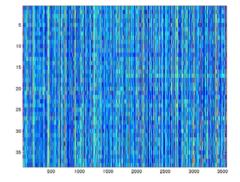
Machine Learning & Artificial Intelligence for Data Scientists: Feature selection and projection

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A problem - too many features

- Aim: To build a classifier that can diagnose leukaemia using Gene expression data.
- ▶ Data: 27 healthy samples,11 leukaemia samples (N = 38). Each sample is the expression (activity) level for 3751 genes. (Also have an independent test set)



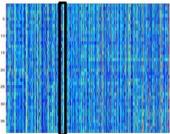
- ▶ In general, the number of parameters will increase with the number of **features** -D = 3751.
 - ▶ e.g. Logistic regression w would have length 3751!
- ► Fitting lots of parameters is hard imagine Metropolis-Hastings in 3751 dimensions rather than 2!

Features

- For visualisation, most examples we've seen have had only 2 features $\mathbf{x} = [x_1, x_2]^T$.
- We sometimes **created** more: $\mathbf{x} = [1, x_1 x_1^2, x_1^3, \dots]^T$.
- ▶ Now, we've been given lots (3751) to start with.
- We need to reduce this number.
- 2 general schemes:
 - Use a subset of the originals.
 - Make new ones by combining the originals.

Finding a subset – example

▶ Take one feature – N values.

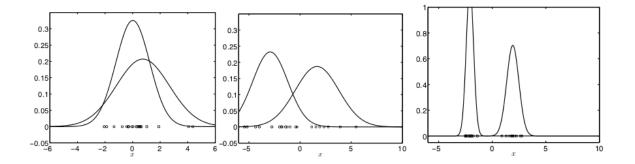


- ▶ Some values from objects in class 1, some from class 0.
- ▶ Split them based on class and compute μ and σ^2 for each class.
- Compute s for each feature:

$$s = \frac{|\mu_1 - \mu_0|}{\sigma_0^2 + \sigma_1^2}$$

Keep features with high s.

Examples



Features get better (higher s) from left to right...

$$s = \frac{|\mu_1 - \mu_0|}{\sigma_0^2 + \sigma_1^2}$$

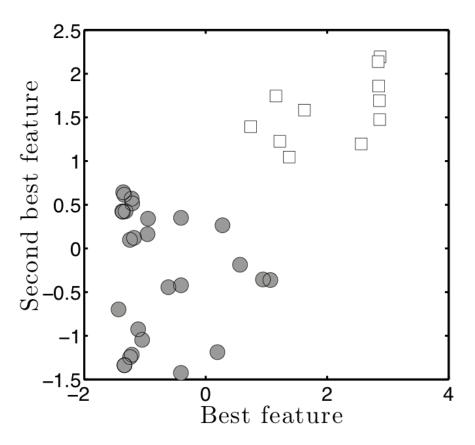
- ▶ Each feature has an *s*-score. The higher the better.
- Use the S features with the highest scores.
- ▶ How to choose *S*?

A feature selection scheme (CV)

- For each candidate S value:
- Split the data into C folds (just as in CV)
- For each fold...
 - 1. Find the feature scores on the **training** data.
 - 2. Train the classifier (whichever we choose).
 - 3. Record the performance.

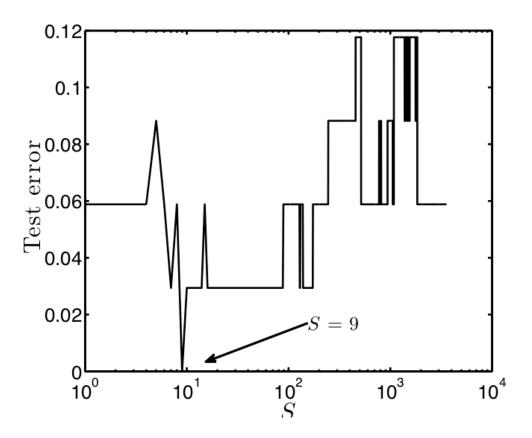
Important: Must only compute scores on training data.
 Otherwise we are implicitly using the test labels for training – biased.

Example



Best two features in our leukaemia data (points labeled by class).

Example



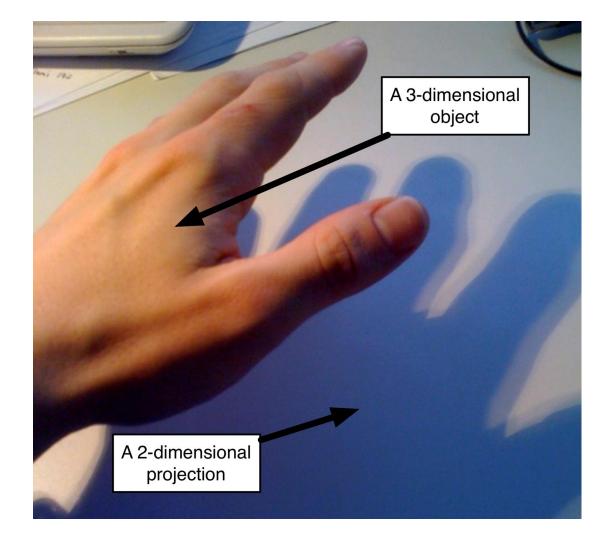
Performance as S increases.

Making new features

- An alternative to choosing features is making new ones.
- Cluster:
 - Cluster the features (turn our clustering problem around)
 - If we use say K-means, our new features will be the K mean vectors.
- Projection/combination
 - Reduce the number of features by projecting into a lower dimensional space.
 - Do this by making new features that are combinations (linear) of the old ones.

Projection

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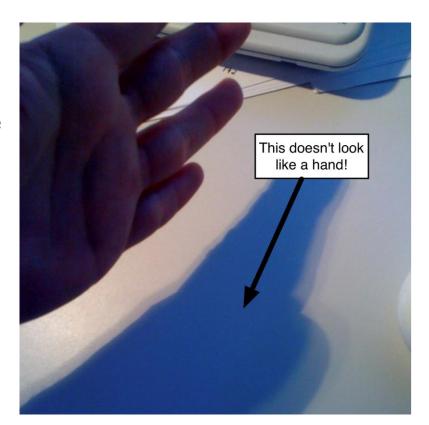


Projection

- ▶ We can project data (D dimensions) into a lower number of dimensions (M).
- ▶ Z = XW
 - \triangleright X is $N \times D$
 - **W** is *D* × *M*
- ► Z is N × M an M-dimensional representation of our N objects.
- ▶ **W** defines the projection
 - Changing W is like changing where the light is coming from for the shadow (or rotating the hand).
 - ► (X is the hand, Z is the shadow)
- ▶ Once we've chosen \mathbf{W} we can project test data into this new space too: $\mathbf{Z}_{new} = \mathbf{X}_{new} \mathbf{W}$

Choosing W

- ▶ Different **W** will give us different projections (imagine moving the light).
- ▶ Which should we use?
- ▶ Not all will represent our data well...



Principal Components Analysis

- Principal Components Analysis (PCA) is a method for choosing W.
- ▶ It finds the columns of **W** one at a time (define the mth column as \mathbf{w}_m).
 - ▶ Each $D \times 1$ column defines one new dimension.
- Consider one of the new dimensions (columns of Z):

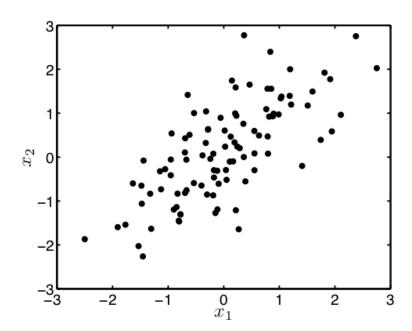
$$z_m = Xw_m$$

ightharpoonup PCA chooses \mathbf{w}_m to maximise the variance of \mathbf{z}_m

$$\frac{1}{N}\sum_{n=1}^{N}(z_{mn}-\mu_m)^2, \quad \mu_m=\frac{1}{N}\sum_{n=1}^{N}z_{mn}$$

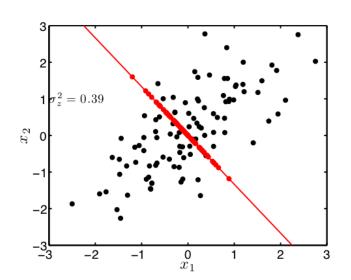
▶ Once the first one has been found, the w₂ is found that maximises the variance and is **orthogonal** to the first one etc etc.

PCA



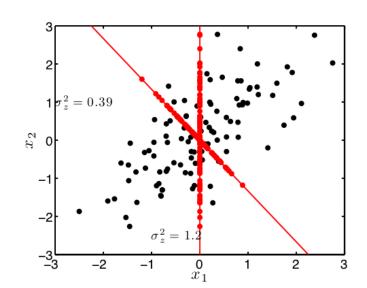
- Original data in 2-dimensions.
- ▶ We'd like a 1-dimensional projection.

PCA– a visualisation



- ▶ Pick some arbitrary w.
- Project the data onto it.
- Compute the variance (on the line).
- ► The position on the line is our 1 dimensional representation.

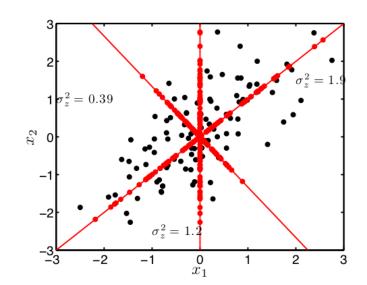
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PCA

a visualisation



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PCA – analytic solution

- ightharpoonup Could search for $\mathbf{w}_1, \ldots, \mathbf{w}_M$
- But, analytic solution is available.
- w are the eignvectors of the covariance matrix of X.
 - You don't need to know this!
- Python: sklearn.decomposition.PCA, R: prcomp

PCA – analytic solution

 $m \times n$ $m \times n$ $m \times m$

https://en.wikipedia.org/wiki/Singular_value_decomposition#/media/File:Singular_value_decomposition_visualisation.svg



Stitch Fix is using something called eigenvector decomposition, a concept from quantum mechanics, to tease apart the overlapping "notes" in an individual's style. Using physics, the team can better understand the complexities of the clients' style minds.



The Style Maven Astrophysicists of Silicon Valley

You know who knows machine learning? People who look at the stars all day. And when it comes to what constellations of clothes and shows and music yo... & wired.com

PCA – analytic solution

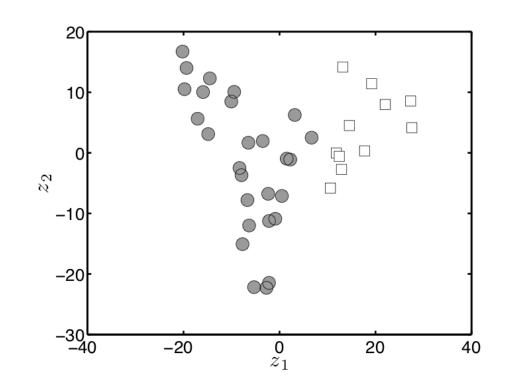
$$C = \frac{X^T X}{n-1}$$

$$C = (U\Sigma V^T)^T U\Sigma V^T / (n-1) =$$

$$= V \Sigma U^T U\Sigma V^T / (n-1) =$$

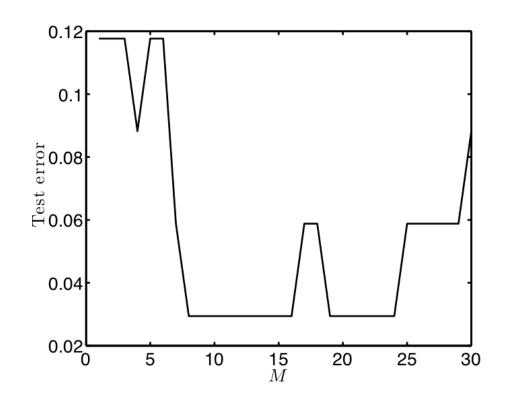
$$= V \frac{\Sigma^2}{n-1} V^T$$

PCA - leukaemia data



First two principal components in our leukaemia data (points labeled by class).

PCA – leukaemia data



Test error as more and more components are used.

Summary

- Sometimes we have too much data (too many dimensions).
- Need to select features.
- Features can be dimensions that already exist.
- Or we can make new ones.
- We've seen one example of each.