

1 Resilience to sensory variance in the primary visual 2 cortex

3 Hugo J Ladret^{1,2,*}, Nelson Cortes², Lamyae Ikan², Frédéric Chavane¹, Christian
4 Casanova², and Laurent U Perrinet¹

5 ¹Institut de Neurosciences de la Timone, UMR 7289, CNRS and Aix-Marseille Université, Marseille, 13005, France

6 ²School of Optometry, Université de Montréal, Montréal, QC H3C 3J7, Canada

7 *Corresponding author : hugo.ladret@univ-amu.fr

8 ABSTRACT

9 Our daily endeavors occur in a complex visual environment, whose intrinsic variability is fundamental to the way we integrate
information to make decisions. By processing myriads of parallel sensory inputs, our brain is theoretically able to compute the
variance of its environment, a cue which is known to guide our behavior. Yet, the neurobiological and computational basis of
such variance computations are still poorly understood. Here, we quantified sensory variance modulations of primary visual
cortex neurons, both in orientation tuning and in dynamics. We report two archetypal responses, one of which is resilient to
changes in variance and can co-encode the sensory feature and its variance, significantly improving the population encoding of
orientation. The existence of these variance-specific responses could be accounted for by a model of intracortical recurrent
connectivity, which furthers our understanding of how the brain performs computations on complex, naturalistic inputs.

10 Introduction

11 Selectivity to the orientation of visual stimuli is an archetypal feature of neurons in the mammalian primary visual cortex (V1)¹.
12 This filter-like property of V1 neurons has been historically described using low-complexity artificial stimuli, such as oriented
13 gratings². These artificial stimuli offer a clear hypothesis as to what neurons are responding to, but only probe for neural
14 correlates of single parameters in V1 (orientation, spatial frequency, ...), which constitutes an experimental limit to our
15 understanding of the rich cortical dynamics involved in natural vision³, which are driven not only by these local parameters, but
16 also by a global context⁴. In orientation space, this means that most of our fundamental knowledge of V1 is currently based on
17 responses to single orientations, rather than more naturalistic responses to many simultaneous orientations.

18 This knowledge gap is not trivial, since the variance between the oriented elements of natural images is a fundamental cue
19 on which our brain relies to produce coherent integration of sensory inputs and prior knowledge of the world^{5,6} in order to
20 drive behaviour⁷. Bayesian inference rules would even posit that low-variance inputs are processed through fast feedforward
21 pathways, whereas higher sensory variance relies on a slower, recurrent integration⁸. How the brain performs computations on
22 variance, however, is still a relatively unexplored terrain. V1 recordings have shown that single neurons undergo non-linear
23 tuning modulations as a function of their input's variance⁹ which can be used as a functional encoding scheme^{10,11}. These
24 results are well accounted for by historical models of recurrent cortical activity of V1^{12,13} and match psychophysical metrics of
25 human subjects^{14–16}. While local intra-V1 interactions are sufficient to encode orientation variance¹⁷, the quantification of
26 single neuron responses and their link to a population encoding of variance remains to be established.

27 Here, we used stimuli which match the orientation content of natural images¹⁸ to study the neural basis of variance processes
28 in V1. We provide a quantitative description of the variance/tuning relationship of single neurons, both in terms of orientation
29 selectivity and dynamical modulation, which falls into two archetypal responses to variance. We report that some neurons are
30 endowed with a degree of invariance to sensory variance, remaining well tuned to orientation even at very high levels of sensory
31 variance. We show that only these "resilient neurons" are performing a co-encoding of both orientation and variance, improving
32 the overall encoding of orientation distributions in V1. A well-established model of intracortical V1 recurrence can account for
33 the presence of these neurons, which matches their predominant locations within supragranular cortical layers. This fits notably
34 well with canonical Bayesian frameworks⁶, potentially establishing uncertainty computations as a new general role for local
35 recurrent cortical connectivity.

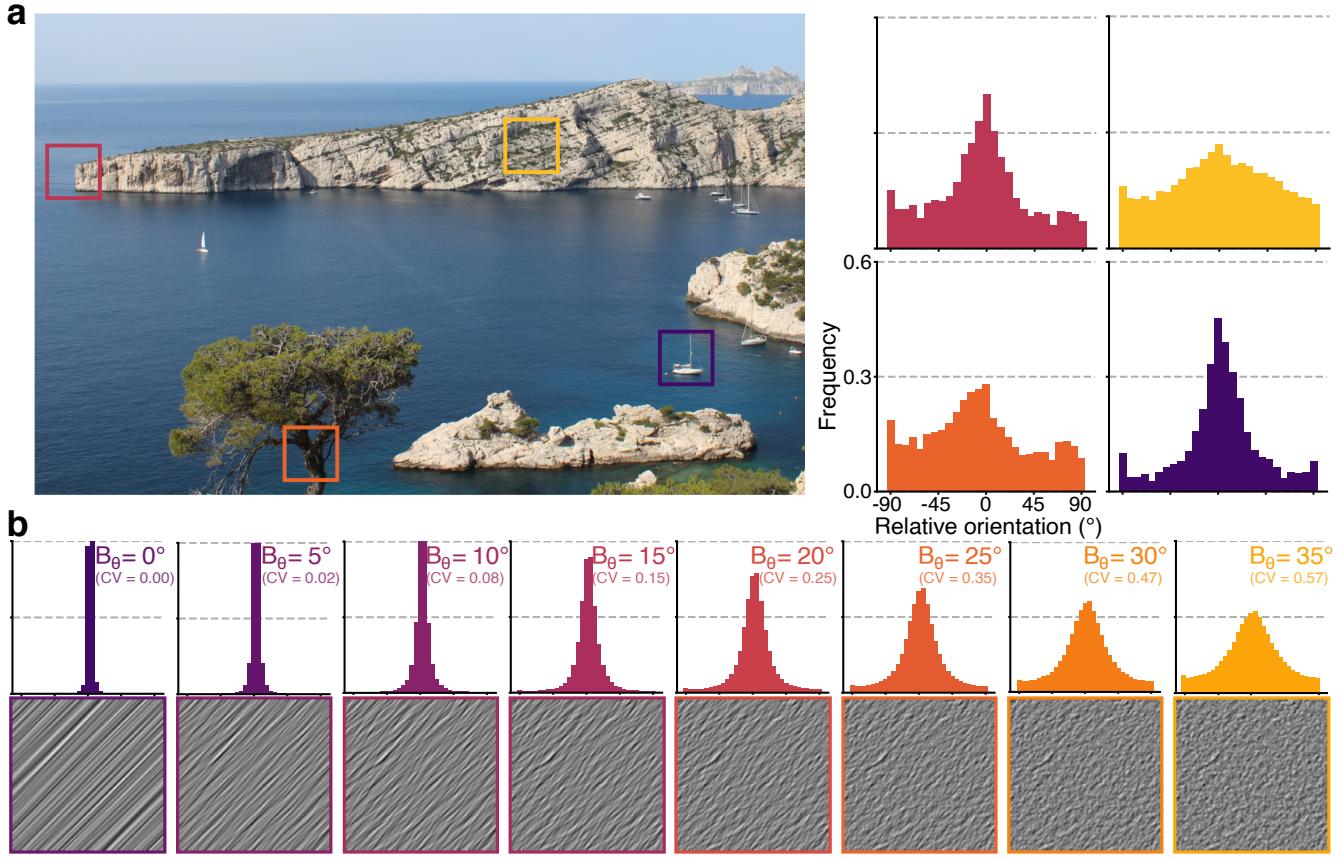


Figure 1. Variance of the orientation distributions characterizes local regions of natural images. **(a)** Distributions of orientation retrieved from four local regions of a natural image (picture taken by H.J.L) using an histogram of oriented gradients (32x32 pixels per cell, 200x200 pixels per region), centered around the most frequent orientation. **(b)** Motion Clouds, naturalistic stimuli (bottom row) oriented at an angle $\theta = 45^\circ$ relative to the horizontal axis, with increasing variance B_θ from left to right. Distributions of orientation retrieved from the stimuli (as done with the natural image) are shown on the upper row. Circular Variance (CV) of the distribution is shown to compare B_θ with a common orientation metric.

36 Results

37 Single neuron response in V1 depends on input variance

38 We recorded neural activity from 249 neurons in anesthetized cat V1, and measured orientation-selective response to naturalistic
 39 images called Motion Clouds¹⁸. These stimuli are band-pass filtered white noise textures that have three major advantages over
 40 both simple grating-like stimuli and complex natural images. First, they rely on a generative framework, which here allows to
 41 finely control both the mean θ and the variance B_θ of the orientation distribution in a stimulus, thereby closely reproducing
 42 the oriented content of natural images (Figure 1). Second, they are stationary in the spatial domain and thus only probe the
 43 orientation space, which removes any potential second-order information found in natural images which could be exploited
 44 by the visual cortex¹⁹. Third, they conform to the statistics of natural images, namely the $1/f^2$ distribution of the power
 45 spectrum²⁰, which gives them an overall ideal compromise between controllability and naturalness²¹. Here, 96 Motion Clouds
 46 were generated by varying the mean orientation θ between 0 and 180° in 12 even steps and the variance B_θ between $\approx 0^\circ$ and
 47 35° in 8 evenly spaced steps.

48 All neurons showed orientation selectivity to Motion Clouds, and almost all of them kept responding to their same preferred
 49 orientation when variance B_θ was increased (98.8% units, $p < 0.05$, Wilcoxon signed-rank test) while markedly diminishing
 50 in peak amplitude of tuning curve (95.1% units, $p < 0.05$, Wilcoxon signed-rank test, 73.1% mean amplitude decrease for
 51 $B_\theta = 36^\circ$), such that only 28.5% of the recorded units were still tuned for $B_\theta = 35.0^\circ$ stimuli ($p < 0.05$, Wilcoxon signed-rank
 52 test). Increments of input variance thus decrease single neuron tuning, which unfolded heterogeneously across neurons, as
 53 evidenced by two representative single units shown in Figure 2a. Neuron A is an example of a single unit which is no longer
 54 orientation-tuned when variance B_θ reaches 35° ($W = 171.0$, $p = 0.24$, Wilcoxon signed-rank test), contrarily to neuron
 55 B ($W = 22.5$, $p = 10^{-6}$) which exemplifies the aforementioned 28.5% of variance-resilient units. Both types of responses

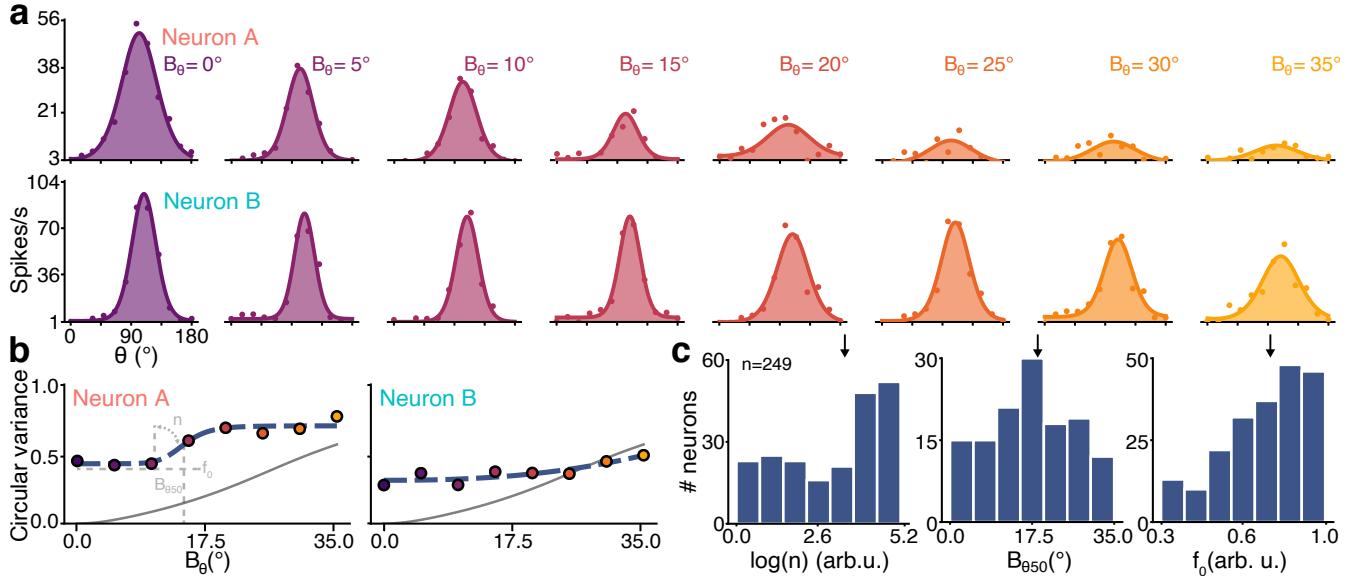


Figure 2. Single neuron tuning correlates with input variance to V1. Additional examples are provided in Figure S 1.

(a) Tuning curves of two example neurons responding to Motion Clouds of increasing variance B_θ from left to right. Dots represent the mean firing rate across trials (baseline subtracted, 300 ms average) and lines a fitted von Mises function.
 (b) Variance-tuning functions (VTF), assessing the change of orientation tuning measured by the circular variance (CV, dots) as a function of variance B_θ , fitted with a Naka-Rushton (NKR) function (dashed curves, parameters shown in light gray). Parameters of the VTF are $\log(n) = 8.4$, $B_{\theta50} = 14.7^\circ$, $f_0 = 0.4$ for neuron A and $\log(n) = 2.4$, $B_{\theta50} = 35.0^\circ$, $f_0 = 0.3$ for neuron B. The CV identity curve is shown in solid gray. (c) Histograms of the NKR parameters (in the [5%;95%] range of possible NKR fitting values) for the 249 recorded units. Median values are indicated by a black arrow ($\log(n) = 3.6$, $B_{\theta50} = 19.2^\circ$, $f_0 = 0.75$).

are further characterized by their B_θ to goodness of tuning (circular variance, CV) relationships. Such "variance-tuning functions" (VTF, Figure 2b) define an input/output ratio of variance in orientation space and were well fitted by a Naka-Rushton function²² (Figure S 2) and summarized the response to changes of sensory variance with three parameters: n , the degree of non-linearity of the VTF ; $B_{\theta50}$, the level of input variance at which a neuron's response transitioned from tuned to untuned state (i.e. the changepoint) and f_0 , the goodness of orientation tuning for inputs with the lowest variance. We also measured variance-bandwidth and variance-response functions (see supfig) which showed XXX These VTF revealed an heterogeneous response to variance between V1 neurons, with median values describing a prototypical VTF that is slightly non-linear with a changepoint at $B_\theta = 19.2^\circ$ (Figure 2c).

Orientation variance impacts not only orientation tuning, but also the dynamics of the response of V1 neurons (Figure 3). Interestingly, both effects are linked, as shown by the two example VTFs : example neuron B, which produced orientation-tuned responses for $B_\theta = 35^\circ$ inputs (Figure 2a), exhibited slower time-dependant change of goodness of tuning (relative min. of reduction of 42% of max. CV at 200 ms post-stimulation onset, $B_\theta = 0^\circ$) compared to neuron A (relative min. of 26% of max. CV at 90 ms post-stimulation onset, Figure 3b). These dynamical modulations were also heterogeneously distributed in the population, significantly more spikes emitted 200 ms after stimulation onset for $B_\theta = 35^\circ$ (Figure 3d, $U = 14936.0$, $p < 0.001$, Mann-Whitney U-test). Overall, orientation variance creates changes in both tuning and dynamics of V1 neurons with seemingly two archetypal types of response : either fast in time and non-linear with respect to variance (neuron A) or slow in time with linear changes of orientation tuning when variance increases (neuron B).

Multiple types of variance responses are found in V1

To properly characterize the two aforementioned types of responses to variance, we separated the recorded neurons in two groups, by K-means clustering the Principal Components (PC, Figure 4) of the neuronal responses. Rather than raw data, we performed clustering on the VTFs (Figure 4b), the statistical measurements of tuning (Figure 4c,d) and on the dynamics of response (Figure 4e,f). We used the first 2 PC for separating the data, which accounted for 39.1% of cumulative variance (Figure S 4a), and chose two clusters for the K-means algorithm, both with regard to the number of example responses shown before and due to the empirical absence of an "elbow"²³ in the Within-Clusters-Sum-of-Squares (WCSS) curve, which, when present, is typically used to select the number of clusters (Figure S 4b). This split the data into a cluster of 164 neurons, to which

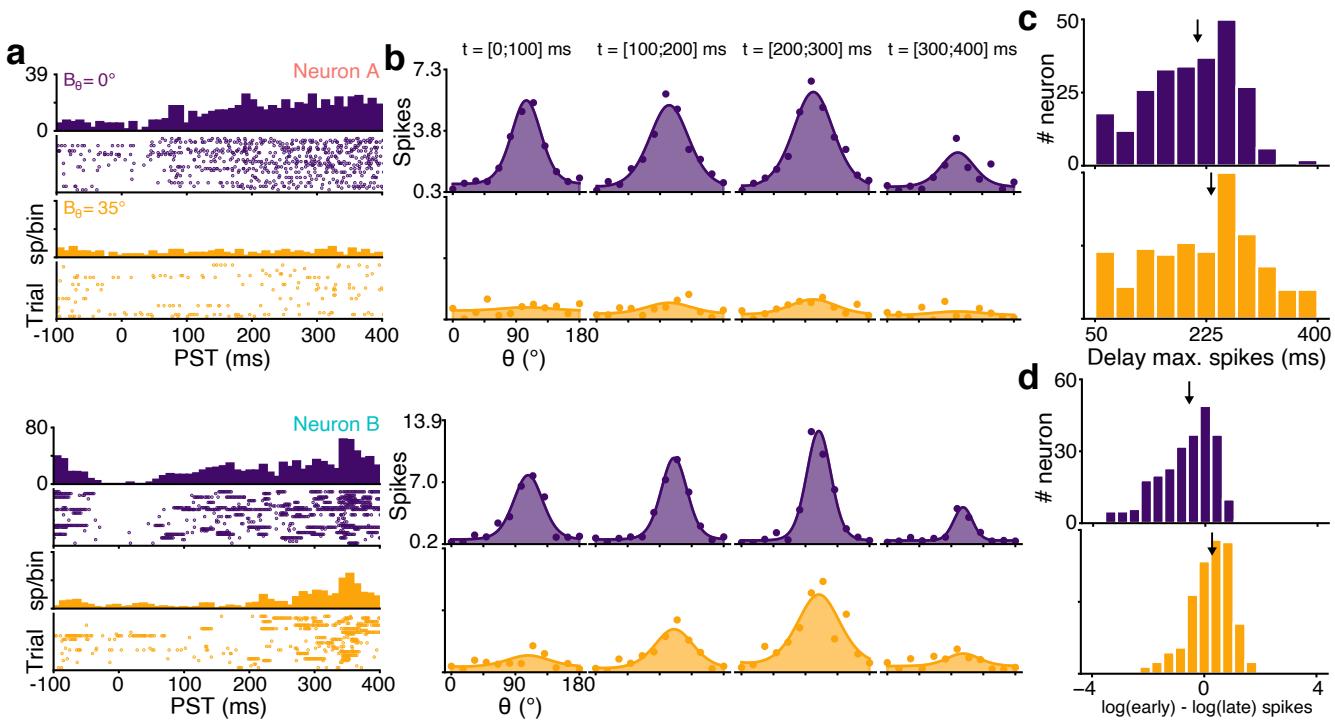


Figure 3. Dynamics of orientation response depend on input variance to V1. Additional examples are provided in Figure S 3. **(a)** Peristimulus time (PST) histogram and associated rasterplot of the two previous example neurons, for input variance $B_\theta = 0^\circ$ (purple) and $B_\theta = 35^\circ$ (yellow). **(b)** Dynamics of the tuning curves in 100 ms windows, starting at the time labelled atop of each column. **(c)** Delay to maximum peak amplitude of tuning curves for $B_\theta = 0^\circ$ (purple, median = 210 ms) and $B_\theta = 35^\circ$ (yellow, median = 233 ms) for the whole population. Median values are indicated by a black arrow. **(d)** Log ratio of early (< 100 ms post-stimulation) and late (> 200 ms) spike counts for $B_\theta = 0^\circ$ (median = -0.54) and $B_\theta = 35^\circ$ (median = 0.27) for the whole population.

belonged neuron A, whilst the rest of the data (85 neurons) was associated with the response types of neuron B. Given how this neuron displayed resilience to increase of input variance (Figure 2a), its cluster was labelled "resilient neurons". Conversely, the neurons clustered with neuron A were designated as "vulnerable neurons" (blue and red colors, respectively, Figure 4a).

The K-means clustering created a significant difference between the two groups' VTF parameters (Figure 4b) : resilient neurons had significantly more linear modulations ($\log(n)$, $U = 4029.0.0$, $p < 0.001$, Mann-Whitney U-test), higher change-points ($B_{\theta50}$, $U = 7854.0$, $p = 0.028$) and better tuning to low-variance inputs (f_0 , $U = 4992.0$, $p < 0.001$), which endows them with the ability to respond to orientation on a wider range of input variance^{24,25}. This is coherent with the clustering on the statistical measurement of orientation tuning, which showed that resilient neurons were still significantly tuned to higher values of B_θ ($B_{\theta max}$, Figure 4c, $U = 9155.0$, $p < 0.001$). However, when B_θ reached 35°, both groups of neurons had similar circular variance (Figure 4d). Taken together, this indicates that both types of neurons were similarly poorly tuned for inputs of highest variance, but underwent difference tuning changes between $B_\theta = 0^\circ$ and $B_\theta = 35^\circ$. In terms of dynamics, the two groups exhibited the same differences that characterized neuron A and B. Resilient neurons discharged significantly later than vulnerable neurons for $B_\theta = 0^\circ$ (Figure 4e, $U = 8455.5$, $p = 0.002$), but both group were on par for inputs of $B_\theta = 35^\circ$ ($U = 7794.5$, $p = 0.063$). Surprisingly, the time to maximum amplitude of the tuning curve was significantly lower in resilient neurons for $B_\theta = 0^\circ$ (Figure 4f, $U = 5542.5$, $p = 0.014$), which is the inverse of the early/late ratio of spikes. Neither group showed variance-dependent modulation of the delay to maximum spike ($U = 3058.0$, $p = 0.084$ and $U = 11545.5$, $p = 0.090$ for resilient and vulnerable, respectively), and both group had similar delay for $B_\theta = 35^\circ$ ($U = 6094.5$, $p = 0.158$).

The existence of these two groups of neurons could not be explained by an integration of the drifting motion of the stimuli (direction selectivity index, Figure 4g, $U = 7031.5$, $p = 0.910$). Rather, the location of the recorded units (unused in the clustering process) predominantly placed the resilient neurons in supragranular layers, which could provide a mechanistic basis for their existence (Figure 4h). Further, these neurons had sharper orientation tuning and slower dynamics, which are distinctive features of supragranular neurons^{26,27}. This, however, does not assign a functional role to these two types of responses in V1.

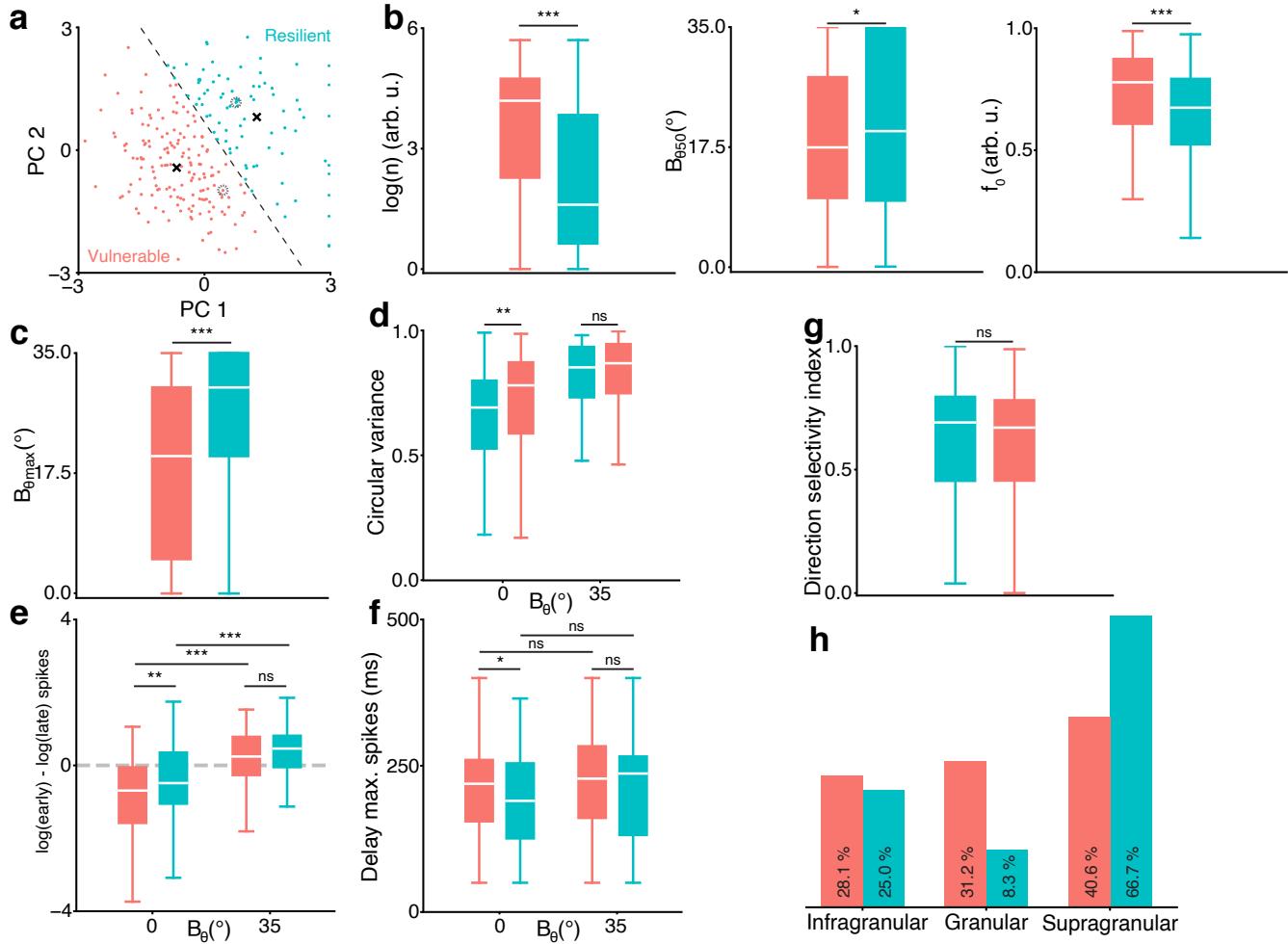


Figure 4. Responses to changes in variance fall into two categories. (a) Principal Components (PC) analysis of the data, K-Means clustered in two groups (2 clusters, centroids shown as black crosses and separatrix as dashed line). 9 resilient neurons with PC1 larger than 3 are plotted at PC1 = 3. Neuron A and B are shown as dashed circles. (b) Boxplot of the VTF parameters $\log(n)$, $B_{\theta 50}$, f_0 (ns, not significant; *, $p < 0.05$; **, $p < 0.01$, ***, $p < 0.001$ Mann-Whitney U-test). (c) Maximum B_θ at which a neuron yields a significant orientation tuning curve. (d) Circular variance for $B_\theta = 0^\circ$ and $B_\theta = 35^\circ$. (e) Log ratio of the early (< 100 ms) and late (> 200 ms) spike counts for $B_\theta = 0^\circ$ and $B_\theta = 35^\circ$. (f) Delay to the maximum peak amplitude of tuning curves for $B_\theta = 0^\circ$ and $B_\theta = 35^\circ$. (g) Direction selectivity index of the two groups (unused in the clustering). (h) Laminar position of the two groups (unused in the clustering).

Population level modulations of the orientation code

To understand the functional role played by resilient and vulnerable neurons, we used a neuronal decoder that probes for population codes in V1, which allowed understanding of what parameters of the stimuli each neuron group is encoding. We trained a multinomial logistic regression classifier²⁸, a probabilistic model that classifies data belonging to multiple classes (see Methods). This classifier was given the firing rate of neurons in a sliding time window (100 ms) and learned, for each neuron, a coefficient that best predict the class (i.e. the generative parameter θ , B_θ or $\theta \times B_\theta$) of the stimulus.

We used this decoder to probe for representation of the stimuli's orientations θ in the population activity. For this purpose, the dataset of trials was separated for each variance, such that 8 independent, B_θ -specific, orientation decoders were learned, with optimal parametrization (Figure S 5). These orientation decoders were able to retrieve the correct stimulus' θ well above the chance level (1 out of 12 orientations, max. accuracy = 10.56 and 4.68 times chance level for $B_\theta = 0^\circ$ and $B_\theta = 35^\circ$, respectively) from the entire population recordings. The temporal evolution of the accuracy of these decoders (Figure 5a) showed that maximally accurate orientation encoding correlates almost linearly with the stimuli's variance (Figure 5e, black). As the decoders are trained independently in each time window, this accumulative process occurs in the recordings themselves, rather than in the decoding algorithm.

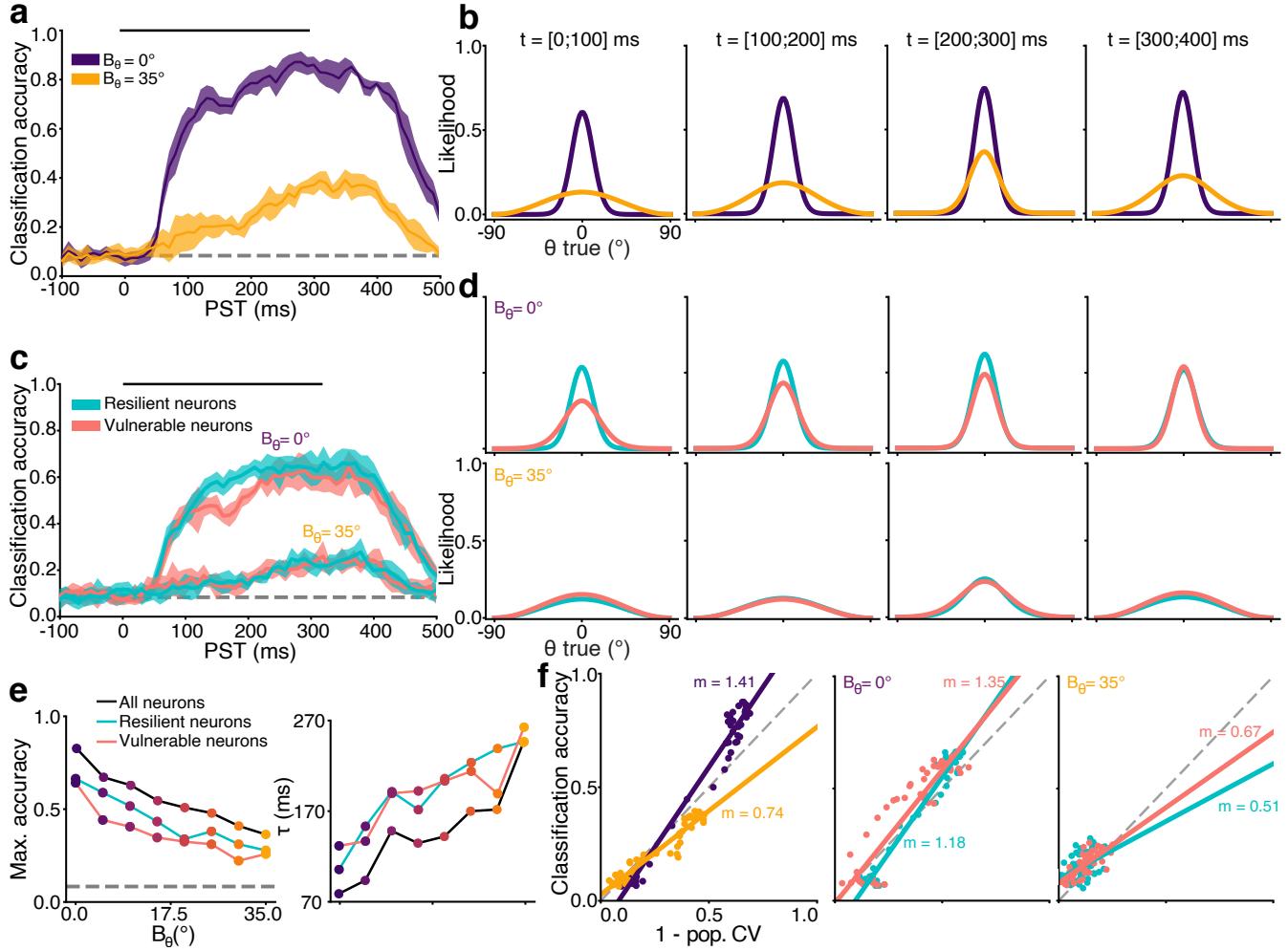


Figure 5. Input variance modulates orientation encoding in V1. (a) Time course of the accuracy for decoding orientation θ at two variances B_θ . Lines are the mean accuracy of a 5-fold cross validation and contour the STD. Decoding at chance level is represented by a gray dashed line and stimulation time by a black line. (b) Population tuning curves fitted with a von Mises function, showing the likelihood of decoding each θ in four time windows, centered around the correct θ . (c) Same as Figure 5a for the two groups of neurons. (d) Same as Figure 5b for the two groups of neurons, with $B_\theta = 0^\circ$ on the upper row and $B_\theta = 35^\circ$ on the lower row. (e) Parameters of the decoding time course the three decoders at all B_θ , estimated by fitting a sigmoid to decoding time course up to PST = 300 ms, where τ is the time constant. (f) Correlation between classification accuracy and population circular variance for the whole population (left), for both groups with $B_\theta = 0^\circ$ (middle) and $B_\theta = 35^\circ$ (right). Linear regression are shown as solid lines with slope m indicated (all significant, $p < 0.001$, Wald Test with t-distribution).

117 The full output of these decoders (see Methods) is a population tuning curve, which shows the likelihood of decoding all
 118 possible input classes (here, all θ , Figure 5b), rather than just the proportion of correct decoding reported by the accuracy metric.
 119 The clear correlation between the sharpness of these population tuning curves (Figure 5f left) and the accuracy of the decoder
 120 shows that improvements of decoding accuracy rely directly on a population-level separation of features within orientation
 121 space²⁸, and even more so at higher B_θ (Figure 5b, third panel). Overall, B_θ modulates the dynamics of the orientation code in
 122 V1. The short delay required to process precise inputs is congruent with the feedforward processing latency of V1²⁹, while
 123 the increased time required to reach maximum accuracy for low precision oriented inputs hints at the presence of a slower,
 124 recurrent, mechanism.

125 We then sought to assert the role of the vulnerable and the resilient neural populations by decoding θ from either group.
 126 The number of neurons in each group was imbalanced (79 more vulnerable neurons), which influences the accuracy of the
 127 decoder (Figure S 5), and we thus removed iteratively vulnerable neurons who had the least effect on maximum accuracy
 128 until both groups had the same number of neurons. This resulted in an overall reduction of only 1.5% maximum accuracy

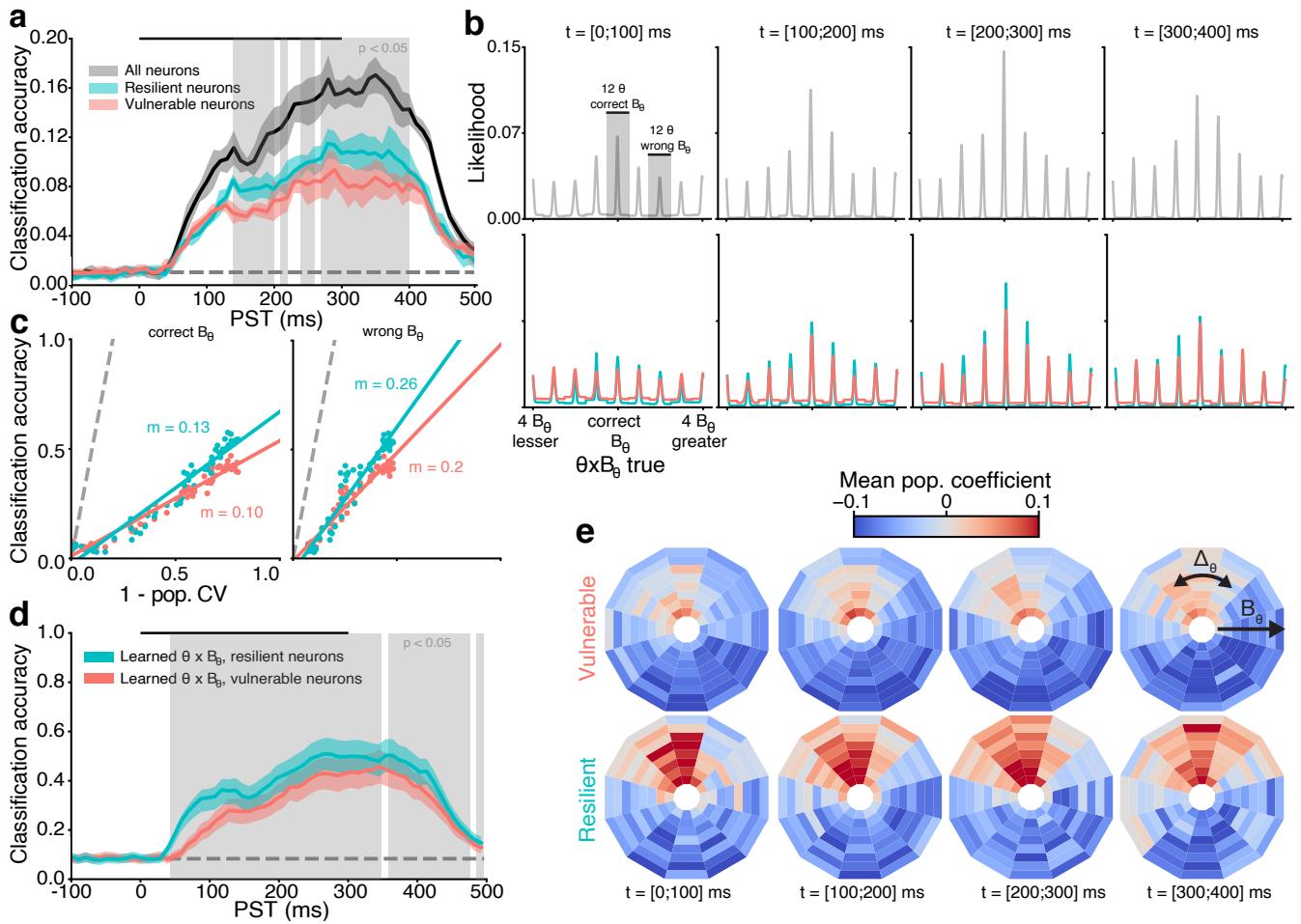


Figure 6. Resilient neurons encode both orientation and its variance. **(a)** Time course of the accuracy for decoding both the orientation and variance $\theta \times B_\theta$ of Motion Clouds. The solid dark line represents the mean accuracy of a 5-fold cross validation and the filled contour the SD. Significantly better decoding from resilient over vulnerable neurons is indicated as a grey overlay (1000 permutations test, $n=6$ cross validations, threshold $p < 0.05$). Decoding at chance level (here, 1/96) is represented by the gray dashed line. **(b)** Population tuning curves, showing the likelihood of decoding each $\theta \times B_\theta$ in four time windows, centered around the correct $\theta \times B_\theta$. Each of the 8 B_θ -specific tuning curve is fitted with an independent von Mises function. **(c)** Correlation between classification accuracy and population circular variance for correct B_θ population tuning curves (left) and averaged across other B_θ tuning curves curves (right). Linear regression are shown as solid lines with slope m indicated (all significant $p < 0.001$, Wald Test with t-distribution). **(d)** Time course of the $\theta \times B_\theta$ decoder, marginalized over B_θ to produce θ -only outputs. **(e)** Mean decoding coefficients of the two groups yielded from the whole population $\theta \times B_\theta$ decoder.

for the vulnerable group, which is thus an upper estimation of this group's decoding capacity. Using the same approach as with the global population decoding, we then trained B_θ -specific orientation decoders on the activity of either group of neurons. The overall decoding performance of both groups were similar (Figure 5c), with identical time courses but slightly (non-significant) better overall accuracy for resilient neurons (Figure 5e). Despite the better tuning of resilient neurons to inputs with higher variance (Figure 4), both groups have overall similar orientation encoding performances, even at $B_\theta = 35^\circ$. Interestingly, the population tuning curves (Figure 5d) reveal that resilient neurons have sharper population tuning curves when $B_\theta = 35^\circ$ (Figure 5f, right), but neither groups do not seem to have a different functional role in encoding orientation at the population level.

137 A subset of V1 neurons co-encode orientation and its variance

Given that orientation encoding did not account for a difference in functional roles of resilient and vulnerable neurons, we then probed for an encoding of the other parameter of stimulus generation, the variance B_θ . The same type of decoder previously used failed to infer the variance B_θ (chance level = 1 out of $8B_\theta$, max. accuracy = 1.91 times chance level) of the stimuli from

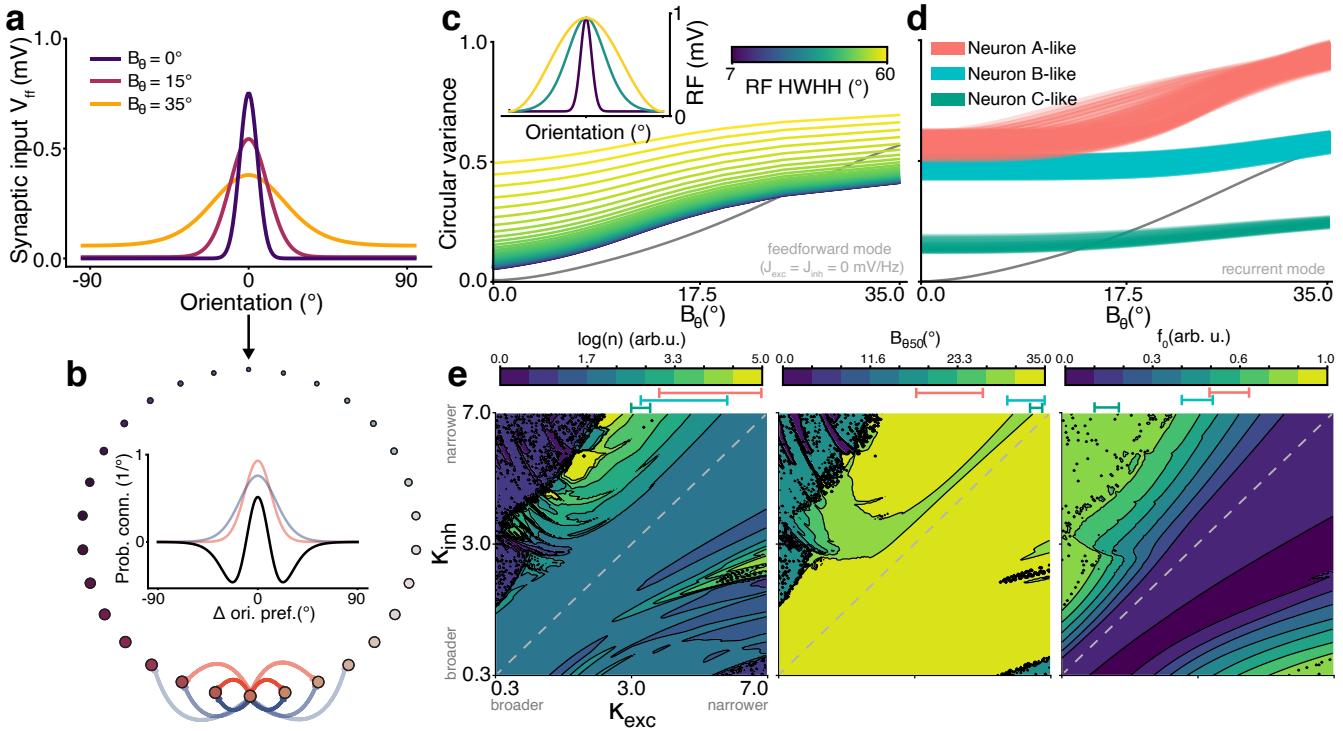


Figure 7. A model of recurrent interaction can explain the existence of resilient and vulnerable neurons. **(a)** Feedforward inputs V_{ff} to the neurons tiling the $[-90^\circ; 90^\circ]$ orientation space, for three values of B_θ . **(b)** Neurons are arranged in a ring topology, dictating the preferred orientation of each neuron. Inset : Recurrent connectivity profile for each neuron, computed as a difference (black) of excitatory (red) and inhibitory (blue) profiles, controlled by a measure of concentration κ_{exc} and κ_{inh} , respectively. **(c)** VTF of the model without recurrent connectivity (i.e. only inputs convolved with receptive fields) and with varying RF's Half-Width at Half-Height (HWHH). The CV identity curve is shown in black. Inset shows examples of receptive fields RF. **(d)** VTF of the model with recurrent connectivity, under two configurations retrieved by searching for VTF parameters close to those of neuron A, B (Figure 2b) and C (Figure S 1). **(e)** VTF parameters obtained from the model for each connectivity profile, shown as a contour plot of κ_{exc} and κ_{inh} . [5%;95%] range of the parameters of the VTFs shown in Figure 7 are displayed below the scale bars.

the population activity (Figure S 7a,b). At the single neuron scale, tuning curves flatten with increments of variance (Figure S 2a), which makes it hard to distinguish activity at the orthogonal orientation with $B_\theta = 0.0^\circ$ from the activity at the preferred orientation with $B_\theta = 35.0^\circ$. This variance decoding also failed to reach more than twice the chance level (max. accuracy = 1.72 and 1.71 times chance level for resilient and vulnerable neurons, respectively) in both resilient and vulnerable neurons (Figure S 7c,d). Thus, neither the encoding of variance B_θ nor the encoding of orientation θ accounts for a different role between resilient and vulnerable neurons.

Rather than assuming that V1 is an encoder of single parameter of a stimulus, a more realistic assumption is to think of the visual system's natural inputs as distribution of information (Figure 1) from which cortical neurons must process contexts over thalamic inputs³⁰, as part of a probabilistic-based computational principle³¹. Here, this means that the natural form of processing for a V1 neuron would co-encoding both the mean feature (θ) and its associated variance (B_θ).

We thus trained a $\theta \times B_\theta$ decoder, i.e. a decoder that retrieves both orientation and variance of the stimulus' simultaneously. This decoder correctly predicted orientation and variance with a maximum accuracy reaching 16.36 times the chance level (1/96, Figure 6a, gray). The particular likelihood structure (Figure 6b, upper row) showed that the correct θ was decoded with many concurrent hypothesis over B_θ . The progressive increase of accuracy stems from the emergence of a dominant encoding of θ at the correct B_θ , which reduces the relative magnitude of representations over other B_θ as time elapses. Interestingly, resilient neurons showed here a significantly different functional role from vulnerable neurons, with markedly better co-encoding of B_θ and θ (max. accuracy = 11.0 and 9.0 times chance level for resilient and vulnerable neurons, respectively, Figure 6a, blue, red). Both groups had similar ambiguity on B_θ (Figure 6b, lower row), and similar sharpening/accuracy ratios on the correct B_θ population curve (Figure 6c, left) or on the off-median population curves (Figure 6c, right).

To understand the usefulness of this co-encoding, we marginalized the decoder over B_θ , creating an orientation-only

encoder which learned both orientation and variance. Data from resilient neurons then provided significantly better encoding of orientation than vulnerable neurons (max. accuracy = 6.0 and 5.4 times the 1/12 chance level for resilient and vulnerable neurons respectively, Figure 6d, gray regions), demonstrating that the overall V1 orientation code improves when co-encoding variance. The difference between resilient and vulnerable neurons is further established by the decoder coefficients, which represent the contributions of each type of neurons towards the overall $\theta \times B_\theta$ code (Figure 6e, for single neuron examples see Figure S 8). Here, these coefficients are represented in the form a polar plot, where the error of the orientation classification Δ_θ is shown as the angle of each bin from the upper vertical and the variance B_θ is representing as the eccentricity of each bin from the center. Visualizing the coefficients of the whole population decoder (trained on the 249 neurons, Figure 6a, gray) shows that the output learned from resilient neurons co-informs about both orientation and variance, as observed by the extent of the bins in the eccentricity (B_θ) axis (Figure 6c, bottom row). Conversely, the decoding process yielded an orientation-only information from the activity of vulnerable neurons (Figure 6c, top row). Although the coefficients are learned independently at each time steps, the difference of information between the two groups of neurons is extremely stable through time.

Resilient neurons can thus co-encode both orientation and its variance simultaneously, while vulnerable neurons can only encode orientation. Given this functional difference between the two groups of neurons, we finally address the question of how both types of neurons can exist in V1.

Recurrent activity can explain the existence of neurons co-encoding orientation and variance

One difference between vulnerable and resilient neurons is their different location within the cortical layers (Figure 4h). This typically implies differences in local circuitry, particularly in the intra-V1 recurrent interactions between cortical columns that are mostly confined to supragranular layers³². Given that resilient neurons are predominantly found in these supragranular layers, we sought a mechanistic explanation for the existence of the two groups of neurons based on local interactions in V1. We derived a novel model from a well-established computational model of recurrent connectivity in V1, which was first used to account for the intracortical activity in cat V1³³ and later simplified as a center-surround filter in the orientation domain²⁷. This model has already accounted for a wide range of emerging properties in cortical circuits^{34,35}. Briefly, this model is built of orientation selective neurons tiling the orientation space and connected amongst themselves via recurrent synapses which follow an excitatory/inhibitory difference of von Mises distributions (Figure 7b). Here, we model inputs with higher variance as more spread in orientation space (Figure 1) and thus in model space (Figure 7a), which hence drives the recurrent dynamics of the model based on B_θ (for a full description, see Methods).

Given that feedforward connectivity with heterogeneous tuning can encode mixtures of orientations and natural images⁹, we first ran our model without recurrent synapses. We reproduced the heterogeneous selectivity by convolving the input with tuning curves of varying bandwidths (Figure 7d, inset). This feedforward mode of the network was only able to produce a limited number of responses (Figure 7d), in which increasing the bandwidth of the tuning curves increased the parameter f_0 of the VTF, but kept n and $B_{\theta50}$ constant.

Barring that explanation, we focused on the role of recurrent synapses and disabled the convolution of inputs. We varied the concentration parameters of the synaptic distributions κ_{inh} and κ_{exc} (Figure 7c) in 200 even steps ranging from 0.35 to 7, yielding 40000 possible configurations of the model. This allowed to manipulate the VTF and to accurately reproduce those of single neurons recorded in V1 (neuron A, B in Figure 2b and C in Figure S 1, modeled in Figure 7e). Varying the type of recurrence between neurons with different orientation preference allowed to reproduce all VTF found in V1. The parameter spaces (Figure 7f) showed a trend for resilient VTFs (low n , high $B_{\theta50}$, low f_0) to be found mostly around the $K_{exc} ; K_{inh}$ identity line, thus produced by a balanced recurrent connectivity. Vulnerable VTFs (high n , low $B_{\theta50}$, high f_0) were, on the contrary, mostly found above the identity line, where the configuration of the network is dominated by excitation over inhibition. Overall, recurrence within V1 neurons is sufficient to explain the existence of vulnerable and resilient neurons, and thus to explain the co-encoding of orientation and variance.

Discussion

The variance of oriented inputs to V1 impacts orientation selectivity⁹ and we have sought an explanation for how V1 could process this variance, from single neurons to population codes. We have shown that variance causes modulations in tuning (Figure 2) and dynamics (Figure 3) of single V1 neurons, which fall into two broad categories, labeled here vulnerable and resilient (Figure 4). Using a decoding approach, we uncovered variance-dependent accumulative dynamics in the two groups of neurons (Figure 5) that are directly tied to a population-level separation of features within orientation space²⁸. Both groups can encode orientation but not variance independently (Figure S 7), yet only resilient neurons are able accurately co-encode orientation and variance of the input to V1 (Figure 6). Based on cortical layer position and on a computational approach (Figure 7), we propose that the overall processing of variance in V1 is performed by recurrent connectivity between local cortical populations with different preferred orientations (Figure 8).

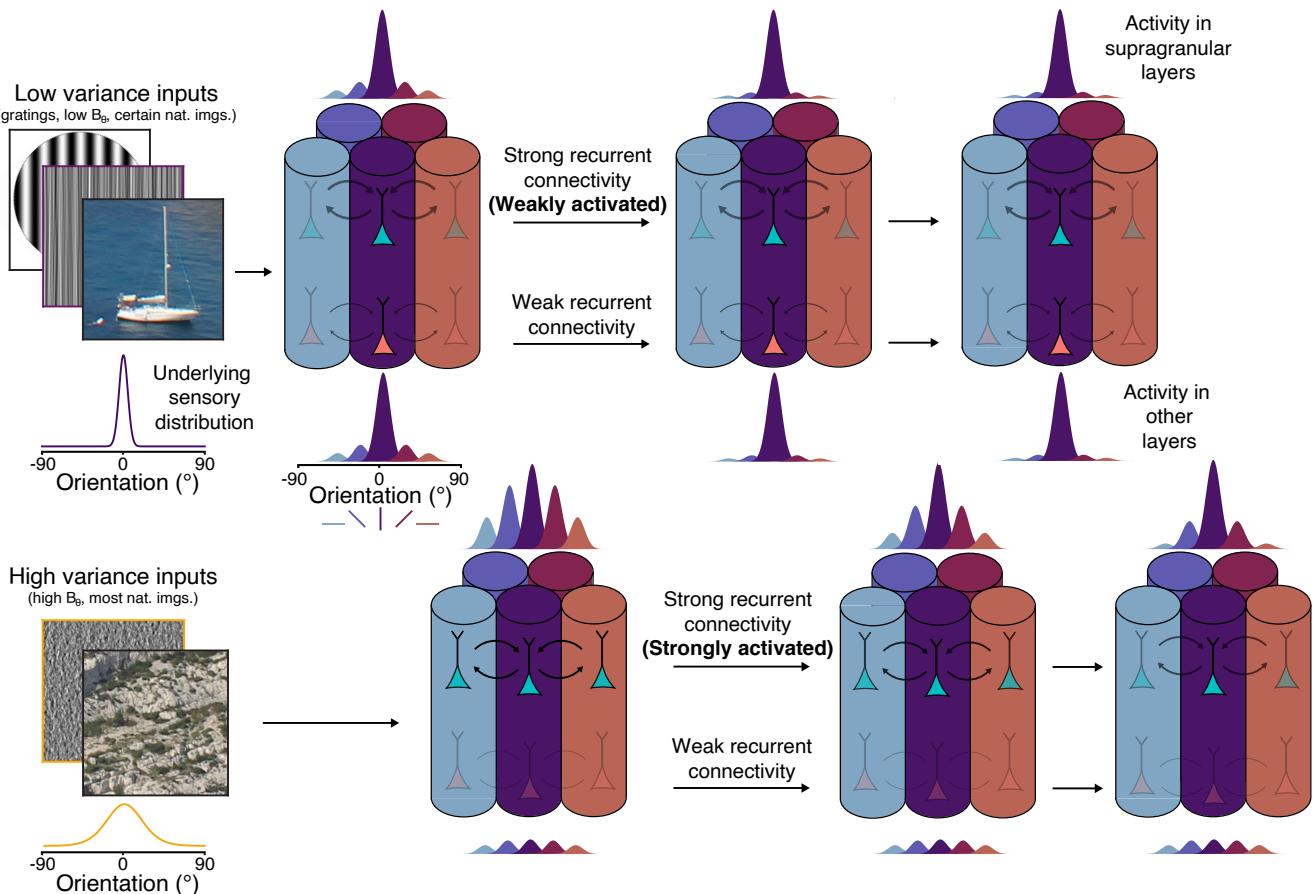


Figure 8. Summary of the findings. The top row is a representation of a set of orientation selective units (here, columns) processing low variance inputs, while the bottom row schematizes the processing of high variance inputs. In the case of low variance inputs to V1, the underlying sensory distribution is sharp in the orientation space, driving mostly a single orientation selective unit which processes orientation in fast feedforward manner. The feature encoded by this activity then stays stable through time (from left to right). For inputs of higher orientation variance, the sensory input is broadly distributed in the orientation space, which drives many dissimilarly tuned units, thus recruiting slow recurrent interactions. The quality of feature encoding progressively increases through time, as recurrent interactions perform computations to represent the most salient oriented feature in the input.

213 Here, we restricted our approach to orientation space, rather than investigating the full extent of spatial relationships which
 214 is present in natural images. As such, full-field stimuli with no second-order correlation were used, which, compared to a
 215 perfectly ecological environment, have likely excluded end-stopped cells³⁶. The time scales reported here might have been
 216 minimally slowed by anesthesia, here halothane, which nonetheless has a limited effect on V1^{37,38}, an area that is also usually
 217 less prone to anesthetics modulations^{39–41}, compared to higher-order areas⁴². Additionally, anesthetized V1 results⁴³ often
 218 correlate extremely well with later studies in awake animals⁴⁴.

219 In limiting our approach to orientation space, we also excluded extrastriate contributions to the variance modulations.
 220 However, there are both neurobiological and computational evidences that V1 alone is sufficient to explain the results, which is
 221 further supported by the domain literature and by theoretical considerations. In line with our results, recurrent excitatory and
 222 inhibitory synaptic connectivity is heterogeneous in V1^{45–48}, which sustains a resilient orientation tuning⁴⁹ that could very well
 223 account for the diversity of single neurons resilience under different connectivity configuration, as shown by our computational
 224 model (Figure 7). The temporal scales of local recurrent connectivity, namely the slowly-conducted horizontal waves in an
 225 orientation map⁴⁴, also fit the view of variance processing as an iterative and accumulative computation implemented by local
 226 recurrent interactions. Supragranular layers, which contained a majority of resilient neurons, are heavily connected through
 227 recurrent interactions with neighbouring cortical columns^{26,27,32,44}, reinforcing the idea that this type of connectivity could
 228 explain the different neuronal response to variance. On the computational side, we used a model specifically designed to
 229 produce intra-V1 data (and here, fitted to cat V1 data³⁵) and showed that it is sufficient to produce various VTF as found in our

recordings. This model relies on pooling information from multiple orientation-tuned units into a single neurons, which could also be viewed as a (flattened across recurrence) converging feedforward model. This would be a canonical model of a complex cell, whose computations can be equivalently implemented by a recurrent network⁵⁰. Motion Clouds are however aperiodic patterns and a modulation "F1/F0" ratio cannot be computed here directly.

Investigation of the electrophysiological basis of variance processing are few, but are all in agreement with local V1 processes. For instance, Goris *et al.*⁹, reported that heterogeneously tuned V1 (and also V2) populations help encode the orientation distributions found in natural images, which can be explained by an L-NL model. Consistent with their findings, we reported a variety of single-neuron variance modulations, that serve as a basis to decode orientation variance at the population level — a requirement for encoding natural images in V1⁵¹. Further analysis of their results has shown a "variance in / variance out" relationship in V1 neurons¹⁷, which is here implemented in the linear VTF neurons.

Theoretical considerations are also in favor of the claim that intra-V1 recurrence is sufficient to explain variance modulation. Bayesian inference , namely under the specific case of predictive coding³¹, canonically assigns (inverse) variance weighting of cortical activity to supragranular recurrent connectivity^{6,8}, without the need for extrastriate computations. This is an interesting perspective which opens up a general interpretation of our results into the broader context of processing variance/precision/uncertainty at different scales of investigations. Extending the present results to other cortical areas or other sensory modalities would be a simple process, given the generative stimulus framework used here¹⁸, which could yield pivotal new insights into our understanding of predictive processes in the brain.

Methods

Visual stimulation

Motion Clouds are generative model-based stimuli¹⁸ which allow for fine parameterized control over a naturalistic stimuli⁵², which is a desirable trait when probing sensory systems under realistic conditions²¹. They are mathematically defined as band-pass filtered white noise stimuli, whose filters in Fourier space are defined as a parameterized distribution in a given perceptual axis (here, only orientation, but can be extended to motion⁵³ and scale⁵⁴). Thus, the Motion Clouds presently used are fully characterized by their mean orientation and their orientation bandwidth, such that a given stimuli S can be defined as:

$$S = \mathcal{F}^{-1}(O(\theta, B_\theta)) \quad (1)$$

where \mathcal{F} is the Fourier transform and O the orientation envelope, characterized by its mean orientation θ and its orientation bandwidth B_θ . A total of 96 different stimuli were generated, with 12 mean orientations θ ranging from 0 to π in even steps, and 8 orientation bandwidth B_θ ranging from ≈ 0 to $\pi/5$ in even steps. The orientation envelope is a von Mises distribution:

$$O(\theta, B_\theta) = \exp \left\{ \frac{\cos(2(\theta_f - \theta))}{4 \cdot B_\theta^2} \right\} \quad (2)$$

where θ_f is the angle of the frequency components of the envelope in the Fourier plane, which controls the spatial frequency parameters of the stimuli, set here at 0.9 cycle per degree. The stimuli were drifting orthogonally in either direction with respect to the mean orientation θ at a speed of 10°/s, which is optimal to drive V1 neurons⁵⁵. For the range of values of B_θ considered here, the orientation envelope approximates a Gaussian distribution and B_θ is thus a measure of the variance of the orientation content of the stimuli.

All stimuli were generated using open-source Python code (see Additional information) and displayed using Psychopy⁵⁶. Monocular stimuli were projected with a ProPixed projector (VPixx Technologies Inc.) onto an isoluminant screen (Da-Lite[®]) covering 104°×79° of visual angle. All stimuli were displayed at 100% contrast for 300 ms, interleaved with a mean luminance screen (25 cd/m²) shown for 150 ms between each trial. Trials were fully randomized, and each stimulus (a unique combination of $\theta \times B_\theta \times$ drift direction) was presented 15 times.

Surgery

Experiments were conducted on 3 adult cats (3.6 - 6.0 kg, 2 males). All surgical and experimental procedures were carried out in compliance with the guidelines of the Canadian Council on Animal care and were approved by the Ethics Committee of the University of Montreal (CDEA #20-006). Animals were initially sedated using acepromazine (Atravet®, 1 mg/kg) supplemented by atropine (0.1 mg/kg). Anesthesia was induced with 3.5% isoflurane in a 50:50 mixture of O₂:N₂O (v/v). Following tracheotomy, animals underwent artificial ventilation as paralysis was achieved and maintained with an intravenous injection of 2% gallamine triethiodide (10 mg/kg/h) diluted in a 1:1 (v/v) solution of 5% dextrose lactated ringer solution. Through the experiment, the expired level of CO₂ was maintained between 35 and 40 mmHg by adjusting the tidal volume and respiratory rate. Heart rate was monitored and body temperature was maintained at 37°C by means of a feedback-controlled heated blanket. Lidocaine hydrochloride (2%) was applied locally at all incisions and pressure points and a craniotomy was

277 performed over area 17 (V1) contralaterally to the stimulated eye (Horsley-Clarke coordinates 4-8P; 0.5-2L). Dexamethasone
 278 (4 mg) was administered intramuscularly every 12h to reduce cortical swelling. Eye lubricant was regularly applied to avoid
 279 corneal dehydration.

280 Electrophysiological recordings

281 During each recording session, pupils were dilated using atropine (Mydriacyl) while nictitating membranes were retracted
 282 using phenylephrine (Mydrin). Rigid contact lenses of appropriate power were used to correct the eyes' refraction. Anesthesia
 283 was changed to 0.5-1% halothane to avoid anesthesia-induced modulation of visual responses³⁸. Finally, small durectomies
 284 were performed before each electrode insertion and a 2% agar solution in saline was applied over the exposed cortical
 285 surface to stabilize recordings. Linear probes ($\approx 1 \text{ M}\Omega$, 1x32-6mm-100-177, Neuronexus) were lowered in the cortical tissue
 286 perpendicularly to the pia and extracellular activity was acquired at 30KHz using an Open Ephys acquisition board⁵⁷. Single
 287 units were isolated using Kilosort 2⁵⁸ and manually curated using Phy⁵⁹. Clusters with low amplitude templates or ill-defined
 288 margin were excluded from further analysis. Additional exclusion was performed if a cluster's was unstable (firing rate below 5
 289 spikes.s⁻¹ for more than 30 seconds), or if the neuron was not deemed sufficiently orientation selective ($R^2 < 0.75$ when fitted
 290 with a von Mises distribution). Laminar positions were determined by the depth of the recording site with respect to the pia,
 291 which was then cross-validated by looking at the evoked Local Field Potential (LFP) using sink/source analysis^{60,61}.

292 Single neuron analysis

293 Orientation tuning curves were computed by selecting a 300 ms window maximizing spike-count variance⁶². The firing rate
 294 was averaged across drift directions and a von Mises distribution⁶³ was fitted to the data:

$$f(\theta_k) = R_0 + (R_{\max} - R_0) \cdot \exp \left\{ \kappa \cdot (\cos(2(\theta_k - \theta_{\text{pref}})) - 1) \right\} \quad (3)$$

295 where θ_k is the orientation of the stimuli, R_{\max} is the response (baseline subtracted) at the preferred orientation θ_{pref} , R_0 the
 296 response at the orientation orthogonal to θ_{pref} and κ a measure of concentration. To control for direction selectivity when
 297 averaging tuning curves across drift direction, we computed a direction selectivity index:

$$D_s = \frac{R_{\text{pref}} - R_{\text{null}}}{R_{\text{pref}}} \quad (4)$$

298 where R_{pref} is the firing rate at the preferred direction (baseline subtracted) and R_{null} the firing rate at the preferred direction
 299 plus π . The quality of each tuning curve was assessed by computing a global metric, the circular variance (CV) of the unfitted
 300 data, which varies from 0 for perfectly orientation-selective neurons to 1 for orientation-untuned neurons²⁷. It is defined as:

$$\text{CV} = 1 - \left| \frac{\sum_k R(\theta_k) \exp \{2i\theta_k\}}{\sum_k R(\theta_k)} \right| \quad (5)$$

301 where $R(\theta_k)$ is the response of a neuron (baseline subtracted) to a stimulus of angle θ_k . The changes of CV as a function of B_θ
 302 were fitted with a Naka-Rushton function²²:

$$f(B_\theta) = f_0 + f_{\max} \frac{B_\theta^n}{B_\theta^n + B_{\theta50}^n} \quad (6)$$

303 where f_0 is the base value of the function, $f_0 + f_{\max}$ its maximal value, $B_{\theta50}$ the stimulus' variance at half f_{\max} and n a strictly
 304 positive exponent of the function.

305 The significance of the tuning to orientation was measured by comparing the unfitted firing rate at the preferred and
 306 orthogonal orientations across trials, using a Wilcoxon signed-rank test correct for continuity, and the maximum value of
 307 B_θ which yielded a significant result was designed as $B_{\theta\max}$ (i.e., the maximum variance at which a neuron is still tuned).
 308 Shifts of the preferred orientation were evaluated as the difference of θ_{pref} between trials where $B_\theta = 0^\circ$ and $B_\theta = B_{\theta\max}$. The
 309 significance of the variation of peak amplitude of the tuning curve was measured as by comparing the unfitted firing rate at the
 310 preferred orientation between trials where $B_\theta = 0^\circ$ and $B_\theta = B_{\theta\max}$.

311 Population decoding

312 The parameters used to generate Motion Clouds were decoded from the neural recordings using a multinomial logistic regression
 313 classifier²⁸. For a given stimulus, the activity of all the recorded neurons was a vector $X(t) = [X_1(t) \ X_2(t) \ \dots \ X_{249}(t)]$,
 314 where $X_i(t)$ is the spike count of neuron i in a time window $[t; t + \Delta T]$. The onset of this window t was slid from -200 ms to 400
 315 ms (relative to the stimulation time) in steps of 10 ms while ΔT was kept constant at 100 ms. It should be noted that merging
 316 neural activity across electrodes or experiments is a common procedure^{64,65}, which we validated in our data by verifying that

317 the electrode or experiment which yielded the data could not be decoded from the neural activity (Figure S 6). Mathematically,
 318 the multinomial logistic regression is an extension of the binary logistic regression²⁸ trained here to classify the spike vector
 319 $X(t)$ between K classes. The probability of any such vector to belong to a given class is:

$$P(y = k|X(t)) = \frac{\exp\{\langle \beta_k, X(t) \rangle\}}{\sum_{k'=1}^K \exp\{\langle \beta_{k'}, X(t) \rangle\}} \quad (7)$$

320 where $\langle \cdot, \cdot \rangle$ is the scalar product over the different neurons, $k = 1, \dots, K$ is the class out of K possible values and β_k are the
 321 coefficients learned during the training procedure of the classifier. Several decoders were trained with classification tasks:
 322 decoding orientation θ ($K = 12$, Figure 5), decoding orientation variance B_θ ($K = 8$, Figure S 7) or both ($K = 12 \times 8 = 96$,
 323 Figure 6). All meta-parameters (the integration window size, penalty type, regularization strength and train/test split size) were
 324 controlled, showing that the decoding performances stem mainly from experimental data rather than fine-tuning of the decoder
 325 parameterization (Figure S 5).

326 The performance of all decoders was reported as the average accuracy across all classes K , known as the balanced accuracy
 327 score⁶⁶. The accuracy for each specific class k can also be reported in the form of a population tuning curve, in which the
 328 likelihood of decoding each possible class K is given by equation 7. To estimate the timecourse of the decoders, they were
 329 fitted in the [0; 300] ms range with a sigmoid function:

$$\sigma = \max_{\text{acc}} \left(\frac{1}{1 + e^{-k\tau}} \right) + \min_{\text{acc}} \quad (8)$$

330 where \max_{acc} and \min_{acc} are respectively the maximum and minimum accuracies of the decoder, k the steepness and τ the time
 331 constant of the function.

332 Computational model

333 We used a recurrent network of orientation-tuned neurons to model responses to increasing orientation variance B_θ . The model
 334 presently used was first used to account for the intracortical activity in the cat primary visual cortex³³, although it was presently
 335 simplified as a center-surround filter in the orientation domain²⁷. Notably, this network has been able to account for numerous
 336 experimental findings, including learning and adaptation of cortical neurons^{34,35}, whose implementations are similar to ours.

337 The model consisted of N orientation-tuned neurons, evenly tiling the orientation space between $-\pi$ and π . Each neuron is
 338 modeled as a single passive unit whose membrane potential obeys the equation:

$$\tau \delta V / \delta T + V = V_{\text{ff}} + V_{\text{exc}} - V_{\text{inh}} \quad (9)$$

339 where τ is the membrane time constant and V_{ff} , V_{exc} , V_{inh} are the synaptic potentials coming from the feedforward input,
 340 recurrent excitatory and recurrent inhibitory connectivity, respectively. The firing rate R at time t of each neuron is computed as
 341 an instantaneous quantity modulated by a gain α :

$$R(t) = \alpha \max(V(t), 0) \quad (10)$$

342 For computational simplicity, the neurons had no spontaneous firing rate and V was measured relative to the firing threshold.
 343 Each neuron could send mixed excitatory and inhibitory synaptic potentials to its neighbour, although this specific model has
 344 been reported to achieve similar behaviour with separate units³⁵. For each stimulus of main orientation θ , the input to a cell
 345 with preferred orientation θ_{pref} is:

$$V_{\text{ff}}(\theta_{\text{pref}}) = J_{\text{ff}} \frac{e^{\kappa_{\text{ff}} \cos(2(\theta - \theta_{\text{pref}}))}}{2\pi I_0(\kappa_{\text{ff}})} \quad (11)$$

346 where J_{ff} is the strength of the input and I_0 is the modified Bessel function of order 0. The right-hand side of the equation
 347 describes a von Mises with mean θ_{pref} and concentration κ_{ff} . This latter parameter is related to the orientation bandwidth B_θ ,
 348 which was varied to yield a model's TVF B_θ/CV curves:

$$B_\theta = \sqrt{\frac{0.5 \arccos((\log(0.5) + \kappa_{\text{ff}}) / \kappa_{\text{ff}})}{2 \log(2)}} \quad (12)$$

349 a total of 20 B_θ spanning the same range used in the experiments were used, each with 32 different θ tiling a $[-75^\circ; 75^\circ]$
 350 orientation space. The recurrent connectivity profile for excitatory (C_{exc}) and inhibitory (C_{inh}) synapses was controlled by
 351 separate von Mises distributions over the orientation space Θ :

$$C_{\text{exc}}(\theta_{\text{pref}}) = \frac{e^{\kappa_{\text{exc}} \cos(2(\Theta - \theta_{\text{pref}}))}}{2\pi I_0(\kappa_{\text{exc}})} \quad (13)$$

$$C_{\text{inh}}(\theta_{\text{pref}}) = \frac{e^{\kappa_{\text{inh}} \cos(2(\theta - \Theta_{\text{pref}}))}}{2\pi I_0(\kappa_{\text{inh}})} \quad (14)$$

353 which are both used to describe an overall connectivity kernel:

$$C_{\text{tot}}(\theta_{\text{pref}}) = J_{\text{exc}} C_{\text{exc}} - J_{\text{inh}} C_{\text{inh}} \quad (15)$$

354 which followed a typical Ricker wavelet (or Mexican hat) shape (Figure 7d). The overall activity of the network is then a
355 weighted sum of the firing rates of all the neurons:

$$V_{\text{exc}} - V_{\text{inh}}(t) = \sum_{\Theta} C_{\text{tot}}(\theta_{\text{pref}}) R(t) \quad (16)$$

356 Parameterization of the model was done to match single V1 neuron recordings of anesthetized cats, in an experimental setup
357 similar to the one used here⁶⁷. The computational procedure to match experimental data was entirely done in a previous
358 publication³⁵. Briefly, it consisted in scanning a range of possible values for each parameters, then fine all possible combinations
359 using a metric of likeliness to single grating response, time-to-peak, peak response and tuning width. The parameters yielded
360 by this procedure were $\tau = 10.8$ ms; $\alpha = 10.6$ Hz/mV; $J_{\text{ff}} = 9.57$ mv/Hz; $J_{\text{exc}} = 1.71$ Hz/mV; $J_{\text{inh}} = 2.0178$ Hz/mV. For the
361 feedforward mode of the model (Figure 7e), J_{exc} and J_{inh} were set to 0 Hz/mV and the input was convolved with a "receptive
362 field" :

$$\text{RF} = \frac{e^{\kappa_{\text{RF}} \cos(2(\theta - \Theta_{\text{pref}}))}}{2\pi I_0(\kappa_{\text{RF}})} \quad (17)$$

363 of which we reported the Half-Width at Half-Height, given by⁶⁸ :

$$\text{HWHH} = 0.5 \arccos\left(\frac{\log(0.5) + \kappa}{\kappa}\right) \quad (18)$$

364 For the recurrent mode (Figure 7f-h), the concentration measures of the recurrent connectivity profiles κ_{exc} and κ_{inh} were
365 both varied from 0.35 to 7, in 200 even steps, and the input was not convolved with a receptive field.

366 Statistical procedures and plotting

367 All data was analyzed using custom Python code (see Additional information). Statistical analysis was performed using non-
368 parametric tests. Wilcoxon signed-rank test with discarding of zero-differences was used for paired samples and Mann-Whitney
369 U test with exact computation of the U distribution was used for independent samples. Due to the impracticality of using error
370 bars when plotting time series, colored contours are used to represent standard deviation values (unless specified otherwise),
371 with a solid line representing mean values. For boxplots, the box extends from the lower to upper quartile values, with a solid
372 white line at the median value. The upper and lower whiskers extend to respectively $Q1 - 1.5 * IQR$ and $Q3 + 1.5 * IQR$, where
373 $Q1$ and $Q3$ are the lower and upper quartiles and IQR is the inter-quartile range.

374 Data availability

375 Data and analysis code that have been used in the present study can be made available from the corresponding author upon
376 reasonable request.

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508 Author contributions

509 L.U.P., C.C. F.C., N.C., and H.J.L. designed the study. H.J.L., N.C. and L.K. collected the data. H.J.L. and L.U.P. analyzed the
510 data. H.J.L. and L.U.P. wrote the original draft of the manuscript. All authors reviewed and edited the manuscript.

511 Competing interests

512 The authors declare no competing interests.

513 **Supplementary information**

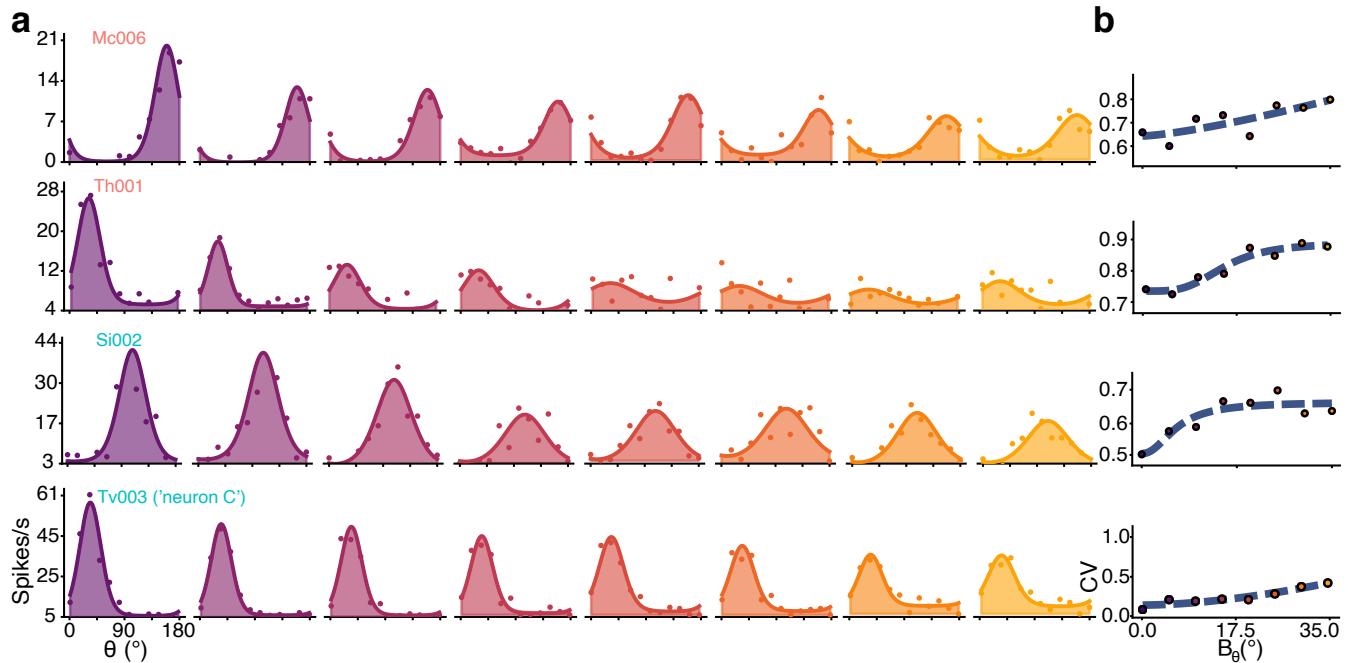


Figure S 1. Additional examples of single neuron tuning curves and VTF. **(a)** Tuning curves of four more neurons, stimulated with Motion Clouds of increasing variance B_{θ} from left to right. Colored dots represent the mean firing rate across trials (baseline subtracted, 300 ms average) and solid line a fitted von Mises function. **(b)** Variance-tuning functions (VTF), measuring the changes of orientation tuning measured by the circular variance (CV, colored dots) as a function of Motion clouds variance B_{θ} , fitted with a Naka-Rushton (NKR) function (dashed curves, parameters shown in light gray).

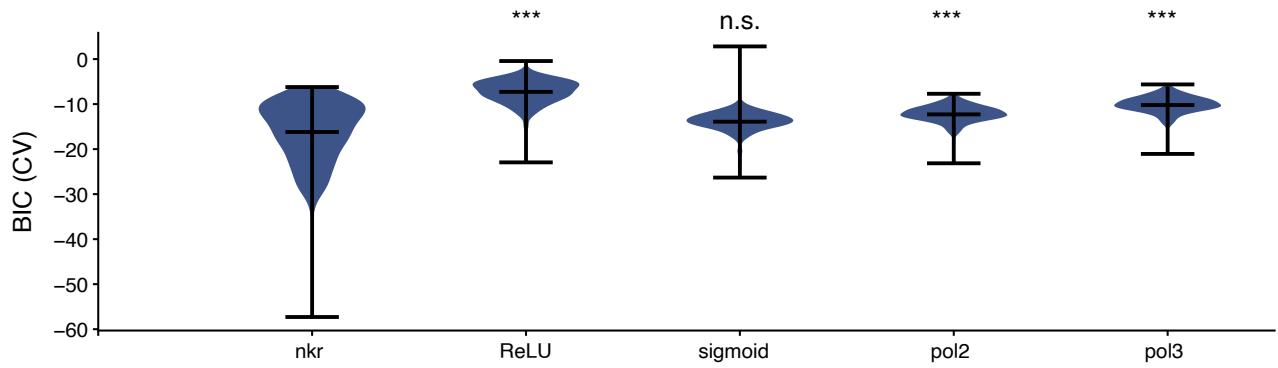


Figure S 2. Naka-Rushton functions are the most adequate fit for PRFs. Violin plot of the Bayesian Information Criterion (BIC) of the CV curves of all recorded neurons. Each violin plot represents a different type of fitted equation, respectively: Naka-Rushton (nkr, see Methods) ; Rectified Linear Unit (ReLU, $f(x) = \max(0, x)$) ; logistic function (sigmoid, $f(x) = \frac{e^x}{e^x + 1}$) ; second degree polynomial function (pol2, $f(x) = ax^2 + bx + c$) and third degree polynomial function (pol3, $f(x) = ax^3 + bx^2 + cx + d$). A lower BIC indicates less information lost in the fitting process, hence a better fitting model. Naka-Rushton curves were chosen over sigmoid functions for the skewness of the BIC distributions towards negative values, as well as the explainability of their parameters. n.s., not significant; *, $p < 0.05$; **, $p < 0.01$; ***, $p < 0.001$ (Kruskal-Wallis H-test, post-hoc Dunn Pairwise test, Bonferroni corrected).

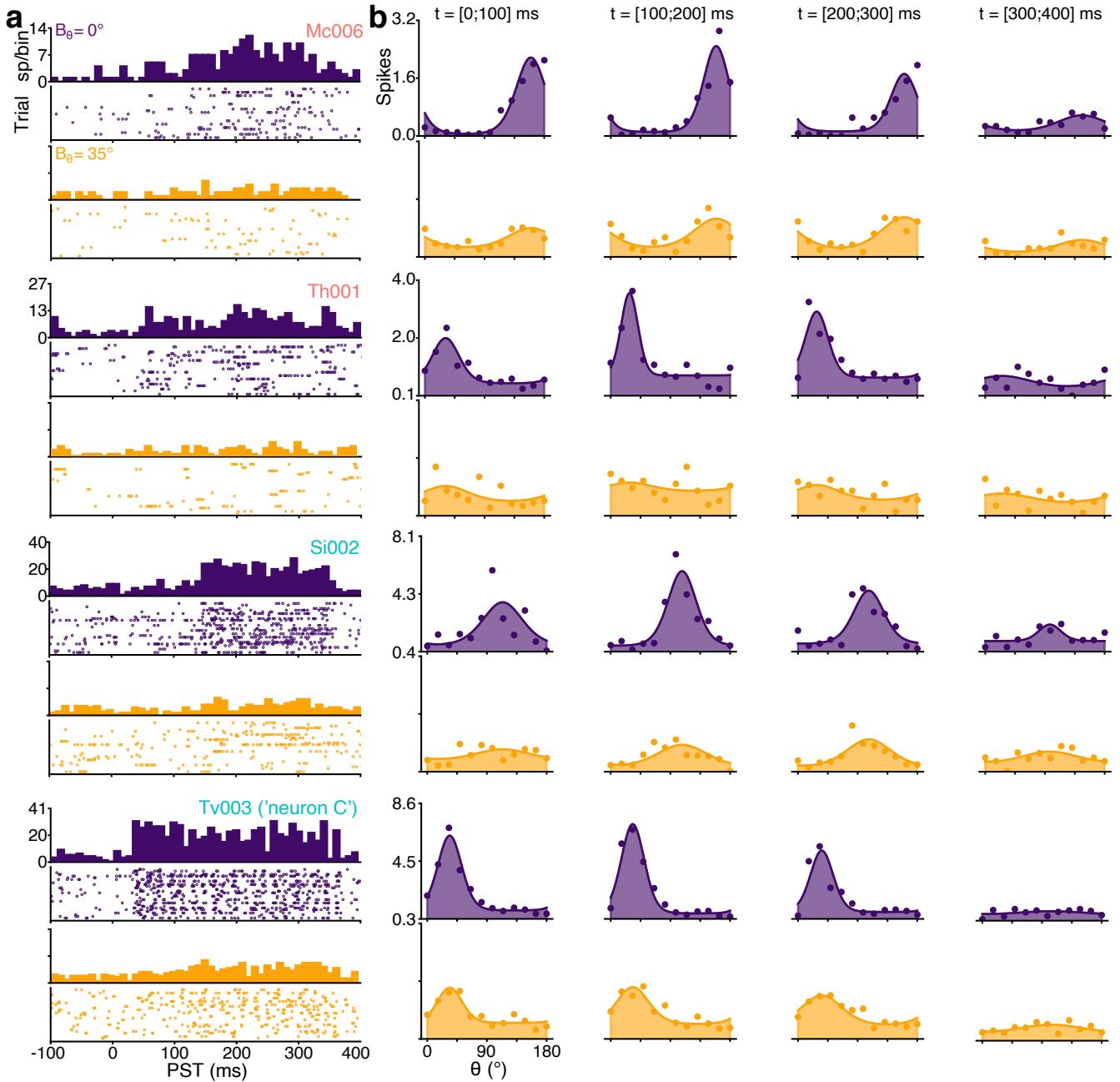


Figure S 3. Additional examples of dynamical properties of neurons. **(a)** Peristimulus time (PST) histogram and associated rasterplot of the four additional example neurons, for Motion Clouds with lowest ($B_\theta = 0^\circ$, purple) and highest ($B_\theta = 35^\circ$, yellow) variance. **(b)** Dynamics of the tuning curves shown in Figure 2 in 100 ms windows, starting at the time labelled atop of each column..

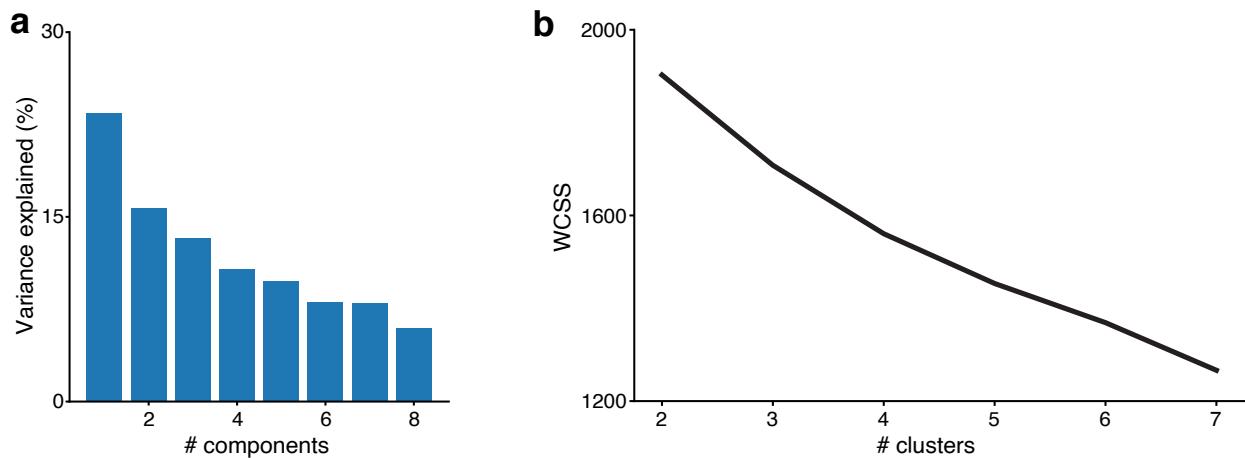


Figure S 4. Graphical reports of clustering analysis. **(a)** Fraction of variance explained as a function of the number of components used in the Principal Component Analysis. **(b)** Within-Cluster-Sum-of-Squares (WCSS) of the K-means clustering, expressed as a function of the number of clusters. An empirical way to select the number of clusters is to typically take the "elbow" of such a curve, if existent.

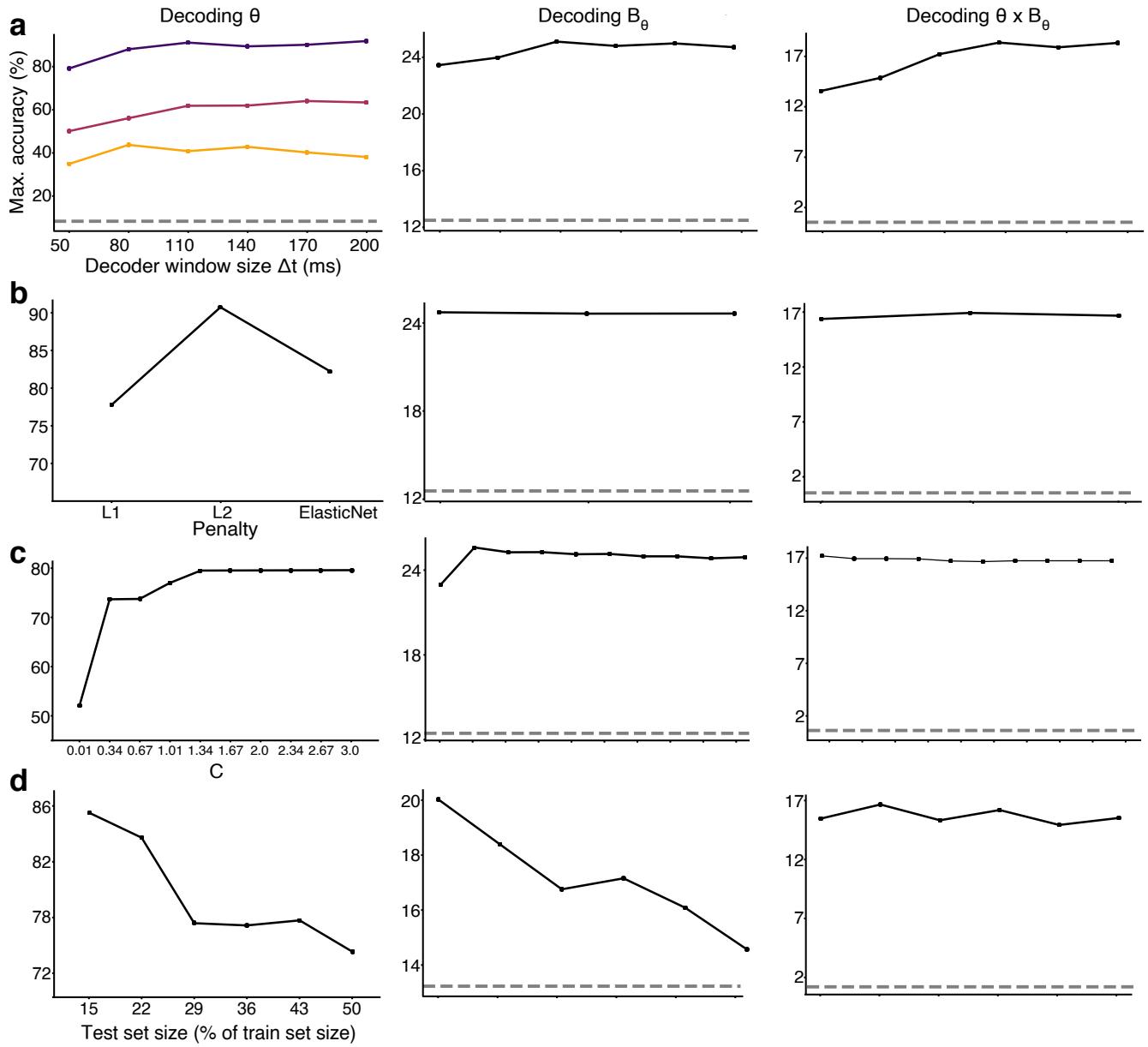


Figure S 5. Parameters evaluation of the decoders. **(a)** Parameter scanning of the orientation θ decoders for three B_θ (chance level 1/12). Maximum accuracy reached during the time course of each decoder is reported for each parameter. On the first row, the parameter optimized is the length of the time window ΔT , which controls the length of the integration time of the decoder (see Methods). On the second row, the type of penalization norm applied to the decoder, which controls the metric by which the classification error is minimized. l_1 and l_2 are defined as $\sum_i |x_i|$ and $\sqrt{\sum_i |x_i|^2}$, where x is a set of features fed to the classifier. The penalization norm "ElasticNet" linearly combines both these norms, here with an equal weighting applied to both norm. On the third row, the parameter C controls the regularization strength, which measure the magnitude of the penalty applied to large parameters, in order to prevent overfitting on reduced sets of data. On the fourth and last row, we optimized the percentage of data kept out of the training set to evaluate the decoder's accuracy. **(b)** Parameters of the orientation variance B_θ decoder (chance level 1/8). **(c)** Parameters of the orientation and orientation variance $B_\theta \times \theta$ decoder (chance level 1/96).

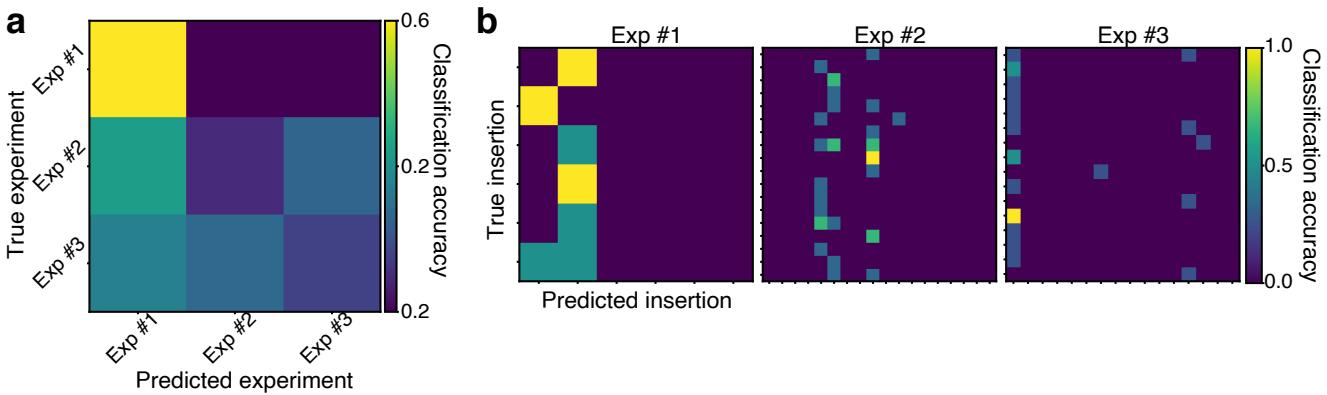


Figure S 6. The experiment identity (the specific experiment which yielded the spikes) and insertion identity (the specific insertion which yielded the spikes) of neurons does not produce any significant result, thereby validating the fusion of multiple datasets in the decoding process. **(a)** Confusion matrix of a decoder trained to retrieve the experiment identity using three groups of 30 neurons (bootstrapped 1000 times). **(b)** Confusion matrices of three decoders trained to retrieve the insertion identity of the neurons recorded in each experiment.

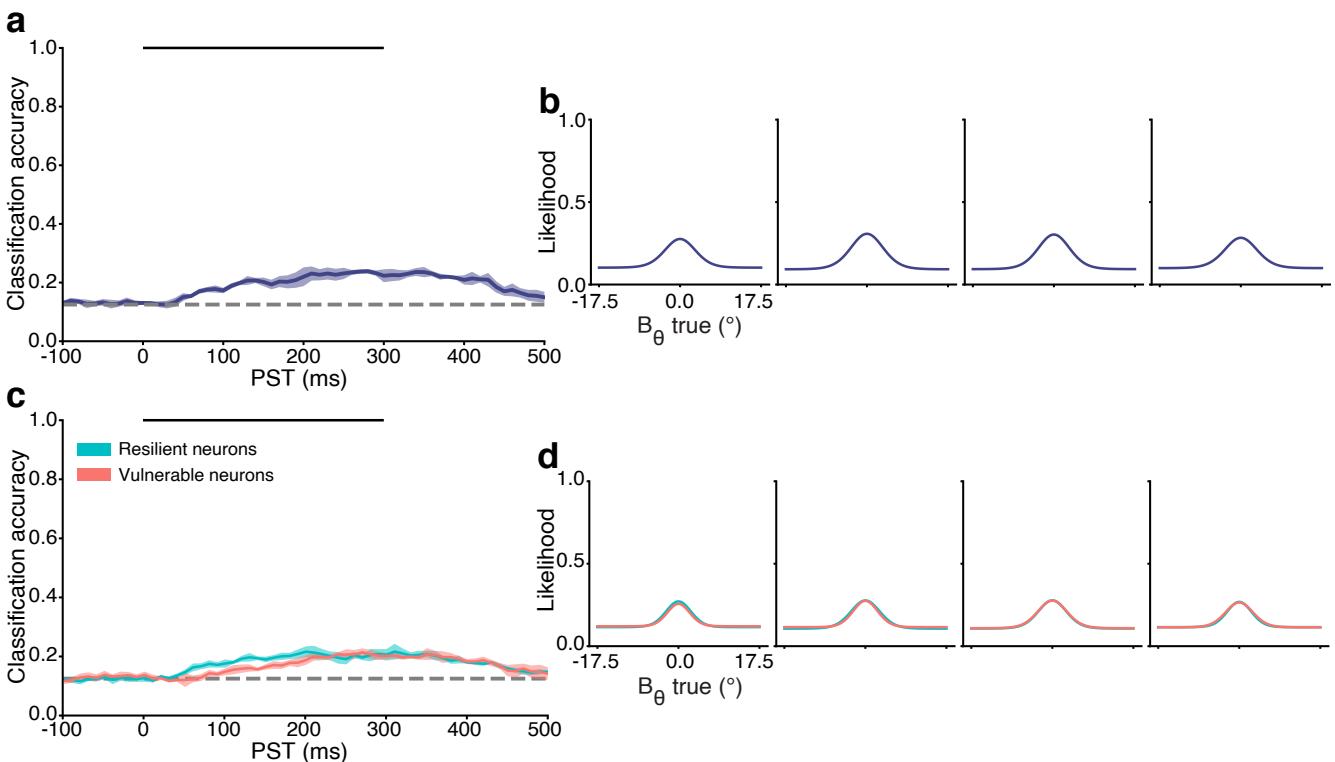


Figure S 7. Orientation variance cannot be accurately decoded from the population activity. **(a)** Time course of a decoder trained to retrieve the variance B_θ of Motion Clouds. Solid dark line represents the mean accuracy of a 5-fold cross validation and filled contour the SD. Decoding at chance level (here, 1/8) is represented by a gray dashed line. **(b)** Population tuning curve of the decoder. **(c)** Same as Figure S7a, trained with spikes from either resilient or vulnerable neurons. **(d)** Same as Figure S7b, trained with spikes from either resilient or vulnerable neurons.

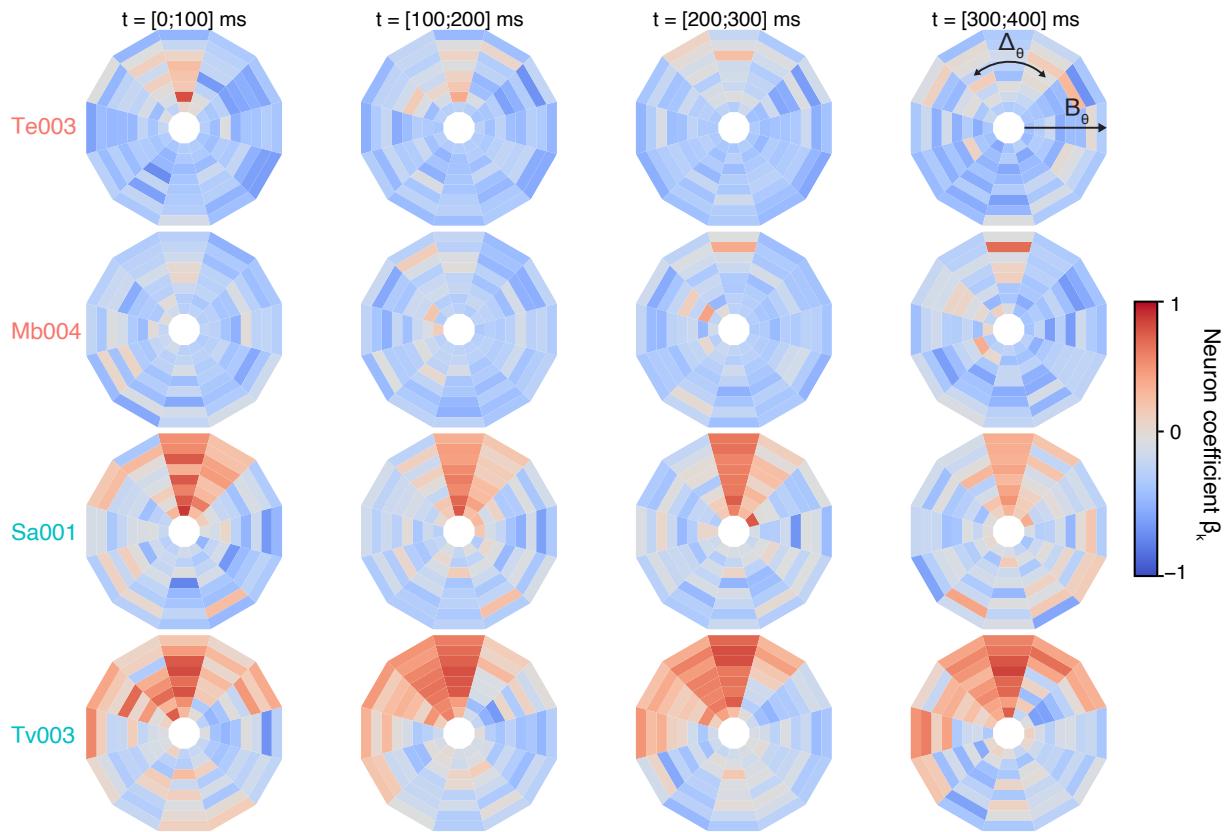


Figure S 8. Polar plot of coefficient matrices from two single vulnerable neurons (top rows) and two single resilient neurons (bottom rows). As in Figure 6e, the angle of each bin represents the error on the θ identity of the stimulus, Δ_θ , and the eccentricity corresponds to the coefficient for each B_θ (highest variance at the edge). The temporal evolution of the coefficients from the decoder is normalized for each neuron by row.