

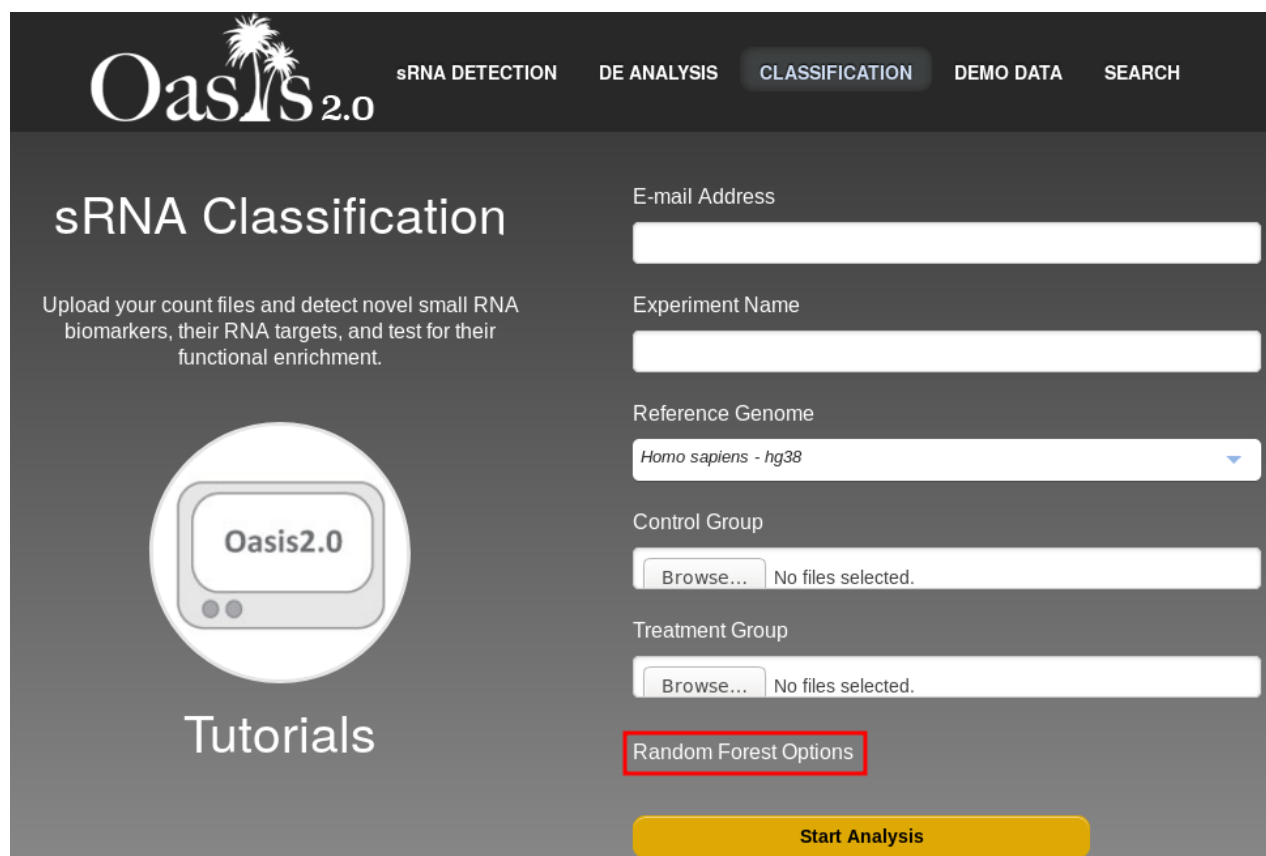
## sRNA-based disease classification tutorial

*Summary:* The third pipeline produces a machine learning model that you can use to predict the disease status of new samples (healthy/disease). With this model you can create a new biomarker for your disease based on sRNA sequence count data.

This tutorial explains how to submit count files (output by [Oasis' sRNA Detection module](#)) to the [Oasis Classification module](#). The Classification module is the third analysis module of Oasis and it performs a binary classification on the count files by applying a Random Forest classifier. This step in the Oasis workflow provides an exploratory analysis of the data, gives classifier's performance measures and a list of the most relevant features (i.e. sRNAs) in classifying the different samples into their equivalent groups. Guidelines on how to interpret the results of the Classification detection module can be found in [Oasis' Classification Output Tutorial](#).

### Submitting a classification job

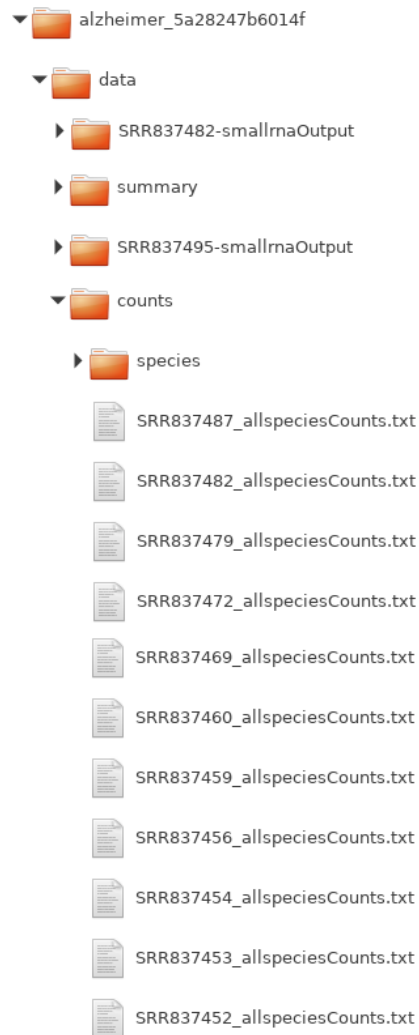
Like any other Oasis job, you need to fill in the fields for the **E-Mail address** where status updates and results should be mailed to. You also have to pick an **Experiment Name** that will later allow you to remember what the Oasis job was about when you submitted it. The **Reference Genome** should correspond to the species from which the samples have been taken.



The screenshot shows the Oasis 2.0 sRNA Classification interface. At the top, there are navigation tabs: sRNA DETECTION, DE ANALYSIS, CLASSIFICATION (selected), DEMO DATA, and SEARCH. The main heading is "sRNA Classification" with a sub-heading "Upload your count files and detect novel small RNA biomarkers, their RNA targets, and test for their functional enrichment." Below this is a circular logo for Oasis2.0 and a "Tutorials" link. The form fields are: E-mail Address (text input), Experiment Name (text input), Reference Genome (dropdown menu showing "Homo sapiens - hg38"), Control Group (file upload field with "Browse..." button and "No files selected." text), and Treatment Group (file upload field with "Browse..." button and "No files selected." text). A red box highlights the "Random Forest Options" link, and a yellow "Start Analysis" button is at the bottom.

**Figure 1: Oasis classification job form. The advanced Random Forest Options are hidden behind the link marked in red.**

For the **control group** and the **treatment group(s)** you need to upload sRNA count files from your computer. You probably have obtained these sRNA count files from downloading your results from the [sRNA detection module](#). If you look at the sRNA detection module results, the folder structure is as depicted in figure 2: You need to access the `data/counts/` subfolder and then pick those `_allspeciesCounts.txt` files that correspond to either the control group or one of the treatment groups. Note that you can select multiple files for each upload field at once. This way you can bundle all control count files and all treatment count files (for each group).



**Figure 2: Folder structure of the results from the Oasis [sRNA detection module](#). This screenshot emphasizes the the `_allspeciesCount.txt` files which contain the counts for each known and predicted small RNA in the Oasis database for the sample in question.**

### Optional parameters

If you click on `Random Forest Options` (marked in a red box in figure 1) you will have access to two more form fields:

`mtry`

Number of variables chosen to build the trees in the forest (can be automatically calculated by Oasis)

`ntree`

The number of trees used to populate the forest.

## References

Capece, V., Garcia Vizcaino, J. C., Vidal, R., Rahman, R.-U., Pena Centeno, T., Shomroni, O., ... Bonn, S. (2015). Oasis: online analysis of small RNA deep sequencing data. *Bioinformatics*, 31 (13), 2205-2207. <http://doi.org/10.1093/bioinformatics/btv113>