A New Kind of Permutation Entropy Used to Classify Sleep Stages from Invisible EEG Microstructure

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Abstract: Permutation entropy and order patterns in an EEG signal have been applied by several authors to study sleep, anesthesia, and epileptic absences. Here, we discuss a new version of permutation entropy, which is interpreted as distance to white noise. It has a scale similar to the well-known $\chi^2$ distributions and can be supported by a statistical model. Critical values for significance are provided. Distance to white noise is used as a parameter which measures depth of sleep, where the vigilant awake state of the human EEG is interpreted as “almost white noise”. Classification of sleep stages from EEG data usually relies on delta waves and graphic elements, which can be seen on a macroscale of several seconds. The distance to white noise can anticipate such emerging waves before they become apparent, evaluating invisible tendencies of variations within 40 milliseconds. Data segments of 30 s of high-resolution EEG provide a reliable classification. Application to the diagnosis of sleep disorders is indicated.

Keywords: permutation entropy; sleep stages; ordinal time series

MSC: 62M10; 62P10; 68T10

1. Introduction

Modern sensor technology makes it possible to monitor vital signs continuously in everyday life, with high precision and without causing much discomfort. This will lead to very personalized, powerful and preventive medical treatment. Sleep medicine is one of the fields where such development is already apparent. While the technical basis is now available, automatic evaluation of the big data series is lagging behind. Classical methods of time series analysis usually require clean data, and preprocessing routines can easily remove information. New methods have to be developed and tested.

Use of order patterns in time series is such a new methodology. Permutation entropy, introduced in [1], has been applied not only to geophysical, financial and machine data but also in biomedical context [2–6]. Here, we are concerned with EEG (electroencephalographic) sleep data only, to which permutation entropy was applied by Ouyang et al. [7], Kuo and Liang [8], Nicolaou and Georgiou [9], and others. We shall introduce a new version of permutation entropy that can be supported by a statistical model which allows for calculating significance, and show how the settings of parameters can be optimized to recognize sleep stages from short time series of a single EEG channel. The presented methodology can be used for other medical applications, such as (see [6,10] for further references):

• Detecting epileptic absences and subclinical activity, cf. [11], or estimating the state of recovery after treatment of diabetic children [12–14];
• Monitoring depth of anesthesia, cf. [15];
• Detecting sleepiness of drivers or workers which may lead to accidents.
In the next section, we define mathematical concepts and compare the new version of permutation entropy with the usual one. A statistical discussion of significance limits for permutation entropy seems to appear here for the first time. In Section 3, we apply our method to a classical database of sleep medicine by Terzano et al. [16], which is available on physionet [17]. The sleep stages annotated by experts apparently coincide with the entropy that is measured on a continuous scale. While the annotation was done by medical doctors on the basis of multichannel data, we use only one EEG channel, and no preprocessing or special treatment of the data, just the simple entropy formula. Segments of 30 s do suffice to evaluate accurately the depth of sleep. While the experts study graphic elements on a scale of several seconds, like delta waves, as recommended by the official guidelines [18], our method analyses elements on a scale of several seconds, like delta waves, as recommended by the official guidelines [18], our method analyses the invisible microstructure in high-resolution measurements. EEG data were recorded with 512 Hz, and patterns of length between 4 and 40 ms were studied. In Section 4, we explain how we found optimal parameters, and Section 5 summarizes our main points.

2. Distance to White Noise—A New Version of Permutation Entropy

We consider a time series with \( T \) values, denoted \( x = (x_1, x_2, ..., x_T) \). In our application, the typical length \( T \) will vary between 500 and 20,000. Any three consecutive values \( (x_t, x_{t+1}, x_{t+2}) \) can form one of the six order patterns, or permutations, shown in Figure 1. We can also consider three values \( (x_t, x_{t+d}, x_{t+2d}) \) with a time distance \( d > 1 \). We say the points represent pattern 231, for instance, if \( x_{t+2d} < x_t < x_{t+d} \). In the context of EEG data, ties \( x_s = x_t \) are very rare. They can be counted as < and will be neglected in the present study. The initial time point \( t \) runs from 1 to \( T - 2d \). The delay parameter \( d \) can vary between 1 and \( d_{\text{max}} \leq T/6 \), and has the same meaning as in classical autocorrelation. Usually, \( m > 3 \) consecutive values are considered for permutation entropy, and there are \( m! \) patterns. In this paper, however, we focus on the case \( m = 3 \), for the following reasons:

- we want to keep things simple;
- for \( m = 3 \), we understand the meaning of each pattern;
- there is a nice statistical theory for patterns of length 3 [19–21];
- results for \( m = 3 \) are good when we consider various delay parameters \( d \). So far, most authors consider only \( d = 1 \) and different \( m \geq 3 \).

![Figure 1. The six order patterns of length 3.](image)

It should also be mentioned that the statistics of order pattern frequencies are excellent, even for short time series like \( T = 300 \), when we have only six patterns. For \( m = 6 \), for instance, we have \( 6! = 720 \) patterns and need a very long time series to estimate all of those pattern frequencies. Permutation entropy will still work since it is an average over all patterns. Here, we prefer the simple setting. Let us explain how frequencies are estimated for a pattern \( \pi \). We count the number of all appearances of the pattern and divide by the number of places where the pattern can occur:

\[
p_{\pi}(d) = \frac{\#\{t \mid (x_t, x_{t+d}, x_{t+2d}) \text{ represents pattern } \pi\}}{(T - 2d)}.
\]

(1)

To understand the method, consider the short time series \( x = (2, 9, 5, 8, 6, 1, 3) \) shown in Figure 2. The table collects the frequencies. As a result, we have \( p_{321}(1) = \frac{1}{7} \), \( p_{321}(2) = \frac{1}{4} \), and \( p_{321}(3) = 0 \), and could draw this as a kind of autocorrelation function.
Frequencies of single patterns have been studied by several authors [4]. For statistical reasons, we prefer to study only certain sums and differences of pattern frequencies [20,21]. The permutation entropy is the Shannon entropy of the distribution of all patterns. It is defined for the set \( S_m \) of all \( m! \) patterns of length \( m \) [1]. For our case \( m = 3 \), the sum involves only the six terms indicated in Figure 1. However, the delay parameter \( d \) can vary again between 1 and \( d_{\text{max}} \) so that permutation entropy also becomes a kind of autocorrelation function:

\[
H(d) = - \sum_{\pi \in S_m} p_{\pi}(d) \log p_{\pi}(d). \tag{2}
\]

Entropy as a measure of disorder is a basic concept in physics. Permutation entropy was introduced as complexity measure for time series. \( H \) assumes its smallest value zero for a monotone series, and its maximum \( \log m! \) for white noise, where there is no dependence among the values. White noise means that all possible permutations appear with the same probability \( 1/m! \). Here, we use a new version of permutation entropy, called “distance to white noise” and defined for \( m \geq 2 \) as

\[
\Delta^2(d) = \sum_{\pi \in S_m} \left( p_{\pi}(d) - \frac{1}{m!} \right)^2 = \sum_{\pi \in S_m} p_{\pi}(d)^2 - \frac{1}{m!}. \tag{3}
\]

We just take the squared Euclidean distance of the observed pattern frequencies from the uniform pattern frequencies \( 1/m! \) in the space of all pattern distributions. Thus, the smallest value of \( \Delta^2 \) is zero and means complete independence of values. Large \( \Delta^2 \) means much dependence among the values of the time series. This is easy to understand, and we cannot become confused by terms like “complexity”, “chaos”, and “disorder”. The sum in Equation (3), as well as in Equation (2), contains \( m! \) terms, which means six terms for \( m = 3 \). The equality on the right side of Equation (3) follows from \( \sum p_{\pi}(d) = \sum 1/m! = 1 \).

There are several reasons to call \( \Delta^2 \) a version of permutation entropy:

- Equation (3) says that we get \( \Delta^2 \) from \( H \) by replacing \( -p \log p \) with the simpler function \( p^2 \), and adding a constant so that the minimum is zero—not a big change!
- Up to a linear transformation, \( \Delta^2 \) is the quadratic Taylor approximation of \( H \) at white noise ([19], see below for \( m = 3 \)).
- For a discrete probability space \( \{\omega_1, \omega_2, \ldots\} \) with probabilities \( p_i = P(\omega_i) \) the quantity \( -\log_2 \sum p_i^2 \) is called Renyi entropy of order 2, or correlation entropy [22], and
- \( 1 - \sum p_i^2 \) is called Tsallis entropy of order 2 or Kendall information content [23].

For the case of two probabilities \( p \) and \( 1 - p \) (length 2 patterns 12 and 21), Figure 3 shows the functions \(-H = p \log p + (1 - p) \log(1 - p)\) and \( 2\Delta^2 = 2 \left[ \left( p - \frac{1}{2} \right)^2 + \left( 1 - p - \frac{1}{2} \right)^2 \right] \). They do not differ much, and agree asymptotically at the point \( p = \frac{1}{2} \). The same holds for the six probabilities \( p_{\pi} \).
of patterns of order 3. At the point $p_{\pi} = \frac{1}{6}$ for all $\pi$, which corresponds to white noise, it can be shown that $H \approx \log 6 - 3\Delta^2$ is the quadratic Taylor approximation of $H$ [19]. Note that EEG data, compared to other time series like ECG (electrocardiogram), are very erratic, close to white noise so that $H$ and $\Delta^2$ will lead to similar results. This will be demonstrated in Figure 10.

![Figure 3. Difference between $H$ and $\Delta^2$ is essentially a scale change, cf. Figure 10.](image)

It turns out that $\Delta^2$ has better statistical properties than $H$. In [19], it was shown that $\Delta^2$ can be separated into different components according to an equation

$$4\Delta^2 = 3\tau^2 + 2\beta^2 + \gamma^2 + \delta^2, \quad (4)$$

which allows a more detailed study with a kind of ANOVA method. For EEG data, only the $\tau$ component is important and will be discussed below.

We need to know the statistics of permutation entropy in order to check whether certain extreme values of $H$ are mere coincidence or really indicate a certain effect: good order or large disorder. Although a few hundred papers deal with permutation entropy, this statistical aspect seems to be discussed here for the first time.

For statistical inference, we always need a null model. In our case, the null hypothesis is that the data are white noise: completely independently chosen random numbers from the same distribution. The type of distribution does not matter for ordinal patterns. We can take the uniform distribution on $[0, 1]$. In a computer simulation, we now take a large number $N$, say 10 million, time series of length $T = 1000$, all made of independent random numbers. We determine the permutation entropy for each sample series, getting $N$ possible values $H$. These $N$ values vary near the maximum value $\log 6$, which is the theoretical value of permutation entropy of length 3 patterns for white noise. In each sample series, the value is somewhat smaller, however.

It is reasonable to consider $H / \log 6$ to get a standard scale with maximum value 1. Figure 4 shows the density of all $N$ sample series. We see that all standardized $H$-values vary between 0.99 and the maximum value 1. The conclusion is that when we observe a time series of length $T = 1000$ in practice, and $H / \log 6$ is less than 0.99, we can be sure that this is not a random deviation from white noise! Even 0.995 would be a significant observation, since the tail probability or $p$-value of 0.995 shown in the lower panel of Figure 4 is about 0.0026. This means that only 0.26% of our 10 million simulations of white noise gave a standardized $H$ value below 0.995. The value 0.99 is more significant, however, since only 100 samples gave a still smaller $H$, which corresponds to a $p$-value of $10^{-5} = 0.001\%$.

The tail probability of $H / \log 6$ in Figure 4 is almost a linear function in semilogarithmic representation, so it is easy to approximate numerically. There are two problems, however. First, the scale between 0.99 and 1 is not very intuitive. It can lead to confusion between quantile values of the $H$-statistics and the $p$-values themselves. Second, and worse, the dependence on $T$ has not been considered. It is clear that the simulation will vary more when we have smaller $T$, that is, shorter time series. The mathematical formula for this dependence is not obvious.
Compared to the other patterns of Figure 1, 123 and 321, white noise is a kind of variance, it can be shown to scale with $1/T$. Phenomena become visible, they manifest themselves in statistical properties of the fine structure of high-resolution panel and in Table 1. Even more importantly, we have drawn $T^2 \Delta^2$ instead of $\Delta^2$. Since distance to white noise is a kind of variance, it can be shown to scale with $1/T$, and the curves of $T \Delta^2$ almost coincide for $T$ larger 1000. Thus, the critical values of $T \Delta^2$ in Table 1 are almost universal while the critical values of $H / \log 6$ are valid for $T = 1000$ only. To give just one example, the value 4.68 for the 0.01% threshold of $\Delta^2$ is 4.67 for $T = 500$ and 4.69 for $T = 2000$, while the 0.01%-quantile 0.9921 of standardized $H$ will change to 0.9843 and 0.9961, respectively. Still more extreme quantiles are harder to simulate and less stable.

To conclude our statistical discussion, let us note that this is just a beginning. More modelling is needed. The white noise hypothesis is not so exciting and would not make sense for heart or respiration data. For EEG data and, in particular, for sleep stages; however, white noise is a reasonable null hypothesis, as we shall see below.

### 3. $\Delta^2$ as a Measure of Sleep Depth

We briefly explain the basic idea for our classification of sleep stages. The brain of a healthy awake adult fulfils a large variety of functions, and each EEG channel covers the activity of millions of neurons in the cortex. Thus, normally, the signal will be almost white noise. With the onset of sleep, neuronal activity will become weaker and less diverse, and global rhythms are taking over. Global phenomena become visible for a human observer of the data as delta waves, sleep spindles, K-complexes, as described in the official guidelines for sleep scoring [18]. However, long before global phenomena become visible, they manifest themselves in statistical properties of the fine structure of high-resolution data. For our application, the main change is an increase of the frequency of patterns 123 and 321, compared to the other patterns of Figure 1.
In other words, the number of local minima and maxima will decrease. Since a change of standardized $H$ from 1 to 0.99 is already highly significant, such a statistical tendency can be swiftly determined with permutation entropy. Using the language of Fourier analysis, we would state that high frequencies become weaker. However, the idea of brain signal as a composition of sine waves is not quite correct. Such waves may develop only partially, for less than a quarter of a wavelength. In such cases, they are detected by permutation entropy before the frequency spectrum shows any changes. Moreover, order pattern statistics is much more stable and less susceptible to data artefacts than the Fourier frequency statistics.

The deeper the sleep, the more our EEG signal will deviate from white noise. Thus, $\Delta^2$ should be a good measure for sleep depth. Figure 5 shows how well this idea works. We have chosen the classical CAP sleep database of Terzano et al. [16] for different reasons: it is freely available at physionet [17]. It contains a number of EEG measurements taken with sample rate 512 Hz while many other datasets contain 128 Hz measurements or low-pass filtered signals. The data quality is good and expert sleep annotation files are provided, still including sleep stage S4, which was later abandoned [18].

For four healthy subjects, Figure 5 shows the expert annotation as a step function on the lower part and $\Delta^2$ as a noisy function on the upper part of the respective panel. It turns out that we almost have mirror symmetry. Whenever the sleep stage increases, $\Delta^2$ increases, and vice versa. The calculation of $\Delta^2$ was done with the data as they were provided, without any preprocessing or selection of "clean segments". Non-overlapping windows of 30 s in length were used to calculate each value.

REM (rapid eye movement) phases, indicated by red lines in every annotation of Figure 5, are not considered here. Figure 6 below shows that with modified delays, a function $\tau$ related with $\Delta^2$ and defined in Equation (5) below can indicate REM phases, but does not classify them accurately. We should admit that this is not a systematic study, which would need tight cooperation with medical experts and could be better done with recent measurements. Moreover, if our primary interest was accurate classification, we had to use the whole power of multivariate datasets. Here, the challenge was to get maximum information from a single EEG channel.

The choice of patients for Figure 5 was based on availability of an EEG channel with 512 Hz frequency in the data, not at all on the quality of the coincidence. There were only four healthy controls with a 512 Hz EEG channel. Figures 7–9 show patients with insomnia, narcolepsy and nocturnal frontal lobe epilepsy with high-resolution EEG channel from the CAP sleep database of Terzano et al. [16].

**Figure 5.** Distance to white noise and expert sleep stage annotation for healthy controls in the CAP sleep database of Terzano et al. [16] available at physionet [17].
available at physionet \cite{17}. The coincidence of annotation and $\Delta^2$ was always excellent, though there were more artefacts. The standard EEG channel was Fp2–F4. On three occasions, another channel was provided, and gave similar results.

Figure 6. An average of $\tau$ in Equation (5) for $d$ between 40 and 70 ms for subject n2 indicates sleep stages and REM phases.

Figure 7. Distance to white noise and expert sleep stage annotation for insomnia patients in the CAP sleep database of Terzano et al. \cite{16}.

For the healthy subjects in Figure 5, maximum values of $\Delta^2$ do not differ much: they are around 0.1. We do not know whether it makes sense to compare sleep depth of different persons just by $\Delta^2$. Maybe individual factors do influence $\Delta^2$. We even had no data to check changes of $\Delta^2$ when one subject is measured repeatedly. We think $\Delta^2$ will not depend much on measurement details, but this was not verified. In Figure 7, showing patients with insomnia, the overall level of $\Delta^2$ is much lower than for the controls in Figure 5. This indicates that $\Delta^2$ can also detect certain sleep disorders, by taking the average $\Delta^2$ and grouped box plots over several hours. In a similar way, average $\Delta^2$ can be used to compare the sleep of one subject in several nights or under different conditions. There are numerous ways to exploit the permutation entropy.

Actually, it does not matter whether we take $\Delta^2$ or the original permutation entropy $H$ as a measure of sleep depth. As Figure 10 shows, they do almost coincide after a linear scale change. Since our data are fairly near to white noise, this can be proved by Taylor’s formula, as mentioned in Section 2. We chose $\Delta^2$ since it has a more natural scale, a nice interpretation, and a familiar statistics. What really matters is the choice of delays $d$ over which we average $\Delta^2$ or $H$, respectively. Now, we explain how we choose those parameters.
Figure 8. Distance to white noise and expert sleep stage annotation for narcolepsy patients in the CAP sleep database of Terzano et al. [16].

Figure 9. Distance to white noise and expert sleep stage annotation for patients with nocturnal frontal lobe epilepsy in the CAP sleep database of Terzano et al. [16].
Two milliseconds and 1000 (two seconds) does not make sense, however. We have to decide whether
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well above the bound
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were based on another idea: we looked for parameter regions that are generally far from white
noise. When there is already some smoothness in the data, a wave is more likely to emerge than in
complete disorder.

Permutation entropy is a simple method. It does not depend too much on long-term experience.
Essentially, with a bit of care, one cannot go wrong with it. Nevertheless, we have to think about some
details in order to optimize performance.

**Window length.** Since we decided to study patterns of length \( m = 3 \), only two parameters can be
chosen: the length of the sliding window and the delay \( d \). For the window, a length of 30 s seemed
to be most appropriate. On one hand, 512 Hz sampling means that we get 15,360 values within
30 s, which provided excellent statistics for order patterns even in the presence of gross artefacts.
On the other hand, we got 120 instances of \( \Delta^2 \) per hour, each obtained independently of all other
data, while expert annotation usually keeps in mind the previous sleep stage. As Figures 5 and 7–9
confirm, there are few outliers, indicating that \( \Delta^2 \) is a reliable and robust measure of sleep depth.
For artefact-free data, shorter windows can be used. It is possible to consider overlapping windows,
which was not needed in this study.

**Delays.** For the choice of delay, some experiments were done. To minimize statistical error,
taking the average over several \( d \) is better than a single \( d \). An average over all possible \( d \) between
1 (two milliseconds) and 1000 (two seconds) does not make sense, however. We have to decide
whether small or large \( d \) will give the most informative \( \Delta^2 \). According to the official guidelines for sleep
annotation [18], we should care mostly for delta and theta waves, that is, \( d \geq 200 \). Our experiments
were based on another idea: we looked for parameter regions that are generally far from white
noise. When there is already some smoothness in the data, a wave is more likely to emerge than in
complete disorder.

In real measurements, true white noise is unlikely to appear. The majority of our \( \Delta^2 \) values was
well above the bound \( \frac{5}{7} = \frac{25}{17500} \approx 0.0003 \), which marks the significance level of 0.01% in Table 1.
Smaller values occur mainly for large \( d \) where measurements at \( t \) and \( t + d \) have nothing to do with
each other (theta and delta waves are exceptions). For small \( d \), however, there are always dependencies
among the values \( x_t \) and \( x_{t+d} \) due to some slowly changing conditions in the environment of the
measuring device. Thus, for small \( d \), there is a kind of smoothness that causes patterns 123 and 321
to occur more often than the other patterns. However, if \( d \) is very small, the smooth component of
\( x_{t+d} - x_t \) will be dominated by noise. This argument says that it is best to take small \( d \), but not too
small \( d \). It seems a general rule for the choice of delays in such applications.

Figure 11 shows large and small values of \( \Delta^2 \) for all windows and all \( d \) between 1 and 768, for the
control n2 shown already in the top panel of Figure 5. Values smaller than \( \frac{15}{2} \approx 0.0015 \) are called small,
and indicated by a dark dot while greater values are left white. Moreover, the \( d \) scale is divided into
an upper part with \( d = 0.25 \text{ s}, ..., 1.5 \text{ s} \) and a lower part \( d = 2 \text{ ms}, ..., 0.25 \text{ s} \), which is magnified in order
to show details. The chosen threshold is three times the significance level of 0.01% of \( \Delta^2 \). Theoretically,
it should correspond to a tiny \( p \)-value (cf. Section 2), but as real data are not white noise, this threshold
seems appropriate [19].

**Figure 10.** Distance to white noise and permutation entropy essentially differ only by a scale change,
as demonstrated here for control n11 in Figure 5. \( H \) varies between 1.4 and 1.8, \( \Delta^2 \) between 0 and 0.1.
Here, we show \( \log 6 - H \) and \( 3\Delta^2 \), which, according to Taylor’s formula, do agree near 0.
In the upper part of Figure 11, the average $\Delta^2$ is 0.0017, and 47% of the places have small $\Delta^2$. These black spots are spread rather uniformly, so there is little chance to get information from this range of $d$. In the lower part, the average $\Delta^2$ is 0.013, and only 10% of the places have small $\Delta^2$. There is some structure related to the sleep annotations in Figure 5. There may be different choices of an interval for $d$. After some experiments, we took the bottom region, $d$ below 40 ms, which is almost completely white. The smallest value $d = 1\leq 2$ ms was excluded since variations within 2 ms are more due to the electronic equipment than to the brain, as we knew from our own measurements. Thus, the $\Delta^2$ for Figures 5 and 7–10 was taken as an average over the values $d = 2, 3, ..., 20$ corresponding to 4, 6, ..., 40 ms.

Checking oscillations. Periodic phenomena have a large influence on the statistics of order patterns. As explained in [20], the persistence

$$\tau(d) = p_{123}(d) + p_{321} - \frac{1}{3}$$

(5)

assumes large negative values for $d = \frac{p}{2}$ and $\frac{3p}{2}$ when $p$ is the period of a periodic component. For our parameters, distance to white noise consists mainly of the persistence part, Equation (4) turns into $\Delta^2 \approx \frac{2}{3}\tau^2$. Thus, it is natural to ask whether our $\Delta^2$ was caused by certain oscillations.

In EEG measurements, a danger is contamination with mains hum, the 50 Hz frequency of the power supply. The corresponding $\frac{\tau}{4}$ is $d = 5$. Since $\tau(5)$ is not particularly small, there seems to be no such contamination. We should also check for alpha waves in the range 8 up to 12 Hz, although they are not likely to appear in the channel Fp2–F4. The corresponding $\frac{\tau}{4}$ is a $d$ between 40 ms and 60 ms and is outside the range of our average. We conclude that our distance to white noise is not caused by oscillations.

This section should demonstrate ways to find good parameters. We do not claim that we made the best choice. Figure 6 shows that for the same person n2 an average of $\tau$ for $d$ between 40 and 70 ms indicates sleep stages as large $\tau$-values and REM phases by negative $\tau$-values.

5. Discussion and Conclusions

Several authors used permutation entropy as a tool for EEG analysis, both in sleep medicine [7–9] as well as in epilepsy [12–14] and anaesthesia research [15]. One advantage is the robustness of ordinal parameters like $H, \Delta^2$ and $\tau$ with respect to motion artefacts and low-frequency perturbations, which often appear in EEG data. While in correlation and spectral analysis, an outlier will cause an error proportional to its size, in ordinal pattern statistics, an outlier is counted as any other value.

In this note, we tried to improve the methodology by introducing distance to white noise, which can be supported by a statistical model. It was shown how good parameters can be determined.
As a result, we defined an average $\Delta^2$ for time spans between 4 and 40 ms, which can be considered as a measure of sleep depth on a continuous scale, very similar to the discrete sleep stages annotated by experts or by automatic scoring. A remarkable coincidence was shown in Figures 5 and 7–9 for 20 subjects in the classical CAP sleep database of Terzano et al. \[16\]. A single EEG channel and short windows of 30 s gave a reliable estimate of sleep depth. Patients with insomnia had much smaller $\Delta^2$ levels than healthy controls.

Although these results have to be checked with other, more recent databases, it could be confirmed that permutation entropy is a very effective tool for distinguishing sleep stages. In the present study, only length 3 patterns were used. The distance between the points, the so-called delay $d$, was varied in a wide range, so that permutation entropy and distance to white noise become functions like classical autocorrelation. Such a function is more meaningful than permutation entropy of patterns of length $m \geq 3$ for delay 1.

On a general level, it was shown that the fine structure of high-resolution measurements can contain invisible information. Routine low-resolution measurement, downsampling or low-pass filtering can destroy this information, while ordinal methods have the capacity to exploit the microstructure of signals. They need to be developed further.

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