RNA-Seq Graph Builder Manual

Stefano Beretta*

March 27, 2012

RNA-Seq Graph Builder is a method to reconstruct the Splicing Graph of a gene from RNA-Seq data, without the genome information, where such a graph is a representation of the variants of alternative splicing of the gene structure.

This program predicts from NGS data the gene structure induced by the different full-length isoforms due to alternative splicing. More precisely, it analyzes RNA-Seq reads that have been sampled from the transcripts of a gene, with the goal of building a graph representation of the variants of alternative splicing corresponding to those full-length isoforms. The novelty of this method relies on the fact that it builds such a graph in absence of the genome.

1 Download and Installation

RNA-seq-Graph-Builder is implemented in C++ and is currently distributed only on source form. It has been developed on Ubuntu Linux machines (v. 10.04 and 10.10) and has been tested on both 32 and 64 bit. The program also requires the C++ library SEQAN available at http://www.seqan.de or in Ubuntu systems it is possible to install the develop package seqan-dev by typing:

\$ sudo apt-get install seqan-dev

Download

RNA-seq-Graph-Builder is developed on the AlgoLab/RNA-seq-Graph Git repository hosted by GitHub. The repository can be explored using the GitHub web interface at https://github.com/AlgoLab/RNA-seq-Graph.

^{*}DISCo, Univ. di Milano-Bicocca, Milan, Italy email: beretta@disco.unimib.it

It is also possible to clone the entire repository using the following command:

\$ git clone git://github.com/AlgoLab/RNA-seq-Graph.git

The source code is available directly in:

zip https://github.com/AlgoLab/RNA-seq-Graph/zipball/v2.0.0
tar.gz https://github.com/AlgoLab/RNA-seq-Graph/tarball/v2.0.0

Compilation

The program can be compiled by issuing the command at the command prompt:

\$ make

2 Usage

The program takes as input a FASTA file with the RNA-seq data of a gene and returns the RNA-Seq Graph. The program is executed by typing the following command:

\$./bin/build_RNA_seq_graph [options] --reads <RNA-Seq_file>
where the possible options are:

```
-o <graphML_out_file> (Default: std output)
```

```
--ref_level {1-5}
```

- 1. Standard Algorithm (Default option)
- 2. Add tiny blocks
- 3. Add linking edges
- 4. Add small blocks
- 5. Refine overlapping nodes

A summary of the available program options can be printed by invoking:

\$./bin/build_RNA_seq_graph

without parameters, or:

\$./bin/build_RNA_seq_graph --help

Alternatively it is possible to view the debug options by typing:

\$./bin/build_RNA_seq_graph --advanced

An example of usage is:

\$./bin/build_RNA_seq_graph --reads Raw_reads_file.fa -o Out_file

3 File Formats

Input: RNA-Seq Dataset

RNA-Seq file is in FASTA format (.fa, .fas or .fasta). In particular, each line of the input file describes a single read and it is composed by at least 2 rows: the first one is the header of the read (that usually starts with '>') and the second one that contains the sequence. For example:

>B2GA004:3:100:1143:752#0/1 /source=region /gene_id=gene /gene_strand=+ GATGAAATACTACTTCTACCATGGCCTTTCCTGGCCCCAGCTCTCTTACATTGCTGAGGACGAGAATGGGAAGAT

Output: RNA-Seq Graph

The program produces as output a file in txt format that contains a list of nodes and arcs of the RNA-Seq graph. It also gives as output the same graph in GDL format (http://www.absint.com/aisee/manual/windows/node58.html). It also print on standard output the graph in GraphML format; this latter can be redirected into a file in order to visualize or export it. By default the files are RNA-seq-graph.txt and RNA-seq-graph.gdl. For example:

- \$./bin/build_RNA_seq_graph --reads Raw_file.fa > RNA-seq-graph.graphml
- If the option -o is specified the 3 files will have the specified name:
- \$./bin/build_RNA_seq_graph --reads Raw_reads_file.fa -o Out-file generates Out-file.txt, Out-file.gdl and Out-file.graphml.

4 Program Code Description

Files

The program is written in C++ and it is organized in the following set of files:

- Main: is the "main" method and contains the initial menu (with the debugging options). In this file all the procedures in the pipeline of the (standard) algorithm are called.
- read_fasta: is the file containing the procedures for building the hash table that index the reads. The function read_fasta() parses the RNA-Seq reads file, creates the entry and insets them into the table.
- build_chains: in this file there are all the procedures for the creation of the chains by using the unspliced reads (i.e. vertices of the graph). There are also the functions to print the built chains (used in debug mode) and to print the reads (in different ways: left/right table, unspliced/spliced/perfectly spliced reads) in the hash table. The function build_unspliced_chains() is invoked as second step of the (standard) algorithm in order to build chains by overlapping half unspliced reads. After that, the procedure merge_unspliced_chains() try to "fuse" chains that overlaps each other (by using the hash table.
- join_chains: in this file there are the procedures for the creation of the links among chains by using perfectly spliced reads (i.e. arcs of the graph). The function link_fragment_chains() creates those links and finally print_graph() creates the graph in output in different formats. In this file there also the following procedures:
 - check_cutted_frags(): try to look if the fragments cut from the chains in the linking phase can be added as graph vertices.
 - confirm_links(): adds the "weight" to the perfectly spliced reads by summing the frequencies of the spliced reads that identify the "same junction".
 - gap_linking(): try to add further links by considering a possible "gap" in the junction.

Data Structures

The main data structures used in the program are the two hash tables used to index the reads. In the program they are implemented as:

```
//Elements of the hash tables
struct element_table{
   table_entry* p;
   bool unspliced;
   bool half_spliced;
};

//Type of Left/Right table
typedef std::map<unsigned long long, element_table> hash_map;

//Hash tables
struct tables{
   hash_map left_map;
   hash_map right_map;
};
```

In tables there are the left_map and the right_map hash maps that are both "map" of the standard library C++ in which:

key is the left (resp. right) fingerprint of the entry reads

value is an element_table in which there is the concatenated list of read entry (table_entry*p) and two flags to indicate that the entry is unspliced or half spliced (i.e. perfectly spliced).

In figure 1 a graphical representation of the types pf the data structures is shown. In this data structures, the reads are added to the hash tables by creating a concatenated list of *table_entry* elements; in fact each *table_entry* element is an instance of the class:

```
class table_entry{
    private:
    //Table List
    table_entry* r_next, r_prev;
    table_entry* l_next, l_prev;
    //Fingerprints
    unsigned long long left_fingerprint, right_fingerprint;
    //Chain List
    table_entry* chain_next, chain_prev;
    //Sequence frequency
    long frequency;
}
```

tables

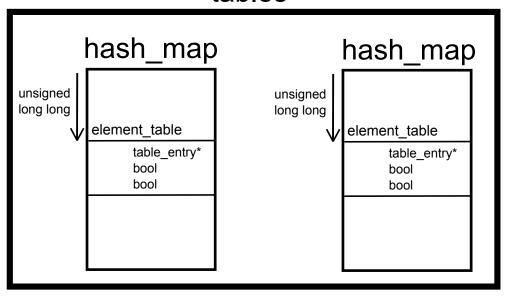


Figure 1: Types of the data structures used in the program

in which the public set/get methods are omitted.

So by using the pointers 1_next and 1_prev it is possible to build a bidirectional list of reads for the *left_map* table, in which all those reads share the same left fingerprints (i.e. the entry collision of the left hash table). The same *table_entry* object is also part of the bidirectional list of the right hash table and this is done by the two pointers r_next and r_prev for the reads that share the same right fingerprint. So each object is part of two list simultaneously (left and right).

The other table_entry pointers (chain_next and chain_prev) are the one used in the chain composition. In particular they link the current (unspliced) read to the (unspliced) one that share its left fingerprint with the right fingerprint of the current read (chain_next), and to the one that share its right fingerprint with the left fingerprint of the current read (chain_prev). The other values are the left and right fingerprints of the read and the frequency (i.e. the number of occurrences) of the read.

5 Memory Profiling

In order to monitor the memory consumption of the program we have used the library for the memory profiling libmemusage.so. This library can be preloaded using LD_PRELOAD and will intercept calls to malloc, free, realloc and various other calls. In short it will trace memory allocations for you:

\$ LD_PRELOAD=/lib/libmemusage.so build_RNA_seq_graph --reads Raw_reads_file.fa

This commando will output on *std error* the memory usage of the program on the Raw_reads_file.fa dataset. An example of the reported output is the following:

```
Memory usage summary: heap total: 3655402386, heap peak: 183838093, stack peak: 7264
        total calls
                     total memory
                                    failed calls
          47158192
                       3655402386
malloc|
realloc|
                 0
                                             0
                                                (nomove:0, dec:0, free:0)
 callocl
                 0
                               0
                                             0
   free
          47158202
                       3655402386
Histogram for block sizes:
   0-15
                  9135
                       <1%
                       <1%
   16-31
                 57130
   32 - 47
               2592145
                        5% =====
   48-63
              22528625
                       64-79
                  5686
                       <1%
  80-95
               5897106
                       12% =======
   96-111
              12973065
                       27% ===========
                       <1%
  112-127
                 14072
  128-143
               2858573
                        6% =====
  144-159
                 53042
                       <1%
  160-175
                  3898
                       <1%
                 10726
  176-191
                       <1%
  192-207
                  1903
                       <1%
  208-223
                  8626
                       <1%
                  2663
                       <1%
  224-239
  240-255
                  6907
                       <1%
  256-271
                  1417
                       <1%
  272-287
                 32463
                       <1%
                  2367
  288-303
                       <1%
  304-319
                  4223
                       <1%
  320-335
                   736
                       <1%
  336-351
                  4181
                       <1%
                       <1%
  352-367
                  1567
  368-383
                  2363
                       <1%
                   500
  384-399
                       <1%
```

The heap peak value indicates the maximum number of Bytes used by the program, i.e. the memory requirement: in the previous example it is 183838093 Bytes, which is $\sim 175 \text{MB}$.

To use libramusage all you have to do is to prepend MEMUSAGE_OUTPUT= mytrace and LD_PRELOAD=/lib/libramusage.so to your application. This will instruct the library to write out a trace to the mytrace file.

This trace file can be converted to a graph using the memusagestat utility. It is not installed by most GNU distributions and can be either build from the glibc sources or from the QtWebKit performance measurement utilities. Using:

\$ memusagestat -o output.png mytrace

an image with memory allocations and stack usage like the one at the end of this post will be created (i.e. ouput.png). The redline is the heap usage, the green one is the stack usage of the application. The x-scale is the number of allocations. The memory occupation of the previous example is reported in figure 2.

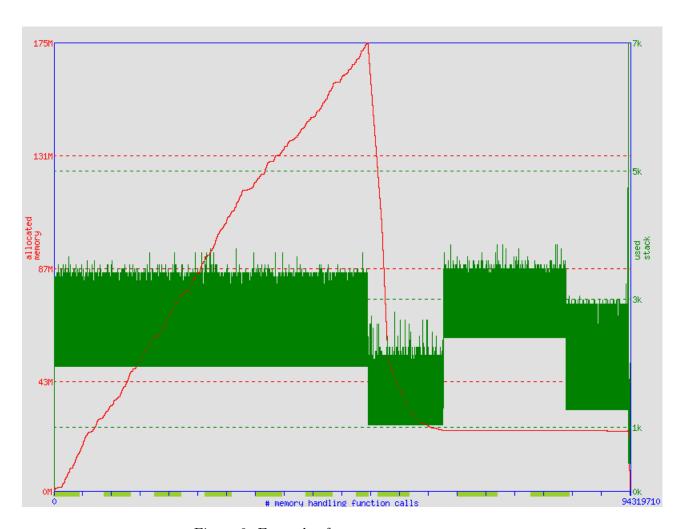


Figure 2: Example of memory usage output