

# ALLPATHS-LG Genome Assembly and Hi-C mapping strategy

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## Abstract

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## Protocol

Quality control of WGS data using NGS QC Toolkit (version 2.3.3) and FastUniq (version 1.1)

### Step 1.

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**FastUniq, 1.1**

WGS Assembly: 1. Calculating insert size using Bowtie 2 (version 2.2.9) and Picard (2.7)

### Step 2.

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**Picard, 2.7** [↗](#)

WGS Assembly: 2. Constructing contigs and scaffolds using ALLPATHS-LG (version r52488)

### Step 3.

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**ALLPATHS-LG, r52488** [↗](#)

WGS Assembly: 3. Linking the scaffolds with SSPACE (version 1)

### Step 4.

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**SSPACE, 1** [↗](#)

WGS Assembly: 4. Filling the gaps with GapFiller (version 1.11) and GapCloser (version 1.12)

### Step 5.

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**GapCloser, 1.12**

Hi-C mapping: 1. Evaluating and qualifying Hi-C data with HiC-Pro (version 2.8.0\_devel)

### Step 6.

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**HiC-Pro, 2.8.0\_devel**

Hi-C mapping: 2. Hi-C assembling with 3D-DNA pipeline (version 170123)

### Step 7.

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**3D-DNA pipeline, 170123**

Hi-C mapping: 3. The contact map was visualized by Juicerbox (version 1.5.2)

### Step 8.

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**Juicerbox, 1.5.2**

Hi-C mapping: 4. To evaluate the Hi-C assembly, the chromosome sequences of moso bamboo were aligned with rice (*Oryza sativa*) genome using Lastz (version 1.02.00) [13] with the parameters: T=2 C=2 H=2000 Y=3400 L=6000 K=2200. Syntenic blocks longer than 2kb were plotted using Circos (version 0.69)

#### Step 9.

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**Circos, 0.69** [↗](#)