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# Development of a Viral Biological-Threat Bioinformatics Resource

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### Abstract

In response to the potential use of viruses as biological weapons, we have established the Viral Biological-threat Bioinformatics Resource (VBBR) that collects, catalogs, annotates, and analyzes genetic information related to potential viral threats. This work expands upon available knowledge of virus replication, pathogenicity, and virus-host interactions on the basis of individual protein domains, individual genes, and whole genomes. To date, we have constructed a genome and gene sequence database that has been populated with the sequence information for viruses currently listed on the NIH and CDC priority pathogen list. We have also developed a variety of analytical and visualization tools that aid in the analysis of the genomic information coded for by these viruses. Finally, the information developed as a result of this work has been made available to the scientific community through a (currently access-controlled) web site (http://vbbr.genome.uab.edu) that supports research efforts to develop environmental detectors, diagnostics, antiviral compounds, new vaccines, and animal models in support of biodefense research goals.

### Subject Terms
Biodefense, biological terrorism, bioinformatics, viruses, sequence database

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INTRODUCTION:

In response to the potential use of viruses as biological weapons, we have established the Viral Biological-threat Bioinformatics Resource (VBBR) that collects, catalogs, annotates, and analyzes genetic information related to potential viral threats. This work expands upon available knowledge of virus replication, pathogenicity, and virus-host interactions on the basis of individual protein domains, individual genes, and whole genomes. This information has been made available to the scientific community to support current research efforts to develop environmental detectors, diagnostics, antiviral compounds, new vaccines, and animal models.

Specifically we are:

1. developing a relational database that supports the data storage, annotation, analysis, and information exchange goals of this proposal;

2. developing a world-wide web site that will allow all information and analytical results compiled and generated by this work to be publicized to the scientific community, providing rapid access to specific information as required by an individual researcher;

3. collecting existing gene and genomic sequences for viral threat agents and importing them into the database for subsequent annotation and analysis;

4. providing computer-automated and human-directed annotation to confirm and update existing biological information in the sequence records for all sequences; and

5. performing a variety of analyses on genome and gene sequences to provide additional information on their structure, function, and evolution.

BODY:

Substantial progress has been made in accomplishing many of the tasks outlined in the original statement of work. Accomplishments extend to database construction, data population, development and implementation of analytical and visualization tools, and publication of all available information on a web site. The accomplishments are listed below, itemized according to the original statement of work and task list.

**Task 1.** To develop a relational database that will support the data storage, annotation, analysis, and information exchange goals of this proposal. (Months 1-6)

- The database has been created based on our previous work on poxviruses and has been updated and refined to better support a wider range of virus threats. An overview of the database schema is provided in figure 1.
Task 2. To develop a world-wide web site that will allow all information and analytical results compiled and generated by this work to be publicized to the scientific community, providing rapid access to specific information as required by an individual researcher. This web site will also provide the necessary security and access controls to ensure that only individuals and groups as designated by the granting agency (USAMRMC) can utilize these resources. (Months 1-12)

- The web site has been established and is accessible at the url: http://vbbr.genome.uab.edu.
- Login to the web site requires registration. Following registration, a user account is established, and access is secured via a username/password. To register, a user needs to
logon to the following web page: https://btd.genome.uab.edu/register/register.asp using the following credentials: Username: genuab; Password: $mgbf2003 and fill out the form. The user will be vetted for access and when approved will then be notified via Email when the account is established.

- The home page for the VBBR web site is shown below in figure 2.

![VBBR Web Site](image)

**VBBR**

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*teams & people | address book | acknowledgements | feedback*

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**Figure 2. VBBR Web Site**

**Task 3.** To collect existing gene and genomic sequences for viral threat agents and import them into the database for subsequent annotation and analysis (Months 1-24).

a. We will obtain existing gene and genomic sequences for all viruses that potentially could be used as agents of biological warfare or terror. The
information collected will include the corresponding descriptive annotation provided with the sequences.

- An automated parser has been developed (XSeq Exchange) that can parse XML-formatted GenBank sequence records and their annotation and load that information into the VBBR database. This application will greatly enhance our ability to add new viral genomes to the VBBR database. The database schema for the parser is shown below in figure 3.

Figure 3. XSeq Exchange Database schema
The genome and gene database is currently populated with most of the viruses currently on the NIAID A-C priority pathogen list. These are shown on the genomes page of the VBBR web site as displayed in figure 4.
b. We will also collect and provide the references for publications in the scientific literature that describe studies on the structure and function of orthopoxvirus genes and genomes. We will link this information on our web site to the appropriate gene records in our database.

- Not yet done. Tasked for year 2.

c. We will initially obtain viral sequence from public databases such as GenBank. We will also obtain unpublished sequence information through inquires and interactions with other scientists that have been initiated by our collaborators.

- Not yet done. Tasked for year 2.

Task 4. To provide computer-automated and human-directed annotation to confirm and update existing biological information in the sequence records for existing as well as newly obtained sequences. (Months 6-24)

- Computer-automated annotation has been completed for the viruses listed in figure 4. Human directed annotation is tasked for year 2. An example annotation record is shown in figure 5.

Task 5. To perform a variety of analyses on genome and gene sequences to provide additional information on their structure, function, and evolution. (Months 6-24)

- Initial computer-aided analysis has been completed. Additional analyses taskd for year 2.

Task 6. To provide through the web site, a set of analytical and visualization tools that will allow users of this resource to mine the data for useful information, to perform comparative analyses between these and other viruses, and to better visualize and assess the significance of their results. (Months 6-24)

- Currently available analytical and visualization tools include:
  a. Genome map visualization (Figure 6.)
  b. Genome protein gene ortholog comparisons
  c. Gene synteny visualization (Figure 7.)
  d. BLAST similarity searches
  e. BLAST search parsing and database storage
  f. Java-based Visualization of BLAST search results (Figure 8.)

- A new database and web server has been installed and is now supporting all VBBR resources. This new server provides added security and has greatly increased performance of the VBBR web site.

- Development of additional analytical and visualization tools are tasked for year 2. These will include tools for creation and visualization of multiple sequence comparisons and tools for the inference and visualization of evolutionary trees.
Figure 5. VBBR Gene Annotation Record.
**Figure 6. VBBR Genome Map.**

**Figure 7. VBBR Gene Synteny Map.**
Figure 8. VBBR BLAST Viewer. The figure shows the results of a BLASTX search of the SARS genome against the Genbank non-redundant protein database.
KEY RESEARCH ACCOMPLISHMENTS:

- Establishment of the VBBR Database
- Establishment of the VBBR Web Site
- Development of the XSeq Exchange GenBank sequence parser
- Population of the VBBR database with virus genomic information
- Computational annotation of virus genes
- Implementation of analytical and visualization tools for BLAST searches
- Implementation of analytical and visualization tools for genome ortholog comparisons

REPORTABLE OUTCOMES:

Informatics:

- VBBR Database
- VBBR Web site (http://vbbr.genome.uab.edu)
- VBBR analytical and visualization tools

Presentations:

- USAMRIID/WRAIR/LANL/UAB Workshop, August 11, 2003 WRAIR
- USAMRMC Bioinformatics Workshop, November 4, 2003 (Poster)

Funding proposals submitted extending this work:

- NIH/NIAID Proposal in response to the RFP: NIH-NIAID-DMID-04-34, "Bioinformatics Resource Centers for Biodefense and Emerging/Re-emerging Infectious Diseases"

CONCLUSIONS:

Development of the Viral Biological-threat Bioinformatics Resource (VBBR) has substantially followed the original task list and essentially all tasks proposed for year 1 have been accomplished. The result is a Bioinformatics Resource Center that can now begin to support the needs of basic and applied biodefense research directed at gaining a better understanding of the pathogenesis, evolution, and overall biology of viral pathogens as well as support the development of detectors, diagnostics, antivirals, and vaccines. Future work will follow the original task list and is aimed at providing human-curated gene records along with additional analytical and visualization tools that can better support understanding the role of individual genes in virus pathogenicity.

REFERENCES:

None other than the VBBR web site.

APPENDICES:

None