UNIVERSITY OF BIRMINGHAM

Research at Birmingham

Prediction of postoperative opioid analgesia using clinical-experimental parameters and electroencephalography

Gram, M.; Erlenwein, J.; Petzke, F.; Falla, Deborah; Przemeck, M.; Emons, M. I.; Reuster, M.; Olesen, S. S.; Drewes, A. M.

Document Version
Peer reviewed version

Link to publication on Research at Birmingham portal

Prediction of Postoperative Opioid Analgesia using Clinical-Experimental Parameters and Electroencephalography

M Gram[1], J Erlenwein[2], F Petzke[2], D Falla[2], M Przemeck[3] , MI Emons[2], M Reuster[2], SS Olesen[1] & AM Drewes[1,4]

1. Mech-Sense, Department of Gastroenterology and Hepatology, Aalborg University Hospital, Aalborg, Denmark
2. Pain Clinic, Department of Anesthesiology, University Hospital, Georg-August-University of Göttingen; Göttingen, Germany
3. Department of Anesthesiology and Intensive Care, Annastift, Hannover, Germany
4. Clinical Institute, Aalborg University Hospital, Aalborg, Denmark

Running head: Prediction of Postoperative Analgesia

Corresponding author: Professor Asbjørn Mohr Drewes, MD, PhD, DMSc

Mech-Sense, Department of Gastroenterology & Hepatology,

Aalborg University Hospital,

Mølleparkvej 4, 9100 Aalborg, Denmark

Telephone +45 99326243

E-mail: amd@mech-sense.com

Conflicts of interest: None

Category: Original article

What does this study add:

- This study is the first clinical investigation using electroencephalography for prediction of postoperative opioid analgesia.
- The combined method of machine learning and electroencephalography offers promising results for future developments of personalized pain treatment

Abstract

Background:

Opioids are often used for pain treatment, but the response is often insufficient and dependent on e.g., the pain condition, genetic factors and drug class. Thus, there is an urgent need to identify biomarkers to enable selection of the appropriate drug for the individual patient, a concept known as personalized medicine. Quantitative sensory testing (QST) and clinical parameters can provide some guidance for response, but better and more objective biomarkers are urgently warranted. Electroencephalography (EEG) may be suitable since it assesses the central nervous system where opioids mediate their effects.

Methods:

Clinical parameters, QST and EEG (during rest and tonic pain) was recorded from patients the day prior to total hip replacement surgery. Postoperative pain treatment was performed using oxycodone and piritramide as patient-controlled analgesia. Patients were stratified into responders and non-responders based on pain ratings 24 hours post-surgery. Parameters were analysed using conventional group-wise statistical methods. Furthermore, EEG was analysed by machine learning to predict *individual* response.

Results:

Eighty-one patients were included, of which 51 responded to postoperative opioid treatment (30 non-responders). Conventional statistics showed that more severe pre-existing chronic pain was prevalent among non-responders to opioid treatment (P=0.04). Preoperative EEG analysis was able to predict responders with an accuracy of 65% (P=0.009), but only during tonic pain.

Conclusions

Chronic pain grade before surgery is associated with the outcome of postoperative pain treatment. Furthermore, EEG shows potential as an objective biomarker and might be used to predict postoperative opioid analgesia.

Significance

Personalized pain medicine based on EEG has received limited attention within the field apart from a single experimental study. The current clinical study demonstrates the viability of EEG as a biomarker and with results in line with previous experimental results.

Introduction

Opioids are the primary analgesic drugs used to treat moderate to severe pain, including pain after surgery (Liu and Wu, 2007). However, treatment of postoperative pain remains in many cases unsatisfactory, leading to unnecessary suffering (Dolin et al., 2002; Drewes et al., 2013; Maier et al., 2010; Sommer et al., 2008). Furthermore, inadequate analgesia carries the risk of developing persistent long-term pain (Gerbershagen et al., 2009; Kehlet et al., 2006; VanDenKerkhof et al., 2012). Better understanding of the inter-individual differences in pain processing is paramount to achieve improved and personalized opioid treatment (Bruehl et al., 2013). However, there is no method to determine if opioid treatment will provide analgesic relief in the postoperative setting. Some studies have focused on quantitative sensory testing (QST), but with contradictory results (Grosen et al., 2013). This indicates a need for improvement in the ability to accurately predict analgesic effect of opioids (Grosen et al., 2013).

Pain processing can be investigated using imaging techniques such as magnetic resonance imaging (Ahmedzai, 2013). However, electroencephalography (EEG) is more clinically feasible to assess pain processing as it has a considerably lower cost and can be used directly at the patient's bedside or in the outpatient clinic. Furthermore, EEG can assess inter-individual differences in pain processing by predicting the experience of pain within a single subject (Schulz et al., 2011). For instance, EEG recorded during rest was used to predict pain perception during experimentally induced pain (Nir et al., 2012) and in the clinic to asses analgesic efficacy (Graversen et al., 2012). We have previously shown that EEG during tonic pain is more reliable and superior than resting EEG for prediction of analgesic efficacy in healthy volunteers (Gram et al., 2014, 2015). The present study now aims to investigate EEG for prediction of analgesic efficacy in clinical population.

We *hypothesized* that risk factors for analgesic inefficacy of postoperative opioid treatment could be identified preoperatively. Additionally, we hypothesized that EEG could be used as an objective biomarker for predicting analgesic efficacy. Thus, the *aims* of this study were to a) determine the effect of the postoperative pain treatment, b) investigate risk factors associated with non-response to analgesic treatment and c) predict the analgesic response based on the pre-operative EEG (during rest and tonic pain) on an individualized basis in patients undergoing hip replacement surgery. Unlike previous EEG prediction studies which used the spectral content within pre-defined frequency bands (Gram et al., 2015) to evaluate brain rhythmicity, we also applied a functional connectivity evaluation which considers the brain as a large inter-connected network and investigates the interactions between different cortical regions (Hardmeier et al., 2014). We also applied machine learning methods which have the ability to assess the interaction between several EEG features at once without making a priori assumptions about the data (Gram et al., 2015).

Methodology

The study was conducted at the Pain Clinic, Center for Anesthesiology, Emergency and Intensive Care Medicine at the University Hospital Göttingen, Germany and the Orthopedic University Hospital of the Medical School Hannover, Germany. The study was approved by the Ethical committees of the University Hospital of Göttingen (No 19/2/13) and conducted according to the recommendations of the Declaration of Helsinki.

Study subjects

Patients admitted to the Orthopaedic University Hospital Annastift in Hannover for total hip replacement were recruited between April and August 2013.

Inclusion criteria were patients above 18 years of age and able to give informed consent. Exclusion criteria were 1) severe neurological disease, which might interfere with the EEG recordings or experimental pain testing (including dementia); 2) Severe psychiatric disease such as major depression or schizophrenia or active drug abuse; 3) High dose of preoperative opioid therapy (> 30 mg/day oral morphine equivalent). In case of post-operative delirium making the subject unable to complete the study due to disorientation or inability to answer questionnaires, the subject was excluded.

Study outcomes and risk factors

Response to post-surgical analgesic treatment was assessed using a response score based on the results of patients outcomes assessed by the "Quality Improvement in Postoperative Pain Management" Questionnaire (QUIPS), which is a validated German outcome measurement instrument for post-operative quality control, see below (Meissner et al., 2008).

Predictors for insufficient analgesic response were investigated and included clinical patient characteristics and standardized assessment of the preoperative pain symptoms, QST parameters, conditioned pain modulation (CPM) effect and EEG during rest and cold pain. Clinical patient data included age, sex and body mass index (BMI) while parameters on the preoperative pain condition included pain duration, chronic pain grade (Von Korff et al., 1992), Mainz Pain Staging System (MPSS) (Schmitt and Gerbershagen, 1990), Western Ontario and McMaster Universities Osteoarthritis Index (WOMAC) (Bellamy, 2005) as well as pain rated on passive rotation of the hip. QST parameters included heat and cold pain thresholds.

Study procedures

Figure 1 shows the experimental setup. Patients were examined one day prior to their operation. Patients were told to refrain from smoking and from drinking coffee and other caffeine-containing beverages 2 hours prior to the clinical testing.

All clinical testing were conducted between 15:00 and 18:00 on the preoperative day by the same examiner. The clinical testing commenced with familiarization of the cold pressor test and the experimental evoked pain conditions applied. Patients were placed on a bed in a semi-recumbent position, and the EEG cap was mounted on their head with conducting gel inserted for each electrode. Resting EEG was recorded followed by EEG recording during the cold pressor test.

Perioperative pain management

The patients preoperative pain medication (type and dosage) was noted and the Medication Quantification Score (MQS), which is a reliable and validated score for quantifying analgesics, was calculated in order to obtain a comparable metric for all different analgesics (Masters Steedman et al., 1992). It enables calculation characterization of analgesics when many analgesics are involved and doses are irregular, as is the case for the patient group in this study. It was calculated for each non-opioid and opioid based on weights assigned by medication class and dosage level (level 1 = sub-therapeutic dosage and/or on demand, level 2 = lower 50% of the therapeutic dose range, level 3 = upper 50% of the therapeutic dose range, level 4 = supra-therapeutic dose) using the 1998 detriment weights (Stormo, Kee, Steedham, & Middaugh, 1998). The detriment weights are summed by the dosage level to provide the final score. To provide a quantitative index for analgesics usage suitable for statistical analysis these scores were summed.

In the evening before and morning prior to surgery, all patients were pre-medicated with 20-30mg dipotassium chlorazepat. General anaesthesia was performed according to the local clinical standards, with remifentanyl (1-1.5 µg/kg /3 min) and propofol (1-2 mg/kg). Orotracheal intubation was facilitated by 0.5 mg/kg atracurium. Propofol (3.5-4.5 mg/kg/h) and remifentanyl (0.15-0.25 µg/kg/min) were used for maintenance of anesthesia. During the implantation of the femoral shaft, intravenous injection of 0.1 mg/kg piritramide and 15 mg/kg metamizol (or paracetamol, if contra-indicated) were administered.

In the recovery phase, patients received 10 or 20 mg slow-release oxycodone (10 mg if their weight was below 60 kg and age above 70 years, otherwise 20 mg was given) along with 600 mg ibuprofen. Pain was titrated using intravenous piritramide bolus application of 3.75mg until pain intensity was 3 or below on a numerical rating scale (NRS; 0 = no pain und 10 = worst imaginable pain) (see section on pain assessments). Then patients received patient-controlled analgesia (PCA) with piritramide for 24 hours using piritramide

(Perfusor fm PCA, B. Braun; single dose 2 mg, lockout 10 min, limit 30 mg/4 hours) at a NRS of 3 or below and as soon as they were able to operate the PCIA system. On the ward, oxycodone (twice a day) and ibuprofen (three times a day) was administered according to the first dosage and followed a standardized post-operative pain treatment protocol (Erlenwein et al., 2012).

Then the 24 hour opioid dose was calculated as oral morphine equivalent (ME; conversion factor to morphine: piritramide 1.5, hydromorphone 0.13, oxycodone 0.75, intravenous vs. oral morphine 3:1).

Chronic pain assessments

Before surgery, patients indicated their duration of pain, and their chronic pain grade was assessed (Von Korff et al., 1992). This grading system divides patients into four grades of chronic pain based on their pain intensity and disability, which can be summarized as:

- Grade I: low pain intensity, low disability
- Grade II: high pain intensity, low disability
- Grade III: high disability – moderately limiting
- Grade IV: high disability – severely limiting

The chronic pain grades I and II are based on the pain intensity, but only for patients with a low disability following their pain. Grades III and IV consists of patients with a certain degree of disability from their pain, and therefore disregards the pain intensity, but solely looks at how limiting the pain is (Von Korff et al., 1992).

Furthermore, the clinician assessed the patient's pain profile using the Mainz Pain Staging System (MPSS), which divides patients into three pain stages of increasing pain chronicity (Frettlöh et al., 2003; Schmitt and Gerbershagen, 1990).

Passive hip rotation

A test was performed to assess the magnitude of evoked pain with passive rotation of the hip scheduled for replacement. First, the leg was brought into a 90° flexed position, with the knee flexed at 90° at the same time, and then slowly rotated interiorly until the patient reported the first onset of pain (pain detection threshold). Then, the passive rotation was continued until the pain threshold was reached and this position was maintained for 30 seconds. Afterwards the patients were asked to rate the evoked hip pain on a NRS.

Quantitative sensory testing (QST)

Perceived pain intensity was rated on a NRS.

Heat pain threshold

Heat pain assessments were performed with $9cm^2$ Peltier Thermode, 10 cm proximal to the wrist of the right volar forearm, using a 'Thermo Tester' (TSA II NeuroSensory analyzer, Medoc Ltd, Ramat Yishai, Israel).The temperature was gradually increased from a baseline of 32°C baseline at a rate $1°Cs^{-1}$ to a maximum temperature of 52°C. The subjects were instructed to press a button when the heat stimulus became painful and this was documented as the heat pain threshold. Four consecutive assessments were performed and the average of the last three threshold estimations was retained for further analysis.

Cold pressor test

Patients were instructed to submerge their non-dominant hand in water until the wrist was covered (~8 °C) and keep it submerged for 2 minutes. The temperature of the water was controlled. The maximum pain intensity perceived during the test was rated on a NRS.

Conditioned pain modulation (CPM)

A CPM paradigm was utilized to investigate the generation of descending pain modulation. The CPM can be measured by application of a test-stimulus (in this case heat pain threshold), during and following the application of a conditioning stimulus (in this case the cold pressor test). An increase of the pain threshold after application of the conditioning stimulus indicates the presence of descending pain modulation (Pud et al., 2009).

In this study, heat pain threshold was repeated at 120 seconds after the cold pressor test was initiated. This was done to assess the conditioned pain modulation of each patient. The difference between the heat pain threshold at baseline and 120 seconds after initiation of the cold pressor test was determined and expressed as a percent to quantify the CPM effect.

Post-operative pain

The outcome quality of post-operative pain management was assessed with the QUIPS-questionnaire which is a validated German outcome measurement instrument for post-operative quality control, and the national register currently includes more than 400,000 patients from around 200 hospitals. It consists of three questions asking for 1) pain intensity during movement, and 2) the least and worst pain over the last 24 hours, all rated on the NRS. Furthermore, it includes 3) questions about patients´ pain intensity and postoperative functional restrictions (mobilization, coughing/deep breathing, night-sleep, yes/no) (W Meissner et al., 2006; Winfried Meissner et al., 2008). The first 24 hours were chosen since it was the most standardized period in the post-surgery treatment, with some patients commencing rehabilitation after 24 hours, while other remained on bed-rest. The questionnaire is in German, but a translation of the questions to English is available (Winfried

Meissner et al., 2008). Hence, it has served as model for the internationally used PAIN OUT questionnaire, which is available in various languages (R. Zaslansky et al., 2015; Ruth Zaslansky et al., 2014).

Since there is no validated definition for a positive opioid response in a perioperative setting, we developed a "response score" to determine analgesic response after surgery based on various patient´s reported outcome-items used in the QUIPS questionnaire. Points indicating an overall successful pain management were awarded up to a maximum of 10 by the following criteria:

- Maximal pain NRS < 5: = + 2 points
- Pain on movement NRS < 5 = + 2 points
- Minimum pain NRS < 3 = + 2 points
- No pain on mobilization = + 1 point
- No pain when coughing = + 1 point
- No pain when waking up = + 1 point
- No effect on mood = + 1 point

Definition of cut off points was based on typical intervention thresholds for postoperative pain management (e.g. pain on movement or maximal pain of 5) (Maier et al., 2010; Rothaug et al., 2013). Interaction of pain with function was included to derive a clinically meaningful response. Patients with a score of 5 or higher were considered opioid responders, those with scores of 4 or lower non-responders to opioids.

EEG recordings

EEG was acquired using a 34-channel cap (34ch prewired cylindrical Ag/AgCl electrodes, MEQNordic A/S, Jyllinge, Denmark) and amplified on a Nuamp system (NuAmp, Neuroscan, El Paso, TX, USA) and recorded for later analysis. The cap was placed symmetrical in a standardized position 3 cm above the nasium. Electrode gel was applied into each of the recording channels to reduce the electrode impedance below 5 kΩ. Recordings were performed in a dimmed light room with all unnecessary electrical equipment turned off to avoid 50 Hz contaminations. Sampling frequency was 1000 Hz.

First resting EEG was acquired and patients were instructed to minimize eye movements and refrain from talking. Open or closed eyes were alternated between in 4 sequences of 2.5 min, starting with eyes open. For this study, only the first recording of 2.5 min eyes open was used.

EEG during the cold pressor test was commenced as soon as the patient immersed their hand into the water. The first 60 seconds of recording after hand immersion was used for analysis to avoid artefacts induced by the conditioned pain modulation procedure.

Pre-processing

The data was pre-processed with Neuroscan software (Neuroscan 4.5, Neuroscan, El Paso, TX, USA) in the following steps: 1) zero-phase shift notch filtering (49–51 Hz) using a finite impulse-response filter with a slope of 24 dB/octave; 2) zero-phase shift band-pass filtering (1-80 Hz) using a finite impulse-response filter with a slope of 12 dB/octave; 3) blinded visual inspection of data quality for all channels using linked-ear reference. Channels with abnormal signals were discarded and replaced by signals interpolated from neighbouring electrodes; 4) Re-referencing to the average electrode.

Spectral analysis

Spectral indices were calculated using Matlab 2012a (The Mathworks, Inc., Natick, MA, USA) in order to obtain the relative EEG amplitude. Calculations were performed using a wavelet transform as this has a superior time-frequency resolution than the more common Fourier transform (Akin, 2002). The continuous wavelet transform was applied to EEG from each channel using the complex Morlet wavelet as a mother wavelet function with a bandwidth of 10 Hz and a center frequency of 1 Hz. Scales for the mother wavelet was chosen to match frequencies ranging from 1 to 32 Hz with a 0.5 Hz between-scale frequency interval. The absolute values of the obtained wavelet coefficients were used for analysis and divided into the following standardized frequency bands: delta (1 – 4 Hz), theta (4 – 8 Hz), alpha (8 – 12 Hz) and beta (12 – 32 Hz). The wavelet coefficients were averaged over time and scales contained within each frequency band were summed together to yield the absolute activity within each frequency band. The relative activity was calculated separately for each channel by dividing each frequency band with the total energy of all bands and multiplying by 100. The values then represent the percentage of total amplitude contained in each frequency band.

Functional connectivity analysis

Functional connectivity is a new approach within EEG, but has been used within structural and functional magnetic resonance imaging (Hardmeier et al., 2014). Functional connectivity considers the brain as a complex network of inter-connected nodes (Hardmeier et al., 2014). Several methods exist for estimation of functional connectivity within EEG analysis and most methods work by investigating the phase-relationships between EEG signals for different electrodes, where electrodes with similar phases are thought to be exchanging information (Nolte et al., 2008). The phase-lag index (PLI) is based on a consistent lag between

9

instantaneous phases between two signals and thus ignores most zero-phase phase-relations in order to discard volume conduction noise (Nolte et al., 2008).

The PLI was calculated for the same frequency bands as used for the spectral analysis, using the implementation from The Neurophysiological Biomarker Toolbox (NBT) (http://www.nbtwiki.net/). The signal was divided into time windows and the PLI was calculated for each window. Lastly, the results for all time windows were averaged. The window width was set to twice the sampling frequency (2000 samples) as this provides a frequency resolution of 0.5 Hz, which is equal to the spectral analysis. The band-pass filter was a 1st order butterworth filter (Lehembre et al., 2012).

Machine learning analysis

A support vector machine (SVM) was used for the machine learning approach. The SVM is a binary classifier which optimizes the decision threshold between two groups without any *a priori* assumptions about the data (Cortes and Vapnik, 1995). The SVM was chosen over other machine learning classifiers since it has previously been used for other prediction studies in pain medicine (Graversen et al., 2012; Olesen et al., 2013a).

To avoid over-fitting, the most discriminative features were selected using the criteria for joint mutual information, as this criterion has been found to provide the best selection for data sets with a limited number of samples (Brown et al., 2012). For both resting EEG and EEG during cold pain, features were selected using the joint mutual information criterion and subsequently used for SVM classification. The number of features to be used was determined by investigating the accuracy of the classifier by gradually increasing the number of features up to 15. Accuracy was defined as the ratio between correctly classified subjects and total number of subjects in percentage. The number of features that yielded the highest accuracy on both study days was chosen for the final analysis due to the importance of reliability in the classifier. Classification was performed using the libSVM toolbox (version 3.20) for Matlab (Chang and Lin, 2011). A linear kernel function was used to avoid over-fitting of the data (Gong et al., 2011). The cost parameter C of the SVM was set to 1. Classification accuracy was determined using leave-one-out cross-validation which consisted of removing one subject before performing feature selection and training the SVM classifier using all remaining subjects. The subject that was removed was then used to test the predictive capability of the classifier. This was repeated until all subjects had been left out, to assess the overall accuracy of the classifier (Gong et al., 2011; Gram et al., 2015; Graversen et al., 2011). Accuracy, positive predictive value and negative predictive value of the classification were calculated.

Lastly, the feature selection method was applied to the complete dataset, in order to indicate which features were most useful for classification.

Statistical analysis

All data are reported as mean ± standard deviation unless otherwise stated. Logistic regression was used to determine the capability of different clinical variables to predict post-operative analgesia. Significant predictors in univariate analysis were included in a multivariate analysis and bootstrapping (1000 samples) was used for internal validation of the multivariate estimates. Analysis of Multichannel EEG was carried out using a mixed analysis of variance (ANOVA) model with electrode (i.e., electrode 1 through 34) as within-subjects factor and responder group (i.e. responder vs. non-responder) as between-subjects factor. The F-test results were corrected for identity covariance matrix by the Greenhouse–Geisser method to take into account possible intragroup correlations. Additionally, data were analyzed using the previously described machine learning approach. SVM classifications were analysed using chi-square. P-values below 0.05 were considered statistically significant.

Results

Out of 175 patients screened for the study, 112 was eligible for the study and signed the informed consent (24 patients declined to participate, 1 was below 18 years of age, 1 was already participating in another study, 1 due to replacement of both hips in the same surgery, 1 because informed consent could not be obtained, 13 for planned spinal anaesthesia during surgery, 4 due to high preoperative opioid use, 1 due to drug abuse and 17 due to a history of neurological conditions). Out of the 112 patients who signed the informed consent, 81 were included in the final analysis (15 excluded because surgery was cancelled or consent was withdrawn, 3 because PCA was not administered, 1 due to postoperative delirium, 1 because of sevoflurane anaesthesia, 1 because of basal-bleeding history, 1 due to insufficient time to complete measurements and 9 patients for having one or more unusable EEG recordings due to poor quality). Demographics and clinical characteristics of the final 81 patients (36 males and 45 females) are reported in Table 1.

Response to post-operative analgesic treatment

Fifty-one patients (63%) were classified as opioid responders and 30 (37%) were classified as opioid non-responders based on the response score from the QUIPS questionnaire. Details of the two groups are reported in Table 2. Mean score was 8.4 ± 1.3 for the responder group and 3.5 ± 1.5 for the non-responder group (P < 0.001).

Prediction of post-operative analgesia

Group-wise analysis

Clinical variables

Clinical and demographic characteristics, pain medications and QST results stratified on responders and non-responders are reported in Table 2. Chronic pain grade ($P = 0.007$) and MPSS stage ($P = 0.01$) were associated with the postoperative analgesic response groups. Non-responders predominantly belonged to more severe chronic pain grades (III and IV) and were skewed towards MPSS stage III (high pain chronicity). Furthermore, responders experienced more pain during the hip rotation test compared to the non-responders ($P = 0.03$). In contrast, none of the QST parameters were associated with post-operative analgesia response (all $P > 0.1$). Results from the multivariate analysis (Table 3) revealed that only the chronic pain grade ($P = 0.02$) remained an independent predictor of postoperative analgesic response and the association remained significant after internal bootstrapping validation. Using the chronic pain grade to classify patients into response groups (Responders: Grade I and II and non-responders: Grade III and IV) yielded an accuracy of 63% (positive predictive value = 81%; negative predictive value = 50%).

Resting EEG

No differences in *resting EEG* were found between responders and non-responders for spectral EEG indices (all $P > 0.1$) or functional EEG connectivity (PLI) (all $P > 0.1$). The topographical plots are shown in Figure 2 for the spectral indices and functional connectivity in Figure 3.

EEG during cold pain

There were no differences in spectral EEG indices (all $P > 0.3$) or functional connectivity (all $P >= 0.1$) between responders and non-responders *during cold pain*. The topographical illustration of the EEG spectral indices for responders and non-responders is shown in Figure 4 while the topography of the functional connectivity is illustrated in Figure 5.

Individual subject analysis (machine learning)

Resting EEG

Machine learning analysis was unable to distinguish between responders and non-responders with any number of resting EEG features (spectral EEG indices and functional connectivity) (maximum accuracy = 58%; $P = 0.2$).

EEG during cold pain

Machine learning analysis discriminated between responders and non-responders when selecting features freely from both spectral indices and functional connectivity measures derived from the EEG. The most accurate classification was achieved using only one feature from the entire dataset. This was the delta spectral index band from frontally placed FP1 electrode with an accuracy of 65 % (positive predictive value = 76%; negative predictive value = 53%; $P = 0.009$).

To ensure that the results were not related to the stratification of subjects, the analysis was also repeated by stratification based on opioid consumption within the first 24 hours after surgery. The cut-off value for opioid consumption was set at 65 mg morphine equivalent units, resulting in 45 patients in the high dose opioid group (i.e. > 65 mg morphine) and 37 patients in the low dose opioid group. Here it was not possible to discriminate between responders and non-responders (maximum accuracy = 57%; $P > 0.05$).

Discussion

This study aimed to investigate response to post-operative analgesic treatment with opioids and investigate factors associated with insufficient response. Severity of the pre-surgical chronic pain condition was a factor associated with post-surgical insufficiency of analgesic treatment, while QST measures were not related. In a first for clinical studies, it was possible to predict analgesic efficacy based on the pre-operative EEG recordings using machine learning, with similar accuracy to the chronic pain grade. In line with previous research, tonic painful EEG proved to be useful in contrast to resting EEG for prediction of opioid analgesia (Gram et al., 2015).

Methodological considerations

For a measurement to be of value within personalized medicine, reliability over longer time periods is a requirement (Bruton et al., 2000; Cannon et al., 2012). This study used spectral analysis and PLI of EEG during both rest and tonic cold pain, which are reliable, and therefore useful as a biomarker for prediction of analgesic efficacy (Gram et al., 2014; Hardmeier et al., 2014).

The only previous study on prediction of analgesic effect of opioids using EEG was performed on healthy volunteers using an experimental pain model (Gram et al., 2015). The current study is clinical and as such, suffers from confounders such as anxiety, and existing medication (Olesen et al., 2012). Firstly, patients were suffering from preoperative chronic pain, which influences EEG (Olesen et al., 2011; Sarnthein et al., 2006). Secondly, patients were already on analgesics before inclusion, which could also affect the EEG (Malver et al., 2014). Since pre-existing analgesic use could not be controlled for, this might have interfered with classification accuracy. On the other hand, since there was no difference in premedication analgesics between the groups and pre-operative opioid doses were low this is likely not of major importance. Lastly, the pain evoked in experimental pain models in our previous study is standardized between subjects, whereas the post-surgical pain is not, even when using a standardized operative procedure. Although the procedure is the same for all patients, there will always be variation in duration and complexity during surgery, which might affect postoperative pain. All these factors could explain why classification accuracy was lower in this clinical study.

It was not possible to distinguish between responders and non-responders based on the resting EEG which is in line with previous research in where the tonic painful EEG was found superior to resting EEG for prediction of analgesic efficacy (Gram et al., 2015). Thus, it seems that the predictive biomarkers in the EEG are only present when subjects experience pain. Previously, tonic pain was shown to result in increases in the delta band, with a simultaneous suppression of the alpha band. This was also seen in the current study, but to

a lesser degree than previously reported (Gram et al., 2014). The water temperature for the cold pressor test was higher than in previous studies (range=1-7 °C) (Mitchell et al., 2004). However, perceived pain during the cold pressor test (~6 NRS) was experienced as intense. It is therefore unlikely that water temperature alone accounts for low accuracy. However, future studies should attempt to use lower temperatures for the cold pressor test.

Within recent years, the gamma band of the EEG has received increasing attention within pain research. While large components of the EEG signal in evoked brain potentials have been shown to be closely related to the saliency of the stimulus rather than pain *per se* (Iannetti et al., 2008), the gamma band contains features specifically associated with pain perception (Tiemann et al., 2015; Zhang et al., 2012). However, most research on the gamma band was done on evoked brain potentials, where potentials contaminated by noise are easily rejected unlike in continuous EEG signals. This study utilized the cold pressor test for evoking pain, since tonic pain models have been shown to more closely mimic the unpleasantness of chronic pain (Rainville et al., 1992). The cold pressor test introduces electromyography artifacts into the gamma band, due to wincing facial expressions that accompanies prolonged and unpleasant tonic pain (Dowman et al., 2008). Taken together, despite its potential relevance for pain, the gamma band was excluded in this analysis to avoid the noise from muscle activity which could potentially also affect the results.

Patients were stratified into two groups based on a combined score of clinical pain ratings in order to account for problems associated with PCA, namely that both pain scores and analgesic consumption varies in a related way in the post-operative setting (Dai et al., 2013). Therefore, the identification of the optimal method of combining pain scores with analgesic consumption is a major focus point for research in analgesic studies (Dworkin et al., 2008; Turk et al., 2008). Attempts have been made to develop integrated scores which encompass both metrics (Silverman, O'Connor, & Brull, 1993), but no method has received widespread acceptance and their validity remains unknown (Grosen et al., 2013). The stratification based on clinical response from the QUIPS questionnaire resulted into a response rate of roughly 65% which is in line with this type of procedure (Maier et al., 2010). It could be argued that opioid consumption should have been incorporated into the response score. However, the fact that stratifying based on opioid consumption did not yield higher prediction accuracies indicates that the stratification method is not the main reason for the relatively low accuracy.

It should also be noted that the opioid medication in this study consisted of a combination of piritramide and oxycodone, which might affect individual patients differently (Drewes et al., 2013). Therefore, several sub-groups might exist within these two response groups, complicating stratification. However, it was not ethically feasible to alter the routine treatment in this study. To further investigate response or non-response to individual opioids, future studies could attempt to treat using only one opioid.

Prediction of post-operative analgesia

Clinical parameters relating to the pre-surgical pain state was associated with post-operative analgesia. Hence, a higher proportion of severe pre-surgical chronic pain grades were associated with poor post-operative analgesia. This corresponds well to the literature which consistently shows pre-operative pain is a strong predictor of acute post-operative pain (Gerbershagen et al., 2010; Pinto et al., 2015; Sommer et al., 2008). Since the responder score in this study is based on the post-operative pain ratings it follows that the pre-surgical pain would be related to this score.

Several studies have investigated prediction of analgesia based on QST and some results have been promising (Grosen et al., 2013). Heat and cold pain was included in this study as QST predictors, and neither proved effective for prediction. This could be due to central sensitization in the patients, for which opioids have limited effect (Olesen et al., 2013b).

A dysfunctional descending inhibition (as assessed by CPM paradigms) has previously been associated with development of post-operative pain in different clinical settings including pain after thoracic and gastrointestinal surgeries (Ip et al., 2009; Yarnitsky et al., 2008). Overall, the CPM effect in patients was weak in both responders and non-responders and this observation likely reflects a dysfunctional descending inhibition.

Group-wise statistical analysis of the EEG data revealed no differences between responders and non-responders, while machine learning was able to discriminate between responder groups at the individual patient level with an accuracy of 65%. This result is comparable to the predictive accuracy of chronic pain grade (63%), but is likely more robust as it is obtained with an objective method. The positive predictive value of EEG based classification was 76%. Hence, if patients had been treated according to the machine learning results based on preoperative EEG during cold pain, response rate to piritramide and oxycodone would have been 76% compared to the actual 65% response rate in this study. The remaining patients would have been treated with an alternative treatment, though only 53% were actually non-responders, meaning that there were a higher proportion of false negatives. However, in a personalized medicine scenario where the EEG and SVM model would be used to decide which patients to switch to secondary analgesic treatment the patients falsely classified as non-responders could still respond to the alternative treatment. Since this study included no alternative treatment the actual response rate in a personalized medicine scenario could not be determined.

The relative delta content from a frontally placed electrode (FP1) was selected as the most discriminate feature. This is comparable with previous research in healthy volunteers where frontal delta activity was

mainly selected by a machine learning approach (Gram, Graversen, Olesen, & Drewes, 2015). However, in this study only one feature was selected in comparison to the previous 7 features.

Conclusions

The novel use of EEG in combination with machine learning allowed for discrimination between responders and non-responders to postoperative analgesic opioid treatment. This shows EEG to be a potential important part in personalized pain medicine, with the potential to reduce suffering and persistent postoperative pain. The use of EEG in daily clinical work is becoming more and more feasible with new devices emerging, offering drastically reduced mounting times while retaining the quality of recordings (Sami et al., 2016). Future studies should work towards optimization of the methods together with inclusion of more variables in combination with the EEG.

Author Contributions

AM Drewes, M Przemeck, F Petzke, D Falla, M Reuster and J Erlenwein initiated and carried out the study. M Reuster and J Erlenwein performed the preoperative assessments. M Gram designed and carried out the EEG analysis. M Gram carried out the machine learning analysis. M Gram, SS Olesen and AM Drewes evaluated results and drafted the manuscript. All authors were involved in reading and approving the final manuscript.

References

Ahmedzai, S.H. (2013). Personalized medicine--one size fits one: tailoring pain therapy to individuals' needs. *J Pain Palliat Care Pharmacother* 27, 83–85.

Akin, M. (2002). Comparison of wavelet transform and FFT methods in the analysis of EEG signals. *J Med Syst* 26, 241–247.

Bellamy, N. (2005). The WOMAC knee and hip osteoarthritis indices: Development, validation, globafization and influence on the development of the AUSCAN hand osteoarthritis indices. *Clin Exp Rheumatol* 23, S148–S153.

Brown, G., Pocock, A., Zhao, M.-J., Luján, M. (2012). Conditional Likelihood Maximisation: A Unifying Framework for Information Theoretic Feature Selection. *J Mach Learn Res* 13, 27–66.

Bruehl, S., Apkarian, a V., Ballantyne, J.C., Berger, A., Borsook, D., Chen, W.G., Farrar, J.T., Haythornthwaite, J. a, Horn, S.D., Iadarola, M.J., Inturrisi, C.E., Lao, L., Mackey, S., Mao, J., Sawczuk, A., Uhl, G.R., Witter, J., Woolf, C.J., Zubieta, J.-K., Lin, Y. (2013). Personalized medicine and opioid analgesic prescribing for chronic pain: opportunities and challenges. *J Pain* 14, 103–113.

Bruton, A., Conway, J.H., Holgate, S.T. (2000). Reliability: What is it, and how is it measured? *Physiotherapy* 86, 94–99.

Cannon, R.L., Baldwin, D.R., Shaw, T.L., Diloreto, D.J., Phillips, S.M., Scruggs, A.M., Riehl, T.C. (2012). Reliability of quantitative EEG (qEEG) measures and LORETA current source density at 30 days. *Neurosci Lett* 518, 27–31.

Chang, C.-C., Lin, C.-J. (2011). LIBSVM. *ACM Trans Intell Syst Technol* 2, 1–27.

Cortes, C., Vapnik, V. (1995). Support-Vector Networks. *Mach Learn* 297, 273–297.

Dai, F., Silverman, D.G., Chelly, J.E., Li, J., Belfer, I., Qin, L. (2013). Integration of pain score and morphine consumption in analgesic clinical studies. *J Pain* 14, 767–777.

Dolin, S.J., Cashman, J.N., Bland, J.M. (2002). Effectiveness of acute postoperative pain management: I. Evidence from published data. *Br J Anaesth* 89, 409–423.

Dowman, R., Rissacher, D., Schuckers, S. (2008). EEG indices of tonic pain-related activity in the somatosensory cortices. *Clin Neurophysiol* 119, 1201–1212.

Drewes, A.M., Jensen, R.D., Nielsen, L.M., Droney, J., Christrup, L.L., Arendt-Nielsen, L., Riley, J., Dahan, A. (2013). Differences between opioids: Pharmacological, experimental, clinical and economical perspectives. *Br J Clin Pharmacol* 75, 60–78.

Dworkin, R.H., Turk, D.C., Wyrwich, K.W., Beaton, D., Cleeland, C.S., Farrar, J.T., Haythornthwaite, J. a, Jensen, M.P., Kerns, R.D., Ader, D.N., Brandenburg, N., Burke, L.B., Cella, D., Chandler, J., Cowan, P., Dimitrova, R., Dionne, R., Hertz, S., Jadad, A.R., Katz, N.P., Kehlet, H., Kramer, L.D., Manning, D.C., McCormick, C., McDermott, M.P., McQuay, H.J., Patel, S., Porter, L., Quessy, S., Rappaport, B. a, Rauschkolb, C., Revicki, D. a, Rothman, M., Schmader, K.E., Stacey, B.R., Stauffer, J.W., von Stein, T., White, R.E., Witter, J., Zavisic, S. (2008). Interpreting the clinical importance of treatment outcomes in chronic pain clinical trials: IMMPACT recommendations. *J Pain* 9, 105–121.

Erlenwein, J., Stüder, D., Lange, J.-P., Bauer, M., Petzke, F., Przemeck, M. (2012). Prozessoptimierung durch zentrale Steuerung der Akutschmerztherapie. *Anaesthesist* 61, 971–983.

Frettlöh, J., Maier, C., Gockel, H., Hüppe, M. (2003). Validität des Mainzer Stadienmodells der Schmerzchronifizierung bei unterschiedlichen Schmerzdiagnosen. *Der Schmerz* 17, 240–251.

Gerbershagen, H.J., Dagtekin, O., Gaertner, J., Petzke, F., Heidenreich, A., Sabatowski, R., Ozgur, E. (2010). Preoperative chronic pain in radical prostatectomy patients: preliminary evidence for enhanced susceptibility to surgically induced pain. *Eur J Anaesthesiol* 27, 448–454.

Gerbershagen, H.J., Özgür, E., Dagtekin, O., Straub, K., Hahn, M., Heidenreich, A., Sabatowski, R., Petzke, F. (2009). Preoperative pain as a risk factor for chronic post-surgical pain - Six month follow-up after radical prostatectomy. *Eur J Pain* 13, 1054–1061.

Gong, Q., Wu, Q., Scarpazza, C., Lui, S., Jia, Z., Marquand, A., Huang, X., McGuire, P., Mechelli, A. (2011). Prognostic prediction of therapeutic response in depression using high-field MR imaging. *Neuroimage* 55, 1497–1503.

Gram, M., Graversen, C., Olesen, A.E., Drewes, A.M. (2015). Machine learning on encephalographic activity may predict opioid analgesia. *Eur J Pain* 19, 1552–1561.

Gram, M., Graversen, C., Olesen, S.S., Drewes, A.M. (2014). Dynamic spectral indices of the electroencephalogram provide new insights into tonic pain. *Clin Neurophysiol* 126, 763–771.

Graversen, C., Brock, C., Drewes, A.M., Farina, D. (2011). Biomarkers for visceral hypersensitivity identified by classification of electroencephalographic frequency alterations. *J Neural Eng* 8, 056014.

Graversen, C., Olesen, S.S., Olesen, A.E., Steimle, K., Farina, D., Wilder-Smith, O.H.G., Bouwense, S. a W., van Goor, H., Drewes, A.M. (2012). The analgesic effect of pregabalin in patients with chronic pain is reflected by changes in pharmaco-EEG spectral indices. *Br J Clin Pharmacol* 73, 363–372.

Grosen, K., Fischer, I.W.D., Olesen, a E., Drewes, a M. (2013). Can quantitative sensory testing predict responses to analgesic treatment? *Eur J Pain* 17, 1267–1280.

Hardmeier, M., Hatz, F., Bousleiman, H., Schindler, C., Stam, C.J., Fuhr, P. (2014). Reproducibility of Functional Connectivity and Graph Measures Based on the Phase Lag Index (PLI) and Weighted Phase Lag Index (wPLI) Derived from High Resolution EEG. *PLoS One* 9, e108648.

Iannetti, G.D., Hughes, N.P., Lee, M.C., Mouraux, a (2008). Determinants of laser-evoked EEG responses: pain perception or stimulus saliency? *J Neurophysiol* 100, 815–828.

Ip, H.Y.V., Abrishami, A., Peng, P.W.H., Wong, J., Chung, F. (2009). Predictors of postoperative pain and analgesic consumption: a qualitative systematic review. *Anesthesiology* 111, 657–677.

Kehlet, H., Jensen, T.S., Woolf, C.J. (2006). Persistent postsurgical pain: risk factors and prevention. *Lancet* 367, 1618–1625.

Von Korff, M., Ormel, J., Keefe, F.J., Dworkin, S.F. (1992). Grading the severity of chronic pain. *Pain* 50, 133–149.

Lehembre, R., Bruno, M.A., Vanhaudenhuyse, A., Chatelle, C., Cologan, V., Leclercq, Y., Soddu, A., Macq, B., Laureys, S., Noirhomme, Q. (2012). Resting-state EEG study of comatose patients: A connectivity and frequency analysis to find differences between vegetative and minimally conscious states. *Funct Neurol* 27, 41–47.

Liu, S.S., Wu, C.L. (2007). The effect of analgesic technique on postoperative patient-reported outcomes including analgesia: A systematic review. *Anesth Analg* 105, 789–808.

Maier, C., Nestler, N., Richter, H., Hardinghaus, W., Pogatzki-Zahn, E., Zenz, M., Osterbrink, J. (2010).

The quality of pain management in German hospitals. *Dtsch Arztebl Int* 107, 607–614.

Malver, L.P., Brokjaer, A., Staahl, C., Graversen, C., Andresen, T., Drewes, A.M. (2014). Electroencephalography and analgesics. *Br J Clin Pharmacol* 77, 72–95.

Masters Steedman, S., Middaugh, S.J., Kee, W.G., Carson, D.S., Harden, R.N., Miller, M.C. (1992). Chronic-pain medications: equivalence levels and method of quantifying usage. *Clin J Pain* 8, 204–214.

Meissner, W., Mescha, S., Rothaug, J., Zwacka, S., Goettermann, A., Ulrich, K., Schleppers, A. (2008). Quality improvement in postoperative pain management: results from the QUIPS project. *Dtsch Arztebl Int* 105, 865–870.

Meissner, W., Ullrich, K., Zwacka, S. (2006). Benchmarking as a tool of continuous quality improvement in postoperative pain management. *Eur J Anaesthesiol* 23, 142–148.

Mitchell, L.A., MacDonald, R.A.R., Brodie, E.E. (2004). Temperature and the cold pressor test. *J Pain* 5, 233–237.

Nir, R.-R., Sinai, A., Moont, R., Harari, E., Yarnitsky, D. (2012). Tonic pain and continuous EEG: prediction of subjective pain perception by alpha-1 power during stimulation and at rest. *Clin Neurophysiol* 123, 605–612.

Nolte, G., Ziehe, A., Nikulin, V., Schlögl, A., Krämer, N., Brismar, T., Müller, K.-R. (2008). Robustly Estimating the Flow Direction of Information in Complex Physical Systems. *Phys Rev Lett* 100, 234101.

Olesen, A., Andresen, T., Staahl, C., Drewes, A. (2012). Human Experimental Pain Models for Assessing the Therapeutic Efficacy of Analgesic Drugs. *Pharmacol Rev* 64, 722–779.

Olesen, S.S., Graversen, C., Bouwense, S. a W., van Goor, H., Wilder-Smith, O.H.G., Drewes, A.M. (2013a). Quantitative sensory testing predicts pregabalin efficacy in painful chronic pancreatitis. *PLoS One* 8, e57963.

Olesen, S.S., Hansen, T.M., Graversen, C., Steimle, K., Wilder-Smith, O.H.G., Drewes, A.M. (2011). Slowed EEG rhythmicity in patients with chronic pancreatitis: evidence of abnormal cerebral pain processing? *Eur J Gastroenterol Hepatol* 23, 418–424.

Olesen, S.S., Juel, J., Graversen, C., Kolesnikov, Y., Wilder-Smith, O.H.G., Drewes, A.M. (2013b). Pharmacological pain management in chronic pancreatitis. *World J Gastroenterol* 19, 7292–7301.

Pinto, P.R., McIntyre, T., Araújo-Soares, V., Costa, P., Almeida, A. (2015). Differential Predictors of Acute Post-Surgical Pain Intensity After Abdominal Hysterectomy and Major Joint Arthroplasty. *Ann Behav Med* 49, 384–397.

Pud, D., Granovsky, Y., Yarnitsky, D. (2009). The methodology of experimentally induced diffuse noxious inhibitory control (DNIC)-like effect in humans. *Pain* 144, 16–19.

Rainville, P., Feine, J.S., Bushnell, M.C., Duncan, G.H. (1992). A psychophysical comparison of sensory and affective responses to four modalities of experimental pain. *Somatosens Mot Res* 9, 265–277.

Rothaug, J., Weiss, T., Meissner, W. (2013). How simple can it get? Measuring pain with NRS items or binary items. *Clin J Pain* 29, 224–232.

Sami, S., Mariella, C., Valeria, D.C., Daniele, A., Giuseppe, S., Michele, D.R., Paolo, A., Piero, A., Sara, M. (2016). A low-cost, user-friendly EEG recording system for the assessment of hepatic encephalopathy. *Hepatology* 00, n/a – n/a.

Sarnthein, J., Stern, J., Aufenberg, C., Rousson, V., Jeanmonod, D. (2006). Increased EEG power and slowed dominant frequency in patients with neurogenic pain. *Brain* 129, 55–64.

Schmitt, N., Gerbershagen, H.U. (1990). The mainz pain staging system (MPSS) for chronic pain. *Pain* 41, 484.

Schulz, E., Zherdin, A., Tiemann, L., Plant, C., Ploner, M. (2011). Decoding an Individual's Sensitivity to Pain from the Multivariate Analysis of EEG Data. *Cereb Cortex* 22, 1118–1123.

Silverman, D.G., O'Connor, T.Z., Brull, S.J. (1993). Integrated assessment of pain scores and rescue morphine use during studies of analgesic efficacy. *Anesth Analg* 77, 168–170.

Sommer, M., de Rijke, J.M., van Kleef, M., Kessels, a G.H., Peters, M.L., Geurts, J.W.J.M., Gramke, H.-F., Marcus, M. a E. (2008). The prevalence of postoperative pain in a sample of 1490 surgical inpatients. *Eur J Anaesthesiol* 25, 267–274.

Stormo, K.J., Kee, W.G., Steedham, S., Middaugh, S. (1998). Medication quantification scores and evaluation of patient pharmacology profiles. *Curr Rev Pain* 2, 171–174.

Tiemann, L., May, E.S., Postorino, M., Schulz, E., Nickel, M.M., Bingel, U., Ploner, M. (2015). Differential neurophysiological correlates of bottom-up and top-down modulations of pain. *Pain* 156, 289–296.

Turk, D.C., Dworkin, R.H., McDermott, M.P., Bellamy, N., Burke, L.B., Chandler, J.M., Cleeland, C.S., Cowan, P., Dimitrova, R., Farrar, J.T., Hertz, S., Heyse, J.F., Iyengar, S., Jadad, A.R., Jay, G.W., Jermano, J. a, Katz, N.P., Manning, D.C., Martin, S., Max, M.B., McGrath, P., McQuay, H.J., Quessy, S., Rappaport, B. a, Revicki, D. a, Rothman, M., Stauffer, J.W., Svensson, O., White, R.E., Witter, J. (2008). Analyzing multiple endpoints in clinical trials of pain treatments: IMMPACT recommendations. Initiative on Methods, Measurement, and Pain Assessment in Clinical Trials. *Pain* 139, 485–493.

VanDenKerkhof, E.G., Hopman, W.M., Goldstein, D.H., Wilson, R. a, Towheed, T.E., Lam, M., Harrison, M.B., Reitsma, M.L., Johnston, S.L., Medd, J.D., Gilron, I. (2012). Impact of perioperative pain intensity, pain qualities, and opioid use on chronic pain after surgery: a prospective cohort study. *Reg Anesth Pain Med* 37, 19–27.

Yarnitsky, D., Crispel, Y., Eisenberg, E., Granovsky, Y., Ben-Nun, A., Sprecher, E., Best, L.-A., Granot, M. (2008). Prediction of chronic post-operative pain: Pre-operative DNIC testing identifies patients at risk. *Pain* 138, 22–28.

Zaslansky, R., Rothaug, J., Chapman, C.R., B??ckstr??m, R., Brill, S., Fletcher, D., Fodor, L., Gordon, D.B., Komann, M., Konrad, C., Leykin, Y., Pogatski-Zahn, E., Puig, M.M., Rawal, N., Ullrich, K., Volk, T., Meissner, W. (2015). PAIN OUT: The making of an international acute pain registry. *Eur J Pain (United Kingdom)* 19, 490–502.

Zaslansky, R., Rothaug, J., Chapman, R.C., Backström, R., Brill, S., Engel, C., Fletcher, D., Fodor, L., Funk, P., Gordon, D., Komann, M., Konrad, C., Kopf, A., Leykin, Y., Pogatzki-Zahn, E., Puig, M., Rawal, N., Schwenkglenks, M., Taylor, R.S., Ullrich, K., Volk, T., Yahiaoui-Doktor, M., Meissner, W. (2014). PAIN OUT: An international acute pain registry supporting clinicians in decision making and in quality improvement activities. *J Eval Clin Pract* 20, 1090–1098.

Zhang, Z.G., Hu, L., Hung, Y.S., Mouraux, A., Iannetti, G.D. (2012). Gamma-Band Oscillations in the Primary Somatosensory Cortex--A Direct and Obligatory Correlate of Subjective Pain Intensity. *J Neurosci* 32, 7429–7438.

Tables

		Patients, n (% of total cohort)
Age (years)	64.5 ± 12.5	81 (100 %)
Male sex, n (%)	36 (44 %)	81 (100 %)
BMI (kg/m^2)	28.1 ± 5.0	79 (98 %)
WOMAC	54.6 ± 18.3	65 (80 %)
Hip-pain duration, n (%)		81 (100 %)
- 0 – 6 months	12 (15 %)	
- 6 – 12 months	11 (14 %)	
- 1 – 2 years	19 (23 %)	
- 2 – 5 years	20 (25 %)	
- More than 5 years	19 (23 %)	
Chronic pain grade, n (%)		78 (96 %)
- Grade I-II	32 (41 %)	
- Grade III-IV	46 (59 %)	
MPSS, n (%)		81 (100 %)
- Stage I	34 (42 %)	
- Stage II	26 (32 %)	
- Stage III	21 (26 %)	
Preoperative MQS - non-opioids	5.1 ± 4.5	79 (98 %)
Preoperative MQS - opioids	0.6 ± 1.6	81 (100 %)

WOMAC = Western Ontario and McMaster Universities Osteoarthritis Index; MPSS = Mainz Pain Staging System; MQS = Medication Quantification Scale.

Table 2: Baseline parameters for the included patients, divided into groups of responders (N = 51) and non-responders (N = 30) to post-operative analgesic treatment with oxycodone and piritramide.

	All patients (N=81)	Responders (N = 51)	Non-responders (N = 30)	OR (95% CI)	P-value
Age (years)		64.2 ± 10.4	64.9 ± 15.7	1.05 (0.73-1.51)	0.81
Male sex, n (%)		25 (49%)	11 (37%)	0.60 (0.24-1.52)	0.28
BMI (kg/m^2)		28.1 ± 4.6	28.1 ± 5.7	1.00 (0.91-1.09)	0.98
WOMAC		51.6 ± 18.9	60.0 ± 16.2	1.03 (1.00-1.06)	0.082
Hip-pain duration, n (%)					
- 0 – 6 months		7 (14%)	5 (17%)	1.00	
- 6 – 12 months		8 (16%)	3 (10%)	0.53 (0.10-3.03)	0.47
- 1 – 2 years		12 (24%)	7 (23%)	0.82 (0.19-3.58)	0.79
- 2 – 5 years		13 (25%)	7 (23%)	0.75 (0.17-3.28)	0.71
- More than 5 years		11 (22%)	8 (27%)	1.02 (0.24-4.41)	0.98
Chronic pain grade, n (%)					
- Grade I-II		26(53%)	6 (21%)	1.00	
- Grade III-IV		23 (47%)	23 (79%)	4.33 (1.50-12.50)	0.007 [a]
MPSS, n (%)					
- Stage I		25 (49%)	9 (30%)	1.00	
- Stage II		18 (35%)	8 (27%)	1.23 (0.40-3.82)	0.71
- Stage III		8 (16%)	13 (43%)	4.51 (1.41-14.46)	0.011 [a]
Pain on hip rotation (NRS)		5.2 ± 2.0	6.2 ± 1.9	1.33 (1.02-1.72)	0.034 [a]
Preoperative MQS - non-opioids		4.6 ± 4.6	5.9 ± 4.3	1.06 (0.96-1.18)	0.24
Preoperative MQS - opioids		0.4 ± 1.3	0.8 ± 1.9	1.14 (0.86-1.52)	0.36
Post-surgery patient controlled analgesia - opioid		114.5 ± 41.2	123.1 ± 47.5	1.00 (0.99-1.02)	0.39

equivalents (mg)

Cold pressor pain (NRS)	6.1 ± 2.3	6.9 ± 2.5	1.16 (0.94-1.42)	0.17
Heat pain threshold (°C)	47.0 ± 2.9	46.3 ± 3.7	0.93 (0.81-1.07)	0.34
Pressure pain threshold (kPa)	71.7 ± 20.4	68.0 ± 22.2	1.01 (0.99-1.03)	0.45
Conditioned pain modulation (%)	2.8 ± 4.3	2.8 ± 4.2	1.00 (0.90-1.12)	0.96

[a] $P < 0.05$

BMI = Body Mass Index; WOMAC = Western Ontario and McMaster Universities Osteoarthritis Index; MPSS = Mainz pain staging system; NRS = Numerical Rating Scale; OE = Opioid equivalent units; MQS = Medication Quantification Score.

Table 3: Multivariate analysis of risk factors associated with inadequate post-operative analgesia identified using logistic regression. Internal validation performed with 1000 bootstrap samples.

	Multivariate analysis		Internal validation	
	OR (95% CI)	P-value	OR (95% CI)	P-value
Chronic pain grade, n (%)				
- Grade I-II	1.00			
- Grade III-IV	3.74 (1.21-11.53)	0.022 [a]	3.74 (1.07-13.14)	0.04 [a]
MPSS, n (%)				
- Stage I	1.00			
- Stage II	1.14 (0.34-3.83)	0.83		
- Stage III	1.99 (0.54-7.30)	0.30		
Pain on hip rotation (NRS)	1.26 (0.95-1.68)	0.11		

[a] $P < 0.05$

MPSS = Mainz Pain Staging System; NRS = Numerical Rating Scale

Figure legends

Figure 1: Experimental setup and study procedures. PCA = Patient-Controlled Anaesthesia; QST = Quantitative Sensory Testing.

Figure 2: Topographical illustration of the spectral indices calculated from the resting EEG in the four frequency bands; delta (1-4 Hz), theta (4-8 Hz), alpha (8-12 Hz) and beta (12-32 Hz). Activity is color coded with higher levels of activity represented in red and yellow colors, while lower activity is represented by blue. Responders to opioid treatment appear to have slightly decreased alpha activity, and increased beta activity.

Figure 3: Topographical illustration of the phase-lag index calculated from the resting EEG in the four frequency bands. The color code shows red and yellow for electrodes that have stronger average connections to all other electrodes, while blue colors represent electrodes, which are poorly connected. Responders to opioid treatment appear to exhibit decreased connectivity of the alpha band compared to non-responders.

Figure 4: Topographical illustration of the spectral indices calculated from EEG during cold pain, in the four frequency bands. Activity is color coded with higher levels of activity represented in red and yellow colors, while lower activity is represented by blue. Increased delta activity appears to be present for responders to opioid treatment.

Figure 5: Topographical illustration of the phase-lag index calculated from the EEG during cold pain, in the four frequency bands. The color code shows red and yellow for electrodes that have stronger average connections to all other electrodes, while blue colors represent electrodes, which are poorly connected. Responders to opioid treatment seems to present with increased connectivity in the delta and theta bands compared to non-responders.