TM10007: Machine Learning Project guidelines

March 12, 2021

1 Introduction

This guide describes how to conduct and report on the final research project for the TM10007 Machine learning course. In this project, you will use one of the provided medical datasets to train and evaluate different classification methods.

2 Goal of the project

The goal of the project is twofold. The first goal is to develop a machine learning method from the dataset that you are given. The second goal is to evaluate this model with the same dataset, in a way that you can quantify the performance as reliably as possible. Since both goals are achieved with the same dataset, you need to design a proper experimental method to achieve both of them. This is a common issue in machine learning research.

In the design of the machine learning method, containing preprocessing, possibly some feature extraction and a classifier, we expect you to motivate your design choices by either experiments or theoretical knowledge, or both. Of course, it is impossible to try every variation of every hyperparameter. Focus, therefore, on a number of relevant choices and evaluate their influence on the performance. In the report, describe the experiments you did, show the results and interpret them. Keep in mind that the final grade does not depend on the performance of the model, or the number of methods presented, but rather on the quality of the experimental work and the insight shown in the report.

3 Deliverables

You will have to hand in a written report similar to a paper, shortly giving an introduction to the problem, and describing your methods, results, and a discussion. The report should have no more than 5 pages, excluding references and appendices, and should contain the following:

- 1. An introduction concerning the (clinical) problem to be solved and a description of the dataset.
- 2. A description of the methods applied, with proper motivation, containing for example:
 - A description of the dataset
 - Preprocessing of the data
 - Classifier(s)
 - Experimental and evaluation setup
 - Statistics

You methods should be sufficiently detailed for another person to replicate your study.

- 3. A results section, providing a comprehensive overview of the results. When possible, try to illustrate your results with plots or tables.
- 4. A discussion, answering the following questions
 - What performance would this method reach on similar unseen data?
 - What are limitations of the study and what could be done to improve it?
 - Can this model be applied in the clinic? If not, what further steps could be taken to make it applicable in a clinical setting?

We do not expect you to perform a statistical analysis of the results, but we do expect you to perform the necessary experiments to reach a conclusion on the most promising method and expected performance. If possible, try to relate to your knowledge of the used methods. If possible, make concrete suggestions for further research.

5. A reflection of the team with the planning, communication strategy, and the division of roles and tasks. Each team member describes his/her individual contribution to both endproducts and his/her opinion on the process of this teamwork, the positive elements and the negative aspects of the combination of this team with this assignment. In total the reflection should be around 1500 words, individual reflection should be around 250 words.

Note that your code is also a deliverable, see the last section of this handout.

4 Grading

When grading the report, we will take the following aspects into account:

• Is a proper motivation supplied for the methodological choices?

- Was a proper experimental setup applied and described in the report?
- Are the results interpreted well, with respect to the different methods?
- Is the clinical relevance of the results discussed?
- Are the conclusions supported by quantitative results and figures?

Note that although a proper analysis and smart choice of methodology will lead to a classifier with high performance, a better performance does not automatically lead to a higher grade.

5 Deadlines

The code and report have to be handed in by April 17th 2020 (23:59 at the latest). The code has to be uploaded to your groups Github reposity and tagged with the tag *final*. Note that the code should be runnable using Google colab.

6 Datasets

All datasets contain quantitative medical image features, on which you are going to apply machine learning to find a relation between these and clinical outcome. This field of research is also known as radiomics.

For each dataset, you will have assess to a set of these quantitative imaging features, on which you have to perform a binary classification problem.

6.1 Prediction of tumor grade in brain cancer [1]

Gliomas are the most common form of brain tumors. The prognosis and treatment is mostly dependent on the tumor grade, which is defined by histological analysis. A rough categorization is that of low-grade (II/III) and high-grade (IV) glioma, the latter of which are also called glioblastoma. For the latter category the median survival is approximately 15 months, while low-grade glioma have a 10-year survival rate of approximately 47%.

The grade is known to affect the presentation of the tumor on magnetic resonance imaging (MRI), but a tissue sample is still needed to get a proper diagnosis. If a diagnosis could be made purely based on imaging, this might eliminate the need for a biopsy in some cases. Note, however, that a tumor resection by craniotomy is the generally preferred treatment for glioma, which automatically results in a tissue diagnosis.

The aim of this study is therefore to predict the tumor grade of glioma's (high or low) before surgery, based on features extracted from a combination of four MRI images: T2-weighted, T2-weighted FLAIR and T1-weighted before and after injection of contrast agent. A good performance on this dataset would be above 80% mean accuracy.

6.2 Distinguishing Alzheimer patients from healthy controls [2]

Dementia is a major problem worldwide, affecting over 36 million. In patients with dementia, the brain shrinks at a higher rate than in healthy people. Dementia consists of several underlying diseases, of which Alzheimer can be seen as one of the extreme forms. As everyone's brain shrinks with age, abnormal shrinkage, i.e. dementia, can be difficult from normal shrinkage.

Recent studies have tried to use quantitative MRI in order to distinguish between patients with normal and abnormal growth. Especially in specific regions such as the hippocampus, abnormal, heterogeneous growth patterns may be observed. To this end, the Alzheimer's Disease Neuroimaging Initiative (ADNI) consortium has build a MRI database of almost 1000 persons, including both patients with Alzheimer and healthy controls.

The aim of this study is therefore to distinguish between Alzheimer patients and healthy controls based features extracted from T1-weighted MRI. A good performance on this dataset would be above 75% mean accuracy. However, you might want to consider a different performance metric to evaluate the results.

6.3 Predicting tumor stage in head and neck cancer [3]

The annual incidence of head and neck (H&N) cancer is around 550.000 cases worldwide per year, with a mortality rate of 300.000 per year. This cancer comes in various forms, and therefore a wide spread in the prognosis and survival of H&N tumors. One of the most important clinical biomarkers is the tumor stage or so called T-stage. This biomarker is used to judge the malignancy of the tumor, and is thus closely related to survival.

While T-stage is highly related with volume, the molecular profile of the tumor does also play a role in the malignancy. However, quantification of the molecular profile can only be obtained after a biopsy or resection. While imaging, mostly computed tomography (CT) may be used to quantify T-stage, manual rating is observer dependent, subjective, and challenging.

The aim of this study is therefore to predict the T-stage (high/low) in patients with H&N cancer based on features extracted from CT. A good performance on this dataset would be above 70% mean accuracy. However, you might want to consider a different performance metric to evaluate the results.

6.4 Diagnosing heart disease in 12-lead ECG data [4]

The electrocardiograms (ECG) is an important diagnostic tool in cardiovascular diseases. The aim of this study is to automatically find abnormalities on 12-lead ECG. The problem is formulated as a binary prediction between normal and abnormal ECG's, where the abnormalities could be a 1st degree AV block, right bundle branch block, left bundle branch block, sinus bradycardia, atrial fibrillation or sinus tachycardia. The majority of samples contain no abnormalities.

Frequency features were extracted from the 12 leads by means of a Fourier transform. From the resulting 9000 features for 827 patients, it is possible to achieve a mean accuracy above 85%. However, you might want to consider a different performance metric to evaluate the results.

References

- [1] Spyridon Bakas, Hamed Akbari, Aristeidis Sotiras, Michel Bilello, Martin Rozycki, Justin S Kirby, John B Freymann, Keyvan Farahani, and Christos Davatzikos. Advancing the cancer genome atlas glioma mri collections with expert segmentation labels and radiomic features. *Scientific data*, 4:170117, 2017.
- [2] R. C. Petersen, P. S. Aisen, L. A. Beckett, M. C. Donohue, A. C. Gamst, D. J. Harvey, C. R. Jack, W. J. Jagust, L. M. Shaw, A. W. Toga, J. Q. Trojanowski, and M. W. Weiner. Alzheimer's Disease Neuroimaging Initiative (ADNI): clinical characterization. *Neurology*, 74(3):201–9, January 2010.
- [3] Hugo J. W. L. Aerts, Emmanuel Rios Velazquez, Ralph T. H. Leijenaar, Chintan Parmar, Patrick Grossmann, Sara Cavalho, Johan Bussink, René Monshouwer, Benjamin Haibe-Kains, Derek Rietveld, Frank Hoebers, Michelle M. Rietbergen, C. René Leemans, Andre Dekker, John Quackenbush, Robert J. Gillies, and Philippe Lambin. Decoding tumour phenotype by noninvasive imaging using a quantitative radiomics approach. Nature Communications, 5, June 2014.
- [4] Antônio H. Ribeiro, Manoel Horta Ribeiro, Gabriela M.M. Paixão, Derick M. Oliveira, Paulo R. Gomes, Jéssica A. Canazart, Milton P.S. Ferreira, Carl R. Andersson, Peter W. Macfarlane, Meira Wagner, Thomas B. Schön, and Antonio Luiz P. Ribeiro. Automatic diagnosis of the 12-lead ECG using a deep neural network. Nature Communications, 11(1):1–9, dec 2020.