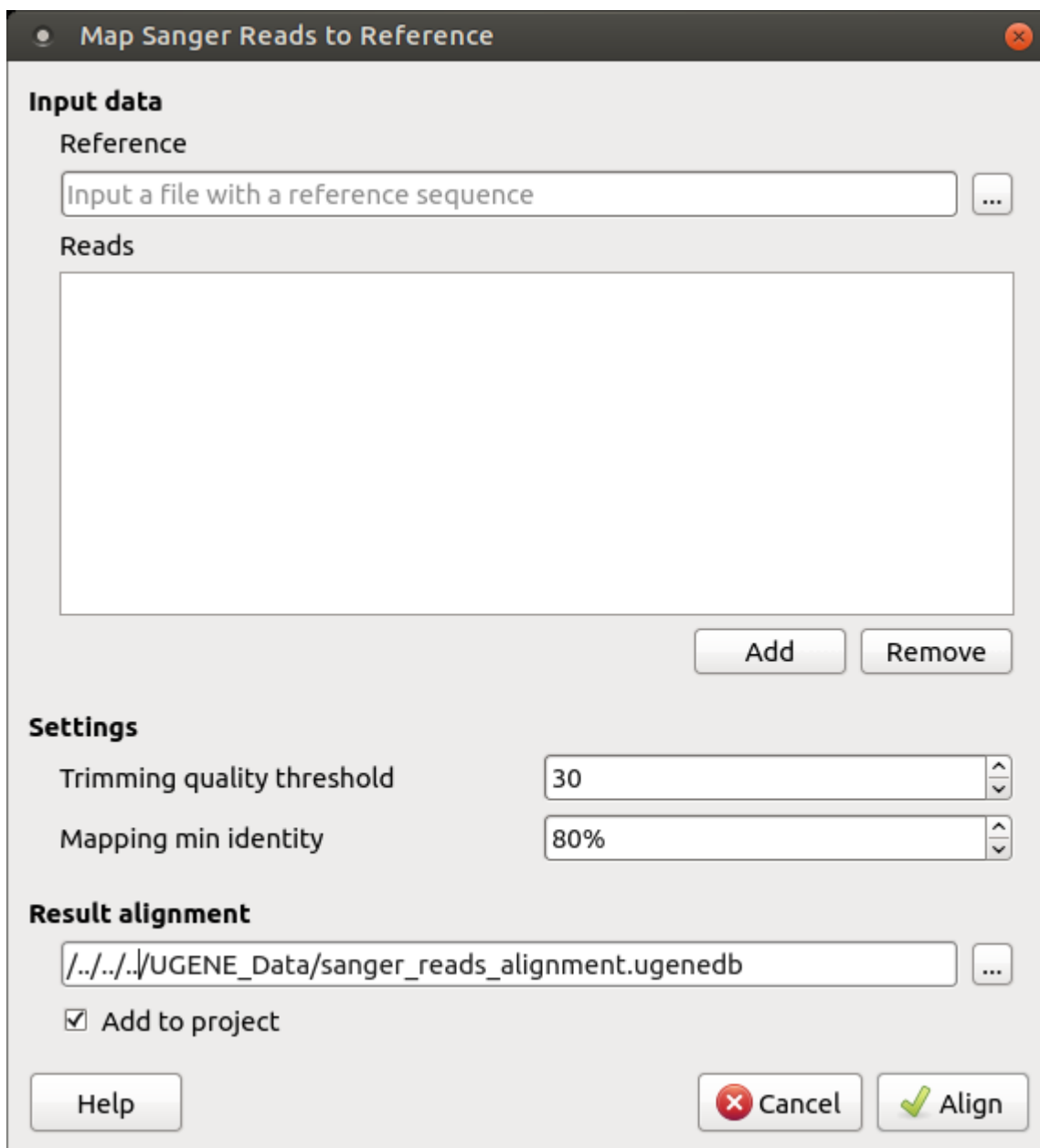


# Mapping Reads to Reference

To map Sanger sequencing reads to a reference use the *Tools*→*Sanger data analysis*→*Map reads to reference* main menu item. The following dialog appears:



The dialog box is titled "Map Sanger Reads to Reference". It contains several sections: "Input data" with fields for "Reference" (a text box with a file icon) and "Reads" (a large empty list box with "Add" and "Remove" buttons); "Settings" with "Trimming quality threshold" (set to 30) and "Mapping min identity" (set to 80%); and "Result alignment" (a text box with a file icon) and a checked "Add to project" checkbox. At the bottom are "Help", "Cancel", and "Align" buttons.

**Map Sanger Reads to Reference**

**Input data**

Reference

Input a file with a reference sequence

Reads

Add Remove

**Settings**

Trimming quality threshold 30

Mapping min identity 80%

**Result alignment**

./.././../UGENE\_Data/sanger\_reads\_alignment.ugenedb

☒ Add to project

Help Cancel Align

It is required to input:

*Reference sequence* — a file with a single DNA sequence of any supported file format (e.g. FASTA or GenBank). This parameter is mandatory.

*Reads* — a set of files in \*.ab1 or \*.scf format with the Sanger sequencing data to be mapped to the reference sequence. It is required to add at least one read. A read orientation (forward or reverse) will be automatically detected during the mapping.

During the mapping task execution these data are processed as follows. First of all, to enhance the further mapping sensitivity, the [low-quality](#) ends of the reads are trimmed off. Then the reads are mapped using a UGENE original method that works in two steps: rough mapping of a read using BLAST+ and enhancement of the read alignment with the Smith-Waterman algorithm. After that the percentage similarity of the two sequences is calculated, that is the number of all edit operations (i.e., insertions, deletions and substitutions) required to transform a read sequence into the corresponding region of the reference sequence. Reads that have low similarity with the reference sequence are filtered out.

One can configure the following parameters of the mapping task

*Trimming quality threshold* — all bases at the ends of the reads with quality lower than the specified value are trimmed. To skip the trimming, set this value to zero.

*Mapping min similarity* — all reads mapped to the reference with lower percentage similarity than the specified value are filtered out.

*Read name in result alignment* — reads in the result alignment can be named either by names of the sequences in the input files or by the input files names. Set this value to "File name", for example, if the sequences in the input \*.ab1 files have the same name, this will help in distinguishing of the reads in the result alignment.

The result alignment is stored in a native UGENEDB format. One can set up the file location and name in the *Result alignment* field. Note that thereafter it is also possible to [export data to standard alignment formats without chromatograms](#) such as FASTA, ClustalW, etc.

To initiate the mapping task execution click the *Map* button in the dialog. Note that when the task is finished, the task statistics can be found in a report, available on clicking the corresponding [notification](#).