## University of Amsterdam

## Master's thesis

# Clustering by file compression

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# Abstract

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# Contents

Abstract														
Contents														
1	Introduction													
	1.1	Simila	arity of data	. 1										
2	Normalized compression distance													
	2.1 Foundations in Kolmogorov complexity													
		2.1.1	Information distance	. 3										
	2.2	Appro	eximating Kolmogorov complexity with a real world compressor	. 3										
		2.2.1	A real world compressor does not exploit all regularity	. 4										
3	Hierarchical clustering of data													
	3.1 Evolution of placental mammals													
		3.1.1	Distance matrix	. 6										
		3.1.2	Clustering method	. 7										
		3.1.3	Dendrogram	. 7										
A	Apj	pendix	Title Here	9										
Bi	bliog	graphy		10										

## Chapter 1

## Introduction

## 1.1 Similarity of data

How do we know that Dutch is more similar to German than it is to French? How do we know that Bob Dylan's music is closer to The Beatles' than it is to Bach's?

Does a computer know?

This thesis concerns a method expressing similarity of data that is feature free: it does not use domain knowledge about the data (for example, word origins or grammar rules in the case of languages.) The method is based on file compression and is rooted in Kolmogorov complexity.

The idea is easy to grasp. If a compressor compresses the concatenation of two files better than it compresses the files separately, it must have found some regularities that appear in both files. This compression gain is used to define a similarity metric that aims to capture the similarity of every dominant feature of the data.

## Chapter 2

# Normalized compression distance

### 2.1 Foundations in Kolmogorov complexity

In this chapter we will make explicit the idea of similarity based on file compression. But first, we explain the notion of Kolmogorov complexity. For a complete reference, see [1].

The Kolmogorov complexity of a string x, written K(x), is the length of the shortest program that outputs x.

Intuitively, 111...111, a string of a million ones, is not very complex. It does not contain much information. Indeed, the Kolmogorov complexity of this string is low. A 27-byte Ruby program produces it:

```
1000000.times { print "1" }
```

I do not claim the Kolmogorov complexity of a string of a million ones is 27 bytes. The true Kolmogorov complexity of a string x cannot be computed in the Turing sense. There is no program that, given x, outputs the (length of) the shortest program that produces x.

The string 011...010, produced by flipping a fair coin a million times, has, with very high probability, a Kolmogorov complexity close to its own length (by counting the number of different bit strings of each length, you can show that the chance to compress a random string by more than c bits is at most  $2^{-c}$ ).

So, printing the literal description may be the best we can do:

print "011...010"

The Kolmogorov complexity of a string x, given a string y, denoted by K(x|y), is the length of the shortest program that outputs x, given y as input.

#### 2.1.1 Information distance

The Kolmogorov complexity is a measure of information content in an individual object. In the same way, in [2] the information distance E(x,y) between two strings x and y is defined as the length of the shortest program that converts x to y and y to x. It is shown that

$$E(x,y) = \max\{K(x|y), K(y|x)\}\$$

E(x, y) is an absolute distance. But similarity is better expressed relatively. To illustrate: if two binary strings of length 100 have a Hamming distance of 50 (i.e. they have different bits in 50 positions), they are not very alike. If, on the other hand, two strings of length  $10^6$  have a Hamming distance of 50, they are very much alike.

In [3], the normalized information distance is defined as

$$NID(x,y) = \frac{\max\{K(x|y), K(y|x)\}}{\max\{K(x), K(y)\}}$$
(2.1)

and it is shown that NID(x, y) minorizes every distance d(x, y) up to a negligible additive term, where d(x, y) belongs to a wide class of normalized distances that includes everything remotely interesting.

This means that if two strings are similar according to some distance (be it Hamming distance, overlap distance, or any other), they are also similar according to the normalized information distance. This is why NID(x, y) is also called *the* similarity distance.

# 2.2 Approximating Kolmogorov complexity with a real world compressor

The remarkable properties of the normalized information distance come at the price of incomputability. But, the Kolmogorov complexity can be approximated by real world (lossless) compression programs like zlib or liblzma. The normalized compression distance [4] is defined as

$$NCD = \frac{C(xy) - \min\{C(x), C(y)\}}{\max\{C(x), C(y)\}}$$
(2.2)

where xy is the concatenation of x and y, and C(x) is the length of x, after being compressed by compressor C.

The NCD is central to this work. It is the real world approximation of the normalized information distance (2.1).

### 2.2.1 A real world compressor does not exploit all regularity

Since K is incomputable, we have no idea how far off the length given by C is. Consider 31415..., the string consisting of the first  $10^9$  digits of  $\pi$ . The Kolmogorov complexity of this string is low: a simple program, perhaps exploiting a converging series formula, will produce it. Any real world compressor C, however, fails to compress this string by even a few bits<sup>1</sup>.

In the following chapters (and extensively in [4]) it is demonstrated that the NCD is adequate for many applications.

<sup>&</sup>lt;sup>1</sup>A textual representation "31415..." can actually be compressed significantly by almost every compressor, but this is an encoding issue. The file consists only of the bytes 0 through 9, which can be more efficiently encoded using, for example, a system where each decimal number is assigned a four bit code (which is still naive.) For clarity, I do not concern myself with encoding in this chapter. Every finite alphabet can be recoded in binary, so it is customary to only think about binary strings in the literature. Conceptually, nothing changes.

## Chapter 3

# Hierarchical clustering of data

In this chapter, we cluster vastly different types of data using the normalized compression distance (NCD) (2.2). To explain the method, we first cluster mitochondrial gene sequences of mammals.

## 3.1 Evolution of placental mammals

Reconstructing an evolutionary tree has intuitive appeal. It should lend itself well to hierarchical clustering, since species emerge from common ancestors. And, within biology, there is agreement on what the true phylogeny tree is: the brown bear and polar bear are closely related, etc. Only higher up on the tree there is ongoing debate. Do the primates first join with the rodents, or are they more closely connected to the ferungulates?

All materials were taken from the GenBank database [5].

In [6], the authors estimate the likelihood of phylogeny trees based on 12 mitochondrial proteins of 20 placental mammals: rat (Rattus norvegicus), house mouse (Mus musculus), grey seal (Halichoerus grypus), harbor seal (Phoca vitulina), cat (Felis catus), white rhino (Ceratotherium simum), horse (Equus caballus), finback whale (Balaenoptera physalus), blue whale (Balaenoptera musculus), cow (Bos taurus), gibbon (Hylobates lar), gorilla (Gorilla gorilla), human (Homo sapiens), chimpanzee (Pan troglodytes), pygmy chimpanzee (Pan paniscus), orangutan (Pongo pygmaeus), Sumatran orangutan (Pongo abelii), using opossum (Didelphis virginiana), wallaroo (Macropus robustus), and the platypus (Ornithorhynchus anatinus).

In [4], 4 more mammals were added: Australian echidna (*Tachyglossus aculeatus*), brown bear (*Ursus arctos*), polar bear (*Ursus maritimus*), and the common carp (*Cyprinus*)

carpio). The common carp is not a mammal and is used as an outgroup. It should join the phylogeny tree at the very top.

The authors of [4] cluster the complete mitochondrial genome sequences with the normalized compression distance. They use a clustering algorithm described by them in [7]. The algorithm tries to optimize a global criterion, namely the (normalized) summed weights of all consistent quartet topologies (layouts of groups of four items). In a binary tree, only one of three possible pairings of four items (ab|cd, ac|bd, ad|bc) is consistent, in the sense that you can connect the two pairs without crossing paths. The sum of the distances (in this case, NCDs) between the items in the consistent pairs is the contribution of quartet abcd to the tree score.

This method works especially well if the items you're trying to cluster result from an evolutionary process, since then (without corruption of data) there should exist an evolutionary tree that embeds all the most likely quartet topologies, and, given enough time, the heuristic presented in [7] finds the true tree.

Now, we will cluster the same 24 genome sequences with a much simpler algorithm (average linkage) and compare results.

### 3.1.1 Distance matrix

The distance matrix contains the distances between all pairs of items. Entry i, j is the distance between item i and item j, i.e. NCD(i, j). The distance matrix for the 24 animals is displayed in table 3.1.

blueWhale	0.01	0.72	0.86	0.67	0.78	0.60	0.83	0.24	0.81	0.80	0.67	0.66	0.62	0.78	0.77	0.82	0.81	0.77	0.83	0.70	0.79	0.78	0.83	0.63
brownBear	0.71	0.01	0.89	0.63	0.84	0.72	0.85	0.69	0.84	0.82	0.56	0.56	0.69	0.81	0.80	0.84	0.84	0.82	0.84	0.11	0.82	0.83	0.84	0.64
carp	0.87	0.88	0.01	0.88	0.89	0.87	0.89	0.89	0.89	0.89	0.89	0.88	0.87	0.88	0.86	0.88	0.89	0.89	0.90	0.88	0.88	0.89	0.88	0.88
cat	0.67	0.59	0.87	0.01	0.79	0.68	0.84	0.69	0.79	0.79	0.59	0.57	0.61	0.81	0.77	0.80	0.80	0.80	0.82	0.60	0.75	0.81	0.78	0.59
chimpanzee	0.78	0.83	0.90	0.81	0.01	0.77	0.87	0.82	0.49	0.34	0.80	0.79	0.79	0.29	0.84	0.88	0.47	0.17	0.89	0.83	0.84	0.47	0.86	0.80
cow	0.61	0.73	0.87	0.68	0.78	0.01	0.84	0.61	0.79	0.78	0.66	0.66	0.62	0.81	0.77	0.82	0.81	0.77	0.84	0.71	0.76	0.81	0.81	0.62
echidna	0.84	0.87	0.89	0.85	0.85	0.83	0.01	0.86	0.85	0.86	0.87	0.87	0.84	0.88	0.81	0.81	0.87	0.85	0.55	0.87	0.84	0.86	0.84	0.83
finWhale	0.25	0.72	0.89	0.71	0.81	0.62	0.85	0.01	0.81	0.82	0.70	0.69	0.64	0.81	0.80	0.84	0.80	0.78	0.85	0.72	0.81	0.81	0.85	0.64
gibbon	0.83	0.85	0.90	0.81	0.51	0.81	0.88	0.85	0.01	0.51	0.82	0.83	0.81	0.50	0.85	0.89	0.53	0.50	0.90	0.85	0.85	0.54	0.87	0.79
gorilla	0.79	0.81	0.90	0.82	0.35	0.79	0.88	0.83	0.50	0.01	0.79	0.82	0.83	0.35	0.83	0.89	0.46	0.35	0.87	0.82	0.82	0.47	0.88	0.80
graySeal	0.69	0.56	0.88	0.60	0.79	0.66	0.84	0.69	0.79	0.80	0.01	0.16	0.64	0.80	0.77	0.82	0.80	0.78	0.85	0.55	0.78	0.80	0.80	0.61
harborSeal	0.67	0.54	0.88	0.58	0.78	0.64	0.85	0.69	0.80	0.80	0.16	0.01	0.61	0.79	0.76	0.83	0.77	0.77	0.84	0.54	0.77	0.77	0.80	0.60
horse	0.63	0.65	0.88	0.63	0.76	0.62	0.85	0.64	0.77	0.79	0.62	0.60	0.01	0.79	0.78	0.81	0.78	0.78	0.83	0.66	0.76	0.79	0.80	0.49
human	0.78	0.83	0.88	0.83	0.30	0.79	0.87	0.80	0.50	0.35	0.82	0.82	0.78	0.01	0.82	0.87	0.46	0.30	0.88	0.83	0.82	0.46	0.86	0.78
mouse	0.77	0.79	0.86	0.76	0.81	0.76	0.81	0.80	0.81	0.82	0.77	0.76	0.75	0.83	0.01	0.78	0.83	0.82	0.83	0.78	0.54	0.83	0.77	0.74
opossum	0.82	0.81	0.87	0.80	0.87	0.81	0.84	0.82	0.87	0.88	0.82	0.82	0.81	0.87	0.80	0.01	0.88	0.87	0.84	0.82	0.82	0.88	0.68	0.83
orangutan	0.80	0.83	0.90	0.82	0.46	0.82	0.88	0.80	0.52	0.46	0.82	0.79	0.83	0.45	0.84	0.88	0.01	0.47	0.89	0.83	0.83	0.25	0.86	0.82
pigmyChimpanzee	0.77	0.81	0.90	0.81	0.17	0.77	0.88	0.81	0.49	0.34	0.82	0.79	0.79	0.29	0.83	0.87	0.47	0.01	0.89	0.81	0.83	0.47	0.86	0.78
platypus	0.83	0.85	0.89	0.86	0.89	0.82	0.56	0.84	0.89	0.87	0.84	0.84	0.87	0.87	0.83	0.82	0.88	0.88	0.01	0.85	0.84	0.89	0.84	0.85
polarBear	0.70	0.10	0.89	0.62	0.81	0.70	0.86	0.71	0.84	0.81	0.56	0.56	0.66	0.82	0.79	0.83	0.82	0.81	0.85	0.01	0.78	0.82	0.82	0.65
rat	0.78	0.82	0.89	0.75	0.80	0.76	0.83	0.78	0.81	0.82	0.79	0.77	0.74	0.82	0.55	0.82	0.81	0.83	0.86	0.82	0.01	0.81	0.84	0.75
sumatranOrangutan	0.78	0.82	0.89	0.80	0.47	0.80	0.86	0.79	0.52	0.47	0.80	0.80	0.77	0.44	0.84	0.87	0.24	0.47	0.88	0.82	0.82	0.01	0.87	0.78
wallaroo	0.81	0.82	0.87	0.80	0.85	0.81	0.85	0.83	0.86	0.86	0.80	0.79	0.81	0.85	0.81	0.68	0.86	0.85	0.83	0.83	0.82	0.87	0.01	0.80
whiteRhinoceros	0.64	0.66	0.89	0.65	0.78	0.64	0.86	0.65	0.78	0.79	0.61	0.61	0.51	0.79	0.78	0.84	0.82	0.79	0.86	0.68	0.77	0.81	0.82	0.01

TABLE 3.1: Distance matrix of genome sequences of 24 animals. To obtain the NCD, I used ruby-xz, the Ruby binding to compressor liblzma, with settings compression\_level = 9, extreme = true, check = :none.

### 3.1.2 Clustering method

In hierarchical clustering, the goal is to build a hierarchy of clusters from a distance matrix. The hierarchy can be displayed as a binary tree. One of the simplest ways to do this is to put every object in its own cluster, and then merge greedily based on a linkage criterion, until all items have been merged into a single cluster.

In single linkage, at each step the two clusters are merged with the smallest pairwise distance. This is also called *nearest neighbor* clustering.

In complete linkage, or *farthest neighbor* clustering, at each step the two clusters with the largest pairwise distance are merged.

Average linkage is a compromise between single and complete linkage. The distance between two clusters is defined as the average between the pairwise distances, i.e.

$$d(A,B) = \frac{1}{|A||B|} \sum_{a \in A} \sum_{b \in B} d(a,b)$$

### 3.1.3 Dendrogram

Figure 3.1 shows the phylogeny tree obtained by average link clustering.

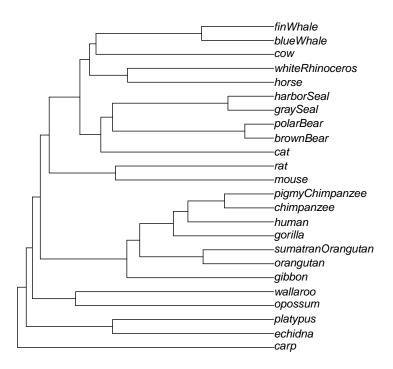


FIGURE 3.1: Result of average linkage clustering of the distance matrix shown in table 3.1. The height at which clusters join is proportional to the distance between them.

# Appendix A

# Appendix Title Here

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