Prod.Type:FTP 8.07f/W(Jul162004) ED:SreejaGA NEUCOM : 10600 + model PAGN:Sakthi SCAN: pp.1-5(col.fig.:NIL) ARTICLE IN PRESS Available online at www.sciencedirect.com ScienceDirect 1 **NEUROCOMPUTING** 3 **FLSEVIER** Neurocomputing **I** (**IIII**) **III**-**III** www.elsevier.com/locate/neucom 5 7 Importance of electrophysiological signal features assessed by 9 classification trees 11 Andreea Lazăr^{a,d}, Raul Mureșan^{b,d}, Ellen Städtler^c, Matthias Munk^c, Gordon Pipa^{b,c,*} 13 ^aFrankfurt International Graduate School for Science, Johann Wolfgang Goethe University, Max-von-Laue-Str. 1, 60438 Frankfurt am Main, Germany ^bFrankfurt Institute for Advanced Studies, Johann Wolfgang Goethe University, Max-von-Laue-Str. 1 60438 Frankfurt am Main Germany 15 ^cMax-Planck-Institute for Brain Research, Deutschordenstraße 46, D-60528 Frankfurt am Main, Germany ^dCenter for Cognitive and Neural Studies (Coneural), Str. Saturn 24, 400504 Cluj-Napoca, Romania 17 19 Abstract 21 Sustained activity in prefrontal cortex is associated with the maintenance of information during short-term memory (STM). We have 23 used impurity reduction criteria of classification trees to investigate how the behavioral performance of a monkey during STM is reflected in the information content of three features of recorded signals: rates of individual neurons, oscillations in the LFP, and oscillations in the spiking activity. The LFP power in all bands, but in the α and β bands in particular, is more informative than the firing rate of 25 neurons and the spike power with respect to the monkey's performance. © 2006 Published by Elsevier B.V. 27

29 *Keywords:* Classification trees; Impurity reduction; Short-term memory; Oscillations; Monkey prefrontal cortex

³¹ 1. Introduction

33 Sustained activity in prefrontal cortex is associated with the maintenance of information during short-term memory 35 (STM) [3–7]. Motivated by the controversial discussion of the rate coding hypothesis [10] and the assembly hypothesis 37 [12] we investigated how the behavioral performance of a monkey performing an STM paradigm reflects in the 39 information content of three features: rates of individual neurons, oscillations in the LFP, and oscillations in the 41 spiking activity. For this purpose we used classification trees, a method that identifies structure in the feature space 43 and ranks features according to their information content Fig. 1.

45

55

47 2. Behavioral task and electrophysiological data

- 49 We recorded multi-unit ('MUA', 32 kHz sampling rate) and field potential ('LFP', 1 kHz sampling rate) activities 51
- *Corresponding author. Frankfurt Institute for Advanced Studies, Johann Wolfgang Goethe University, Max-von-Laue-Str. 1 60438 Frankfurt am Main Germany.

E-mail address: pipa@mpih-frankfurt.mpg.de (G. Pipa).

0925-2312/\$ - see front matter © 2006 Published by Elsevier B.V. doi:10.1016/j.neucom.2006.10.136 simultaneously from up to 16 fiber microelectrodes arranged in a 4×4 matrix with 500 µm spacing in the prefrontal cortex of two monkeys. Signals were filtered (.5-5 kHz (MUA) and 5-150 Hz (LFP) 3 dB/octave) and digitized, preprocessed by rejecting artifacts (movements, licking) and removing line noise at 50 ± .5 Hz. In total we analyzed four sessions with 1319 trials altogether (Sessions (1) 227; (2) 505; (3) 332; (4) 255).

The behavioral task of the monkeys was a visual short 67 term memory task. The task consisted of a sample period (first 500 ms) during which a sample stimulus was 69 presented, followed by 3 s of delay. After the delay a test stimulus that was either a matching or non-matching visual 71 object to the sample was presented. The monkey's task was to discriminate between matching and non-matching 73 stimuli and indicate its decision by a button press (match = left, non-match = right) on each trial. On 75 average, the monkeys gave correct responses in 80% of the trials. 77

3. Method

We compared neuronal activity recorded during trials in $_{81}$ which the monkey gave a correct response with activity

83

79

Please cite this article as: A. Lazăr, et al., Importance of electrophysiological signal features assessed by classification trees, Neurocomputing (2006), doi:10.1016/j.neucom.2006.10.136





A. Lazăr et al. / Neurocomputing I (IIII) III-III



5 Fig. 1. Time course of the visual short term memory task: after a -0.5-0 s baseline, a sample stimulus is presented for 500 ms(starting at time 0), followed 63 by a 3s delay, and a second matching/nonmatching test stimulus. Time line is divided in 10 consecutive windows of 500 ms each, from which the central 300 ms are used (gray rectangles) to extract features of the signal: rate, spike power and LFP power.

65

67

2

9

11 recorded during trials with a wrong response. For each recording session we matched trials with correct and 13 incorrect responses with respect to their number and

temporal proximity. We divided the signal of each trial 15 into 10 windows of 500 ms length from which we only considered the central 300 ms to make the windows

17 mutually independent (200 ms sliding window). For each window, we derived the spike rate by dividing spike counts

19 by the window size (300 ms), and computed the spike power and LFP power based on a multi-taper method for

21 frequencies of interest between 5 and 100 Hz ([8], frequency steps 5 Hz, smoothing frequency of 10 Hz). Based on the

23 spectral power estimated for each frequency of interest we extracted the average power in four frequency bands (1)

25 5-10 Hz; (2) 15-25 Hz; (3) 30-50 Hz; (4) 55-100 Hz). Thus, we derived a total of nine different features for each

27 channel, trial and temporal window: rate, four bands of LFP power and four bands of spike power. To allow for 29 compatibility between sessions we ensured the same

number of extracted features per session by randomly 31 selecting the smallest number of channels existing in all

four experiments, which was seven, leading in total to 63 33 features per session.

To assess the discriminative performance concerning 35 correct and incorrect responses per session of each feature, we employed classification trees based on the Gini Index

37 and entropy estimation [1,9]. Classification trees have been widely recognized as an effective techniques for classifica-

39 tion in data mining. They were designed to explore data in search of consistent patterns and relationships between

41 variables. A tree is constructed by recursively partitioning a learning sample of data. Making use of the class informa-

43 tion for this learning sample, the splits are selected in such a way, that for each step, the maximum separation between

45 different classes is achieved. The ideal split would divide the data so that all items belonging to one class would be

completely separated from the items belonging to other 47 classes.

49 Different measurements have been proposed for evaluating splits [1,9,11,13], but they all have the same basic goal

51 which is to favor homogeneity within each child node and heterogeneity between the child nodes. The goal of splitting

53 is to produce child nodes with minimum impurity (heterogeneity within a node) so that the difference

55 between the impurity of the parent node and those of the

children (impurity reduction: Γ) is maximized

$$\Gamma = I_{\text{parent}} - \sum_{i=1}^{\kappa} p(i)I(i), \qquad (1) \qquad 69$$

71 where k is the number of child nodes (k = 2 for a binary)tree) and p(i) is the fraction of items belonging to each child 73 node after the split. In our case, the impurity was assessed using the Gini index $(I_{\rm G})$ and entropy $(I_{\rm E})$ measurements 75

$$I_{\rm G}(i) = 1 - \sum_{j=1}^{C} f(i,j)^2,$$
(2) 77
79

 $I_{\rm E}(i) = -\sum_{i=1}^{C} f(i,j) \log_2 f(i,j),$ 81 (3) 83

where C is the number of classes and f(i, j) is frequency of 85 value *i* in node *i*.

One of the main advantages of using a decision tree 87 technique for classification is that this method inherently estimates the suitability of features for the separation of 89 items belonging to different classes. This property can be easily exploited when aiming for feature selection. Our goal 91 was to determine a ranking of the features of our extracted signals in different periods of the STM task. Selection and 93 ranking of features is emergent from the classification tree structure. 95

We investigated whether any of the described features was informative in distinguishing between the two classes: 97 correct and incorrect responses. To this end we built a classification tree for each of the 10 analysis windows that 99 were 300 ms long and covered the baseline, sample, delay and test period of the task. For each time window, one 101 decision tree was used to classify all trials according to the two classes.

103 To build the classification tree, our approach was to use one set of observations (learning sample) and cross validate 105 it with another completely independent set (testing sample). The prediction for the testing sample gives information on 107 the generality of the tree. Having a similar performance in training and test implies the extraction of the relevant 109 information describing the data and prevents overfitting. To find the 'right size' of the tree that maximizes generality 111 we started with a maximally detailed tree and used pruning of leaves until classification performance during test was 113 close to the one during training.

57

Please cite this article as: A. Lazăr, et al., Importance of electrophysiological signal features assessed by classification trees, Neurocomputing (2006), doi:10.1016/j.neucom.2006.10.136

NEUCOM : 10600

ARTICLE IN PRESS

A. Lazăr et al. / Neurocomputing ■ (■■■) ■■■=■■

1 4. Results

Since our results of the optimal tree size based on pruning and cross validation indicated that only small trees
can be reliably built on the analyzed data set, we assessed the impurity reduction (Γ) based on the first level of the
trees corresponding to the first split. We assessed Γ for each of 63 features of the spike and LFP signal in four
individual sessions with each 200 to 500 trials and in nine epochs that covered the baseline, sample, delay and test periods. Results based on the Gini Index and entropy were similar.

In Fig. 2, results from one of the four sessions are presented. The session contains 505 trials (258 correct and 59 247 incorrect). For each of the nine temporal windows (time line on the vertical as indicated on the right side of 61 the image), we computed the impurity reduction using rate (left column), LFP power (center), and spike power (right 63 column). Since we considered seven randomly selected recording channels, we obtain seven Γ values for rate, 7×4 65 Γ values for LFP power and $7 \times 4 \Gamma$ values for spike power. The four frequency bands for power considered are: 67 (a) 5-10 Hz; (b) 15-25 Hz; (c) 30-50 Hz and (d) 55-100 Hz.

13



Fig. 2. Impurity reduction, Γ, for separation between correct/incorrect responses for nine periods in time (see Fig. 1) and for signal features of seven Channels (A) rate, (B) LFP power, and (C) spike power. In A individual channels are indexed from 1 to 7. In B and C, four different frequency bands are 113 displayed (a) 5–10 Hz; (b) 15–25 Hz; (c) 30–50 Hz; (d) 55–100 Hz).

Please cite this article as: A. Lazăr, et al., Importance of electrophysiological signal features assessed by classification trees, Neurocomputing (2006), doi:10.1016/j.neucom.2006.10.136

3

69

NEUCOM : 10600

ARTICLE IN PRESS

A. Lazăr et al. / Neurocomputing [(IIII) III-III



Fig. 3. Average impurity reduction, $\bar{\Gamma}$, (using Gini index) for separation between correct/incorrect responses assessed for nine features and nine periods in time (see Fig. 1)

²³ Thus, we have in total 63 Γ values for each of the time windows.
²⁵ For individual channels and features the impurity.

For individual channels and features the impurity reduction ranges between 0 and 0.035 across baseline and all periods of the task. Impurity reduction (Γ) values increased during the test period for most LFP frequency

²⁹ bands and channels, but also for rate on channel 5, and for spike power on channel 2. Impurity reduction of different footures analyzed for the same channels are more similar

features analyzed for the same channels are more similar than Γ of the same feature from different channels. This

³³ implies that all considered features reflect on some level the same underlying mechanisms. Overall, low frequency
 ³⁵ millitation of the LEB here the birth discrimination.

³⁵ oscillations of the LFP have the highest discriminative power between correct and incorrect trials.
 37 To compare the discriminative power sense different.

To compare the discriminative power across different periods of the task irrespective of the channel's identity we assessed the average impurity reduction $\bar{\Gamma}$ across all channels (Fig. 3). In two of the four sessions we observed

⁴¹ an increased average impurity reduction $\overline{\Gamma}$ during the test period for LFP power between 5 and 50 Hz. The effect is

⁴³ strongest for α and β bands and more moderate for the γ band while the impurity reduction for the spike rate and

⁴⁵ spike power features are comparably low. Remarkably, the time course of $\overline{\Gamma}$ indicates that the impurity reduction

⁴⁷ during the test period is 4 to 5 times higher than during the rest of the task.

51 5. Conclusions

Δ

53 By using classification trees we intended to rank the importance of signal features for discriminative perfor-

55 mance. We found that the reliable size of extracted decision trees was rather small (in the order of one to a few splits),

57 and that the classification performance was low. The

reason was, first, that the data did not form well separated clusters, second, that single features were not highly discriminative, and third, that differences between features were rather small. Taken together this might either mean that encoding and maintenance of information in the prefrontal cortex relies on complex signals that express multiple features which are only weakly modulated by behavior, or, that signals which are strongly correlated to behavior were not described by the signal we extracted and analyzed. 81 82 83 83 84 85 85 86 87 89

6. Uncited reference

93

Acknowledgments

This work was in part funded by the Hertie Foundation (AL, RM, GP) and the Volkswagen Foundation (MM, GP).

101

103

79

91

95

97

99

References

- L. Breiman, J.H. Friedman, R.A. Olshen, C.J. Stone, Classification and Regression Trees, Rapport technique, Wadsworth International, Monterey, CA, 1984.
- [2] G. Buzsaki, A. Draguhn, Neuronal oscillations in cortical networks, Science 304 (2004) 1926–1929.
- [3] A. Compte, N. Brunel, P.S. Goldman-Rakic, X. J Wang, Synaptic mechanisms and network dynamics underlying spatial working memory in a cortical network model, Cereb. Cortex 10 (2000) 910–923.
- [4] J.M. Fuster, G. Alexander, Neuron activity related to short-term memory, Science 173 (1971) 652–654.
- [5] J.M. Fuster, Memory in the Cerebral Cortex, MIT Press, Cambridge, 113 MA, 1995.

Please cite this article as: A. Lazăr, et al., Importance of electrophysiological signal features assessed by classification trees, Neurocomputing (2006), doi:10.1016/j.neucom.2006.10.136

NEUCOM : 10600

ARTICLE IN PRESS

A. Lazăr et al. / Neurocomputing I (IIII) III-III

- 1 [6] P.S. Goldman-Rakic, Cellular basis of working memory, Neuron 14 (1995) 477–485.
- [7] E.K. Miller, C.A. Erickson, R. Desimone, Neural mechanisms of visual working memory in prefrontal cortex of the macaque, J. Neurosci. 16 (1996) 5154–5167.
 [8] P.P. Micon P. P. Andréa C. La contract of the macaque of
 - [8] P.P. Mitra, B. Pesaran, Analysis of dynamic brain imaging data, Biophys. J. 76 (1999).
- 7 [9] J.R. Quinlan, C4.5: Programs for Machine Learning, Morgan Kaufmann, San Mateo, CA (1993).
- [10] M.N. Shadlen, W.T. Newsome, Neural basis of a perceptual decision
 9 in the parietal cortex (area LIP) of the rhesus monkey, J. Neurophysiol. 86 (2001).
 11
- [11] Y.S. Shih, Families of splitting criteria for classification trees, Statist. Comput. 9 (1999) 309–315.
- [12] W. Singer, Neuronal synchrony: a versatile code for the definition of relations, Neuron 24 (1999).
- [13] P.C. Taylor, B.W. Silverman, Block diagrams and splitting criteria 15 for classification trees, Statist. and Comput. 3 (1993) 147–161.

NCORPERING