Case Studies in Data Intensive Computing: Large Scale DNA Sequence Analysis as the Million Sequence Challenge and Biomedical Computing

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Abstract

Many areas of science are seeing a data deluge coming from new instruments, myriads of sensors and exponential growth in electronic records. We take two examples – one the analysis of gene sequence data (35339 Alu sequences) and other a study of medical information (over 100,000 patient records) in Indianapolis and their relationship to Geographic and Information System and Census data available for 635 Census Blocks in Indianapolis. We look at initial processing (such as Smith Waterman dissimilarities), clustering (using robust deterministic annealing) and Multi Dimensional Scaling to map high dimension data to 3D for convenient visualization. We show how scaling pipelines can be produced that depending on data set size, either use multicore laptop or desktop clients or supercomputer or modest clusters for the computer intensive sections. This study illustrates challenges in integrating data exploration tools with a variety of different architectural requirements and natural programming models. We present preliminary results for end to end study of two complete applications.

# 1. Introduction

Data Intensive Computing is very popular at this time. Partly this is due to the well understood data deluge with all activities including science, government and modern Internet (Web 2.0) systems all generating exponentially increasing data. One special driver is that Web Search and related data mining can use an especially simple programming model MapReduce of which there are now several implementations. It is attractive to understand how generally applicable MapReduce is to other data intensive problems as one can expect excellent commercial support for software in this area. In previous papers we have looked at the impact of clouds and compared Yahoo (Hadoop) and Microsoft (Dryad) implementations of the MapReduce step. These technologies are still maturing and their performance may not be so important at this stage compared to the overall architecture of a complete system extending from raw data to scientific discovery. We choose two biomedical applications. One studies the structure of Gene families and the processing steps involve sequence alignment, clustering and visualization after projecting sequences to 3 dimensions using Multidimensional scaling MDS. The second application involves correlating electronic patient records with environmental information (from Geographical Information Systems)associated with the patient location. Here the end to end study involves substantial data validation, processing with many standard tools such as those in R but also many possible other applications such as Multidimensional Scaling dimension reductions.

We present performance results from Tempest – An Infiniband connected 32 node system running Windows HPCS with each node having 24 cores spread over 4 Intel chips. Such a modest cluster can fully process all stages of the 35,000 element Alu study in less than a day and is suitable for up to 200,000 sequences even though all steps in analysis are of O(N2) time complexity. We estimate that a 1024 node Tempest architecture cluster would tackle well our million sequence goal. We find systems easy to use and and program as well as giving good wall clock execution time. Some of our studies used a slightly older cluster Madrid with 8 nodes each with four AMD Opteron chips with 4 cores each.

Section 2 presents some overall architecture comments while sections 3 and 4 describe the two main applications. Section 5 has conclusions and future work.

# 2. Data intensive computing architecture

Table 1. Hardware and software configurations of the clusters used in this paper. In addition a traditional 8-node Linux Cluster “Gridfarm” was used to run statistics package R in section 4.



The computer architecture needed to support data intensive computing is obviously complex and varied. Here we do not discuss virtualization or issues of distributed systems which although important are not the topic of this paper. We abstract many approaches as a mixture of pipelined and parallel (good MPI performance) systems, linked by a pervasive storage system. Here we have many interesting possibilities including Amazon and Azure “Blob” storage, traditional supercomputer environment like Lustre plus importantly the file systems (such as Cosmos from Microsoft or HDFS from Hadoop) supporting the new MapReduce systems.



Figure 1. A Data intensive computing architecture

These cloud/Web 2.0 technologies support a computing style where data is read from one file system, analyzed by one or more tools and written back to a database or file system. An important feature of the newer approaches is explicit support for data parallelism which is needed in our applications.

In figure 1, we abstract this disc/database-compute model and assume it will underlie many applications even when some of resources will be local and others in the cloud or part of a large grid. In figures 3 and 2 we give in more detail the data pipelines used in the applications of sections 3 and 4 respectively.

Finally we record in table 1, the major facilities used in this study. Note they run Windows (HPC Edition) and stress both multicore and traditional parallelism.  
  
 Figure 2. Stages of Gene sequencing application



Figure 3. Stages of health application

The largest Tempest cluster has 768 Intel Cores spread over 32 nodes while the smaller one Madrid has 128 Opteron cores spread over 8 nodes. Our work [5, 19, 21] stresses both Windows and Linux so we can explore Hadoop, Dryad and the emerging cloud approaches. This paper focuses on results from the Windows clusters.

# 3. Gene Sequencing Applications

**3.1. Alu Sequencing Studies**

The Alu clustering problem [13] is one of the most challenging problem for sequencing clustering because Alus represent the largest repeat families in human genome. There are about 1 million copies of Alu sequences in human genome, in which most insertions can be found in other primates and only a small fraction (~ 7000) are human-specific insertions. This indicates that the classification of Alu repeats can be deduced solely from the 1 million human Alu elements. Notable, Alu clustering can be viewed as a classical case study for the capacity of computational infrastructures because it is not only of great biological interests, but also a problem of a scale that will remain as the upper limit of many other clustering problem in bioinformatics for the next few years, e.g. the automated protein family classification for a few millions of proteins predicted from large metagenomics projects.

**3.2. Smith Waterman Dissimilarities**

The first step is to identify human Alu gene sequences which were obtained by using Repeatmasker [14] with Repbase Update [15]. We have been gradually increasing the size of our projects with the current sample having 35339 sequences the largest and requires a modest cluster such as Tempest (768 cores). Note from the discussion in section 3.1, we are aiming at supporting problems with a a million sequences -- quite practical today on TeraGrid and equivalent facilities given basic analysis steps scale like O(N2).

We used open source version [16] of the Smith Waterman – Gotoh algorithm SW-G [17, 18] modified to ensure low start up effects by each thread/processing large numbers (above a few hundred) at a time. Memory bandwidth needed was reduced by storing data items in as few bytes as possible.

**3.2.1 Performance of Smith Waterman Gotoh SW-G Algorithm**

The calculation of the 624 million independent dissimilarities is of course architecturally simple as each computation is independent. Nevertheless it shows striking structure shown in figure 4. As in previous papers, we look at different patterns denoted a (Thread per process) x (MPI process per 24 core node) x (Number of Nodes). In pattern ***t***x***m***x***n***. We have for Tempest defined in table 1, ***n*** <=32 and ***t***x***m*** <= 24. We present results in terms of parallel overhead f(P) defined for Parallelism P by

f(P) = [PT(P) –T(1)] /T(1) (1)

Where T(1) is replaced in practice by T(on smallest number of processes that can run job).



Figure 4.Performance of Alu Gene Alignments for different parallel patterns

The striking result for this step is that MPI easily outperforms the equivalent threaded version of this embarrassingly parallel step. In figure 4, all the peaks in the overhead correspond to patterns with large values of thread count ***t***. On figure 4, we note that MPI intranode 1x24x32 pattern completes the full 624 billion alignments in 2.33 hours – 4.9 times faster than threaded implementation 24x1x32. This 768 core MPI run has a parallel overhead of 1.43 corresponding to a speed up of 316.

The SW-G alignment performance is probably dominated by memory bandwidth issues but we are still pursuing several points that could effect this but not at our highest priority as SW-G is not a dominant step. We have tried to identify the reason for the comparative slowness of threading. Using Windows monitoring tools, we see in figures 5 and 6 that threaded version has about a factor of 100 more context switches (note different scale factors used) than case where in MPI we have one thread per process.

Figure 5.Paging and Context Switching for a pure Threaded SW-G



Figure 6:Paging and Context Switching for an MPI SW-G

This could lead to a slow down of threaded approach and correspond to Windows handing of paging of threads with large memory footprints. However there is also an important data transfer effect that we discuss in the following subsection.

**3.2.2 The O(N2) Factor of 2 and data transfer**

There is a well known factor of 2 in many O(N2) parallel algorithms such as those in direct simulations of astrophysical stems. We initially calculate in parallel Distance D(i,j) between points (sequences) i and j and as discussed above this is done in parallel over all processor nodes selecting criteria i < j to avoid calculating both D(i,j) and the identical D(j,i). This can require substantial file transfer as it is unlikely that nodes requiring D(i,j) in a later step, will find that it was calculated on nodes where it is needed.

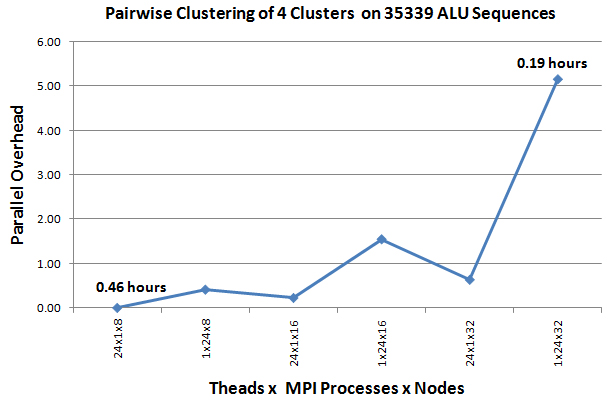
For example the MDS and PW(PairWise) Clustering algorithms described in next 2 sections, require a parallel decomposition where each of N processes (MPI processes, threads) has 1/N of sequences and for this subset {i} of sequences stores in memory D({i},j) for all sequences j and the subset {i} of sequences for which this node is responsible. This implies that we need D (i,j) and D (j,i) (which are equal) stored in different processors/disks). This is a well known collective operation in MPI called either gather or scatter. Note that we did NOT get good performance for data transfer of D(i.j) to its needed final processor from either MPI (it should be a seconds on Petabit/sec Infiniband switch) or Dryad. We intend to make the needed collective (reduction) primitives more precise and expect substantial performance improvement. However, for the results presented here the timings include the I/O necessary to write results from each process to local disk. An additional step was necessary in our processing workflow to combine the results into a single file used in downstream processing such as clustering and MDS.

**3.2.3 Relevance of Hadoop and Dryad**

We expected to use Dryad for this initial SW-G computation and data transfer but were not able to complete this work in time for this paper. Clearly this step fits MapReduce very well and as technology and our experience with it improves [19], we expect to include Dryad (on Tempest) and Hadoop (Linux) evaluations here and it is quite likely that they will deliver preferred implementation.

**3.3 Pairwise Clustering**

As data sets increase in size, we expect some applications to require particularly robust algorithms that are as insensitive as possible to well known difficulties such as “trapping in local minima”. This increases computing challenge which grows to accommodate data set and increase robustness of results. For example, clustering methods like Kmeans are very sensitive to false minima but some 20 years ago a more robust EM (Expectation Maximization) method using annealing (deterministic not Monte Carlo) was developed by Ken Rose (UCSB) [1], Fox and others [4].

Here the annealing is in distance (as represented by D(i,j) ) resolution. One slowly lowers a Temperature T that implements an algorithm sensitive to distance scales of order T0.5. This method has interesting feature that it automatically splits clusters when instabilities detected. Further it has a highly efficient parallel algorithm which we have studied in detail in earlier papers on smaller problems [5]. These clustering approaches are fuzzy methods where points are assigned probabilities for belonging to a particular cluster. There are striking differences between pattern dependence of figures 4, 7 and 8. In all cases MPI is used as communication mechanism between nodes but we can use any mix of threading and MPI on a single node. For figure 4 intranode MPI always gave best performance but in figures 7 and 8, intranode threading is the best. We have analyzed this in detail elsewhere and found it is a consequence of MPI communication overheads that increase as data parallel unit (of size 35339/(m n)) decreases. For large data parallel units MPI is fastest but for smaller ones used here, threading is superior.Figure 7: Performance of Pairwise Clustering for 4 clusters on Tempest. 10 Clusters take about 3.5 times longer

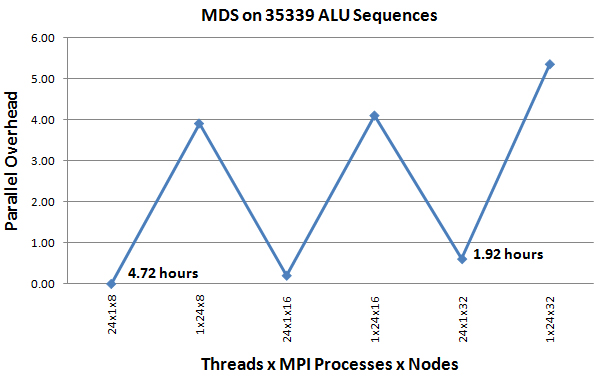
The original clustering work was based in a vector space (like Kmeans) where a cluster is defined by a vector as its center. However in a major advance 10 years ago [2, 3], it was shown how one could use a vector free approach and operate with just the distances D(i,j). This unfortunately does increase the computational complexity from O(N) to O(N2) for N sequences. It appears however more natural for studies of sequences which do not have Euclidean vectors easily associated with them. We completed these pairwise vector free algorithms and implemented them in parallel. We have discussed elsewhere detailed algorithm and performance issues. Here we report the clustering as part of a large end to end component of our “Million Sequence Analysis as a Service project. All capabilities discussed in this paper will be made available as cloud or TeraGrid services over next 3-12 months.

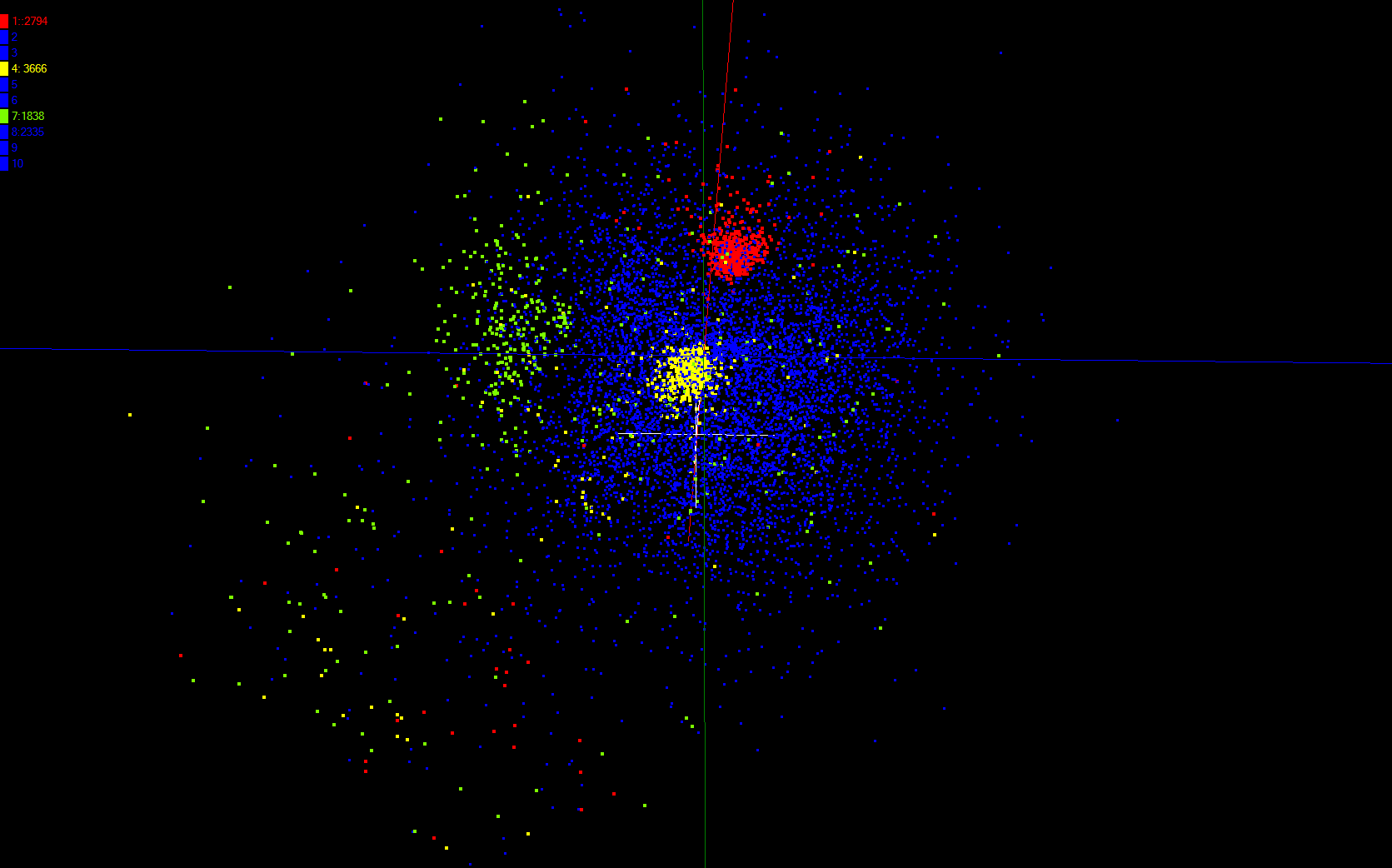
**3.4 Multidimensional Scaling MDS**

Given dissimilarities D(i,j), MDS finds the best set of vectors **x***i* in any chosen dimension d minimizing   
 Σ*i,j* weight(i,j) (D(i,j) – |**x***i* – **x***j*|n)2 (2)

The weight is chosen to reflect importance of point or perhaps a desire (Sammon’s method) to fit smaller distance more precisely than larger ones. The index n is typically 1 (Euclidean distance) but 2 also useful

We have previously reported results using Expectation Maximization and we are exploring adding to this deterministic annealing to improve robustness. Here we use a different technique exploiting that (2) is “just” χ2 and one can use very reliable nonlinear optimizers to solve it [20]. We have implemented and got good results with the Levenberg–Marquardt approach (adding suitable multiple of unit matrix to nonlinear second derivative matrix) to χ2 solution.

Figure 8: Performance of Pairwise Clustering on Tempest

 Figure 9: Clustering of Alu Sequences

This “MDS as χ2” approach allows us to incorporate some powerful features including very general choices for the weight(i,j) and n. Our MDS service is fully parallel over unknowns **x***i*. Further it allows “incremental use”; fixing an MDS solution from a subset of data and adding new points at a later time. One can also optimally align different versions of MDS (e.g. different choices of weight(i,j) to allow precise comparisons. All our MDS services feed their results directly to powerful Point Visualizer. Figure 9 shows the end to end Alu study after SW-G alignments, pairwise clustering and MDS projection. One sees three small clusters red (2794 points), yellow (3666) and green (1838 sequences) isolated from larger (27041) collection of blue sequences that are presumably older. Note that total time for all 3 steps on the full Tempest system is about 6 hours and clearly getting to a million sequences is not unrealistic and would take around a week on a 1024 node cluster.

# 4. Linking Environment and Health Data

**4.1. Introduction**

Another area where our tools are naturally used comes in Geographical information systems where we have already presented results [21]. Here we link environmental and patient (health) data. This is challenging as a community’s vulnerability and impact may depend on special concerns like environmentally sensitive areas or historical structures, socioeconomic conditions, and various social concerns such as the degree of public trust, education levels, literacy, and collective action and solidarity. The event impact must account for a blend of physical and social measures.

One example is the SAVI Community Information System (www.savi.org)1 is one of the nation’s largest community information systems [22]. SAVI, designed to improve decision-making in Central Indiana communities. SAVI includes over a ~22 million individual data values, provides over 161,322 event datasets, 3,099 basic indicators on the socio-economic conditions, health, economy, housing, and many other aspects of the community and makes them available for 11 types of geographic areas, such as census tracts, neighborhoods, and school corporations. The SAVI system now is being used by a variety of other sectors for community development, public health research, education, program planning, disaster mitigation planning and more. Only recently has the field of social epidemiology begun to develop the theoretical tools that make possible the identification of explanatory pathways from the physical and social infrastructure to health-related behaviors, which then lead to adverse health outcomes [23-25]. We see geographic clustering in many health outcomes because social environment has an effect on health and/or health behaviors [26-28]

**4.2. Obesity**

We used an ongoing childhood obesity study as our first application to test the relevance of our tools in the area of linking environment and social/health data. [6-7] Obesity is presently one of the most pervasive, serious, and challenging health problems facing the world. Over the past 30 years, the obesity rate has nearly tripled for children ages 2 to 5 years (from 5 to 14 percent) and tripled for youth ages 12 to 19 years (from 5 percent to 17 percent). The obesity rate for children 6 to 11 years of age has quadrupled from 4 to 19 percent. What is causing the dramatic and threatening rise in obesity? Bray concisely captured the etiology of obesity in metaphor: “Genes load the gun, the environment pulls the trigger.” 23 Genetic factors are thought to account for 25-40% of the variance in BMI (Body Mass Index) by determining differences in such things as resting metabolic rate and weight gain in response to overfeeding. However, it is highly improbable that changes in genetic factors explain the rapid increases in obesity prevalence over the past two decades. [26] Rather the obesity epidemic is almost certainly rooted in environmental factors that promote excessive caloric intake and sedentary lifestyle [8].

In addition to physical environmental factors, social environmental factors also have bearing on obesity by facilitating or constraining behavior. Specific social environmental factors that have been examined include crime, safety, social support, social networks, and neighborhood socioeconomic status. Perceived (or actual) lack of a safe environment is a significant barrier to physical activity. According to a study conducted by the Centers for Disease Control in 2004, persons who perceived their neighborhoods as less than extremely safe were more than twice as likely to have no leisure-time physical activity, and those who perceived their neighborhoods as not at all safe were nearly three times as likely to have no leisure-time physical activity. Research also indicates that parental concerns about traffic and crime have a strong influence on children’s physical activity levels and that child and parent perceptions of the environment are as important as the actual environment.

This motivates studies that study linkage between patient health and environment factors.. We can examine urban planning data that provides information on characteristics of the built environment, such as street features, land use mix, and neighborhood greenness. We examine insurance information from patient medical records as an indicator of family-level social environment. We examine U.S. Census and Uniform Crime Report information for areas surrounding patients’ residential addresses as indicators of neighborhood social environment. Here we are setting up the infrastructure linking the tool R with our other tools described in section 3 and only have preliminary results on this use case for a new generation of large scale data analysis tools. As there are some 30 patient attributes and over one hundred environmental attributes, tools like MDS that reduce dimensionality were our first focus.

**4.3. Canonical Correlation Analysis**

The canonical correlation analysis (CCA) is a tool of multivariate statistical analysis for finding correlations between two sets of variables [9, 10]. We are applying CCA to study how childhood obesity is mostly related with what kinds of environmental factors. Our full data set we used for this research consists of over 314,000 real-life patient records collected over 15 years (some records are 20 year old) and measured on about 180 variables, mostly related with biological and environmental factors. We stored our full data set (size of 832 MB) in a database system for easy exploration and fast extraction. Among the full data set, we only used the cleanest data for our initial studies.

For performing CCA over the patient data set and conducting various kinds of statistical analysis, we used R, one of the most well-known statistical computing environments, to expedite complicated statistical data manipulations with ease and utilize highly optimized and multi-threaded numeric packages, such as BLAS, Goto-BLAS [11], and ATLAS [12]. Another advantage in using R is that we can use various open-source based add-on packages for additional functionalities. For example, with the help of packages for databases, such as PostgreSQL and MySQL, we can directly access the data stored in the database system. Here we focus on integrating R with our specialized tools and present initial MDS analysis here.

**4.3. Multi Dimensional Scaling and Visualization**

The core of CCA is to find an optimal linear projection of two sets of data in a sense that the correlation of them in the projected space, also called “canonical space”, is maximized. More specifically, for the given two sets of data matrix X and Y, the CCA seeks two optimal projection vectors *a* and *b*, which make the following correlation maximum:

,

where  and are vectors in the canonical space. One can see that the vector U and V, known as *canonical correlation variables*, are the new representation of the data matrix X and Y in the canonical space, transformed by the projection vector a and b respectively. By further investigating U and V as a product of the CCA, one may infer cross-relationships exist between two sets of variables, which are not directly available from the row data set.

In our project, the CCA is the best match for our purpose. We have two sets of data – patient and environmental data – and want to find out which variables of environmental data have some connections to patient’s obesity or more generally health. For this purpose, we can use X as an environmental data and Y as a patient data into the CCA to find the best optimal canonical variables U and V, which maximize the correlation between the patient and the environmental data in the canonical space. Figure 10 presents one of the CCA results on our data set as an example.

As an alternative to CCA, which maximizes vector in both data sets, one can find the vectors a and b by using the Principle Component Analysis (PCA) within a single sector. For an example, with our health data set, we can find new projection vector a by fixing b in terms of Principle Components (PC) of the patient data matrix Y.

Since the well known CCA algorithm itself is not our focus in this paper, we will not present more details in As an example of CCA results to the patient data set, we found the optimal correlation in the canonical space (Figure 10). Those results can feed in to the MDS to find more robust structures in 3-dimension (Figure 11).

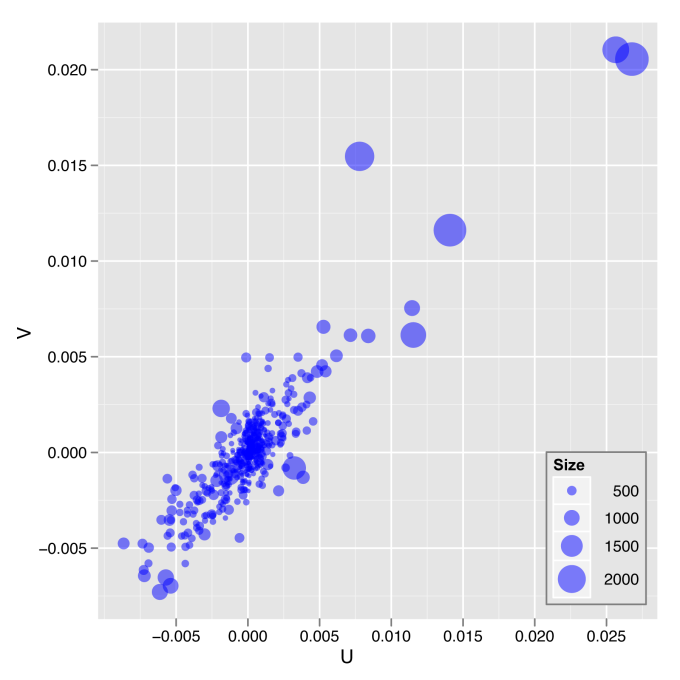


Figure 10. The plot of the first pair of canonical variables for 635 Census Blocks

Figures 11(a) and 11(b) show this CCA analysis projected on an MDS representation of environment data. this paper. More details can be found in [9, 10]. Each point corresponds to one of 635 Census blocks. We color projections on a green (lowest) to red/mauve (highest) scale and see clear clustering of the different colors in different regions of MDS. The low (green) values occur together and are well separated from the high weighted red and mauve points. In these plots the MDS was weighted (using weight(i,j) in equ. (2) proportionally to number of patients in block. Figures 11(c) and (d) show correlations for a pure principal component analysis PCA; this is high for environmental PCA (as one would expect) but still present for PCA in patient health space.

In processing CCA in our project, we have used R as statistical computing environments to utilize various matrix manipulation and linear algebra packages with efficiency. Also, by building R with multi-threaded enabled BLAS libraries, we got parallel speed up in our 8 core Linux cluster nodes “Gridfarm”. As a result, we have applied the CCA to our data set with various parameter settings and visualized them by using our parallel MDS projection described earlier. Figure 11 illustrate the MDS visualization of PCA and CCA analyses. This clearly illustrates that correlations can be seen when projecting societal data on MDS of environmental information. We are following on with both obesity and SAVI datasets with different selections of environmental data.

Figure 11. MDS visualization of PCA and CCA analyses for correlations between patient and environmental data described below.

# 5. Conclusion and Future Work

This paper examines the technology to support rapid analysis of million sequence problems that appear to typify today’s high end challenges. As well as our local sample problems, we would like to refine and test the technology on a broader range of problems. To encourage this, we will make key capabilities available as services that we eventually be implemented on virtual clusters (clouds) to address very large problems. This will require work we are doing now on Hadoop and Dryad – can they be a single unifying technology? Relevant services we will make available include the basic Pairwise dissimilarity calculations, R (done already by us and others), MDS in EM and χ2 forms; the vector and pairwise deterministic annealing clustering including support of fingerprints and other ”unusual” vectors. Our point viewer (Plotviz) will be made available either as download (to Windows!) or as a Web service. We note all our current code is written in C# (high performance managed code) and runs on Microsoft HPCS 2008 (with Dryad extensions)

We’ve shown two examples of data intensive science applications in area of biology and health using several modern technologies. We suggest that these ideas will support new generations of large scale data analysis systems for patient records, demographic data and next generation gene sequencers.

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