Metric Methods in Mining

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Content

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- Conclusions and Future Work
Preliminaries
**Dissimilarities and Metrics**

Let $S$ be a set (patients, phenotypes, clinics, etc).

**Dissimilarity:** a function

$$d : S \times S \longrightarrow \mathbb{R}_+$$

such that $d(p, q) = 0$ if and only if $p = q$.

$d(p, q)$ measures the dissimilarity between two objects; if $d(p, r) < d(p, q)$ this means that $r$ resembles more to $p$ then $q$ does.
If $d$ is a dissimilarity on $S$ such that

$$d(p, q) = d(q, p)$$

for every objects $p, q$, then $d$ is a **symmetric dissimilarity**.

A **metric** on the set $S$ is a symmetric dissimilarity that satisfies the triangular inequality:

$$d(p, q) + d(q, r) \geq d(p, r)$$

for every $p, q, r \in S$. 

Examples

- Standard distance on real line:

\[ d(p, q) = |p - q| \]

- Minkowski distance in \( \mathbb{R}^n \):

\[
 d_k(p, q) = \left( \sum_{i=1}^{n} |p_i - q_i|^k \right)^{\frac{1}{k}}
\]

for \( p = (p_1, \ldots, p_n) \) and \( q = (q_1, \ldots, q_n) \in \mathbb{R}^n \).
Examples

In $\mathbb{R}^2$:

\[
d_1(p, q) = |p_1 - q_1| + |p_2 - q_2|
\]
(Manhattan distance)

\[
d_2(p, q) = \sqrt{|p_1 - q_1|^2 + |p_2 - q_2|^2}
\]
(Euclidean distance)

\[
d_\infty(p, q) = \lim_{k \to \infty} d_k(p, q)
\]

\[
= \max\{|p_1 - q_1|, |p_2 - q_2|\}
\]
(Canberra distance)
\[ p = (p_1, p_2) \quad |p_1 - q_1| \]

\[ q = (q_1, q_2) \quad |p_2 - q_2| \]
Partitions

\[ \text{PART}(S) : \text{set of partitions of set } S \]

Partition \( \pi = \{B_1, \ldots, B_7\} \)

\( B_1, \ldots, B_7 \) are the blocks of \( \pi \)
Partitions Partial Order

\[ \sigma \leq \pi \text{ if each block } C \text{ of } \sigma \text{ is included in a block of } \pi. \]

Partition \( \sigma = \{C_1, \ldots, C_{12}\} \leq \pi \)
Let $L \subseteq S$ and $\pi = \{B_1, \ldots, B_n\}$. The trace of the partition $\pi$ on $L$ is:

$$\pi_L = \{B_i \cap L \mid 1 \leq i \leq k \text{ and } B_i \cap L \neq \emptyset\}.$$ 

Trace of partition $\pi = \{B_1, \ldots, B_7\}$ on set $L$
Key Issue for Data Mining:

Defining Dissimilarities and Metrics for Partitions
Shannon’s Entropy

For random variables...
The Shannon entropy is introduced for a random variable distribution

\[ X : \begin{pmatrix} x_1 & \cdots & x_n \\ p_1 & \cdots & p_n \end{pmatrix} \]

is \( \mathcal{H}(X) = - \sum_{i=1}^{n} p_i \log_2 p_i \).
Shannon entropy

... for partitions
A partition $\pi = \{B_1, \ldots, B_m\}$ on a finite, nonempty set $A$ generates naturally a random variable:

$$X_{\pi} : \begin{pmatrix} B_1 & \cdots & B_m \\ \frac{|B_1|}{|S|} & \cdots & \frac{|B_m|}{|S|} \end{pmatrix}$$

We define the Shannon entropy of $\pi$ as the Shannon entropy of $X_{\pi}$. 
Measuring concentration of values

\[ H_1(\pi_4) = 1.9609 \]

\[ H_1(\pi_3) = 2.0464 \]

\[ H_1(\pi_2) = 2.1709 \]

\[ H_1(\pi_1) = 2.3219 \]
Gini’s Index

\[ \mathcal{H}_2(\pi) = 1 - \sum_{i=1}^{n} p_i^2 \]

\[ \mathcal{H}_1(\pi_4) = 0.68 \]

\[ \mathcal{H}_1(\pi_3) = 0.72 \]

\[ \mathcal{H}_1(\pi_2) = 0.79 \]

\[ \mathcal{H}_1(\pi_1) = 0.80 \]
Generalized Entropy of Partitions

Daróczy’s $\beta$-generalized entropy of $\pi = \{B_1, \ldots, B_n\}$:

$$
\mathcal{H}_\beta(\pi) = \frac{1}{1 - 2^{1-\beta}} \left( 1 - \sum_{i=1}^{n} \left( \frac{|B_i|}{|S|} \right)^\beta \right).
$$

For $\beta = 2$ we obtain the Gini index. Also, $\lim_{\beta \to 1} \mathcal{H}_\beta(\pi)$ is Shannon’s entropy

$$
\mathcal{H}(\pi) = - \sum_{i=1}^{n} \frac{|B_i|}{|S|} \log_2 \frac{|B_i|}{|S|}.
$$
Set Purity and Entropy

\( \mathcal{H}(\pi_L) \) measures the impurity of the set \( L \) relative to the partition \( \pi \): the larger the entropy, the more \( L \) is scattered among the blocks of \( \pi \).

If \( \pi, \sigma \in \text{PART}(S) \), the average impurity of the blocks of \( \sigma \) relative to \( \pi \) is the conditional entropy of \( \pi \) relative to \( \sigma \):

\[
\mathcal{H}(\pi|\sigma) = \sum_{j=1}^{m} \frac{|Q_j|}{|S|} \mathcal{H}(\pi_{Q_j}),
\]

where \( \sigma = \{Q_1, \ldots, Q_m\} \).
Generalized Conditional Entropy

For $\pi, \sigma \in \text{PART}(S)$ such that

$$\pi = \{P_1, \ldots, P_k\}$$
$$\sigma = \{Q_1, \ldots, Q_m\}$$

the conditional $\beta$-entropy $\mathcal{H}_\beta(\pi|\sigma)$ is:

$$\mathcal{H}_\beta(\pi|\sigma) = \sum_{j=1}^{m} \left(\frac{|Q_j|}{|S|}\right)^\beta \mathcal{H}_\beta(\pi_{Q_j})$$

$$= \frac{1}{(2^{1-\beta} - 1)|S|^{\beta}} \left(\sum_{i=1}^{k} \sum_{j=1}^{m} |P_i \cap Q_j|^{\beta} - \sum_{j=1}^{m} |Q_j|^{\beta}\right).$$
Metrics on Partition Sets

López de Mántaras:

\[ d(\pi, \sigma) = \mathcal{H}(\pi|\sigma) + \mathcal{H}(\sigma|\pi) \]

Simovici and Jaroszewicz:

\[ d_\beta(\pi, \sigma) = \mathcal{H}_\beta(\pi|\sigma) + \mathcal{H}_\beta(\sigma|\pi) \]
\[ = \frac{1}{(2^{1-\beta} - 1)|S|^\beta} \left( 2 \cdot \sum_{i=1}^{k} \sum_{j=1}^{m} |P_i \cap Q_j|^\beta \right. \\
\left. - \sum_{i=1}^{n} |P_i|^\beta - \sum_{j=1}^{m} |Q_j|^\beta \right) \cdot \]

\[ \lim_{\beta \to 1} d_\beta(\pi, \sigma) = d(\pi, \sigma) \]
A database table $\tau$ is a triple $\tau = (T, H, \rho)$

The header:

$H = A_1 \cdots A_n$

Dom($A_i$): domain of $A_i$

The content of the table: $\rho = \{t_1, \ldots, t_m\}$ where $\rho \subseteq \text{Dom}(A_1) \times \cdots \times \text{Dom}(A_n)$.
Every attribute set $K \subseteq H$ induces a partition $\pi_K$:
Classification in Data Mining

Training Set
Initial Data Set

CLASSIFIER

Test Data

Classification of test data
Goodman-Kruskal Association Index and Metric
The Goodman-Kruskal Coefficient

Let $X$, $Y$ be two discrete random variables. The Goodman-Kruskal coefficient of $X$ and $Y$ is defined by

$$\text{GK}(X, Y)$$

$$= \sum_{i=1}^{l} P(X = a_i) \left( 1 - \max_{1 \leq j \leq k} P(Y = b_j | X = a_i) \right)$$

$$= 1 - \sum_{i=1}^{l} P(X = a_i) \max_{1 \leq j \leq k} P(Y = b_j | X = a_i).$$

Classification rule: an elementary event is classified in the class that has the maximal probability.
GK Classification Rule

- $P(Y = b_j | X = a_i)$: the probability of predicting the value $b_j$ for $Y$ when $X = a_i$

An event that has the component $X = a_i$ is classified in the $Y$-class $b_j$ if $j$ is the number for which $P(Y = b_j | X = a_i)$ has the largest value.

- The probability of misclassification:

$$1 - \max_{1 \leq j \leq k} P(Y = b_j | X = a_i).$$
\( \text{GK}(X, Y) \) is the expected probability that in a randomly chosen case the value of \( Y \) will be incorrectly predicted from \( X \).

\( \lambda_{Y|X} \) is the relative reduction in the probability of prediction error:

\[
\lambda_{Y|X} = 1 - \frac{\text{GK}(X, Y)}{1 - \max_{1 \leq j \leq k} P(Y = b_j)}
\]

\( \lambda_{Y|X} \) is the proportion of the relative error in predicting the value of \( Y \) that can be eliminated by knowledge of the \( X \)-value.
The Goodman-Kruskal Coefficient for Partitions

Consider two partitions

\[ \pi = \{B_1, \ldots, B_l\} \text{ and } \sigma = \{C_1, \ldots, C_k\}. \]

Define the *Goodman-Kruskal coefficient* of these partitions \( \text{GK}(\pi, \sigma) \) as the number:

\[
\text{GK}(\pi, \sigma) = 1 - \sum_{i=1}^{l} \max_{1 \leq j \leq k} \frac{|C_j \cap B_i|}{|S|}.
\]
Partitions and Random Variables

The partitions $\pi, \sigma$ define two random variables

$$X : \left( \begin{array}{ccc}
1 & \cdots & l \\
|B_1| & \cdots & |B_l|
\end{array} \right) \text{ and } Y : \left( \begin{array}{ccc}
1 & \cdots & k \\
|C_1| & \cdots & |C_k|
\end{array} \right)$$

such that conditional probability $P(Y = j | X = i)$ is given by:

$$P(Y = j | X = i) = \frac{P(Y = j \land X = i)}{P(X = i)} = \frac{|C_j \cap B_i|}{|B_i|}.$$
Interpretation of GK

For a fixed $i$, the largest error in predicting $Y$ is:

$$1 - \max_{1 \leq j \leq k} P(Y = j | X = i) = 1 - \max_{1 \leq j \leq k} \frac{|C_j \cap B_i|}{|B_i|}.$$ 

The expected value of the largest error in predicting $Y$ is:

$$\sum_{i=1}^{l} \frac{|B_i|}{|S|} \cdot \left(1 - \max_{1 \leq j \leq k} \frac{|C_j \cap B_i|}{|B_i|}\right)$$

$$= 1 - \sum_{i=1}^{l} \max_{1 \leq j \leq k} \frac{|C_j \cap B_i|}{|S|},$$

which is exactly $\text{GK}(X, Y)$. 

Properties of GK

- We have $GK(\pi, \sigma) = 0$ if and only if $\pi \leq \sigma$.
- The function $GK$ is monotonic in its first argument and dually monotonic in its second:
  - If $\pi, \pi', \sigma$ are three partitions of the set $S$ such that $\pi \leq \pi'$, then $GK(\pi, \sigma) \leq GK(\pi', \sigma)$.
  - If $\pi, \sigma', \sigma$ are three partitions of the set $S$ such that $\sigma \leq \sigma'$, then $GK(\pi, \sigma) \geq GK(\pi, \sigma')$.
- $GK$ satisfies a triangular inequality:
  $GK(\pi, \sigma) \leq GK(\pi, \tau) + GK(\tau, \sigma)$. 
Metric Associated to GK

The Goodman-Kruskal coefficient allows us to define a metric on $\text{PART}(S)$.
Let $d_{GK} : \text{PART}(S) \times \text{PART}(S) \longrightarrow \mathbb{R}$ be

$$d_{GK}(\pi, \sigma) = \text{GK}(\pi, \sigma) + \text{GK}(\sigma, \pi).$$

for $\pi, \sigma \in \text{PART}(S)$.

The function $d_{GK}$ is a metric on the set $\text{PART}(S)$.
Goodman-Kruskal Coefficient for Attribute Sets

Let $K, L$ be two sets of attributes of a table. Define $\text{GK}(K, L) = \text{GK}(\pi_K, \pi_L)$: the expected error that occurs when we try to predict the value of $t[L]$ from the value of $t[K]$.

- If $K_1 \subseteq K_2$, then $\pi_{K_2} \leq \pi_{K_1}$, so $\text{GK}(K_2, L) \leq \text{GK}(K_1, L)$.
- If $L_1 \subseteq L_2$, then $\text{GK}(K, L_2) \leq \text{GK}(K, L_1)$. 
Goodman-Kruskal Metric on Attribute Sets

Define $d_{GK}(K, L) = d_{GK}(\pi_K, \pi_L)$ for any two sets of attributes $K, L$.

The new metric can be used for:

- constructing classifiers;
- discretization of continuous attributes;
- attribute clustering, feature selection and data compression.
\( \epsilon \)-predictors

An \( \epsilon \)-predictor for a set of attributes \( Y \) is a set of attributes \( K \) such that \( G_K(K, Y) \leq \epsilon \).

- If \( K \) is an \( \epsilon \)-predictor for \( Y \), then any superset \( K' \) of \( K \) is also a \( \epsilon \)-predictor for \( Y \).
- An \( \epsilon \)-predictor such that no of its proper subsets is an \( \epsilon \)-predictor is called \emph{minimal}.
An Apriori-like Algorithm for $\epsilon$-predictors

**Input:** A set of attributes $H$, a target attribute $Y$, $Y \notin H$ and an error level.

**Output:** Set $P$ of all minimal $\epsilon$-predictors from $H$.

1. $\text{Cand} = \{ \{ A \} : A \in H \}$;
2. $P = \emptyset$;
3. $P = P \cup \{ K \in \text{Cand} : \text{GK}(K, Y) \leq \epsilon \}$;
4. $\text{Cand} = \text{Cand} \setminus P$;
5. $\text{Cand} = \{ L \subseteq H : \text{for all } K \subset L, |K| = |L| - 1 \text{ we have } K \in \text{Cand} \}$;
6. goto (3);
• If a set is a nonminimal predictor, so are all of its supersets, which can thus be skipped.
• Initialize candidate set of predictors $\text{Cand}$ to include one-set attributes.
• The set of minimal predictors $\mathcal{P}$ is constructed starting from $\text{Cand}$.
• Initialize $P$ to include all singleton predictors whose error is below the threshold $\epsilon$. Remove those from $C$ and the search for minimal two-attribute predictors makes use of the remaining candidate attributes, etc.

• The stopping condition could be exceeding the maximum predictor size or finding a predictor with desired prediction error.
Experimental Results – KHAN


Differential diagnosis of four small round blue cell tumors of childhood (SRBCTs):

**NB:** neuroblastoma

**RMS:** rhabdomyosarcoma

**BL:** Burkitt lymphoma

**EWS:** Ewing family of sarcomas
Previous work:

- single layer neural networks (Khan)
- logistic regression model (Weber)
- SVMs (Mukerjee)
- combined classifiers (Yeo)
Khan Data

- 2308 genes were measured using cDNA microarrays
- Training Data: 63 cases (12 NB, 20 RMS, 8 BL, and 23 EWS)
- Test Data: 25 cases (6 EWS, 5 RMS, 6 NB, 3 BL, and 5 non-SRBCTs)
- The test cases include 5 cases which do not belong to any of the predicted SRBCT types. Such cases are not present in the training set.
Preprocessing

Replace each class attribute with 4 binary attributes, one for each cancer type.

<table>
<thead>
<tr>
<th>original attribute</th>
<th>computed attributes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cancer type</td>
<td>NB</td>
</tr>
<tr>
<td>NB</td>
<td>1</td>
</tr>
<tr>
<td>EWS</td>
<td>0</td>
</tr>
<tr>
<td>RMS</td>
<td>0</td>
</tr>
<tr>
<td>other</td>
<td>0</td>
</tr>
</tbody>
</table>
• A separate predictor is built for each binary attribute to allow for handling of cases of type ‘other’ present in the test set, but absent in the training set.

• We expect that for ‘other’ cancer type all of the predictors will give the value of 0 thus indicated that none of the 4 cancer types is present.
• Predictors may contradict each other (infrequently, because low error rate of individual classifiers).

• If presence of more than one cancer type is predicted consider it misclassified.

• Small predictors decrease the risk of overfitting (small number of training cases!)
Discretization

- Every gene expression level $X$ attribute is discretized into two intervals: $X \leq T$ and $X > T$.
- $T$ is chosen such that the Shannon entropy $H(Y|X')$ of the target $Y$ conditional on the discretized attribute $X'$ is minimal.
- A separate discretization (using Fayyad-Irani) has been performed with respect to each cancer type.
Limitations on the Computation

- We find all predictors with 1 or 2 attributes, allowing up to one misclassified instance on the training set.
- The stopping rule: reaching the maximum prescribed size of the predictor, or obtaining an error rate less than to \( \frac{1}{t} \), where \( t \) is the size of the training set.
- All but 30 most predictive attributes are discarded.
- For each cancer type the first predictor with minimal training error is manually picked at random (without looking at its test set performance to avoid bias in the choice).
<table>
<thead>
<tr>
<th>Cancer type</th>
<th>selected predictor</th>
<th>image ids</th>
<th>mtr</th>
<th>mte</th>
<th>1GP</th>
<th>2GP</th>
</tr>
</thead>
<tbody>
<tr>
<td>BL</td>
<td>WAS ≤ 0.69 ⇒ BL</td>
<td>236282</td>
<td>0</td>
<td>1</td>
<td>15</td>
<td>5</td>
</tr>
<tr>
<td>EWS</td>
<td>FCGRT ≤ 1.59 ⇒ EWS</td>
<td>770394</td>
<td>1</td>
<td>3</td>
<td>2</td>
<td>10</td>
</tr>
<tr>
<td>NB</td>
<td>MAP1B &gt; 2.17 or RCV1 &gt; 1.98 ⇒ NB</td>
<td>629896 - 383188</td>
<td>0</td>
<td>0</td>
<td>2</td>
<td>28</td>
</tr>
<tr>
<td>RMS</td>
<td>TNNT2 &gt; 0.55 or SGCA &gt; 0.44 ⇒ RMS</td>
<td>298062 - 796258</td>
<td>0</td>
<td>2</td>
<td>0</td>
<td>25</td>
</tr>
</tbody>
</table>

Legend:

- mte: misclassified cases in test set
- mtr: misclassified cases in training set
- 1GP: number of one-gene predictors
- 2GP: number of two-gene predictors
• A fairly large number (12–30) of very simple predictors have been found for each cancer type.
• Each of those predictors has very good classification rate on the training set: up to one misclassified case is allowed.
• The results show that there are many genes based on which a diagnosis can be made for each cancer type.
• All genes except for the one that predicts BL were reported among the 96 selected in Khan.
Bonferroni Correction

The probability that a single gene expression predicts BL perfectly on training set when there is no correlation between the gene and the tumor type:

$$2 \cdot \frac{55! \cdot 8!}{63!} = 5.16 \cdot 10^{-10}$$

much less than $0.05/2308 = 2.16 \cdot 10^{-5}$, the 5% significance level after Bonferroni correction. This shows that selected gene is with very high probability related to BL.
• If a classifier for only one type of tumor gave a positive prediction, then the instance was classified as this type of tumor.

• If none of them gave positive prediction we declared the case as ‘other tumor type’.

• If more than one classifier was active the case was considered a prediction error.

• The combined classifier used a total of 6 genes and classified correctly 19 out of 25 test cases.

• Out of the 6 misclassified cases, 2 gave classifications when the real outcome was ‘other’, 3 SRBCT cases were undetected, and there was 1 conflict.
Experimental Results - GOLUB

- Training data: 38 cases (27 acute lymphoblastic leukemia and 11 acute myelocytic leukemia)
- Test data: 34 cases (20 ALL and 14 AML);
- Data involves 6817 genes.
- We discretized the gene expression levels using Fayyad-Irani
- 20 genes were retained for which the Goodman-Kruskal coefficient was below 0.04.
• Five single-genes predictors and 66 two-gene predictors were identified.

• We identified two two-genes predictors (MGST1, APLP2 and CD33, CystatinA) for which the errors on the test set are 0 and 0.0294118, respectively.

• CD33 was among the 50 genes selected by Golub et al.
Distribution of the errors on the test set for the remaining set of minimal two-genes predictors:

<table>
<thead>
<tr>
<th>Error Interval</th>
<th>Number of 2-attribute predictors</th>
</tr>
</thead>
<tbody>
<tr>
<td>[0.0, 0.05]</td>
<td>2</td>
</tr>
<tr>
<td>(0.05, 0.10]</td>
<td>9</td>
</tr>
<tr>
<td>(0.10, 0.15]</td>
<td>10</td>
</tr>
<tr>
<td>(0.15, 0.20]</td>
<td>7</td>
</tr>
<tr>
<td>(0.20, 0.25]</td>
<td>13</td>
</tr>
<tr>
<td>(0.25, 0.30]</td>
<td>14</td>
</tr>
<tr>
<td>(0.30, 0.35]</td>
<td>3</td>
</tr>
<tr>
<td>(0.35, 0.40]</td>
<td>4</td>
</tr>
<tr>
<td>(0.40, 0.45]</td>
<td>3</td>
</tr>
</tbody>
</table>
Voting Mechanism

- We retained 19 one-attribute predictors whose prediction error on the training set did not exceed 5.3% (that is, two errors out of the 38 training cases).
- A vote was taken, and the instance was classified according to the majority vote.
- We obtained 3 errors on the test set of 34 cases. Namely, the errors occurred on the 57th, 60th and 66th cases of the original Golub test set ("unclassifiable" in the original study (Golub)).
Conclusions - 1A

- The Goodman-Kruskal dissimilarity $GK$ is a simple, but powerful measure of predictive power that can be used to produce robust classifiers.
- The small number of training cases makes reliable construction of more complex models like Bayesian networks or C4.5 trees very hard or even impossible.
- Naive Bayesian classifiers suffer from independence assumptions which may not be satisfied in the microarray setting where most genes are correlated with each other.
- The Bonferroni correction, though conservative, yields valid results.
Conclusions - 1B

The fact that GK generates a metric suggests one could use this metric to cluster attributes such that those that belong to a cluster have similar predictive power and, thus, are interchangeable in classifiers. The clustering structure could be used in forming the voting committees, and in simplifying and increasing the robustness of predictive algorithms.
A New Metric Discretization Algorithm
From numerical to nominal

Previous work on discretization:

- fixed $k$-interval discretization (J. Dougherty, R. Kohavi, M. Sahami, 1995)
- fuzzy discretization (Kononenko 1992-1993)
- Shannon-entropy discretization (Fayyad and Irani, 1993)
- highly dependent attributes (M. Robnik and I. Kononenko, 1995)
Basic Results

- a generalization of Fayyad-Irani discretization technique
- a geometric criterion for halting the discretization process
- better results in building
  - naive Bayes classifiers
  - decision trees
Discretization of a numeric attribute $B$

Set of cutpoints: $S = \{t_1, \ldots, t_\ell\}$ in $\text{aDom}(B)$, where $t_1 < t_2 < \cdots < t_\ell$.

Discretization partition of $\text{aDom}(B)$:

$$\pi^S = \{Q_0, \ldots, Q_\ell\}$$
Boundary Points

\( \pi_A \) the partition of the set of tuples of a table determined by the values of an attribute \( A \):
\[
\text{select } * \text{ from } T \text{ group by } A
\]
t\(_1, \ldots, t_n\): the list of tuples sorted on the values of an attribute \( B \).

\( \pi_{B,A} \) is the partition of \( \text{aDom}(B) \) that consists of the longest runs of \textit{consecutive} \( B \)-components of the tuples in this list that belong to the \textit{same block} \( K \) of the partition \( \pi_A \).
The \textit{boundary points} of the partition \( \pi_{B,A} \) are the least and the largest elements of each of the blocks of the partition \( \pi_{B,A} \).
We have \( \pi_{B,A}^* \leq \pi_A \) for any attribute \( B \).
Main Results -I

Theorem: Let $T$ be a table where the class of the tuples is determined by the attribute $A$ and let $\beta \in (1, 2]$.

If $S$ is a set of cutpoints such that the conditional entropy $\mathcal{H}_\beta(\pi_A | \pi_*^S)$ is minimal among the set of cutpoints with the same number of elements, then $S$ consists of boundary points of the partition $\pi_{B,A}$ of $\text{aDom}(B)$. 
Main Results -II

Theorem: Let $\beta \in (1, 2]$. If $S$ is a set of cutpoints such that the distance $d_\beta(\pi_A, \pi^S_*)$ is minimal among the set of cutpoints with the same number of elements, then $S$ consists of boundary points of the partition $\pi_{B,A}$ of $a\text{Dom}(B)$. 
To discretize $a\text{Dom}(B)$ we seek a set of cutpoints such that

$$d_\beta(\pi_A, \pi^S) = \mathcal{H}_\beta(\pi_A|\pi^S) + \mathcal{H}_\beta(\pi^S|\pi_A)$$

is minimal.

Seek a set of cutpoints $S$ such that the partition $\pi^S$ induced on the set of rows is as close as possible to the target partition $\pi_A$. 
Discretization Algorithm

**Input:** A table $T$, a class attribute $A$, and a real-valued attribute $B$.

**Output:** A discretized attribute $B$.

$BP$ is the set of boundary points of partition $\pi_{B,A*}$.
Method:

sort $T$ on $B$;
compute BP;
$S = \emptyset$; $d = \infty$;
while BP $\neq \emptyset$ do

let $t = \arg \min_{t \in BP} d_\beta(\pi_A, \pi^S \cup \{t\})$;
if $d \geq d_\beta(\pi_A, \pi^S \cup \{t\})$ then
begin

$S = S \cup \{t\}$; BP = BP $- \{t\}$;
$d = d_\beta(\pi_A, \pi^S)$

end

else exit while loop;
end while

for $\pi^S = \{Q_0, \ldots, Q_\ell\}$ replace every value in $Q_i$ by $i$ for $0 \leq i \leq \ell$. 

Metric Methods in Mining – p.66/79
$d_\beta(\pi_A, \pi_*^S)$ as a function of $|S|$

78% of the total time is spent on decreasing the distance by the last 1%
\[ d_\beta(\pi_A, \pi^S) = \mathcal{H}_\beta(\pi_A | \pi^S) + \mathcal{H}_\beta(\pi^S | \pi_A) \]

If \( S \subseteq S' \) then \( \pi^S \geq \pi^{S'} \) and

\[
\begin{align*}
\mathcal{H}_\beta(\pi_A | \pi^S) &\geq \mathcal{H}_\beta(\pi_A | \pi^{S'}) \\
\mathcal{H}_\beta(\pi^S | \pi_A) &\leq \mathcal{H}_\beta(\pi^{S'} | \pi_A).
\end{align*}
\]

Process starts with \( S = \emptyset \), so \( \pi^S = \omega \).

Practical halting criterion:

\[ |d - d_\beta(\pi_A, \pi_*^{S \cup \{t\}})| > 0.01d. \]
Experimental Results

- Accuracy measured in stratified 10-fold cross-validation
- UCI datasets with $\beta \in \{1.5, 1.8, 1.9, 2\}$
### Experimental Results - I

<table>
<thead>
<tr>
<th>Method</th>
<th>Size</th>
<th>Leaves</th>
<th>Accuracy</th>
</tr>
</thead>
<tbody>
<tr>
<td>standard</td>
<td>51</td>
<td>30</td>
<td>79.20</td>
</tr>
<tr>
<td>(\beta = 1.5)</td>
<td>20</td>
<td>14</td>
<td>77.36</td>
</tr>
<tr>
<td>(\beta = 1.8)</td>
<td>28</td>
<td>18</td>
<td>77.36</td>
</tr>
<tr>
<td>(\beta = 1.9)</td>
<td>35</td>
<td>22</td>
<td>76.01</td>
</tr>
<tr>
<td>(\beta = 2.0)</td>
<td>54</td>
<td>32</td>
<td>76.01</td>
</tr>
</tbody>
</table>

#### heart-c:

<table>
<thead>
<tr>
<th>Method</th>
<th>Size</th>
<th>Leaves</th>
<th>Accuracy</th>
</tr>
</thead>
<tbody>
<tr>
<td>standard</td>
<td>57</td>
<td>30</td>
<td>57.28</td>
</tr>
<tr>
<td>(\beta = 1.5)</td>
<td>32</td>
<td>24</td>
<td>71.02</td>
</tr>
<tr>
<td>(\beta = 1.8)</td>
<td>56</td>
<td>50</td>
<td>77.10</td>
</tr>
<tr>
<td>(\beta = 1.9)</td>
<td>64</td>
<td>58</td>
<td>67.57</td>
</tr>
<tr>
<td>(\beta = 2.0)</td>
<td>92</td>
<td>82</td>
<td>66.35</td>
</tr>
</tbody>
</table>

#### glass:
# Experimental Results - II

## ionosphere:

<table>
<thead>
<tr>
<th>Standard</th>
<th>( \beta = 1.5 )</th>
<th>( \beta = 1.8 )</th>
<th>( \beta = 1.9 )</th>
<th>( \beta = 2.0 )</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>15</td>
<td>19</td>
<td>15</td>
<td>15</td>
</tr>
<tr>
<td></td>
<td>8</td>
<td>12</td>
<td>10</td>
<td>10</td>
</tr>
<tr>
<td></td>
<td>( 90.88 )</td>
<td>( 88.31 )</td>
<td>( 90.02 )</td>
<td>( 90.02 )</td>
</tr>
</tbody>
</table>

## iris:

<table>
<thead>
<tr>
<th>Standard</th>
<th>( \beta = 1.5 )</th>
<th>( \beta = 1.8 )</th>
<th>( \beta = 1.9 )</th>
<th>( \beta = 2.0 )</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>7</td>
<td>7</td>
<td>7</td>
<td>7</td>
</tr>
<tr>
<td></td>
<td>5</td>
<td>5</td>
<td>5</td>
<td>5</td>
</tr>
<tr>
<td></td>
<td>( 95.33 )</td>
<td>( 96 )</td>
<td>( 96 )</td>
<td>( 96 )</td>
</tr>
</tbody>
</table>
### Experimental Results - III

<table>
<thead>
<tr>
<th>diabetes:</th>
<th>standard</th>
<th>43</th>
<th>22</th>
<th>74.08</th>
</tr>
</thead>
<tbody>
<tr>
<td>$\beta = 1.8$</td>
<td></td>
<td>5</td>
<td>3</td>
<td>75.78</td>
</tr>
<tr>
<td>$\beta = 1.9$</td>
<td></td>
<td>7</td>
<td>4</td>
<td>75.39</td>
</tr>
<tr>
<td>$\beta = 2.0$</td>
<td></td>
<td>14</td>
<td>10</td>
<td>76.30</td>
</tr>
</tbody>
</table>
Heart-c

![Bar chart showing tree size, number of leaves, and accuracy for different values of \(\beta\).]
Glass

Tree size

Number of leaves

Accuracy

- **standard**
- $\beta = 1.5$
- $\beta = 1.8$
- $\beta = 1.9$
- $\beta = 2.0$
## Naive Bayes Classifiers

### Error Rate

<table>
<thead>
<tr>
<th>Discretization Method</th>
<th>Diabetes</th>
<th>Glass</th>
<th>Ionosphere</th>
<th>Iris</th>
</tr>
</thead>
<tbody>
<tr>
<td>$\beta = 1.5$</td>
<td>34.9</td>
<td>25.2</td>
<td>4.8</td>
<td>2.7</td>
</tr>
<tr>
<td>$\beta = 1.8$</td>
<td>24.2</td>
<td>22.4</td>
<td>8.3</td>
<td>4</td>
</tr>
<tr>
<td>$\beta = 1.9$</td>
<td>24.9</td>
<td>23.4</td>
<td>8.5</td>
<td>4</td>
</tr>
<tr>
<td>$\beta = 2.0$</td>
<td>25.4</td>
<td>24.3</td>
<td>9.1</td>
<td>4.7</td>
</tr>
<tr>
<td>weighted prop</td>
<td>25.5</td>
<td>38.4</td>
<td>10.3</td>
<td>6.9</td>
</tr>
<tr>
<td>prop.</td>
<td>26.3</td>
<td>33.6</td>
<td>10.4</td>
<td>7.5</td>
</tr>
</tbody>
</table>
Conclusions

An appropriate choice of $\beta$ that defines the metric used in discretization, yields better classifiers (decision trees and naive Bayes)

Open issues:

- identifying simple parameters of data sets that inform the best choice of $\beta$;
- metric discretization for data with missing values.
Future Directions of Work
The metric space of attributes can be used to cluster attributes.

- Similar attributes are grouped in clusters, that may have biological significance.
- Retaining one attribute per cluster (e.g., the centroid) allows for data compression and for simplification of decision techniques.
Our metric approach to discretization can be used to provide “quasi-optimal” discretization to microarray data. It would be interesting to compare this approach with a previous fuzzy discretization approach (Ohno-Machado).

Microarray data have high dimensionality (in the thousands or tens of thousands). By focusing on partitions induced by genes rather than on individual phenotypes we hope to avoid the difficulties related to high dimensionality.