


Guest Editorial for the 17th Asia Pacific Bioinformatics Conference

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THIS special section of *IEEE/ACM Transactions on Computational Biology and Bioinformatics (TCBB)* is a collection of eight papers presented at the 17th Asia Pacific Bioinformatics Conference (APBC), which was held in Wuhan, China, 14–16 January 2019. The conference aims to bring together researchers, academics, and industrial practitioners to interact. Its scientific program included three keynote, eight highlight, 51 contributed oral and 33 contributed poster presentations. The conference drew close to 150 attendees from all over the world.

The seven papers in this special section cover diverse topics, including gene tree-species tree reconciliation, genome evolution, data structures for genome assembly, protein binding-site inference and Boolean network modelling of protein-protein networks.

Gene tree-species tree reconciliation is an approach to inferring evolutionary history for gene families. It is also used to investigate the evolutionary histories of parasites and hosts as well as of species and their biogeographical habitats. In this approach, the discordance of a gene tree and a species tree is visualized by mapping the gene tree onto the species tree and is measured by the minimum number of postulated evolutionary events (e.g., gene duplication, gene loss, horizontal gene transfer) that are necessary for reconciling the trees, called the reconciliation cost of the two trees. In “The Unconstrained Diameters of the Duplication-Loss Cost and the Loss Cost,” Gorecki, Eulenstein, and Tiuryn study a few mathematics and algorithmic issues of the maximum reconciliation cost between two arbitrary binary trees in the duplication-loss model and the gene loss model.

In “Reconciliation Reconsidered: In Search of a Most Representative Reconciliation in the Duplication-Transfer-Loss Model,” Grueter *et al.* studies how well a single reconciliation can represent the entire space of optimal reconciliations.

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They answer an open problem of Huber *et al.* (Inform. Process. Letts, vol. 136, 2018) by developing an efficient algorithm for computing a median reconciliation under the path distance metric in the duplication-transfer-loss model. They also present an efficient algorithm for determining the maximum possible average support value that an optimal reconciliation can achieve.

In “Multiple Optimal Reconciliations under the Duplication-Loss-Coalescence Model,” Du *et al.* presents new algorithms for computing the size of the space of the parsimony reconciliations under the duplication-loss-coalescence model and for uniformly sampling from this space. The proposed algorithms are efficient, with runtime polynomial in the sizes of the species tree and gene tree when the number of genes that map to any given species is fixed. De Bruijn graph is a data structure that is frequently used to design efficient computer assembler for de novo genome sequencing. It also plays an important role in metagenomics analysis. In “[D]eGSM: Memory Scalable Construction of Large Scale de Bruijn Graph,” Guo *et al.* develops a lightweight parallel approach to constructing de Bruijn graph of millions of reads. With its high scalability and efficiency, the construction method has great potential in large scale genomics study.

Boolean networks that are used to model different biological systems are usually large, dense and are of modular structure. In “Towards Optimal Decomposition of Boolean Networks,” Su, Pang, and Paul propose a novel method for decomposition of BNs to balance the relation between structure and dynamics of a network.

Inverse tandem duplication random loss (iTDR) event is proposed by researchers to model the gene order evolution of the mitochondrial genome, in which a continuous segment of a gene order is duplicated and inverted, followed by the random loss of one of the redundant copies for each gene. In “Sorting Signed Permutations by Inverse Tandem Duplication Random Losses,” Hartmann, Banach, and Middendorf study the algorithmic issues of genome rearrangement in iTDR model. They show that a shortest rearrangement scenario that transforms a gene order into another can be obtained in quasilinear time and the length of such a scenario can be calculated in linear time.

Flavin mono-nucleotide (FMN) is a protein that is involved as a co-factor in carrying and transferring electrons in cellular respiration. In “Prediction of FMN Binding Sites in Electron Transport Chain based on 2-D CNN and PSSM Profiles,” Nguyen and Le propose a deep learning model for FMN binding sites using a two-dimensional

convolutional neural network and position specific scoring matrices profiles.

We would like to thank the program committee members and external researchers who reviewed the selected papers: Tatsuya Akutsu, Paola Bonizzoni, Pawel Gorecki, Katharina Huber, Sun Kim, Manuel Lafond, Kui Lin, Minsik Oh, Laxmi Parida, Mingfu Shao, Jens Stoye, Glenn Tesler, Lusheng Wang, and Shuqin Zhang. Lastly, special thanks go to the editorial staff of the *IEEE/ACM TCBB* for assistance in publishing this special section of the APBC'2019.

Guest Editors

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Shaoliang Peng is currently the executive director and a Changjiang Scholars professor with the National Supercomputing Center and College of Computer Science and Electronic Engineering, Hunan University, Changsha, China. He has authored or coauthored dozens of academic papers in several internationally influential journals, including *Science*, *Nature Communications*, *Cell*, *The American Journal of Human Genetics*, *Nucleic Acids Research*, *Genome Research*, *Bioinformatics*, *ACM/IEEE Transactions*, *Bioinformatics*, and

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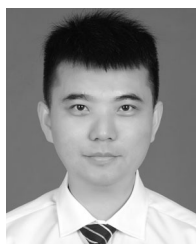
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