Instructions for scaffolding MIRA 454 contigs using 454 paired-end data and BAMBUS

Before you begin you should have a fasta file of the paired-end (PE) adaptor sequence, the PE and shotgun SFF files, and know the mean and standard deviation of PE construct (get from lab, likely a sizing gel).

Prepare fasta, qual, xml files using paired-end SFF file

Follow the instructions for combined paired-end /shotgun MIRA assembly. Issue something the something like the following command:

```
sff_extract -l ../paired_end_adaptor.fasta F04Y2AI01.sff -i "insert_size: 25000,insert_stdev:9000" -o non_0157
```

to create the following files:

```
-rw-rw-r-- 1 greg greg 125438250 Aug 24 16:59 non_0157.xml
-rw-rw-r-- 1 greg greg 391429721 Aug 24 16:59 non_0157.fasta.qual
-rw-rw-r-- 1 greg greg 136943770 Aug 24 16:59 non_0157.fasta
```

To check if everything went well., type

```
> less non_0157.xml
```

to get:

```
<?xml version="1.0"?>
<trace_volume>
	<trace>
		<trace_name>F04Y2AI01AIS9Z.fn</trace_name>
		<insert_stdev>9000</insert_stdev>
		<insert_size>25000</insert_size>
		<clip_vector_left>5</clip_vector_left>
		<clip_vector_right>58</clip_vector_right>
		</trace>
		<trace>
			<trace_name>F04Y2AI01AJLJM.r</trace_name>
			<insert_stdev>9000</insert_stdev>
			<insert_size>25000</insert_size>
			<clip_vector_right>2</clip_vector_right>
			<template_id>F04Y2AI01AJLJM</template_id>
			<trace_end>r</trace_end>
			</trace>
	</trace>
</trace_volume>
```

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looks good, PE data in XML file.

Now, work on shotgun SFF files and append (use `-a`) to PE data in fasta, qual, xml files:

```
sff_extract -a FAPPU2401.sff FA79DFG01.sff E95K6EU01.sff -o non_0157
```

to get something like

```
-rw-rw-r-- 1 greg greg 230827474 Aug 24 17:06 non_0157.xml
-rw-rw-r-- 1 greg greg 924673698 Aug 24 17:06 non_0157.fasta.qual
-rw-rw-r-- 1 greg greg 322941190 Aug 24 17:06 non_0157.fasta
```

Notice how the file sizes have changed to accommodate the shotgun seqs? Change names to make them mira friendly:

```
>mv non_0157.xml  non_0157_traceinfo_in.454.xml
>mv non_0157.fasta.qual non_0157_in.454.fasta.qual
>mv non_0157.fasta non_0157_in.454.fasta
```

Fire off mira:

```
mira --project=non_0157 --job=denovo,genome,accurate,454 >&log_assembly &
```

or

```
mira --project=non_0157 --job=denovo,genome,accurate,454 -GE:not=6:kcim=yes >&log_assembly &
```

Generates

```
-rw-rw-r-- 1 greg greg 1612330844 Aug 25 04:49 non_0157_out.caf
-rw-rw-r-- 1 greg greg 19702684 Aug 25 04:49 non_0157_out.wig
-rw-rw-r-- 1 greg greg 20298482 Aug 25 04:49 non_0157_out.unpadded.fasta.qual
-rw-rw-r-- 1 greg greg 6820644 Aug 25 04:49 non_0157_out.unpadded.fasta
-rw-rw-r-- 1 greg greg 20342658 Aug 25 04:49 non_0157_out.padded.fasta.qual
-rw-rw-r-- 1 greg greg 6841175 Aug 25 04:49 non_0157_out.padded.fasta
-rw-rw-r-- 1 greg greg 708184237 Aug 25 04:50 non_0157_out.ace
```

You’ll be using the ace file with BAMBUS

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Install AMOS and BAMBUS

Install [AMOS](download and use amos-2.0.8, because the ace2contig script created by the installer is necessary to convert MIRA ace file into a contig file that can be used by BAMBUS; later AMOS-shotr-tgz versions of AMOS don’t create this script). Follow directions for installing amos 2.0.8. I am installing on a 64-bit Redhat Enterprise Linux 5. The configure script complains that Qt doesn’t work correctly and GUI apps won’t run properly. I ignore this because I’m really only interested in the ace2contig script. For BAMBUS to run correctly, the BAMBUS team recommends you should pre-install:

- [XML::Parser perl module](XML::Parser perl module)
- [Config::IniFiles perl module](Config::IniFiles perl module)
- [GraphViz package](GraphViz perl module)

Just to be safe I also pre-installed

- [GraphViz perl module](GraphViz perl module)

Download bambus-2.33 and decompress. Edit Makefile file to set environment variables (within script) for BASEDIR & PERL

**There is a bug in the installation source scripts goBabmus.pl**

bambus-2.33/src/goBabmus.pl line 22

```bash
$ENV{PERLLIB} .= "::$BAMBUS_BASE/lib";
```

**should change PERLLIB to PERL5LIB**

```bash
$ENV{PERL5LIB} = "::$BAMBUS_BASE/lib";
```


Run the test case as directed in the manual to make sure BAMBUS is installed properly.

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For Ubuntu 9 users (anonymous user contribution, added 3/10)

There are problems making the grommit executable. They are remedied by the following procedure:

In the bambus-2.33/src/TIGR_Foundation_CC directory apply the following changes

ConfigFile.hh (add the line #include <string.h>)

Logger.hh  (add the line #include <stdlib.h>)

OptionResult.hh (add the line #include <stdlib.h>)

Options.hh (add the line #include <string.h>)

After changes, remake bambus from the the bambus-2.33 directory run

make clean

make all

make install

For Suse 11.2 from Lionel Guy guy.lionel@gmail.com, added 7/27/10)

To be able to compile, had to further modify the following files (in the bambus-2.33/src/

TIGR_Foundation_CC directory

FileSystem.cc: in function FileSystem::isCreatableFile (line 58)

    Changed

char * end_of_path = strrchr(filename, PATH_DELIMINATOR);

to

const char * end_of_path = strrchr(filename, PATH_DELIMINATOR);

Also, in Options.cc: added #include <cstdio>

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**Generate scaffolds**

Convert mira ace file to contig file for BAMBUS using AMOS's ace2contig

```
ace2contig -i non_0157_out.ace -o non_0157_out.contig
```

To produce contig file **non_0157_out.contig**

**Setup bambus configuration file**

Read the [configuration file help](#). Note: The mingroupsize switch/parameter doesn’t appear to work. This switch is supposed to control the minimum size scaffold the application returns. Currently, it returns all scaffolds with lots of cruft. Bug report is in about this.

Configuration file is **non_O157_out.conf**, contents below

```
# Priorities  
priority ALL 1

# The following lines can be un-commented to specify certain  
# per-library settings

# Redundancies  
# redundancy lib_some 1

# allowed error  
# error MUMmer 0.5

# min group size  
mingroupsize 50000

# Global redundancy  
redundancy 2

# overlaps allowed  
# overlaps MUMmer Y
```

**Setup mates file**

Follow [directions](#) for generating mates file. Mates file is **non_0157_out.mates** (tab delimited!), contents below

```
library twentyfiveKB    9000    40000   (.........).*
```

```gregory.harhay@ars.usda.gov```
Run BAMBUS

goBambus -c non_0157_out.contig -m non_0157_out.mates -C non_0157_out.conf -o non_0157_bambus

Once this has completed, BAMBUS will have produce a bunch of files looking similar to this

If you want a fasta file produced of the scaffolds read [http://sourceforge.net/apps/mediawiki/amos/index.php?title=Bambus_Manual - Getting more (or less) information from the output](http://sourceforge.net/apps/mediawiki/amos/index.php?title=Bambus_Manual - Getting more (or less) information from the output). A fasta file of the contig sequences (unpadded) generated by MIRA is required. This file is non_0157_out.unpadded.fasta, and is used in the command below:

```
printScaff -e non_0157_bambus.evidence.xml -s non_0157_bambus.out.xml -l non_0157_bambus.lib -f non_0157_out.unpadded.fasta -merge -o non_0157_bambus_scaffold -page -dot
```

You can edit the fasta and dot files to keep the desired scaffolds and get rid of the cruft, in this case, 1 giant 5.4 MB scaffold was produced (this is what we want) and hundreds sub-1 KB less desirable shorter scaffolds of little interest. You can use GraphVis’s dot and dotty to visualize the scaffolding dot plots.

```
untangle.pl (added 3/10)
```

It has been reported that in some installations that the untangle script doesn’t run properly and generates the following error:

```
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```
Can't locate TIGR/Foundation.pm in @INC
to locate Fountation.pm type:
locate Foundation.pm
the output of which will be the path to your Foundation.pm. In my case, it is
/usr/local/share/apps/bambus-2.33/lib/TIGR/Foundation.pm
Include this path at the beginning of the untangle.pl script as in
use lib "/usr/local/share/apps/bambus-2.33/lib/TIGR/Foundation.pm";

Hope this helps:

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