## Input YAML

```
assembly:
 assem level: {scaffold|chromosome}
 assem version: {Version number of the assembly}
 sample id: {Name of the assembly}
 latin name: {Scientific Name}
 defined class: {User defined class of the input assembly}
 project id: {Project ID of assembly} #Optional
reference file: {Path to .f{a|n|asta}{.gz} formatted input}
map order: length
assem reads:
read type: hifi
 read data: {Folder containing longread data in .fasta.gz
format }
hic data:
hic cram: {Folder containing HiC reads in cram format with
.crai}
hic aligner: {minimap2|bwamem2}
kmer profile:
 kmer length: {Default to 31}
 dir: {Path to pre-existing FKPROF files if they exist}
#Optional
alignment:
data dir: {Path to the top level gene alignment data folder}
 geneset id: {A csv delimited list of geneset data to align}
self comp:
motif len: 0
mummer chunk: 10
intron:
 size: "50k"
telomere:
 teloseq: {The expected telomeric sequence}
synteny:
synteny path: {Folder of FASTA files used for alignments}
 synteny genomes: {Specify files in the above path}
busco:
lineages path: {Path to the busco database e.g. /busco/v5}
 lineage: {The odb10 lineage to use}
```

## Run:

nf-core run sanger-tol/treeval -r 1.1.0 -input TreeVal.yaml profile {singularity|docker} -output {OUTDIR} -entry RAPID

## or

nf-core run sanger-tol/treeval -r 1.1.0 -input TreeVal.yaml profile {singularity|docker} -output {OUTDIR}



## Usage Cheatsheet

```
(OUTDIR)
                                        Output files
 – treeval_upload/
                      # GENERATE GENOME*
    - mv.aenome
    coverage.bw
                      # READ COVERAGE
                        # READ COVERAGE
    coverage log.bw
     * repeat density.bw # REPEAT DENSITY
    *gap.bed.gz
                      # GAP FINDER
    *gap.bed.gz.tbi
                      # GAP FINDER
    *telomere.bed.gz + .tbi # TELO FINDER
     * buscogene.bigbed
                         # BUSCO ANALYSIS*
     * ancestral.bigbed
                       # BUSCO ANALYSIS*
     *.qff.qz + .tbi
                       # GENE ALIGNMENT-PEPTIDE*
                       # GENE ALIGNMENT-NUCLEAR*
     * cdna.bigBed
    * cds.bigBed
                      # GENE_ALIGNMENT-NUCLEAR*
     * rna.bigBed
                      # GENE ALIGNMENT-NUCLEAR*
    BSPQ1.bigBed
                        # INSILICO DIGEST*
    BSSS1.bigBed
                       # INSILICO DIGEST*
    DLE1.biaBed
                       # INSILICO DIGEST*
    * selfcomp.bigBed
                        # SELFCOMP*
    - *.paf
                   # SYNTFNY*
    *.ref.spectra-cn.ln.png # KMER*
    * {kmer size} .bw
                        #KMER COVERAGE
    punchlists

    halfcoverage.bigbed # READ COVERAGE

                       # READ COVERAGE
     zerodepth.biabed
     - maxdepth.bigbed
                        # READ COVERAGE
      * pep_punchlist.bed # GENE_ALIGNMENT-PEPTIDE*
      * cdna punchlist.bed # GENE ALIGNMENT-NUCLEAR*
      cds punchlist.bed # GENE ALIGNMENT-NUCLEAR*
      * rna punchlist.bed # GENE ALIGNMENT-NUCLEAR*
  hic files
    - *gap.bed
                     # GAP_FINDER
     * repeat density.bw # REPEAT DENSITY
                    # HIC MAPPING
     *.mcool
    * pretext normal.pretext # HIC MAPPING + PRETEXT INGESTION
    * pretext hr.pretext # HIC MAPPING + PRETEXT INGESTION
   - *telomere.bed
                      #TELO FINDER
   pipeline info
   - TreeVal Runs*.txt
                       # TreeValProject.Summary (Groovy Function)
                      # STANDARD OUTPUT
    execution*{html|txt}
                     # STANDARD OUTPUT
    pipeline*{html|txt}
    software versions.yml # STANDARD OUTPUT
```