# Accurate online training of dynamical spiking neural networks through Forward Propagation Through Time

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### **10** ABSTRACT

With recent advances in learning algorithms, recurrent networks of spiking neurons are achieving performance competitive with vanilla recurrent neural networks. Still, these algorithms are limited to small networks of simple spiking neurons and modest-length temporal sequences, as they impose high memory requirements, have difficulty training complex neuron models, and are incompatible with online learning. Here, we show how recently developed 'Forward-Propagation-Through-Time' (FPTT) learning combined with novel Liquid Time-Constant spiking neurons resolves these limitations. Applying FPTT to networks of such complex spiking neurons, we demonstrate online learning of exceedingly long sequences while outperforming current online methods and approaching or outperforming offline methods on temporal classification tasks. FPTT's efficiency and robustness furthermore enables us to directly train a deep and performant spiking neural network for joint object localization

<sup>11</sup> and recognition, demonstrating the ability to train large-scale dynamic and complex spiking neural network architectures. With recent advances in learning algorithms, recurrent networks of spiking neurons are achieving performance competitive with vanilla recurrent neural networks. Still, these algorithms are limited to small networks of simple spiking neurons and modest-length temporal sequences, as they impose high memory requirements, have difficulty training complex neuron models, and are incompatible with online learning. Here, we show how recently developed 'Forward-Propagation-Through-Time' (FPTT) learning combined with novel Liquid Time-Constant spiking neurons resolves these limitations. Applying FPTT to networks of such complex spiking neurons, we demonstrate online learning of exceedingly long sequences while outperforming current online methods and approaching or outperforming offline methods on temporal classification tasks. FPTT's efficiency and robustness furthermore enables us to directly train a deep and performant spiking neural network for joint object localization and recognition, demonstrating the ability to train large-scale dynamic and complex spiking neural network architectures.

### 12 Introduction

The binary, event-driven and sparse nature of communication between spiking neurons in the brain holds great promise for flexible and energy-efficient AI. Recent work has demonstrated effective and efficient performance from spiking neural networks (SNNs)<sup>1</sup>, enabling competitive and energy-efficient applications in neuromorphic hardware<sup>2</sup> and novel means of investigating biological neural architectures<sup>3,4</sup>. This success stems principally from the use of approximating surrogate gradients<sup>5,6</sup> to integrate networks of spiking neurons into auto differentiating frameworks like Tensorflow and Pytorch<sup>7</sup>, enabling the application of standard learning algorithms and in particular Back-Propagation Through Time (BPTT).

The imprecision of the surrogate gradient approach however expounds on the existing drawbacks of BPTT. In particular, BPTT has a linearly increasing memory cost as a function of sequence length T,  $\Omega(T)$  and suffers from vanishing or exploding backpropagating gradients, which limits its applicability on long time sequences<sup>8</sup> and large-scale SNN models<sup>9</sup>. Alternative approaches like real-time recurrent learning (RTRL)<sup>10</sup> similarly exhibit excessive computational complexity, and low complexity approximations to BPTT like e-prop<sup>11</sup> or OSTL<sup>12</sup> at best approach BPTT performance. Training on long temporal sequences in SNNs is of particular importance when the tasks require a high temporal resolution, for instance to match the physical characteristics of low-latency clock-less neuromorphic hardware<sup>2,13</sup>.

Kag et al.<sup>8</sup> recently introduced an algorithm for online learning in recurrent networks, Forward Propagation Through Time
 (FPTT), demonstrating better generalization on many temporal classification benchmark tasks compared to BPTT. In particular,
 FPTT improved over BPTT on long sequence training in Long Short-Term Memory networks (LSTMs). FPTT differs from

<sup>29</sup> BPTT in that it does not calculate a gradient through time, and instead considers learning-through-time as a coordinated

30 consensus problem. Using regularized synaptic tags, FPTT enables immediate, online learning in RNNs similar to feedforward

networks, eliminating the problematic dependence of the gradient calculation in BPTT on the products of partial gradients

<sup>32</sup> along the time dimension: FPTT exhibits linear  $\Omega(T)$  computational cost per sample. For training recurrent SNNs however, as

 $_{33}$  we demonstrate, a straightforward application of FPTT on long sequence training does not improve performance as it does in<sup>8</sup>,

and we observed this also with vanilla RNNs. Therefore, we deduce that FPTT particularly benefits from the gating structure

<sup>35</sup> inherent in LSTM-style gated RNNs, which is lacking in vanilla RNNs and SRNNs.

Taking inspiration from the concept of Liquid Time-Constant  $(LTCs)^{14}$ , we introduce a class of spiking neurons, the Liquid

<sup>37</sup> Time-Constant Spiking Neuron (LTC-SN), where time-constants internal to the neuron are dynamic and input-driven in a

<sup>38</sup> learned fashion, resulting in functionality similar to the gating operation in LSTMs. We integrate these neurons in networks

that are trained with FPTT and demonstrate that the resulting LTC-SNNs outperform various SNNs trained with BPTT on long sequences while enabling online learning and drastically reducing memory complexity. We show this for several classical

<sup>41</sup> benchmarks that can easily be varied in sequence length, like the Add-task and the DVS-GESTURE benchmark<sup>15, 16</sup>. We

<sup>41</sup> also show how FPTT-trained LTC-SNNs can be applied to large convolutional SNNs, where we demonstrate state-of-the-art

43 for online learning in SNNs on a number of standard benchmarks (S-MNIST, R-MNIST, DVS-GESTURE) and to near

44 (Fashion-MNIST, DVS-CIFAR10) or exceeding (PS-MNIST, R-MNIST) state-of-the-art performance as obtained with offline

45 BPTT.

Finally, the training and memory efficiency of FPTT enables us to directly train SNNs in an end-to-end manner at network sizes and complexity that was previously infeasible. We demonstrate this in a large-scale You-Look-Only-Once (YOLO)

LTC-SNN architecture for object detection on the Pascal Visual Object Classes (Pascal VOC) dataset<sup>17</sup>. Object detection is a challenging task, as it involves accurate multi-object identification and precise bounding box coordinate computation;

<sup>49</sup> is a challenging task, as it involves accurate multi-object identification and precise bounding box coordinate computation; <sup>50</sup> previous SNN approaches have been limited to either ANN-to-SNN conversions<sup>18–20</sup>, requiring many thousands of time-steps

<sup>50</sup> previous SNN approaches have been limited to either ANN-to-SNN conversions<sup>10–20</sup>, requiring many thousands of time-steps <sup>51</sup> at inference time, or small scale and inefficient SNNs with performance far removed from that of modern ANNs <sup>21</sup>. Our

at inference time, or small scale and inefficient SNNs with performance far removed from that of modern ANNs<sup>21</sup>. Our FPTT-trained YOLOv4<sup>22</sup> implementation – SPYv4 – uses 21 layers, 6.2M LTC spiking neurons, and 14M parameters to

<sup>32</sup> achieve state-of-the-art for SNNs, exceeding the performance of converted ANNs while achieving extremely low latency.

With FPTT and LTC spiking neurons, we demonstrate end-to-end online training of large and high-performance SNNs

<sup>55</sup> comprised of complex spiking neuron models that were previously infeasible.

### 56 Related Work

The problem of training recurrent neural networks has an extensive history $^{23-25}$ . To account for past influences on current 57 activations in a recurrent network, the network can be unrolled, and errors are computed along the paths of the unrolled network. 58 The direct application of error-backpropagation to this unrolled graph is known as Backpropagation-Through-Time<sup>23</sup>. BPTT 59 needs to wait until the last input of a sequence before being able to calculate parameter updates and, as such, cannot be applied 60 in an online manner. Alternative online learning algorithms for RNNs have been developed, including Real-Time Recurrent 61 Learning (RTRL)<sup>10</sup> and approximations thereof<sup>26</sup>. RTRL however is prohibitive in time and memory complexity, and while 62 approximations improve complexity (see<sup>12</sup>, and Table 1a), they yield variable and task-dependent accuracy deficits compared 63 to exact gradients<sup>11,27</sup>. 64

Spiking neural networks are neural networks composed of spiking neurons: stateful neural units that communicate using 65 binary values, i.e. spikes. Their state is determined by current and past inputs, and this state then determines the (binary) value 66 of the emitted output through a spiking mechanism. The discontinuity of the spiking mechanism challenges the application 67 of error-backpropagation, which can be overcome using continuous approximations<sup>5,28</sup>, so-called "surrogate gradients"<sup>6</sup>. 68 Recurrent SNNs trained with surrogate gradients and BPTT now achieve competitive performance compared to classical 69 RNNs<sup>1, 16, 29</sup>. In these and other studies, more intricate spiking neuron models, like those including adaptation, outperformed 70 less complex models such as standard Leaky-Integrate-and-Fire neurons<sup>11</sup>. Additionally, training internal spiking neuron's 71 model parameters like the time-constants of adaptation and membrane-potential decay then further improves performance<sup>1,29</sup>. 72 Still, the application of BPTT in SNNs has several drawbacks: in particular, BPTT accumulates the approximation error of 73 surrogate gradients along time. Moreover, the spike-triggered reset of some state variables in typical spiking neuron models (e.g. 74 the membrane potential) causes a vanishing gradient when applying BPTT. We found these effects to be particularly problematic 75 when training networks with complex and more biologically detailed neuron models like Izhikevich and Hodgkin-Huxley 76 models (unpublished). Furthermore, because the SNN training accuracy heavily depends on hyperparameters, obtaining 77 convergence using BPTT in SNNs is non-trivial. 78

For spiking neural networks, approximations to BPTT like e-prop<sup>11</sup> and Online Spatio-Temporal Learning (OSTL)<sup>12</sup> achieve linear time complexity and have proven effective for many small-scale benchmark problems, ATARI GAMES and also large-scale networks like cortical microcircuits<sup>30</sup>. In terms of trained accuracy, however, none of these approximations have been shown to outperform standard BPTT and, applications to deeper networks use approximate spatial credit assignment approaches like learning-to-learn<sup>11</sup>.



**Figure 1. a**, Computational complexity of gradients, parameter updates and memory storage per sample, with *N* the batch-size. Computational expense increases as the length *T* of the sequence grows. **b**, Roll-out of the computational graph of a spiking neuron as used for BPTT (left) and FPTT (right); **c**, Illustration of LIF (left) and LTC (right) spiking neurons as recurrent network structures. **d**, F-I curve of a LTC spiking neuron for combinations of effective LTC weights  $W_{\tau} = \{W_{\tau_m}, W_{\tau_{adp}}\}$ :  $W_{\tau_m} = \{-1 - 0.5, 0.5, 1\}$  and  $W_{\tau_{adp}} = \{-1 - 0.5, 0.5, 1\}$  associated with respective dynamic time-constant functions  $\sigma$  subject to input current *I*. **e**, Response firing rate *Fr* of LTC spiking neuron to varying current input *I* (top) for combinations of effective LTC weights  $W_{\tau_m} = \{-1, 1\}$  and  $W_{\tau_{adp}} = \{-1, 1\}$ .

### 84 FPTT in SNNs

As introduced in<sup>8</sup>, FPTT can be implemented directly on the computational graph of SNNs, similar to BPTT approaches. This is illustrated in Fig. 1b; the gradient calculation and corresponding algorithms are given in the Methods section and Alg. SA1. FPTT intuitively provides a more robust and efficient gradient approximation for spiking recurrent neural networks than BPTT, as FPTT simplifies the complex gradient computation path in SNNs. This potentially lessens surrogate gradients' cumulative effect, avoiding or reducing the gradient vanishing or explosion problem.

As we will show, FPTT applied directly to SNNs like the Adaptive Spiking Recurrent Neural Networks (ASRNNs)<sup>1</sup> 90 converges but without the learning improvements reported for RNN architectures<sup>8</sup> – and we observed this also for vanilla 91 non-spiking RNNs (not shown). As FPTT was successfully applied to RNNs with gating structures (LSTM), we introduce the 92 Liquid Time-Constant Spiking Neuron (LTC-SN) as a spiking neuron model with a similar gating structure. We observe that in 93 spiking neurons, the time-constant of the membrane potential acts similar to the forget-gate in LSTMs; the LSTM forget-gate 94 however is dynamically controlled by learned functions of inputs. Inspired also by Hasani et al.<sup>14</sup>, the LTC-SN's internal 95 time-constants are a learned function of the inputs and hidden states of the network (illustrated in Fig. 1c; see also Methods). 96 I.e., for adapting spiking neurons, the time-constant of the membrane potential decay,  $\tau_m$  and the time-constant of the adaptation 97 decay,  $\tau_{adp}$  can be made dynamic. A spiking neuron with such varying internal dynamics can respond in flexible and unexpected 98 ways to input currents: as shown in Fig. 1d, depending on the effective weights  $W_{\tau}$ , the current-spike-frequency response curve 99 can be muted-and-saturating, near-linear, or rapidly increasing-and-then-saturating; when subject to a dynamically varying 100

<sup>101</sup> input current, a higher input current into LTC-SN units can result in a reduced firing rate, and transient dynamics can be absent

<sup>102</sup> or present (Fig. 1e, samples of trained behavior are shown in Fig. S5).

### 103 Experiments

We demonstrate the effectiveness of FPTT training for LTC-SNNs on several well-known temporal classification benchmarks,
 including the Add-task as in<sup>8</sup> and several established SNN benchmarks (the DVS-GESTURE and DVS-CIFAR10 classification
 tasks, and the Sequential, Sequential-Permuted, rate-based and Fashion MNIST classification tasks). We moreover demonstrate
 how the memory efficiency of FPTT enables training large-scale LTC-SNNs for applications like object localization.

The Add-task<sup>31</sup> is used to evaluate the ability of RNN architectures to maintain long range memory. An example data 108 point consists of two sequences  $(x_1, x_2)$  of length T and a target label y. The sequence  $x_1$  contains values sampled uniformly 109 from [0, 1],  $x_2$  is a binary sequence containing only two 1s, and the label y is the sum of the two entries in the sequence  $x_1$ , 110 where  $x_2 = 1$ . The **IBM DVS Gesture** dataset<sup>15</sup> consists of 11 kinds of hand and arm movements of 29 individuals under 111 three different lighting conditions captured using a DVS128 camera. DVS-CIFAR10 is a widely used dataset in neuromorphic 112 vision, where the event stream is obtained by displaying moving images of the CIFAR-10 dataset<sup>32</sup>. The Sequential and 113 Permuted-Sequential MNIST (S-MNIST, PS-MNIST) datasets were developed to measure sequence recognition and memory 114 capabilities of learning algorithms: the S-MNIST dataset is obtained by reshaping the classical MNIST 28x28 pixel images 115 into a set of one-dimensional sequences consisting of 784 time-steps per sample, where pixels are then sequentially entered 116 one-by-one as input to the network; the PS-MNIST dataset is generated by performing a fixed permutation on the S-MNIST 117 dataset. Theoretically and in practice, PS-MNIST is a more complex classification task than S-MNIST because it lacks 118 temporally correlated patterns. The rate-coded MNIST (R-MNIST) is an SNN-specific benchmark where a biologically 119 inspired encoding method is used to generate the network input that produces streaming events (a spike train), by encoding 120 the grey values of the image with Poisson rate-coding<sup>33</sup>. We also applied FPTT-trained LTC-SCNNs to the traditional static 121 MNIST and Fashion-MNIST datasets for comparison with other models trained offline. Here, we input pixel values directly 122 as injected current into the spiking input layer of the network, repeated 20 times to mimic a constant input stream. For object 123 localization, we used the PASCAL VOC benchmark<sup>17</sup> which is comprised of standard 416×416 RBG images with 20 different 124

<sup>125</sup> objects and corresponding bounding box locations.

FPTT-SNN requires Liquid Time-Constant Spiking Neurons. We apply both BPTT and FPTT to the Add-task as originally studied in<sup>8</sup> to illustrate the need for more complex spiking neurons like LTC-SNs when applying FPTT learning. We do this for various network types, including non-spiking LSTMs as a baseline, ASRNNs<sup>1</sup>, and LTC-SNNs.

For a long adding sequence of length 1000, example loss-curves are plotted in Fig. 2a and averaged converged losses in Fig. 2b. We find that as in<sup>8</sup>, a standard LSTM network trained with BPTT fails to converge to zero loss, while the same network does converge using FPTT. For SNNs, we find that ASRNNs trained with either FPTT or BPTT do not fully converge, and for the LTC-SNNs trained with BPTT learning rapidly diverges due to exploding gradients. LTC-SNNs trained with FPTT however successfully minimize the loss: ablating the LTC-SN dynamics, we find that the input-dependent dynamic gating of the membrane-potential time-constant is critical for convergence (Tab. S4), and the memory provided by the LTC-SN self-recurrence does in fact suffice for solving this task for shorter sequences, though not for longer ones (Fig. S4).

**FPTT allows for longer sequence training.** Next, we study the ability of FPTT-trained LTC-SNNs to learn increasingly long sequences. For this, we use the DVS-GESTURE dataset and systematically investigate the performance of a fixed architecture shallow SRNN model on increasingly many frames sampled from the same gesture signals as in<sup>16</sup>, ranging from 20 to 1000 frames. For each frame-encoded dataset, we train various networks types for a fixed number of epochs with an identical number of neural units, and report the best performance for BPTT and FPTT trained networks; networks either converged before the final epoch or diverged (Fig S1a,b).

The results for different sampling frequencies are shown in Fig. 2c. Of all methods and networks, the LTC-SNN trained using FPTT achieves the best accuracy in all cases, also outperforming standard (BPTT-trained) LSTMs. Furthermore, the FPTT-trained LTC-SNNs and the ASRNNs exhibit constant accuracy over the whole range of sequence lengths, where the LTC-SNNs consistently outperform the ASRNNs.

In contrast, the accuracy of both LTC-SNNs and ASRNNs trained with BPTT quickly deteriorates as sequence length increases. For the LTC-SNNs, the networks failed to converge for frame-lengths 200 and 500, and best validation accuracy is reported in Fig. 2c. For the baseline standard LSTM, this effect is also there, albeit more moderate, as performance decreases from 88.9% at 100 frames to 82.5% at 500 frames. This suggests that indeed the gradient approximation errors in SNNs add up when training with BPTT. The plot also illustrates the memory-intensiveness of BPTT: when applied to the 1000 frame-length data, GPU-memory (24GB) was insufficient for training LSTM, ASRNN and LTC-SNN.

<sup>152</sup> Comparing sparsity (average firing rate) for the different SNN models, we find no meaningful differences (Fig. S1c); <sup>153</sup> parameter-matched networks showed similar performance as unit-matched networks, and the inclusion of an auxiliary loss<sup>8</sup>



**Figure 2. a**, Example loss-curves for networks trained on the Add-task with sequence length T = 1000. The loss of BPTT-trained LTC-SNN becomes *NaN* after 2000 iterations. **b**, corresponding average loss and std of the last 100 training iterations (obtained from 5 runs). **c**, Plot of test accuracy for BPTT and FPTT trained shallow networks on the DVS-GESTURE dataset. Accuracy bar average and std over 3 runs and individual data points are inserted; (\*): training diverged; reported accuracy is the best accuracy before divergence; (+): out-of-GPU-memory when training. **d**,**e**, GPU-memory use when training the DVS-GESTURE dataset (d) or R-MNIST dataset (e) for different sequence lengths with BPTT or FPTT.

only aided BPTT-trained LSTMs but not BPTT-trained SRNNs. Making only the membrane-decay time-constant dynamic has
 a small negative impact (ASRNN- Table S1).

FPTT Requires Less Memory. FPTT-trained LTC-SNNs require increasingly less memory as sequence length increases as measured on GPU. For the DVS-GESTURE dataset (Fig. 2d), FPTT memory-use is both less and increases less rapidly compared to BPTT as a function of the number of frames used per data sample. FPTT's increasing memory use can be attributed to the rapidly inflating size of the frame-encoded dataset, increasing in size from 7.4 GB for 20 frames to 368.1 GB for 1000 frames. We validated this by training an LTC-SNN network on the R-MNIST classification problem, where longer sequences are simulated by showing the same sample for an increasing number of frames. We then find, as expected, that the memory required for FPTT training remains fixed. At the same time, BPTT memory-use linearly increases (Fig. 2e).

FPTT with LTC Spiking Neurons improves over Online Approximate BPTT. To demonstrate the power of FPTT as an online
 training method, we used state-of-the-art deep spiking convolutional network architectures (SCNNs) for standard sequential
 benchmarks from the literature and trained these architectures with LTC-SN neurons and FPTT.

In Table 1, we compare the LTC-SNNs to existing state-of-the-art online and offline SNNs. We find that LTC-SCNNs

**Table 1.** Test accuracy of deep SRNNs/SCNNs on various tasks. **Bold-faced** denotes state-of-the-art (SoTa) online performance, *slanted bold* denotes overall SNN state-of-the-art. We apply FPTT-trained LTC-SNNs with one recurrent layer comprised of 512 neurons on the (P)S-MNIST tasks. For RMNIST, an identical network structure as in<sup>12</sup> is used, and for the remaining benchmarks the network structures match<sup>16</sup>.

Task	Online baseline		This work		Offline SoTa	
S-MNIST	-		FPTT+LTC	97.37%	BPTT+ASRNN <sup>1</sup>	<b>98.7</b> %
PS-MNIST	-		FPTT+LTC	93.23%	BPTT+ASRNN <sup>1</sup>	94.3%
RMNIST	$OSTL + SNU^{12,34}$	95.34%	FPTT+LTC	98.63%	BPTT+SNU <sup>34</sup>	97.72%
MNIST	DECOLLE+SNN <sup>35</sup>	98.0%	FPTT+LTC	99.62%	BPTT+PLIF <sup>16</sup>	<b>99.72%</b>
Fashion MNIST	EMSTDP+SNN <sup>36</sup>	85.3%	FPTT+LTC	93.58%	BPTT+PLIF <sup>16</sup>	94.38%
DVS-GESTURE	DECOLLE+SNN <sup>37</sup>	95.54%	FPTT+LTC	97.22%	BPTT+PLIF <sup>16</sup>	97.57%
DVS-CIFAR10	-		FPTT+LTC	73.2%	BPTT+PLIF <sup>16</sup>	74.8%



**Figure 3. a**, Mean Average Precision (mAP) and components for classification and recognition of 20 different kinds of objects in SPYv4 on the PASCAL VOC07 dataset. **b**, An example of object recognition with Spiking-YOLO on a sequence of images. Objects are localized and identified for each image independently. **c**, SPYv4 applied to streaming video. Top: example images, bottom: raster plot of spiking activity for 100 spiking neurons randomly drawn from the shallow and deep layer.

trained with FPTT consistently and substantially outperform SNNs trained with online BPTT approximations like OSTL and

e-prop. Compared to offline BPTT approaches, the FPTT-trained LTC-SNNs achieve state-of-the-art for SNNs (R-MNIST)

<sup>169</sup> or achieve close to similar performance (PS-MNIST,S-MNIST, DVS-GESTURE), DVS-Cifar10); additionally, the memory

requirements for FPTT vs. BTTP trained networks were lower by up to a factor of 5 (Table S3) while the training time was only

<sup>171</sup> slightly longer (Table S4).

### 172 Large-scale Object-detection: Spiking YOLO

<sup>173</sup> The memory efficiency of FPTT-trained LTC-SNNs enables us to train SNNs of comparable complexity as modern ANNs: we

demonstrate this by training a large spike-based object-detection model based on the Tiny YOLO-v4 architecture<sup>22,38</sup>. The

'You-Only-Look-Once' (YOLO) architecture calculates both bounding boxes locations and object identities for all identifiable 175 objects in an image using a single pass through a deep neural network. 176

Our SPiking tiny Yolo-v4 network - SPYv4 - has 19 spiking convolutional layers with about 6.2M spiking neurons, 2 177 convolutional output layers and 14M parameters in total, illustrated in Fig. S2a. This makes it both larger and deeper than 178 previous end-to-end trained spiking models. Training a single time-step in the network requires less than 14GB of GPU-memory, 179 and as BPTT scales linearly with the number of time-steps, learning with BPTT in such a large network over multiple time-steps 180 is infeasible. 181

To carry out object detection, the network uses multiple reads of the input image, e.g. 4 or 8 times, as time-steps in the 182 network to obtain the final result; trained with FPTT, SPYv4 achieves a mean Average Precision (mAP) of 51.38% at 4 reads 183 184 and 53.25% at 8 reads (Fig. 3a) on the VOC dataset (see Methods); Fig. 3b shows examples of the detected and classified objects. Neural activation in the network is highly sparse, with about 10% of neurons active on average at each time-step. 185 When receiving direct camera inputs images, inference achieves about 60 time-steps per second on an NVIDIA RTX3090 186 equipped workstation, corresponding to processing 7 or 15 images per second. Fig. 3c shows example activity from neurons 187 from respectively a shallow (near input) and deep network layer: neurons in the deeper layer fall silent when irrelevant stimuli 188 are shown, while neurons closer to the inputs remain active. Earlier work like Spiking-YOLO<sup>18</sup> achieved mAP 51.83% with 189 8000 simulation time-steps; our SPYv4 network thus outperforms these networks in performance, sparseness, and latency. 190

#### Discussion 191

We showed how a recently proposed training approach for recurrent neural networks, FPTT, can be successfully applied to 192 long sequence learning with recurrent SNNs using Liquid Time-Constant Spiking Neurons. Compared to BPTT, FPTT is 193 compatible with online training, has constant memory requirements, and can learn longer sequences. The increased memory 194 efficiency of FPTT allows for training much larger SNNs as was previously feasible, as we demonstrated in the SPYv4 network 195 for object detection. In terms of accuracy, FPTT outperforms online approximations of BPTT like OSTL and e-prop, and 196 enabled a demonstration of online learning in tasks like DVS-CIFAR10. When training large convolutional LTC-SNNs with 197 FPTT, excellent performance is achieved, approaching or exceeding offline BPTT-based solutions using corresponding network 198 architectures - LTC-SNN specific architecture searches may improve results further. 199

To achieve efficient and accurate online learning with FPTT, we introduced Liquid Time-Constant Spiking Neurons 200 (LTC-SNs), where the neuron's time-constants are calculated as a learned dynamic function of the current state and input. 201 When training on various tasks, BPTT failed to converge when applied to LTC-SNNs on long sequences due to diverging 202 gradients, while FPTT consistently converged. As we speculated, this suggests that FPTT provides for a more robust learning 203 signal. While LTC-SNs provide additional memory in the networks, and LTC-SNNs without recurrent connections were able to 204 solve shorter sequences, though not longer (Fig. S4), the optimal degree of network recurrency remains to be determined and 205 may allow for further efficiency and accuracy improvements<sup>39</sup>. LTC-SNs maintain binary communication between neurons, 206 but impose additional calculations to determine the neuron state. In neuromorphic implementations, LTC-SNs could be 207 implemented as multi-compartment neurons or require novel spiking neuron model implementations. 208

The LTC-SN is inspired by multi-compartment modeling of pyramidal neurons in brains. Pyramidal neurons are known to 209 have complex non-linear interactions between different morphological parts far exceeding the simple dynamics of LIF-style 210 neurons<sup>40,41</sup>, where the neuron's apical tuft may calculate a modulating term acting on the computation in the soma<sup>42</sup> that could 211 act similar to the trainable Liquid time-constants used in this work. In a similar vein, learning rules derived from weight-specific 212 traces may relate to synaptic tags<sup>43,44</sup> and are central to biologically plausible theories of learning working memory<sup>45</sup>. In 213 general, we find that FPTT, unlike BPTT, can also train networks of complex biologically realistic spiking neuron models, like 214 Izhikevich and Hodgkin-Huxley models (e.g. for the DVS-GESTURE task, Table \$5). These considerations suggest variations 215 of FPTT may be potential candidates for temporal credit assignment mechanisms in the brain. As a candidate for biologically 216 plausible learning, the spatial error-backpropagation employed in FPTT would need to be replaced with a plausible spatial credit 217 assignment solution, where we anticipate that at least some of the current proposals<sup>46,47</sup> may be compatible. With such local 218 spatial credit-assignment, FPTT-training of LTC-SNNs can also likely be implemented efficiently on neuromorphic hardware. 219 Taken together, our work suggests that FPTT is an excellent training paradigm for large-scale SNNs comprised of complex 220 spiking neurons, with implications for both decentralized AI based on local neuromorphic computing and investigations of 221 biologically plausible neural processing. 222

## Methods

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Forward Propagation through Time. FPTT considers learning as a consensus problem between the network updates at 224 different time-steps, where the network update at each single time-step needs to move toward the same converged optimal 225 weights. To achieve this, FPTT updates the network parameters W by optimising the instantaneous risk function  $\ell_r^{dyn}$ , 226

which includes the ordinary objective  $\mathcal{L}_t$  and also a dynamic regularisation penalty  $\mathcal{R}_t$  based on previously observed losses  $\ell_t^{dyn} = \mathcal{L}_t + \mathcal{R}_t$  (see Appendix A in the SI for details). In FPTT, the empirical objective  $\mathcal{L}(y_t, \hat{y}_t)$  is the same as for BPTT, representing a function of the gap between target values  $y_t$  and real time predictions  $\hat{y}_t$ .

As in<sup>8</sup>, the FPTT-specific dynamic regularization is controlled by a form of running average of all the weight-updates calculated so far  $\bar{W}$ , where the update schema of this regularizer  $\mathcal{R}_t$  is as follows:

$$\mathscr{R}(W_t) = \frac{\alpha}{2} \parallel W_t - \bar{W}_t - \frac{1}{2\alpha} \nabla l_{t-1}(W_t) \parallel^2$$
(1a)

$$W_{t+1} = W_t - \eta \nabla_W l(W_t) \tag{1b}$$

$$\bar{W}_{t+1} = \frac{1}{2}(\bar{W}_t + W_{t+1}) - \frac{1}{2\alpha}\nabla l_t(W_{t+1}).$$
(1c)

Here, the state vector  $\bar{W}_t$  summarises past losses: the update is first a normal update of parameters  $W_t$  based on gradient optimization with fixed  $\bar{W}_t$ , after which  $\bar{W}_t$  is optimized with fixed  $W_t$ . This approach allows the RNN parameters to converge to a stationary solution of the traditional RNN objective<sup>8</sup>. Note that in Eq. 1c, the loss  $\nabla l_t(W_{t+1})$  is estimated as in<sup>8</sup>, avoiding

propagating the gradient through Eq. 1b, where the  $\nabla_W l(W_t)$  calculates the direct spatial gradient.

The plain FPTT learning process requires the acquisition of an instantaneous loss  $l_t$  at each time step. This is natural for sequence-to-sequence modeling tasks and streaming tasks where a loss is available for each time step; for temporal sequence classification tasks however, the target value is only determined after processing the entire time series. To apply FPTT to learning such tasks in an online manner, a divergence term was introduced<sup>8</sup> in the form of an auxiliary loss to reduce the distance between the prediction distribution  $\hat{P}$  and target label distribution Q:

$$l_{t} = \beta l_{t}^{CE}(\hat{y}_{y}, y) + (1 - \beta) l_{t}^{div},$$
(2)

where  $\beta \in [0,1]$ ;  $l_t^{CE}$  is the classical cross-entropy for a classification loss and  $l_t^{div} = -\sum_{\bar{y}} Q(\bar{y}) \log \hat{P}(\bar{y})$  is the divergence term. We use the auxiliary loss in all experiments as in<sup>8</sup> with  $\beta = \frac{t}{T}$ , where *T* is the sequence length. Note that variants of FPTT, FPTT-K<sup>8</sup>, update *K* times during the sequence rather than every time-step, plain FPTT corresponds to FPTT-T. Absent the dynamic regularization through  $\bar{W}$  FPTT-T amounts to spatial error-backpropagation (BP) without backpropagation-throughtime – the absence of these regularization terms drastically reduces accuracy, both for plain BP as well as for truncated BPTT (Fig. S6).

Training networks of spiking neurons. To apply FPTT to SNNs, we define the spiking neuron model and specify how BPTT
 and FPTT are applied to such networks. All networks were trained using batches as in<sup>8</sup> to exploit GPU parallelism; reduction
 to batch-size=1 yielded similar results (e.g. Fig. S3).

An SNN is comprised of spiking neurons which operate with non-linear internal dynamics. These non-linear dynamics consist of three main components:

(1) Potential Updating: the neurons' membrane potential  $u_t$  updates following the equation:

$$u_t = f(u_{t-1}, x_t, s_{t-1} \| W, \tau)$$
(3)

where  $\tau$  is the set of internal time constants and *W* is the set of associated parameters, like synaptic weights. The membrane potential evolves based on previous neuronal states (e.g. potential  $u_{t-1}$  and spike-state  $s_{t-1} = \{0, 1\}$ ) and current inputs  $x_t$ . Training the time constants  $\tau$  in the spiking neurons is known to optimize performance by matching the neuron's temporal dynamics of the task<sup>16,29</sup>.

(2) Spike generation: A neuron will trigger a spike  $s_t = 1$  when its membrane potential  $u_t$  crosses a threshold  $\theta$  from below, described as a discontinuous function:

$$s_t = f_s(u_t, \theta) = \begin{cases} 1, & \text{if } u_t \ge \theta \\ 0, & \text{otherwise} \end{cases}$$
(4)

(3) Potential resting: When a neuron emits a spike ( $s_t = 1$ ), its membrane potential will reset to resting potential  $u_r$ :

$$u_t = (1 - s_t)u_t + u_r s_t,\tag{5}$$

where in all experiments, we set  $u_r = 0$ .

**BPTT for SNNs** BPTT for SNNs amounts to the following: given a training example  $\{x, y\}$  of T time steps, the SNN generates 250 a prediction  $\hat{y}_t$  at each time step. At time t, the SNN parameters are optimized by gradient descent through BPTT to minimize 251 the instantaneous objective  $\ell_t = \mathscr{L}(y_t, \hat{y}_t)$ . The gradient expression is the sum of the products of the partial gradients, defined 252 by the chain rule as 253

$$\frac{\partial \ell_t}{\partial W} = \frac{\partial l_t}{\partial \hat{y}_t} \frac{\partial \hat{y}_t}{\partial s_t} \frac{\partial s_t}{\partial u_t} \sum_{j=1}^t \left( \prod_{m=j}^t \frac{\partial u_m}{\partial u_{m-1}} \right) \frac{\partial s_{m-1}}{\partial W},\tag{6}$$

where the partial derivative of spiking  $\frac{\partial s_t}{\partial u_t}$  is calculated by a surrogate gradient associate with membrane potential  $u_t$ . Here, we 254 use the Multi-Gaussian surrogate gradient function  $\hat{f}'_{s}(u_{t},\theta)^{1}$  to approximate this partial term. 255

The computational graph of BPTT is illustrated in Fig1a and shows that the partial derivative term depends on two pathways, 256  $\frac{\partial u_m}{\partial u_{m-1}} = \frac{\partial u_m}{\partial u_{m-1}} + \frac{\partial u_m}{\partial s_{m-1}} \frac{\partial s_{m-1}}{\partial u_{m-1}}$ . The product of these partial terms may explode or vanish in RNNs, and this phenomenon becomes even more pronounced in SNNs as the discontinuous spiking process is approximated by the continuous surrogate gradient and 257 258 the incurred gradient error accumulates and amplifies. 259

FPTT for SNNs. FPTT can be used for training SNNs by minimizing the instantaneous loss with the dynamic regularizer  $\ell_t^{dyn} = \mathscr{L}(\mathbf{y}_t, \hat{\mathbf{y}}_t) + \mathscr{R}(\bar{W}_t)$ . The update function Equation (6) then becomes:

$$\frac{\partial \ell_t^{dyn}}{\partial W} = \frac{\partial l_t}{\partial \hat{y}_t} \frac{\partial \hat{y}_t}{\partial s_t} \frac{\partial s_t}{\partial u_t} \frac{\partial u_t}{\partial W}.$$
(7)

Compared to Equation (6), Equation (7) has no dependence on a chain of past states, and can thus be computed in an online 260 manner. 261

Liquid Time-Constant Spiking Neurons The LTC-SN is modeled as a standard adaptive spiking neuron<sup>1,11</sup> where the time 262 constants  $\tau$  (here, membrane time constant  $\tau_m$  and adaptation time constant  $\tau_{adp}$ ) are a dynamic and learned function of internal 263 dynamic state variables like membrane potential u and deviation<sup>11</sup> b. In the network, time-constants are either calculated as 264 a function  $\alpha = \exp(-dt/\tau_m) = \sigma(Dense[x_t, u_{t-1}])$ , for non-convolutional networks, or using a 2D convolution for spiking 265 convolutional networks,  $\alpha = \exp(-dt/\tau_m) = \sigma(Conv([x_t, u_{t-1}]))$ , where we use a sigmoid function  $\sigma(\cdot)$  to scale the inverse of 266 the time constant to a range of 0 to 1, ensuring smooth changes when learning. This results in a Liquid Time-Constant Spiking 267 Neuron *i* defined as: 268

$$\tau_{adp} update: \rho^{i} = \exp(-dt/\tau^{i}_{adp}) = \sigma(Dense_{adp}[x_{t}, b^{i}_{t-1}])$$
(8a)

$$\tau_{adp} update : \rho = \exp(-dt/\tau_{adp}^{i}) = O(Dense_{adp}[x_{l}, \sigma_{l-1}])$$

$$\tau_{m} update : \alpha^{i} = \exp(-dt/\tau_{m}^{i}) = \sigma(Dense_{m}[x_{l}, u_{l-1}^{i}])$$
(8b)

$$\theta_t update \quad : b_t^i = \rho^i b_{t-1}^i + (1 - \rho^i) s_{t-1}^i; \ \theta_t^i = 0.1 + 1.8 b_t^i \tag{8c}$$

$$u_t u_t u_t u_t = -u_{t-1}^i + x_t; u_t^i = \alpha^i u_{t-1}^i + (1 - \alpha^i) du^i$$
(8d)

spikes<sub>t</sub> : 
$$s_t^i = f_s(u_t^i, \theta^i)$$
 (8e)

resetting : 
$$u_t^i = u_t^i (1 - s_t^i) + u_{rest} s_t^i$$
, (8f)

where the neuron uses an adaptive threshold  $\theta_t$  as in the Adaptive Spiking Neurons<sup>11</sup>, resting potential  $u_{rest} = 0$ , and time-269 constants  $\tau_m^i$  and  $\tau_{adp}^i$  are computed as liquid time-constants of neuron *i*. 270

Datasets We focus on a number of datasets suitable for temporal classification, where the goal is to obtain high accuracy 271 read-out on test-samples at the final time-step T of the sequence. For the Add-task, the trained networks consist of 128 272 recurrently connected neurons of respective types LTC-SN (LTC-SNN), LSTM, or Adaptive Spiking Neuron<sup>11</sup> (ASRNN), and 273 a dense output layer with only 1 neuron. 274

For the DVS-GESTURE dataset, each frame is a 128-by-128 size image with 2 channels. Each sample in the DVS-275 GESTURE dataset was split into fixed-duration blocks as in<sup>16</sup>, where each block is averaged to a single frame. This conversion 276 results in sequences of up to 1000 frames depending on block length. As input for the shallow SRNN as used in the increasingly 277 long sequence setting, we first down-sample the frame of a 128-by-128 image into a 32-by-32 image by averaging each 4-by-4 278 pixel block. The 2D image at each channel is then flattened into a 1D vector of length 1024. For each channel of the image, 279 the network consists of a spike-dense input layer consisting of 512 neurons as an encoder, where the information of each 280 channel is then fused into a 1D binary vector through concatenation. This 1D vector is then fed to a recurrently connected 281

layer with 512 hidden neurons. Finally, a leaky integrator is applied to generate predictions, resulting in a network architecture
 [1024,1024]-[512D,512D]-512R-111<sup>1</sup>. All networks were trained for 30 epochs using the Adam optimizer with initial learning
 rate 3e-3, using the same initialization schemes and learning-rate scheme for all networks to compare the effect of neural units.

<sup>285</sup> Hyperparameters for ASRNNs were taken from<sup>1</sup>.

To achieve high performance with SCNNs, we applied FPTT with LTC-SNs to high-performance architectures from the 286 literature, where we used hyperparameter settings for the surrogate gradient from<sup>29</sup> and from<sup>8</sup> for FPTT training. Specifically, 287 in (P)S-MNIST, we used a shallow network with one recurrent layer comprised of 512 hidden neurons and an output layer 288 consisting of 10 (number of classes) leaky integrator neurons. The FPTT-trained LTC-SNNs are optimized using Adam<sup>48</sup> with a 289 batch size of 128 using 200 training epochs. We set the initial learning rate to 3e-3 and decay by half after 30, 80, and 120 epochs. 290 For the S-MNIST and PS-MNIST tasks, we also find that the leak time-constants of the output units after training averaged 291  $87 \text{ms} \pm 13 \text{ms}$  (S-MNIST) and  $65 \text{ms} \pm 12 \text{ms}$  (PS-MNIST), substantially shorter than the sequence length and demonstrating that 292 the recurrent network maintains working memory. For R-MNIST, we follow the architecture from<sup>12</sup>: an SNN with two hidden 293 layers of 256 neurons, each followed by 10 output neurons. The SNN is given 20 presentations of the image, after which the 294 classification is determined. For the static MNIST and Fashion MNIST datasets, we apply the architecture from<sup>16</sup>: an SCNN 295 with 3 convolutional layers, 1 Dense layer and 1 leaky Integrator output layer (ConvK3C32S1P1-MPK2S2-ConvK3C128S1P1-296 MPK2S2-ConvK3C256S1P1-MPK2S2-512D-10I. To describe the network structure, we follow standard conventions as follows: 297 ConvK7C64S1P1 represents the convolutional layer with *out put channels* = 64, *kernel size* = 7, *stride* = 1 and *padding* = 3. 298 MPK2S2 is the max-pooling layer with kernel size = 2 and stride = 2. 512D and 512R represents the fully connected 299 spiking layer and recurrent spiking layer respectively with *out put features* = 512. 10I indicates the leaky integrator with the 300 out put feature = 10.). The resulting LTC-SCNN network was then optimized using FPTT and Adamax with a batch size of 64 301 and an initial learning rate of 1e-3. For the DVS-GESTURE and DVS-CIFAR10 datasets, we also follow the high-performance 302 architecture from<sup>16</sup> and use 20 sequential frames, where the network makes a prediction only after reading the entire sequence. 303 The LTC-SCNN thus has a structure ConvK7C64S1P3-MPK2S2-ConvK7C128S1P3-MPK2S2-ConvK3C128S1P1-MPK2S2-304 ConvK3C256S1P1-MPK2S2-ConvK3C256S1P1-MPK2S2-ConvK3C512S1P1-MPK2S2-512D-11I. These networks were 305 optimized through Adamax<sup>48</sup> with a batch size of 16 and initial learning rate of 1e-3. 306

<sup>307</sup> For all networks, to measure GPU memory consumption, the GPU management interface "nvidia-smi" was used.

**Spiking Tiny YOLOv4 – SPYv4.** The SPYv4 network follows the Tiny YOLO-v4 architecture<sup>22, 38</sup>. The backbone of the network consists of three CSP-Blocks in the cross-stage partial network. In contrast to regular additive residual connections, the CSP-Block is spike-based rather than event-based as residual connections are constructed by concatenation rather than using an adding operation, ensuring that only binary spikes are used for information transfer between layers. Raw pixel values are fed directly into the network as input currents. As the object recognition task necessitates a more precise output vector to draw the bounding box, we use a single standard conventional convolutional layer instead of a spiking convolutional layer.

We trained and evaluated our model on Pascal VOC dataset<sup>17</sup> as was done in<sup>18</sup>: the network was trained using a combination of VOC2007 and VOC2012 (16551 images), and evaluated on the validation dataset of VOC2007 (4952 images).

For target detection, we use a threshold on the Intersection Over Union (IOU), i.e., the ratio of the intersection of the prediction box and the ground truth box of the target, to indicate whether the detection is correct. A target is considered to be correctly detected when the IOU exceeds the threshold. The average precision (AP) is then computed as the average of all the precision for all possible recall values, and the mAP is the average of the AP of multiple classification tasks. Training maximizes the mAP@ $\vartheta$ : the mean average precision of the calculated bounding box exceeding an overlap threshold  $\vartheta$  over the actual boundaries of the object in the dataset, where we use  $\vartheta = 0.5$ .

We applied both mosaic data augmentation<sup>38</sup> and label smoothing during training to obtain better performance. In the mosaic enhancement, the network uses a mosaic of 4 images during training instead of a single image; this is applied only during the first 200 training cycles. The network was optimized by Adagrad with a batch size of 32 and an initial learning rate of 1e-3.

### **Code Availability**

The code used in the study is publicly available from the GitHub repository https://github.com/byin-cwi/sFPTT/ tree/v1.0.0<sup>49</sup>.

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### **419** Supplementary Information

420

Algorithm 1: Training SNN with BPTT	Algorithm 2: Training SNN with FPTT
// Dataset $B = \{x_t, y_t\}_{t=0}^T$ # Epochs E	// Dataset $B = \{x_t, y_t\}_{t=0}^T$ , # Epochs E
// Set Optimizer and learning rate $\eta$	// Set Optimizer and learning rate $\eta$
// Initialize Weight $W = \{W^h, W^y\}$	// Initialize weight $W = \{W^h, W^y\}$ , and
for $i = 1$ to $E //$ Training Loop	$\bar{W} = W$
do	for $i = 1$ to $E / /$ Training Loop
Initialize neuron states $u_t, s_t$ ; Randomly Shuffle B	do
for $t = 0$ to $T //$ For each sample	Initialize neuron states $u_t$ , $s_t$ ; Randomly Shuffle B
do	for $t = 0$ to $T / /$ For each sample
$  s_{h,t}, u_{h,t} = \hat{f}_s(x_t, [u_{h,t-1}, s_{h,t-1}]    W^h)$	do
$\hat{y}_t = \hat{f}_s(s_{h,t}, [u_{o,t}, s_{o,t}]    W^y)$	$  s_{h,t}, u_{h,t} = \hat{f}_s(x_t, [u_{h,t-1}, s_{h,t-1}]    W_t^h)$
Loss:	$\hat{y}_t = \hat{f}_s(s_{h,t}, [u_{o,t}, s_{o,t}]    W_t^y)$
$\ell(W) = \sum_{t=1}^{T} \ell(\mathbf{y}_t, \hat{\mathbf{y}}_t)$	Loss: $\ell_t(W_t)$ : $\ell_t(W_t) = \ell(y_t, \hat{y}_t)$
Update: $-i = 1 + 0 + 1 + 1 + 1 + 1 + 1 + 1 + 1 + 1 +$	Dynamic Loss: $\ell^{dyn}(W_t) =$
$W = W - \eta \nabla_W \ell(W) _W$	$\ell_t(W_t) + \frac{\alpha}{2} \ W_t - \bar{W}_t - \frac{1}{2\alpha} \nabla \ell_{t-1}(W_t)\ ^2$
	Update W:
	$W_{t+1} = W_t - \eta \nabla_W \ell^{dyn}(W_t)$
	Update $\overline{W}$ :
	$\bar{W}_{t+1} = \frac{1}{2}(\bar{W}_t + W_{t+1}) - \frac{1}{2}\nabla \ell_t(W_{t+1})$
Algorithms CA1 DDTT (1-ft) and EDTT algorithms Adapted for	$\nabla I(\mathbf{W})$ ) is estimated as in 8 second s

Algorithms SA1. BPTT (left) and FPTT algorithm. Adapted from<sup>8</sup> where  $\nabla l_t(W_{t+1})$ ) is estimated as in<sup>8</sup>, avoiding propagating the gradient through Equation 1b. The algorithms are described for batch-size 1, while in our experiments we relax this to larger batch-sizes as in<sup>8</sup>.

Frames	BPTT						FPTT		
Traines	LSTM+Aux	LSTM	LTC-SRNN+Aux	LTC-SRNN	ASRNN+Aux	ASRNN <sup>+</sup> +Aux	$LTC^{-}$	LTC-SRNN	ASRNN
20	86.69±0.43	82.29±2.46	83.42±1.35	84.37±2.27	86.82±0.31	80.78±1.99	87.83±1.21	89.31±0.59	87.51±0.85
40	88.77±1.71	84.95±0.71	85.96±1.16	84.37±1.24	87.29±0.87	82.99±0.70	88.53±0.57	90.39±0.71	87.61±0.43
60	87.61±0.86	85.15±0.75	85.62±1.18	83.91±0.71	87.02±1.19	81.77±0.34	$88.08 \pm 1.40$	<b>90.74</b> ±0.16	87.62±1.15
80	87.97±0.14	$84.83 \pm 1.42$	85.30±0.71	80.44±3.6	86.34±0.87	$76.84{\pm}0.85$	88.66±1.14	<b>91.31</b> ±0.98	87.60±1.06
100	88.89±0.49	83.79±0.71	83.21±0.43	78.70±0.91	86.22±0.71	74.61±1.25	89.93±1.13	<b>91.89</b> ±0.16	$87.40 {\pm} 0.28$
200	85.76±0.49	$81.87 {\pm} 2.58$	51.39±6.0	(43.98±2.35)*	79.62±1.89	65.89±1.69	89.24±1.56	<b>92.13</b> ±0.87	88.31±1.33
500	82.52±1.82	78.81±1.5	38.89±3.22	(36.46±1.5)*	49.03±1.52	52.43±1.32	86.69±0.43	90.64±1.56	85.76±1.30
1000	+	+	+	+	+	+	90.05±1.56	91.28±1.05	84.24±1.23
Param	4.2M	4.2M	4.7M	4.7M	1.6M	4.7M	3.2M	4.7M	1.6M

**Table S1.** Performance comparison between BPTT and FPTT on the DVS gesture dataset. Each number in the table is the average of three runs. All networks have an equal number of neural units unless indicated otherwise. (\*) denotes that training diverged; reported accuracy is the best accuracy before divergence. (+) denotes out-of-GPU-memory when training. The *ASRNN*<sup>+</sup> is an ASRNN with the same number of parameters as the LTC-SRNN. *LTC*<sup>-</sup> denotes an LTC-SRNN network where only the membrane time constant is dynamic, and the adaptation time constant is a learned fixed parameter. We remark that with a parameter-matched single recurrent layer ASRNN with 1400 neurons trained w/ BPTT, we achieved accuracy of  $81.134\% \pm 0.71$  on 100 frames DVS-gestures dataset, while the same network trained with eProp achieved 75.45% ± 1.15. Results for Truncated BPTT including plain spatial error-backpropagation are illustrated in Fig S6.

Loss	$ au_m$	$\tau_m(x)$	$\tau_m(x,u)$
$ au_{adp}$	0.17	0.0027	0.0025
$\tau_{adp}(x)$	0.17	0.0024	0.003
$\tau_{adp}(x,b)$	0.16	0.0021	0.0019

**Table S2.** Ablation study of LTC-SN performance on Add Task where either  $\tau_m$ ,  $\tau_{adp}$ , or both, are trained ( $\tau_m$ ,  $\tau_{adp}$ ), dynamic from external input x ( $\tau_m(x)$ ,  $\tau_{adp}(x)$ ) or both external and internal input ( $\tau_m(x,u)$ ,  $\tau_{adp}(x,b)$ ).



**Figure S1.** Learning curve of ASRNN trained via FPTT and BPTT on **a** 100 frames, and **b** 500 frames. Plotted is average and std over 3 run; **c**, Bar chart of mean firing rate (fr) comparison between BPTT and FPTT on the DVS gesture dataset. Each number in the table is the average of three runs.



**Figure S2.** Spiking YOLOv4 (SPYv4) neural network architecture. The dash line read the "Feature" tensor which calculated before the last concatenation in CSP BLOCK



**Figure S3.** Example learning curves on the Add-task of LTC-SRNN trained with BPTT, FPTT, FPTT and batch-size 1 ("LTC-SRNN\_bz1", red curve).



**Figure S4.** Ablation study of recurrent connections on the DVS Gesture dataset. Plot for test accuracy of LTC-SRNN on DVS Gesture dataset. The label "W/O Rec" denotes the LTC-SNN corresponding to the baseline recurrently connected LTC-SRNN but with the recurrent connections in the recurrent layer removed; "W/O Rec++" denotes an LTC-SNN with the same number of parameters as the LTC-SRNN (increasing the number of neural units and without recurrent connections). Test accuracy bars show mean  $\pm$  std over 3 runs and individual data points are inserted.

**Table S3.** Memory efficiency. (\*): the reported number is obtained using a halved batch size compared to the other entries to fit into GPU memory,

	S-MNIST	R-MNIST	MNIST	DVS-Gesture
BPTT	11.1GB	1.5GB	9.67GB	15.72GB(*)
FPTT	1.9GB	1.4GB	2.23GB	3.75GB

Table S4.	. Total	training	time	per	epoch
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	S-MNIST	R-MNIST	MNIST	DVS gesture
BPTT	2518s	192s	362s	108s
FPTT	1233s	204s	384s	112s

Frames	Izhikevich	Hodgkin Huxley
100	84.03%	87.50%

**Table S5.** Performance of SNNs with Izhikevich and Hodgkin Huxley models on DVS-Gesture dataset. The network structure is same as the network in Table S1, where for the Izhikevich the a and b parameters are trainable and we kept c and d fixed, for the Hodgkin-Huxley model all neuron parameters were kept fixed; the network structure and the training-related hyperparameters were not further fine-tuned.



**Figure S5.** Dynamic neural responses of LTC-SNs. (a) an MNIST sample is sequentially fed into the LTC-SNN, pixel-by-pixel along the row-direction. (b) for illustration, we calculated the histograms of the resulting average  $dt/\tau$  values after training for this single sample (c) average responses of neurons in terms of firing rate binned by  $dt/\tau_m$ ,  $exp(-dt/\tau_{adp})$  values (d) Example of dynamics of inverse time constant  $dt/\tau_m$  for four randomly selected neurons during presentation of the sequence.



**Figure S6.** Ablation study of Truncated-BPTT-k (TBPTT - k) on DVS Gesture dataset of 200 frames where *k* is number of updates in the sequence resulting in truncation period *subL*. TBPTT-1, with truncation length 1, is functionally equivalent to BP. (a) Appropriate segmentation in time by truncation assists the network to achieve better results in long sequence classification problems. Compared to BPTT, T-BPTT has fixed consumption proportional to the truncation length. (b) For TBPTT-1, performance on the same network architecture is essentially constant as a function of sequence length.

### **424** Appendix A: FPTT theory

 $_{425}$  For conciseness, we briefly summarize the theory underlying FPTT as developed by Kag et al.<sup>8</sup>.

**Back-propagation-through-time** Back-propagation-through-time (BPTT) uses backpropagation to calculate the gradient of the accumulated loss along the spatial-temporal dimension with respect to the parameters of the recurrent networks. Let us define a recurrent network described by differential equation  $(\hat{y}^t, h_t) = NN(x_t, h_{t-1})$  where  $x_t$  is the input,  $\hat{y}^t$  is the prediction and  $h_t$  is the hidden states. The gradient of time *t* is then computed by considering the effect of the state  $x_t$  on all future losses  $l_{30}^t$   $l^t, l^{t+1}, \dots, l^T$ :

$$\frac{\partial L}{\partial w} = \sum_{t=1}^{T} \frac{\partial l^{t}}{\partial w} = \sum_{t=1}^{T} \sum_{i=1}^{t} \frac{\partial l^{t}}{\partial h_{i}} \frac{\partial h_{i}}{\partial w} = \sum_{t=1}^{T} (\sum_{i=t}^{T} \frac{\partial l^{i}}{\partial h_{t}}) \frac{\partial h_{t}}{\partial w} = \sum_{t=1}^{T} (\sum_{i=t}^{T} \frac{\partial l^{i}}{\partial h_{t}}) \frac{\partial h_{t}}{\partial w} = \sum_{t=1}^{T} \{\sum_{i=t}^{T} (\prod_{j=i}^{T-1} \frac{\partial l^{j+1}}{\partial l^{j}}) \frac{\partial l^{i}}{\partial h_{t}}\} \frac{\partial h_{t}}{\partial w},$$
(A.1)

and a weight is then updated as:  $w_{new} \leftarrow w_{old} - \frac{\partial L}{\partial w}$ . At the end of training, the loss *L* will be minimized via optimal solution  $w^*$ , where  $\frac{\partial}{\partial w}L(w^*) \cong 0$ .

For online update based on BPTT, we will have  $w_{t+1} \leftarrow w_t - \sum_{i=1}^t \frac{\partial}{\partial w} l^i(\hat{y}^i, y^i, w_t)$  where  $l^i(\hat{y}^i, y^i, w_t)$  is the cost of time step *i* with parameter  $w_t$ ,  $\hat{y}^i$  and  $y^i$  are the prediction and target label of the time step *i*. When the algorithm converges to an optimal solution  $w^*$  at time step  $\varphi$ , we will have an optimal solution where:

$$w^* - w_t = -\nabla_w(l^t) = \frac{\partial}{\partial w} l^{\varphi}(\hat{y}^{\varphi}, y^{\varphi}, w^*) - \frac{\partial}{\partial w} l^t(\hat{y}^t, y^t, w_t)$$
(A.2)

and, for one step optimization:

$$w_{t+1} - w_t = \nabla_w l^{t+1} - \nabla_w l^t.$$
(A.3)

This demonstrates that for any timestep, the change of weight update is proportional to the change of the gradient; this observation (Equation (A.3)) is the foundation of Forward Propagation Through Time.

**Forward Propagation Through Time** FPTT aims to derive an online weight update mechanism with guaranteed convergence to optimal solution  $w^*$ . To have a smooth solution, FPTT learns from the historical information of weight changes by introducing a running mean  $\bar{w}_t$  to summarize the historical information of weight evolution:

$$w_{t+1} - w_t = \nabla_w(l^{t+1}) - \nabla_w(l^t)$$
(A.4)

$$\Rightarrow \bar{w}_t - w_t \sim \nabla_w(l^{t+1}) - \nabla_w(l^t) \tag{A.5}$$

$$\Rightarrow \nabla_{w}(l^{t+1}) - \nabla_{w}(l^{t}) = \alpha[(\bar{w}_{t} - w_{t}) - (w_{t+1} - w_{t})] = \alpha(\bar{w}_{t} - w_{t+1})$$
(A.6)

From this, the convergence-guaranteed loss function for online update is derived, based on Eq. (A.6).

$$\nabla_{w}(l^{t+1}) - \nabla_{w}(l^{t}) = \alpha(\bar{w}_{t} - w_{t+1}) \qquad \Leftrightarrow \qquad \nabla_{w}(l^{t+1}) - \nabla_{w}(l^{t}) - \alpha(\bar{w}_{t} - w_{t+1}) = 0 \tag{A.7}$$

we define the constraint into the function  $f(w_{t+1}) = \nabla_w(l^{t+1}) - \nabla_w(l^t) - \alpha(\bar{w}_t - w_{t+1})$ . We now consider a convex function F(w) which approached its minimum when  $f(w_{t+1}) = 0$ ; we then have

$$F(w) = \int_{w} f(w)dw - \text{searching } w_{t+1} \text{ over parameter space}$$
(A.8)  
$$= l^{t}(w) + \frac{\alpha}{2} \|w - \bar{w}_{t} - \frac{1}{2\alpha} \nabla_{w}(l(w_{t}))\|^{2}$$
(A.9)

In this form, Eq A.7 is the first order condition for F(w). So, the weight optimization is to minimize the new objective function

$$w_{t+1} = \underset{w}{\arg\min} l^{t}(w) + \frac{\alpha}{2} \|w - \bar{w}_{t} - \frac{1}{2\alpha} \nabla_{w}(l(w_{t}))\|^{2}$$
(A.10)