

scripties voor de repository van FSE, FA <theses-fse@rug.nl>

Re: Thesis deposit from: Kaitlin Vos | degree programme: Computing Science

1 message

Michael Biehl <m.biehl@rug.nl>

Wed, Jul 19, 2023 at 4:42 PM

To: theses-fse@rug.nl

The thesis may be publicly accessible. Best wishes, Michael

On Wed, Jul 19, 2023 at 4:32 PM <theses-fse@rug.nl> wrote:

Deposit Form Student Theses Faculty of Science and Engineering

We've received your thesis successfully. After review, the library will provide access to it through our theses database: ([https://fse.studenttheses.ub.rug.nl/id/eprint/30800])

If you see incorrect information below, or need further information, please contact the Library of FSE (theses-fse@rug.nl)

The provided information:

: 19-07-2023

Dear sir/madam,

On 19-07-2023 Kaitlin Vos uploaded a paper (Research Report / Essay) in the FSE Thesis Repository. He/she indicated that this document should be publicly accessible on the internet.

Would you give permission for this?

X Yes

Please reply to theses-fse@rug.nl. A non-readable copy of this email correspondence will be saved in PDF with the related thesis.

Author(s)

Student number Family name First name Email address
S4266811 Vos Kaitlin k.x.vos@student.rug.nl

Degree programme

Degree programme Thesis type Computing Science Bachelor's Thesis

Supervisor(s) at UG

Family name First name, prefix

Email address

BiehlM.Bernoulliborg, Intelligente SystemenM.Biehl@rug.nlLovdalS.S.Medische Wetenschappen, Radiologie Research-Developm. S.S.Lovdal@rug.nl

Original title

Iterative Subspace Correction Procedure for Generalized Matrix Learning Vector Quantization in Biomedicine

Abstract of thesis

Neurodegenerative diseases are challenging to diagnose accurately due to their incurable nature and overlapping symptoms. One potential solution is to use a trained Generalized Matrix Learning Vector Quantization (GMLVQ) system on patients' FDG-PET scans to differentiate between different neurodegenerative diseases. In this thesis, we explore a modified method incorporating iterative subspace corrections. The GMLVQ system is trained on homogeneous healthy control cohorts, allowing extraction of vectors highlighting centre differences. By ignoring these in our feature space, we aim to eliminate the centre-specific variance present in the data. The method's

performance is assessed by using artificial data sets of increasing complexity, generated as Gaussian clusters, and evaluating it based on the known characteristics used to generate them. The results showed that the iterative method successfully aligned the subspace with the ground truth centre direction in simple cases, maintaining a high accuracy in disease classification and arbitrary classification in centre identification. In complex cases, the iterative method struggled to accurately capture the underlying differences. Larger data sets generally improved performance, but complexity posed challenges in classification accuracy. The centre-wise z-scored data approach demonstrated better adaptability, achieving high disease classification accuracy and arbitrary centre classification across different complexities.

Number of pages 111 Language of the thesis English Year of publication 2023

Additional comments

Indicate whether the document should be publicly accessible.: Yes (we'll ask your supervisor to confirm this)

File(s)

bCS_2023_VosK.pdf

Browser info: Mozilla/5.0 (Windows NT 10.0; Win64; x64) AppleWebKit/537.36 (KHTML, like Gecko) Chrome/114.0.0.0 Safari/537.36

Prof. Dr. Michael Biehl Bernoulli Institute for Mathematics, Computer Science & Artificial Intelligence P.O. Box 407, 9700 AK Groningen, NL https://www.cs.rug.nl/~biehl m.biehl@rug.nl