## Mapping File

(Accession, NCBI taxon ID)

NOTE: New marker packages are named according to input filenames (e.g. MarkerAlignment.fasta). Core marker data will be overwritten during new marker builds if input files do not have unique names compared to existing PhyloSift markers.

PhyloSift markers.

Quantitative metric (minimum hamming distance) used to match edges between NCBI taxon tree and molecular phylogeny



Locally indexed marker packages will not interfere with automatic updates to PhyloSift core markers PD cutoff

## Alignment File

(Marker sequences in FASTA format)

Execute build\_marker mode

HMMbuild

Unalign sequences and create profile HMM out of input data



Generate unique IDs for input sequences

FastTree

Build tree and collapse topology according to a user-specified PD cutoff (e.g. 99%)



Sequence accessions mapped to phylogeny via a dummy pplacer file

Tree Reconciliation Reconcile NCBI taxonomy IDs with phylogenetic topology



Clean and package new marker genes

Built Marker
Packages

New marker gene packages placed into shared PhyloSift marker directory

HB: Want to mention the dummy

readers). Is this appropriate?

pplacer step here but don't want to go

into too much detail (might confuse



Execute index mode

Index Marker
Database

Indexes the marker databases needed for LAST and Bowtie

## Built PhyloSift Marker package

Tree

Hmm profile

Taxon map

Representative sequences

Alignment

HB: Creepy monkey (top) or non creepy monkey (below)?? Or both (since they fill out the white space nicely)?