

2 **Evolving, dynamic clustering of spatio/spectro-temporal data**
3 **in 3D spiking neural network models and a case study on EEG**
4 **data**

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6 Received: 12 June 2016 / Accepted: 14 February 2017
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8 **Abstract** Clustering is a fundamental data processing
9 technique. While clustering of static (vector based) data and
10 of fixed window size time series have been well explored,
11 dynamic clustering of spatiotemporal data has been little
12 researched if at all. Especially when patterns of changes
13 (events) in the data across space and time have to be cap-
14 tured and understood. The paper presents novel methods
15 for clustering of spatiotemporal data using the NeuCube
16 spiking neural network (SNN) architecture. Clusters of
17 spatiotemporal data were created and modified on-line in
18 a continuous, incremental way, where spatiotemporal rela-
19 tionships of changes in variables are incrementally learned
20 in a 3D SNN model and the model connectivity and spik-
21 ing activity are incrementally clustered. Two clustering
22 methods were proposed for SNN, one performed during
23 unsupervised and one—during supervised learning mod-
24 els. Before submitted to the models, the data is encoded as
25 spike trains, a spike representing a change in the variable
26 value (an event). During the unsupervised learning, the
27 cluster centres were predefined by the spatial locations of
28 the input data variables in a 3D SNN model. Then clusters
29 are evolving during the learning, *i.e.* they are adapted con-
30 tinuously over time reflecting the dynamics of the changes
31 in the data. In the supervised learning, clusters represent
32 the dynamic sequence of neuron spiking activities in a
33 trained SNN model, specific for a particular class of data
34 or for an individual instance. We illustrate the proposed
35 clustering method on a real case study of spatiotemporal

EEG data, recorded from three groups of subjects during a
cognitive task. The clusters were referred back to the brain
data for a better understanding of the data and the processes
that generated it. The cluster analysis allowed to discover
and understand differences on temporal sequences and spa-
tial involvement of brain regions in response to a cognitive
task.

Keywords Dynamic spatiotemporal streaming data
clustering · EEG data · NeuCube · Spiking neural
networks · Unsupervised learning · Supervised learning ·
Personalised clustering

1 **Introduction**

Clustering aims at objectively organise data samples into
homogenised groups, where the data samples within a
group are similar. So far many clustering methods have
been developed to identify structures in different data
types, such as static, temporal, etc. Data is static when the
feature values do not change over time, and it is time series
(temporal) if the features change their values over a contin-
uous time. With respect to different data types, clustering
methods differ significantly in the notion of the similarity
or distance measures.

Clustering can be classified according to several criteria:

- i. Clustering of raw data *versus* clustering of pre-pro-
cessed data (such as encoded into spikes, as the case in
the paper is);
- ii. Clustering of absolute values of variables *versus* clus-
tering of changes in the variables (also called events, as
the case in this paper is);

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- 65 iii. Direct clustering of data *versus* clustering of a model
66 that is being created to learn and capture the essential
67 characteristics of interest from the data (as the case in
68 this paper is);
69 iv. Clustering of vector based, static data *versus* clustering
70 of dynamic, streaming data (as the case in the paper is).

71 We can distinguish the following classes of clustering
72 methods for static, vector based data:

- 73 a) Clustering of static, vector based data, where the
74 number of clusters is pre-defined (such as C-means,
75 K-Mean clustering methods);
76 b) Clustering of time series data, where the number of the
77 time points are fixed and the time series samples are
78 represented as vectors, applying the above methods.
79 c) Evolving clustering methods, where a stream of vec-
80 tors are clustered incrementally without pre-defining
81 the number of clusters such as DENFIS (Kasabov
82 and Song 2002) and Evolving Fuzzy Neural Network
83 (EFuNN) (Kasabov 2001);

84 In this paper, we propose clustering methods for whole
85 spatiotemporal data samples from a data stream. In prin-
86 ciple, these methods can be classified into the following
87 groups:

- 88 d) Two tier clustering of the data: first according to the
89 spatial information, and then according to the temporal
90 information;
91 e) Integrated clustering of both spatial and temporal
92 information;
93 f) In case of both static and spatiotemporal data are avail-
94 able, clustering of the static information first, and
95 then—the spatiotemporal information. This can be
96 applied for personalised modelling (Gholami Doborjeh
97 and Kasabov 2016).
98 g) Integrated clustering of both static and spatiotemporal
99 data for personalised modelling.

100 The paper presents two methods in the groups (d) and (e)
101 from above that are also defined in the categories (i)–(iv).

102 1.1 Overview on static data clustering, temporal data 103 clustering and evolving clustering methods

104 Clustering methods for various static data are classified
105 into five major categories: hierarchical methods (Johnson
106 1967); partitioning methods; density-based methods (Ester
107 et al. 1996); grid-based methods; and model-based meth-
108 ods. Partitioning methods construct k partitions of data,
109 where each partition is represented by a similarity metric
110 of the objects in the partition. A partition is hard if each

object belongs to only one cluster [k-means (Hartigan and
Wong 1979)], or fuzzy if one object is allowed to be in
more than one cluster by different degree [fuzzy c-means
(Bezdek et al. 1984)]. A hierarchical clustering represents
groups of data objects using a tree of clusters, where the
similarity is measured using a pairwise distance matrix
of the objects. In density-based clustering, a cluster is
continuously growing as long as the density in the neigh-
bourhood surpasses a threshold. In grid-based clustering,
the object space is quantized into a finite number of cells,
where the operations for clustering are performed. A com-
mon example of the grid-based approach is STING (Wang
et al. 1997). Model-based clustering undertakes a model for
each cluster and aims at best fit of data to the model. One
major method of model-based clustering is neural network
approach. Two prominent clustering methods of the neural
network are competitive learning, including ART (Carpenter
and Grossberg 1987) and self-organizing maps (SOM)
(Kohonen 1998).

Massive amount of temporal data (time-series data) has
been recorded so far in various areas, such as electronic,
video/audio, biologic, neurology, etc. In case of cluster-
ing of such data, given a set of individual time series val-
ues, the objective is to group similar patterns into the same
cluster. This task demands a measure notion to estimate
the level of similarity between time series. However, the
known Euclidean distance and other typical measures used
for non-temporal data are unsuitable metrics to evaluate the
similarity between time series, because they are unable to
deal with temporal interaction between time series data fea-
tures. Various temporal data clustering methods have been
introduced so far, such as biclustering (Mirkin 1998) that
has been used for clustering the time series gene expres-
sion data (Tanay et al. 2002). Any data that can be repre-
sented as a matrix is amenable to biclustering. Biclustering
methods simultaneously clusters both rows and columns of
a matrix.

Evolving clustering methods represent incremental
growth of clusters and creation of new clusters from a
stream of vector based data. While SOM assigns similar
input vectors into topologically close neurons, evolving
Self-organizing maps (ESOM) (Deng and Kasabov 2000)
and the DENFIS evolving clustering method (Kasabov and
Song 2002) were introduced for online unsupervised data
clustering. EFuNN (Kasabov 2001) was introduced for
evolving supervised clustering related to classification or
regression. In (Katwal et al. 2013), a graph-based visualisa-
tions of SOM has been used for clustering fMRI data. In
another research (Liao et al. 2008), an integrated SOM and
hierarchical clustering architecture was designed to detect
activation patterns of fMRI data. While using these meth-
ods for clustering of time series data, the temporal com-
ponents of each sample data variables are transposed into

164 feature vectors, where the time is hidden and no temporal
165 interaction of time series within a sample can be learned
166 anymore.

167 A method of spatiotemporal clustering has been pro-
168 posed in (Deng et al. 2013) for clustering input variables of
169 meteorological data. In this research the input variables can
170 be clustered if they are within a certain neighbourhood and
171 their dynamics are auto-correlated.

172 All the above methods deal with vector based data and
173 do not reveal any spatiotemporal information related to the
174 processes that generated the data. This is a significant con-
175 trast to the clustering methods that we propose in this paper
176 as they deal with whole spatiotemporal patterns of data.

177 Unlike the above methods that aimed to directly cluster
178 raw data, we propose to dynamically cluster not the raw
179 data, but the changes in the data that are dynamically cap-
180 tured in a brain-like evolving SNN computational model.
181 The model is dynamically evolving from the input spatio-
182 temporal data after the data is encoded into spikes repre-
183 senting changes in the data, and the clusters in the model
184 are evolving too. These clusters represent: (1) dynamic
185 measures of spatiotemporal similarity in the input variables
186 in respect to their changing values in time; (2) sequentially
187 activated areas of the SNN model that capture changes in
188 the data over time.

189 1.2 Spatiotemporal data clustering

190 Spatiotemporal clustering is a procedure of grouping data
191 samples based on their spatial and temporal similarity.
192 Learning dynamic patterns of spatiotemporal data is a chal-
193 lenging task, as temporal features may manifest complex
194 interaction that may also change dynamically over time.
195 Developing new clustering methods that can capture these
196 “hidden” interactions and interrelationships among multi-
197 variate data, is of crucial importance not only for the spatio-
198 temporal data analysis and data understanding, but also
199 for future events prediction based on captured complex spatio-
200 temporal patterns from Spatio-Spectro-Temporal Data
201 (SSTD). A variety of techniques have been developed for
202 SSTD recordings, such as EEG, fMRI, DTI, etc. (Nieder-
203 meyer and da Silva 2005; Ogawa et al. 1992).

204 SNN methods have been developed for SSTD learning,
205 some of them are implemented in neuromorphic hardware
206 systems (Indiveri et al. 2011). The challenge now for infor-
207 mation science and artificial intelligence is to develop new
208 computational methods that utilize SNN and neuromorphic
209 hardware systems for efficient processing of SSTD, includ-
210 ing clustering of such data.

211 NeuCube (Kasabov 2014) is a generic evolving Spatio-
212 Temporal Data Machine (STDM) based on SNN for learn-
213 ing, classification/regression, visualisation and interpreta-
214 tion of spatiotemporal data. It is able to capture both *time*

and *space* features of SSTD in a SNN architecture for the
215 sake of understanding the data. When compared with tradi-
216 tional statistical analysis methods and methods of artificial
217 intelligence when dealing with SSTD, NeuCube models
218 resulted in significantly higher accuracy of classification
219 results, faster data processing, and a better visualisation and
220 interpretation of the SSTD. This is due to the ability of a
221 NeuCube model to learn and capture spatiotemporal inter-
222 action between the data variables. So far NeuCube has been
223 successfully used for EEG data modelling, learning, and
224 classification (Gholami Dobarjeh et al. 2016; Kasabov and
225 Capecci 2015; Capecci et al. 2015; Schliebs et al. 2013).

226 This paper now contributes to the NeuCube computa-
227 tional framework with dynamic clustering of spatiotempo-
228 ral connectivity and spiking activity of the spiking neurons
229 in a NeuCube model, while it is learning from streaming
230 data.

231 This paper introduces new methods for dynamic, evolu-
232 ting clustering of spatiotemporal streaming data through
233 encoding the data into spikes, creating dynamic, evolving
234 clusters of spiking neurons in a 3D SNN reservoir, such
235 as the NeuCube system, both in space and time. The main
236 purposes of the introduced methods, that have not been
237 achieved so far, are the following:
238

- 239 1. To detect similar spatiotemporal patterns of changes in
240 the streaming data, which are *dynamically* generated
241 with respect to the interaction between input variables.
242 The dynamic visualisation of the clusters captures the
243 time in which a cluster is created and it demonstrates
244 how this cluster is changed over time. It enables us for
245 the first time to study the dynamics of such clusters.
- 246 2. To understand hidden spatiotemporal patterns of
247 changes in the data by pursuing the trend of the clus-
248 ter creation. For the case study shown in the paper, this
249 relates to brain activities.

250 The clustering methods are applied to a case study of
251 EEG data, but can be used for other spatiotemporal data,
252 such as fMRI (Gholami Dobarjeh and Kasabov 2015), seis-
253 mic and other data (Kasabov et al. 2016).

254 This paper is structured as follows: Sect. 2 describes the
255 NeuCube architecture for SSTD analysis (Kasabov 2014).
256 Section 3 introduces the proposed methods for cluster-
257 ing of SSTD in an unsupervised and supervised learning
258 modes. Section 4 describes the case study data. Section 5
259 describes the details of the proposed dynamic unsupervised
260 clustering methods and illustrates them on the case study
261 data. Section 6 describes the details of the personalised
262 clustering methods at supervised learning stage and illus-
263 trates them on the case study data. Section 7 represents a
264 comparison of the proposed clustering methods with SOM
265 clustering. Section 8 discusses results and future directions.

2 The NeuCube spiking neural network architecture for SSTD

Spiking Neural Network (SNN) models can learn both space and time components from data and are considered as suitable models to process SSTD (Maass et al. 2002). The NeuCube-based SNN architecture (shown graphically in Fig. 1) consists of: input data encoding module; that encodes multivariable continuous temporal stream data into spike trains; a 3D recurrent SNN cube (SNNcube), where input data are mapped and learned in an unsupervised mode; and an SNN classifier that learns in a supervised mode to classify the spatiotemporal patterns of the SNNcube activities which represent patterns from the input data (Kasabov 2007, 2012, 2014; Kasabov et al. 2013).

2.1 Input data encoding, mapping, and model initialisation

2.1.1 Input spatiotemporal data encoding

In one of the implementations of the NeuCube encoding module, a Threshold-Based Representation method (TBR) (Delbruck 2007) is applied to the SSTD to produce spike trains. Generated spikes represent the form of the SSTD wave signal in terms of amplitude changes. Once a signal change exceeds the threshold TBR_{thr} , one spike occurs. Therefore, in the case of brain data, the spike trains

represent real brain activity patterns that will be used to train a SNNcube.

2.1.2 SSTD mapping to the SNNcube

A 3D brain-like SNN cube is created to map a relevant brain data template, such as Talairach (Talairach and Tournoux 1988), MNI (Lancaster et al. 2007), etc. The size of the SNNcube is controlled by three parameters: n_x, n_y, n_z representing the number of spiking neurons along x, y and z coordinates. Every neuron in this cube is a computational unit that is implemented based on the Leaky-Integrate and Fire Model (LIFM) of a spiking neuron as one implementation (Abbott 1999). Input neurons are allocated to the input SSTD variables for transferring their spike trains to the SNNcube. In order to preserve the spatial information of the SSTD, each neuron in the SNNcube represents a spatial location from the brain template, therefore—an area of the brain. Also each allocated input neuron in the SNNcube has the same (x, y, z) coordinates as the corresponding input data variable in the used brain template.

2.1.3 SNNcube initialisation

The SNNcube is initialised with the use of the “small world” connectivity. Each neuron in the SNNcube is connected to its nearby neurons, which are within a maximum distance d . The initial connection weight between neurons i and j , denoted by $w_{i,j}$, is defined as follow:

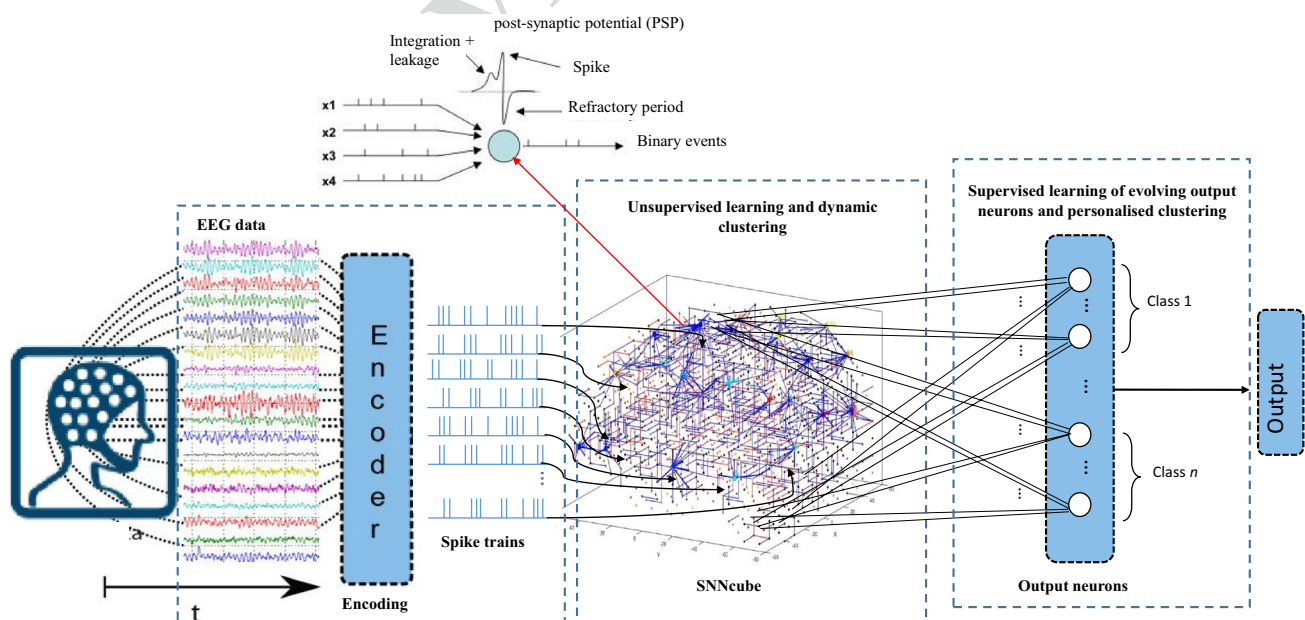


Fig. 1 A schematic block diagram of the NeuCube Architecture illustrated on EEG brain streaming data

$$w_{ij} = \frac{r_{ij}}{d_{ij}}; \quad r \in \mathbb{R}_{[-1,1]} \quad (1)$$

2.2 Unsupervised learning in the SNNcube

After SNNcube initialisation, unsupervised learning is performed using Spike-Timing Dependent Plasticity (STDP) learning rule (Song et al. 2000) as one implementation. STDP is one of the most popular example of Hebbian learning (Hebb 1949), which depends on the relative timing of pre- and post-synaptic action potentials. STDP learning rule is defined using the following relation:

$$F(\Delta t) = \begin{cases} A_+ \exp(\Delta t / \tau_+) & \text{if } \Delta t < 0 \\ -A_- \exp(-\Delta t / \tau_-) & \text{if } \Delta t \geq 0 \end{cases} \quad (2)$$

where $F(\Delta t)$ defines the synaptic modification elicited from a single pair of pre- and post-synaptic spikes separated by a time Δt . The parameters A_+ and A_- define the maximum quantities of synaptic modification, which transpire when ≈ 0 . The parameters τ_+ and τ_- determine the ranges of pre-to-postsynaptic inter spike intervals over which the synaptic strengthening and weakening occur (Song et al. 2000),

STDP learning stage is performed through transferring spikes (in time) across spatially located synapses and modifying the synapses over time. In this learning, a neuron's Post Synaptic Potential (PSP) increases by every input spike at time t to reach the firing threshold which can be also a dynamical threshold as proposed in (Schliebs and Fiasche 2012). Once the PSP exceeds this threshold, the neuron fires and sends a spike to the other neurons that are connected to it. In STDP learning, if neuron i fires before neuron j , the connection weight from neuron i to neuron j will increase, otherwise it will decrease. As a result, STDP adjusts the connection weights between neurons based on the relative timing of a particular neuron's output and input spikes. STDP learning encodes the 'hidden' spatiotemporal relations between SSTD streams in the form of "neuronal connections" and "spiking activities" in the SNNcube model. This information is used to define spatiotemporal similarity in the proposed dynamic clustering method.

2.3 Supervised learning of evolving output neurons for classification

For data classification/regression, dynamic evolving SNNs (deSNNs) (Kasabov et al. 2013) is used to train an output classifier based on the association between class labels and training samples. For each training sample, an output neuron is created and connected to each neuron in the SNNcube (shown in Fig. 1). After the unsupervised learning stage in the SNNcube is finished, the same data is propagated again through the trained SNNcube sample

by sample. The spatiotemporal pattern of activation of the trained SNNcube evoked by a particular sample is used as input data to train an output neuron to recognize this pattern. The initial connection weight between a neuron i from the SNNcube and an output neuron j is defined by using the Rank-Order (RO) learning rule (Thorpe and Gautrais 1998). Through this rule, the first arrived spike to the output neuron j would have the highest value:

$$W_{ij}(t) = \sum \text{mod}^{\text{order}(i)} \quad (3)$$

where mod is a modulation factor and $\text{order}(i)$ is the order of the coming spikes to the connection between neurons i and j .

While the RO learning will set the initial values of the connection weights W_{ij} , the STDP rule will adjust these connection weights based on further incoming spikes. The connection weight W_{ij} is further modified by a small *drift* parameter value, so that at a next time t if a spike arrives from neuron i to neuron j , the weight W_{ij} will increase by a positive *drift* parameter value and if not, it will decrease by the *drift* value. The potential of neuron j at time t is calculated as follow:

$$PSP(j, t) = \sum \text{mod}^{\text{order}(i)} W_{ij} \quad (4)$$

The details of the deSNN learning are published in (Kasabov et al. 2013).

3 The proposed methodology for clustering of SSTD in a NeuCube SNN model

In contrast to traditional clustering techniques, which are based on either spatial or temporal components and corresponding similarity measures (Kohonen 1998; Kasabov and Song 2002; Deboeck and Kohonen 1998; Kasabov 2007), we propose here a new approach for dynamic data clustering with respect to spatiotemporal similarity measures including both spatial and temporal components from the data that is learned in a SNN model. The proposed methodology is performed in both unsupervised and supervised learning modes and consists of the following clustering methods:

1. *Dynamic clustering in an unsupervised learning mode in a 3D SNN model*: dynamic spatiotemporal clusters, represented as either similar neuronal connections or similar temporal interactions between the spiking neurons in a SNN model during the unsupervised learning (Sect. 3.1).
2. *Dynamic clustering in supervised learning mode related to individual outputs ("personalised")*: (a) clusters of connections between the 3D SNN model

407 from above and each individual output neuron, repre- 442
 408 senting one class or one prototype from a given class; 443
 409 (b) clusters of spike time activity in the 3D SNN model 444
 410 from above related to each individual output neuron
 411 activation (Sect. 3.2).

412 These clustering phases are followed by analysis and 445
 413 interpretation of the created clusters for the discovery of 446
 414 new information about the SSTD and the processes that 447
 415 generated it. 448
 449
 450
 451

416 3.1 Dynamic clustering based on spiking activity 445 417 and neuronal connectivity during unsupervised 446 418 learning 447

419 In order to apply the dynamic clustering of spatiotempo- 448
 420 ral data in a SNNcube, the cluster centres are predefined 449
 421 by the spatial locations of the data sources used as input 450
 422 variables (e.g. EEG channels in the case study presented 451
 423 later). During the unsupervised STDP learning in the SNN- 452
 424 cube, spikes are transmitted between neurons across syn- 453
 425 apses that cause modifications of the connection weights. 454
 426 The more spikes are transmitted between two neurons i and 455
 427 j , (S_{ij}), the stronger the connection (W_{ij}) becomes between 456
 428 them. During the clustering procedure, each neuron 457
 429 belongs to a cluster centre (which is the input neuron, cor- 458
 430 responding to an input variable) that has the strongest con- 459
 431 nection or the most spikes received from this input neuron 460
 432 when compared with other input neurons. 461

433 Such created clusters are three dimensional and have 462
 434 different shapes as illustrated in Fig. 2a. The size and the 463
 435 shape of a cluster represent the importance of the cluster 464
 436 centre in the trained SNNcube model and therefore— 465
 437 the importance of the corresponding input variable in the 466
 438 SSTD. 467

439 In the SNNcube, the input neurons are allocated to the 468
 440 cluster centres and labelled by the input variables (e.g. 469
 441 EEG channels). The rest of the neuros in the SNNcube 470
 442 are unlabelled. The goal is to assign the cluster labels of 471
 443 the unlabelled neurons in the SNNcube. This procedure is 472
 444 addressed through the following steps:

1. An adjacency graph $G(V, E)$ is defined on the SNN- 445
 cube, where vertex set V represents the spiking neurons 446
 of the SNNcube and edges E are weighted by either 447
 S_{ij} or W_{ij} when the clustering is based on the *spiking* 448
activity or *neuronal connectivity* respectively. Note that 449
 stronger E_{ij} represents stronger intraction and more 450
 information shared between neurons i and j in SNN- 451
 cube. 452
2. Given graph G and starting vertex v_s , a random walk 453
 (Tu et al. 2014) in G of length n is defined as a ran- 454
 domised process in which, starting from the vertex v_s , 455
 we repeat n times a step that consists of choosing a ran- 456
 dom neighbour of the vertex v_s . We define p_v^n to be the 457
 probability that we chose vertex v after n steps of ran- 458
 dom walk. If random walk starts from vertex v_s , which 459
 represents one input neuron in the SNNcube, the initial 460
 distribution of the random walk can be defined as: 461

$$462 \quad p_v^0 = \begin{cases} 1, & \text{if } v = v_s \\ 0, & \text{otherwise} \end{cases} \quad (5)$$

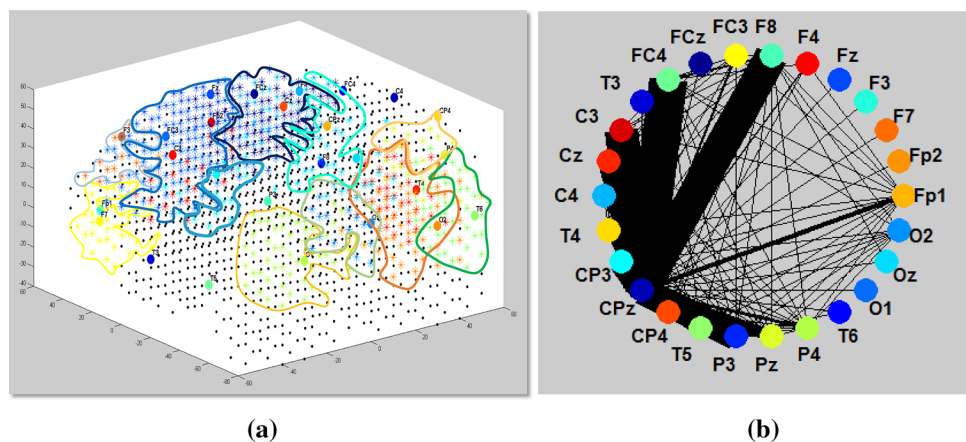
463 At a further step n of this random walk, the probability 464
 465 distribution is defined as:

$$466 \quad \forall v \in V: P_v^{n+1} = \sum_{(u,v) \in E} \frac{E_{uv}}{d(u)} P_u^n \quad (6)$$

467 where $d(u)$ is the weighted degree of vertex u , that is, the 468
 469 sum of weights of adjacent edges. 470

3. The evolution of the probability P_v^n continues as a func- 468
 tion of n until it converges, i.e. $P_v^{n+1} = P_v^n$. For every 469
 vertex $v \in V$, the obtained p_v represents the member- 470
 ship to the input vertex. At the end, each vertex $v \in V$ 471
 will belong to a starting vertex (input neuron) from 472

Fig. 2 **a** Neuronal clusters created through unsupervised clustering in a SNNcube; **b** measuring spike exchanges between neuronal clusters (the thicker the lines, the more spikes are exchanged between the clusters). The example uses results from the case study simulation of brain spatiotemporal data measured as EEG through 26 channels



473 which it obtained the highest membership p_i . This
 474 algorithm is mathematically implemented in (Tu et al.
 475 2016; Zhou et al. 2004).

476 In this paper, the above technique is applied at each time
 477 unit t of the STDP learning process to generate dynamic
 478 clusters which are formed with respect to S_{ij} or W_{ij} of the
 479 SNNcube. This clustering also allows to represent the level
 480 of interaction between clusters (corresponding to brain
 481 areas in the example of Fig. 2a) measured as number of
 482 spikes exchanged between the clusters—see Fig. 2b. In the
 483 graph presented in Fig. 2b, the thicker the connected line
 484 between two variables is, the more the spikes exchanged.

485 3.2 Dynamic clustering of the SNNcube connectivity 486 and temporal activity based on supervised learning 487 of individual evolving outputs (“Personalised”)

488 As discussed in Sect. 2, after the unsupervised learning in
 489 the SNNcube is finished, the same data is propagated again
 490 through the trained SNNcube sample by sample for super-
 491 vised learning. For each training sample, we create an out-
 492 put neuron and connect it to each neuron in the SNNcube.
 493 An output neuron is associated with a single input spatio-
 494 temporal sample or prototypes of samples from a given
 495 class. As a partial case it can represent brain or other data
 496 from one person. Initial connection weights of the neurons
 497 from the SNNcube to an output neuron j are set according
 498 to RO rule in which the earlier spikes arriving to a neuron
 499 j resulting in a higher initial connection weight. The output
 500 neuron is trained to recognize the spatiotemporal pattern
 501 of activity in the already trained SNNcube that is triggered
 502 when an input pattern corresponding to this individual is
 503 propagated through the SNNcube.

504 Through the clustering, neurons in the SNNcube are
 505 labelled by their connection weights to the output neuron j .

506 The stronger the connection from a neuron i from the
 507 SNNcube to an output neuron j , the higher the impact of
 508 the spiking activity of neuron i is on the activation of the
 509 output neuron j representing the SSTD related to brain
 510 activity of an individual subject j . In order to define the
 511 cluster centres in SNNcube, we can choose a number of
 512 neurons with different label values (between the minimum
 513 and the maximum connection weights). This clustering is
 514 performed through grouping the neurons in the SNNcube
 515 into clusters of similar connection weights to neuron j . In
 516 this way the importance of the brain areas can be ranked
 517 according to the performance of an individual j during a
 518 cognitive task.

519 During the supervised learning, when an input data sam-
 520 ple related to the performance of an individual j is entered
 521 into the already trained SNNcube, the order in which
 522 the neurons in the SNNcube spike reflects the temporal

523 activities captured in the SSTD of the individual j . Captur-
 524 ing this order in groups of neurons that spike at a similar
 525 time period results in clusters of neurons in the SNNcube.
 526 In this procedure, the neurons of SNNcube will be ranked
 527 by the temporal order in which they spike. The minimum
 528 neuron rank is 1 and the maximum rank is when the last
 529 SSTD time point comes into the SNNcube for supervised
 530 learning. We selected a number of neurons from the ranked
 531 SNNcube as cluster centres to capture groups of neurons
 532 with similar spiking orders to the centres. The cluster mem-
 533 bers are selected if they have close proximity (more than
 534 a similarity threshold) to the cluster centre. These clus-
 535 ters can be used to understand the timing of the sequential
 536 brain activities when a particular individual j is performing
 537 a cognitive task. Sect. 3.2 is illustrated in Sect. 6 on three
 538 individuals, from the case study data, belonging to three
 539 classes. Figure 3 shows a block diagram of the proposed
 540 clustering methods.

541 4 Feasibility study of the proposed clustering 542 methods on a case study EEG data

543 In order to validate the proposed methods, we demonstrate
 544 it on a case study of EEG data which was recorded dur-
 545 ing a cognitive GO-NOGO task performed by three groups
 546 of subjects: healthy subjects (H), opiate user subjects (OP),
 547 and opiate users under methadone maintenance treatment
 548 (M). Prior to commencing this research, ethical approval
 549 was granted by the “Northern X Regional Ethics Commit-
 550 tee of New Zealand” and informed consent was given by all
 551 participants. Identifying information of participants includ-
 552 ing names, initials, etc. are not reported in the paper.

553 A cognitive GO-NOGO task has been used, in which
 554 participants were repeatedly presented with the word
 555 ‘PRESS’ (for 500 ms). The colour of the word ‘PRESS’
 556 was presented randomly in either red or green. Participants
 557 were instructed to respond by pressing a button in response
 558 to the word that appeared in green (GO) and not respond to
 559 the word that appeared in red (NOGO). The EEG data was
 560 recorded via 26 EEG channels: Fp1, Fp2, Fz, F3, F4, F7,
 561 F8, Cz, C3, C4, CP3, CPz, CP4, FC3, FCz, FC4, T3, T4,
 562 T5, T6, Pz, P3, P4, O1, O2, and Oz.

563 The GO-NOGO task was displayed and simultane-
 564 ously the event-related potentials (ERPs) were recorded.
 565 We extracted 75 EEG time points per subject, each corre-
 566 sponds to grand mean average. In this case the EEG data
 567 analysis is in time domain. Based on the most literature
 568 about GO/NOGO task, the brain response inhibition, as
 569 a core executive function, is expected to be observed in
 570 prefrontal, frontal, dorsal, ventral, and parietal regions,
 571 which are related to human response inhibition. On the
 572 other hand, psychological reports showed there is a direct

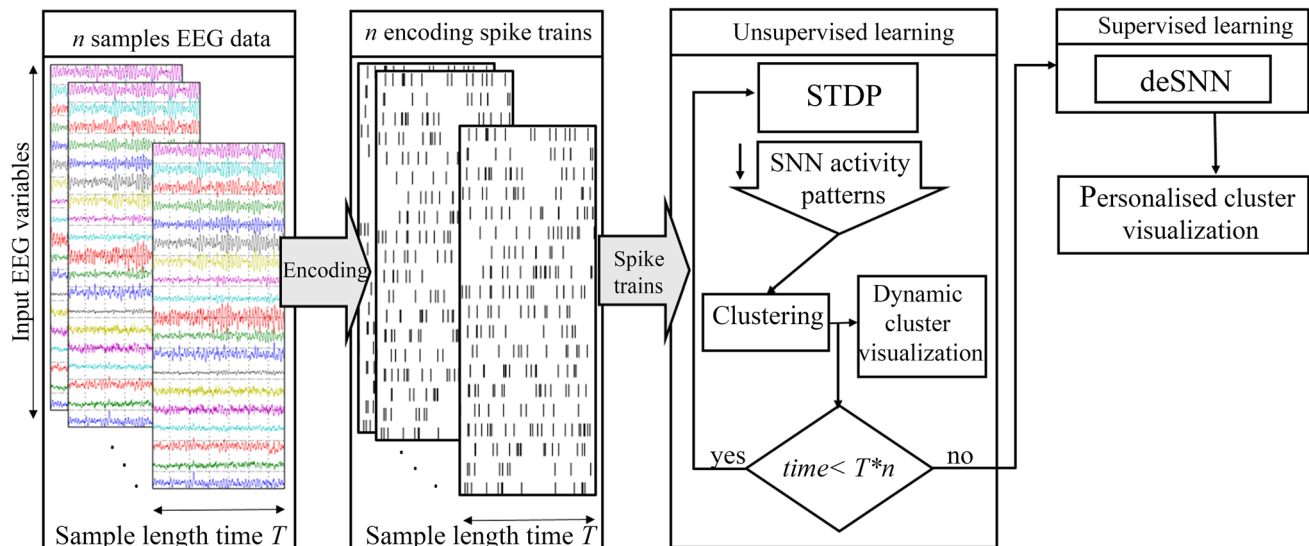


Fig. 3 A block diagram of the clustering methods in the NeuCube SNN architecture

573 relation between the response reduction in prefrontal cortex
574 magnitude and addictive behaviour, due to the drug
575 effects on these particular brain functions.

576 In this study, we aim to apply the proposed clustering
577 methods to the EEG case study, to illustrate the differ-
578 ences between the clusters in terms of the cluster evolu-
579 tion pattern, cluster creation time, and cluster size across
580 the subject groups when performed GO *versus* NOGO
581 trials.

582 We performed six EEG sample files, each containing
583 EEG data captured from one group (M/OP/H) per cog-
584 nitive task (GO *versus* NOGO). Each sample file was
585 entered separately into the SNNcube for unsupervised
586 training. Every learning was started by entering the first
587 EEG time point to train the SNNcube and it was finished
588 after entering the final EEG time point. Simultaneously,
589 neuronal clusters were created with respect to the neuron
590 spiking activity and connectivity in the SNNcube.

591 **5 Application of the proposed dynamic** 592 **clustering in the unsupervised learning mode** 593 **of the SNNcube on EEG data for the case study** 594 **problem**

595 **5.1 Dynamic clustering of EEG data in a SNNcube** 596 **model for the case study problem**

597 The SNNcube clusters were formed and updated with
598 every new input EEG time point entered, frame by frame.
599 This process can be traced and analysed in terms of:

1. The order in which input EEG channels formed the clusters, related to the order of activity of the corresponding areas of the brain. 600 601 602
2. The evolution of the size of the clusters, related to the importance of the activity of brain areas over time. 603 604

605 Figure 4 shows step-wise visualisation of the neuronal
606 cluster evolution corresponding to the 26 EEG channels
607 in the SNNcube models. Figure 4a shows how the input neu-
608 rons of the SNNcube are allocated to the respective EEG
609 channels for transferring the spike trains. It also shows the
610 evolution of the clusters for four selected time points dur-
611 ing the unsupervised learning of the SNNcube with EEG
612 data of H group when performing a GO task. Cluster crea-
613 tion started from predefined centroids and clusters were
614 changed after every input EEG time point is entered into
615 the SNNcube. Since there were 21 healthy subjects, and 75
616 EEG time points were captured from every subject, the last
617 time point of the training data was $21 \times 75 = 1575$.

618 Figure 4b, c represent the dynamic clustering of the EEG
619 data related to M and OP subjects in GO trials with a total
620 number of $29 \times 75 = 2175$ and $18 \times 75 = 1350$ EEG time
621 points respectively. The reason that we have chosen differ-
622 ent time frames in our visualisation is the time differences
623 in clusters creation across the subject groups with respect
624 to their EEG data. Once new clusters were created during
625 the NeuCube training, a new figure was captured to display
626 the step-wise changes in the cluster evolution. These results
627 show that when a SNNcube is training with EEG data of H
628 group in GO task, the first created clusters correspond to
629 Fz and FCz channels after entering the 8th EEG time point
630 to the learning process. Those neurons that are clustered

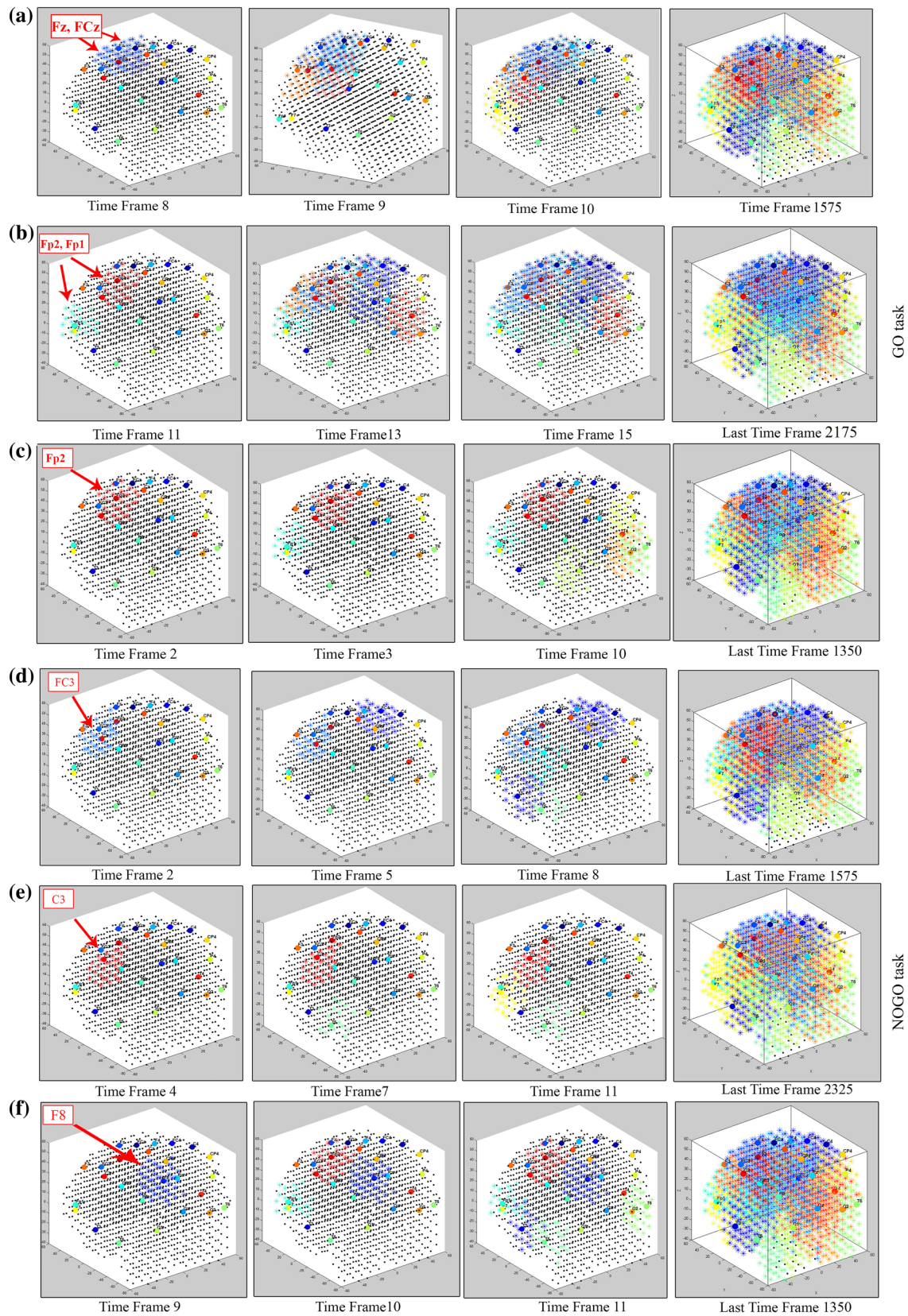


Fig. 4 a–c The cluster evolution during the unsupervised learning with EEG data of H, M and OP subjects in the GO task, d–f the neuronal cluster evolution of H, M and OP subjects in the NOGO task

by Fz and FCz channels have received the most number of spikes from these corresponding channels by this moment of the learning process. On the other hand, for M group, the first neuronal clusters are created by FP1 and FP2 channels at 11th time points. In case of OP subjects, the first cluster is created by FP2 at the 2nd EEG time point.

Figure 4d–f are related to the NOGO trials. For H group, there are a total number of $21 \times 75 = 1575$ EEG time points and the first created cluster is related to the FC3 channel at 2nd EEG time point. It represents that more spikes were transferred into the SNNcube *via* FC3 channel at this time. Therefore, those neurons located around this channel are clustered faster than the other neurons in the SNNcube. As the frontal areas of the brain are involved in inhibition functions, our finding may indicate that H group has successful inhibition with less response time in performing the NOGO trials. However, the first clusters for M and OP groups were generated later than the clusters in H group and they belonged to the C3 and F8 at 4th and 9th EEG time points respectively. These results show that in M and OP groups, slower response was observed from frontal regions and consequently less number of spikes were entered into the SNNcube when compare it with H group. There were $31 \times 75 = 2325$ and $18 \times 75 = 1350$ EEG time points for M and OP group respectively.

The neuronal clusters evolved in the SNNcube during unsupervised learning can be also statistically compared in terms of the size (number of neurons that belong to each cluster) and also in terms of the cluster creation time. The clusters were scaled up or down with respect to the number of neurons associated with every input EEG variable. A bigger cluster contains larger number of spiking neurons around the centre, which means more spikes transmitted via this centre to the SNNcube. By comparing the number of neurons that belong to each cluster centroid, we can differentiate the dynamic brain activity captured *via* different EEG channels across the subject groups in GO *versus* NOGO trials. Figure 5a visualises how the cluster size of two EEG variables change during the unsupervised learning in SNNcube. The horizontal axis represents the number of EEG data points entered to the SNNcube training via input neurons corresponding to the EEG channels. The vertical axis represents the number of neurons that belong to each cluster at each time unit of the SNNcube learning process.

As observed in prefrontal electrodes (as the potentially best candidate to reflect inhibition-related cognitive activity), there is a specific reduction in the prefrontal and frontal activity in the M group in NOGO task (illustrated by red plot). Less number of neurons in F3 and FP2 clusters may represent that M group exhibited significant reduction of their attention within the frontal region. This clustering trend allows us to understand the differences between

subject groups by distinguishing their brain activities as different clusters.

Using the proposed clustering methods, we can also extract important EEG variables that contribute to generating large clusters.

Figure 5a showed the dynamics of the cluster size changes during the presentation of the EEG data of a particular order of the subject data. Two questions may arise in this regards.

Would cluster evolution be different within a subject group?

Would the order of presentation of subject data influences the final clustering?

To address these questions, we performed the clustering experiment ten times for each subject group using random order of the subject data as shown in Fig. 6. It can be seen that different clusters based on EEG channels have different variability across subject groups and also within a subject group.

In the SNNcube, the neuron's Post Synaptic Potential (PSP) increases by every input spike arrived to the neuron at time t . Once the PSP reaches a threshold, neuron emits the output spike. The total spiking rate emitted by the neurons within a cluster as well as their total PSP rate are reported in Fig. 5b for one randomly selected cluster (FP1 channel). This result allows us to study the pattern of the cluster creation more in details by looking at the number of spikes produced within a cluster and also the pattern of the neurons' PSP during the learning.

5.2 Analysis of dynamic brain processes through mapping neuronal clusters from the NeuCube model back to brain areas

Using the NeuCube models, we can map the obtained neuronal clusters into the corresponding brain areas for a better localization of the brain activity sources and to precisely show which areas of the brain are associated to each cluster. In order to interpret the dynamic clusters in terms of associated brain areas, we have used the generic Talairach brain template (Koessler et al. 2009; Lancaster et al. 2000; Talairach and Tournoux 1988). In Fig. 7, a trained SNNcube is labelled by eight Talairach brain areas, namely: frontal lobe in yellow-green; temporal lobe in pink; parietal lobe in light-blue; occipital lobe in red; posterior lobe in light yellow; sublobar region in orange; limbic lobe in green; and anterior lobe in blue.

Figure 7a illustrates the connected neurons in two randomly selected clusters (FC3 and FC4) in healthy versus OP groups in GO task. The stronger the connections created between neurons, the more spikes transmitted between them. Figure 7b shows the strongest connections (larger than a threshold) within the clusters. Less connections

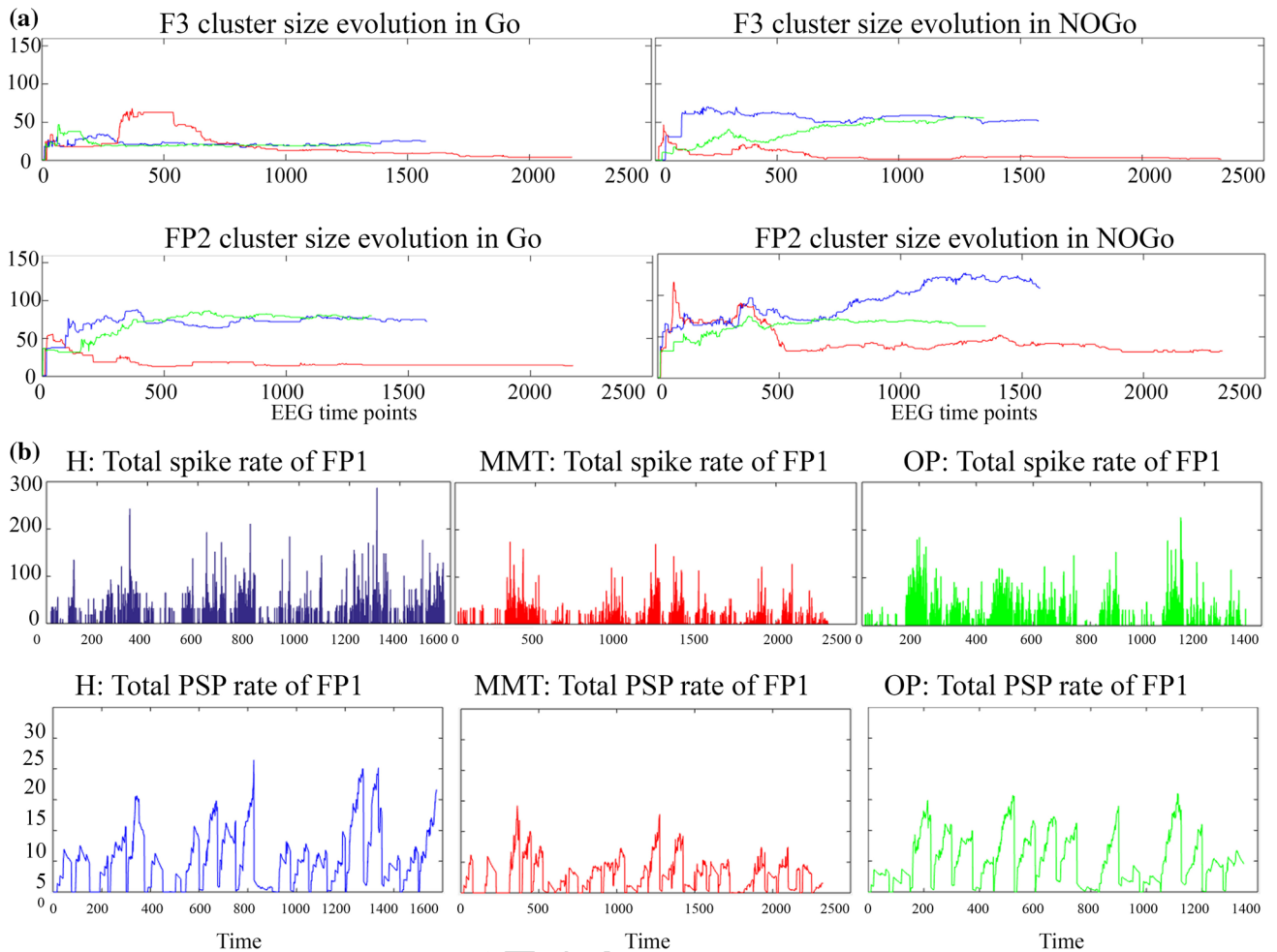


Fig. 5 **a** The size of the clusters, centred at the input neurons corresponding to EEG channels (F3, FP2), changes during the cluster evolution in the SNNcube while training on EEG data of H (blue), M (red), OP (green) subjects. **b** Total spike rate and PSP rate of the

neurons within the FP1 cluster during the unsupervised learning of EEG data in NOGO task. Blue H, red M, and green OP. The Leaky-Integrate and Fire (LIF) patterns of spiking neurons can be captured from both spiking rate and PSP rate

735 captured for OP group represents less functional brain
 736 activities observed when OP group was expected to
 737 response in the GO task in comparison with healthy group.
 738 As it is seen from Fig. 7b, c, greater connections were
 739 detected in the neuronal clusters of the healthy group, sig-
 740 nificantly in the frontal areas. Our findings represent that
 741 healthy group was able to perform more accurate responses
 742 in the GO task. Figure 7d shows the static information of
 743 dynamic cluster creation (size and time) in one particular
 744 order of subject data. It allows us to trace the sequence
 745 of the functional brain activities associated with differ-
 746 ent brain areas. In this figure we can see when the healthy
 747 group deals with GO task, the clustering of the neurons
 748 starts from the functional brain activities generated in mid-
 749 dle frontal lobe (37 and 26 neurons of SNNcube belong to
 750 the Fz and FCz clusters respectively at 8th time point of
 751 learning process) and then followed by the inferior and

the superior frontal gyrus (23, 35, 43, 27, 26, 29 neurons
 belong to F3, Fz, F4, Fc3, Fcz and C3 clusters respectively
 at 9th time point).

6 Dynamic clustering of SNNcube patterns in a supervised mode for individual output neurons of a NeuCube model on the case study problem

As described in Sect. 2.3, the second training stage is to train the output classifier using class label information associated with the training samples. The deSNN is used as output classifier is illustrated in Fig. 8 that shows for every training sample representing EEG data of one subject, one output neuron is evolved and labelled by its class label (red: healthy subjects; green: M subjects; and

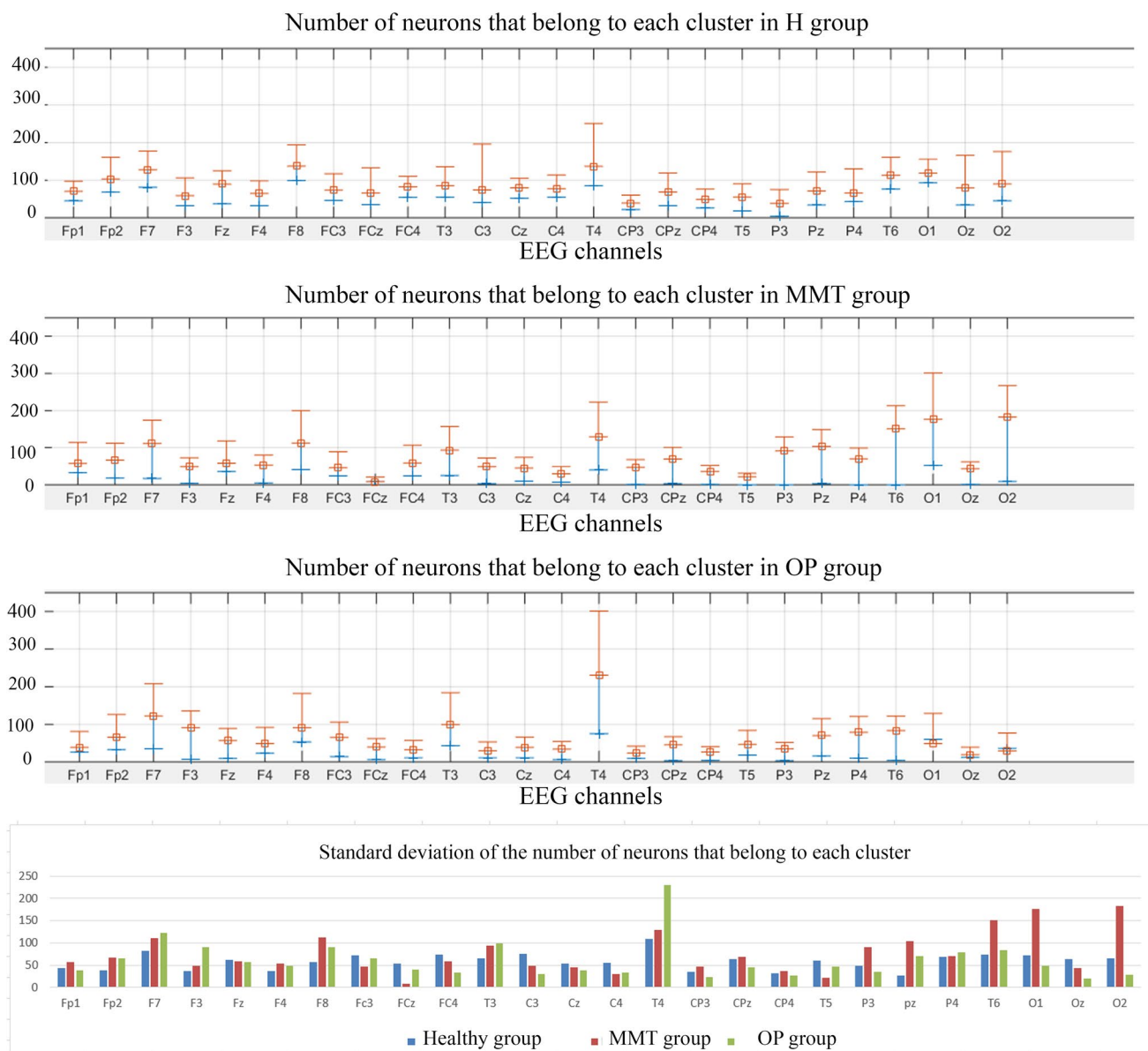


Fig. 6 Minimum, maximum and mean of the number of neurons that belong to each cluster. The dynamic clustering is run 10 times for each group using random order of the subject data presentations.

The standard deviation σ is reported in the *last row*. Higher σ mostly observed in M and OP groups

766 blue: OP subjects). The output neurons are connected to
 767 all neurons of the SNNcube and the connection weights
 768 between them are initialised based on the RO rule. This
 769 learning rule endows a higher priority to the first spike
 770 coming to the output neuron. After the first spikes
 771 entered, the connection weights are modified accord-
 772 ing to the further incoming spikes to the output neurons.
 773 When the supervise learning is completed, the connec-
 774 tion weights between SNNcube and the output neurons
 775 are established. These connection weights are visual-
 776 ised by coloured neurons. Brighter neurons mean larger

connection between neuron i from SNNcube to the output
 neuron j .

In order to represent an individual performance, we clus-
 tered the connections weights between the SNNcube’s neu-
 rons and an output neuron j based on two aspects:

- a) Clusters reveal the importance of the areas of neurons
 in the SNNcube for the activation of this output neu-
 ron.
- b) Clusters reveal the order of activity of neurons in the
 SNNcube according to their time of spiking.

Figure 8 shows for an individual subject (selected output neuron j), each neuron i from the SNNcube is labelled by the connection weight value between i and the output neuron j . In the labelled SNNcube, a number of neurons with different label values (selected between the minimum and the maximum connection weights) are selected as the cluster centres. Those neurons that have similar label value to the centre (less than a distance threshold) are selected as the cluster members.

In this example, for one individual healthy subject in NOGO task, the biggest cluster is located in the frontal areas which are associated with brain cognitive inhibition. The personalised clusters represent the importance of the brain areas corresponding to this particular subject's performance during the cognitive NOGO task. This finding is consistent with the literature that cognitive inhibition processes are ascribed to the frontal and prefrontal cortex that is fundamental for healthy neuropsychological functions. However, this observation is different for the selected M and OP individuals.

Figure 8b shows that the SNNcube is labelled by the temporal order in which the neurons spike.

For one individual healthy subject in NOGO task, we selected 5 neurons with different label values between 1 and 75 (there are 75 EEG time points per subject). In this case, we can see different clusters are captured based on the similarity in the neuron temporal spiking time. The brighter neurons mean they spike earlier. Cluster members are those neurons with close distance (less than a distance threshold) to the centres. The clusters can be interpreted to discover which areas of the brain were activated earlier. For example, for the selected healthy subject, we can see those neurons located in the visual cortex are brighter, which means they spiked earlier. These cognitive processes were then followed by spikes at neurons located in central parietal and superior frontal areas. Our findings in Fig. 8b prove that the NeuCube clustering is supported by the neuroscience literature that reported visual perception initiates as soon as the eye transfers light to the retina, where it is observed by a layer of photoreceptor cells (Carter 2014). Compared to the M and OP individuals, we conclude that the visual cortex is activated first, but not many early spikes were captured in the frontal areas. This result may suggest the cognitive inhibition impairment in those subjects with the history of drug usage.

7 Comparative analysis of the proposed dynamic, evolving clustering methods with SOM

Here we demonstrate the principle differences between the proposed clustering methods and the method of clustering in Self-Organizing Maps (SOM) (Kohonen 1998;

Deboeck and Kohonen 1998)—one of the most popular method in computational intelligence. SOM are topological maps trained on a sequence of vector based data. Here we have applied SOM [available at (Hassinen 2015)] to the case study used in the paper of EEG data of healthy, M and OP subjects in GO task. In the healthy group, there are 21 subject data samples, each contains 26 EEG channels measured during 75 time points. We reorganized the EEG data of every subject as a vector of 130 elements. For each EEG channel, the 75 time points are aggregated into 5 time points. Therefore, $26 \text{ channel} \times 5 \text{ time points} = 130$ elements involved for each sample. The training data used for SOM has 21 vectors representing 21 healthy subjects, each vector of 130 elements (Fig. 9). The neurons in the trained SOM are clustered into nine clusters. These clusters represent which input vectors are similar to each other, but it does not reveal any spatiotemporal information related to the brain processes. A single individual sample will be mapped as a point on the SOM that does not reveal any dynamic information related to the performance of this individual. This is a significant contrast to the proposed in this paper clustering methods.

8 Discussions and future work

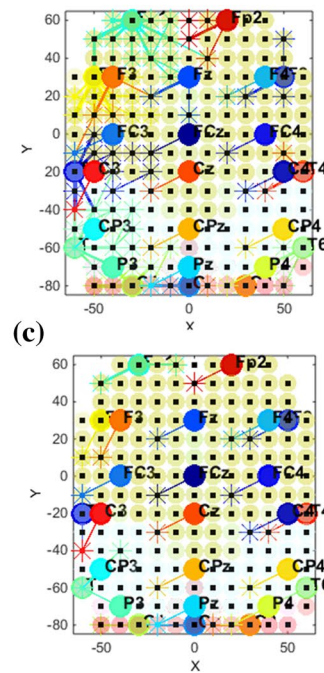
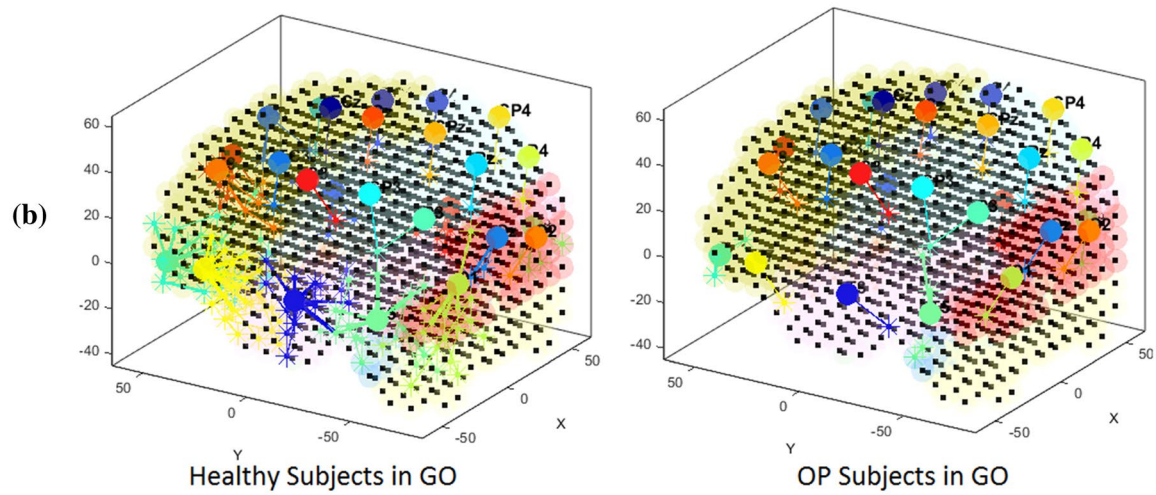
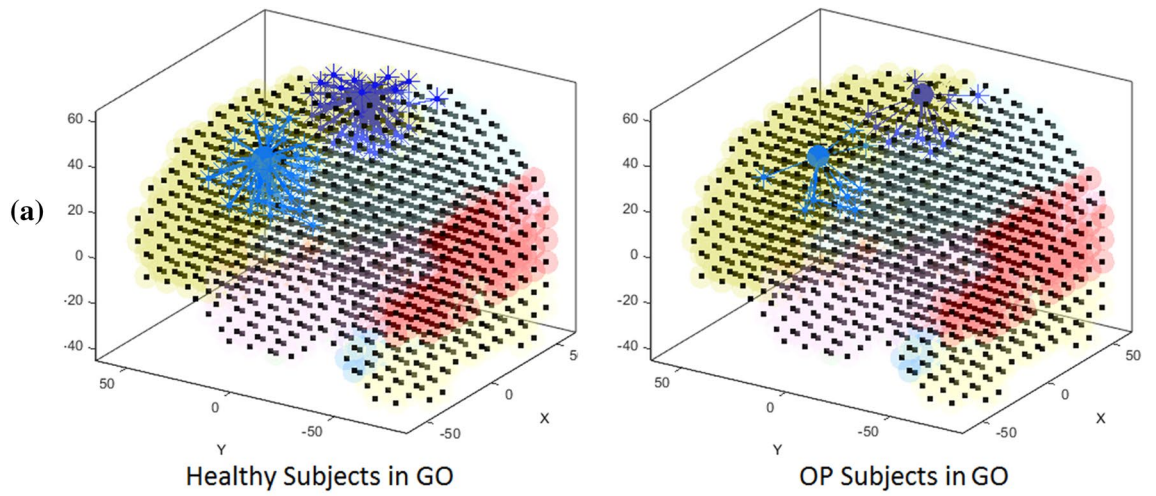
In this study, a generic SNN methodology was proposed as a novel approach to dynamic clustering of SSTD. The methods are based on the following scheme:

Dynamic processes (e.g. brain processes; seismic activities, etc.) → *Spatio/Spectro Temporal Data (SSTD)* → *3D NeuCube model creation* → *NeuCube model clustering* → *Analysis of the data and the processes that generated it* → *Cluster update on new data.*

The paper introduces two new methods for: (1) *dynamic clustering in an unsupervised learning mode in 3D SNN model*; (2) *dynamic clustering in a supervised learning mode for individual outputs ("personalised")*.

Dynamic clusters represented similar spatiotemporal changes (events) in a SNN model during unsupervised learning. This method captures significant information about SSTD as it records the exact time in which a cluster was formed and it reveals how this cluster's shape changes over time. The cluster *size* and the *time* of creation represent the importance of the input variables (e.g. EEG channels) at different time t of the learning process, providing insights into the input data structures and processes. This clustering also was used to investigate the dynamic patterns of *spiking activity* rate and *membrane potential* inside of each cluster over time.

The proposed supervised learning clustering method for individual output neurons (representing class or prototype labels) reveals two aspects of the data: (1) the importance



	Fp1	Fp2	F7	F3	Fz	F4	F8	Fc3	Fc2	Fc4	T3	C3	Cz	C4	T4	Cp3	Cp2	Cp4	T5	P3	Pz	P4	T6	O1	Oz	O2	
1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	
2	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	
3	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	
4	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	
5	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	
6	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	
7	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	
8	1	1	1	1	37	1	1	1	26	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	1	
9	1	1	1	23	35	43	1	27	26	1	1	29	1	1	1	1	1	1	1	1	1	1	1	1	1	1	
10	1	1	36	20	35	43	1	27	26	1	1	29	1	1	1	1	1	1	1	1	1	1	1	1	1	1	
11	1	1	36	28	35	43	1	33	26	1	1	20	1	1	45	19	1	1	1	1	1	1	1	1	1	1	
12	1	1	36	28	35	43	1	33	26	1	1	20	1	1	45	19	1	1	1	1	1	1	1	1	1	1	
13	1	1	36	28	35	43	1	33	26	1	1	20	1	1	45	19	1	1	1	1	32	33	35	36	28	18	
14	1	1	36	28	35	43	1	33	26	1	1	20	1	1	39	19	1	1	1	1	25	35	41	36	33	18	
15	1	1	36	28	35	43	1	33	26	1	1	20	1	1	39	19	1	1	1	1	25	27	41	36	33	26	
16	1	1	36	28	35	43	1	33	26	1	1	20	1	1	39	19	1	1	1	1	21	27	41	36	51	41	
17	1	37	36	20	35	40	1	41	26	1	40	20	1	1	39	19	1	1	1	1	21	26	41	36	53	43	
18	1	36	36	28	37	34	49	31	26	1	40	20	24	1	39	19	1	1	1	1	21	26	41	36	53	43	
19	1	36	36	28	37	34	49	33	12	1	40	20	36	1	39	19	1	1	1	1	21	25	41	35	54	44	
20	1	36	36	28	37	34	49	33	12	1	40	20	36	1	39	19	1	1	1	1	21	19	35	36	42	71	
21	15	36	36	28	37	34	49	31	14	1	40	20	36	1	39	19	1	1	1	8	22	24	37	38	53	68	
22	15	36	36	28	37	34	49	31	14	1	40	20	36	1	39	8	13	17	8	7	16	21	59	90	77	85	
23	15	36	36	28	37	34	49	33	12	1	31	20	36	1	39	8	13	17	8	7	16	21	59	90	77	85	
24	33	36	36	28	35	36	49	33	24	1	31	20	24	1	39	8	11	17	6	6	21	27	66	96	81	83	
25	34	36	36	28	35	36	49	33	24	1	31	20	24	1	39	8	12	17	4	7	17	34	72	98	85	87	
...																											
1575	166	92	89	30	47	35	117	52	16	73	83	43	36	27	95	10	85	13	35	41	10	30	97	67	43	53	

(d)

Fig. 7 **a** Clusters of connected neurons generated by FC3 and FC4 channels in two SNNcubes trained on EEG data of healthy versus OP subjects in GO task; **b** the strongest connected neurons in 26 clusters; **c** 2D visualisations of **b**; **d** static information of the dynamic cluster creation, The EEG time points are represented as rows and cluster centres are represented as columns. Each cell represents the number of neurons that belong to the cluster at this time of the learning process in the SNNcube

of the spatially located neurons in a trained SNNcube to activate an output neuron; (2) the temporal order in which neurons in the SNNcube spike to activate the output neuron.

The proposed clustering methods were applied to a case study of EEG data that measured the brain activity during a cognitive task performed by three groups of subjects.

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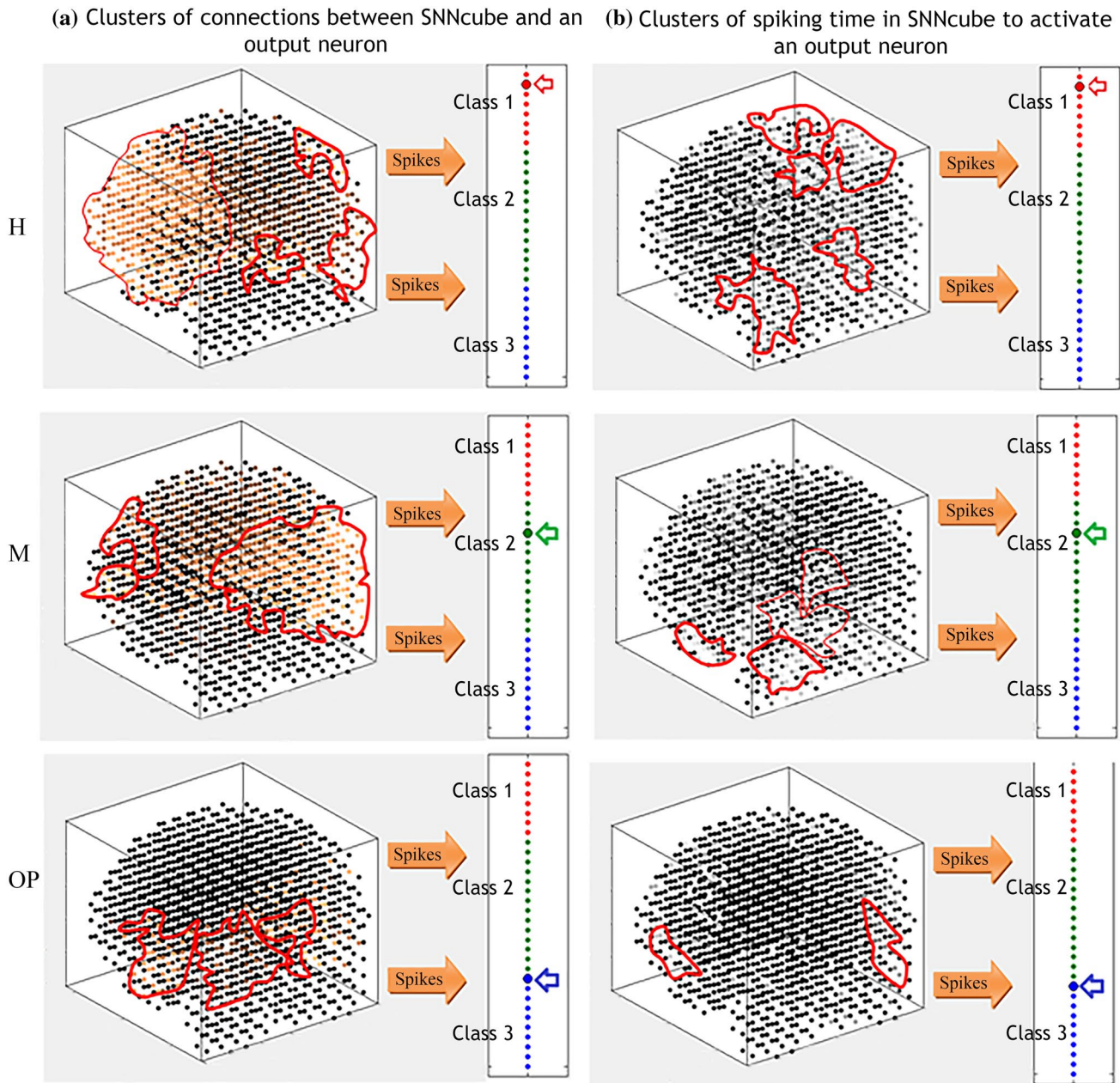


Fig. 8 In the output layer, for each training sample, one output neuron is evolved and connected to the neurons of the trained SNNcube. The output neurons are labelled by their class information in different colours (Class 1: healthy subjects is red, Class 2: M subjects in green, and Class 3: OP subjects in blue). **a** Personalised clusters of three selected subjects, belonging to each of the three groups H, M,

OP. Brighter neurons mean larger connection between neuron i from SNNcube to the output neuron j . **b** Personalised clusters represent the similar temporal order in which the neurons in SNNcube spike to activate the corresponding output neuron. The brighter the colour of the neurons, the earlier they spike

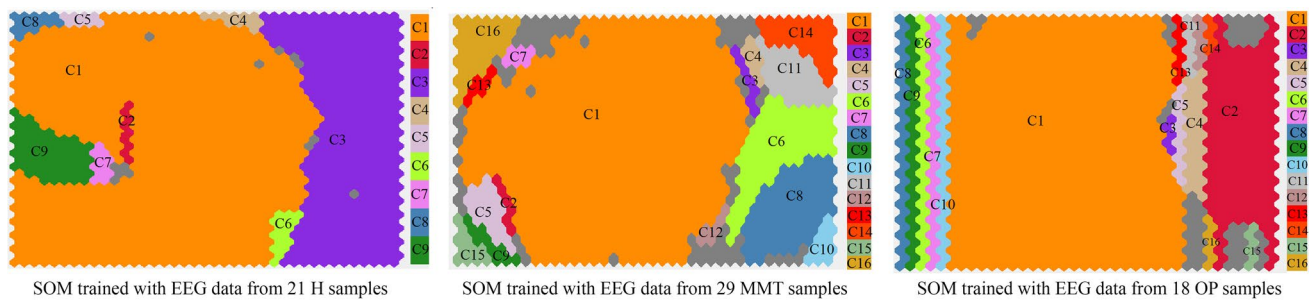


Fig. 9 SOM map generated from EEG data of 21 healthy subjects, 29 M subjects, and 18 OP subjects in a GO task. Similar healthy individual samples are clustered into nine clusters. Similar M and OP individual samples are clustered into 16 clusters each. Clusters represent

similarity between subjects across all channels and the whole time period, but not the dynamic spatiotemporal differences. No new information can be revealed about the dynamic processes captured in the EEG data for specific individuals or groups of individuals

894 Formation of the dynamic clusters of SSTD revealed the
 895 activity of the corresponding EEG channels at each time t
 896 of the SNNcube learning. The time of cluster creation and
 897 the evolution of the cluster size were found to be mean-
 898 ingful in terms of EEG channel activity over time. Through
 899 tracing the sequential spikes in the SNNcube, a direct rela-
 900 tion between the cluster creation sequence and the con-
 901 secutively activated brain areas can be revealed. Personal-
 902 ised clusters of spiking activities in the trained SNNcube
 903 revealed temporal patterns as trajectories of functional
 904 brain activities. The proposed clustering methods enabled
 905 us to comparatively analyse the EEG data recorded from
 906 different subject groups. We found greater clusters created
 907 in frontal areas on healthy group in NOGO task. It repre-
 908 sents that inhibition responses of the healthy group were
 909 stronger in comparison with the M and OP groups.

910 As a future work, we will develop an integrated cluster-
 911 ing of both static and spatiotemporal data for personalised
 912 modelling.

913 **Acknowledgements** The research is supported by the Knowledge
 914 Engineering and Discovery Research Institute of the Auckland Uni-
 915 versity of Technology (<http://www.kedri.aut.ac.nz>). M. GD was also
 916 supported by a summer research grant from the faculty of Design and
 917 Creative Technology of AUT. The authors would like to acknowledge
 918 Professor Robert Kydd and Dr. Bruce Russell from the University of
 919 Auckland and Dr. Grace Wang from AUT for providing us with the
 920 EEG data. We also acknowledge the assistance of Joyce D'Mello, Dr.
 921 Enmei Tu, Dr. Elisa Capecchi, Lei Zhou, Israel Espinosa Ramos and
 922 Akshay Gollahalli. We are indebted to the reviewers for their detailed,
 923 precise and constructive comments and suggestions that helped us
 924 tremendously. The NeuCube software along data are available free at
 925 <http://www.kedri.aut.ac.nz/neucube>.

926 References

927 Abbott LF (1999) Lapicque's introduction of the integrate-and-fire
 928 model neuron (1907). *Brain Res Bull* 50:303–304
 929 Bezdek JC, Ehrlich R, Full W (1984) FCM: the fuzzy c-means clus-
 930 tering algorithm. *Comput Geosci* 10:191–203

Capecchi E, Kasabov N, Wang GY (2015) Analysis of connectivity
 in NeuCube spiking neural network models trained on EEG
 data for the understanding of functional changes in the brain:
 a case study on opiate dependence treatment. *Neural Netw*
 68:62–77
 Carpenter GA, Grossberg S (1987) A massively parallel architecture
 for a self-organizing neural pattern recognition machine. *Comput*
Vis Gr Image Process 37:54–115
 Carter R (2014) *The human brain book*. Penguin books
 Deboeck G, Kohonen T (1998) *Visual explorations in finance: with*
self-organizing maps. Springer
 Delbruck T (2007) jAER open source project. <http://sourceforge.net/p/jaer/wiki/Home/>. Accessed 15 May 2016
 Deng D, Kasabov N (2000) ESOM: an algorithm to evolve self-organ-
 izing maps from on-line data streams. *IJCNN* 6:3–8
 Deng M, Liu Q, Wang J, Shi Y (2013) A general method of spatio-
 temporal clustering analysis. *Sci China Inf Sci* 56:1–14
 Ester M, Kriegel H-P, Sander J, Xu X (1996) A density-based algo-
 rithm for discovering clusters in large spatial databases with
 noise. In: *Kdd* 96(34):226–231
 Gholami Dobarjeh M, Kasabov N (2015) Dynamic 3D clustering of
 spatio-temporal brain data in the NeuCube spiking neural net-
 work architecture on a case study of fMRI data. In: *ICONIP*,
 pp 191–198
 Gholami Dobarjeh M, Kasabov N (2016) Personalised modelling on
 integrated clinical and EEG spatio-temporal brain data in the
 NeuCube spiking neural network system. In: *WCCI: IJCNN*,
 pp 1373–1378
 Gholami Dobarjeh M, Wang GY, Kasabov N, Kydd R, Russell B
 (2016) A spiking neural network methodology and system for
 learning and comparative analysis of EEG data from healthy ver-
 sus addiction treated versus addiction not treated subjects. *IEEE*
Trans Biomed Eng 63(9):1830–1841
 Hartigan JA, Wong M (1979) Algorithm AS 136: a k-means cluster-
 ing algorithm. *J R Stat Soc Ser C Appl Stat* 28:100–108
 Hassinen P (2015) SOM analyzer. <http://som-analyzer.software.informer.com/>. Accessed 15 May 2016
 Hebb DO (1949) *The organization of behavior: a neuropsychological*
approach. Wiley
 Indiveri G et al (2011) Neuromorphic silicon neuron circuits. *Front*
Neurosci 5:1–23, Article 73
 Johnson SC (1967) Hierarchical clustering schemes. *Psychometrika*
 32:241–254
 Kasabov N (2001) Evolving fuzzy neural networks for supervised/
 unsupervised online knowledge-based learning. *IEEE Trans Syst*
Man Cybern Part B Cybern 31:902–918
 Kasabov N (2007) *Evolving connectionist systems*. Springer

- 978 Kasabov N (2012) NeuCube EvoSpike architecture for spatio-temporal. In: Artificial neural networks in pattern recognition, lecture notes in computer science. Springer, pp 225–243 1018
- 979
- 980
- 981 Kasabov N (2014) NeuCube: a spiking neural network architecture for mapping, learning and understanding of spatio-temporal brain data. *Neural Netw* 52:62–76 1019
- 982
- 983
- 984 Kasabov N, Capecchi E (2015) Spiking neural network methodology for modelling, classification and understanding of EEG spatio-temporal data measuring cognitive processes. *Inf Sci* 294:565–575 1020
- 985
- 986
- 987
- 988 Kasabov N, Song Q (2002) DENFIS: dynamic evolving neural-fuzzy inference system and its application for time-series prediction. *IEEE Trans Fuzzy Syst* 10:144–154 1021
- 989
- 990
- 991 Kasabov N, Dhoble K, Nuntalid N, Indiveri G (2013) Dynamic evolving spiking neural networks for on-line spatio-and spectro-temporal pattern recognition. *Neural Netw* 41:188–201 1022
- 992
- 993
- 994 Kasabov N, Scott NM, Tu E, Marks S (2016) Evolving spatio-temporal data machines based on the NeuCube neuromorphic framework: design methodology and selected applications. *Neural Netw* 78:1–14 1023
- 995
- 996
- 997
- 998 Katwal SB, Gore JC, Marois R, Rogers BP (2013) Unsupervised spatiotemporal analysis of fMRI data using graph-based visualizations of self-organizing maps. *IEEE Trans Biomed Eng* 60:2472–2483 1024
- 999
- 1000
- 1001
- 1002 Koessler L, Maillard L, Benhadid A, Vignal JP, Felblinger J, Vespignani H, Braun M (2009) Automated cortical projection of EEG sensors: anatomical correlation via the international 10–10 system. *Neuroimage* 46:64–72 1025
- 1003
- 1004
- 1005 Kohonen T (1998) The self-organizing map. *Neurocomputing* 21:1–6 1026
- 1006
- 1007 Lancaster JL et al (2000) Automated Talairach atlas labels for functional brain. *Hum Brain Mapp* 10:120–131 1027
- 1008
- 1009 Lancaster JL et al (2007) Bias between MNI and Talairach coordinates analyzed using the ICBM-152 brain template. *Hum Brain Mapp* 28:1194–1205 1028
- 1010
- 1011
- 1012 Liao W, Chen H, Yang Q, Lei X (2008) Analysis of fMRI data using improved self-organizing mapping and spatio-temporal metric hierarchical clustering. *IEEE Trans Med Imaging* 27:1472–1483 1029
- 1013
- 1014
- 1015 Maass W, Thomas N, Henry M (2002) Real-time computing without stable states: a new framework for neural computation based on perturbations. *Neural Comput* 14:2531–2560 1030
- 1016
- 1017
- Mirkin B (1998) Mathematical classification and clustering. Springer 1031
- Niedermeyer E, da Silva FL (2005) Electroencephalography: basic principles, clinical applications, and related fields. Lippincott Williams and Wilkins 1032
- Ogawa S, Tank DW, Menon R, Ellermann JM, Kim SG, Merkle H, gurbil K (1992) Intrinsic signal changes accompanying sensory stimulation: functional brain mapping with magnetic resonance imaging. *Proc Natl Acad Sci* 89:5951–5955 1033
- Schliebs S, Fiasche M (2012) Constructing robust liquid state machines to process highly variable data streams. In: International Conference on Artificial Neural Networks, Springer, pp 604–611 1034
- Schliebs S, Capecchi E, Kasabov N (2013) Spiking neural network for on-line cognitive activity classification based on EEG data. In: International Conference on Neural Information Processing, Springer, pp 55–62 1035
- Song S, Miller KD, Abbott LF (2000) Competitive Hebbian learning through spike-timing-dependent synaptic plasticity. *Nat Neurosci* 3:919–926 1036
- Talairach J, Tournoux P (1988) Co-planar stereotaxic atlas of the human brain. 3-Dimensional proportional system: an approach to cerebral imaging. Thieme Medical Publishers, New York 1037
- Tanay A, Sharan R, Shamir R (2002) Discovering statistically significant biclusters in gene expression data. *Bioinformatics* 18:136–144 1038
- Thorpe S, Gautrais J (1998) Rank order coding. In: Computational neuroscience, Springer, pp 113–118 1039
- Tu E, Cao L, Yang J, Kasabov N (2014) A novel graph-based k-means for nonlinear manifold clustering and representative selection. *Neurocomputing* 143:109–122 1040
- Tu E, Kasabov N, Yang J (2016) Mapping temporal variables into the NeuCube for improved pattern recognition, predictive modeling and understanding of stream data. *IEEE Trans Neural Netw Learn Syst*. doi:10.1109/TNNLS.2016.2536742 1041
- Wang W, Yang J, Muntz R (1997) STING: a statistical information grid approach to spatial data mining. *VLDB* 97:186–195 1042
- Zhou D, Bousquet O, Lal TN, Weston J (2004) Learning with local and global consistency. *Adv Neural Inf Process Syst* 16:321–328 1043