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Sonic hedgehog (Shh) regulates neural progenitor cells in the adult brain but its role in postmitotic mature neurons is not well understood. Using immunoelectron microscopy, we have recently demonstrated the postsynaptic distribution of Patched (Ptc) and Smoothened (Smo), the receptors for Shh, in hippocampal neurons of the adult rat brain. In this study, we describe the distribution of Shh protein in these adult hippocampal neurons. We find that Shh is present in both presynaptic and postsynaptic terminals. In presynaptic terminals, Shh is located either at the center or on the side of the synaptic junction. In postsynaptic terminals, Shh is mostly located on the side of the synaptic junction. We also find Shh in dendrites. Synaptic and dendritic Shh often reside in or are associated with vesicular structures that include dense-cored vesicles, synaptic vesicles and endosomes. Thus, our subcellular map of Shh and its receptors provides a foundation for elucidating the functional significance of Shh signaling in mature neurons.

Sonic hedgehog (Shh) plays at least two important roles in the nervous system. One is to stimulate the production of stem/progenitor cells, which include granule cell precursors in the young cerebellum\(^1\)-\(^3\) and neural stem cells in the specific brain regions of the adult brain.\(^4\)-\(^8\) The other is to promote axon growth of young neurons, which include spinal cord commissural neurons,\(^9\),\(^10\) retinal ganglion cells,\(^11\) olfactory sensory neurons,\(^12\) and midbrain dopaminergic neurons.\(^13\) Evidence has indicated that Shh and its signaling components also exist in mature neurons, neurons that do not have progenitor properties.\(^14\)-\(^16\) However, where Shh signaling takes place in these mature neurons and how Shh signaling affects them remain largely unknown.

We have recently described the subcellular distribution of Patched (Ptc) and Smoothened (Smo), the receptor and transducer for Shh respectively, in hippocampal neurons of adult rats.\(^17\) Multiple types of hippocampal neurons express Ptc and Smo, which are particularly concentrated in their dendrites, spines and postsynaptic terminals.\(^17\) Here, we studied the distribution of Shh protein within adult hippocampal neurons.

We used the same hippocampal tissue samples from adult rats that were used in our previous ultrastructural analysis of Ptc and Smo.\(^17\) We performed postembedding immunogold labeling (two animals) using monoclonal anti-Shh antibody (5E1; Developmental Studies Hybridoma Bank). The 5E1 antibody was generated against the N-terminus of Shh (aa 1–198 of rat Shh)\(^18\) and its specificity has been characterized.\(^10\),\(^15\),\(^18\),\(^19\)

Because of the preferential distribution of Ptc and Smo in the postsynaptic terminals of hippocampal neurons,\(^17\) we wondered whether Shh also was located near or at the synapse. We examined several hippocampal regions that include the CA1 stratum pyramidale and stratum radiatum, the molecular layer of the dentate gyrus, and the CA3 stratum lucidum. Synapses from these regions exhibited different morphological characteristics. Nevertheless, we found Shh labeling present in all of these synapses (Fig. 1). Figure 1A is an example of an inhibitory synapse based on its symmetric appearance.
with pit-like structures, which were situated opposite or across from the presynaptic Shh labeling (Fig. 1A).

Figure 1B–H show typical excitatory synapses based on their prominent postsynaptic density. In these synapses, presynaptic Shh labeling displayed a slightly different pattern from postsynaptic Shh labeling. Presynaptic Shh labeling could be found either directly at the membrane of the synaptic junction (Fig. 1B–D) or on the side of the terminal (Fig. 1H). Postsynaptic Shh labeling, on the other hand, was found mostly on the side, away from the center of the synaptic junction (Fig. 1E–G).

Figure 1I–M shows examples of mossy fiber synapses. As for other types of synapses, Shh was found in both the presynaptic mossy terminals and the postsynaptic thorny excrescences. Within the presynaptic mossy terminals, most Shh labeling was clearly seen associated with either dense-cored vesicles (arrows in Fig. 1J–M), or other smaller vesicular structures (arrowheads in Fig. 1I, J and M). Within the postsynaptic thorny excrescences, Shh was associated with tubulovesicular organelles in some cases (arrowheads in Fig. 1I), or positioned quite close to the postsynaptic membrane in other cases (arrowhead in Fig. 1L).

In addition to synaptic localizations, Shh labeling was found in various tubulovesicular organelles in the soma and dendrites of hippocampal neurons (Fig. 2). Interestingly, some of these Shh-labeled organelles made direct contact with the cell membrane surface, typically near contacts with adjacent processes (Fig. 2A–C).

Our previous immunoelectron microscopic work has described the postsynaptic localization of Ptch and Smo in adult hippocampal neurons. The present findings showing the presence of Shh in the presynaptic terminal, in particular localized in close vicinity to or even directly at the synaptic contact, raises the possibility that Shh signaling may occur across the synapse in these neurons. It is then tempting to ask what form of Shh protein is being released from the presynaptic terminal. In the photoreceptor neurons of the developing Drosophila retina, while the Hedgehog (Hh) C-terminus harbors the axonal targeting signal, the N-terminal

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Shh labeling was seen in the presynaptic compartment as well as in the postsynaptic compartment. Within both compartments, the Shh labeling was similarly distributed toward the side rather than the center of the synaptic junction (Fig. 1A). Upon closer examination, the postsynaptic Shh labeling appeared to be associated...
domain, and a small amount of the full-length Hh, also travel along the axon.20
Our results obtained using the 5E1 antibody—specific to the Shh N-terminus, could reflect the full-length as well as the N-terminal domain of Shh. To definitively identify the Shh forms that are present and possibly released from the presynaptic terminal will require further studies, including the use of the Shh C-terminus specific antibody.

We also observed an interesting subcellular distribution of Shh in postsynaptic spines and dendrites of hippocampal neurons. Several studies have shown that the dendrite and the postsynaptic terminal of neurons can release transmitters21 or growth factors.22 Moreover, studies of other Hh-producing cells have shown that the release of Hh occurs on the apical and the basal side of the Drosophila photoreceptor neurons.20 Likewise, in Drosophila wing disk epithelium, Hh is found in the apical and basolateral plasma membranes.23 It will be interesting to investigate whether Shh in mammalian hippocampal neurons is also released from both pre- and post-synaptic sites. The mapping of Shh protein and its receptor at the subcellular level will advance the understanding of where and how Shh signaling occurs, and what roles Shh plays in the function and plasticity of mature neurons.

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Figure 2. Shh (arrowheads; 10 or 15 nm gold particles) is found in tubulovesicular organelles that are located in soma (so) or dendrites (de). These Shh-containing organelles often make direct contact with the membrane surface of the neuron, near adjacent cell processes (pr in A–C). (A) the CA1 stratum pyramidale; (B) the CA1 stratum radiatum; (C) the molecular layer of the dentate gyrus; (D) the CA3 stratum lucidum. Scale bars are 100 nm.