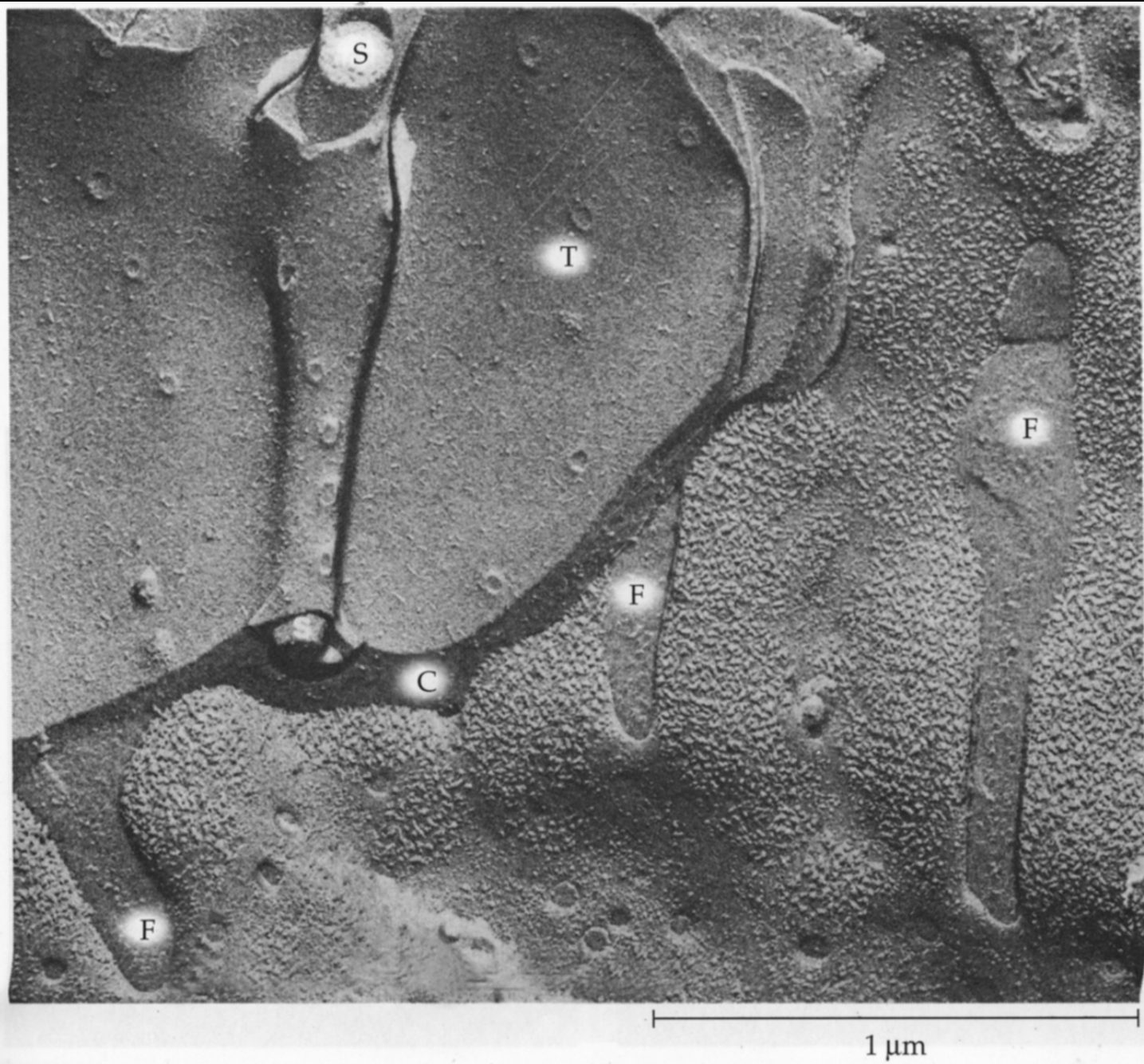
20 μ m

McMahan, 1984

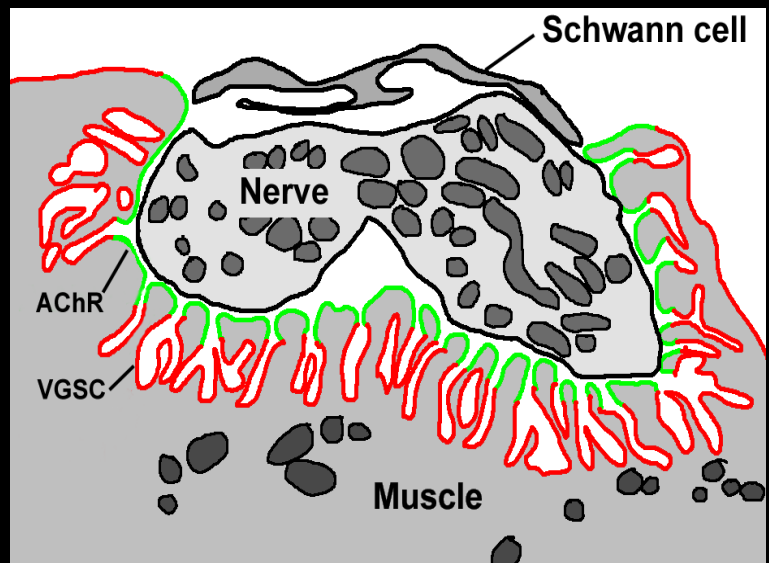
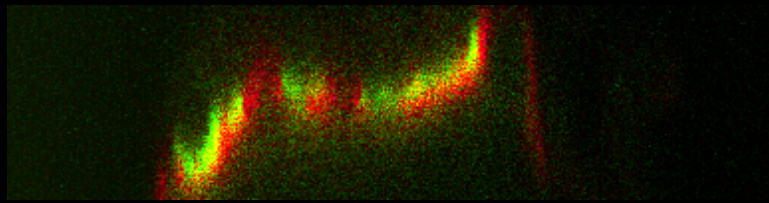
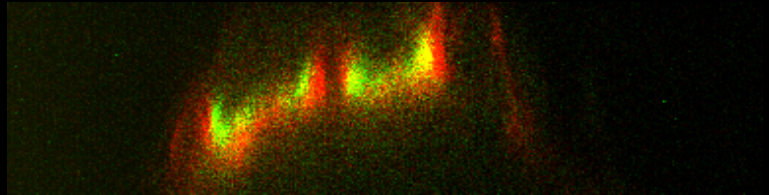
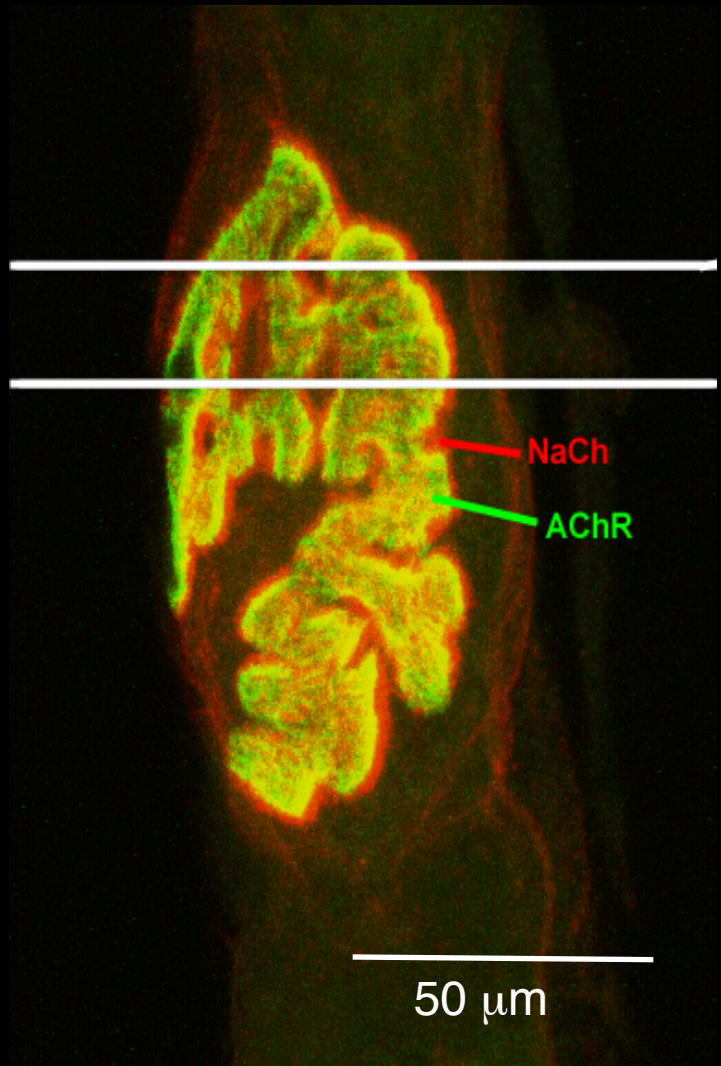


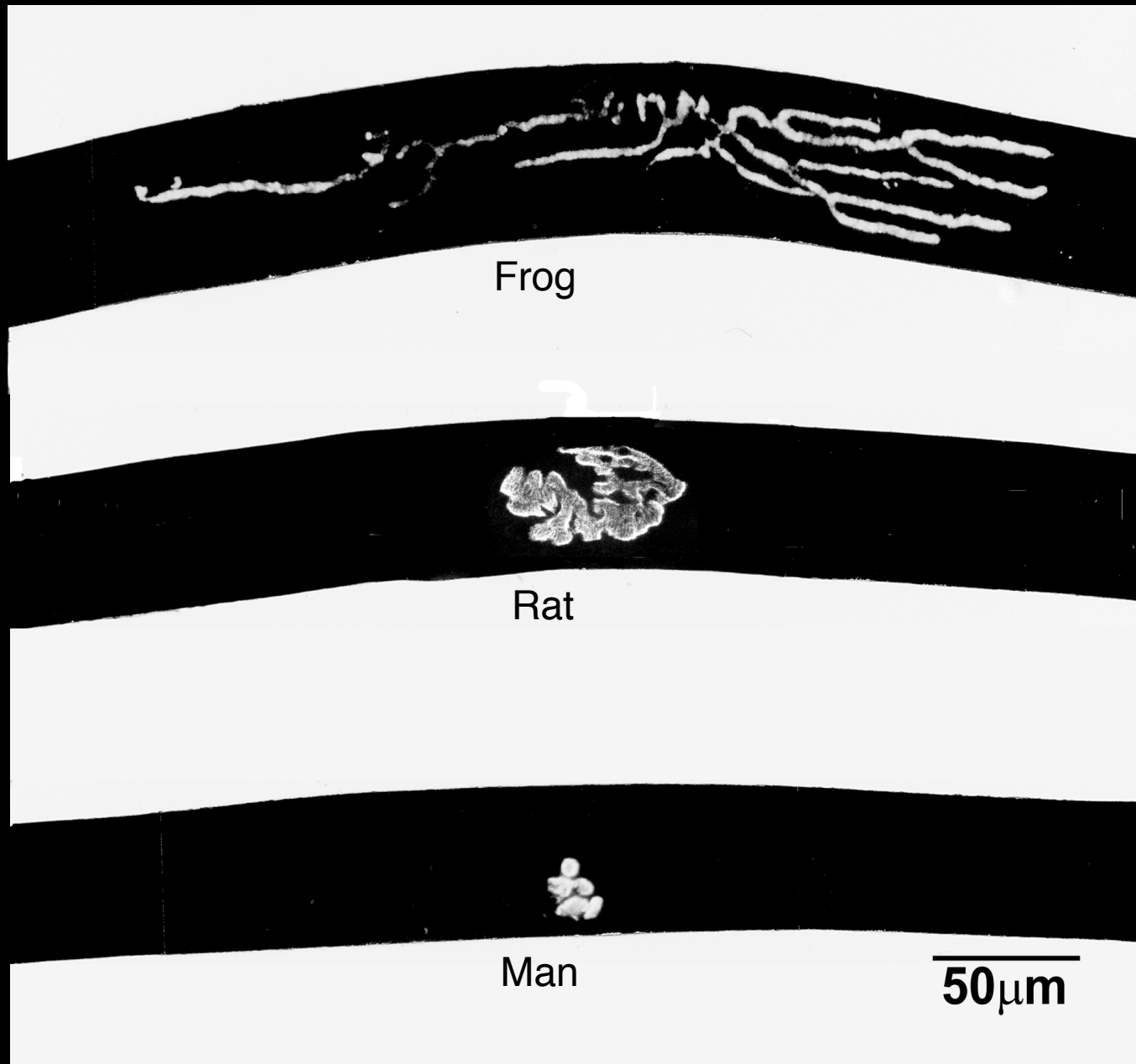


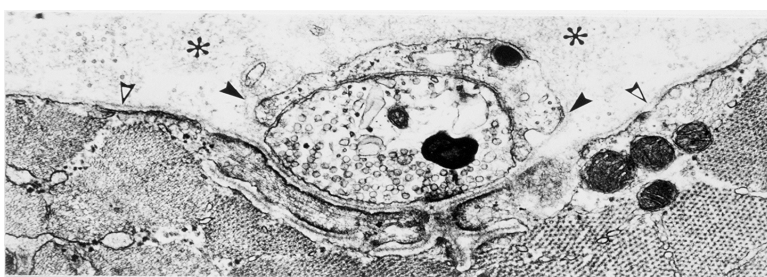
Fertuck & Salpeter, J. Cell Biol. 69: 144-158, 1976



Heuser, Reese, Llandis (1974), J. Neurocytol 3:109-131.







Frog

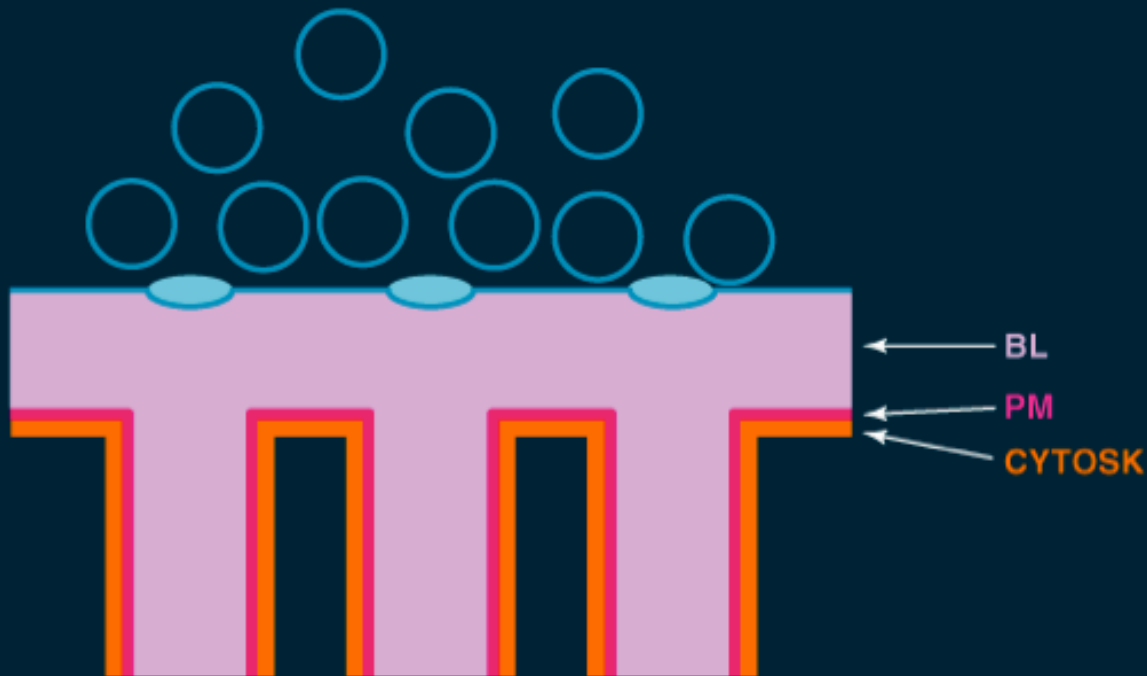


Rat



Man

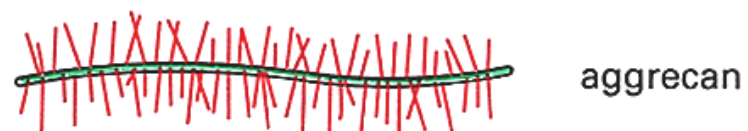
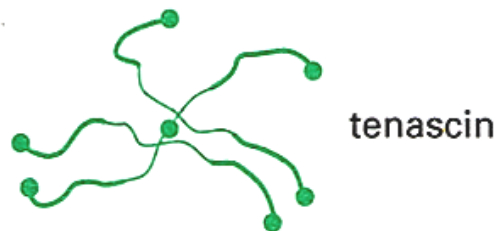
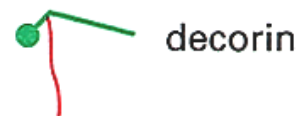
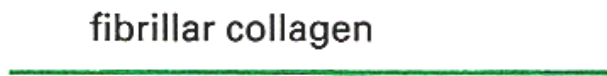
Axon-induced Postsynaptic Apparatus



Basal Lamina
 acetylcholinesterase
 agrin
 collagens
 heparin sulphate proteoglycan
 laminin A
 neuregulin
 s-laminin

Postsynaptic Membrane
 acetylcholine receptor ϵ
 neuregulin A
 erb B receptors 2, 3 & 4
 integrin
 MuSK
 N-CAM
 sodium channels

Cytoskeleton
 rapsyn
 vinculin
 talin
 paxillin
 filamin
 α -actinin
 tropomyosin 2
 58k protein
 87k protein
 utrophin
 acetylated tubulin
 ankyrin
 lamin B
 actin
 β -spectrin



100 nm

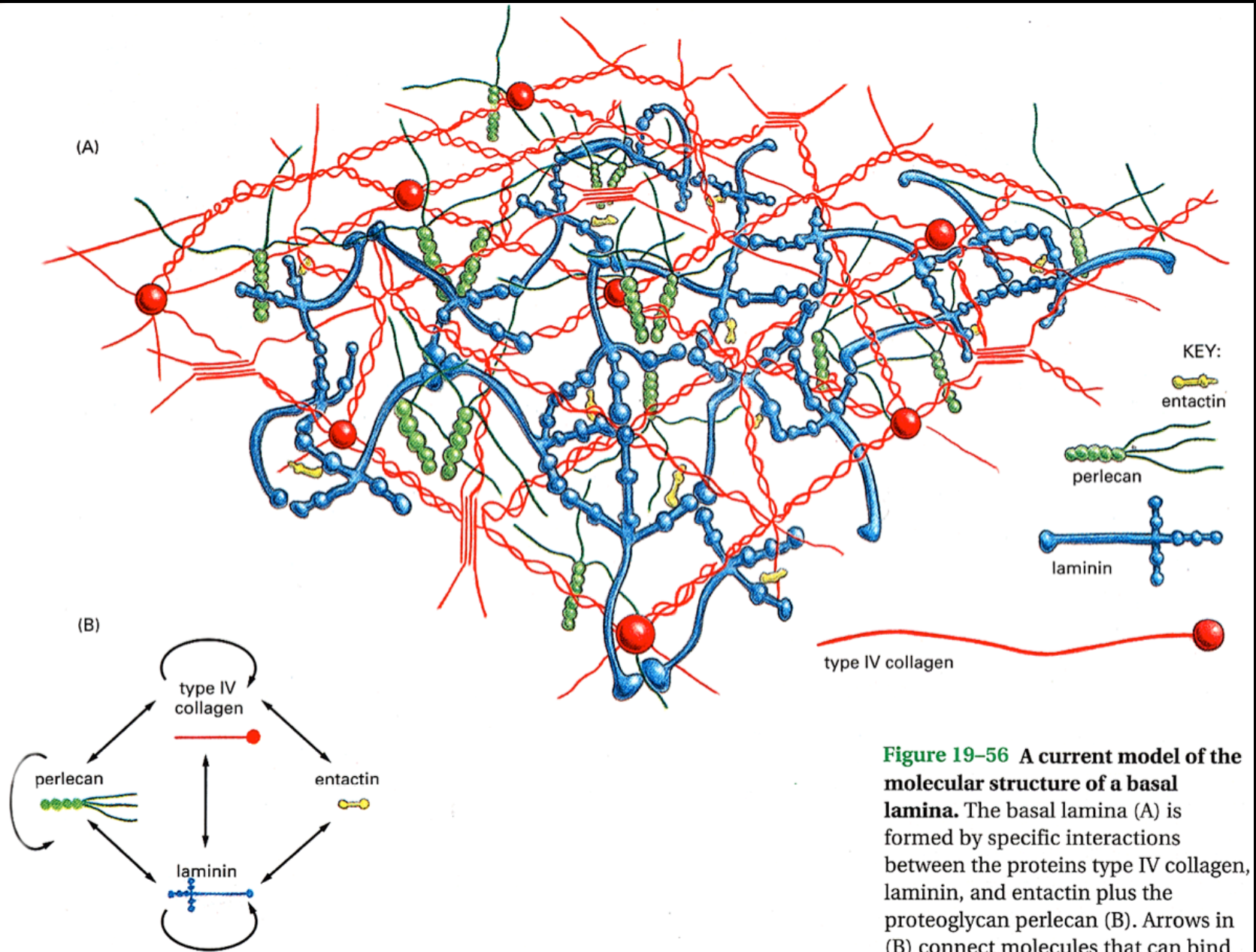


Figure 19-56 A current model of the molecular structure of a basal lamina. The basal lamina (A) is formed by specific interactions between the proteins type IV collagen, laminin, and entactin plus the proteoglycan perlecan (B). Arrows in (B) connect molecules that can bind directly to each other. (Based on P.D. Yurchenco and J.C. Schittny, *FASEB J.* 4:1577-1590, 1990.)

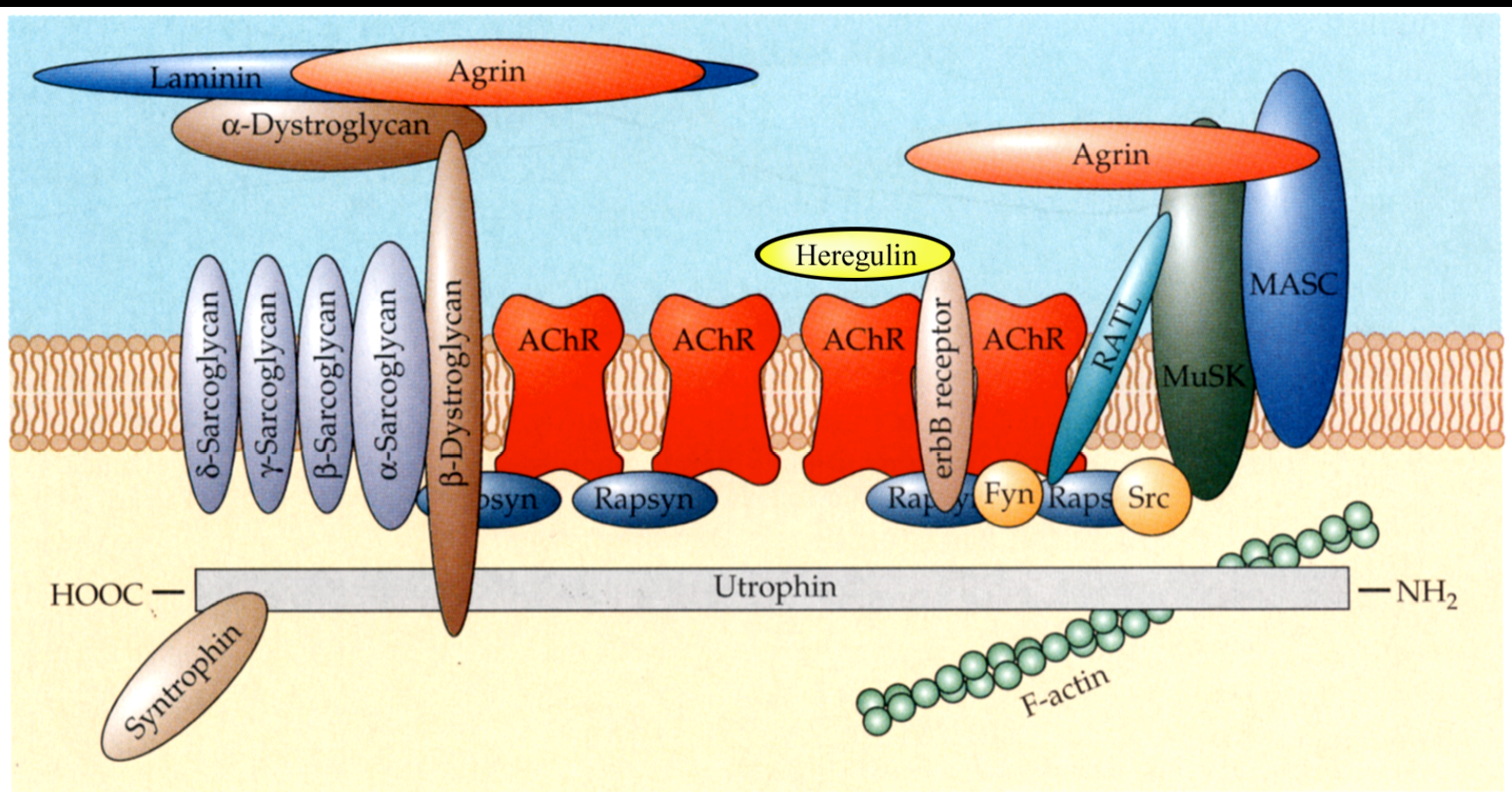
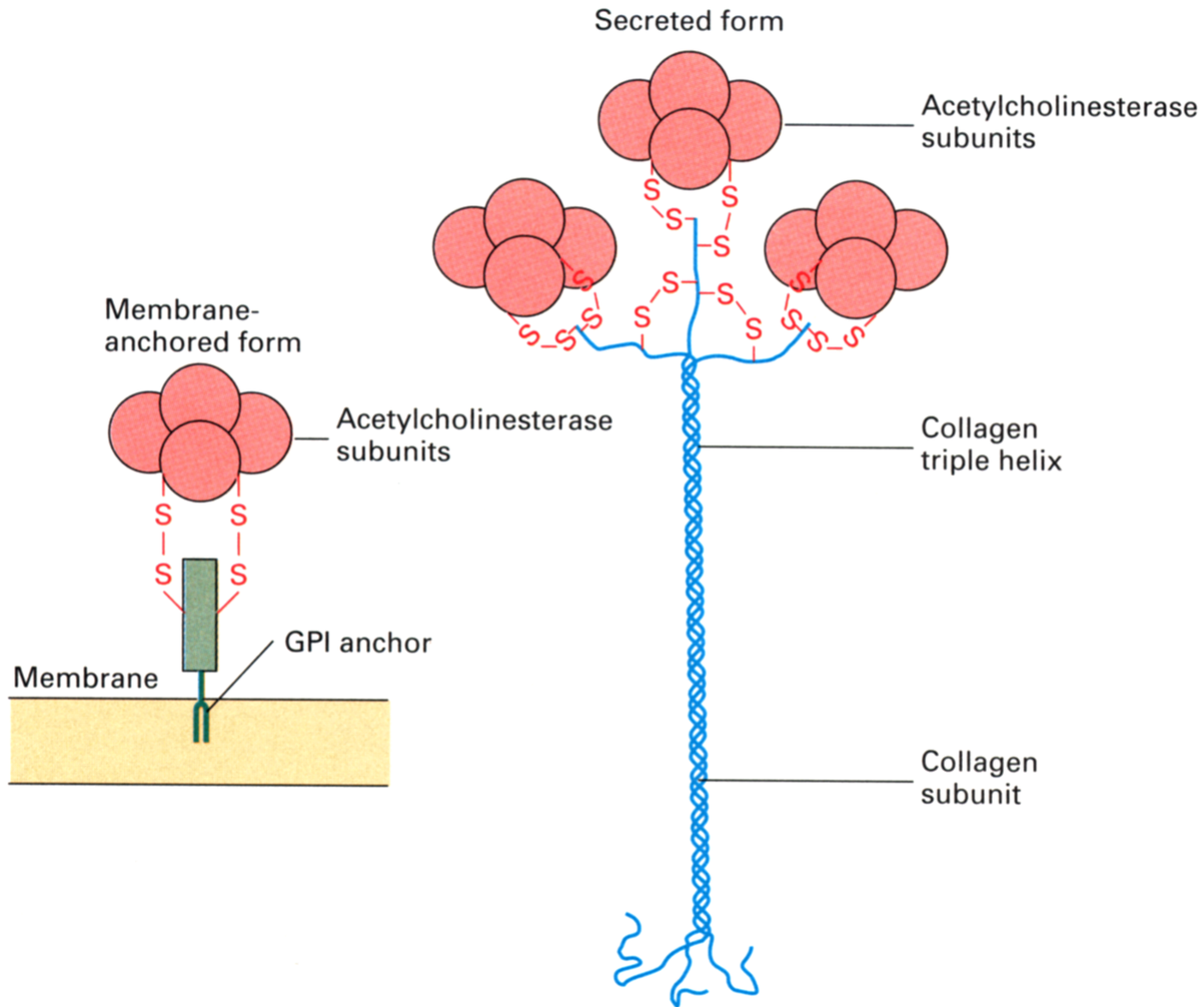
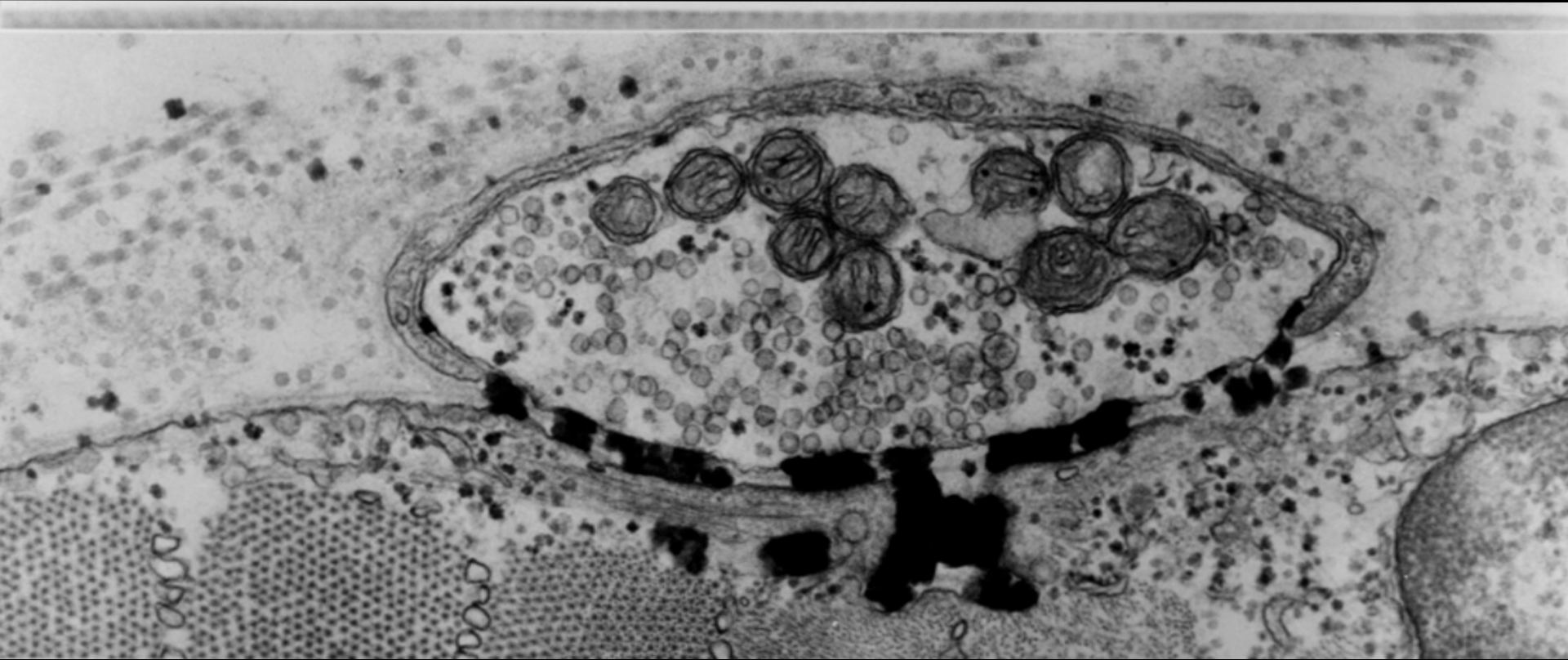
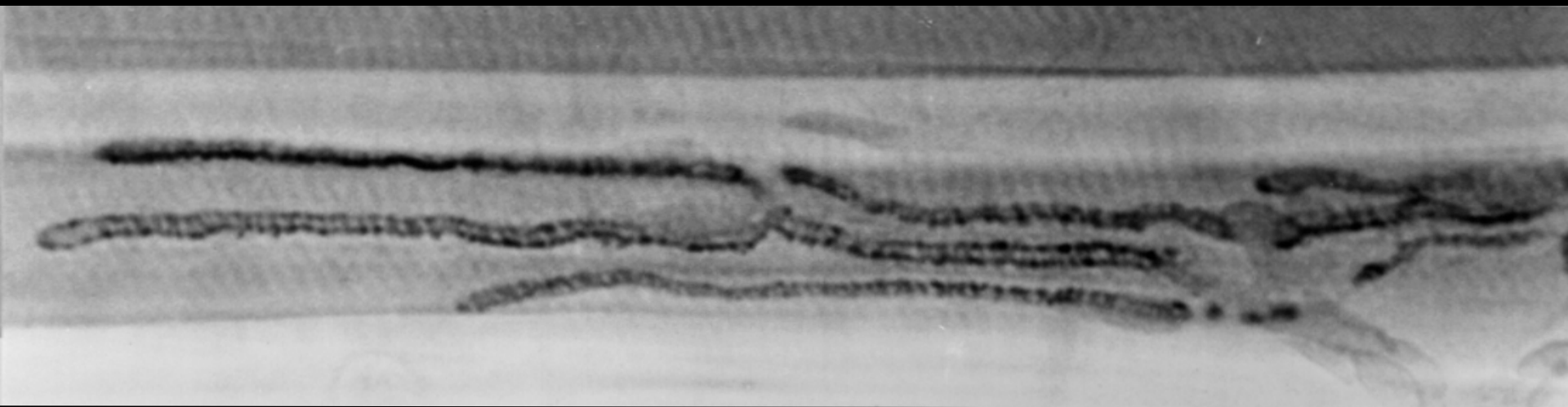
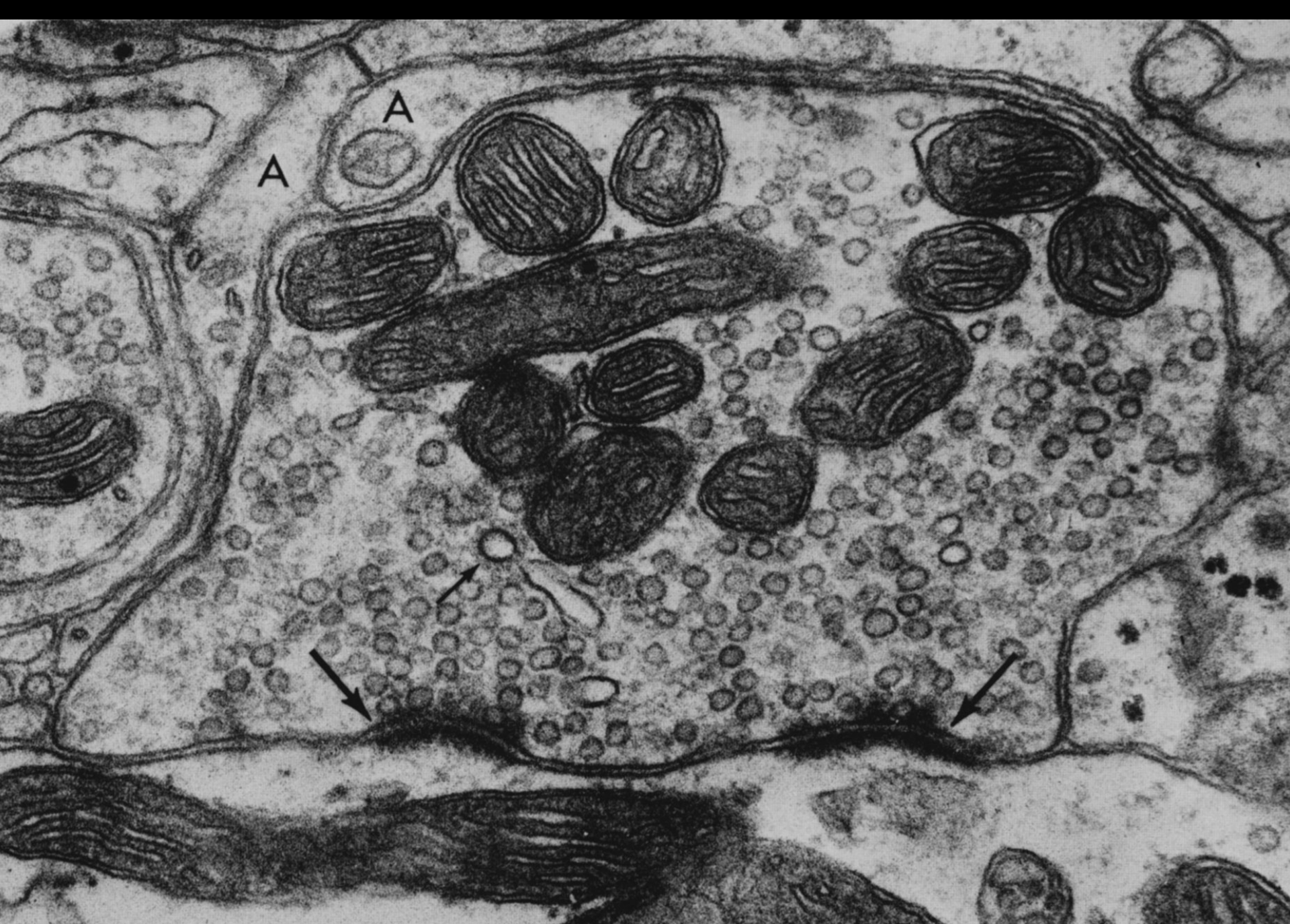


FIGURE 13.20 Postsynaptic Components of AChR-Rich Regions at the vertebrate skeletal neuromuscular junction. The dystrophin glycoprotein complex (utrophin, α - and β -dystroglycan, and the sarcoglycans) links together the actin cytoskeleton, the membrane, and the extracellular matrix. Agrin binds to laminin and α -dystroglycan and signals through the receptor tyrosine kinase MuSK to trigger formation of the postsynaptic apparatus during development (Chapter 23). Rapsyn plays a key role in linking MuSK and AChRs to the cytoskeleton. RATL and MASC are as yet unidentified components that mediate interaction of MuSK with rapsyn and agrin, respectively.

Principal Forms of Acetylcholinesterase



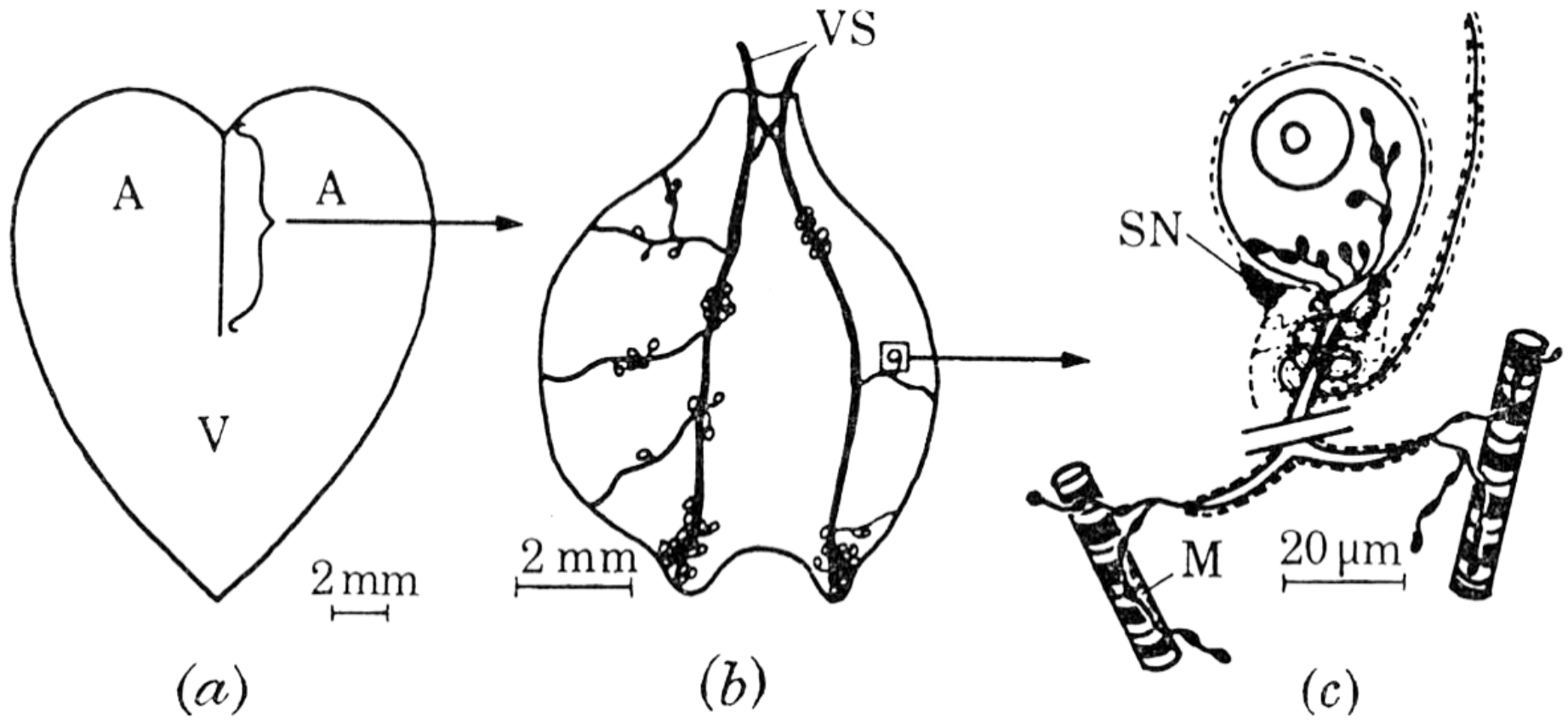


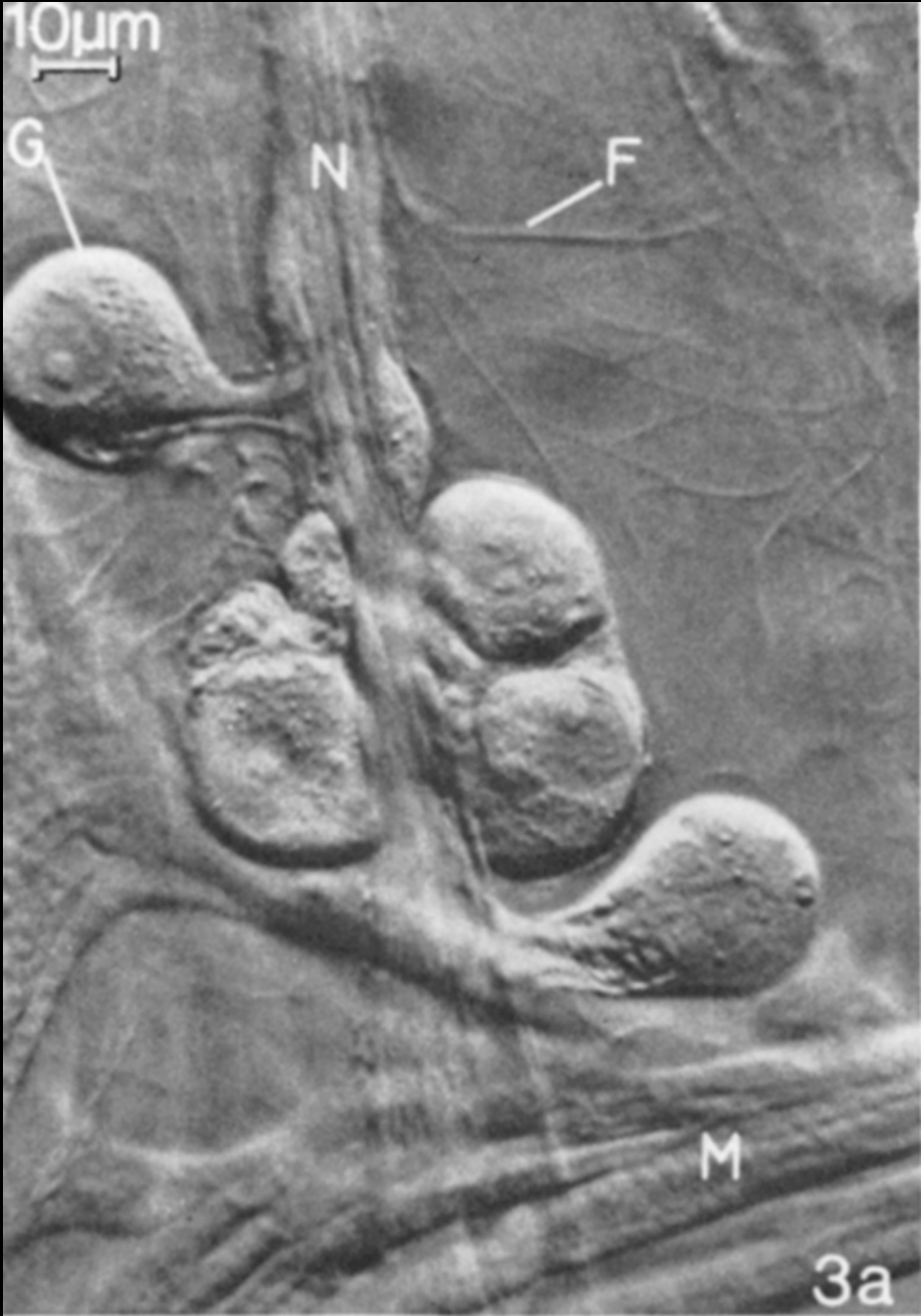


A

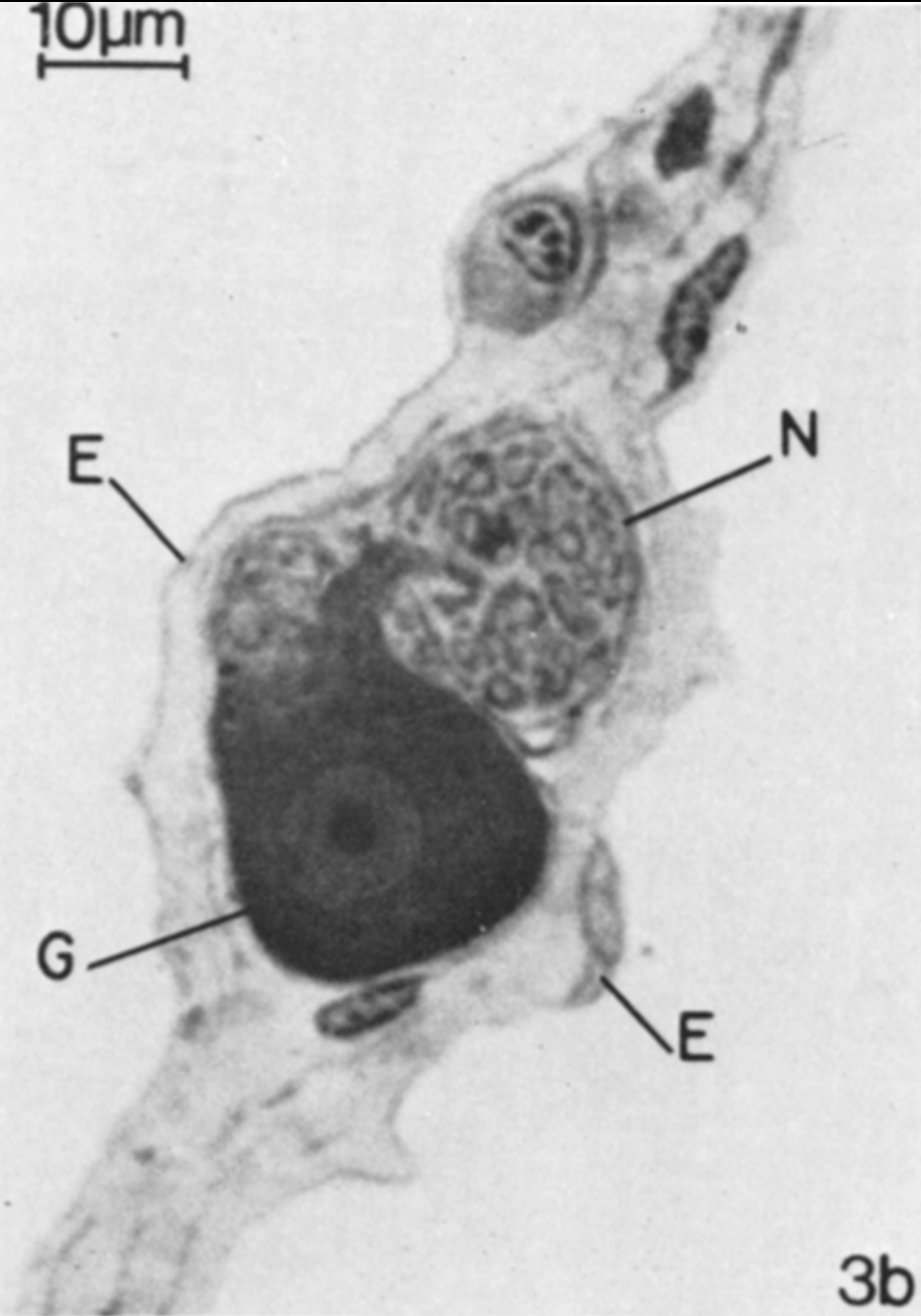
A



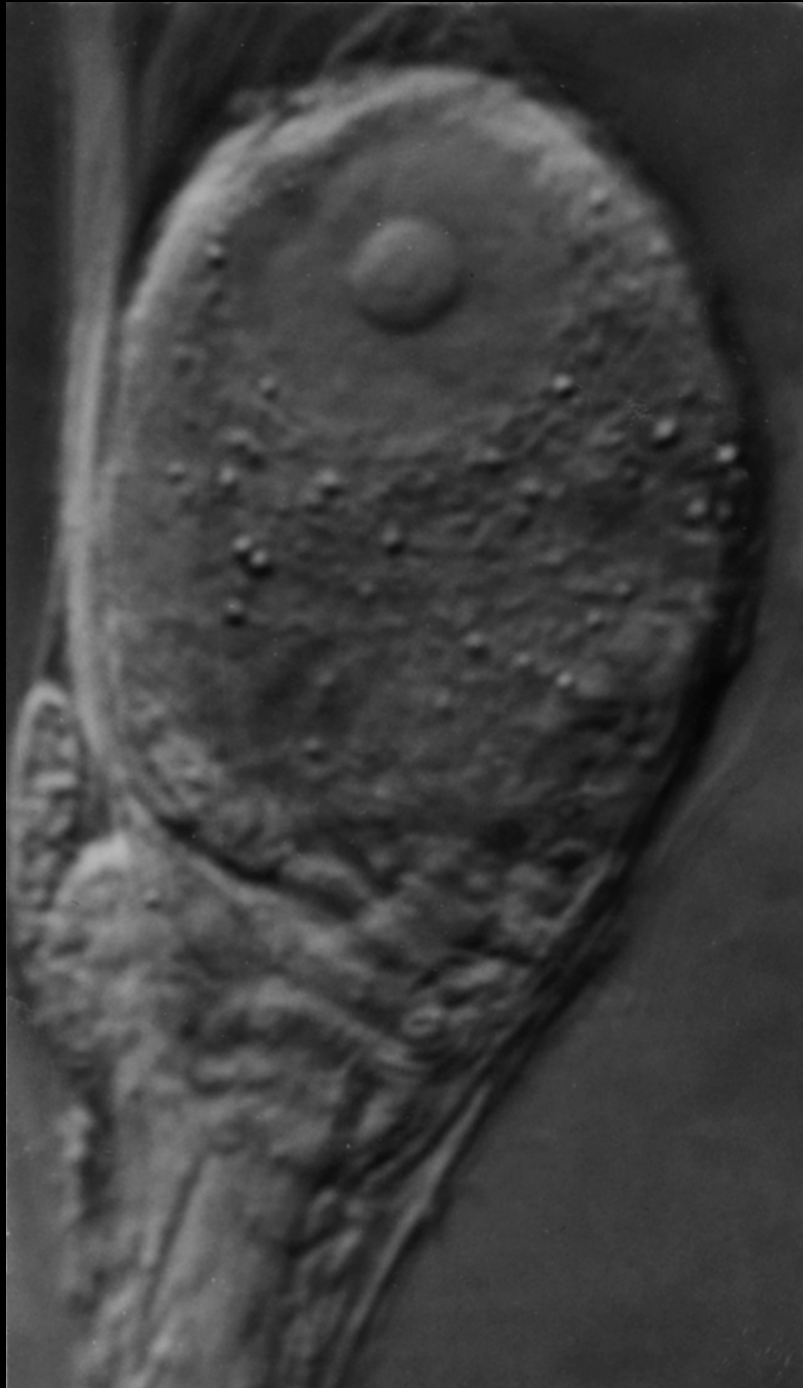




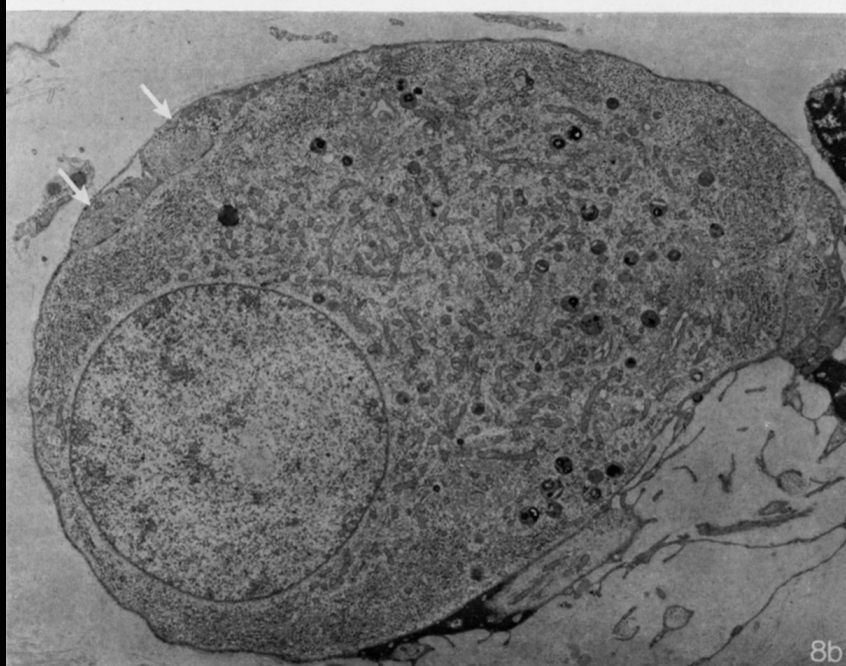
3a

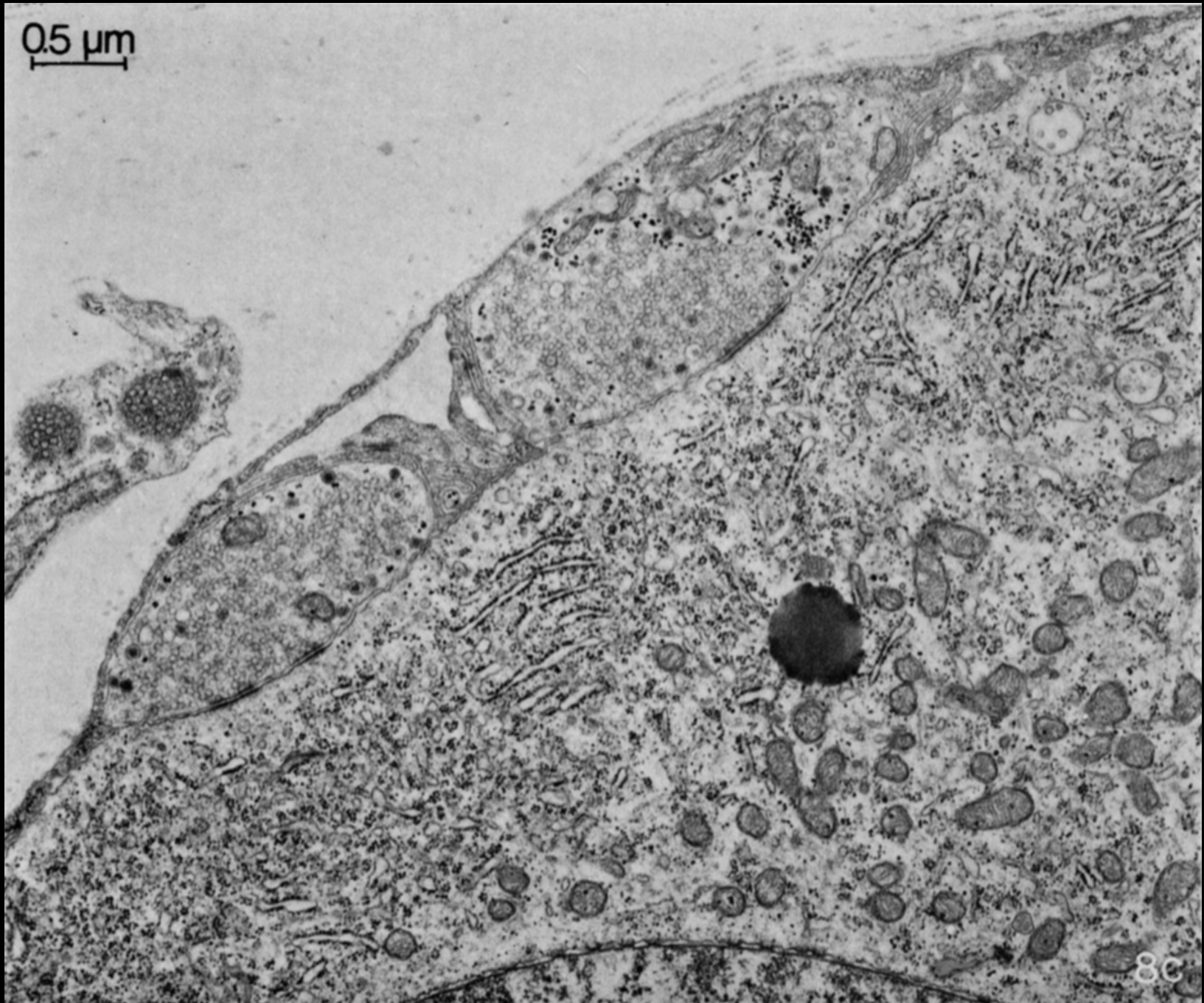


3b

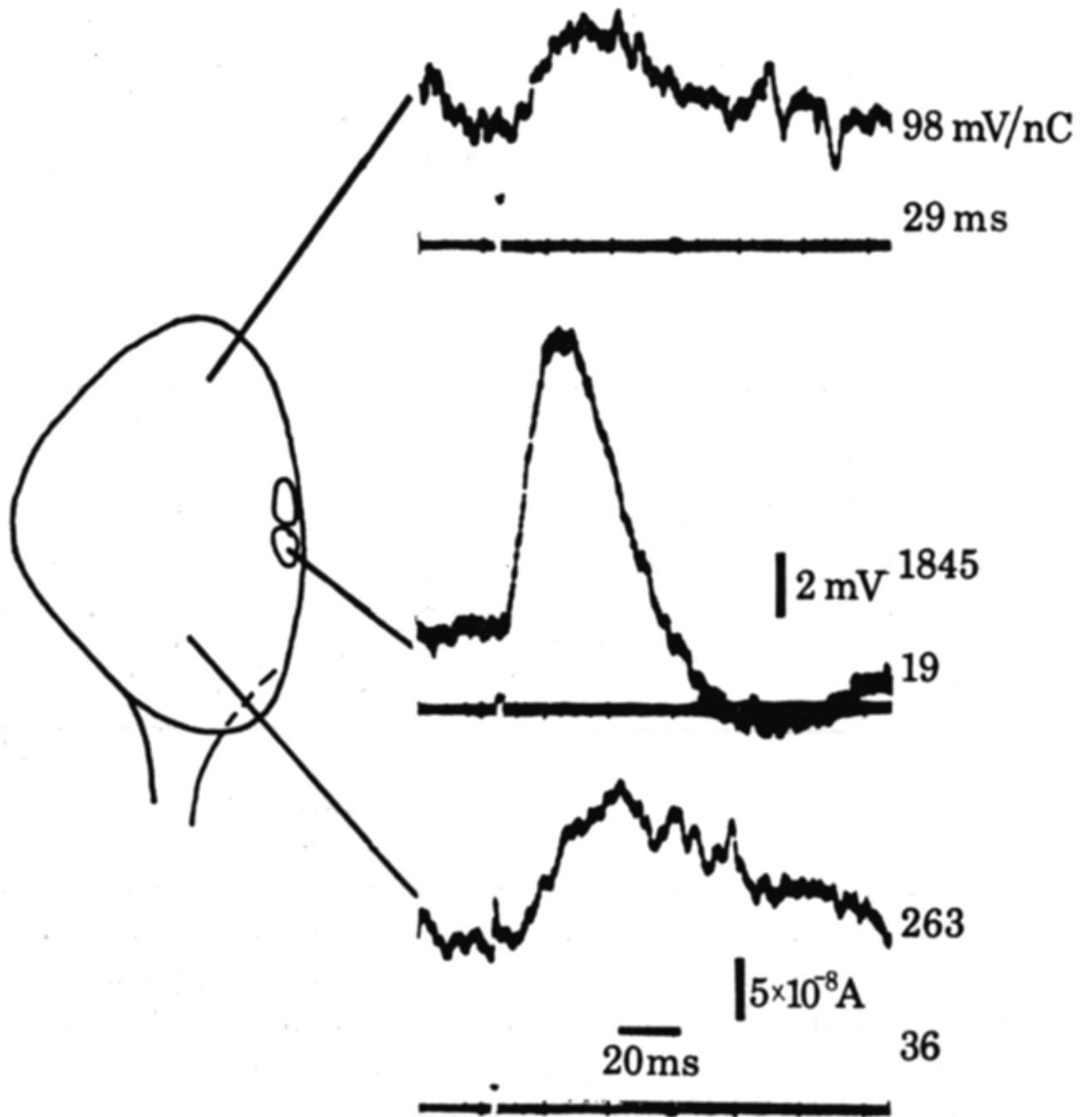


McMahan, U.J. & Kuffler, S.W. Proc. Roy. Soc. Lond. B. 177:485-508, 1971.





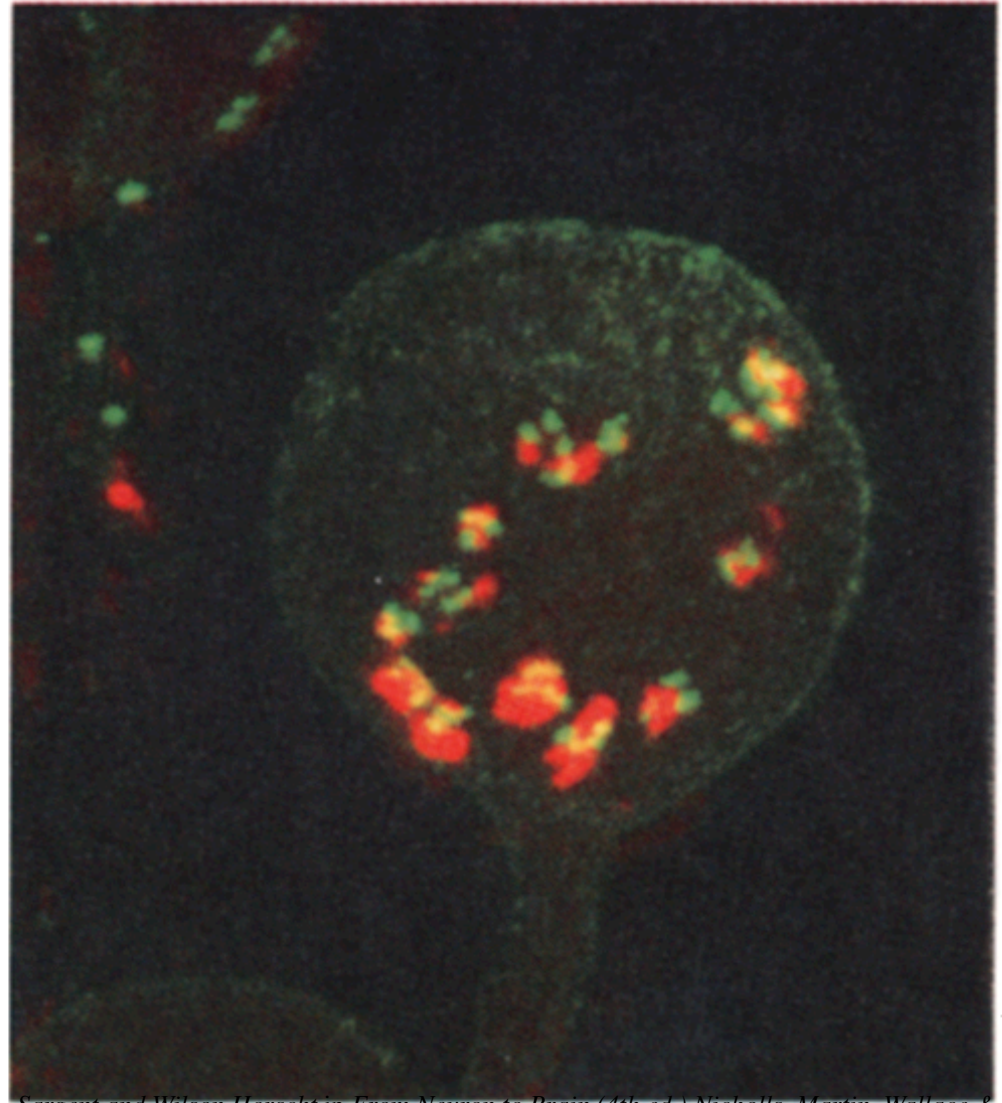
McMahan, U.J. & Kuffler, S.W. Proc. Roy. Soc. Lond. B.



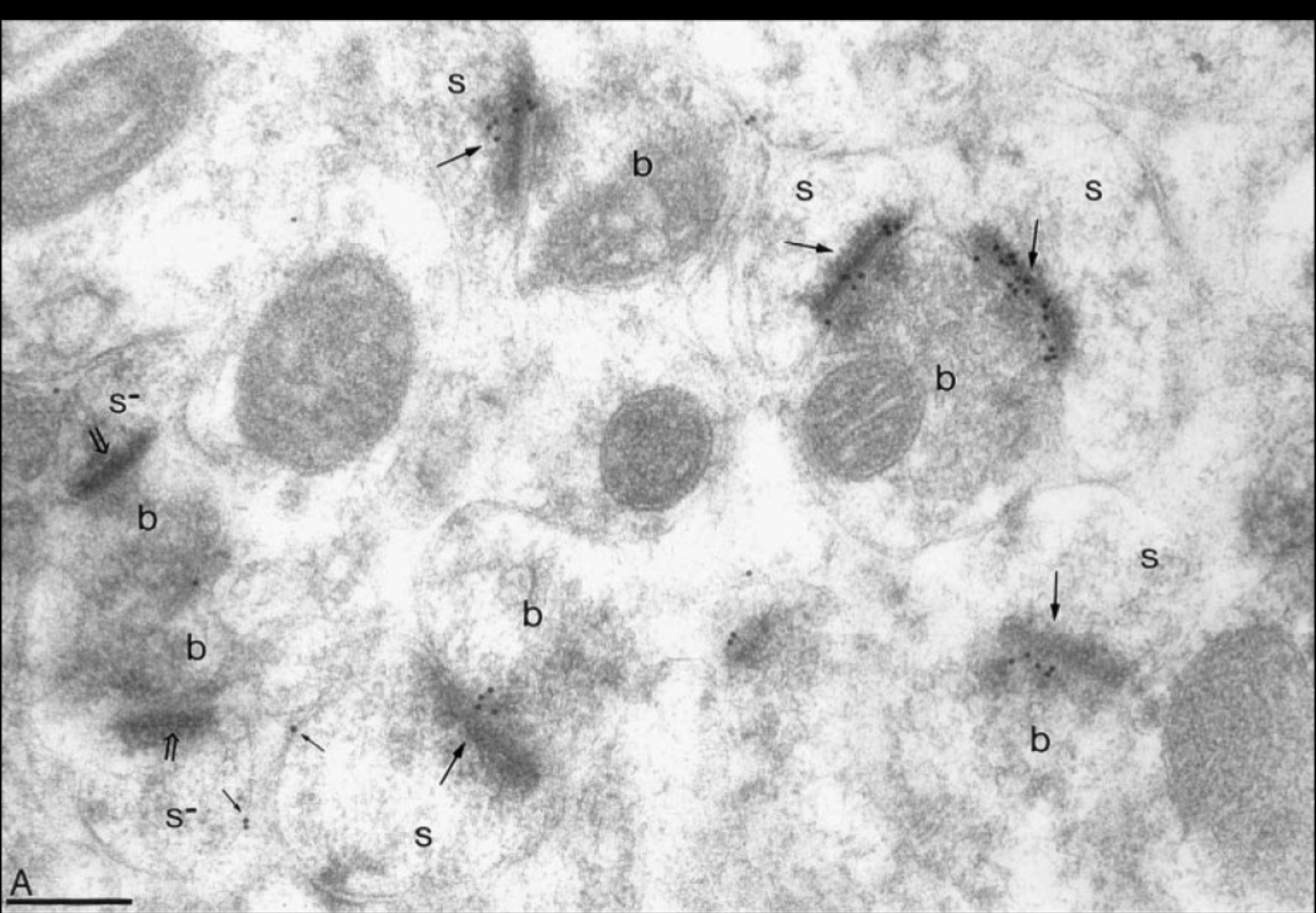
(A)



(B) Normal

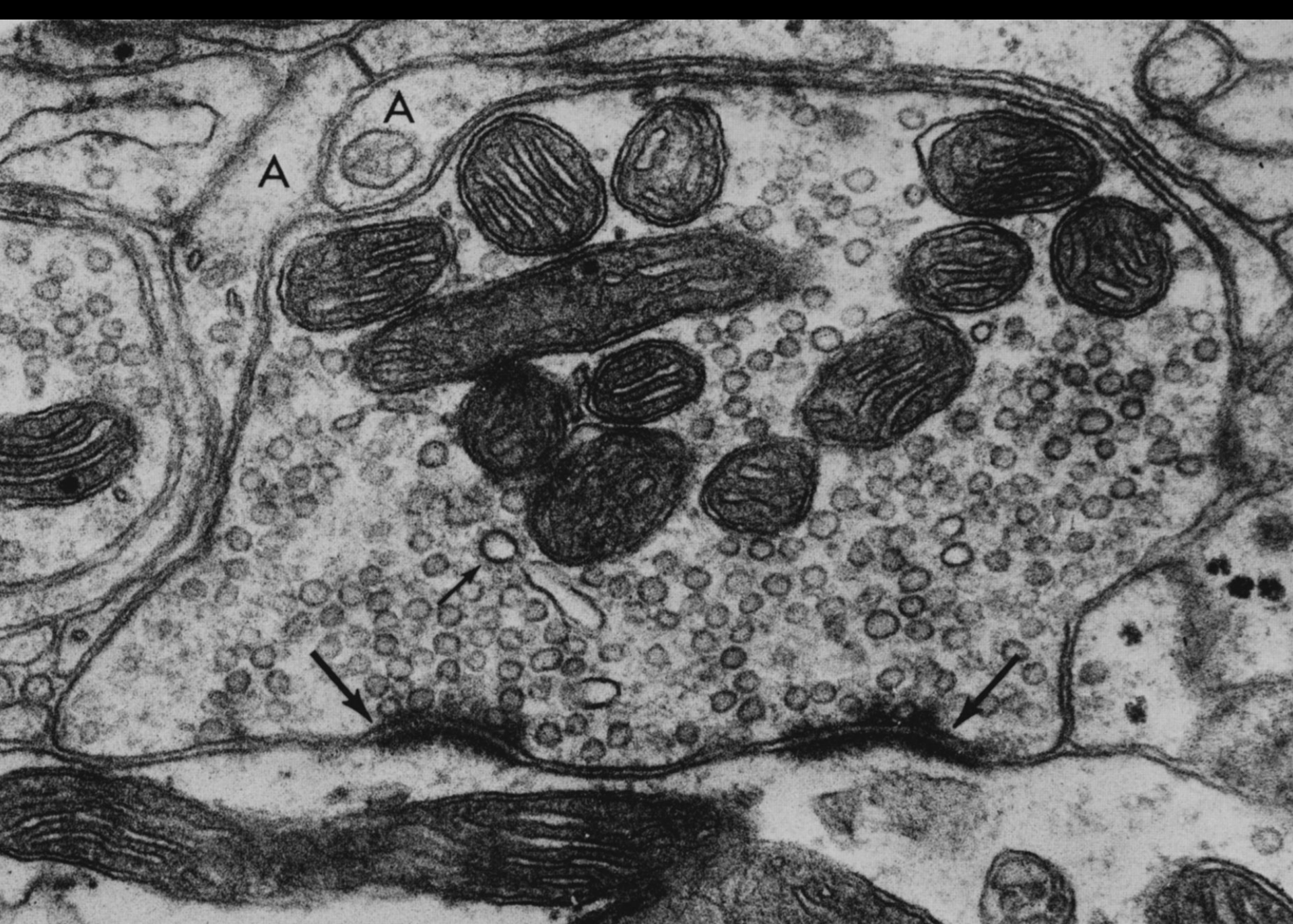


Sargent and Wilson Horscht in From Neuron to Brain (4th ed.) Nicholls, Martin, Wallace & Fuchs(2001), Sinauer Associates, Inc., Sunderland.

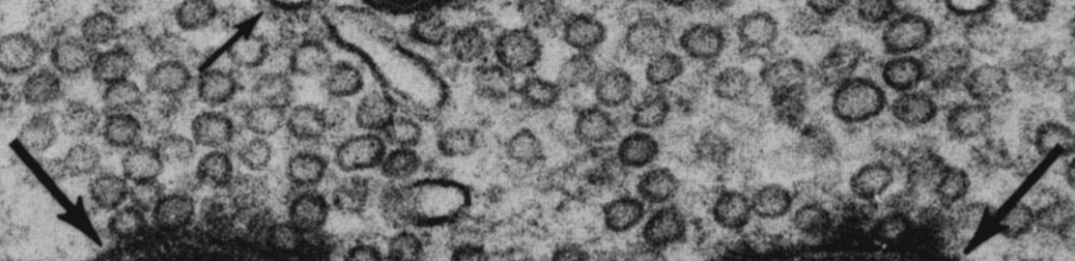




Landis & Reese, 1974, J. Comp. Neurol. 155: 93-125.



A A



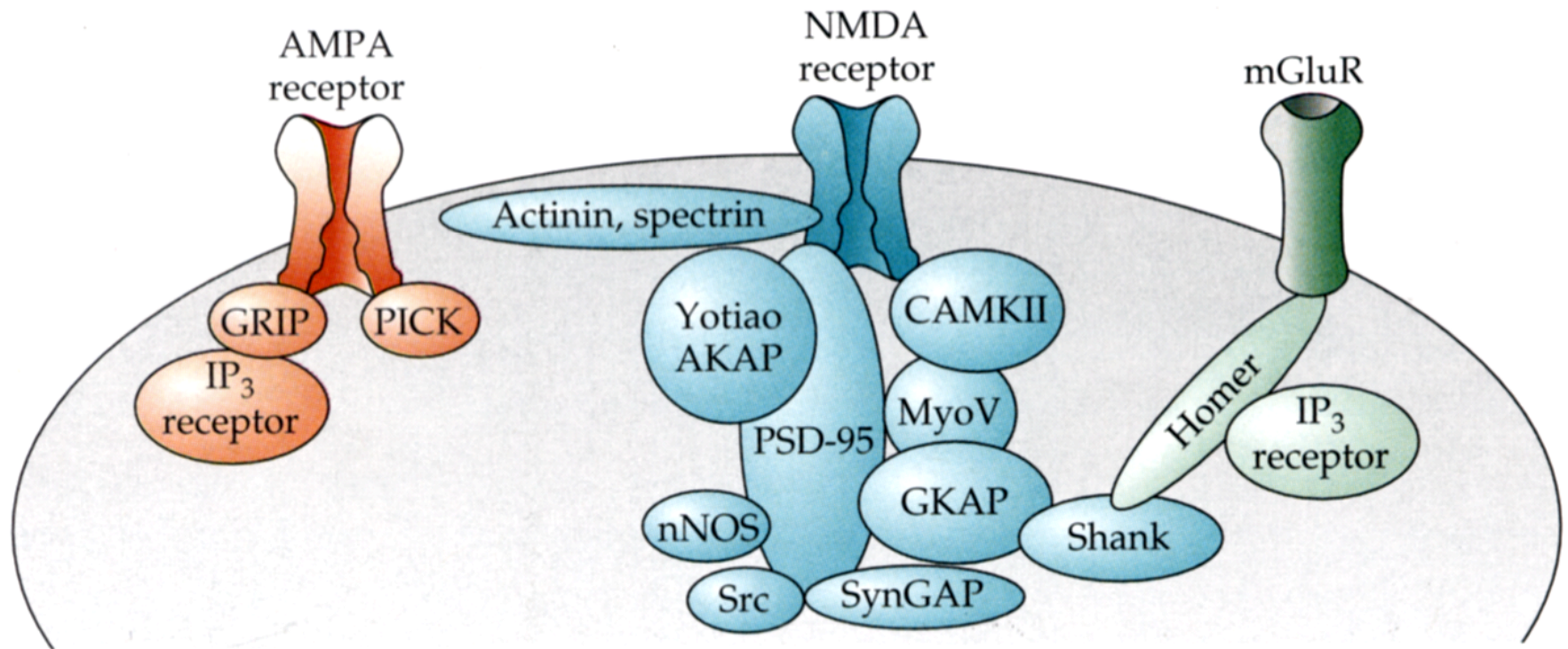
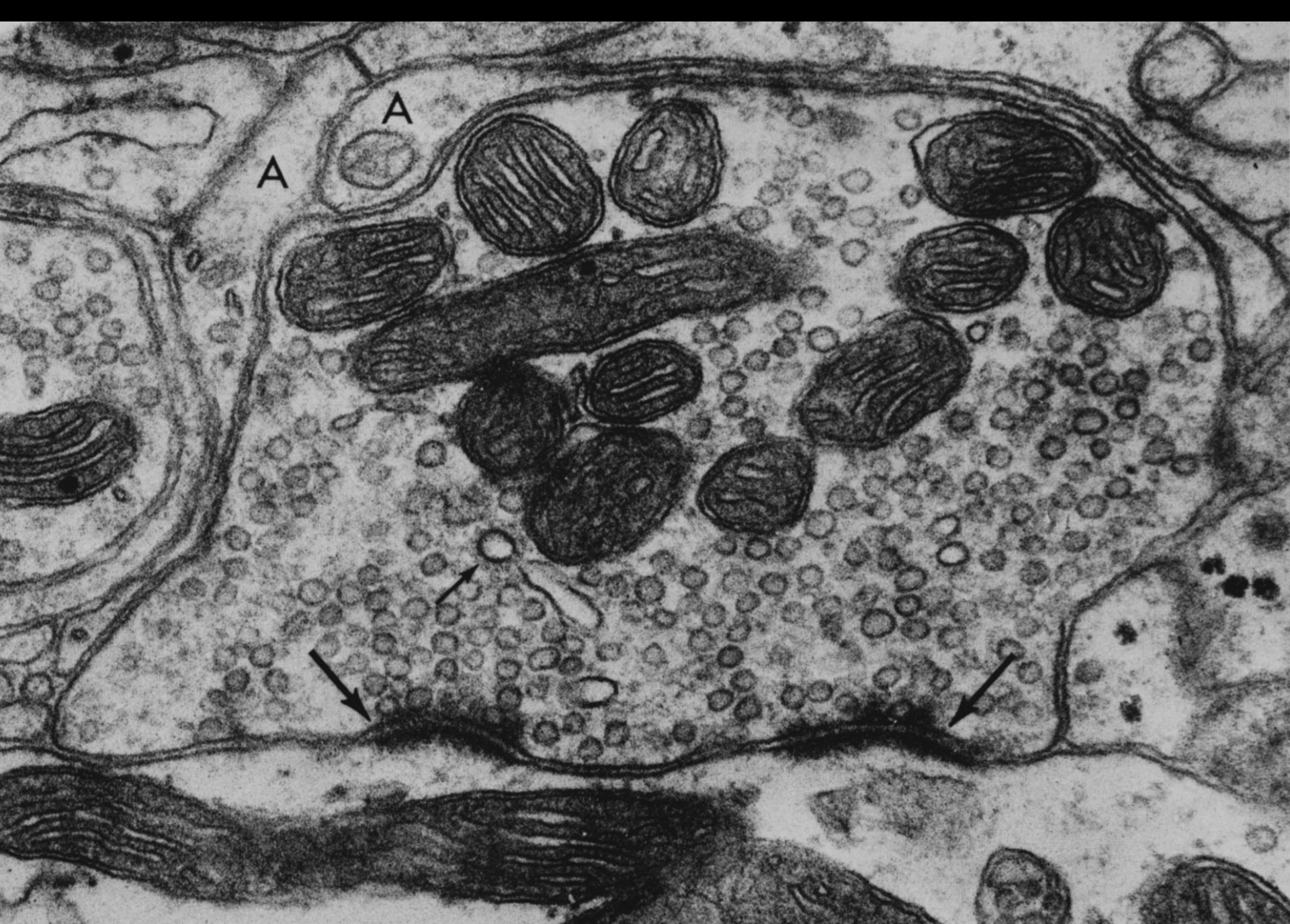
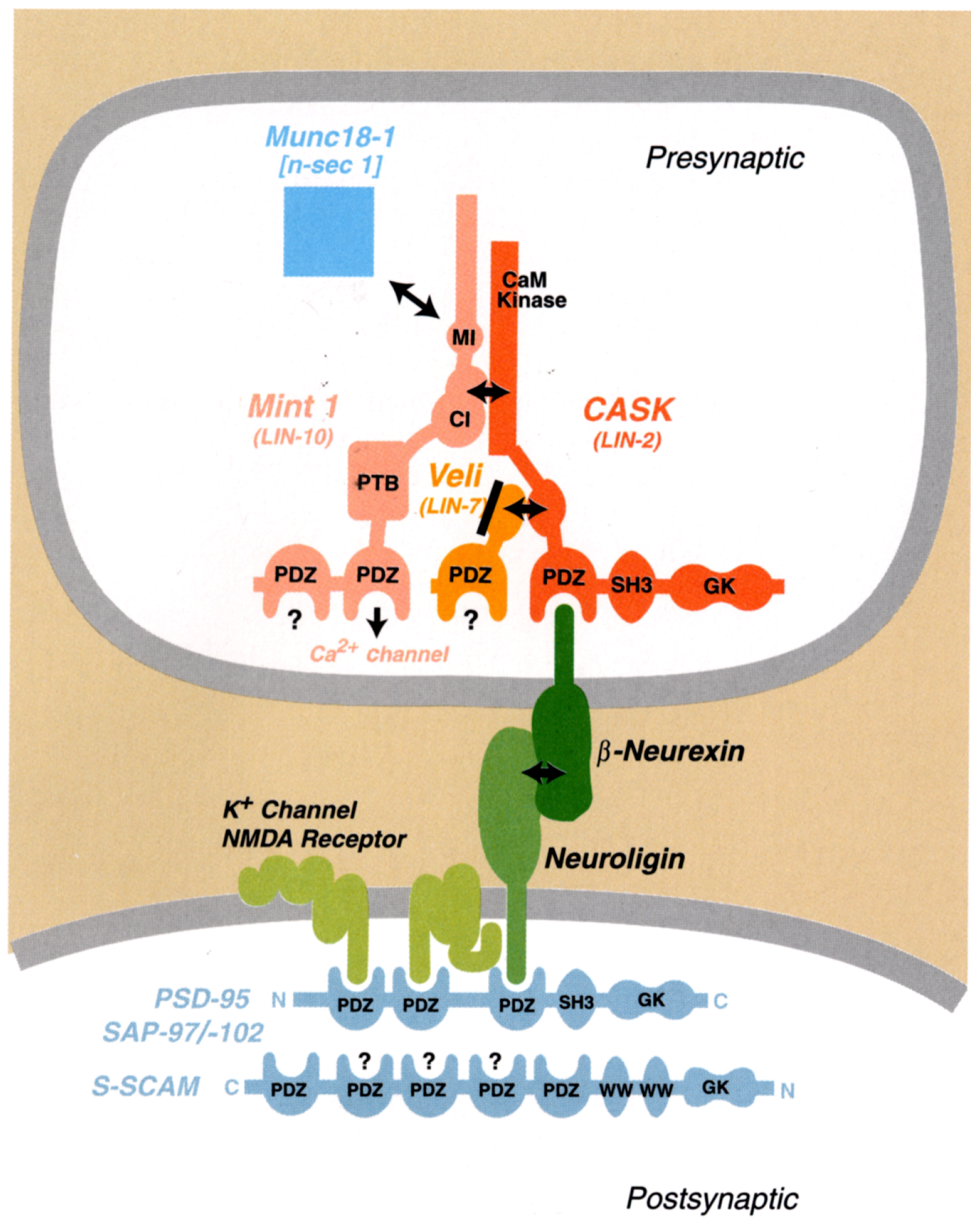


FIGURE 13.21 Glutamate Receptors Are Linked to a Postsynaptic Scaffold that includes proteins involved in intracellular signaling cascades. Members of the GRIP protein family link AMPA receptors to the IP₃ receptor. PSD-95 and its homologues connect NMDA receptors to Yotiao, nNOS, Src, SynGAP, and GKAP. CaMKII binds to NMDA receptors and to MyoV. The Homer protein family links metabotropic glutamate receptors to Shank and thereby to the NMDA receptor complex. (After Sheng and Lee, 2000.)





Südhof, T.C. (2001) *The Synaptic Cleft and Synaptic Cell Adhesion in Synapses* ed. by Cowan, Südhof & Stevens. The Johns Hopkins University Press/ Baltimore

Figure 6.6. Model of the intercellular junctions formed by the β-neurexin-neuroligin interaction and coated by PSD-95 and S-SCAM on the postsynaptic side and by CASK, mint 1, and velis on the presynaptic side. The localization of this junction to synapses is a model based on the presence of CASK, neuroligin 1, and PSD-95 at this structure. Modified from Butz et al. (1998).

