Simple-Cell-Like Receptive Fields Maximize Temporal Coherence in Natural Video

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Recently, statistical models of natural images have shown the emergence of several properties of the visual cortex. Most models have considered the nongaussian properties of static image patches, leading to sparse coding or independent component analysis. Here we consider the basic time dependencies of image sequences instead of their nongaussianity. We show that simple-cell-type receptive Delds emerge when temporal response strength correlation is maximized for natural image sequences. Thus, temporal response strength correlation, which is a nonlinear measure of temporal coherence, provides an alternative to sparseness in modeling simple-cell receptive Deld properties. Our results also suggest an interpretation of simple cells in terms of invariant coding principles, which have previously been used to explain complex-cell receptive Delds.

1	Introduction			

The functional role of simple cells has puzzled scientists since the structure of their receptive Delds was Drst mapped by Hubel and Wiesel in the 1950s (Palmer, 1999). The Drst hypothesis concerning their role was based on their visual appearance, that is, their similarity with edges and bars. The second major theory was the local spatial frequency analysis theory, which is based on generally applicable signal processing principles. The current view of the functionality of sensory neural networks emphasizes learning and the relationship between the structure of the cells and the statistical properties of the information they process (see, e.g., Field, 1994; Simoncelli & Olshausen, 2001). A major advance was achieved when Olshausen and Field (1996) showed that simple-cell-like receptive Delds emerge when sparse coding is applied to natural image data. Similar results were obtained with independent component analysis (ICA) shortly after (Bell & Sejnowski, 1997; van Hateren & van der Schaaf, 1998). In the case of image data, ICA is closely related to sparse coding (Hyvärinen, Karhunen, & Oja, 2001; Olshausen & Field, 1997).

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In this article, we show that an alternative principle, temporal coherence (Becker, 1993; Földiák, 1991; Kayser, Einhäuser, Dümmer, König, & Körding, 2001; Mitchison, 1991; Stone, 1996; Wiskott & Sejnowski, 2002), leads to the emergence of simple-cell receptive Delds from natural image sequences. This Dinding is signid cant because it means that temporal coherence provides a complementary theory to sparse coding as a computational principle behind the formation of simple-cell receptive Delds. The results also link the theory of achieving invariance by temporal coherence (Földiák, 1991) to real-world visual data and measured properties of the visual system. Whereas previous research has focused on establishing this link for complex cells, we show that such a connection exists even on the simple-cell level.

Temporal coherence is based on the idea that when processing temporal input, the representation changes as little as possible over time. Földiák (1991) was one of the \Box rst to suggest the usefulness of temporal coherence in computational neuroscience. He developed a two-layer network that was able to learn to identify a \Box xed feature, such as a line with a \Box xed orientation, even if the way the feature was expressed in the data changed, for example, if the line was translated. Földiák used temporal coherence as a tool to learn translation invariances: arti \Box cially generated input data were temporally coherent (consecutive input frames contained translated versions of a line with the same orientation), and by using competition and short-term memory, the output was also taught to be temporally coherent. This associated translated versions of a feature with each other.

Several other researchers have studied temporal coherence and other forms of coherence, such as coherence with respect to different views of the same scene. For example, in Becker and Hinton (1992) and Stone (1996), surface depth was discovered from stereograms by a multiple layer nonlinear network. In Becker and Hinton (1992), this was done by using stereograms representing different views of the same randomly generated scene and maximizing mutual information between outputs. In Stone (1996), learning was achieved by using a temporal sequence of slightly different stereograms and maximizing temporal smoothness of output while preserving variability in output. In these studies, the input data sets were generated so that there was an underlying coherent parameter in the data, and the objective was to \Box nd that parameter by using coherence. Therefore, the main result was the demonstration of the usefulness of coherence using simulated data.

The contribution of this article is to show that when the input consists of natural image sequences, the linear leters that maximize temporal response strength correlation are similar to simple-cell receptive leds. We let describe temporal response strength correlation, which is a measure of temporal coherence, and an algorithm capable of optimizing the measure. In section 3, we apply this algorithm to natural image sequences. In addition to the main results, we describe several control experiments that were made to ensure the validity and novelty of our results. Finally, in section 4, we give an intuitive explanation of why optimization of the objective function pro-

duces such results, discuss the spatiotemporal and nonlinear (nonnegative) extensions of the model, and conclude by discussing the implications of this work.

2 Temporal Response Strength Correlation ______

In the basic model, we restrict ourselves to consider linear spatial models of simple cells. Linear simple-cell models are commonly used in studies concerning the connections between visual input statistics and simple-cell receptive Delds (Bell & Sejnowski, 1997; Olshausen & Field, 1996; van Hateren & van der Schaaf, 1998), because linearity seems to approximately characterize most simple cells (DeAngelis, Ohzawa, & Freeman, 1993b). Extensions of this basic framework are discussed in sections 4.4 and 4.5.

The model uses a set of spatial \square Iters (vectors) $w_1; \ldots; w_K$ to relate input to output. Let signal vector x.t/ denote the input of the system at time t: A vectorization of image patches can be done by scanning images column-wise into vectors. For windows of size N Σ N; this yields vectors with dimension N²: The output of the kth \square Iter at time t, denoted by signal y_k .t/; is given by the dot-product

$$y_k.t/D w_k^T x.t/$$
: (2.1)

Let matrix W D $[w_1; ...; w_K]^T$ denote a matrix with all the \square Iters as rows. Then the input-output relationship can be expressed in vector form by

$$y.t/DWx.t/;$$
 (2.2)

where signal vector y.t/ D $[y_1.t/; ...; y_K.t/]^T$:

Temporal response strength correlation, the objective function, is defined by

f.W/D
$$\underset{kD1}{\times}$$
 E_tfg.y_k.t//g.y_k.t i 1 t//g; (2.3)

where the nonlinearity g is strictly convex, even (rectifying), and differentiable. The symbol 1 t denotes a delay in time. The nonlinearity g measures the strength (amplitude) of the response of the \Box lter and emphasizes large responses over small ones (see section 4). Examples of choices for this nonlinearity are $g_1.x/Dx^2$; which measures the energy of the response, and $g_2.x/Dx^2$ in cosh x; which is a robust version of $g_1:A$ set of \Box lters with a large temporal response strength correlation is such that the same \Box lters often respond strongly at consecutive time points, outputting large (either positive or negative) values. This means that the same \Box lters will respond strongly over short periods of time, thereby expressing temporal coherence of a population code.

To keep the outputs of the \Box Iters bounded, we enforce the unit variance constraint on each of the output signals $y_k.t/$; that is, we enforce the constraint $E_tfy_k^2.t/g$ D $w_k^TC_xw_k$ D 1 for all k; where matrix C_x D $E_tfx.t/x^T.t/g$ Additional constraints are needed to keep the \Box Iters from converging to the same solution. Standard methods (Hyvärinen et al., 2001) are either to force the set of \Box Iters to be orthogonal or to force their outputs to be uncorrelated, from which we choose the latter. This introduces additional constraints $w_i^TC_xw_j$ D 0; i D 1; ...; K; j D 1; ...; K; j D i: These uncorrelatedness constraints limit the number of \Box Iters K we can \Box Ind so that K · N²: The unit variance constraints and the uncorrelatedness constraints can be expressed by the single matrix equation,

$$WC_xW^T D I$$
: (2.4)

Note that if we use a nonlinearity g.x/ D x^2 and 1 t D 0; the objective function becomes f.W/ D $_{\text{kD1}}^{\text{K}} \text{E}_{\text{t}} \text{f} y_{\text{k}}^4.\text{t/g}$ In this case, the optimization of the objective function under the unit variance constraint is equivalent to optimizing the sum of kurtoses of the outputs. Kurtosis is a commonly used measure in sparse coding. Similarly, in the case of nonlinearity g.x/ D In cosh x and 1 t D 0; the objective function can be interpreted as a nonquadratic measure of the nongaussianity of \square ter outputs. We return to this issue in section 3.

Thus, the receptive \square elds are learned in our model by maximizing the objective function 2.3 under the constraint 2.4. The optimization algorithm used for this constrained optimization problem is a variant of the gradient projection method of Rosen (for the original algorithm, see Luenberger, 1969). The optimization approach employs whitening, that is, a temporary change of coordinates, to transform the constraint 2.4 into an orthonormality constraint. Then a gradient projection algorithm employing optimal symmetric orthogonalization can be used. See the appendix for details.

3 Experiments on Natural Image Sequences _

3.1 Data Collection. The natural image sequences used as data were a subset of those used in van Hateren and Ruderman (1998). The original data set consisted of 216 monochrome, noncalibrated video clips of 192 seconds each, taken from television broadcasts. More than half of the videos feature wildlife; the rest show various topics, such as sports and movies. Sampling frequency was 25 frames per second, and each frame was block-averaged to a resolution of 128 £ 128 pixels. For our experiments, this data set was pruned to remove the effect of human-made objects and artifacts. First, many of the videos feature human-made objects, such as houses and furniture. Such videos were removed from the data set, leaving us with 129 videos. Some of these 129 videos had been grabbed from television broadcasts, and there was a wide black bar with height 15 pixels at the

top of each image, probably because the original broadcast had been in wide screen format. Our sampling procedure never took samples from this topmost part of the videos. If these artifacts were not removed from the data set, the static ICA results computed for comparison showed longer horizontal and vertical receptive Delds than results obtained from scenes without the artifacts. The Dnal, preprocessed (see below) data set consisted of 200,000 pairs of consecutive 11 $\mathfrak L$ 11 image windows (patches) at the same spatial position, but 1 t milliseconds apart from each other. Depending on the experiment, 1 t varied between 40 ms and 960 ms: However, because of the temporal Ditering used in preprocessing, initially 200,000 longer image sequences with a duration of 1 t C 400 ms; and the same spatial size 11 $\mathfrak L$ 11; were sampled with the same sampling rate.

A second data set was needed for computing the corresponding (static) ICA solution for comparison. This data set consisted of 200,000 11 \pounds 11 images sampled from the same video data.

3.2 Preprocessing. The preprocessing in the main experiment consisted of three steps: temporal decorrelation, subtraction of local mean, and normalization. (The same preprocessing steps were applied in the control experiments. Whenever preprocessing was varied in control experiments, it is explained separately below.) Temporal decorrelation can be motivated in two ways. First, it can be motivated biologically as a model of temporal processing at the lateral geniculate nucleus (Dong & Atick, 1995). Second, for 1 t D 0; the objective function can be interpreted as a measure of sparseness. Therefore, it is important to rule out the possibility that there is barely any change in short intervals in video data, since this would imply that our results could be explained in terms of sparse coding or ICA. To make the distinction between temporal response strength correlation and measures of sparseness clear, temporal decorrelation was applied because it enhances temporal changes. Note, however, that this still does not remove all of the static part in the video, an issue addressed in the control experiments that follow.

To investigate the effect of temporal decorrelation, we Drst examined the histogram of the distances between subsequent image windows separated by 1 t D 40 ms; shown in Figure 1A. (This histogram also shows that there are indeed large changes between subsequent time points even without temporal decorrelation.) The local mean has been removed from these windows, and they have been normalized (see below); note that 2 is maximal distance because of normalization. Temporal decorrelation was performed with a temporal Diter, shown in Figure 1B. The Fourier magnitude of the Diter was computed by inverting the amplitude spectrum of input data. A Wiener Diter approach was used in order not to amplify high-frequency noise. (We determined the Diter directly from data—see Dong & Atick, 1995—for an analytic solution.) Noise power was estimated by assuming that it is equal to signal power at 5:5 Hz; avalue also used in Dong and Atick (1995). Phases

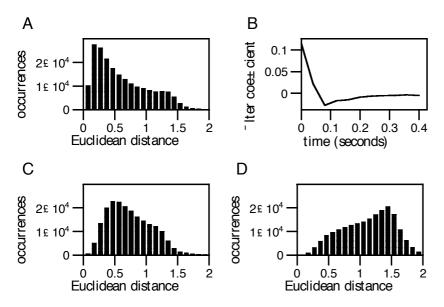


Figure 1: Temporal decorrelation enhances temporal changes. (A) Distribution of Euclidean distances between consecutive samples at the same spatial position, but 40 ms apart from each other, without temporal decorrelation. Note that two is the maximum value because of normalization. (B) The temporally decorrelating Illter. (C) Distribution of Euclidean distances between consecutive samples at the same spatial position, but 40 ms apart from each other, after temporal decorrelation. (D) Distribution of Euclidean distances between consecutive samples at the same spatial position, but 120 ms apart from each other, after temporal decorrelation. Note that removal of DC component and normalization were also performed in all of these cases.

were determined by adding a minimum energy delay constraint on the Ilter (see Oppenheim & Schafer, 1975). Filter length was originally 2500 ms but was truncated to 400 ms because this part contains over 99% of Ilter energy. When short image sequences of length 1 t C 400 ms are Iltered with this temporal Ilter, the resulting sequences are of length 1 t: From each of these temporally Iltered short sequences, two windows separated by 1 t were taken. Figure 1C shows the distribution of the distances between such temporally decorrelated windows, after removal of local mean and normalization (see below), when 1 t D 40 ms: This histogram shows that temporal decorrelation enhances temporal changes in the data. Figure 1D shows the distribution of the distances between temporally decorrelated windows, after removal of local mean and normalization, when 1 t D 120 ms: In this case, consecutive windows are, on average, quite far from each other when measured with the Euclidean norm. In fact, the peak of the histogram is

approximately at 1.4, which for normalized vectors means that a large part of the consecutive windows are approximately orthogonal to each other.

After temporal decorrelation, the local mean (DC component) was subtracted from each window. This reduces the number of dimensions of the data by one, so with image window size N D 11; the number of Elters K · 120: Finally, the sample vectors (vectorized windows) were normalized to have unit Euclidean norm, which can be considered a form of contrast gain control (Carandini, Heeger, & Movshon, 1997; Heeger, 1992). Note that no spatial low-pass Eltering or dimensionality reduction was performed during preprocessing.

In the case of the second data set, which was used to compute the corresponding static ICA solution, preprocessing consisted of removal of the local mean, followed by normalization. Temporal decorrelation was not performed here, since it has no meaning in the case of static image data.

3.3 Results: Temporally Coherent Filters of Natural Image Sequences. The main experiment consisted of running the symmetric gradient projection algorithm 50 times using different random initial values, followed by a quantitative analysis of these results, as well as results obtained with a corresponding ICA algorithm. In this experiment, 1 t was 40 ms: The number of extracted 🛘 Iters was set at the maximum value K D 120: Nonlinearity g in objective function 2.3 was chosen to be g.x/ D In cosh x because of its robustness against outliers (Hyvärinen et al., 2001).

Figure 2 shows the resulting \Box Iters (i.e., rows of matrix W) of the \Box rst run. The \Box Iters have been ordered according to $E_t fg. y_k. t//g. y_k. t_j$ 1 t//g. that is, according to their "contribution" into the \Box nal objective value (\Box Iters with largest values top left). The \Box Iters resemble Gabor \Box Iters. They are localized and oriented and have different scales. These are the main features of simple-cell receptive \Box elds (Palmer, 1999).

When compared qualitatively with earlier reported results, obtained with sparse coding and independent component analysis (Bell & Sejnowski, 1997; Olshausen & Field, 1996; van Hateren & van der Schaaf, 1998), our results show a larger variety of different spatial scales. To compare the results quantitatively, we extracted a corresponding set of 50 ICA separation matrices using the symmetric Dxed-point ICA (FastICA) algorithm, with robust nonlinearity tanh (Hyvärinen et al., 2001). This algorithm is a symmetric version of that used in van Hateren and van der Schaaf (1998). The symmetric nature of the algorithm facilitates extracting a balanced set of Diters (Hyvärinen et al., 2001). Filters maximizing temporal response strength correlation were compared against the rows of the separating matrix (the ICA Diters), since these Diters are the natural counterparts in ICA (van Hateren & van der Schaaf, 1998). Figure 3 shows the ICA Diters obtained from the Drst run.

In the quantitative comparison of the results, we measured the most important properties of the receptive Delds. The results are shown in Figure 4. The measured properties were peak spatial frequency (see Figures 4A and

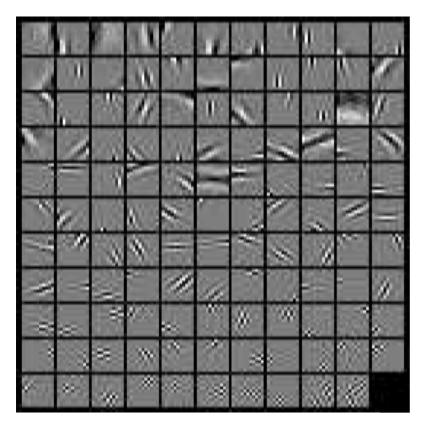


Figure 2: Temporally coherent \square Iters of natural image sequences, given by the \square rst run of the main experiment. The \square Iters were estimated from natural image sequences by optimizing temporal response strength correlation with the symmetric gradient projection algorithm (here nonlinearity g.x/ D In cosh x). The \square Iters have been ordered according to $E_tfg.y_k.t/g.y_k.t_i$ 1 t//g that is, according to their contribution into the \square nal objective value (\square Iters with largest values at top left).

4B; note the logarithmic scale and units cycles per pixel), peak orientation (see Figures 4C and 4D), spatial frequency bandwidth (see Figures 4E and 4F), and orientation bandwidth (see Figures 4G and 4H). See van Hateren and van der Schaaf (1998) for de⊡nitions of these measures. Although there are some differences, the most important observation here is the similarity of the histograms. This supports the idea that ICA =sparse coding and temporal coherence are complementary theories, in that both result in the emergence of simple-cell-like receptive □elds. A sfor the differences, the results obtained using temporal response strength correlation have a slightly

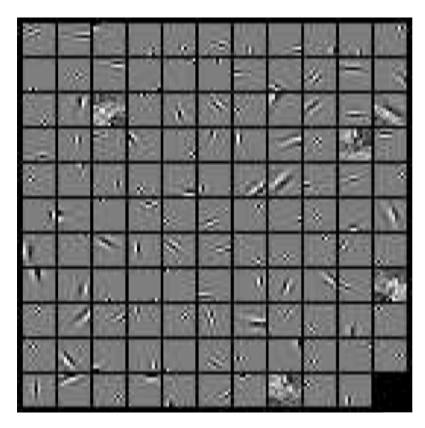


Figure 3: For comparison, ICA 🗆 ters estimated from natural image sequences by using the symmetric 🗆 xed-point algorithm. Note that these 🗀 ters have a much less smooth appearance than in most published ICA results; this is because for comparison, we show here the 🗀 ters and not the basis vectors, and further, no low-pass 🗀 tering or dimension reduction was applied in the preprocessing.

smaller number of high-frequency receptive Delds. Also, temporal response strength correlation seems to produce receptive Delds that are somewhat more localized with respect to both spatial frequency and orientation.

When the results are compared against the results in van Hateren and van der Schaaf (1998), the most important difference is the peak at zero bandwidth in Figures 4E and 4F. This difference is probably a result of the fact that no dimensionality reduction, antialiasing, or noise reduction was performed here, which results in the appearance of very small, checkerboard-like receptive Delds. This effect is more pronounced in ICA, which also explains the stronger peak at the 45 degree angle in Figure 4D.

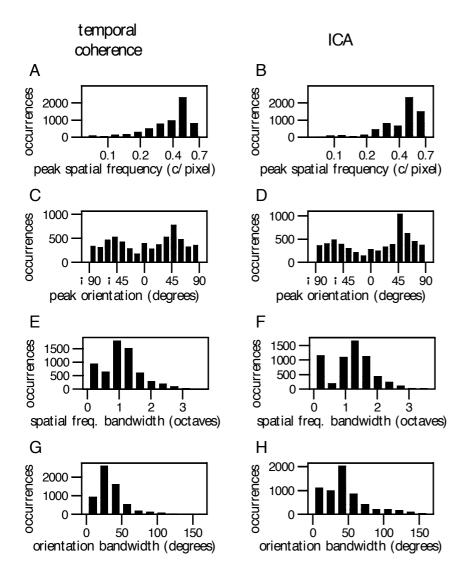


Figure 4: Comparison of properties of receptive Delds obtained by optimizing temporal response strength correlation (left column, histograms A, C, E, and G) and estimating ICA Diters (right column, histograms B, D, F, and H). See the text for details.

- 3.4 Control Experiments. To ensure the novelty and validity of our results, we made eight control experiments. All other aspects, except those speci□cally mentioned here, were similar in these experiments as in the main experiment.
- 3.4.1 Control Experiment I: No Temporal Decorrelation. In the main experiment, we used temporally decorrelated data. However, as was seen in Figure 1, there is considerable temporal change in natural video data even without temporal decorrelation. Therefore, it is reasonable to ask whether temporal decorrelation is necessary for achieving results like those shown in Figure 2. To answer this question, the algorithm was run for data that were not temporally decorrelated. The results are shown in Figure 5A. Although the Diters with highest-frequency components seem to be somewhat less localized, the results remain qualitatively very similar to those in Figure 2 in that they also resemble Gabor Diters. This suggests that the simple-cell-like properties of the results of the main experiment are not a consequence of temporal decorrelation.
- 3.4.2 Control Experiments II—IV: Longer 1 t. In the main experiment, consecutive samples were separated by 40 ms: Does the phenomenon found in the main experiment hold only for very small 1 t? To answer this question, we examined the case 1 t D 120 ms in control experiment II. As can be seen in Figure 1D, with this time separation, preprocessed consecutive sample windows are typically much farther from each other than when 1 t D 40 ms: Figure 5B shows the results of applying the symmetric gradient projection algorithm to this data. Although the receptive □elds are now larger than in the main experiment, the results are still qualitatively similar. This implies that the discovered relationship applies to short-time natural image sequences in general, not only for some speci□c value of 1 t:

It seems natural that there would be an upper limit for which the results are no longer qualitatively similar to those in Figure 2. This is indeed the case. The degree of spatial localization decreases when 1 t increases. This can already be seen to some degree in Figure 5B and is more pronounced in the results of control experiment III, in which 1 t D 480 ms (see Figure 5C). In the 480 ms case, most of the Diters are poorly localized, resembling Fourier basis vectors corresponding to different frequencies. As 1 t becomes large enough, spatial localization disappears, and Diters also start to lose their orientation selectivity. This is illustrated in Figure 5D for 1 t D 960 ms (control experiment IV).

3.4.3 Control Experiment V: Randomly Selected Consecutive Windows. Control experiment V was made to ensure that the results reDect the dynamics of natural image sequences, and not just the relationship between any arbitrary image patches. Instead of using samples separated by 1 t; random image samples were chosen as consecutive window pairs. No temporal

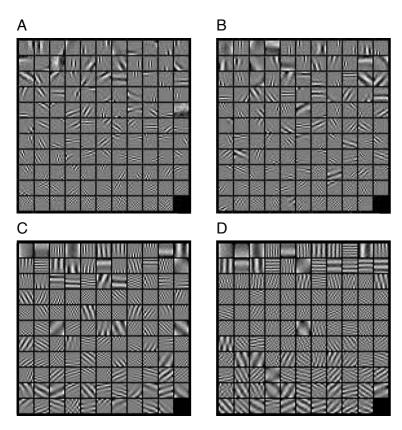


Figure 5: Results of control experiments I–IV. (A) Results of control experiment I in which no temporal decorrelation was performed. (B) Results of control experiment II in which 1 t D 120ms: (C) Results of control experiment III in which 1 t D 480ms: (D) Results of control experiment IV in which 1 t D 960ms:

decorrelation was done here, since random window pairs do not have a temporal relationship. Figure 6A shows the resulting spatial □lters, which correspond to noise patterns, indicating that the original results in Figure 2 do re□ect natural image sequence dynamics.

3.4.4 Control Experiment VI: Complete Removal of the Static Part of Video. In most experiments, the data were temporally decorrelated, leading to enhanced temporal change. For example, note from Figure 1D that in the case 1 t D 120 ms; after preprocessing (including the normalization step), the peak of the histogram of distances between preprocessed consecutive windows is at approximately 1.4. Remembering that the samples have been normalized, this indicates that a large part of the preprocessed sample con-

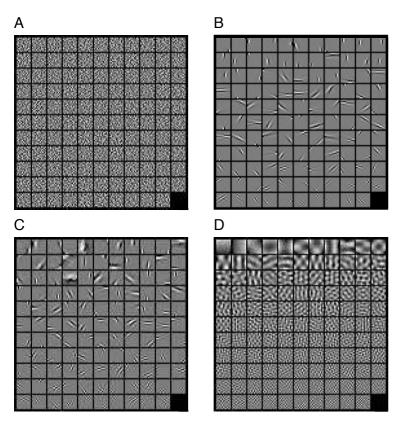


Figure 6: Results of control experiments V–VIII. (A) Results of control experiment V in which consecutive window pairs were chosen randomly. (B) Results of control experiment VI in which the static part of natural image sequences was removed altogether by employing Gram-Schmidt orthogonalization to consecutive windows. (C) Results of control experiment VII in which observer (camera) movement was compensated by using a tracking mechanism. (D) Results of control experiment VIII in which ordinary linear correlation between output values was maximized.

sists of consecutive windows that are approximately orthogonal to each other. This supports the idea that our results are not a consequence of the static part of natural image sequences.

However, since preprocessing does not remove the static part of the video completely, we made another control experiment (control experiment VI), with 1 t D 120 ms; in which the static part was removed altogether. This was done by modifying the preprocessing step so that no temporal decorrelation was done; instead, after removal of the local mean, consecutive windows

were orthonormalized using the Gram-Schmidt procedure. This removes the static part, the part present already at time t; 1 t; from the window at time t completely. No other form of temporal \square tering was performed in this experiment.

Figure 6B shows the results of this control experiment. The resulting Diters are still localized and oriented and have different scales. This shows that the qualitative nature of our results is not a consequence of the static part of natural image sequences. The Diters are more localized than in the corresponding experiment with temporal decorrelation, probably because removal of the static part is more likely to remove large features from the data (see section 4). If the same orthogonalization procedure is applied in the case 1 t D 40 ms; the results (not shown) are even more localized. This suggests that here too, as in the case of temporally decorrelated data, the degree of spatial localization is a function of 1 t:

3.4.5 Control Experiment VII: Compensation of Observer Movement. Control experiment VII was made to study the role of observer (camera) movement. To compensate for this movement, a simple correlation-based tracking mechanism was implemented into the sampling procedure. Tracking was applied before temporal Ditering (temporal decorrelation), so each 440 ms (D 1 t C 400 ms) sequence was tracked. Let x_1 be the Drst vectorized sample window in a 440 ms sequence, and x_n be the set of candidate windows in the nth video frame of the sequence, differing at most 10 pixels from the spatial position of the Drst window x_1 : The nth window x_n was chosen by x_n D arg max $_{x_2x_n} \frac{x_{n_1}^T x}{kx_{n_1} + kkxk}$: The results, shown in Figure 6C, are qualitatively similar to the original results, showing that the main results are not a consequence of observer movement. The number of low-frequency receptive Delds seems to be smaller in the control results. This change is probably caused by decreased large-scale movement (see section 4).

3.4.6 Control Experiment VIII: Ordinary Linear Correlation. Finally, the purpose of control experiment VIII was to show that higher-order correlation is indeed needed for the emergence of simple-cell-like Diters. To study this, we computed the optimal Diter solutions for maximizing linear correlation $f\cdot.w_k/$ D $E_tfy_k.t/y_k.t$; 1 t/g (see also Mitchison, 1991). The unit variance constraint is used here again, so the problem is equivalent to minimizing $E_tf.y_k.t/$ j $y_k.t$ j 1 t//²g with the same constraint. A

¹ Note that this objective function is de□ned for single □lters. A similar single-unit rule for optimizing $E_tfg.y_k.t'/g.y_k.t_i$ ¹ t'/gcan be de□ned and optimized for temporal response strength correlation. The optima for this objective function are similar to those in Figure 2, but the receptive □elds are more elongated. In addition, there are problems with obtaining a complete basis because a de□ationary algorithm (Hyvärinen et al., 2001), in which □rst-extracted solutions dominate, has to be used. In the case of linear correlation, we do not have this problem since a closed-form solution can be found.

closed-form solution can be derived from necessary Karush-Kuhn-Tucker (KKT) conditions and an additional zero DC constraint. Let C_x D ED E^T : from which the DC eigenvector corresponding to zero eigenvalue has been dropped out. The necessary conditions are fulllled by those eigenvectors of $ED^{\dagger} E^{T} E_{t} f. x. t/ j x. t j 1 t //. x. t/ j x. t j 1 t //^{T} g that correspond to nonzero$ eigenvalues. Analysis of suf□cient KKT conditions reveals that the □lters ful Illing the Irst-order conditions behave as in the case of principal components; that is, after the minimizing Diter is found and removed from the set of eigenvectors, another eigenvector from the set will be the next minimum, assuming that the output of the next selected later has to be uncorrelated with the outputs of the previously selected ones. The eigenvector corresponding to the smallest eigenvalue is the global minimum, and the next minimum is the one with the next smallest eigenvalue. Figure 6D shows the resulting Elters, sorted according to the corresponding eigenvalue (smallest eigenvalue top left). These Elters resemble Fourier basis vectors, and not simple-cell receptive □elds. Thus, we see that emergence of $localized\ receptive\ \square eld\ s\ requires\ the\ use\ of\ a\ nonlinear\ temporal\ correlation$ measure.

4 Discussion __

- 4.1 Temporal Coherence of Large Responses. Temporal response strength correlation, as delined by equation 2.3, does not explicitly measure the rate of change of the output signal. Therefore, it is important to examine what type of temporal coherence the objective function measures. In order to do this, consider three different temporally uncorrelated signals, y₁.t/; y₂.t/; and y₃.t/; depicted in Figure 7. (Note that in the main experiment, the output signals yk. t/ are temporally uncorrelated because they are spatially linearly Eltered from temporally decorrelated input data.) All of these signals have unit energy and a gaussian marginal distribution. Signals y_2 . t/ and y_3 . t/ have been created by reordering the samples of y_1 . t/ so that they contain two intervals of high amplitude (see the □gure caption for details). Signal y₃.t/ has the largest temporal response strength correlation of these signals, as measured by equation 2.3. This is because the objective function emphasizes the temporal coherence of large amplitudes. This is not true for an arbitrary measure of amplitude correlation. For example, for g.x/ D $\overline{|x|}$ (concave on interval]0; 1]), signal y_2 .t/ has a larger measure than $y_3.t/(f.y_2.t// \frac{1}{4} 0.73; f.y_3.t// \frac{1}{4} 0.71)$.
- 4.2 Temporal Coherence vs. Sparseness. We saw in the previous section that our objective function gives large values when both y_k . t/ and y_k . t/ 1 t/ have large amplitudes, thus emphasizing the correlation of large activations. This property must not, however, be confused with sparseness. Sparseness of y_k . t/ means that very large amplitudes, as well as very small ones, are relatively common. It is thus a property of the marginal distribution of

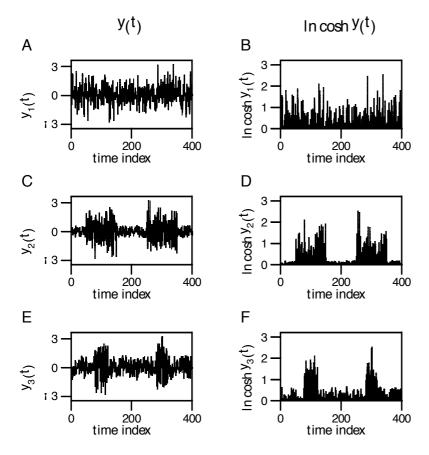


Figure 7: Temporal response strength correlation emphasizes temporal coherence of large amplitudes. Three temporally uncorrelated signals y.t/ with unit energy and gaussian marginal distributions (A,C,E) and their recti \Box ed magnitudes In cosh y.t/ (B,D,F). The signals in C and E were obtained by rearranging the time indices of the signal in A. This was done so that the two intervals of high amplitude in these signals contain the samples with the largest amplitudes, in random order. In C, the total length of the intervals is half of the total signal length; in E, this ratio is $\frac{1}{5}$: The signal in E has the largest temporal response strength correlation, as measured by equation 2.3 (f.y₁.t// ¼ 0:13; f.y₂.t// ¼ 0:23; f.y₃.t// ¼ 0:29). This illustrates the fact that the objective function emphasizes the temporal correlation of large amplitudes. See the text for discussion.

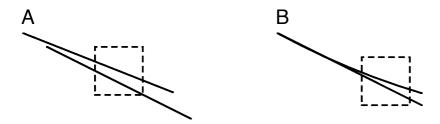


Figure 8: Examples of transformations inducing approximate local translations. The local area is marked with a dashed square. (A) Planar rotation. (B) Bending.

 y_k .t/: The temporal correlation of large amplitudes says nothing about their frequency or any other aspect of the marginal distribution of y_k .t/: All the signals in Figure 7 (on the left) have a gaussian marginal distribution, yet the signals vary considerably in their temporal coherence, as measured by our objective function.

Our measure of temporal coherence could indirectly measure the marginal distribution (sparseness) only where there is little change in the data, that is, if the static part of the image sequence dominates. However, this possibility was ruled out by the use of temporal decorrelation, which ensures that there is quite a large amount of change in the data, as shown in Figure 1. Ultimately, control experiment VI showed that the use of temporal coherence produced similar results even if the static part was removed completely, thus proving that the principle of temporal coherence is distinct from sparse coding.

4.3 An Intuitive Explanation of Results.

4.3.1 The Importance of Translation in Natural Image Sequences. The most universal local visual properties of objects are edges and lines, so we limit the discussion here to their dynamic properties. Objects can undergo a number of transformations in image sequences: translation, rotation, occlusion, and, for nonrigid objects, deformation. A transformation in the 3D space can induce a different transformation in an image sequence. For example, a translation toward the camera induces a change of object size in the image sequence. Our hypothesis is that for local edges and lines and during short time intervals, most 3D object transformations result in local translations of these elements in image sequences. This is, of course, true for 3D translations of objects. Figure 8 illustrates this phenomenon for two other transformations: a planar rotation and bending of an object. Note that the effect illustrated in Figure 8A is even more pronounced if object rotation is not purely planar.

4.3.2 Why Gabor-Like Filters Maximize Correlation of Square-Recti⊡ed Responses. In order to demonstrate the correlation of square-recti⊡ed responses at consecutive time points, we will consider the interaction of features and □lters in one dimension (orthogonal to the orientation of the □lter). This allows us to consider the effect of local translations in a simpli⊡ed setting. Figure 9 illustrates, in a simpli⊡ed case, why the temporal response strengths of lines and edges correlate positively as a result of Gabor-like □lter structure. Prototypes of two different types of image elements—the pro□les of a line and an edge—which both have a zero DC component, are shown in the topmost row of the □gure. The leftmost column shows the pro□les of three different □lters with unit norm and zero DC component: a Gabor-like □lter, a sinusoidal (Fourier basis-like) □lter, and an impulse □lter. The rest of the □gure shows the square recti□ed responses of the □lters to the inputs as functions of spatial displacement of the input.

Consider the recti ded response of the Gabor-like diter to the line and the edge. The squared response at timet; 1 t (spatial displacement zero) is strongly positively correlated with response at timet; even if the line or edge is displaced slightly. This shows how small local translations of basic image elements still yield large values of temporal response strength correlation for Gabor-like diters. If you compare the responses of the Gabor-like diter to the responses of the sinusoidal diter, you can see that the responses to the sinusoidal diter are typically much smaller. This leads to a lower value of our measure of temporal response strength correlation that emphasizes large values. Also, while the response of an impulse diter to an edge correlates quite strongly over small spatial displacements, when the input consists of a line, even a very small displacement will take the correlation to almost zero.

Thus, we can see that when considering three important classes of leters—leters that are maximally localized in space, maximally localized in frequency, or localized in both—the optimal leter is a Gabor-like leter localized in both space and frequency. If the leter is maximally localized in space, it fails to respond over small spatial displacements of very localized image elements. If the the leter is maximally localized in frequency, its responses to the localized image features are not strong enough.

Figure 10 shows why we need nonlinear correlations instead of linear ones: raw output values might correlate either positively or negatively, depending on the displacement. Thus, we see why ordinary linear correlation is not maximized for Gabor-like Diters, whereas the rectiDed (nonlinear) correlation is.

4.3.3 Emergence of Simple-Cell-Like Filters. Future research is needed to provide a detailed analysis of which properties of natural image sequence data are needed for the emergence of simple-cell-like □lters. However, at

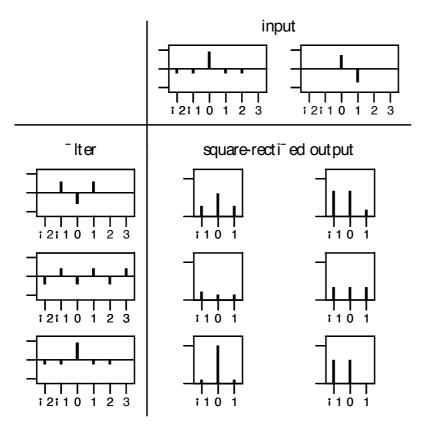


Figure 9: A simpli Ded illustration of why a Gabor-like Diter, localized in both space and frequency, yields larger values of temporal response strength correlation than a Diter localized only in space or only in frequency. Top row: cross sections of a line (left) and an edge (right) as functions of spatial position. Leftmost column: cross sections of three Diters with unit norm and zero DC component: a Gabor-like Diter (top), a sinusoidal Diter (middle), and an impulse Diter (bottom). The other plots in the Digure show the responses of the Diters to the inputs as a function of spatial displacement of the input. The Gabor-like Diter yields fairly large positively correlated values for both types of input. The sinusoidal Diter yields small response values. The impulse Diter yields fairly large positively correlated values when the input consists of an edge, but when the input consists of a line, even a small displacement yields a correlation of almost zero.

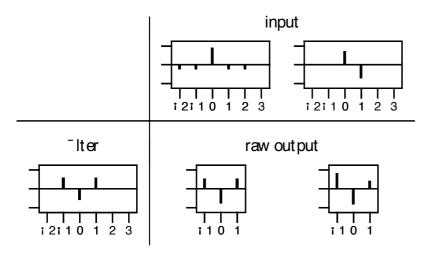


Figure 10: A simpli ded illustration of why nonlinear correlation is needed for the emergence of the phenomenon. Raw response values of the Gabor-like deter to the line and edge may correlate positively or negatively, depending on the displacement. (See Figure 9 for an explanation of the layout of the digure.)

this point, we can provide a set of hypotheses as to why oriented, localized Ulters with multiple scales emerge from the data:

Orientation. The 🗆 ters are oriented because the contours (edges and lines) in image sequences are oriented. Because our objective function emphasizes strong responses, the 🗀 ters need to be matched to the dominant features in the data (see Figure 9).

Localization. As illustrated in Figure 9, the features are limited in width because lines and edges are mostly limited in width as well, and because short-time translations are very local. This second point is supported by the results of control experiments II through IV (see Figures 5B–5D), which showed that when 1 t is increased, localization decreases. The features are limited in length for the same reason that they are limited in ICA or sparse coding: contours in real images have some curvature. Even a weak curvature makes the match (dot-product) with an elongated Gabor quite weak, and gives poor temporal coherence (Hoyer & Hyvärinen, 2002).

Multiple scales. The Ilters respond to multiple scales for two reasons. The Irst reason is the scale invariance of natural image sequences. This explanation is supported by the results of control experiment VI (Figure 6B), in which the static part of image sequences was removed completely. The results exhibit a narrower range of different scales because removal of the

static part of the sequences is more likely to remove large features. The second reason is that the features move with many different velocities in the data. This is supported by the results of control experiment VII (see Figure 6C), in which tracking was used to compensate for observer movement. The results exhibit a narrower range of different scales, because tracking is likely to reduce the velocity of moving features in the data.

4.4 The Case of Spatiotemporal Receptive Fields. Due to limited computational resources, we are currently unable to estimate the most temporally coherent spatiotemporal receptive □elds. However, argumentation similar to the intuitive explanation provided above can be given to illustrate a similar phenomenon in the spatiotemporal case. Figure 11 illustrates a case in which a vertical line is moving in the image sequence. The response of a simple-cell-likemotion-selective spatiotemporal □lter (DeA ngelis, Ohzawa, & Freeman, 1993a), whose spatiotemporal position and orientation match the initial position of the line and its direction of movement, is large in magnitude at consecutive time points. This illustrates how large temporal response strength correlation could arise in the case of spatiotemporal receptive □elds.

4.5 The Case of a Nonlinear (Nonnegative) Cell Model. Linear □lters with negative coef□cients or negative-valued data have signed outputs. Their widespread use as simple-cell models is often motivated by an interpretation in which the term simple cell in fact refers to a unit of two cells. These two positive-output cells, with reversed polarities, are modeled using a single linear □lter with signed output.² This two-cell approach will be explained in detail below. At this point, notice that this coupling is very different from complex-cell pooling of simple cells. In complex-cell pooling, the coupled cells respond to similar features at different spatial positions, whereas two cells with opposite polarities respond to similar features at the same spatial position.

For single simple cells, a more realistic basic model of the mapping from input to output via a receptive \square eld is a combination of linear \square ltering and a nonlinearity called half-wave recti \square cation (Heeger, 1992). Using the same notation as above, the output of the cell, y_k . t/; is computed by

$$y_k.t/D maxf0; w_k^T x.t/g,$$
 (4.1)

instead of the purely linear input-output relationship, equation 2.1. In this model, the output of a cell is never negative.

² Another possibility would be to consider the negative and positive values as changes from maintained □ring rate. However, simple cells have a low maintained □ring rate, which makes this approach undesirable (Heeger, 1992).

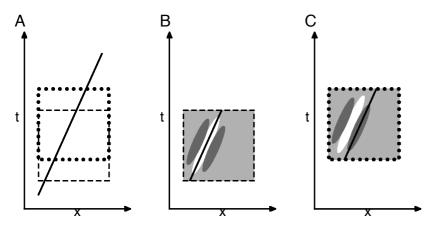


Figure 11: An illustration of how temporal response strength correlation could be exhibited by the outputs of simple-cell-like spatiotemporal receptive □elds. A phenomenon analog to translation in the spatial case can be observed in the spatiotemporal case. Let x and y denote the horizontal and vertical spatial coordinates, respectively, and let t denote the temporal coordinate. (A) The spatiotemporal trace (solid line) of a moving vertical line is shown here in the x-t coordinate system. The plot is similar for all y-coordinates because the moving line is vertical. Two different overlapping spatiotemporal input windows, separated by a small time difference, are also marked: one with dashed line and the other with dotted line. (B) A simple-cell-like spatiotemporal receptive \square eld, with position and orientation that match the initial position of the line and its direction of movement, responds strongly to the moving line. Here the spatiotemporal Diter has been superimposed over the dashed temporal window. White indicates large, positive values in the 🗆 ter, dark indicates large negative values, and middle gray indicates zero values. (C) When the same spatiotemporal receptive Deld, at the same spatial position, is applied to the same input a moment later (dotted spatiotemporal input window), the response is still strong, but the sign changes. Therefore, the temporal response strength correlation of the outputs of the simple-cell-like spatiotemporal receptive □eld would be large for this kind of input.

is de □ned as

$$y_{k}.t/D y_{k;1}.t/j y_{k;2}.t/$$
: (4.2)

It is straightforward to show that this model is equivalent to a purely linear model, that is, that $y_{k;1}.t/j$ $y_{k;2}.t/D$ $w_kx.t/$:

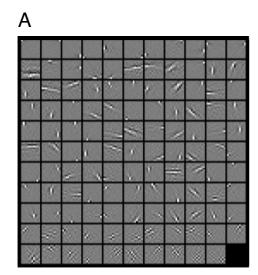
Therefore, one might interpret our model as measuring the temporal coherence of this two-cell unit, where the cells have similar receptive Delds with reversed polarities. A large value of the objective function indicates that either of the two cells responds strongly to the input. As mentioned above, using the linear model is a common approach. The same model is used, for example, in Bell and Sejnowski (1997), Olshausen and Field (1996), and van Hateren and van der Schaaf (1998), which makes the comparison of results obtained with these different models feasible.

However, a natural question is whether the same principle applies to the outputs of individual, half-wave recti ded simple cells. To address this issue, we computed the optimal solution for half-wave rectided cell outputs, that is, replaced the original linear input-output relationship (see equation 2.1) with the half-wave rectided relationship (see equation 4.1), and computed the optimal solution for this model. The same constraint (see equation 2.4) was used in this case for simplicity. Instead of using the temporally decorrelated data set, we used the orthonormalized data set with 1 t D 120 ms; also used in control experiment VI (see section 3.4.4). This is because by using the orthonormalized data set, we excluded the possibility of obtaining simple-cell-like receptive delds because of the static part of the video. This is important here since the static part might yield positive responses at consecutive time instances.

The results of this experiment are shown in Figure 12A. Although the features are not as well de □ned as before, the resulting □lters still show orientation, localization, and different spatial scales.

What is the reason for the emergence of such features even in this case when the negative part of the linear later response has been discarded and the static part of the video has been removed completely? First, as in the purely linear case, the objective function emphasizes the temporal correlation of large responses, and oriented and localized laters respond strongly to lines and edges. But how does temporal correlation arise? An illustration of a possible explanation is shown in Figure 12B. When a Gabor-like later is applied to a line, the half-wave rectiled output still correlates positively over time, but more weakly and over longer spatial displacements.

Finally, let us note that some form of temporal coherence of simple-cell outputs is implicitly assumed in many studies. The \Box ring rate of a simple cell is typically assumed to code for the output of a linear \Box lter, possibly after some simple nonlinear transformations, such as half-wave recti \Box cation. This requires that the output of the linear \Box lter has some temporal coherence. If the output of the linear \Box lter changes quite randomly, the \Box ring rate cannot



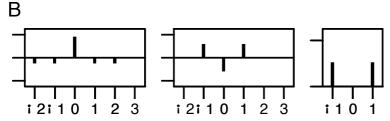


Figure 12: Temporal response strength correlation for a half-wave recti ded (non-linear, nonnegative) cell model. (A) The results of running the algorithm for a nonlinear cell model with half-wave rectidation of cell output. (B) When a simple-cell-like diter is applied to an input containing a moving line, the half-wave rectided output is still correlated positively over time, but the correlation is weaker, and the spatial displacement needed for the correlation is larger. Cross sections of a line (left) and a diter (middle), and the half-wave rectided output (right). See also Figure 9.

provide much information on the output, because then the \Box ring rate is very noisy when computed over short time intervals. Thus, maximization of temporal coherence could have a relation to maximization of the ef \Box ciency of a code based on \Box ring rates.

4.6 Implications of Our Results. In this article, we have shown that simple-cell-type receptive □elds maximize temporal response strength correlation at cell output when the input consists of natural image sequences. Temporal response strength correlation, or temporal correlation of recti-

□ed cell outputs, can be considered as a measure of temporal coherence. Our □ndings have several important implications. First, temporal coherence provides an alternative or complementary theory to sparse coding as a computational principle behind the formation of simple-cell receptive □elds. The application of either one of these principles results in the emergence of simple-cell properties from natural data.

Second, Földiák and others (Földiák, 1991; Kayser et al., 2001; Kohonen, Kaski, & Lappalainen, 1997; Wiskott & Sejnowski, 2002) have proposed that invariant visual representations, such as those found in complex cells, can be found by maximizing temporal coherence. (For an alternative complex-cell model using sparse coding, see Hyvärinen & Hoyer, 2001.) Our results show that this principle is applicable to the visual system even on the level of simple cells, which usually are not considered as invariant detectors. Although in some complex-cell models (Kayser et al., 2001; Kohonen et al., 1997) simple-cell receptive Delds are obtained as by-products, the learning is strongly modulated by the complex cells and therefore is very different from our model, which considers only the statistics of simple-cell outputs. Moreover, the simple-cell receptive Delds learned in Kayser et al. (2001) and Kohonen et al. (1997) do not seem to have the important properties of spatial localization and multiresolution (different scales).

Furthermore, whereas most earlier research results linking temporal coherence and properties of visual system have been based on theoretical considerations and simulated data, the results published in this article have been computed from natural image sequence data. To our knowledge, this is the Drst time that localized and oriented receptive Delds, with different scales, have been shown to emerge from natural data using the principle of temporal coherence. A step like this is important for any theory that tries to explain the structure and functionality of sensory neural networks using the statistical properties of natural input data.

Appendix: Details of the Symmetric Gradient Projection Algorithm ____

This appendix gives a detailed description of the optimization algorithm. First, the maximization of objective function 2.3 under constraint 2.4 can be made easier by employing whitening, a temporary change of coordinates, so that constraint 2.4 is transformed into an orthonormality constraint. Let equation C_x D EDE^T denote the eigenvalue decomposition of matrix C_x : If an eigenvalue of C_x is zero, then that eigenvalue can be dropped out of eigenvalue matrix D; and the corresponding eigenvector can be removed from eigenvector matrix E: This was the case in our experiments because preprocessing reduced the dimensionality of input data x.t/ by one when the DC component was removed from data. This dimensionality reduction also means that the number of \Box Iters that can be extracted is $K \cdot N^2$; 1:

De[□]ning matrix

$$U D WED^{1=2}; (A.1)$$

constraint 2.4 can be expressed as an orthonormality constraint,

$$UU^{\mathsf{T}} \mathsf{D} \mathsf{I}; \tag{A.2}$$

This simpler constraint will be easier to handle in the optimization algorithm below.

To express objective function 2.3 using the same transformed \Box Iter matrix U; we have to solve equation A.1 for matrix W: This is equivalent to solving matrix equation $E^TD^{1=2}W^T$ D U^T : In the case where the DC component has been removed, this is an underdetermined set of linear equations. By imposing the additional zero DC constraint on the \Box Iters in rows of W; the solution is given by

$$W D UD^{\dagger}^{1=2}E^{T}$$
 (A.3)

(see Hurri, 1997, for details). Substituting this into equation 2.2 gives

y.t/ D UD
i
 1=2E T x.t/ D Uz.t/; (A.4)

where signal vector $z.t/DD^{i-1=2}E^Tx.t/$: This transformation from input data x.t/t to z.t/t is called PCA whitening (Hyvärinen et al., 2001).

The above shows that after input data x.t/ are whitened, we can optimize

f.U/D
$$E_t fg.y_k.t//g.y_k.t$$
; 1 t//g; (A.5)

where output y.t/ is given by equation A.4, with respect to orthonormality constraint A.2. When the solution to this problem is found, the solution to the original problem is given by equation A.3.

The actual optimization algorithm used for this constrained optimization problem is a variant of the gradient projection method of Rosen (for the original algorithm, see Luenberger, 1969). Let ® m/ be a nonnegative decreasing sequence of real numbers, which converges to zero (initial step length ® 0/ is changed adaptively to speed up convergence). Let U 0/ be a random orthonormal matrix, and let U n/ be the value of matrix U at iteration n: The algorithm \Box nds a new candidate point by projecting matrix U n/ C ® m/ $\frac{\text{df.U.n//}}{\text{dU}}$ onto the constraint surface de \Box ned by equation A 2. If the candidate point is not an improvement, m is increased by one to \Box nd a new candidate point.

The critical step in the algorithm is the projection onto the constraint surface. This is achieved by optimal symmetric orthogonalization. Let A

be a square matrix with full rank. Then B D A.A^TA/i ¹⁼² is the nearest orthonormal matrix to A with respect to Frobenius matrix norm (see Fan & Hoffman, 1955, theorem 1, for a generalized proof of this). This is exactly the property required to achieve the projection step.

The resulting algorithm for maximizing temporal response strength correlation (TRSC) is shown below. The input arguments of the program are whitened data vectors z.t/; convergence tolerance 2 ; and initial step length @.0/ (in this version @.m/ D $\frac{@.0/}{2m}$). A daptation of initial step length @.0/ is not included in this description. The algorithm assumes convergence when the objective function changes very little between successive steps. In our experiments, convergence tolerance 2 varied between $10^{i\ 3}$ and $10^{i\ 6}$: Note that denoting U D $[u_1\ \mbox{$\omega\omega_{\rm K}]T ; the kth row of $\frac{df.U/}{dU}$ is given by the transpose of

```
\frac{@!.U'}{@!_k} D \frac{@}{@!_k} E_t fg.y_k.t//g.y_k.t; 1 t//g
       D E_t f g^0.y_k.t//g.y_k.t_i 1 t//z.t/C g.y_k.t//g^0.y_k.t_i 1 t//z.t_i 1 t/g.
funct U D max TRSC.z.t/;2; ®.0//
   U.0/:D rand./ comment: A random initial starting point.
   U.0/:D symmetricOrth.U.0//
   fOld:D0
   n:D0
  while f.U.n//; fOld > 2
          fOld:D f.U.n//
          m :D 0 ±
          m:DmC1
          <u>end</u>
          \frac{\text{end}}{\text{U.n C 1/:D symmetricOrth U.n/C}} \pm \frac{2}{2^m} \frac{\text{dt.U.n//}}{\frac{\text{dl.U.n//}}{\text{dl.U.n/}}}
          n:DnC1
   <u>end</u>
   U:DU.n/
funct B D symmetricOrth. A/
   B:D A.A^{T}A/i 1=2
```

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References _____

- Becker, S. (1993). Learning to categorize objects using temporal coherence. In S. J. Hanson, J. D. Cowan, & C. L. Giles (Eds.), Advances in neural information processing systems, 5 (pp. 361–368). San Mateo, CA: Morgan Kaufmann.
- Becker, S., & Hinton, G. (1992). Self-organizing neural network that discovers surfaces in random-dot stereograms. Nature, 355(6356), 161–163.
- Bell, A., & Sejnowski, T.J. (1997). The independent components of natural scenes are edge Elters. Vision Research, 37(23), 3327–3338.
- Carandini, M., Heeger, D. J., & Movshon, J. A. (1997). Linearity and normalization in simple cells of the macaque primary visual cortex. Journal of Neuroscience, 17(21), 8621–8644.
- DeAngelis, G. C., Ohzawa, I., & Freeman, R. D. (1993a). Spatiotemporal organization of simple-cell receptive Delds in the catts striate cortex. I. General characteristics and postnatal development. Journal of Neurophysiology, 69(4), 1091–1117.
- DeAngelis, G. C., Ohzawa, I., & Freeman, R. D. (1993b). Spatiotemporal organization of simple-cell receptive Delds in the catts striate cortex. II. Linearity of temporal and spatial summation. Journal of Neurophysiology, 69(4), 1118–1135.
- Dong, D. W. & Atick, J. (1995). Temporal decorrelation: A theory of lagged and nonlagged responses in the lateral geniculate nucleus. Network: Computation in Neural Systems, 6(2), 159–178.
- Fan, K., & Hoffman, A. J. (1955). Some metric inequalities in the space of matrices. Proceedings of the American Mathematical Society, 6, 111–116.
- Field, D. (1994). What is the goal of sensory coding? Neural Computation, 6(4), 559–601.
- Földiák, P. (1991). Learning invariance from transformation sequences. Neural Computation, 3(2), 194–200.
- Heeger, D. J. (1992). Normalization of cell responses in cat striate cortex. Visual Neuroscience, 9, 181–198.
- Hoyer, P.O., & Hyvärinen, A. (2002). A multi-layer sparse coding network learns contour coding from natural images. Vision Research, 42(12), 1593–1605.
- Hurri, J. (1997). Independent component analysis of image data. Master thesis, Helsinki University of Technology. Available on-line: http://www.cis.hut.□/jarmo/publications/.
- Hyvärinen, A., & Hoyer, P. O. (2001). A two-layer sparse coding model learns simple and complex cell receptive □elds and topography from natural images. Vision Research, 41(18), 2413–2423.
- Hyvärinen, A., Karhunen, J., & Oja, E. (2001). Independent component analysis. New York: Wiley.
- Kayser, C., Einhäuser, W., Dümmer, O., König, P., & Körding, K. (2001). Extracting slow subspaces from natural videos leads to complex cells. In G. Dorffner, H. Bischof, & K. Hornik (Eds.), Arti⊡cial neural networks—ICANN 2001 (pp. 1075–1080). New York: Springer-Verlag.
- Kohonen, T., Kaski, S., & Lappalainen, H. (1997). Self-organized formation of various invariant-feature □lters in the adaptive-subspace SOM. Neural Computation, 9(6), 1321–1344.

- Luenberger, D. G. (1969). Optimization by vector spacemethods. New York: Wiley. Mitchison, G. (1991). Removing time variation with the anti-Hebbian differential synapse. Neural Computation, 3(3), 312–320.
- Olshausen, B. A., & Field, D. (1996). Emergence of simple-cell receptive Deld properties by learning a sparse code for natural images. Nature, 381(6583), 607–609.
- Olshausen, B. A., & Field, D. (1997). Sparse coding with an overcomplete basis set: A strategy employed by V1? Vision Research, 37, 3311–3325.
- Oppenheim, A., & Schafer, R. (1975). Digital signal processing. Upper Saddle River, NJ: Prentice Hall.
- Palmer, S. E. (1999). Vision science—Photons to phenomenology. Cambridge, MA: MIT Press.
- Simoncelli, E. P., & Olshausen, B. A. (2001). Natural image statistics and neural representation. Annual Review of Neuroscience, 24, 1193–1216.
- Stone, J. (1996). Learning visual parameters using spatiotemporal smoothness constraints. Neural Computation, 8(7), 1463–1492.
- van Hateren, J. H., & Ruderman, D. L. (1998). Independent component analysis of natural image sequences yields spatio-temporal □lters similar to simple cells in primary visual cortex. Proceedings of the Royal Society of London B, 265(1412), 2315–2320.
- van Hateren, J. H., & van der Schaaf, A. (1998). Independent component Diters of natural images compared with simple cells in primary visual cortex. Proceedings of the Royal Society of London B, 265(1394), 359–366.
- Wiskott, L., & Sejnowski, T. J. (2002). Slow feature analysis: Unsupervised learning of invariances. Neural Computation, 14(4), 715–770.

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