

Victorian Centre for Biostatistics**Seminar****Thursday 21st August 2014****9.30am to 10.30am****Melbourne School of Population and Global Health, Melbourne University****Room 515, Level 5,****207 Bouverie Street, Carlton****Exploring the structure of whole-genome conservation profiles using Bayesian segmentation****Dr Jonathan Keith****Monash University**

Conservation is a key indicator of function in genomes, and can potentially be used to discover novel functional non-protein-coding RNAs and regulatory sequences. However, recent investigations have demonstrated that a simple dichotomy between conserved and non-conserved sequence is too naïve a distinction to reflect the full complexity of the numerous types of structural and functional constraints acting on genomes. This presentation will discuss recent investigations into the detailed structure of whole-genome conservation profiles, using Bayesian segmentation techniques to identify multiple classes of conservation level. By integrating information about conservation with profiles of other properties indicative of function, including GC content and transition/ transversion ratios, a much finer level of structure can be detected. The method has been applied to a range of species including *Drosophila*, zebrafish, malaria and bacterial genomes, and results from each of these will be presented. One key implication of these results is that the proportion of functionally constrained sequence in eukaryotic genomes may be very much larger than previously supposed. Another key implication is that genomic sequences may be subject to ephemeral functional constraints that act on too short a time scale to be detected in most comparative genomic studies. The functional content of various classes of conserved sequence will also be discussed.

Dr Jonathan Keith works in the School of Mathematical Sciences at Monash University as a Senior Lecturer. He is a researcher in Bayesian methods, bioinformatics and genetic epidemiology. His current main research interest is in statistical methods for the detection of novel non-protein-coding functional elements in genomes. He also has current and recent projects in phylogenetics, whole genome association studies and identification of quantitative trait loci.

He is currently a chief investigator on two ARC Discovery grants: Statistical methods for detection of non-coding RNAs in Eukaryote Genomes and Statistical methods for discovering RNAs contributing to human diseases and phenotypes.

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VicBiostat is a Centre of Research Excellence in biostatistics funded by Australia's National Health & Medical Research Council (NHMRC). The Centre is a collaboration between biostatistical researchers at the Murdoch Childrens Research Institute, the Department of Epidemiology & Preventive Medicine at Monash University, and the Centre for Epidemiology & Biostatistics (CEB) at The University of Melbourne.