

LIF neuron model
$$C_{\rm m} \frac{du}{dt} = g_{\rm l}(E_{\rm l} - u) + I^{\rm syn} +$$

$$\theta = \tau_{\rm syn} \rightarrow {\rm synaptic memory!}$$

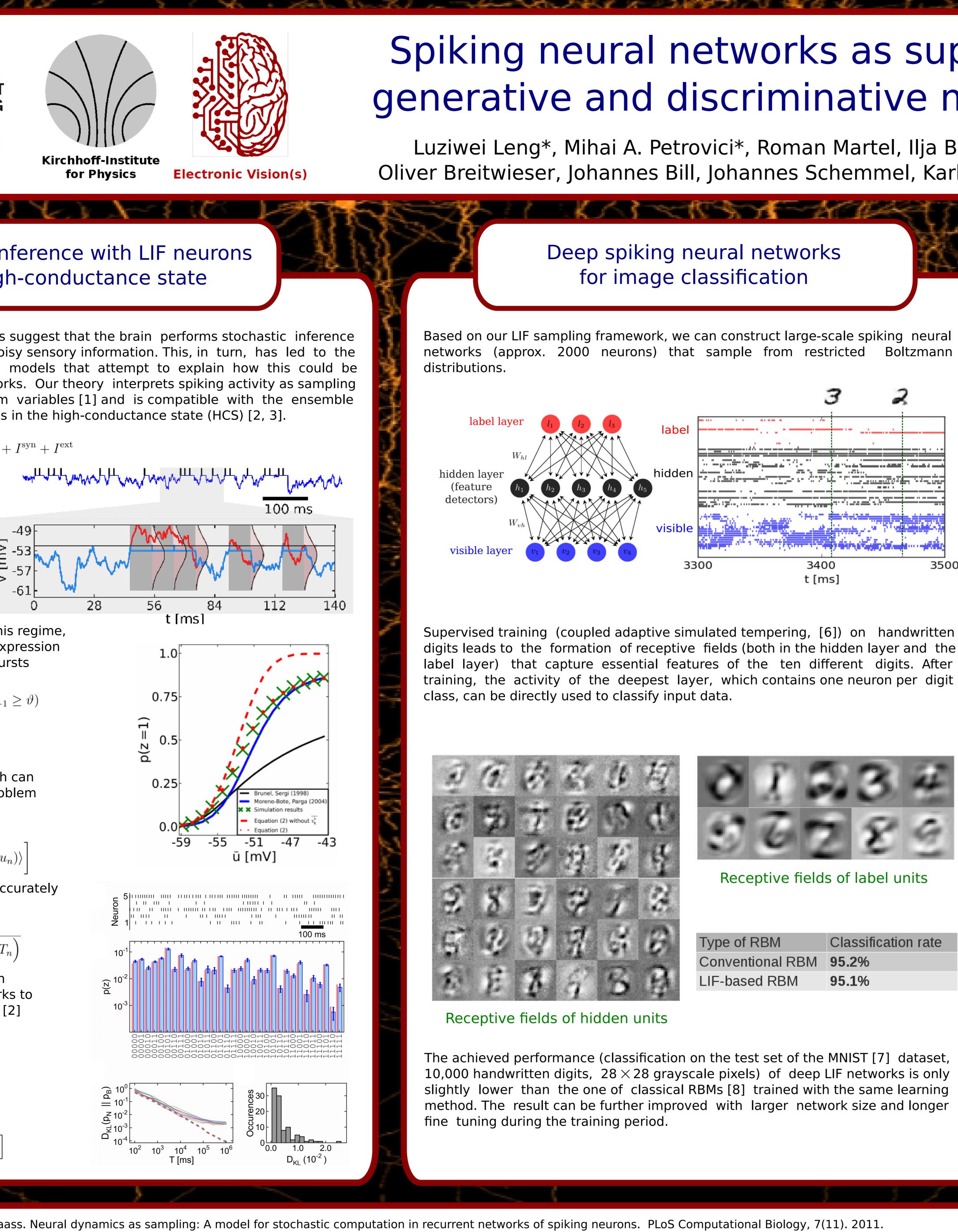
$$P_{n} = \left(1 - \sum_{i=1}^{n-1} P_{i}\right) \int_{\vartheta}^{\infty} du_{n-1} p(u_{n-1} | u_{n-1} \ge \vartheta)$$
$$\left[\int_{-\infty}^{\vartheta} du_{n} p(u_{n} | u_{n-1})\right]$$

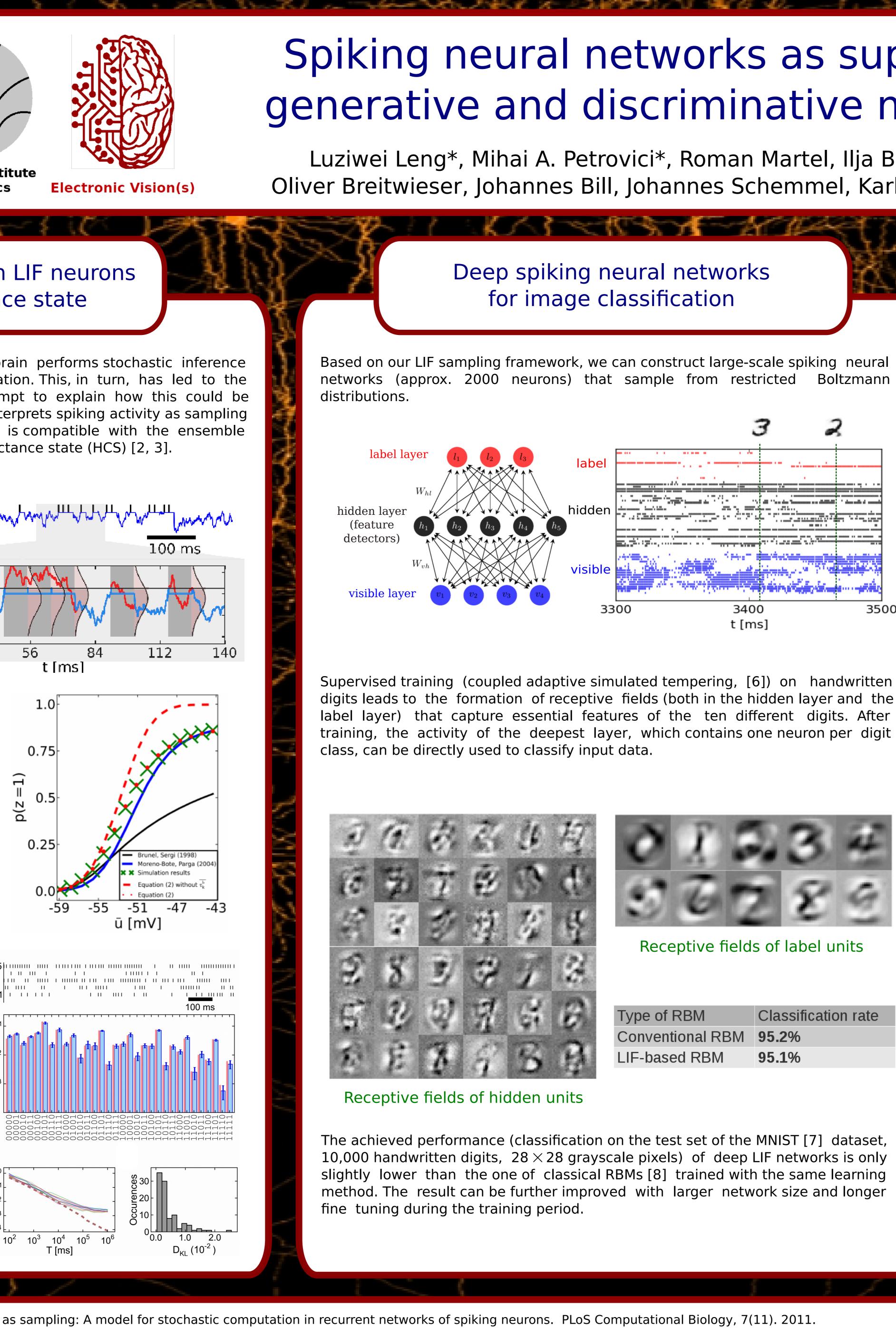
$$T_{n} = \int_{\vartheta}^{\infty} du_{n-1} p(u_{n-1} | u_{n-1} \ge \vartheta) \\ \left[\int_{-\infty}^{\vartheta} du_{n} p(u_{n} | u_{n} < \vartheta, u_{n-1}) \langle T(\vartheta, u_{n}) \rangle \right]$$

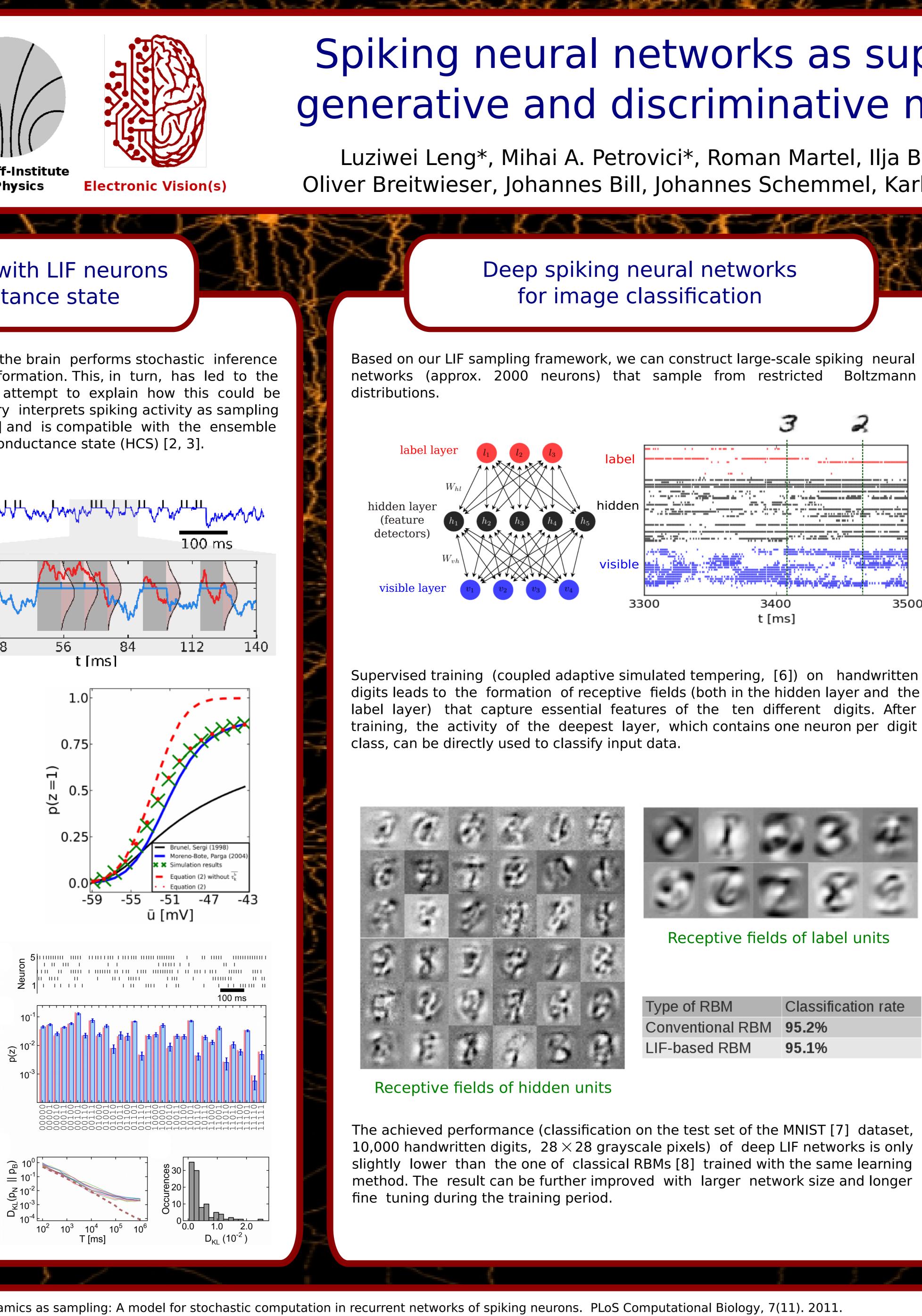
$$p(z=1) = \frac{\sum_{n} P_n \cdot n \cdot \tau_{\text{ref}}}{\sum_{n} P_n \cdot \left(n\tau_{\text{ref}} + \sum_{k=1}^{n-1} \overline{\tau_k^{\text{b}}} + T_n\right)}$$

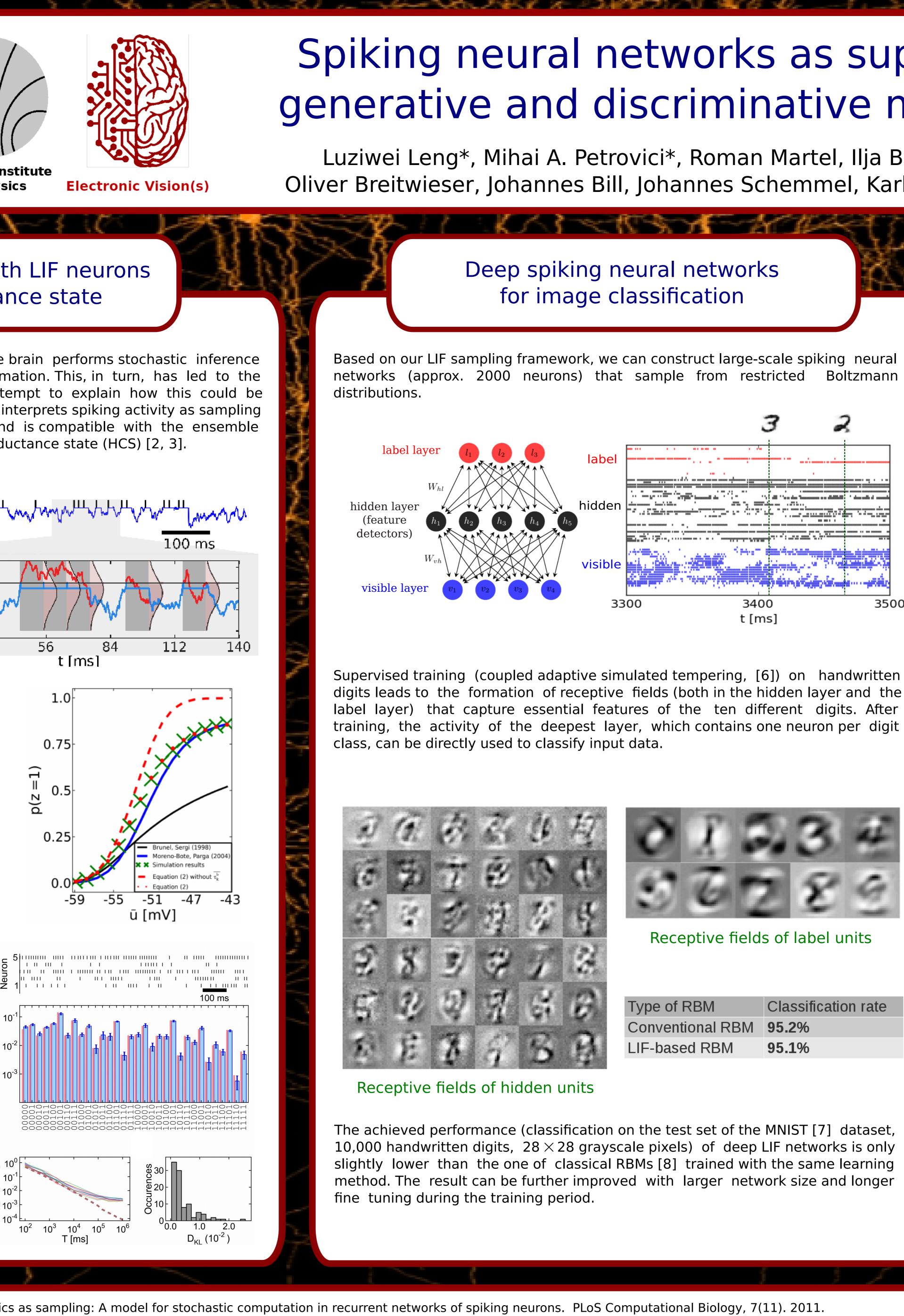
$$u_{k}(t) = \log \frac{p(z_{k} = 1 | z_{\backslash k})}{p(z_{k} = 0 | z_{\backslash k})}$$
$$b_{k} = (\bar{u}_{k}^{b} - \bar{u}_{k}^{0})/\alpha$$
$$W_{kj} = \frac{1}{\alpha C_{m}} \frac{w_{kj} \left(E_{kj}^{rev} - \mu\right)}{1 - \frac{\tau_{syn}}{\tau_{eff}}}$$
$$\left[\tau_{syn} \left(e^{-1} - 1\right) - \tau_{eff} \left(e^{-\frac{\tau_{syn}}{\tau_{eff}}} - 1\right)\right]$$

[2] M. A. Petrovici, J. Bill, I. Bytschok, J. Schemmel, and K. Meier, Stochastic inference with deterministic spiking neurons. arXiv preprint arXiv:1311.3211, 2013. [3] M. A. Petrovici, Bytschok, I., Bill, J., Schemmel, J. & Meier, K. The high-conductance state enables neural sampling in networks of LIF neurons. 24th Annual Meeting of the CNs, 2015. [4] R. Moreno-Bote and N. Parga. Role of synaptic filtering on the firing response of simple model neurons. Physical Review Letters, 92(2):028102, 2004. [5] N. Brunel and S. Sergi. Firing frequency of leaky integrate-and-fire neurons with synaptic current dynamics. Journal of Theoretical Biology, 195:87–95, 1998. [6] R. Salakhutdinov, Learning deep Boltzmann machines using adaptive MCMC, in Proceedings of the 27th International Conference on Machine Learning (ICML-10), pp. 943–950, 2010. [7] Y. LeCun, and C. Cortes, The MNIST database of handwritten digits, 1998. [8] G. E. Hinton, Training products of experts by minimizing contrastive divergence. Neural Computation, 14 (8), 17711800, 2002. [9] G. Fuhrmann, I. Segev, H. Markram, and M. Tsodyks. Coding of temporal information by activity-dependent synapses. Journal of Neurophysiology, 87(1): 140–148, 2002.









Brain Scales ScaleS

	Classification rate
I RBM	95.2%
BM	95.1%

In classical RBMs, statistics are typically gathered by Gibbs sampling:

$$p(z_k^{(t)} \to z_k^{(t+1)}) = p(z_k | z_{\backslash k})$$

This algorithm has a distinct disadvantage when dealing with high-dimensional multimodal distributions, where it often gets trapped in local minima due to deep troughs in the energy landscape that appear during training. It is for thi reason that conventional RBMs that may perform very well as discriminative models are, at the same time, rather poor generative models of the learned data. While methods exist that alleviate this problem [6], they usually come at a highly increased computational cost.

	-53
5	-54

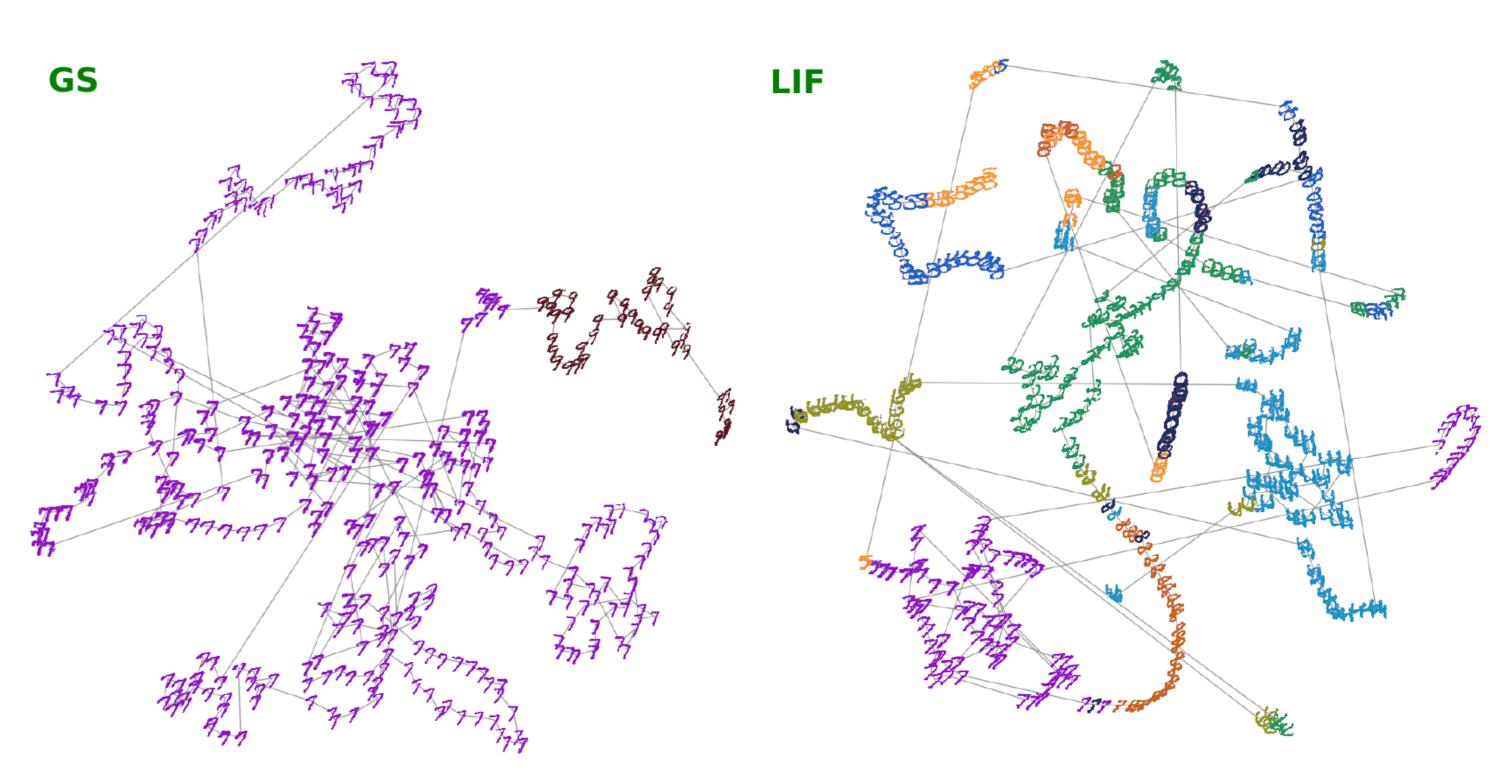
$$\frac{\mathrm{d}R}{\mathrm{d}t} = \frac{(1-R)}{\tau_{\mathrm{rec}}} - U_{SE}R\delta(t-t_{sp})$$
-54
$$\frac{\mathrm{d}U_{SE}}{\mathrm{d}t} = -\frac{U_{SE}}{\tau_{\mathrm{rec}}} + U1 \cdot (1-U_{SE}) \cdot \delta(t-t_{sp})$$
-55

Short-term plasticity [9] allows the

activity-dependent modulation of

synaptic interaction.

With an appropriate choice of parameters, the expectation value of synaptic weights can be maintained, ensuring that the energy landscape remains, on average, unchanged, while at the same time modulating the strength of the active local attractor of the network. The initial potentiation deepens the local energy well, allowing the visible layer of the network to produce a clearer image. The subsequent depression increases the energy of the network, thereby catapulting it out of the local minimum.



Short-term plasticity enables LIF networks to travel efficiently through the energy landscape and thereby attain a generative performance that significantly surpasses the one achievable by Conventional Gibbs sampling. This distinct feature of biological neural networks allows them to Simultaneously become good generative and discriminative models of learned data. Our finding points towards an important advantage of spike-based computation and communication, which is Relevant in any scenario where spiking neural networks need to be able to escape local attractors.

