Cytoarchitecture and Connectivity of the Superior Colliculus in Mouse Brain

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Abstract: Superior Colliculus (SC) plays a vital role in visual target selection and attention shifting; it also an important structure in studying the central nervous system. However, the cytoarchitecture and connectivity of the SC in mouse brain have not been explicitly explored. In this paper, to investigate the structural delineations and connectivity SC, we first delineated the morphology of the SC by Nissl stain, and further explored different genes expressed in the SC. It demonstrates that gene Tpd52l1 is densely expressed in the SC, which helps describe the borders of the SC. In addition, we explore the connectivity of Superior Colliculus (validating the projections from other structures to SC and examining the projections from SC to other structure). The anterograde and retrograde projection circuitries between SC and IC are particularly addressed, which indicates that the SC is involved in the visual, auditory and other somatosensory physiological activities.

Keywords: Superior Colliculus, Cytoarchitecture, Connectivity, Gene Expression, Mouse Brain

1. Introduction

The SC (Superior Colliculus) is a superstructure of the mammalian midbrain quadrigeminal body [1-2]. It is also known as the optic tectum, or simply “tectum” in vertebrates. The SC is a relay nucleus in the mammalian visual pathway and the primary target of the primary visual projection. It is not only an important reflex center of the visual pathway, but also involved in many physiological activities, such as auditory, somatosensory and other activities [3-5]. Therefore, the SC is considered to be an essential structure in the study of the central nervous system.

The SC is structurally divided into seven layers in mammals: the zonal layer (ZO), superficial gray layer (SGR), optic layer (OP), intermediate gray layer (IGR), intermediate white layer (IW), deep gray layer (DGR), and deep white layer (DW) (figure 1). The ZO, SGR and OP are three superficial layers (SL) which receive most of the afferent fibers from the retina and the visual cortex and are mainly involved in visual information formation. While IGR, IW, DGR, and DW are four deep layers (DL), which receive visual projections from cortical and subcortical regions, they also receive auditory and somatosensory projections. At the same time, it also issues motion instructions and controls the eyes and ears and head toward the significant stimuli in animals [6-9]. Therefore, the SC is considered recognized as a visual motor integration center of the central nervous system. The superficial layers receive direct visual inputs, but the deep layers receive indirect visual inputs [10-11]. The superficial layer and deep layer cells in morphology differ with respect to two main points: the superficial layer cells have short and dense but not overlapping dendrites, similar to the structure found in the perception of the neurons, with significant differences compared to the relatively long and overlapping neuronal dendrites in the deep layer. The second is that the difference of superficial layer neurons in cell size is not as significant as deep layer neurons [12].

The origin of the superior colliculus fibers is complex, there are many studies about this, and the afferent fiber mainly of SC can be classified into two categories. One is related to the visual function of the nucleus, called visual structure, and the other is independent of the visual function of nuclei, called non-visual structure. There are several different types of projections in visual and non-visual structures. The projections of visual structure include the projection from the retina to SC (retinal
ganglion cells projecting to the superior colliculus are one of the largest sources of afferent fibers in the superior colliculus [13-15], the projection from the visual cortex to SC (in mammals, the visual cortex is generally divided into the primary visual area (area 17), the secondary visual area (area 18) and the third area (area 19). The primary visual area is mainly reacted to the contrast of retina graphics and movements, secondary visual area and the third area is processed, analyzed and integrated the visual information [16-17]. The lateral geniculate nucleus is a relay station of the visual pathway in the thalamus, the projection from lateral geniculate nucleus to SC is composed of several layers of cells and fibers. The projections of the non-visual structure include the projection from parabigeminal nucleus (Pbg) to SC, the projection from Substantia nigra (Sn) to SC, the projection from Inferior colliculus (IC) to SC and the projection from the dorsal nucleus of the lateral lemniscus (DII) to SC [18-19]. The projections from the efferent fibers include the projection from the SC to the lateral geniculate nucleus (LGN), the projection from the SC to Pbg, and the projection from the SC to the auditory cortex [20-22]. Additionally, the superior colliculus also sends fibers to the nucleus of the Sn, the anterior tectum, the superior nucleus of the thalamus, the nucleus of the thalamus, and other structures. Moreover, there is a complex fiber connection between both sides of the colliculus.

The purpose of this study is two-fold. First, the study aims to investigate the structural delineations of SC, especially the sub-regions which can reflect the functions. Secondly, this paper validates the projections from other structures to SC and examines the projections from SC to other structures. The hypothesis is that gene expression in the SC can also be found in other structures to illustrate the role the current gene. In addition, the other hypothesis is the projections of SC can demonstrate the connectivity between SC and different structures.

2. Method

To delineate the cytoarchitecture of SC, two public online sources are used: the Allen Brain Atlas (ABA) and the Mouse Connectome Project [23]. The brainmaps.org website is more used to explore sub-regions and the gene expression in the SC, and mouseconnectome.org website is used to find connectivity information about SC.

2.1. Nissl Staining

Based on the Allen Brain Atlas, six coronal slices (83, 86, 88, 91, 94 and 103) and one sagittal slice representative images have been selected to illustrate the different regions in the SC.

2.2. Gene Expression

Because not all the listed gene in the ABA can be clearly observed in the SC, eight representative genes (Gabbr1, Tpd52l1, LOC433228, Mgl2, Lats2, Dgkh, Cpn5 and LOC433258) are selected.

2.3. Connectivity

Both the ABA and the mouseconnectome website are used to explore the connectivity, and several tracers are used including AAV, Ctb555, PHAL and fg tracer.

3. Results and Discussion

3.1. Cytoarchitecture

As noted from figure 1 and figure 2, the different regions of the SC in the coronal view can be clearly distinguished. In addition to the seven layers in the SC, it shows that the SC can be divided into sensory related (SCs), motor related (SCm) and two fiber tracts (brachium of the superior colliculus (bsc) and superior colliculus commissure (csc)). The sensory related structures in SC can be subdivided into ZO layer, SGR layer, OP layer, IGR layer, IW layer, DGR layer, and DW layer. The motor related structures in SC can also be classified in to IGR layer, IW layer, DGR layer, and DW layer, which are also called deep layers.

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3.1.1. Nissl Staining

Table 1 shows the location and cytoarchitecture of SC. Six coronal slices (83, 86, 88, 91, 94 and 103) and one sagittal slice representative images have been selected to illustrate the different regions in SC. Each slice contains two images; the left is named SC sub-regions and the right is the corresponding Nissl stain images. The purple highlight on the right Nissl stain images is the region which can be identified easily. In addition, the manual delineated lines are the boundaries between different sub-regions. The Nissl stain in slice 83 can hardly distinguish the different sub-regions in the SC; the slice 86 and 88 can easily find the ZO layer in the superficial layers; the slice 91, 94 and 103 can identify more regions, especially slice 94, which can substantially identify all the sub-regions in the SC. It is impossible to identify the seven sub-regions in the sagittal slice 16, since these sub-regions in SCs and SCM are completely compacted with each other.

The superior colliculus is not a homogenous structure. The superficial layers (ZO, SGR and OP) receive information from visual systems, while the deep layers (IGR, IW, DGR, and DW layer) receive visual, auditory, somatosensory signal from cortex, and inputs from the frontal eyes fields. Therefore, the SC is almost multimodal and associational [10-24]. As shown in the six slices, the superficial layers are significantly different from the deep layers. The area of deep layers is obviously larger than the superficial layers, in that they are more pronounced with a higher cell density in the deep layer compared to the superficial layers. These two layers can be divided by the OP layer which is a slender region. The deep layers are compacted below the OP layer. The sub-regions in SC are actually difficult to distinguish either the superficial layers or the deep layers; only a small number of Nissl stain image can be identified, for example, slice 83 is difficult to identify the seven layers in the SC. Besides, two sub-regions (bsc and csc) are not further explored, since these two fibers tracts are not the region of interests (ROIs). In the slice 86 and 88, the first layer ZO can be facile identified, since this layer is just below the Retrosplenial area (RSP). The OP and DW layers have relatively low cell density than the other 5 layers. In addition, SGR has the highest number of the cell density among these regions.

3.1.2. Gene Expression

Table 2 illustrates the distinctively expressed gene in the SC. Eight genes have been selected to show the differentiation in SC. There are 21 genes in the mouse.brain-map.org, but some of the gene expression is not visible in the images, so eight representative genes (Gabrr1, Tpd52l1, LOC433228, Mgll, Lats2, Dgkh, Cqne5 and LOC433258) are selected. Four genes (Tpd52l1, Dgkh, Cqne5 and LOC433258) are densely expressed within the SC, and the other four genes are sparsely expressed in the SC.
The name of Tpd52l1 gene is tumor protein D52-like 1, it also can be found to be highly expressed in thalamus, and it is a protein coding gene. As shown in table 2, the Tpd52l1 gene has a number of dense black dots in the IGR layer, which means that there may be more cell proliferation in IGR layer [25]. The name of Dgkh gene is diacylglycerol kinase Eta. It is also a protein coding gene which encodes for the diacylglycerol kinase (DGK) enzyme and is densely expressed in the OP layer (table 2), and densely expressed in several structures (Isocortex, Olfactory areas, Hippocampal formation, Cortical subplate, Striatum, Thalamus, Midbrain and Cerebellum). It has been observed that the DGKH gene is associated with several complex diseases include Bipolar Disorder and Schizophrenia. Therefore, there are some

| Gene      | Slices          |
|-----------|-----------------|
| Gabbr1    | Slice: 39, 41, 43 |
| Tpd52l1   | Slice: 71, 74, 80 |
| LOC433228 | Slice: 70, 75, 77 |
| Mgl1      | Slice: 36, 39, 42 |
| Lats2     | Slice: 37, 39, 40 |
| Dgkh      | Slice: 37, 39, 41 |
| Cpne5     | Slice: 54, 56, 61 |
| LOC433258 | Slice: 34, 36, 39 |

**Table 2.** Gene expression of the SC (Macaca mulatta), the left column is the expressed gene and three different slices about that gene in the SC region.
changes in the structures that the DHKH expressed when the disease symptoms are manifested [26]. The name of CpnE5 gene is copine V, it is one of the genes which encodes the calcium-dependent protein, a protein that can regulate molecular events and play role in calcium-mediated intracellular processes and the formation of the dendrites (RefSeq, Sep 2015). The gene CpnE5 is expressed in the OP layer and can also be found in other structures (Isocortex, Olfactory areas, Hippocampal formation, cortical subplate, and Pallidum) which have calcium-dependent protein. The name of LOC433258 is attractin-like 1. It is highly expressed in OP and SGR layers, and can also be found in other structures including Isocortex, Olfactory areas, Hippocampal formation, Cortical subplate, Thalamus, Midbrain. For the other four genes (Gabrr1, LOC433228, Mgl2 and Lats2) that are sparsely expressed in the SC also play an important role in the protein expression. For example, the Gabrr1 (gamma-aminobutyric acid (GABA) C receptor, subunit rho 1), a locus for GABA receptor subunit encoding, has been identified to be slightly expressed in the superficial layers. GABA is the major inhibitory neurotransmitter in the mammalian brain. On the whole, the superior colliculus has a relative higher volume of GABA, while the GABA volume in SGS layer is highest in the central nervous system. GABAc receptors are first found in the retina, but in a few of the brain regions that contain this receptor, SGS is the most strongly labeled [28].

There are still thirteen genes expressed in the SC which are not as specifically explored as the above eight genes, but these thirteen genes are also play a role in the SC, this research does not cover these genes.

3.2. Connectivity

Table 3 illustrates the anterograde projection from SC to other regions based on the Allen Brain Atlas. The left column displays the projection structures, and the right column shows the four different slices overlaid on the ABA. There are six representative projections are shown in the table 3, and more projections are described on connectivity. brain-map. org.

Hughes et al., 1984 observed that superior colliculus projected to the LGN, and the projection cells of the SC were mainly labeled in superficial layer (Ⅰ and Ⅱ, ZO and SGR), some of the labeled cells were located in the layer Ⅲ (OP) [29]. Several studies showed that the projection from the superior colliculus to the lateral geniculate nucleus is relatively simple and direct [30-32]. The nerve impulse of parabigeminal nucleus (Pbg) mainly comes from the superficial layer of SC. Baleydier et al 1979 injected HRP into the Pbg of the cat, they found that there were more than 60% in the deep layers of the SC, 30% in the IGR layer, and there were about 10% of the cells in the superficial layer. In the experiments, it was confirmed that the SC had fibers connecting directly to Pbg [33]. It has been mentioned that the function of the SC is complex, and it is also involved in the physiological activities of hearing, somatosensory and other activities. The common projection site is IC [34-35].

Table 3. The projection from SC to other structures using AAV tracer based on ABA.

| SCs → SCm | SCs → SCm and RSP | SCm → IC | SCm → MRN |
|-----------|-------------------|---------|-----------|
| Figure 3. The anterograde projections are from the SC to different regions of the mouse brain using mouseconnectome. org. It used the Chb555 and phal as the tracers. The left image is composed of injection sites in the SC to other structures, and the right image is the zoom in (100% larger) projected structures (PAG: Periaqueductal Gray, RN: Red Nucleus, CLI: Central linear nucleus raphe). |
Figure 4. The retrograde projections from other regions to the SC using the fg tracer.

As shown in figure 4, the left image is composed of injection sites from other structures to the SC, and the right image is the zoom in (100% larger) projected structures. These two representative projections are used the fg tracer. The yellow dots illustrate that the projections from PSPv and PBG are relatively obvious. In addition, there are more small structures, which can also project to SC are not mentioned in the figure 4, such as Red Nucleus (RN), PAG, IC, Primary visual area (VIS), Midbrain reticular nucleus (MBN), RL, Sn, Pbg, Rostral linear nucleus raphe (RLN), and Central linear nucleus raphe (CLI).

There are more projections from other regions that are explored in the past studies. Several studies inferred that the retina can project to SC, since the projection from the retina to SC is one of the largest sources of afferent fibers in the SC [36-37]. The visual cortex (including the area of 17, the area of the 18, and the area of the 19) is mainly projected into the visual layer and the IGR layer of the SC and most of them are projected to the layer IV and V (IGR and IW) [38-39]. There is a fiber linkage between the Pbg and the SC, which means Pbg can project to the SC and the SC can also project to the Pbg. Bickford et al., 1992 studied the projection from SNr to SC by HRP combining with GABA immunohistochemistry, suggested that the projection from SNr to the SC is GABA-ergic neurons [18]. The IC is an important relay nucleus and auditory reflex center in the auditory system and it is usually divided into three parts: the central, peripheral and central nuclei. SC and IC are collectively known as the Corpora Quadrigemina (CQ), and the pathway of the SC and IC is relatively important (appendix figure 2 showed the scheme of the anterograde and retrograde projection circuitries between SC and IC). The SC can project into the SC, and the IC can also project into the deep layers of the SC, which indicates that in addition to being responsible for visual stimulation, the SC are responsible for auditory and other sensory stimulation [41-43].

4. Strengths and Weaknesses

The strength of this paper is that it delineates for the first time the sub-regions of the SC using Nissl stain, gene expression and the different tracers based on the two online sources. However, there are still some weaknesses in this study. Firstly, this study relies on the ABA too much, and some of the information may not complete since the ABA is still not fully complete, thus it still requires more information. For example, the connectivity of SC with other structures has not been fully described in the atlas. Furthermore, the resolution of some images is still not high enough, and it is difficult to distinguish different layers. For example, it is hard to find the seven layers of SC in Nissl stain images. Future studies should concentrate on the connectivity of SC (the connectivity of SC is limited by the sources, especially in the mouseconnectome website, where only two structures can be found). There are actually more projections related to SC which are not delineated in the sources, but past studies show these projections. It may also be ideal to further explore the Tpd521gene in the SC, since it is densely expressed in the SC, so may play a greater role in the SC than expected.

5. Conclusion

The Superior Colliculus plays an important role in visual target selection and attention shifting (both overt and covert attention). In this study, it addresses the cytoarchitecture and connectivity of the SC in mouse brain. This study demonstrated that gene Tpd521 is densely expressed in the SC, which helps delineate the borders of the SC. In addition, the anterograde and retrograde projection circuitries between SC and IC are particularly addressed, which indicated that the SC is involved in the visual, auditory and other somatosensory physiological activities. However, this study does not clarify the entire complexity of the SC. For future work, we will investigate more genes which is expressed in SC, and explore more connectivities of the SC.

Appendix
Figure 6. The projections involved in the IC and the SC (dorsal view of the surface of the corpora quadrigemina). 1: Intrinsic projections of the SC; 2: Intrinsic projections of the IC; 3: Projections from the IC to the SC; 4: Projections from the SC to the IC; 5: Commissural projections of the SC; 6: Commissural projections of the IC. For easy understanding, some of the contralateral SC/IC projections are not shown in this scheme [44].

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