Multi-context blind source separation by error-gated Hebbian rule

Running title: Multi-context error-gated Hebbian rule

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Abstract

Animals need to adjust their inferences according to the context they are in. This is required for the multi-context blind source separation (BSS) task, where an agent needs to infer hidden sources from their context-dependent mixtures. The agent is expected to invert this mixing process for all contexts. Here, we show that a neural network that implements the *error-gated Hebbian rule* (EGHR) with sufficiently redundant sensory inputs can successfully learn this task. After training, the network can perform the multi-context BSS without further updating synapses, by retaining memories of all experienced contexts. Finally, if there is a common feature shared across contexts, the EGHR can extract it and generalize the task to even inexperienced contexts. This demonstrates an attractive use of the EGHR for dimensionality reduction by extracting common sources across contexts. The results highlight the utility of the EGHR as a model for perceptual adaptation in animals.

Introduction

Inference of the causes of a sensory input is one of the most essential abilities of animals (*Helmholtz, 1925; Knill, Pouget, 2004; DiCarlo et al, 2012*) — a famous example is the cocktail party effect, i.e., the ability of a partygoer to distinguish a particular speaker's voice against a background of crowd noise (*Brown et al., 2001; Mesgarani, Chang, 2012*). This ability has been modelled by blind source separation (BSS) algorithms (*Cichocki et al., 2009; Comon, Jutten, 2010*), by considering that several hidden sources (speakers) independently generate signal trains (voices), while an agent receives mixtures of signals as sensory inputs. A neural network, possibly inside the brain, can invert this mixing process and separate these sensory inputs into hidden sources using a BSS algorithm. Independent component analysis (ICA), achieves BSS by minimizing the dependency between output units (*Comon, 1994; Comon, Jutten, 2010*). Numerous ICA algorithms have been proposed for both rate-coding (*Bell, Sejnowski, 1995; Bell, Sejnowski, 1997; Amari et al., 1996; Hyvarinen, Oja, 1997*) and spiking neural networks (*Savin et al., 2010*).

Previously, we developed a biologically plausible ICA algorithm, referred to as the *error-gated Hebbian rule (EGHR)* (*Isomura, Toyoizumi, 2016*). This learning rule can robustly estimate the hidden sources that generate sensory data without supervised signals. Importantly, it can reliably perform ICA in undercomplete conditions (*Lee, Girolami et al., 2000*), where the number of inputs is greater than that of outputs. A simple extension of the EGHR can separate sources while removing noise within a single-layer neural network (*Isomura, Toyoizumi, 2018*), by simultaneously performing principal component analysis (PCA) (*Pearson, 1901; Oja, 1989*)

and ICA. The EGHR is expressed as a product of pre- and post-synaptic neuronal activities and
a third modulatory factor, each of which can be computed locally (i.e., local learning rule (*Lee,*
Girolami et al., 2000)). In this sense, the EGHR is more biologically plausible than non-local
engineering ICA algorithms (*Bell, Sejnowski, 1995; Bell, Sejnowski, 1997; Amari et al., 1996*).
Because of these desirable properties, the EGHR is considered as a candidate mechanism for
neurobiological BSS (*Kuśmierz et al., 2017; Avitan, Goodhill, 2018*), as well as a next-generation
neuromorphic implementation (*Neftci, 2018*) for energy efficient BSS.

The optimal inference and behavior often depend on context. Indeed, our perception and
decisions reflect this context-dependence, i.e., cognitive flexibility (*Dajani, Uddin, 2015*).
Studies in primates have suggested that a contextual-cue-dependent dynamic process in the
prefrontal cortex controls this behavior (*Dehaene, Changeux, 1991; Gilbert, Sigman, 2007;*
Mante et al., 2013), and several computational studies have modeled it (*Mante et al., 2013;*
Song et al., 2016; Song et al., 2017; Miconi, 2017; Chaisangmongkon et al., 2017). Likewise,
context dependence of auditory perceptual inference has been modeled (*Ahrens et al., 2008*).
In addition to experimental evidence, recent progress in machine learning has also addressed
this multi-context problem, in an attempt to create artificial general intelligence (*Yu et al.,*
2010; Kirkpatrick et al., 2017; Zenke et al., 2017). By implementing (task-specific) synaptic
consolidation, a neural network can learn a new environment, while retaining past memories,
by protecting synaptic strengths that are important to memorizing past environments. Those
findings indicate the importance of multi-context processes for cognitive flexibility.

Unlike the above-mentioned tasks, BSS in several different contexts has some difficulty.
Conventional ICA algorithms assume the same number of input and output neurons (*Bell,*
Sejnowski, 1995; Bell, Sejnowski, 1997; Amari et al., 1996, Foldiak, 1990; Linsker, 1997) and,
thus, cannot straightforwardly perform a multi-context BSS. After learning, the synaptic
strength matrix of these algorithms converges to the inverse of the mixing matrix of the
current context (or its permutation or sign-flip), which is generally different from that in the
previous context. Hence, when the network subsequently encounters a previously learnt
context, it needs to relearn the synaptic strengths from the very beginning. More involved
engineering ICA algorithms, such as the nonholonomic ICA algorithm (*Amari et al., 2000*) and
the ICA mixture algorithm (*Lee, Lewicki et al., 2000; Hirayama et al., 2015*), are expected to
perform the multi-context BSS. However, we argue that EGHR has the advantage of being a
biologically plausible local learning rule.

Here we show that the EGHR can perform multi-context BSS when a neural network receives
redundant sensory inputs. It can retain memories of previously experienced contexts and
process the BSS right after contextual switching to a previously learnt context. This suggests
that the EGHR can also be used as a powerful data compression method (*Cunningham,*
Ghahramani, 2015), since it extracts hidden sources shared across contexts, despite the
proportional increase in data dimensions to the number of contexts. Moreover, when a

common feature is shared across contexts, the EGHR can extract it to perform BSS, while
filtering out features that vary among contexts. Once the learning is achieved, the network can
perform BSS even in an inexperienced context, indicating some generalization capability or
transfer learning. We demonstrate that the EGHR with sufficiently redundant sensory inputs
learns to distinguish birdsongs from their superpositions and retains this ability even after
learning different sets of birdsongs. The rule finds a general representation that is capable of
separating an unheard set of birdsongs. Finally, possible neurobiological implementations of
the EGHR are discussed.

Results

Error-gated Hebbian rule (EGHR)

In a BSS task, several hidden sources (s) independently generate signal traces, while our
agent receives their mixtures as sensory inputs (x). In this study, we consider a multi-context
BSS task, in which a set of contexts with different mixing weights is used. Sensory inputs are
randomly generated from one of these contexts for a period of time. Let k (= 1, …, C) be an
index of context. Our experimental setup consists of an N_s-dimensional vector of hidden

sources $s \equiv \left(s_1, \ldots, s_{N_s}\right)^T$ whose elements s_i independently follow a non-Gaussian

distribution $p(s_i)$, an N_x-dimensional vector of sensory inputs $x \equiv \left(x_1, \ldots, x_{N_x}\right)^T$, and an N_u-

dimensional vector of neural outputs $u \equiv \left(u_1, \ldots, u_{N_u}\right)^T$ (Fig. 1). The sensory inputs in the k-

th condition are generated by transforming the hidden sources, i.e., the so-called generative
process:

$$\textbf{Sensory inputs} \quad x = A^{(k)}s. \tag{1}$$

Here $A^{(k)}$ is the $N_x \times N_s$ mixing matrix for the k-th context that defines the magnitude of
inputs, when each source generates a signal. To ensure that each $A^{(k)}$ represents a different
context and that each context has an ICA solution, column vectors of a block matrix
$\left(A^{(1)}, A^{(2)}, \ldots, A^{(C)}\right)$ are supposed to be linearly independent of each other. We design the
task such that these contexts appear sequentially or randomly. The neural outputs are
expressed as sums of inputs weighted by an $N_u \times N_x$ synaptic strength matrix W, and
calculated by:

$$\textbf{Neural outputs} \quad u = Wx. \tag{2}$$

It is well known that when a presynaptic neuron (x_j) and a postsynaptic neuron (u_i) fire

together, Hebbian plasticity occurs, and the synaptic connection from x_j to u_i, denoted by
W_{ij}, is strengthened (Hebb, 1949; Bliss, Lomo, 1973). Because this constitutes associative
learning, correlations between $x_1, \dots, x_{N_x}$ and u_i are usually enhanced; thereby,
correlations among neural outputs also increase. This is opposite to separation of signals (i.e.,
BSS) because each neural output is expected to encode a specific source to perform BSS. To
separate signals, we introduce a global scalar factor (i.e., a third factor) given by the sum of
nonlinearly-summed output units (*Isomura, Toyoizumi, 2016*):

$$\textbf{Global factor} \qquad E(u) \equiv -\log p_0(u) = -\sum_{i=1}^{N_u} \log p_0(u_i). \qquad (3)$$

Here $p_0(u)$ is the prior distribution that the agent expects the hidden sources to follow; e.g.,
when $p_0(u)$ is a Laplace distribution of mean zero and unit variance, $E(u) = \sqrt{2}\big(|u_1| +$
$\cdots + |u_{N_u}|\big) + \text{const}$. We suppose that this global factor modulates Hebbian plasticity. Recent
experimental studies have reported that synaptic plasticity can be modulated by various
neuromodulators (*Reynolds et al., 2001; Zhang et al., 2009; Salgado et al., 2012; Yagishita et*
al., 2014; Johansen et al., 2014), GABAergic inputs (*Paille et al., 2013; Hayama et al., 2013*), or
glial factors (*Ben Achour, Pascual, 2010*). Possible neurobiological implementations of the
global factor are further discussed in the Discussion section. Overall, the synaptic strength
matrix W is updated by the EGHR in the following way:

$$\textbf{Synaptic plasticity (EGHR)} \qquad \dot{W} \propto \langle \underbrace{\big(E_0 - E(u)\big)}_{\text{global factor}} \underbrace{g(u)}_{\text{post}} \underbrace{x^T}_{\text{pre}} \rangle, \qquad (4)$$

where $\dot{W}$ is the derivative of *W* with respect to time, and $g(u) \equiv dE(u)/du$ is a non-linear
function usually associated with a nonlinear activation function. A constant E_0 scales the
neural outputs; the output scale becomes equivalent to the source scale when $E_0 =$
$\text{E}[-\log p_0(s)] + 1$, where $\text{E}[\cdot]$ is the expectation over the input distribution. In short, the
EGHR constitutes a Hebbian learning rule when the global factor is smaller than the threshold
$(E(u) < E_0)$, otherwise $(E(u) > E_0)$, it becomes an anti-Hebbian rule. This mechanism makes
output neurons independent from each other. The detailed derivation and theoretical proofs
of the EGHR have been described in previous reports (*Isomura, Toyoizumi, 2016; Isomura,*
Toyoizumi, 2018).
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Figure 1. Model setup for multi-context BSS task. In this model, $s_1, \dots, s_{N_s}$ are hidden sources (e.g., birdsongs); $x_1, \dots, x_{N_x}$ are sensory inputs that an agent receives; $u_1, \dots, u_{N_u}$ are neural outputs; $A_{11}^{(k)}, \dots, A_{1N_s}^{(k)}, A_{21}^{(k)}, \dots, A_{N_x N_s}^{(k)}$ are elements of the k-th-context mixing matrix; $W_{11}, \dots, W_{1N_x}, W_{21}, \dots, W_{N_u N_x}$ are synaptic strengths; and E is a scalar global factor that mediates synaptic plasticity. Synaptic strengths are adjusted to perform multi-context BSS by the EGHR.

Memory capacity of the EGHR

First, we analytically show the memory capacity of a neural network established by the EGHR. As the number of contexts increases, larger dimensions of inputs are needed to retain information pertaining to past contexts in the neural network. For simplicity, let us suppose that N_u = N_s. Because the network represents a linear inverse model of the generative processes, the goal of the multi-context BSS is generally given by:

$$W\left(A^{(1)}, \dots, A^{(C)}\right) = \left(\Omega^{(1)}, \dots, \Omega^{(C)}\right), \tag{5}$$

where $\Omega^{(k)}$ is an $N_u \times N_s$ matrix equivalent to the identity matrix, up to permutations and sign-flips. This is because the success of BSS is defined by one-by-one mapping from sources to outputs. Thus, if and only if the pseudo inverse matrix of a set of mixing matrices $\left(A^{(1)}, \dots, A^{(C)}\right)$ is a full-rank, the multi-context BSS is successful. Hence, we find that the following conditions are necessary to achieve the multi-context BSS for a generic $\left(A^{(1)}, \dots, A^{(C)}\right)$: (1) the input dimension needs to be equal to or larger than the number of contexts times the number of sources, $N_x \geq CN_s$; and (2) the output dimension needs to be

equal to or larger than the source dimension, $N_u \geq N_s$. Note that the neural network learns

the information representation compressing the sensory inputs because we consider much

greater input than output dimensions.

The memory capacity of the EGHR was empirically confirmed by numerical simulations (Fig.

2). Here, we supposed that two contexts generated inputs alternately. In each context, six-

dimensional inputs were generated from two-dimensional sources with different mixing

weights, as denoted by $A^{(1)}$ and $A^{(2)}$ (see top and middle rows in Fig. 2A). A neural network

consisting of six input and two output neurons received the inputs and changed its synaptic

strengths through the EGHR (i.e., training). After training, each neural output came to

selectively respond to (i.e., encode) one of the two sources (bottom row in Fig. 2A). Thus, the

network achieved separation of the sensory inputs into their sources without being taught the

mixing weights (i.e., BSS).

Crucially, the neural network was able to retain the information learnt for all past contexts

if provided with sufficiently redundant sensory inputs. This property is illustrated by the

trajectories of the BSS error and EGHR cost function in Fig. 2B. We defined the BSS error for

context *k* as the ratio of first to second maximum absolute values averaged for every row and

column of matrix $K^{(k)} \equiv W A^{(k)}$, which expresses the mapping from sources to outputs

(equivalent to the covariance matrix between hidden sources and neural outputs $K^{(k)} =$

$\mathrm{Cov}(u, s)$). This definition was made to ensure that it took zero if and only if one source

mapped onto one output, and *vice versa*; otherwise it took a positive value less than one.

Moreover, the cost function of the EGHR was defined as the expectation of the square of the

global factor: $L \equiv E\left[\left(E_0 - E(u)\right)^2 / 2\right]$. Context 1 (red in Fig. 2A) was provided in the first

session. Since synaptic strengths started from a random initial state, the BSS error at the

beginning of the first session was large; then, the network learned an optimal set of synaptic

strengths, and the error became zero. Notably, the EGHR was defined to reduce the EGHR cost

function by gradient descent updates. However, when context 2 (blue in Fig. 2A) was provided

for the first time in the second session, the EGHR cost function transiently increased because

it needed to learn the new mixing matrix. An important point was revealed at the first step of

the third session, in which context 1 was provided again. The BSS error was significantly smaller

than that in the first session and close to zero from the beginning of this session, indicating

that the network retained synaptic strengths that were optimized for context 1 even after

learning context 2. After several iterations, the BSS error for both contexts converged to zero.

The success of learning was also confirmed by the trajectory of the EGHR cost function that

also converged to the minimum value.

These results show that an undercomplete EGHR increased the speed of re-adaptation to

previously experienced contexts, suggesting that memory of past experiences was preserved

within the network. Moreover, the network learned the optimal set of synaptic strengths that

entertained both contexts after several iterations. A key feature for this ability is the "null
space" in the synaptic strength matrix. Since only 2 × 2 dimensions were required to express a
mapping from two-dimensional sources to two-dimensional outputs in one context, the
synaptic strength matrix comprised eight (2 × 6 – 2 × 2)-dimensional degrees of freedom. This
freedom spanned a null space in which synaptic strengths were equally optimized with a zero
BSS error. Similarly, when two different contexts were considered, four-dimensional degrees
of freedom remained, as an overlap between the two eight-dimensional null spaces. To
visualize such a null space, we projected synaptic strengths onto a subspace spanned by the
first (PC1) and second (PC2) principal components of the trajectory of synaptic strengths. On
this PC1-PC2 plane, a null space was illustrated as a nullcline. Since the dynamics of synaptic
strengths were determined to go down the slope of a cost function for either context 1 or 2,
synaptic strengths were started from a random initial state and reached the nullcline of either
context 1 or 2, in turn. Crucially, this trajectory converged to the cross point of two nullclines,
where the synaptic strengths entertained both contexts. Because of this, the BSS error reached
zero after iterative training, i.e., an ICA solution that was optimized for both contexts was
found.
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Figure 2. Results of multi-context BSS. (**A**) Distributions of sources, inputs, and outputs for context 1 and 2. (**B**) Trajectories of BSS error (top) and cost function (bottom). (**C**) Visualization of null spaces. The panel illustrates the shapes of the cost function under each context (left and right) and the trajectory of synaptic strengths (*W*) projected in a subspace spanned by the first (PC1) and second (PC2) principal components (center). The trajectory is determined by the gradient of either cost function, depending on the context. On the PC1-PC2 plane, null spaces are illustrated as nullclines; red and blue curves are nullclines for contexts 1 and 2, respectively. Low cost areas (i.e., valleys of the cost functions) are highlighted by red or blue shading. The synaptic strength matrix starts from a random initial state (star mark), shifts to the nullcline of context 1 or 2, and eventually converges to the cross point of the two nullclines, where the synaptic strengths perform the BSS for both contexts. Each source was randomly generated by the unit Laplace distribution. A learning rate of $\eta = 4 \times 10^{-6}$ was used. The MATLAB source code for this simulation is appended as Supplementary Source Codes.

Furthermore, we examined the multi-context BSS by the EGHR using a large number of
contexts (Fig. 3). Our agent received redundant (2000-dimensional) sensory inputs, comprising
100 sets (contexts) of mixtures of ten hidden sources (1000 sources in total), that were
generated as products of the context-dependent mixing matrix and sources. Ten outputs
neurons learned to infer each source from their mixtures by updating synaptic strengths
through the EGHR. After training, we found that they successfully represented the ten sources
for every context, without further updating synaptic strengths, as illustrated by the reduction
of the BSS error for all 100 contexts (Fig. 3A) and the convergence of the covariance between
sources and outputs to a diagonal matrix (up to permutations and sign-flips) (Fig. 3B). This was
because synaptic strengths had sufficient capacity and were formed to express the inverse of
the concatenated mixing matrices from all contexts, which was further confirmed by the
convergence of the synaptic strength matrix in the null space (Fig. 3C).

Figure 3. BSS with large number of contexts. One of 100 different contexts was randomly
selected for each session. Each session contained $T = 10^5$ time steps and the training continued
600 sessions. In each session, the 2000-dimensional sensory inputs (x) were generated from
ten-dimensional hidden sources (s), which independently followed the unit Laplace
distribution, through a context-dependent random mixing matrix $A^{(k)}$. The neural network
consisting of ten-dimensional neural outputs (u) was trained with a learning rate of $\eta = 1 \times 10^{-5}$.
(**A**) Trajectories of BSS error for context 1 and 100 and the average BSS error over contexts 1
to 100. The shaded area shows the standard deviation. (**B**) Mappings from ten sources to ten
outputs in contexts 1 and 100 after training. Elements of matrix $K^{(k)} = W A^{(k)}$ with $k = 1$ and 100
are illustrated by the heat map. Only one element in each row and column takes ±1, indicating
the one-to-one mapping from sources to outputs, i.e., the success of multi-context BSS. (**C**)
The dynamics of synaptic strength matrix W projected in the three-dimensional space spanned
by the first to third principal components (PC1 to PC3). The matrix starts from a random initial
point (star mark) and converges to the null space, in which synaptic strengths are optimized
for all trained contexts. The C code for this simulation is appended as Supplementary Source
Codes.

BSS in constantly time-varying environments

In the previous section, we described a general condition for the neural network to achieve
the multi-context BSS. In special cases, where the mixing matrices in each context have
common features, the neural network can perform BSS beyond the capacity mentioned above.
Here we show that when contexts are generated from a low-dimensional subspace of mixing
matrices and thereby are dependent on each other, the EGHR can find the common features
and use them to perform the multi-context BSS.

As a corollary of the property of the EGHR when provided with redundant inputs, the EGHR
can perform the BSS even when the mixing matrix changes constantly as a function of time
(Fig. 4A). Without loss of generality, a time-dependent mixing matrix is expressed by the sum
of time-invariant and time-variant components, as follows:

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$$A(t) \equiv \underbrace{A^{(0)}}_{\text{time-invariant}} + \underbrace{A^{(1)}R(t)}_{\text{time-variant}} , \qquad (6)$$

where $A^{(0)}$ is a constant matrix with the same size as $A(t)$, $A^{(1)}$ is a constant vertically-long
rectangular (or square) matrix, and $R(t)$ is a matrix composed of either smoothly or
discontinuously changing functions in time. Each component of $R(t)$ is supposed to on
average slowly change, i.e., their time-derivatives are typically much smaller in magnitude
than those of $s(t)$. This condition is required to distinguish whether changes in inputs are
caused by changes in the mixing matrix $A(t)$ or the hidden sources $s(t)$. Formally, $A(t)$
expresses infinite contexts along the trajectory of $R(t)$. This is a more complicated setup than
the standard BSS in the sense that both sources and the mixing matrix change in time.
Nonetheless, the EGHR can achieve BSS for all contexts if a solution of the synaptic strength
matrix that satisfies $W\big(A^{(0)}, A^{(1)}\big) = (\Omega, O)$ exists. Here, Ω represents the identity matrix
up to permutations and sign-flips. Such a solution generally exists if $A^{(0)}$ is a full rank matrix,
and N_x is greater than the sum of the source dimension and the row dimension of $R(t)$. The
above condition means that the network performs BSS based on the time-invariant features
$A^{(0)}$ of the mixing matrix, while neglecting the time-varying features $A^{(1)}R(t)$. This can be
viewed as a way to compress high-dimensional data. This is distinct from the standard
dimensionality reduction approach by PCA, which would preferentially extract the time-
variant features due to their extra variances.

In the simulation, we supposed $R(t)$ to be a two-dimensional rotation matrix, $R(t) =$
$(\cos \omega t, -\sin \omega t; \sin \omega t, \cos \omega t)$, with an angular frequency of $\omega = \sqrt{2}\pi/100$. The
simulation showed a reduction in the BSS error (Fig. 4B). At the same time, $K = WA$
converged to the identity matrix up to permutations and sign-flips, $K \rightarrow (0,1; -1,0)$ in this
case, although A continuously changed in time (Figs. 4C and 4D). As illustrated in Fig. 4E, the
synaptic matrix W became perpendicular to the time-varying features $A^{(1)}$ (i.e., $WA^{(1)} = O$),

by a monotonic reduction of the overlap between W and $A^{(1)}$ (defined by the Frobenius
norm of their product). After training, the overlap converged to zero. Hence, synaptic strengths
were optimized regardless of $R(t)$ at this solution, which enabled the network to perform
BSS with a virtually infinite number of contexts.

Figure 4. BSS with time-varying mixing matrix. (**A**) Schematic image of sensory inputs
generated from two sources through time-varying mixing matrix $A(t)$. The mixing matrix is
controlled by the low-dimensional rotation matrix $R(t)$. (**B**) Trajectory of BSS error. (**C**)
Trajectories of mapping weights from sources to outputs, i.e., matrix $K = W A(t)$. (**D**) Dynamics
of matrix K projected on the first two-dimensional PCA subspace of K's trajectory over training.
The matrix starts from a random initial point (star mark) and follows a spiral trajectory as it
converges to a subspace in which synaptic matrix W is perpendicular to the time-varying
component $A^{(1)}$. (**E**) Overlap of synaptic matrix W with time-invariant component $A^{(0)}$ and time-
variant component $A^{(1)}$. The overlap between two matrices was defined by the Frobenius norm

of their product, i.e., $\left|WA^{(k)}\right|_F \equiv \sqrt{\sum_{ij}(WA^{(k)})_{ij}^2}$. Each source was randomly generated by

the unit Laplace distribution. The learning rate of $\eta = 1 \times 10^{-5}$ was used. The C code for this
simulation is appended as Supplementary Source Codes.

Next, we demonstrated the utility of the EGHR, when supplied with redundant inputs, by
using natural birdsongs and a time-variant mixing matrix that expressed a natural contextual

change. Figure 5 illustrates the BSS task of two birdsongs when birds moved around the agent; thereby, the mixing matrix changed in time according to the positions of the birds. Please see also Supplementary Movie S1 for a comparison of the neural outputs before and after training. To obtain time-independent features, we assumed that the two birds moved around in non-overlapping areas. For simplicity, we also assumed that the two birds moved around at different heights. The agent received mixtures of the two birdsongs through six microphones with different direction preferences regarding the z-axis (i.e., time-invariant features) and the x- and y-axes (i.e., time-variant features). By tuning synaptic strengths by the EGHR, neural outputs were established to infer each birdsong, while the mixing matrix changed continuously. Crucially, after training, the mapping from the sources to the outputs ($K = W A$) became constant with time, although matrix A was time-dependent. More precisely, the EGHR found a representation where W satisfied $W(A^{(0)}, A^{(1)}) = (\Omega, O)$. Hence, neural outputs could separate the two birdsongs, although the amplitudes of the songs recorded by the microphones continuously changed depending on the positions of birds.

Figure 5. BSS of birdsongs when two birds move around the agent. A snapshot of the simulation overview movie after training is shown. Songs (or sources) generated by two birds s_1, s_2 (right top) are mixed with a time-varying mixing matrix A, resulting in six-dimensional sensory inputs x_1, …, x_6 (right middle). The mixed signals correspond to the recording through six microphones with different preferences. The neural network converts the six inputs into two neural outputs u_1, u_2 (right bottom) using the synaptic strength matrix W. The synaptic updates by the EGHR enable the outputs to encode each birdsong. Matrix $K = W A$ represents the mapping from sources to outputs. See Methods for the detailed simulation setup.

Generalization for inexperienced environments

Finally, we examined the generalization capability of the multi-context BSS by the EGHR
using natural birdsongs. For the sake of simplicity, we reduced Eq. (6) by considering $R(t)$ that
changes discontinuously at the beginning of each session but otherwise is constant.
Specifically, we considered the mixing matrix

$$A(v) = \underbrace{A^{(0)}}_{\text{context-independent}} + \underbrace{A^{(1)}v_1 + \cdots + A^{(n)}v_n}_{\text{context-dependent}}, \qquad (7)$$

written using context-independent matrix $A^{(0)}$, context-dependent matrices $\{A^{(1)}, \dots, A^{(n)}\}$,
and context vector $v \equiv (v_1, \dots, v_n)$ that discontinuously changes at the beginning of a new
session. The first term in the right-hand side of Eq. (7) corresponds to the context-independent
(i.e., constant) component, which should be a full-rank matrix to provide an ICA solution.
Similarly to the case with the continuously time-varying mixing matrix, the EGHR can establish
synaptic matrix W that expresses the pseudo inverse of $A^{(0)}$ up to permutations and sign-
flips, while keeping W perpendicular to $A^{(1)}, \dots, A^{(C)}$, i.e., $W\big(A^{(0)}, A^{(1)}, \dots, A^{(C)}\big) =$
$(\Omega, O, \dots, O)$. Notably, the EGHR can establish such W by using only a handful samples of v
out of combinatorially many possibilities. This is because the mappings from sources to inputs
are restricted to be a linear transformation and thereby observations with the polynomial
(probably quadratic) order number of contexts can identify the mapping for all contexts. This
property is particularly useful when *v* is high dimensional.

In this demonstration, ten sets (contexts) of mixtures of ten birdsongs were introduced to
our agent, with redundant sensory inputs composed of 100 mixed sound waves (Fig. 6A).
Those contexts were defined by random mixing matrices $A^{(0)}, A^{(1)}, \dots, A^{(4)}$. We trained the
network using only 10 contexts: *v* = (1,0,0,0), (½,½,0,0), (0,1,0,0), (0,½,½,0), (0,0,1,0), (0,0,½,½),
(0,0,0,1), (½,0,0,½), (½,0,½,0), (0,½,0,½). At the beginning of each session, *v* was randomly
selected from the above ten vectors, which provided a discrete random transition among 10
contexts. Ten output neurons learned to infer each birdsong from their mixtures, by updating
synaptic strengths through the EGHR. After training, they successfully represented the ten
birdsongs without further updating synaptic strengths. Crucially, the network could perform
BSS even in an inexperienced context (for example, in *v* = (¼,¼,¼,¼); the most right column in
Fig. 6A). This speaks to the generalization of the multi-context BSS for unseen test contexts.

Next, we quantitatively showed that, as learning progresses, the BSS error for test contexts
(defined using 20 randomly sampled *v* from a uniform distribution [0,1] that were
inexperienced in the training), as well as trained contexts, decreased (Fig. 6B). Regardless of
the given context, matrix *K* converged to a constant matrix that was the same as the identity
matrix up to permutations and sign-flips, as illustrated by the convergence of principal
components of *K* into a fixed point representing the context-independent null space (Fig. 6C).

This convergence was further validated by plotting the trajectories of the overlaps between W

and A components (Fig. 6D). While the overlap between W and $A^{(0)}$ increased as learning

progressed, the overlap with context-dependent components ($A^{(1)}, ..., A^{(4)}$) decreased and

converged to zero, showing that W became perpendicular to $A^{(1)}, ..., A^{(4)}$ by the EGHR.

Hence, this learnt network could perform BSS with $A(v)$ determined by arbitrary v in the four-

dimensional space, without further synaptic updating or transient error, while the network

was trained only with 10 contexts. Those results highlight the significant generalization

capability of the neural network established by the EGHR and the robustness against

inexperienced environments for performing BSS.

Figure 6. Generalization of multi-context BSS. (**A**) Spectrograms of ten hidden sources (birdsongs), 100 sensory inputs, and ten neural outputs. Each context is illustrated in a different color. (**B**) Trajectories of the BSS error with four trained contexts and the average BSS error over 20 inexperienced test contexts, created using randomly sampled v. (**C**) Dynamics of matrix K projected on the first two-dimensional PCA subspace. The matrix starts from a

random initial point (star mark) and converges to a fixed point, at which the synaptic matrix

entertains every trained context. (**D**) Overlap of synaptic matrix W with context-independent

component $A^{(0)}$ and context-dependent components $A^{(1)}$, …, $A^{(4)}$. Overlap between two

matrices is defined by $\left| W A^{(k)} \right|_F$, as described in Fig. 4. See Methods for the detailed

simulation setup.

Discussion

While a real environment comprises several different contexts, humans and animals retain

the experience of past contexts to perform well when they find themselves in the same context

in the future. This ability is known as conservation of learning or cognitive flexibility (*Mante et*

al, 2013; Dajani, Uddin, 2015). Although analogous learning is likely to happen during BSS, the

conventional biological BSS algorithms (*Foldiak, 1990; Linsker, 1997*) must forget the memory

of past contexts to learn a new one. Thereby, when the agent subsequently encounters a

previously experienced context, it needs to relearn it from the very beginning. We overcame

this limitation by using the described algorithm, the EGHR. The crucial property of the EGHR

is that when the number of inputs is larger than the number of sources, the synaptic matrix

contains a null space in which synaptic strengths are equally optimized for performing BSS.

Hence, with sufficiently redundant inputs, the EGHR can make the synaptic matrix optimal for

every experienced context. This is an ability that the conventional biologically plausible BSS

algorithms do not have, due to the constraint that the number of inputs and outputs must be

equal; however, this ability is crucial for animals to perceive and adapt to dynamically changing

multi-context environments. It is also crucial for animals to generalize past learning to

inexperienced contexts. We also found that, if there is a common feature shared across the

training contexts, the EGHR can extract it and generalize the BSS result to inexperienced test

contexts. This speaks to a generalization capability and transfer learning, i.e., this result implies

prevention of overfitting to a specific context; alternatively, one might see this as an extraction

of a general concept across the contexts. Therefore, we argue that the EGHR is a good

candidate model for describing the neural mechanism of conservation of learning or cognitive

flexibility for BSS.

Moreover, the process of extracting hidden sources in the multi-context BSS setup can be

seen as a novel concept of dimensionality reduction (*Cunningham, Ghahramani, 2015*). If the

dimensions of input are greater than the product of the number of sources and the number

of contexts, the EGHR can extract the low-dimensional sources (up to context-dependent

permutations and sign-flips), while filtering out a large number of context-dependent signals

induced by changes in the mixing matrix. ICA algorithms for multi-context BSS (*Amari et al.,*

2000; Lee, Lewicki et al., 2000; Hirayama et al., 2015) and undercomplete ICA for compressing

data dimensionality (*Porill and Stone, 1997; Isomura and Toyoizumi, 2016; Isomura and*

Toyoizumi, 2018) have been separately developed. Nevertheless, to our knowledge, our study

is the first to attempt a dimensionality reduction in the multi-context BSS setup. This method

is particularly powerful when a common feature is shared across the contexts, because the

EGHR can make each neuron encode an identical source across all the contexts. Our results

are different from those obtained using standard dimensionality reduction approaches by PCA

(*Pearson, 1901; Oja, 1989*), because PCA is used for extracting subspaces of high-variance

principal components and hence would preferentially extract the context-dependent varying

features, given that each source has the same variance. Therefore, our study proposes an

attractive use of the EGHR for dimensionality reduction.

We demonstrated that a neural network learns to distinguish individual birdsongs from their

superposition. Young songbirds learn songs by mimicking adult birds' songs (*Tchernichovski et*

al., 2001; Woolley, 2012; Lipkind et al., 2013; Lipkind et al., 2017). A study reported that

neurons in songbirds' higher auditory cortex exhibit a teacher specific activity (*Yanagihara,*

Yazaki-Sugiyama, 2016). One can imagine those neurons correspond to the expectation of

hidden sources (u), as considered in this study. Importantly, the natural environment that

young songbirds encounter is dynamic, as we considered in Fig. 5. Therefore, the conventional

BSS setup, which assumes a static environment or context, is not suitable for explaining this

problem. It is interesting to consider that young songbirds might employ some computational

mechanism similar to the EGHR to distinguish a teacher song from other songs in a dynamically

changing environment.

It is worth noting that the application of standard ICA algorithms to high-pass filtered inputs

cannot solve the multi-context BSS problem. This is because context-dependent changes in

the mixing matrix not only change the means of the inputs, which can be removed by high-

pass filtering, but also change the gain of how each source fluctuations are propagated to input

fluctuations. Hence, the difference in contexts cannot be expressed as a linear ICA problem

after high-pass input filtering. Therefore, selective extraction of context-invariant features is

an advantage of the EGHR. Moreover, if provided with redundant input, the EGHR can solve

multi-context BSS even if the context changes continuously in time, as we demonstrated in

Figs. 4, 5.

Biological neural networks implement an EGHR-like learning rule. The main ingredients of

the EGHR are Hebbian plasticity and the third scalar factor that modulates it. Hebbian plasticity

occurs in the brain depending on the activity level (*Bliss, Lomo, 1973; Dudek and Bear, 1992*),

spike timings (*Markram et al., 1997; Bi, Poo, 1998; Zhang et al. 1998; Feldman, 2012*), or burst

timings (*Butts et al. 2007*) of pre- and post-synaptic neurons. In contrast, the third scalar factor

can modify the learning rate and even invert Hebbian to anti-Hebbian plasticity (*Paille et al.,*

2013), similarly to what we propose for the EGHR. In general, such a modulation forms the

basis of a three-factor learning rule, a concept that has recently received attention, see
(*Pawlak et al., 2010; Frémaux, Gerstner, 2016; Kuśmierz et al., 2017*) for reviews, and is
supported by experiments on various neuromodulators and neurotransmitters, such as
dopamine (*Reynolds et al., 2001; Zhang et al., 2009; Yagishita et al., 2014*), noradrenaline
(*Salgado et al., 2012; Johansen et al., 2014*), and GABAergic input (*Paille et al., 2013; Hayama
et al., 2013*), as well as glial factors (*Ben Achour, Pascual, 2010*). Those third factors may
encode reward (*Izhikevich, 2007; Florian, 2007; Legenstein et al., 2008; Urbanczik, Senn, 2009;
Frémaux et al., 2010*), likelihood (*Brea et al., 2013*), novelty/surprise (*Rezende, Gerstner, 2014*),
or error from a prior belief (*Isomura, Toyoizumi, 2016; Isomura, Toyoizumi, 2018*). Importantly,
the EGHR only requires such local signals, which neurons can access via their synaptic
connections. Furthermore, a study using *in vitro* neural networks suggested that neurons
perform simple BSS using a plasticity rule that is different from the most basic form of Hebbian
plasticity, by which synaptic strengths are updated purely as a product of pre- and postsynaptic
activity (*Isomura et al., 2015*). A candidate implementation of the EGHR can be made for
cortical pyramidal cells and inhibitory neurons; the former constituting the EGHR output
neurons and encoding the expectations of hidden sources, and the latter constituting the third
scalar factor and calculating the nonlinear sum of activity in surrounding pyramidal cells. This
view is consistent with the circuit structure reported for the visual cortex (*Hofer et al., 2011;
Harris, Mrsic-Flogel, 2013*). These empirical evidences support the biological plausibility of the
EGHR as a candidate model of neuronal BSS.

A local computation of the EGHR is highly desirable for neuromorphic engineering (*Chicca
et al., 2014; Merolla et al., 2014; Neftci, 2018*). The EGHR updates synapses by a simple
product of pre- and postsynaptic neurons' activity and a global scalar factor. Because of this,
less information transfer between neurons is required, compared to conventional ICA methods
that require non-local information (*Bell, Sejnowski, 1995; Bell, Sejnowski, 1997; Amari et al.,
1996*), all-to-all plastic lateral inhibition between output neurons (*Foldiak, 1990; Linsker, 1997*),
or an additional processing step for decorrelation (*Hyvarinen, Oja, 1997*). The simplicity of the
EGHR is a great advantage when implemented in a neuromorphic chip because it can reduce
the space for wiring and the energy consumption. Furthermore, unlike the conventional ICA
algorithms that assume an equal number of input and output neurons, a neuromorphic chip
that employs the EGHR with redundant inputs would perform BSS in multiple contexts, as
allowed by the network memory capacity, without requiring readaptation. The generalization
capability of the EGHR, as demonstrated in Fig. 6, is an additional benefit, as the EGHR captures
the common features shared across training contexts to perform BSS in inexperienced test
contexts.

Notably, although we considered the linear BSS problem in this study, multi-context BSS can
be extended to a non-linear BSS, in which the inputs are generated through a non-linear
mixture of sources (*Lappalainen, Honkela, 2000; Karhunen, 2001*). To solve this problem, a
promising approach would be to use a linear neural network. A recent study showed that when

the ratio of input-to-source dimensions and source number are large, a linear neural network

can find an optimal linear encoder that separates the true sources through PCA and ICA, thus

asymptotically achieving zero BSS error (*Isomura, Toyoizumi, 2018b*). Because both the

asymptotic linearization and multi-context BSS by the EGHR are based on high-dimensional

sensory inputs, it would be useful to combine them to solve the multi-context, non-linear BSS

problem.

In summary, we demonstrated that the EGHR can retain memories of past contexts and,

once the learning is achieved for every context, it can perform multi-context BSS without

further updating synapses. Moreover, the EGHR can find common features shared across

contexts, if present, and uses them to generalize the learning result to inexperienced contexts.

Therefore, the EGHR will be useful for understanding the neural mechanisms of flexible

inference and sensory representation under dynamically changing environments.

Methods

Model and learning rule

The neural network model and used learning rule (the EGHR) are described in the Results

section.

BSS solution

Suppose that $N_u = N_s$. We define a transform matrix $K^{(k)}$ by

$$K^{(k)} \equiv W A^{(k)}. \tag{8}$$

543 For $N_x \geq N_s$, the ICA for context k is achieved when $K^{(k)}$ is the identical matrix up to permutations

544 and sign-flips. Hence, when column vectors of a block matrix $(A^{(1)}, …, A^{(C)})$ are independent of

545 each other, if and only if $(A^{(1)}, …, A^{(C)})$ is a full-rank matrix, an ICA solution that separates all

546 sources for context $1, …, C$ exists. Namely, W achieves the multi-context BSS when it satisfies

$$\left(K^{(1)}, …, K^{(C)}\right) = W\left(A^{(1)}, …, A^{(C)}\right) = \left(\Omega^{(1)}, …, \Omega^{(C)}\right), \tag{9}$$

where $\Omega^{(k)}$ is an $N_u \times N_s$ matrix equivalent to the identity matrix up to permutations and sign-

flips. Regarding the *i*-th row of matrix $K^{(k)}$, as denoted by a row vector $\left(K_{i1}^{(k)}, …, K_{iN_s}^{(k)}\right)$, the

achievement of ICA is justified when one element is one and the others are zero. Thus, there

are many candidate sets of $\left(W_{i1}, …, W_{iN_x}\right)$ that can achieve ICA, because N_x is larger than N_s.

Our numerical analyses showed that among these potential solutions, the one that is the

nearest to the solution for the previous context is likely to be chosen. This can be understood
as follows: when the network finds an ICA solution for all contexts, the error (i.e., cost function
of the EGHR) is minimized including transient periods between two contexts; hence, according
to gradient descent, synaptic strengths converge to such a solution as training progresses.
Owing to this mechanism, the initial errors converge to zero when previously experienced
environments are provided as stimuli.

Simulation protocols

For figure 5: Two birdsongs were downloaded from Xeno-canto ([https://www.xeno-](https://www.xeno-canto.org/132149)
[canto.org/132149](https://www.xeno-canto.org/132149), https://www.xeno-canto.org/133054). Two hidden sources were created
by trimming the first 60 s of these songs (with 4410-Hz time resolution) and normalizing them,
to ensure each source sequence had zero mean and unit variance. During the training, the
song sequences were repeated. To add stochasticity, a hidden source was defined by the sum
of a song and a white-noise sequence generated by a Laplace distribution. The mixing matrix
was defined by 6 × 2 random matrices, $A^{(0)}, A^{(1)}$, and a rotation matrix, $R(t) \equiv$
$(\cos \omega t, -\sin \omega t; \sin \omega t, \cos \omega t)$. The angular frequency ω was randomly set as -0.1π, 0,
or 0.1π [rad/s], by following Poisson process with a transition probability of 1/8820. The
training time and learning rate were defined by T = 4410 × 300 [step] and $\eta = 1 \times 10^{-7}$.

For figure 6: Ten birdsongs were downloaded from Xeno-canto (the URLs are
https://www.xeno-canto.org/****** where ****** was replaced with the following numbers:
27060, 64735, 67307, 110303, 121326, 121691, 126481, 132149, 133054, 133862). Ten
hidden sources were created in the same manner as described above. The mixing matrix was
defined by 100 × 10 random matrices $A^{(0)}, A^{(1)}, A^{(2)}, A^{(3)}, A^{(4)}$, where $\hat{A}^{(1)}, \dots, \hat{A}^{(4)}$ were
randomly generated and $A^{(0)}, \dots, A^{(4)}$ were defined by $A^{(0)} = (\hat{A}^{(1)} + \hat{A}^{(2)} + \hat{A}^{(3)} + \hat{A}^{(4)})/$
4 and $A^{(k)} = \hat{A}^{(k)} - A^{(0)}$, for k = 1, …, 4. This treatment served to ensure that $A^{(1)}, \dots, A^{(4)}$
do not involve common features across contexts. The training comprised 120 sessions, with
each session continued for T = 4410 × 600 [step]. The context vector v randomly chose one of
the following ten vectors, v = (1,0,0,0), (½,½,0,0), (0,1,0,0), (0,½,½,0), (0,0,1,0), (0,0,½,½),
(0,0,0,1), (½,0,0,½), (½,0,½,0), (0,½,0,½), at the beginning of each session and maintained the
value during the session. The learning rate was defined by $\eta = 2 \times 10^{-7}$. For the test, 20
randomly generated vectors were used, and their elements were randomly sampled from [0,1]
and then normalized to satisfy $v_1 + v_2 + v_3 + v_4 = 1$.

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Author Contributions

593   Conceived the idea: T.I. and T.T. Performed the analyses: T.I. and T.T. Wrote the paper: T.I.
594 and T.T.

Additional Information

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