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# A Guide on Modelling Synapses in CellBlender and MCell

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## **Abstract**

This guide aims to improve on the work by Czech et. al., by updating the guide for the MCell - CellBlender software.

# 1 Pre- and Postsynaptic Geometry with Blender

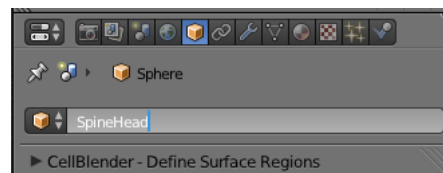
This section follows the paper written by Czech et. al. closely. It is expected that users will have familiarised themselves with Blender, and have read the MCell - CellBlender guide online.

## 1.1 Creating a Spine Head

1. Open Blender. In the '3D View' pane, delete the default object (shortcut: x)
2. Create a sphere. Select the 'UV Sphere' option from the sidebar. Below a pane called 'Add Circle' appears; set 'Segments' and 'Rings' to 16, 'Radius' to '0.25'.



3. Rename the sphere. Double-click the entry box below to change the default name to 'SpineHead'.



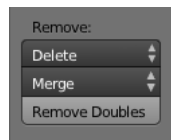
4. Change view to see the sphere from the 'Front' view. (shortcut: 1 on numpad)
5. Deselect the sphere (shortcut: a) and make it transparent (shortcut: z)
6. Select the vertices to be removed. First, switch from 'Object Mode' to 'Edit Mode'. Ensure that 'Edge select' is enabled. Then use box select (shortcut: b) to capture only the faces that make up the top half of the sphere. Delete these faces (shortcut: x) and select the 'Faces' option in the delete menu.



7. Close the opening. Select the topmost vertices (remaining in 'Edge select' mode) using box select (shortcut: b). Then, extrude (shortcut: e) and click 'Edges Only' under 'Extrude' in 'Mesh Tools'.

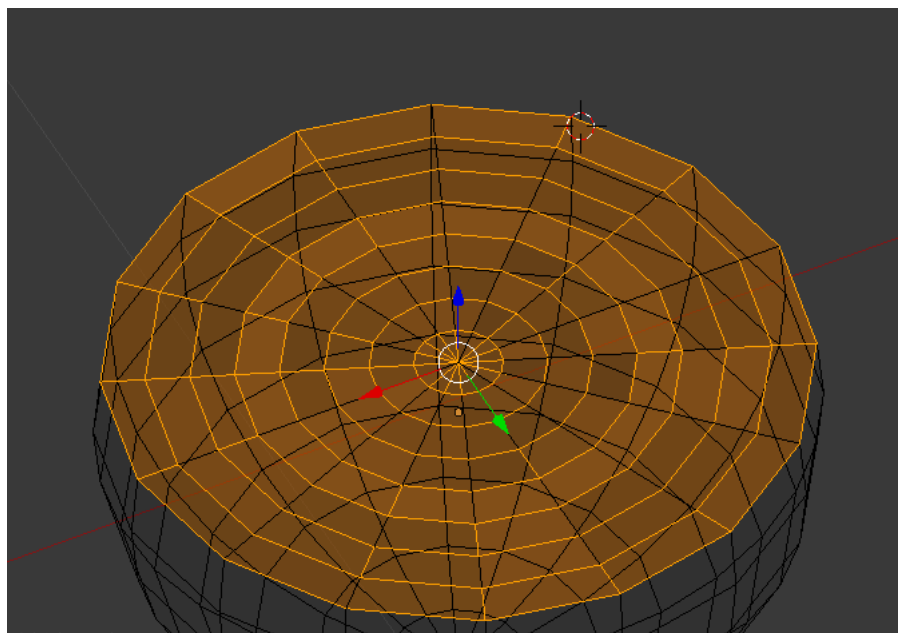


Set the extrude distance by pressing 0 and Enter to confirm. Scale the extrusion by pressing s, 0 and Enter to confirm. Select 'Remove Doubles' under 'Mesh Tools' to remove the duplicated vertices and reconnect the triangles. Blender should note that you remove 15 vertices as a result.



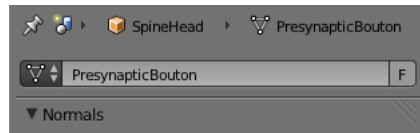
The object should now be closed by a flat top.

8. Subdivide triangles at the top. Unfortunately the 'Multicut tool' has been deprecated. To create set of concentric rings, first select the topmost vertices (shortcut: b). Then enter 'Knife' mode (shortcut: k), and hold down *control*. This locks the knife cut to the midpoint. Click around to make a circle cut, and then press *Enter* to complete the cut. Continue these cuts until the picture looks as below:

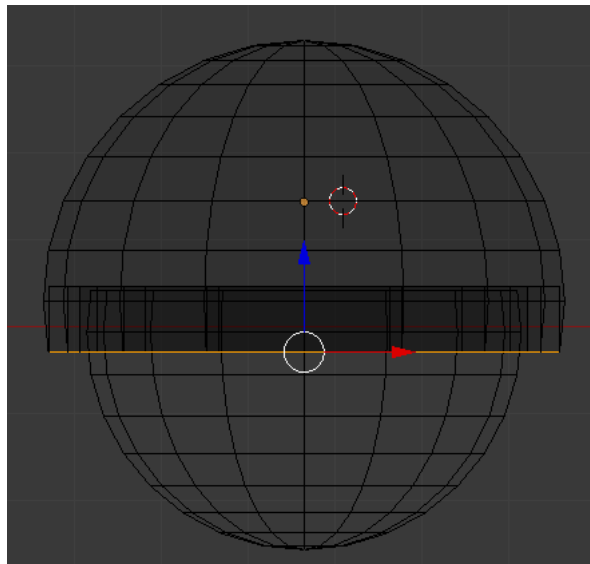


## 1.2 Creating a presynaptic bouton

1. Duplicate the *spine.blend* file as *bouton.blend*. in *bouton.blend* deselect all (shortcut: a).
2. Duplicate the spine head and rotate. First, switch to 'Front' view (shortcut: 1 on numpad). Duplicate (shortcut: Shift-d, Enter) and rotate 180 degrees (shortcut: r, 180, Enter). To separate the selection of the spine heads, press p and click 'Selection'.
3. Rename the item. Switch to 'Object Mode' (shortcut: Tab) and edit the name field to 'PresynapticBouton'.



4. Shift and scale the bouton. Grab the object (shortcut: g), constrain movement to the z-axis (shortcut: z) and type 0.15, Enter to move the object. Scale the object to be 20% larger by typing s, 1.2, Enter.
5. Creating the invagination. Switch to 'Edit Mode' (shortcut: Tab) and deselect everything (shortcut: a). Select the lowermost vertices (see below) and perform the 'Select Less' operation (shortcut: Control-Minus on numpad). Click the 'Extrude Region' button under 'Add' in 'Tools', press z to constrain, and type -0.075 to extrude it upwards.



## 1.3 Adding Axonal and Dendritic Extensions

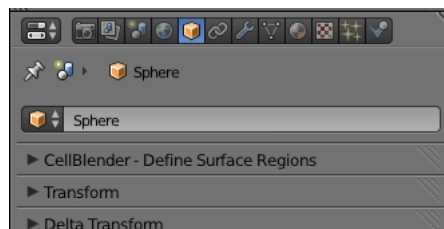
1. Enter 'Edit Mode' (shortcut: Tab), and switch view to front (shortcut: 1 on numpad)
2. Zoom in on the top of the vertex. Select the vertex by right-clicking on it, ensuring that 'Vertex mode' selection is enabled. Perform two 'Select More' operations (shortcut: Control-Plus on numpad) until two rings are highlighted. Press x, select 'Faces' on the menu,

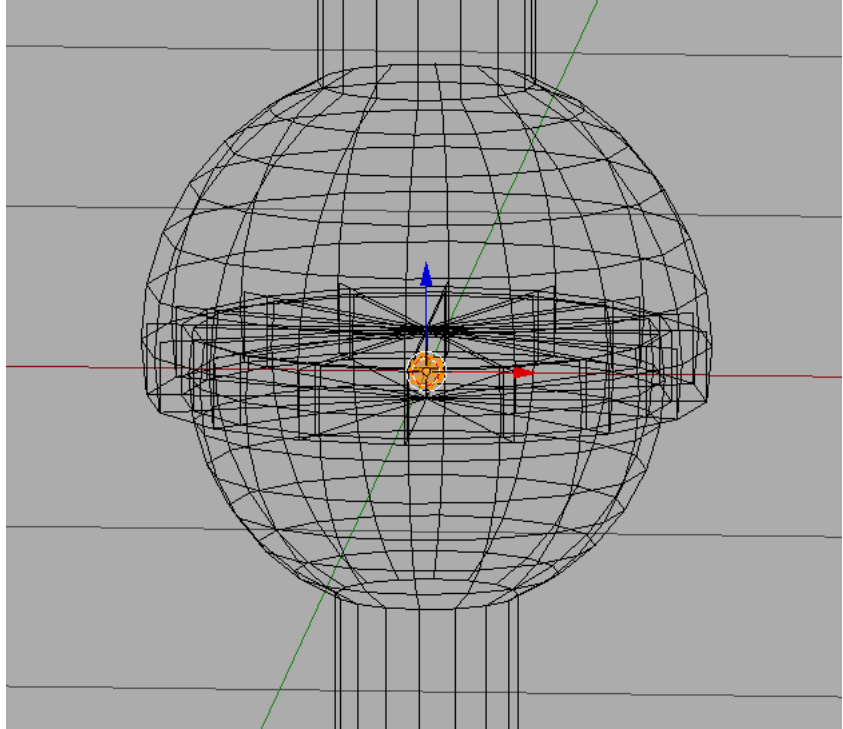
*Enter* to confirm. Press *b* and select the vertices on the top edge using 'Box select'. Press *e*, click on 'Edges Only' in 'Tools' (left pane), then type *z, 3.0, Enter*. This should produce an axon on the presynaptic bouton.

3. Select the spine head. Hit *Tab* to enter 'Object Mode' and right click the spine head (the bottom half-sphere) to select it. *Tab* back into 'Edit Mode'
4. Create a cylindrical spine on the spine head. As in the earlier step, select the bottom vertex, perform two 'Select More' operations, press *x* and select 'Faces' on the 'Erase' menu. Hit *b* and select the vertices that line the hole in the bottom. Press *e*, select 'Edges Only' in 'Tools', press *z*, type *-2.0, Enter*.

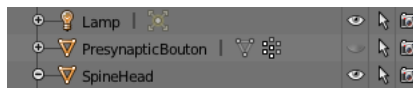
#### 1.4 Add Synaptic Vesicles with Regions for Calcium Binding

1. Create the first vesicle. First, ensure that the object is at the origin by moving it if necessary (shortcut: *g*). Move the cursor to the origin (shortcut: *Shift-c*). Ensure that 'Object Mode' is enabled. Under the 'Create' tab at left, select Ico-Sphere. A pane appears at lower left - change the default settings to 1 subdivision a size of 0.02. A small sphere should appear at the origin.
2. It is necessary to rotate the vesicle. With the sphere selected, press *r, x, 90, Enter* to align the sphere with the x-axis. Enter the 'Object' tab at right, and edit the name of the sphere to 'Vesicle'.
3. Define the calcium binding region. With the vesicle still selected, change to 'Edit Mode' (shortcut: *Tab*) and triangulate the faces (shortcut: *Control-t*). First, select the 'Define Surface Region' tab which is under the 'Object' tab. Add a new region and name it 'vesicle1'. With the vesicle selected, click 'Assign'. We must now add the vesicle to the list of mesh objects to be included in the simulation. Under the 'Scene' tab, select 'Model Objects' and add the vesicle to this list. A green tick should appear.





4. Move the vesicle. Tab into 'Object Mode', press g to grab the vesicle, type x, -0.108 *Enter* to move the vesicle along the x-axis. Hit g, 0.105, *Enter* to move it 0.105 units along the z-axis.
5. Duplicate the vesicle. Hit *Shift-d*, g, 0.216, *Enter* to move the vesicle the appropriate distance. Then, restrict visibility of the Presynaptic Bouton, select both vesicles, and under 'Define Surface Region' hit the assign button again.



## 1.5 Defining region for voltage-gated calcium channels (VGCC)

1. Clip the existing mesh objects. Use *Alt-b* to draw a box around the presynaptic terminal and the vesicles. This will only show those items and hide the rest of the model.
2. Switch to overhead view. Hit 7 on the numpad to look down from above on the vesicles. We want to align the vesicles over a few faces (note how they sit at the intersection of four faces). Select the vesicles and press r, 10 to rotate them over the appropriate faces.
3. Select the faces. Enter 'Edit Mode' with 'PresynapticBouton' selected to observe the faces. While holding *Shift* click on the faces directly under the vesicle (4 total).
4. Assign the selected regions. In the bottom right pane under 'Objects', go to 'Define Surface Region', and label it 'vgcc.region'. We want this region to be an object in its own right for later selection and editing. With the region selected, press p and hit 'Selection'.

Rename the object created to 'VGCC' in the top right pane. Remove the clipping box by pressing *Alt-b*.

## 1.6 Defining region for postsynaptic receptors

1. Hide the presynaptic bouton and the VGCC's. Click the eye button on the top right pane for the appropriate objects.
2. Define the postsynaptic receptor region. Right click on the postsynaptic dendrite and select the top face using the bounding box (shortcut: *b*). Use two 'Select Less' operations (shortcut: *Control - minus* on numpad) to deselect two outer rings. With this selection, create a new object by hitting *p* and clicking on 'Selection'. Then name this new object 'PostsynapticReceptor'.
3. Assign region. Under 'Objects' go to 'Define Surface Region' and label the receptor as 'postsynaptic.receptor'. Press *Control - t* to triangulate the faces, and then add it under 'Scene' in 'Model Objects'.

## 2 Simulation using the CellBlender Interface

### 2.1 Specification of Molecules

The following molecules will need to be added to our simulation under 'Scene' in 'Define Molecules'. The format is name of molecule, type of molecule, and diffusion constant.

1. Ca, Volume, 1e-6
2. VGCC\_C, Surface, 0
3. VGCC\_O, Surface, 0
4. CaBS, Surface, 0
5. CaBS-Ca, Surface, 0

### 2.2 Define Reactions between Molecules

Given a set of molecules, we must define the interactions that occur when molecules meet. Under 'Define Reactions', enter the following (consists of reaction, rate constant):

1. VGCC\_C'  $\rightarrow$  VGCC\_O', 5e5
2. VGCC\_O'  $\rightarrow$  VGCC\_C', 500
3. VGCC\_O'  $\rightarrow$  VGCC\_O' + Ca', 1e3
4. Ca' + CaBS'  $\rightarrow$  CaBS-Ca', 1e7
5. CaBS-Ca'  $\rightarrow$  Ca' + CaBS', 500

The apostrophes are used to indicate to CellBlender the geometry at which the reaction occurs.

### 2.3 Adding Molecules to Regions

We only want the closed VGCC's and the calcium binding sites 'CaBS' to be present when the model is initialised. All other molecules will be generated by the reactions during simulation. The following items will need to be added under 'Modify Surface Regions'.

## 1. 'PresynapticBouton