

# **POSTER PRESENTATION**

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# Understanding molecular recognition and epitope prediction from Information Theoretic approach

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From UT-ORNL-KBRIN Bioinformatics Summit 2010 Cadiz, KY, USA. 19-21 March 2010

## **Background**

Cellular immunity is dependent on T-cell recognition of peptide/major histocompatibility complex (MHC) and is a critical molecular recognition component [1]. A large class of bioinformatics tools facilitates the identification of T-cell epitopes to specific MHC alleles. However, not all peptide residues contribute equally or are relevant to binding due to polymorphism of genes encoding MHC, making development of statistical methods difficult. Information Theory has proved to be one of the most universal mathematical theories that governs virtually all processes [2]. The success of this approach in analyzing a huge range of engineering, technological and natural processes is impressive. In Molecular Biology the applications have been very successful at the sequence level, many sequence comparison and binding site identification methods now boasts a sound information theoretic foundation.

### Materials and methods

In this work we have developed a mathematical formalism for applying information theory in identifying an explicit computational strategy and developing algorithms for the study of peptide/MHC interactions through epitope predictions. A sampling method has been initiated to circumvent the binding problem. Comparisons have been made with existing Machine Learning Methods and a validation of the efficiency of the model may be tested [3,4]. The results will have significant impact for understanding the immune system and for rational drug design [5].

### Acknowledaments

This work was partially supported by DOD grant W81XHW-05-01-0227 received by YC. Authors would also like to thank Dr. IrisAntes, Technical University, Munich for helpful discussions.

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Published: 23 July 2010

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### doi:10.1186/1471-2105-11-S4-P22

Cite this article as: Mitra and Cui: Understanding molecular recognition and epitope prediction from Information Theoretic approach. *BMC Bioinformatics* 2010 **11**(Suppl 4):P22.

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