

# Computer lab 1 block 1

## Instructions

- Create a report to the lab solutions in PDF.
- Be concise and do not include unnecessary printouts and figures produced by the software and not required in the assignments.
- **Include all your codes as an appendix into your report.**
- A typical lab report should 2-4 pages of text plus some amount of figures plus appendix with codes.
- The lab report should be submitted via LISAM before the deadline.

## Assignment 1. Handwritten digit recognition with K-nearest neighbors.

The data file **optdigits.csv** contains information about normalized bitmaps of handwritten digits from a preprinted form from a total of 43 people. The data were first derived as 32x32 bitmaps which were then divided into nonoverlapping blocks of 4x4 and the number of on pixels are counted in each block. This has generated the resulting image of size 8x8 where each element is an integer in the range 0..16. Accordingly, each row in the data file is a sequence corresponding to 8x8 matrix, and the last element shows the actual digit from 0 to 9.

1. Import the data into R and divide it into training, validation and test sets (50%/25%/25%) by using the partitioning principle specified in the lecture slides.
2. Use training data to fit 30-nearest neighbor classifier with function *kknn()* and kernel="rectangular" from package *kknn* and estimate
  - Confusion matrices for the training and test data (use *table()*)
  - Misclassification errors for the training and test dataComment on the quality of predictions for different digits and on the overall prediction quality.
3. Find any 2 cases of digit "8" in the training data which were easiest to classify and 3 cases that were hardest to classify (i.e. having highest and lowest probabilities of the correct class). Reshape features for each of these cases as matrix 8x8 and visualize the corresponding digits (by using e.g. *heatmap()* function with parameters *Colv=NA* and *Rowv=NA*) and comment on whether these cases seem to be hard or easy to recognize visually.
4. Fit a K-nearest neighbor classifiers to the training data for different values of  $K = 1, 2, \dots, 30$  and plot the dependence of the training and validation misclassification errors on the value of  $K$  (in the same plot). How does the model complexity change when  $K$  increases and how does it affect the training and validation

- errors? Report the optimal  $K$  according to this plot. Finally, estimate the test error for the model having the optimal  $K$ , compare it with the training and validation errors and make necessary conclusions about the model quality.
5. Fit  $K$ -nearest neighbor classifiers to the training data for different values of  $K = 1, 2, \dots, 30$ , compute the error for the validation data as cross-entropy (when computing log of probabilities add a small constant within log, e.g.  $1e-15$ , to avoid numerical problems) and plot the dependence of the validation error on the value of  $K$ . What is the optimal  $K$  value here? Assuming that response has multinomial distribution, why might the cross-entropy be a more suitable choice of the error function than the misclassification error for this problem?

## Assignment 2. Linear regression and ridge regression

The data file **parkinson.csv** is composed of a range of biomedical voice measurements from 42 people with early-stage Parkinson's disease recruited to a six-month trial of a telemonitoring device for remote symptom progression monitoring. The purpose is to predict Parkinson's disease symptom score (motor UPDRS) from the following voice characteristics:

- Jitter(%),Jitter(Abs),Jitter:RAP,Jitter:PPQ5,Jitter:DDP - Several measures of variation in fundamental frequency
  - Shimmer,Shimmer(dB),Shimmer:APQ3,Shimmer:APQ5,Shimmer:APQ11,Shimmer:DDA - Several measures of variation in amplitude
  - NHR,HNR - Two measures of ratio of noise to tonal components in the voice
  - RPDE - A nonlinear dynamical complexity measure
  - DFA - Signal fractal scaling exponent
  - PPE - A nonlinear measure of fundamental frequency variation
1. Divide it into training and test data (60/40) and scale it appropriately. In the coming steps, assume that motor\_UPDRS is normally distributed and is a function of the voice characteristics, and since the data are scaled, no intercept is needed in the modelling.
  2. Compute a linear regression model from the training data, estimate training and test MSE and comment on which variables contribute significantly to the model.
  3. Implement 4 following functions by using basic R commands only (no external packages):
    - a. *Loglikelihood* function that for a given parameter vector  $\theta$  and dispersion  $\sigma$  computes the log-likelihood function  $\log P(T|\theta, \sigma)$  for the stated model and the training data
    - b. *Ridge* function that for given vector  $\theta$ , scalar  $\sigma$  and scalar  $\lambda$  uses function from 3a and adds up a Ridge penalty  $\lambda \|\theta\|^2$  to the minus log-likelihood

- c. *RidgeOpt* function that depends on scalar  $\lambda$ , uses function from 3b and function `optim()` with `method="BFGS"` to find the optimal  $\theta$  and  $\sigma$  for the given  $\lambda$ .
- d. *DF* function that for a given scalar  $\lambda$  computes the degrees of freedom of the Ridge model based on the training data.
4. By using function *RidgeOpt*, compute optimal  $\theta$  parameters for  $\lambda = 1, \lambda = 100$  and  $\lambda = 1000$ . Use the estimated parameters to predict the `motor_UPDRS` values for training and test data and report the training and test MSE values. Which penalty parameter is most appropriate among the selected ones? Compute and compare the degrees of freedom of these models and make appropriate conclusions.

### Assignment 3. Logistic regression and basis function expansion

The data file **pima-indians-diabetes.csv** contains information about the onset of diabetes within 5 years in Pima Indians given medical details. The variables are (in the same order as in the dataset):

1. Number of times pregnant.
  2. Plasma glucose concentration a 2 hours in an oral glucose tolerance test.
  3. Diastolic blood pressure (mm Hg).
  4. Triceps skinfold thickness (mm).
  5. 2-Hour serum insulin (mu U/ml).
  6. Body mass index (weight in kg/(height in m)<sup>2</sup>).
  7. Diabetes pedigree function.
  8. Age (years).
  9. Diabetes (0=no or 1=yes).
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1. Make a scatterplot showing a Plasma glucose concentration on Age where observations are colored by Diabetes levels. Do you think that Diabetes is easy to classify by a standard logistic regression model that uses these two variables as features? Motivate your answer.
  2. Train a logistic regression model with  $y = \text{Diabetes}$  as target  $x_1 = \text{Plasma glucose concentration}$  and  $x_2 = \text{Age}$  as features and **make a prediction for all observations** by using  $r = 0.5$  as the classification threshold. Report the probabilistic equation of the estimated model (i.e., **how** the target depends on the features and the estimated model parameters probabilistically). Compute also the training misclassification error and make a scatter plot of the same kind as in step 1 but showing the predicted values of Diabetes as a color instead. Comment on the quality of the classification by using these results.

3. Use the model estimated in step 2 to a) report the equation of the decision boundary between the two classes b) add a curve showing this boundary to the scatter plot in step 2. Comment whether the decision boundary seems to catch the data distribution well.
4. Make same kind of plots as in step 2 but use thresholds  $r = 0.2$  and  $r = 0.8$ . By using these plots, comment on what happens with the prediction when  $r$  value changes.
5. Perform a basis function expansion trick by computing new features  $z_1 = x_1^4$ ,  $z_2 = x_1^3 x_2$ ,  $z_3 = x_1^2 x_2^2$ ,  $z_4 = x_1 x_2^3$ ,  $z_5 = x_2^4$ , adding them to the data set and then computing a logistic regression model with  $y$  as target and  $x_1, x_2, z_1, \dots, z_5$  as features. Create a scatterplot of the same kind as in step 2 for this model and compute the training misclassification rate. What can you say about the quality of this model compared to the previous logistic regression model? How have the basis expansion trick affected the shape of the decision boundary and the prediction accuracy?

## Submission procedure

**First read ‘Course Information.PDF’ at LISAM, folder ‘Course documents’**

**When submitting the report, remember to specify in LISAM your group name and all group members that wrote the report!**

Group information

Group name

Group A1

Group members

Search for student to add

Anders Andersson (anand111) Remove

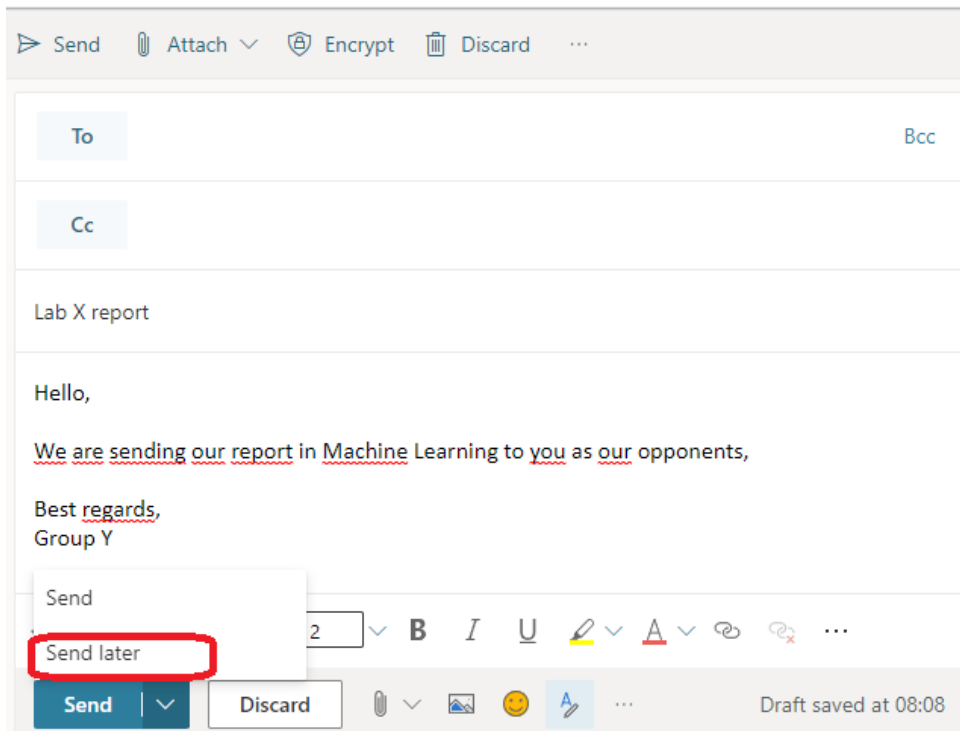
Per Persson (peper222) Remove

**If you are neither speaker nor opponent for this lab,**

- Make sure that you or your group mate submits the group report using *Lab X* item in the *Submissions* folder before the deadline. Make sure that the report contains the Statement Of Contribution describing how each group member has contributed into the group report.

**If you are a speaker for this lab,**

- Make sure that you or your group mate does the following before the deadline:
  1. submits the group report using *Lab X* item in the *Submissions* folder before the deadline. Makes sure that the report contains the Statement Of Contribution describing how each group member has contributed into the group report.
  2. Goes to LISAM→Course Documents→Deadlines.PDF, finds the deadline (date and time) for the current lab.
  3. Goes to LISAM→Course Documents→Seminars.PDF and find the group number of your opponent group
  4. Goes to LISAM→Course Documents→Groups.PDF and finds email addresses of the students in the opponent group
  5. Go to LISAM→Outlook app and in the Outlook web client creates a new message where you
    - Specify Lab X report as a title (X is lab number)
    - Specify email addresses of the opponents in the “To:” field
    - Attach your group PDF report.
    - **Important:** Click on arrow next to “Send” button, choose “Send Later” and specify the lab deadline as the message delivery time stamp (see figure)



**If you are opponent for this lab,**

- Make sure that you or your group mate submits the group report using *Lab X* item in the *Submissions* folder before the deadline. Make sure that the report contains the Statement Of Contribution describing how each group member has contributed into the group report.
- After the deadline for the lab has passed you should be able to receive the PDF report of the speakers per email. Compile it, read it carefully and prepare (in cooperation with your group comrade) **at least three questions/comments/improvement suggestions per lab assignment** in order to put them at the seminar.