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

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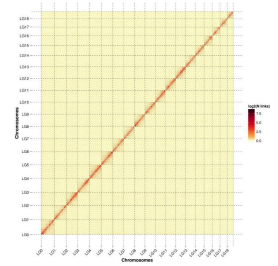
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Protocol status: Working

We use this protocol and it's working

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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.

Note

```
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).

Note

```
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.

Note

```
(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.
```