



Apr 21, 2019

Working

The pipeline of Hi-C assembly of the *Scapharca broughtonii* genome

In 1 collection

Chang-Ming Bai¹¹Yellow Sea Fisheries Research Institute, Chinese Academy of Fishery Sciences[dx.doi.org/10.17504/protocols.io.z8cf9sw](https://doi.org/10.17504/protocols.io.z8cf9sw)

Chang-Ming Bai

ABSTRACT

This protocol include the detailed methods of Hi-C assembly of the *Scapharca broughtonii* genome

- 1 Run BWA (v0.7.10-r789) to align the Hi-C reads to the initially assembled *S. broughtonii* genome, and found the Hi-C reads mapped to the assembled genome.



```
bwa index -a bwtsw fasta
bwa aln -M 3 -O 11 -E 4 -t 2 fq1
bwa aln -M 3 -O 11 -E 4 -t 2 fq2
```

- 2 Filter the mapped Hi-C reads obtained in the step 1 using HiC-Pro (v. 2.10.0).



```
mapped_2hic_fragments.py -v -S -s 100 -l 1000 -a -f -r -o
```

- 3 Extract valid interaction pair reads according the HiC-Pro results.

- 4 Break the initial assembly to 300 bp, and then run LACHESIS (v2e27abb) for assembling based on Hi-C data.

- 5 Run LACHESIS (v2e27abb) to assemble corrected contigs obtained in step 4 into chromosome and modified manually.



```
(1) CLUSTER_MIN_RE_SITES = 22
(2) CLUSTER_MAX_LINK_DENSITY=2
(3) CLUSTER_NONINFORMATIVE_RATIO = 2
(4) ORDER_MIN_N_RES_IN_TRUN=10
(5) ORDER_MIN_N_RES_IN_SHREDS=10.
```



This is an open access protocol distributed under the terms of the [Creative Commons Attribution License](https://creativecommons.org/licenses/by/4.0/), which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited