

| SeqQual pipeline / program (prog) name* | usage** | arguments (arg) | description (Record of actions and parameters printed into log.txt) | Shell examples *** |
|--|---|---|---|---|
| print_log-ace_changename.pl | (perl ~/SeqQual/prog arg >> log.txt) = (A) | | change ace name | 2.1-ace-only.sh |
| print_log-largeace_split.pl | (A) | | split large ace | 2.3-ace-qual.sh |
| print_log-largephd_split.pl | " | | split large phd files | 2.3-ace-qual.sh |
| print_log-acedealing-qual.pl | " | | dealing with ace and qual files | 2.3-ace-qual.sh |
| print_log-aceonly.pl | " | | dealing with ace files which have no other related files | 2.3-ace-qual.sh |
| print_log-aln-diploid-ab1.pl | " | polyphred score, polyphred quality and 18 parameters for phrap | diploid_ab1; phrap parameters used in phrap, polyphred pamaters used in polyphred | 1.1-diploid-ab1.sh |
| print_log-aln-haploid-ab1.pl | " | 18 parameters from Phrap | haploid_ab1; phrap parameters used in phap | 1.2-haploid-ab1.sh |
| print_log-arlequin-diploid.pl | " | up to 20 group names | writing arlequin input files from diploid data; and up to 20 group numbers or names used | *fasta-*data.sh |
| print_log-arlequin-haploid.pl | " | up to 20 group names | write arlequin input files for haploid data; and up to 20 group numbers or names used | *fasta-*data.sh |
| print_log-fasta.pl | " | | dealing with fasta alignment files | *fasta-*data.sh |
| print_log-remove1.pl | " | | remove bad positions with only "?" and "-" | 1.1-diploid-ab1.sh, 1.2- haploid-ab1.pl |
| print_log-renamephd.pl | " | | rename phd files | 1.1-diploid-ab1.sh, 1.2- haploid-ab1.pl |
| print_log-replace.pl | " | max length of "isolated nucleotides" | replace isolated nucleotides which are surrouned by "?"; the max length of "isolated nucleotides" (which is used by the script) | 1.1-diploid-ab1.sh |
| print_log-truncate.pl | " | number of nucleotides in the column from which truncate process is done | truncate; the number of nucleotides in the column from which the truncate process is done | 1.1-diploid-ab1.sh |
| print_log-userphd-diploid-ab1.pl | " | | inserting user phd/poly files | 1.1-diploid-ab1.sh |
| print_log-userphd-haploid-ab1.pl | " | | inserting user phd files | 1.2-haploid-ab1.sh |
| print_log-write_aln-diploid.pl | " | phred score, genotype score | write fasta alignment from ace, phd, polyphred.out; phred score and genotype score used | 1.1-diploid-ab1.sh |
| print_log-write_aln-qual.pl | " | phred score | write fasta alignment , phd score used | 2.3-ace-qual.sh |
| print_log-write_aln-haploid.pl | " | phred score | write fasta alignment , phd score used | 1.2-haploid-ab1.sh |
| print_log-write_haplotypesaln.pl | " | | write haploid phase unknown alignment files from diploid alignment files | *fasta-*data.sh |
| print_log-write_SNP-diploid.pl | " | phred score, genotype score | write SNP alignment; phred score, genotype score | *fasta-*data.sh |
| print_log-write_SNP-fasta_aln.pl | " | | write SNP alignment | *fasta-*data.sh |
| print_log-write_unaln-diploid.pl | " | phred score, genotype score | write unaligned fasta files; phred score, genotype score | *fasta-*data.sh |
| print_log-write_unaln-haploid.pl | " | phred score | write unaligned fasta files; phred score | *fasta-*data.sh |
| print_log-write_unaln-fasta.pl | " | | write unaligned fasta files | *fasta-*data.sh |
| take_log_to_output.pl | " | | take log.txt to folder Output, check name to log_data&time.txt | 1.1-diploid-ab1.sh, 1.2- haploid-ab1.sh, *fasta- *data.sh |

*all print_log scripts relate to the print_source scripts of the same names and can be used along with them

**all scripts append log information to the log.txt file

***all shell scripts can be run by typing "source *.sh"