

数学与系统科学研究院

计算数学所学术报告

报告人: **Prof. Steve Qin**

**(Department of Biostatistics and
Bioinformatics Emory University,
USA)**

报告题目:

**Bayesian model-based methods for
analyzing ChIP sequencing data**

邀请人: 唐贻发研究员

报告时间: **2010年7月26日(周一)**

上午 10: 00

报告地点: **科技综合楼三层 311**

计算数学所报告厅

Abstract:

Protein-DNA interaction constitutes a basic mechanism for genetic regulation of target gene expression. Deciphering this mechanism is challenging due to the difficulty in characterizing protein-bound DNA on a genomic scale. The recent arrival of ultra-high throughput sequencing technologies has revolutionized this field by allowing quantitative sequencing analysis of target DNAs in a rapid and cost-effective way.

ChIP-Seq, which couples chromatin immunoprecipitation (ChIP) with next-generation sequencing, provides millions of short-read sequences, representing tags of DNAs bound by specific transcription factors and other chromatin-associated proteins.

The rapid accumulation of ChIP-Seq data has created a daunting analysis challenge. Here we propose a hidden Markov model (HMM)-based algorithms to detect genomic regions that are significantly enriched by ChIP-Seq. We also propose a multi-level hierarchical HMM that will allow integration of data from both ChIP-Seq and ChIP-chip experiments. Finally, we will discuss some issues related to post-processing ChIP-Seq data to obtain new biological insights.

欢迎大家参加!