

# How to use Genome Browser

## Getting Genome Browser

To get Jsorolla you can add to your package.json dependency:

## Using JavaScript version

First we should define our genome browser. This will contain the general structure of the genome browser and the karyotype and chromosome panels. To draw these panels, genome browser will use the database "Cellbase" to which also it should be indicated which species we are using. You can see all the species available on Cellbase [here](#).

The target parameter indicates the id of the component wherer the genome browser will be created, in the example there is a "div" called "application". Autorender to draw the only one. If this variable is false, it will be necessary to call the render function of the genome browser. Another parameter of configuration is if we want that the genome browser can be resizable or on the contrary, once defined the size this will be fixed.

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### Example: Define Genome-Browser

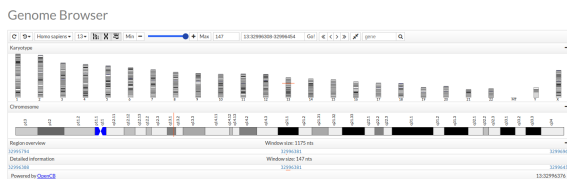
```
let species = {id:"hsapiens", scientificName:"Homo sapiens", assembly:
{name:"GRCh37"}};
let region = new Region({chromosome: "13", start: 32996311, end:
33000856}); //initial region
let genomeBrowser = new GenomeBrowser({
  client: cellbaseClient,
  cellBaseHost: CELLBASE_HOST,
  cellBaseVersion: CELLBASE_VERSION,
  target: 'application',
  width: document.querySelector('#application').getBoundingClientRect().
width,
  region: region,
  species: species,
  autoRender: true,
  resizable: true,
  karyotypePanelConfig: {
    collapsed: false,
    collapsible: true
  },
  chromosomePanelConfig: {
    collapsed: false,
    collapsible: true
  },
  navigationBarConfig: {
    componentsConfig: {
    }
  },
  handlers: {
    'region:change': function(e) {
      console.log(e)
    }
  }
});
```

The karyotype and chromosome panels can be passed the configuration if the default is closed or not, or if it is possible to close it or it will always be open.

Line 22-24 are optional, and are useful if you want to add some custom settings to the navigation bar.

Finally we can add to the genome-browser handlers as shown in the example.

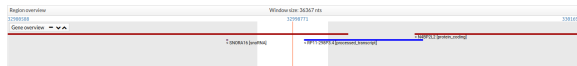
At this moment your genome browser would be started and it would look like this:



This would be basic form of the genome browser, to enrich it we can define a series of tracks or zones of visualization with different features as explained below.

## Overview Track

The particularity of this track is that it has a different zoom to the rest. This smaller zoom (more remote), serves to have an overview of what is in that area sought while in the rest of tracks you see in more detail the zone.



We will define a `FeatureTrack` to which we will pass the renderer, the adapter and other parameters of configuration of the track like the title, the minimum size where it changes histogram (`minHistogramRegionSize`), the maximum size where the names of the components display (`maxLabelRegionSize`) or height.

`FeatureRenderer` is the renderer in charge of painting a box for each feature. Adding the gene type (`FEATURE_TYPES.gene`) knows how to draw the color for each feature and the tag among other things.

The data will be collected with a `CellBaseAdapter`, indicating that we want to collect the genes and exclude the transcripts since being a distant zoom if we painted them would be difficult to visualize.

### Example: Overview Track

```
let gene = new FeatureTrack({
  title: 'Gene overview',
  minHistogramRegionSize: 20000000,
  maxLabelRegionSize: 10000000,
  height: 80,
  renderer: new FeatureRenderer(FEATURE_TYPES.gene),
  dataAdapter: new CellBaseAdapter(cellbaseClient, "genomic", "region",
"gene", {
    exclude: 'transcripts,chunkIds'
  }, {
    chunkSize: 100000
  })
});
genomeBrowser.addOverviewTrack(gene);
```

## Sequence Track

The sequence track shows that nucleotide matches in the position we are looking for.



This track accepts in this definition the title, the height of the track, the maximum size of nucleotides in which the track continues painting (`visibleRegionSize`), the renderer and the `CellBaseAdapter`. We recommend that the value of the `visibleRegion` not be very high because the sequence would be clearly seen when painting thousand of letters.

### Example:Sequence Track

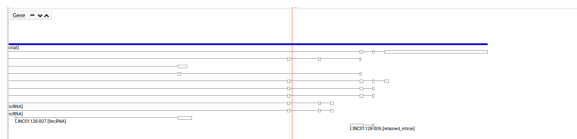
```
let tracks = [];  
this.sequence = new FeatureTrack({  
  title: 'Sequence',  
  height: 20,  
  visibleRegionSize: 200,  
  renderer: new SequenceRenderer(),  
  dataAdapter: new CellBaseAdapter(cellbaseClient, "genomic", "region",  
"sequence", {}, { chunkSize: 100})  
});  
tracks.push(this.sequence);  
genomeBrowser.addTrack(tracks);
```

To add this track to the genome browser, we will define an array of tracks that we will add all the track we want to add (except overviewtrack), and later add this to the genome browser as it does on line 10.

## Gene Track

Adding a gene track is similar to making a sequence track since we will also define it and add to the track array which we then add to the genomeBrowser.

Gene track shows the genes and transcripts that are in the region visualized, for this we consult the cellbase database to which we pass which species we are consulting.



### Example: Gene Track

```
this.gene = new GeneTrack({  
  title: 'Gene',  
  minHistogramRegionSize: 20000000,  
  maxLabelRegionSize: 10000000,  
  minTranscriptRegionSize: 200000,  
  height: 120,  
  cellbase: {  
    "host": CELLBASE_HOST,  
    "version": CELLBASE_VERSION,  
    "species": "hsapiens"  
  }  
});  
tracks.push(this.gene);
```

If we want to change the renderer or default adapter of the gene track we can indicate it with the parameters `renderer` and `adapter` respectively.

### Example: Gene Track Advanced Mode

```
this.gene = new GeneTrack({
  title: 'Gene',
  minHistogramRegionSize: 20000000,
  maxLabelRegionSize: 10000000,
  minTranscriptRegionSize: 200000,
  height: 120,
  cellbase: {
    "host": CELLBASE_HOST,
    "version": CELLBASE_VERSION,
    "species": "hsapiens"
  }
  renderer: new GeneRenderer({
    handlers: {
      'feature:click': function(e) {
        console.log(e)
      }
    }
  })
  dataAdapter: new CellBaseAdapter(cellbaseClient, "genomic",
"region", "gene", {
  exclude: 'transcripts.tfbs,transcripts.xrefs,transcripts.
exons.sequence'
}, {
  chunkSize: 100000
})
});
tracks.push(this.gene);
```

## Variant Track

The variant track has two operating modes: individual or family. If we pass name of one of ours will understand that these samples correspond to a family and will activate the family mode, otherwise the individual mode will be activated.

### Individual mode:

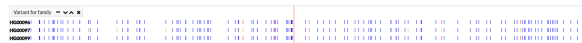


In the individual mode each variant is and individual and so is shown. In the example as they are variants of type snp, we pass the configuration of the painting that we want to the renderer. This configuration is defined in the config file of the source code, inside the genome-browser.js then the library is downloaded with npm.

### Example: Individual Variation

```
let variant = new VariantTrack({
  title: "Variant",
  closable: true,
  minHistogramRegionSize: 20000000,
  maxLabelRegionSize: 10000000,
  minTranscriptRegionSize: 200000,
  visibleRegionSize: 100000000,
  height: 300,
  opencga: {
    client: opencgaClient,
    studies: "platinum:illumina_platinum"
  }
});
tracks.push(variant);
```

## Family mode:



When we define the samples in the variant track the family mode is activated. This shows in the left side the name of the sample and in horizontal we can see the variants of each sample.

### Example: Family Variant

```
let variant = new VariantTrack({
  title: "Variant for family",
  closable: true,
  minHistogramRegionSize: 20000000,
  maxLabelRegionSize: 10000000,
  minTranscriptRegionSize: 200000,
  visibleRegionSize: 100000000,
  height: 300,
  opencga: {
    client: opencgaClient,
    studies: "reference_grch37:1kG_phase3",
    samples: ["HG00096", "HG00097", "HG00099"]
  }
});
tracks.push(variant);
```

When you use the opencga parameter, the genome browser itself understands that the adapter we want to user is the OpencgaAdapter. The renderer to use is the one that comes by default defined in the variant track. This renderer is the FeatureRender using the FeatureType.Variant configuration. This FeatureType, there in config archive in the code or genome-browser.js when you getting jsorolla using npm.

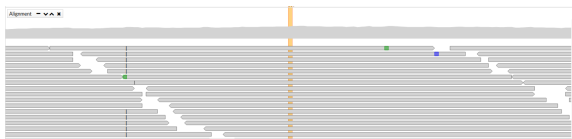
The code equivalent to the above but using the adapter definition and renderer would be as follows:

### Example: Individual Variant using adapater and renderer definition

```
let variant = new VariantTrack({
  title: "Variant",
  closable: true,
  minHistogramRegionSize: 20000000,
  maxLabelRegionSize: 10000000,
  minTranscriptRegionSize: 200000,
  visibleRegionSize: 100000000,
  height: 300,
  renderer: new FeatureRenderer(FEATURE_TYPES.variant),
  dataAdapter: new OpencgaAdapter(opencgaClient, "analysis
/variant", undefined, "query",
  {
    studies: "platinum:illumina_platinum",
    exclude: "studies.files,studies.stats,annotation"
  }, {
    //Exclude for samples mode
    exclude: "studies,annotation" //Exclude for individual
    variation mode
  }, {
    chunkSize: 100000
  })
});
tracks.push(variant);
```

It is important for the developer to understand that the genome browser always uses a track, an adapter, and a renderer even though it does not look the same in its definition. If the developer wants to change the renderer or the adapter using the default tracks should include the adapter or renderer they want in the definition of the track.

## Alignmet Track



The alignment track is made up at the top of the coverage and at the bottom of the bam reads.

#### Example: Alignment Track

```
let alignment = new AlignmentTrack({
  title: "Alignment",
  closable: true,
  minHistogramRegionSize: 5000,
  maxLabelRegionSize: 3000,
  visibleRegionSize: 100000000,
  height: 300,
  renderer: new AlignmentRenderer(FEATURE_TYPES.bam),
  dataAdapter: new OpencgaAdapter(opencgaClient, "analysis
/alignment", undefined, "query",
  {
    fileId: fileId,
    study: study
  })
});

tracks.push(alignment);
```

When you use the adapter of Opencga to visualize the bam you must pass the name of the files to him by the parameter fileId.

## Custom Track

You may want to collect data from another database but you don't want to implement a custom adapter for yourself. The solution to this problem is to use the adapter template.

In the following example, instead we create a variant track using a template adapter instead of our adapter for the cellbase database.

### Example: Template Adaptor

```
let t = new FeatureTrack({
  title: 'Variant',
  closable: true,
  minHistogramRegionSize: 20000000,
  maxLabelRegionSize: 10000000,
  minTranscriptRegionSize: 200000,
  histogramColor: '#92de47',
  height: 100,
  renderer: new FeatureRenderer(FEATURE_TYPES.snp),
  dataAdapter: new FeatureTemplateAdapter({
    multiRegions: false,
    histogramMultiRegions: false,
    uriTemplate: "http://ws.babelomics.org/cellbase/webservices/rest/v4
/hsapiens/genomic/region/13:32990000-32999999/snp?          exclude=
{exclude}",
    templateVariables: {
      exclude: 'annotation.populationFrequencies,annotation.
additionalAttributes,transcriptVariations,xrefs,samples'
    },
    species: genomeBrowser.species,
    cacheConfig: {
      chunkSize: 100000
    },
    parse: function(response) {
      var chunks = [];
      for(var i = 0; i< response.response.length; i++){
        var res = response.response[i].result;
        chunks.push(res);
      }
      return chunks;
    }
  })
});
tracks.push(t);
```

Param	Type	Description
multiRegions	boolean	Indicates if large regions will make a single call to the server or several.
histogramMultiRegions	boolean	Indicates whether large regions will make a single call to the server or multiple calls when the display zone is in histogram mode.
uriTemplate	string	Uri of the server to which calls will be made. The value of the query params will be indicated by a "{variable}". This value will be declared in templateVariables.
templateVariables	object	Object that includes the value of each query param of the uriTemplate. As query param will be defined as: in the key, the name of the query params that we have put in the uriTemplate and as value, the value to be replaced in the uriTemplate.
species	object	Specie object, This parameter is optional and servers to add name of the species in the cache. It will be necessary if your viewer has the option to change species.
cacheConfig	object	Object to define the configuration of the cache whose main element is the chunkSize.
parse	function	Since the render expects an array of chunks to paint, you may need to parse the result of the response. Defining this function will parse the result when making calls and before passing the result to the renderer.

# Using Polymer web component

To facilitate the use of the Genome Browser component, a WebComponent has been created that you can import into your html page.  
In this tutorial you will learn how use this component for visualize differents tracks.

In first time, you must import the component in your page.

## import

```
<link rel="import" href="jsorolla/dist/core/webcomponent/genome-browser.html">
```

Subsequently, we will declare the genome component browser in our html code. For this we must know the list of parameters that the component accepts.

Here is a list of parameters that the component accepts and uses:

Param	Type	Description
cellbase-client	Object	Client for use Cellbase. Necessary for: Gene track (using cellbase adapter, default).
opencga-client	Object	Client for use opencgaDB. Necessary for: Varriant Track and Aligment Track (using opencga adapter, default).
project	Object	To use opencga webservices, you will need to know which project the data to display.
study	Object	To use the opencga webservices, this will need to know in which study within a project, are the data to be visualized.
region	Object	Region where the genome-maps will initially be positioned.
species	Object	Species to draw. Default: homo sapiens.
width	Number	Component width.
active	Boolean	Enables or disables the component. If it is disabled the component will not be displayed.  Default: false.
settings	Object	This parameter sets other settings of the genome maps that the user wants to pass.

Finally, declare the component. In this case, we only defined the GeneTrack and Sequence track:

```
<genome-browser cellbase-client="{{cellbaseClient}}" region="{{region}}"
tracks="{{tracks}}" active="true">
</genome-browser>
```

In the javascript part we will give value to the variables:



```
CELLBASE_HOST = "bioinfo.hpc.cam.ac.uk/cellbase";
CELLBASE_VERSION = "v4";

let cellBaseClientConfig = new CellBaseClientConfig
(CELLBASE_HOST, CELLBASE_VERSION);
this.cellbaseClient = new CellBaseClient(cellBaseClientConfig);

this.region = new Region({chromosome: "11", start:
68177378, end: 68177510});
this.tracks = {sequence: {type: "sequence"}, gene: {type:
"gene"}};
```

We have created a cellbase client that will connect to the Host "[bioinfo.hpc.cam.ac.uk/Cellbase](http://bioinfo.hpc.cam.ac.uk/Cellbase)". If we do not want to change the installation of Cellbase, it is not necessary that we define the host and the version, but we initialize a client. With the parameter "region", we ask that the display is at "11:68177378-68177510". It's possible to not see these positions in the display where the painted region is shown, but you will see a window frame containing the region. This is because the genome browser adjusts the display to optimize the cache.

Finally, the last param is "tracks", in this case we want to see the track "sequence" and "gene". We can define other types of tracks as follows:

- Individual Variant Track (for example SNP): {trackname: {type: "variant", config: {}} }.
- Family Variant Track: {trackname: {type: "variant", config: {samples: "sample1", "sample2"}}}.
- Alignment Track: {trackname: {type: "alignment", config: {files: "bam1", "bam2"}}}.