## Genome analysis

Advance Access publication March 3, 2011

# **GBOOST:** a GPU-based tool for detecting gene–gene interactions in genome–wide case control studies

Ling Sing Yung\*, Can Yang, Xiang Wan and Weichuan Yu\*

Laboratory for Bioinformatics and Computational Biology, Department of Electronic and Computer Engineering, The Hong Kong University of Science and Technology, Clear Water Bay, Kowloon, Hong Kong, China

Associate Editor: Alfonso Valencia

### ABSTRACT

**Motivation:** Collecting millions of genetic variations is feasible with the advanced genotyping technology. With a huge amount of genetic variations data in hand, developing efficient algorithms to carry out the gene–gene interaction analysis in a timely manner has become one of the key problems in genome-wide association studies (GWAS). Boolean operation-based screening and testing (BOOST), a recent work in GWAS, completes gene–gene interaction analysis in 2.5 days on a desktop computer. Compared with central processing units (CPUs), graphic processing units (GPUs) are highly parallel hardware and provide massive computing resources. We are, therefore, motivated to use GPUs to further speed up the analysis of gene–gene interactions.

**Results:** We implement the BOOST method based on a GPU framework and name it GBOOST. GBOOST achieves a 40-fold speedup compared with BOOST. It completes the analysis of Wellcome Trust Case Control Consortium Type 2 Diabetes (WTCCC T2D) genome data within 1.34 h on a desktop computer equipped with Nvidia GeForce GTX 285 display card.

**Availability:** GBOOST code is available at http://bioinformatics.ust .hk/BOOST.html#GBOOST.

Contact: timyung@ust.hk; eeyu@ust.hk

**Supplementary information:** Supplementary data are available at *Bioinformatics* online.

Received on December 30, 2010; revised on February 24, 2011; accepted on February 26, 2011

## **1 INTRODUCTION**

Interaction patterns of single-nucleotide polymorphism (SNP) can be used to interpret genetic disease risks in individuals. With the advances of technologies, the burden of collecting genome-wide DNA sequence variations has been lifted. The burst of genotype data leads to an urge of efficient algorithms to analyze many genome-wide association studies (GWAS) data in a reasonable period of time. A comprehensive review (Cordell, 2009) summarized some popular methods in detecting gene–gene interactions. PLINK was recommended as the most computationally feasible method for detecting gene–gene interaction in genome-wide data. It was reported that PLINK finished the pairwise interaction examination of 89 294 SNPs selected from the WTCCC Crohn disease dataset with ~5000 samples in 14 days (Cordell, 2009). Recently, Wan *et al.* (2010) proposed a fast method, named BOOST, to examine all pairwise interactions in genome-wide case–control studies. BOOST completed the pairwise interaction analysis of human genome data with  $\sim$ 350 000 SNPs and  $\sim$ 5000 samples on a computer with 3 GHz central processing unit (CPU) and 4 GB memory in 60 h.

However, we can foresee that the growth of data will overwhelm BOOST in the near future. Ma *et al.* (2008) suggested that the analysis time in GWAS can be largely reduced by parallel computing. The development of graphical processing units (GPUs) enables modern display cards to have hundreds of cores, providing a high memory bandwidth at a low price. A recent GPU implementation of the multifactor dimensionality reduction (MDR) method (Greene *et al.*, 2010) has significantly reduced the time required for detecting gene–gene interactions. The demand of massive memory operations in collecting contingency tables and independence of analyzing different SNP pairs in BOOST make it suitable to implement BOOST in GPU. Here we propose GBOOST, a GPU implementation of BOOST. GBOOST is able to finish the genome-wide interaction analysis of a typical dataset on a single workstation within a few hours.

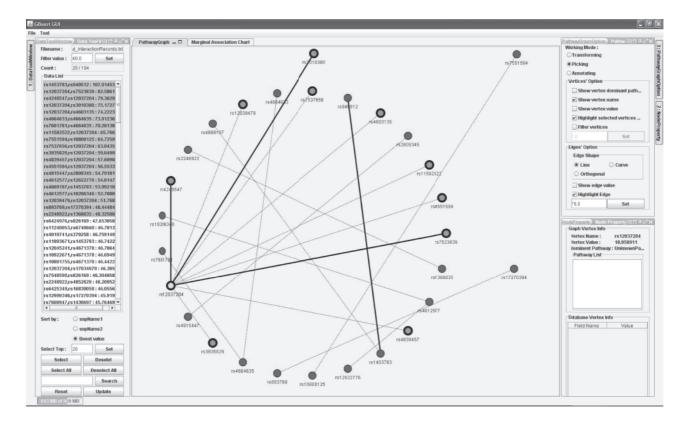
# 2 METHODS

GBOOST is a software package targeting at gene–gene interaction analysis of large genome data. It is a C++ parallel implementation of the BOOST method using Compute Unified Device Architecture runtime application programming interface (Corporation, 2008).

BOOST	GBOOST
Input Stage <ul> <li>Load and convert data into bit representa</li> <li>Screening and Testing</li> </ul>	tion.
<ul> <li>For each SNP pair :</li> <li>(Step 1) Collect the contingency table by bitwise operations with a pre-computed look-up table.</li> <li>(Step 2: Screening) Approximate the likelihood ratio test statistic. If the score is smaller than a pre-defined threshold, then skip step 3.</li> <li>(Step 3: Testing) Evaluate the exact solution using the classical likelihood ratio test.</li> <li>Output Stage</li> </ul>	<ul> <li>This screening step is parallelized by applying the following GPU techniques :</li> <li>Maximize coalesced memory access for the data in global memory.</li> <li>Use Hamming weight for bit string counting.</li> <li>Use branch prediction for consistent computer instructions.</li> <li>The testing step is the same as that in BOOST.</li> </ul>
<ul> <li>Output stage</li> <li>Output the SNP pair indexes and interacti</li> </ul>	on scores.

Fig. 1. Main steps in BOOST and GBOOST. GBOOST parallelizes the screening step in BOOST to achieve a speedup of 40.

<sup>\*</sup>To whom correspondence should be addressed.



**Fig. 2.** A snapshot of a pathway graph generated from 20 interaction pairs from one GBOOST result. Each node is labeled by its SNP name and the node value is the marginal association score from the association analysis. The edge value is the interaction score of the linked nodes (i.e. SNP pairs). Scaling, rotation and translation are available in the pathway graph. Various functions are also available to remove or highlight components in the pathway graph. The layout can be interactively customized.

The computational burden of BOOST lies in the screening stage. Thus, GBOOST modifies input data structures and parallelizes computations in the screening stage. Figure 1 gives the key differences between BOOST and GBOOST. Please refer to the Supplementary data for the detailed implementation of GBOOST.

## **3 RESULTS AND DISCUSSION**

Table 1 shows the performance of BOOST and GBOOST on different datasets. It also provides basic visualization by using two publicly available libraries JFreeChart (http://www.jfree .org/jfreechart) and JUNG (http://jung.sourceforge.net/). Figure 2 presents a pathway graph example generated from one GBOOST result.

In our future work, we plan to extend GBOOST to support execution on multiple GPUs and explore new memory optimization techniques.

#### ACKNOWLEDGEMENTS

We thank the editor and the anonymous reviewers for their constructive suggestions and comments.

*Funding*: This work was partially supported with grants RPC10EG04 and PCF004.09/10 from the Hong Kong University of Science and Technology.

Conflict of Interest: none declared.

Table 1. Running time of BOOST and GBOOST on different datasets

	BOOST	GBOOST
n = 5000, p = 5000	42 s	1.04 s
n = 5000, p = 10000 n = 5003, p = 351542	170 s 60 h	4.11 s 1.34 h

BOOST is tested on a computer with 3 GHz CPU. GBOOST is tested on a computer with a GTX 285 display card. Here n denotes sample size and p denotes number of SNPs.

## REFERENCES

- Cordell,H. (2009) Detecting gene–gene interactions that underlie human diseases. *Nat. Rev. Genet.*, **10**, 392–404.
- NVIDIA. (2008) NVIDIA compute unified device architecture programming guide version 2.1. *Technical report*. Availabel at: http://developer.download.nvidia.com/ compute/cuda/2\_1/toolkit/docs/NVIDIA\_CUDA\_Programming\_Guide\_2.1.pdf.
- Greene,C. et al. (2010) Multifactor dimensionality reduction for graphics processing units enables genome-wide testing of epistasis in sporadic ALS. *Bioinformatics*, 26, 694.
- Ma,L. et al. (2008) Parallel and serial computing tools for testing single-locus and epistatic SNP effects of quantitative traits in genome-wide association studies. BMC Bioinformatics, 9, 315.
- Wan,X. et al. (2010) BOOST: A boolean representation-based method for detecting SNP-SNP interactions in genome-wide association studies. Am. J. Hum. Genet., 87, 325–340.