SPIKE-SSM: A SPARSE, PRECISE, AND EFFICIENT SPIKING STATE SPACE MODEL FOR LONG SEQUENCES LEARNING

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ABSTRACT

Spiking neural networks (SNNs) provide a low-power, energy-efficient solution by utilizing the spike-based and sparse nature of biological systems. Since the advent of Transformers, SNNs have struggled to compete with artificial networks on long sequential tasks, until the recent emergence of state space models (SSMs), which offer superior computational efficiency and modeling capability. However, applying the highly capable SSMs to SNNs for long sequences learning poses three major challenges: **1** The membrane potential is determined by the past spiking history of the neuron, leading to reduced efficiency for sequence modeling in parallel computing scenarios. Complex dynamics of biological spiking neurons are crucial for functionality but challenging to simulate and exploit effectively in large networks. 3 It is arduous to maintain high sparsity while achieving high accuracy for spiking neurons without resorting to dense computing, as utilized in artificial neuron-based SSMs. To address these challenges, we propose a sparse, precise and efficient spiking SSM framework, termed SPikE-SSM. For $\mathbf{0}$, we propose a boundary compression strategy (PMBC) to accelerate the inference of the spiking neuron model, enabling parallel processing for long sequence learning. For **2**, we propose a novel and concise neuron model incorporating reset-refractory mechanism to leverage the inherent temporal dimension for dynamic computing with biological interpretability. For **③**, we hierarchically integrate the proposed neuron model to the original SSM block, and enhance the dynamics of SPikE-SSM by incorporating trainable thresholds and refractory magnitudes to balance accuracy and sparsity. Extensive experiments illustrate the effectiveness and robustness of SPikE-SSM on the long range arena benchmarks and large language dataset WikiText-103, showing the potential of dynamic spiking neurons in efficient long sequence learning. The code will be publicly available.

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1 INTRODUCTION

Spiking neural networks (SNNs) recently emerged as a competitive paradigm to improve AI energy efficiency. SNNs transmit information as binary spikes between synapses to perform sparse and event-driven computation. Despite being increasingly more competitive with artificial neural networks (ANNs) in vision tasks, SNNs still struggle with long-sequence modeling – a critical task for a wide range of temporal or sequential data-driven machine learning applications, such as text comprehending (Zhou et al., 2023), electroencephalograms spanning (Tang et al., 2023), etc.

Transformer (Vaswani et al., 2017) and its variants (Kitaev et al., 2020; Zaheer et al., 2020; Katharopoulos et al., 2020) have been developed for sequential tasks. However, their architectures are not suitable for SNN-based long sequence learning as SNN requires a time window-based simulation to enhance spike-based representation, resulting in slow inference compared to their ANN counterparts (Zhou et al., 2022; Yao et al., 2024). Moreover, the self-attention mechanisms (Vaswani et al., 2017) in Transformers are computationally intensive, contrasting with the energy-efficient properties of event-based representations and the sparse computation inherent to SNNs. As a competitive alternative to Transformer, state space models (SSMs) have garnered significant attention due to their long sequence modeling capabilities, such as S4 (Gu et al., 2021), DSS (Gupta et al., 2022), S5 (Smith et al., 2022) and Mamba (Gu & Dao, 2023). Notably, SSMs can achieve fast in-



Figure 1: Main ideas of SPikE-SSM for long-sequence modeling. (*Left*) Overview: A parallel max-min boundary compression (PMBC) strategy is proposed to address ① (§ 3.2); a new refractory neuron model with trainable dynamics is developed to address ② (§ 3.3). We integrate the proposed refractory neuron with *soft reset* within SSMs to address ③ (§ 3.4). (*Right*) An example showing that the relevant information for the task at hand is often sparse in long-sequence inputs.

ference and parallel training by incorporating dynamic hidden states for handling long-range dependencies (LRDs), inspired by the low-complexity inference mechanism of recurrent neural networks (RNNs) (Sherstinsky, 2020; Schuster & Paliwal, 1997). Meanwhile, the sequential computing nature of SSMs is also more compatible with SNNs as the dynamics of spiking neurons can be inherently exploited in the temporal dimension. Furthermore, for tasks with long-sequence inputs, it is often the case that the *relevant information* to the problem at hand is *inherently sparse* (see Figure 1 *Right* for an example), aligning well with the sparse representation of SNNs.

073 Therefore, spiking SSMs naturally emerge as a promising paradigm for efficient long-sequence 074 modeling. Recent works have highlighted notable advancements in capturing LRDs using spiking 075 SSMs (Stan & Rhodes, 2023; Bal & Sengupta, 2024; Shen et al., 2024). However, these exist-076 ing methods are still inadequate in addressing the following challenges when applying spike-based 077 computation to SSMs: **1** The membrane potential of a neuron in SNNs depends on its past spiking history, making parallel processing infeasible and, in turn, hindering the efficiency of sequence modeling. ⁽²⁾ Biological neuron models exhibit complex dynamics that are essential for function-079 ality (Urbanczik & Senn, 2014; Mikulasch et al., 2021; Capone et al., 2023) but challenging to simulate efficiently in large networks – an issue often overlooked by existing methods. Sparse 081 representation is key for efficient computation in SNNs (Olshausen & Field, 2004; Jiao et al., 2022; 082 Raposo et al., 2024); however, balancing the trade-off between sparsity (i.e., *spiking rate*) and ac-083 curacy remains challenging for spiking SSMs, as SSMs were originally designed on top of artificial 084 neurons with dense computations. 085

In this work, we propose a novel spiking SSM model, termed **SP**ikE-SSM, to exploit the intricate dynamics of Leaky Integrate-and-Fire (LIF) neuron (Gerstner et al., 2014) in SSMs for sparse, 087 **p**arallel, and **e**fficient long-sequence modeling. First, to address $\mathbf{0}$, we propose a parallel max-min 088 boundary compression strategy (PMBC) to accelerate the inference of the LIF neuron, enabling par-089 allel processing for long sequence modeling. Second, to address **2**, we propose a refined LIF neuron 090 model incorporating a reset-refractory dynamics to fully utilize the inherent temporal dimension for 091 dynamic computing with biological interpretability; in the meantime, the hyperparameters of the 092 proposed neuron model are trained efficiently and explicitly based on PMBC, enabling a systematic study of their functional impacts on the network. Third, to address $\boldsymbol{\Theta}$, we integrate the refractory neuron into an SSM block to adjust the membrane potential with dynamic reset, achieving both high 094 accuracy and low spiking rate (i.e., high efficiency). An overview comparison of our SPikE-SSM with existing spiking SSMs is presented in Table 1. The main contributions are as follows: 096

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- In this paper, we propose SPikE-SSM to effectively model the long sequence with SNNs. In contrast to existing spiking SSMs, our method can realize comprehensive parallel acceleration with trainable temporal dynamics, facilitating sparse, precise, efficient training and inference for long-range dependencies learning.
- To tackle the dilemma of event-driven neuronal dynamics with parallel processing for long sequence modeling, we propose a max-min boundary compression (PMBC) strategy to facilitate an efficient inference of SPikE-SSM. We empirically demonstrate that PMBC is versatile and effective for accelerating neuronal dynamics for parallel computing of SNNs.
- A new LIF neuron model with a refractory mechanism is proposed to fully utilize the inherent temporal dimension for biologically interpretable dynamic computation, achieving both high accuracy and sparsity with the trainable dynamics.

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Table 1: Comparison of our model with existing spiking SSMs. Previous methods mostly apply 109 binary activation to SSMs without considering the intricate neuronal dynamics. [†] SpikingSSM ap-110 proximates the LIF neuron with hard reset dynamics by using a surrogate model, which is subject 111 to approximation errors. [‡] SpikingSSM has partial trainable dynamics since hard reset is rough and 112 simplified with limited dynamic variables. In contrast, our method can train the neuron hyperparam-113 eters and temporal dynamics efficiently and explicitly in a parallel manner with the proposed PMBC, 114 enabling a functional study of their impact on the network. In SPikE-SSM, a more interpretable soft 115 reset mechanism is employed, incorporating additional trainable dynamic variables and parameters. 116

Method	Reset Mechanism	Trainable Dynamics
Binary-S4D (Stan & Rhodes, 2023)	×	×
S6-based SNN (Bal & Sengupta, 2024)	×	X
SpikingSSM (Shen et al., 2024)	\checkmark^{\dagger}	partial [‡]
SPikE-SSM (ours)	1	· /

 Extensive experiments are conducted on LRA benchmarks and the large-scale WikiText-103 language modeling databases, the results of which validate the effectiveness and efficiency of the proposed SPikE-SSM for long-range dependencies learning.

2 RELATED WORKS

129 130 2.1 Long Sequences Learning Models

131 Long sequence modeling has gained significant attention recently due to its widespread applica-132 tion across different domains such as text comprehending (Zhou et al., 2023), computer vision (Shi 133 et al., 2024; Zhong et al., 2024) and electroencephalograms spanning (Tang et al., 2023). The key 134 challenge in long-sequence modeling lies in efficiently compressing context into a manageable state while capturing information spread across observations separated by thousands of timesteps. To 135 address them, Transformer and Attention (Vaswani et al., 2017; Dao et al., 2022; Dao, 2023) are 136 proposed to retain the entire context during auto-regressive inference, which is effective but requires 137 quadratic-time computational complexity. Although some Transformer variants (Kitaev et al., 2020; 138 Katharopoulos et al., 2020) are proposed to reduce the compute and memory requirements, their 139 performances on long-range reasoning remain considerably suboptimal (Gu et al., 2021). Inspired 140 by RNNs, RWKV (Peng et al., 2023) combines the parallel training of transformers with the effi-141 cient inference of RNNs. Similarly, other recurrent models aim to compress context into a finite 142 state, offering constant-time inference and linear-time training, but their effectiveness is limited by 143 the quality of compression and a fixed representation space (Qin et al., 2023). More recently, SSM-144 based methods (Smith et al., 2022; Fu et al., 2022; Mehta et al., 2022) have emerged as a promising 145 alternative to sequence models such as RNNs and Transformers. For example, HiPPO (Gu et al., 2020) pioneered compressing long inputs into dynamic representations using orthogonal polynomi-146 als, while S4 (Gu et al., 2021) advanced this with low-rank corrections for stable diagonalization 147 and simplified Cauchy kernel operations. Mamba (Gu & Dao, 2023) focuses on selective state 148 representations to optimize efficiency and effectiveness, using a selection mechanism and hardware-149 optimized algorithms to maintain robust contextual information capture. All above methods are 150 based on artificial neurons with analog-valued output, resulting in dense vector-matrix multipli-151 cation (VMM) and huge computational costs. In contrast, the proposed SPikE-SSM utilizes the 152 compatibility between the remarkable LRDs modeling ability of SSMs and the intrinsic dynamics 153 of SNNs, promoting sparse training and fully parallel inference with trainable temporal dynamics.

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2.2 SNNs-based Sequence Modeling and Applications

SNNs (Ghosh-Dastidar & Adeli, 2009) have gained attention as a compelling bio-plausible and computational efficient substitute for traditional artificial neural networks (ANNs) in many vision tasks. However, SNNs have struggled to make significant progress in long-sequence modeling tasks due to the inherent serial computing nature. Therefore, to train SNNs in parallel, PSN (Fang et al., 2024) simplifies spiking neuron by omitting the reset mechanism, leading to reduced sparsity. To handle this issue, a probabilistic reset mechanism is proposed in PSU (Li et al., 2024) to achieve

162 parallel computing with elevated sparsity by decoupling the integration-spiking-resetting process, 163 which comes at the expense of higher computational complexity. With the recent resurgence of 164 SSMs, there has been a renewed focus on applying efficient parallel computing to SNNs. For ex-165 ample, SpikeS4 (Du et al., 2024) integrates LIF neurons with S4 layers for speech learning. Binary 166 S4D builds a binary SSM by applying a spiking activation directly to the sum of hidden states, enabling parallel training but neglecting neuronal dynamics (Stan & Rhodes, 2023). To further en-167 hance sparsity, a stochastic spiking neuron is proposed in S6-based SNN (Bal & Sengupta, 2024), 168 which is trained with stochastic noises in gradients, resulting in accuracy degradation. More recently, SpikingSSMs (Shen et al., 2024) utilizes a surrogate dynamic network (SDN) to approxi-170 mate the dynamics of LIF neurons, which extremely accelerates the training and inference by par-171 allel computing. However, the pre-training requirement of SDN could constrain its application on 172 more general dynamic spiking neurons which are hard to approximate. Due to the effectiveness of 173 spike-based sequence learning, some SNNs-based language models are proposed for more efficient 174 language modeling, such as SpikeGPT (Zhu et al., 2023) and SpikeBERT (Lv et al., 2023). In con-175 trast to existing spiking SSMs, SPikE-SSM proposed in this paper realizes comprehensive parallel 176 acceleration with trainable temporal dynamics, efficiently achieving both high sparsity and excel-177 lent accuracy for long-range dependencies learning, which possesses the potential and prospects for constructing low-energy language models and enabling widespread applications. 178

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3 Method

3.1 PRELIMINARIES OF SSMS AND LIF NEURON

SSMs. According to (Gupta et al., 2022) and (Gu et al., 2021), SSMs provide a framework for long sequences modeling with lower computational complexity, which aims to transform an input sequence $x(t) = (x_0, \dots, x_{L-1}) \in \mathbb{R}^{1 \times L}$ into an output sequence $y(t)(y_0, \dots, y_{L-1}) \in \mathbb{R}^{1 \times L}$, where *L* is the length of sequence. This transformation occurs with the aid of an implicit latent state $h(t) \in \mathbb{R}^{N \times 1}$, which captures the underlying dynamics and relationships between the input and output sequences. The continuous representation of this model is formulated as:

$$\frac{dh(t)}{dt} = h'(t) = Ah(t) + Bx(t), y(t) = Ch(t),$$
(1)

where the state matrix $A \in \mathbb{R}^{N \times N}$ and vectors $B \in \mathbb{R}^{N \times 1}$, $C \in \mathbb{R}^{1 \times N}$ are the parameters. To adapt SSM to real-world discrete data, one can discretize the continuous formulation Eq. (1) with discretization rules such as zero-order hold (Gupta et al., 2022; Voelker et al., 2019). Then x(t) can be mapped to y(t) in a recurrent view:

$$\bar{A} = e^{\Delta A}, \bar{B} = A^{-1}(\bar{A} - I)B, \bar{C} = C \implies h_t = \bar{A}h_{t-1} + \bar{B}x_t, y_t = \bar{C}h_t,$$
 (2)

where $\Delta \in \mathbb{R}^+$ is the sample time, and $h_{-1} = 0$ for convenience. Note that the recurrence operation in Eq. (2) can be explicitly unrolled as a kernel view:

$$y_k = \sum_{j=0}^{k} \bar{K}_j \cdot x_{k-j}, \quad \bar{K} = \left(\overline{CB}, \overline{CAB}, \dots, \overline{CA}^{L-1}\bar{B}\right) \in \mathbb{R}^{1 \times L}, \tag{3}$$

which requires $\mathcal{O}(L^2)$ multiplications despite all the elements of y can be expediently computed in parallel by computing the kernel \overline{K} first. Fortunately, Eq. (3) can be accelerated by Fast Fourier Transform (FFT) (Duhamel & Vetterli, 1990) with time complexity $\mathcal{O}(L \log L)$ (Gupta et al., 2022).

LIF Neuron. The LIF neuron is widely used in spiking networks (Eshraghian et al., 2023), as it can capture the "leaky-integrate-fire-reset" process and balances ease of implementation with temporal dynamics by simplifying an RC circuit dynamical system (Gerstner et al., 2014). Let t denote the time step, the input currents I are linearly integrated into the membrane potential u in LIF neuron, the process of which can be formulated as follows.

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$$\tau \frac{du(t)}{dt} = -u(t) + IR, \quad u'_t = \beta u_{t-1} + (1-\beta)I_t, \quad s_t = H_s \left(u'_t - v_{\rm th}\right), \tag{4}$$

where $\tau \in \mathbb{R}$ is the time constant and β is its discrete-time equivalent. R denotes the resistivity. u'_t and u_t are the membrane potentials before and after the trigger of a spike. H_s denotes the the

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Alg	gorithm 1 The Optimization Process of Parallel Max-min Boundary Compression (PMBC)
Inj	put: Parameters τ , v_{th} , U_{th} ; Input signal $I \in \mathbb{R}^{1 \times L}$; Maximum of iterations M .
Ot	itput: Spiking signals $s \in \mathbb{R}^{1 \times L}$.
1:	Define $p = (\tau^0, \tau^1, \cdots, \tau^{L-1}); k = iFFT (FFT (I) \cdot FFT (p)).$
2:	Initialize $s^{up} = (1, \dots, 1) \in \mathbb{R}^{1 \times L}$ and $s^{low} = (0, \dots, 0) \in \mathbb{R}^{1 \times L}$.
3:	Repeat up to <i>M</i> times:
4:	$\hat{m}^{up} = U_{\text{th}} \cdot \text{iFFT} (\text{FFT}(p) \cdot \text{FFT}(s^{up})) + v_{\text{th}};$
5:	$m^{low} = U_{\rm th} \cdot {\rm iFFT} \left({\rm FFT} \left(p \right) \cdot {\rm FFT} \left(s^{low} \right) \right) + v_{\rm th};$
6:	If $k_t > m_t^{up}$, then $s_t^{low} = 1$; If $k_t < m_t^{low}$, then $s_t^{up} = 0$;
7:	Until convergence of spike rate $\frac{1}{L} \sum_{i} s_{i}^{low}$.
8:	Return $s = s^{low}$.

Heaviside function of LIF. As spikes are discrete events highly localized in time, a spike s is emitted when the membrane potential exceeds the firing threshold ($v_{th} \in \mathbb{R}$), that is $s_t = 1$, otherwise $s_t = 0$. After firing, the membrane voltage is adjusted by the reset mechanism, making subsequent spiking more difficult. Specifically, the membrane voltage is either reset to a specific value u_r (hard reset) or reduced by subtracting the same value v_{th} as the firing threshold (soft reset), that is:

soft reset:
$$u_t = u'_t - s_t v_{\text{th}}$$
, hard reset: $u_t = u'_t (1 - s_t) + u_r$. (5)

From Eq. (5) we can observe that the hard reset clears all historical membrane voltage signals, while the soft reset retains a proportion of them after spiking, which is more bio-plausible. Furthermore, we creatively decouple the firing threshold value and soft reset magnitude into $v_{\rm th}$ and $U_{\rm th}$ respectively, which can promote the representation capability of LIF neuron. However, all the reset mechanisms introduce unavoidable iterative computations due to the form of temporal dependence and Heaviside function, similar to the nonlinearities in RNN.

3.2 PARALLEL MAX-MIN BOUNDARY COMPRESSION (PMBC)

This subsection aims to address Challenge **0**. According to discretizing the LIF neuron with the soft reset mechanism in Eq. (5) combined with a decoupling reset magnitude $U_{\rm th}$, we can obtain the following formula:

$$u_t = \tau u_{t-1} - s_{t-1} U_{\text{th}} + I_t, \quad s_t = H_s \left(u_t - v_{\text{th}} \right).$$
(6)

The output membrane voltage u is iteratively computed by Eq. (6) since u_t depends on the spiking history from the previous time steps, notwithstanding the input current I can be obtained in parallel. This leads to a significant reduction in computational efficiency, especially for long sequence inputs. To solve this problem, we propose the following assertion, which lays the foundation for subsequent parallel computation to accelerate training and inference (See Appendix A.1 for the proof).

Assertion 3.1. The historical input signal I and spiking information s are deconstructed in the iteration process of Eq. (6), which is equivalent to:

$$u_t = k_t - m_t + v_{\text{th}}, \quad s_t = H_s \left(k_t - m_t \right),$$
(7)

where
$$k_t = \sum_{i=1}^{l} \tau^{t-i} I_i, \quad m_t = U_{\text{th}} \sum_{i=1}^{l-1} \tau^{t-1-i} s_i + v_{\text{th}}.$$
 (8)

261 Note that τ and $v_{\rm th}$ are fixed in one training step. Given I_i at all times, notice that the form of k is 262 the convolution of input sequence I and the exponential sequence of τ , we can obtain k_t at all times 263 in parallel with accelerated calculation through FFT. The question of whether we can obtain s_t in 264 parallel becomes how to obtain m_t at different times in parallel. To this end, we propose the PMBC 265 strategy to address Challenge \bullet . It can be observed that the spiking signal s is a binary variable, taking a value of either 0 or 1. Thus we can initialize the upper and lower bounds of m_t^{up} and m_t^{low} 266 by setting all the spiking signals $s_i = 1, (i = 0, \dots, L-1)$ and $s_j = 0, (j = 0, \dots, L-1)$ respec-267 tively. The two bounds can be utilized to compare with k_t simply, obtaining most spiking signals s_t 268 by parallel computation, and then update bounds values using these new s_t . This process can be it-269 erated until convergence in order to obtain all spiking states as shown in Figure 2(a). We summarize



283 the process of parallel computation of PMBC as Algorithm 1. After finite iterations of PMBC, there 284 may exist still a few fuzzy spiking signals s_i unidentified, which can be assigned randomly or based 285 on a prior distribution. The detailed discussion about the fuzzy spiking signals is provided in the Appendix C.3.2. To promote a lower spiking rate, we choose $s = s^{low}$ as the final output of spiking 286 signals. To accelerate training and inference, we implement the FFT and inverse FFT operations 287 with only setting M = 3, and the experimental results have proven that this configuration is capable 288 of identifying around 99% of the spiking signals without compromising accuracy. Our method can 289 determine the majority of spikes in the initial iterations, as shown in Figure 2(b). This is because the 290 distribution of k_t is closely tied to I, which is influenced by the normalization process before. With 291 a proper initialization of v_{th} , the first PMBC iteration effectively identifies that most spiking signals 292 are zero. This significantly reduces the number of required iterations and improve training efficiency 293 (e.g., M = 3 vs. L = 1024). Figure 2(c) provides an intuitive comparison between traditional serial computing and PMBC. The detailed analysis of the boundary evolution and convergence process of 295 PMBC in Figure 2 are described in the Appendix B.1. Particularly, we have the following assertion 296 (see Appendix A.2 for proof):

Assertion 3.2. For the input signal $y \in \mathbb{R}^{1 \times L}$, all the spiking signals can be identified with finite iterations of PMBC ($\leq L$), achieving significant acceleration compared to original serial computing.

300 3.3 REFRACTORY LIF NEURON MODEL

In biological neurons, spiking is usually followed by a refractory period during which new spiking is
 more difficult. This mechanism improves the overall sparsity of the network and could substantially
 reduce its energy consumption. Therefore, to simulate the intrinsic temporal dynamics of realistic
 neurons and further improve network sparsity, we introduce an innovative refractory LIF neuron
 model based on the soft reset mechanism, which effectively *addresses Challenge* . The LIF neuron
 with a refractory period can be mathematically described as:

$$u_t = \tau u_{t-1} + I_t - R_t U_{\text{th}}, \quad s_t = H_s(u_t - v_{\text{th}}),$$
(9)

where
$$R_t = \tau_r R_{t-1} + s_{t-1},$$
 (10)

In our refractory neuron model, τ_r is the refractory magnitude. R_t denotes the refractory periodbased sliding pulse, which is determined by both spiking signal s_{t-1} and R_{t-1} in the last time step. From Eq. (10) we can observe that the larger the value of the previous sliding pulse R_{t-1} , the greater R_t becomes, causing membrane voltage u_t to decrease accordingly, which makes it harder for the neuron to spike again during the refractory period. Similar to Assertion 3.1, we have the following results for the proposed refractory neuron model (see Appendix A.3 for the proof):

Assertion 3.3. In the refractory LIF neuron, the historical input signal I and spiking information s is deconstructed in the iteration process of Eq. (9), which is equivalent to:

$$u_t = k_t - m_t + v_{\text{th}}, \quad s_t = H_s \left(k_t - m_t \right),$$
(11)

where
$$k_t = \sum_{i=1}^t \tau^{t-i} I_i$$
, $m_t = U_{\text{th}} \sum_{i=1}^{t-1} \sum_{j=0}^{t-1-i} (\tau/\tau_r)^j \tau_r^{t-1-i} s_i + v_{\text{th}}$. (12)

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The PMBC algorithm of the refractory LIF neuron is summarized as Algorithm 2 in Appendix B.2, which only differs from the LIF neuron with soft reset in the representation of m_t .



Figure 3: The SPikE-SSM block. (Left) Forward computation graph of a single SPikE-SSM layer. (**Right**) Comparison of SSMs. The original SSM outputs floating-point numbers, while SPikE-SSM replaces its non-linearity with the proposed refractory neuron model, which can incorporate higher-level neuronal dynamics for long sequence modeling. *D*, *N*, and *L* represent the model dimension, SSM hidden dimension, and sequence length, respectively. SAF is the spiking activation function.

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3.4 THE BLOCK OF SPIKE-SSM

For Challenge 0, due to the exceptional long sequence modeling capability of SSMs, we integrate the proposed refractory neuron with soft reset mechanism and PMBC to the inherent SSM block, which aims to maintain both the high sparsity and excellent accuracy in the inference progress. In the proposed SPikE-SSM, we choose the original block of S4D model (Gu et al., 2022) as the backbone since it can achieve pragmatic simplification to enhance model efficiency as the latest diagonal version of SSM. Then the output y of the S4D block is activated by the proposed refractory neuron, hence Eq. (9) is rewritten as follows with Eq. (10) unchanged:

$$y_t = \bar{C}h_t, \quad u_t = \tau u_{t-1} + y_t - R_t U_{\text{th}}, \quad s_t = H_s(u_t - v_{\text{th}}).$$
 (13)

352 Inspired from (Rathi & Roy, 2021), we render $v_{\rm th}$ and $U_{\rm th}$ as trainable parameters within the SPikE-353 SSM block. This approach is motivated by their pivotal role in regulating the neuron's spiking 354 rate, thereby not only bolstering the SPikE-SSM's capability to attain exceptional performance and 355 expedite the convergence of PMBC, but also serving as a further stepping stone to tackle Challenge 356 **③** with greater efficacy. The results of Eq. (13) are fed into a linear layer that comprises a Conv1D 357 operation followed by a GLU activation function (Dauphin et al., 2017). The Conv1D enables 358 efficient local feature extraction, while the GLU activation selectively gates the information flow, 359 improving the model's ability to capture critical patterns in sparse binary data. Since the Heaviside 360 function H_s is non-differentiable at x = 0, we adopt the surrogate gradient (SG) method in SPikE-SSM. The details of SG in our method are provided in the Appendix B.3. 361

Figure 3 presents the forward computation graph of the SPikE-SSM block and the comparison with the original S4 blocks. Notably, from a neurobiological perspective, the SPikE-SSM block resembles a multi-time scale dendritic neuron (London & Häusser, 2005; Zheng et al., 2024), where hrepresents the dendrites and y the soma, both showing self-recurrent temporal dynamics.

4 EXPERIMENTS

In this section, we conduct extensive experiments to validate the superiority of our method, including the testing of long-range modeling capabilities on the sequential LRA and WikiText-103 tasks, with ablation studies and other related analyses. More experiments are shown in the Appendix C.

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4.1 DATASETS AND EXPERIMENTAL SETTINGS

Datasets. In this paper, we perform experiments on extensive long sequence databases, including
sequential MNIST (sMNIST) (Le et al., 2015), LRA benchmarks (comprising six tasks) (Tay et al.,
2020) and WikiText-103 (one large Wikipedia text data) (Merity et al., 2016). The Details of these datasets are shown in the Appendix C.1.

Table 2: Accuracy performance comparison of SPikE-SSM and state-of-the-art methods on the LRA benchmarks. Since the original S4D-Lin failed on the Path-X task, we report the results of its close variant, S4D-Inv. Following S4D, we assume 50% accuracy for Path-X when not available and calculate the overall average (AVG) across all tasks. The best two results are highlighted in bold.
For SpikingSSM and SPikE-SSM, the spiking rates (↓) of each task are highlighted in shaded gray areas. "—" indicates not applicable or unworkable, same for the other tables in this paper.

Method	SNN	ListOps	Text	Retrieval	Image	Pathfinder	Path-X	AVG
Transformer (Vaswani et al., 2017)	No	36.37	64.27	57.46	42.44	71.40	_	53.66
LMUFormer (Liu et al., 2024)	No	34.43	68.27	78.65	54.16	69.90	_	59.24
S4D-Lin (Gu et al., 2021)	No	60.52	86.97	90.96	87.93	93.96	92.80	85.52
Spiking LMUFormer (Liu et al., 2024)	Yes	37.30	65.80	79.76	55.65	72.68		60.20
Binary S4D (Stan & Rhodes, 2024)	Yes	54.80	82.50	85.03	82.00	82.60	61.20	74.69
S6-based SNN (Bal & Sengupta, 2024)	Yes	55.70	77.62	88.48	80.10	83.41	_	72.55
SpikingSSM (Shen et al. 2024)	Vac	59.93	82.35	88.20	86.81	93.68	94.80	84.30
Spikingssivi (Shen et al., 2024)	105	(13%)	(10%)	(6%)	(22%)	(7%)	(10%)	(11%)
SDile SSM (ours)	Vac	60.17	82.43	88.82	87.23	92.04	94.37	84.18
SFIKE-SSIVI (OUIS)	168	(12%)	(3 %)	(7%)	(10%)	(9 %)	(7%)	(8 %)

Table 3: Perplexity performance comparison of SPikE-SSM with SOTA methods on WikiText-103. The symbol \downarrow indicates that a smaller value for this metric is better, the same for other tables.

Method	SNN	Perplexity (\downarrow)	Parameters	Layer Count	Spiking Rate (\downarrow)
Transformer (Vaswani et al., 2017)	No	20.51	231M	48	_
S4 (Gu et al., 2021)	No	20.95	249M	48	—
SpikeGPT (Zhu et al., 2023)	Yes	39.75	213M	48	_
SpikingSSM (Shen et al., 2024)	Yes	33.94	75M	16	26.4%
SPikE-SSM (ours)	Yes	33.18	75M	16	24.5%

Implementation Details. The hyper-parameters τ and τ_r are set to 0.1 and 0.9, respectively. To ensure the threshold and refractory magnitude are positive during training, the trainable parameters v_{th} and U_{th} are computed by $\exp(v_{th})$ and $\exp(U_{th})$ with zero initialization (i.e. $\exp(v_{th})$ and $\exp(U_{th})$ are initialized as 1). Other parameters of SPikE-SSM blocks are initialized same as S4D-Lin (Gu et al., 2022). SPikE-SSM is trained with Pytorch library on four NVIDIA A100-SXM4-80GB GPUs and AMD EPYC 7642 48-core CPUs, using AdamW optimization (Loshchilov, 2017). For sCIFAR10, sMNIST, psMNIST and LRA benchmarks, the model is trained by the cross-entropy loss (Mao et al., 2023) with accuracy (Acc) results reported, while the Perplexity results are reported for WikiText-103. The division of training and test data is consistent with (Shen et al., 2024). The details of settings on nine different tasks are described in Table 7 in Appendix C.2, including six LRA benchmarks, three sequential vision tasks, and a large text dataset (WikiText-103).

4.2 PERFORMANCES COMPARISONS

Results on LRA Benchmarks. Table 2 compares SPikE-SSM with both non-spiking and spiking networks using Transformer or SSM architectures . While maintaining accuracy comparable to the original model, SPikE-SSM achieves an average network sparsity of less than 10%. Additionally, our model shows a significant performance improvement over previous SNN sequence models. No-tably, SPikE-SSM successfully tackles the Path-X task with extreme sparsity (only 0.07%). This task, which demands reasoning over long-range dependencies across sequences with 16,384 steps, is highly challenging and unsolvable by S4D-Lin, highlighting the robustness of our method.

Results on WikiText-103. In addition to LRA datasets, we further conduct experiments on the large Wikipedia text data, WikiText-103, to prove the advanced long sequence learning ability of SPikE-SSM against existing SOTA methods. The Perplexity results are shown in Table 3, which can be observed that SPikE-SSM achieves better performance with fewer parameters. Although model sparsity can improve computational efficiency, it is generally observed that achieving high accuracy often conflicts with maintaining strong sparsity, as sparsity typically results in information loss. However, it is particularly noteworthy that SPikE-SSM achieves both higher sparsity and accuracy compared to SpikingSSM, fully validating the effectiveness and superiority of the proposed model.

Table 4: Ablation studies of SPikE-SSM of different variants reported. Acc and SpkR denote Accuracy(%) \uparrow and Spiking Rate(%) \downarrow respectively. Spiking Rate is not applicable for ANN-S4D.

Dataset	sM	NIST	psM	INIST	sCIFAR10		
Criterion	Acc (%)	SpkR (%)	Acc (%)	SpkR (%)	Acc (%)	SpkR (%)	
ANN-S4D	99.50	_	98.20	_	87.11	_	
Spiking-S4D	99.46	7.81	97.68	7.73	85.34	12.70	
SPikE-SSM-SR	99.50	7.23	97.61	6.81	85.29	12.56	
SPikE-SSM-SRT	99.51	6.09	96.97	5.65	85.61	11.03	
SPikE-SSM-SRR	99.39	5.07	96.25	4.57	84.35	10.26	
SPikE-SSM-Full	99.53	5.56	97.89	5.13	85.67	9.85	

Table 5: Comparison of training speed of different methods. Training with the PMBC strategy achieves significant acceleration, the speed-up ratio amplifies with increasing sequence length.

Method	Speed (iterations / s) \uparrow						
	L = 1K	L = 2K	L = 4K	L = 8K			
Training with BPTT (Mozer, 2013)	0.60	0.29	0.11	0.03			
Training with SLTT (Meng et al., 2023)	0.73	0.33	0.12	0.03			
Training with PMBC (ours)	17.1	10.1	5.28	2.63			
Speed-up Ratio	25.6×	32.2×	$47.9 \times$	81.7×			

4.3 ABLATION STUDY

We conduct ablation studies to verify the design rationality of SPikE-SSM following the same experimental setups as Table 2. The variants with different levels of biological interpretability include:

• ANN-S4D. ANN-based SSM (S4D) model.

- Spiking-S4D. LIF-based spiking SSM without reset mechanism and refractory period.
- SPikE-SSM-SR. Only the soft reset mechanism is considered in the LIF neuron of SPikE-SSM block with PMBC, as shown in Eq. (6).
- SPikE-SSM-SRR. Both the soft reset mechanism and refractory period are considered in the LIF neuron of SPikE-SSM block with PMBC, as shown in Eq. (9-10).
- SPikE-SSM-SRT. Only the soft reset mechanism is considered in the LIF neuron of SPikE-SSM block with PMBC. Uth and Vth are trainable.
- SPikE-SSM-Full. Both the soft reset mechanism and refractory period are considered in the LIF neuron of SPikE-SSM block with PMBC. U_{th} and V_{th} are trainable.

Note that U_{th} and V_{th} are trainable only in SPikE-SSM-Full and SPikE-SSM-SRT. We compare the performances of different variants of SPikE-SSMs on sMNIST, psMNIST and sCIFAR10. The results are shown in Table 4, from which we can observe that each component designed for three Challenges is effective in SPikE-SSM. Specifically, the proposed refractory neuron model with the soft reset mechanism can optimize both high accuracy and pronounced sparsity with the thresholds v_{th} and refractory magnitudes U_{th} trainable in the SPikE-SSM block. More ablation studies about hyper-parameters τ and τ_r , fire modes of fuzzy spiking signals, and the number of iterations M in PMBC are shown in Tables 8 and 9 in the Appendix C.3.1, Tables 10 and 11 in the Appendix C.3.2, and Tables 12 and 13 respectively in the Appendix C.3.3, where our experiments in Figure 5 illus-trates that SPikE-SSM with fixed τ and τ_r performs better than that with trainable τ and τ_r .

4.4 TRAINING SPEED AND COMPUTATION COST ANALYSE

The Superiority of PMBC on Training Speed. We compare the training speed of SPikE-SSM, enhanced by our PMBC strategy, against traditional methods based on iterative LIF neurons, including Back-Propagation Through Time (BPTT) (Mozer, 2013) and the more recent Spatial Learning Through Time (SLTT) (Meng et al., 2023), which uses an optimized computational graph. The input consists of randomly generated 1-D sequences with various lengths of L = 1K, 2K, 4K, and 8K,



Table 6: Computation cost comparison of SSM with ANN settings, SpikingSSM and SPikE-SSM on WikiText-103. "Ops" is an abbreviation for "operations".

Figure 4: Spiking rate across all layers of SPikE-SSM and SpikingSSM on sCIFAR10 and WikiText-103 datasets. The number following each legend represents the respective average spiking rate.

and a batch size of 64. All time measurements were conducted on a single NVIDIA A100-SXM4-80GB GPU. As shown in Table 5, the speedup ratio using PMBC increases with sequence length, achieving a nearly two-order acceleration at 8K. 509

510 The Energy-efficiency of SPikE-SSM. We compare the energy costs of the proposed SPikE-511 SSM and its corresponding ANN-based version on WikiText-103, the sequence length of which is L = 8192. Spiking networks are considered energy-efficient due to sparse binary activation. The 512 multiplication between a binary activation and a floating-point weight can be performed using only 513 addition operations in some neuromorphic chips (Yao et al., 2024). As a result, the primary oper-514 ation in SNNs, synaptic accumulation (AC), incurs lower energy costs compared to the multiply-515 and-accumulate (MAC) operation in traditional ANNs. Although the hardware specifics and neuron 516 dynamics are not considered here, a theoretical analysis can provide an estimate of SNN efficiency. 517 Following previous studies (Yao et al., 2024; Li et al., 2024), we assume the energy cost of an MAC 518 operation is $E_{MAC} = 4.6pJ$, while an AC operation costs $E_{AC} = 0.9pJ$ (Horowitz, 2014). In this 519 part of the experiment, our model is set to comprise 16 layers, including a linear layer that projects 520 spikes from d = 1024 to d = 2048. For specific quantitative comparison, we first report the spiking 521 rates across different layers of SPikE-SSM in Figure 4. Then we report the MAC, AC, and energy consumption in these feature-mix layers since they occupy the majority of parameters and computa-522 tions. Specifically, if these projections were fully computed via floating-point multiplications (SSM 523 with ANN-based settings), they would require 275.2G MACs, consuming approximately 1.265J. 524 However, in our model (SPikE-SSM with SNN-based settings), the inputs to these layers are binary, 525 with an average spiking rate of less than 25%. Based on the spiking rates in Figure 4, our model per-526 forms 67.42G ACs, consuming 60.68mJ. The results are summarized in Table 6, which illustrate 527 the high energy efficiency of SPikE-SSM compared with ANN-based SSM and SpikingSSM. 528

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CONCLUSION 5

531 In this paper, we introduced SPikE-SSM, a novel spiking state space model designed to address key 532 challenges in long-sequence learning with SNNs. Specifically, we innovatively address the conflict of event-driven neuronal dynamics with parallel computing in long sequence modeling by the PMBC 534 method, enabling explicit and efficient training of neuronal dynamics. Subsequently, a concise resetrefractory neuron model is proposed to exploit the functionality of biological-plausible temporal 536 dynamics. Its effective integration with the SSM block and incorporation of trainable thresholds and refractory magnitudes realize a balance between sparsity and accuracy. Extensive experiments on sequential vision tasks, LRA benchmarks, and WikiText-103 language modeling validate the 538 superior efficiency, accuracy, and sparsity of SPikE-SSM. Our work shows the potential of dynamic spiking neurons in efficient long sequence learning.

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756 A THE PROOFS

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A.1 The Proof of Assertion 3.1

We utilize the Mathematical Induction Theory (Bather, 1994) to prove Assertion 3.1. As the presupposition, we set $s_0 = 0$ and $u_0 = 0$, which is reasonable since there's no signal at t = 0 with the time step of the spiking signal s, membrane voltage u, refractory momentum R and input current Isignal all considered as 1 to L, that is s_t , u_t , R_t and I_t , $t = 1, \dots, L$. On this basis, according to Eq. (6) ($u_t = \tau u_{t-1} - s_{t-1}U_{th} + I_t$, $s_t = H_s (u_t - v_{th})$), we have the following derivation.

Firstly, consider t = 1, we can obtain: 766

$$u_1 = \tau u_0 - s_0 U_{\rm th} + I_1 \xrightarrow{s_1 = H_s(u_1 - v_{\rm th})} k_1 = I_1, \quad m_1 = v_{\rm th}, \tag{14}$$

which is congruent to Eqs. (7-8) with t = 1. Then, to verify that Eqs. (7-8) hold for all $t \in [1, L]$, we assume that Eqs. (7-8) hold for $t = t_1 \in [1, L - 1]$, that is

$$u_{t_1} = k_{t_1} - m_{t_1} + v_{th}, \quad s_{t_1} = H_s \left(k_{t_1} - m_{t_1} \right), \tag{15}$$

where
$$k_{t_1} = \sum_{i=1}^{t_1} \tau^{t_1 - i} I_i, \quad m_{t_1} = U_{\text{th}} \sum_{i=1}^{t_1 - 1} \tau^{t_1 - 1 - i} s_i + v_{\text{th}},$$
 (16)

based on this we have:

$$u_{t_{1}+1} = \tau u_{t_{1}} - s_{t_{1}} U_{\text{th}} + I_{t_{1}+1}, \quad s_{t_{1}+1} = H_{s} \left(u_{t_{1}+1} - v_{\text{th}} \right).$$
(17)

According to substituting the u_{t_1} in Eq. (15) into Eq. (17), we have:

$$u_{t_{1}+1} = \tau \left(k_{t_{1}} - m_{t_{1}} + v_{\text{th}} \right) - s_{t_{1}} U_{\text{th}} + I_{t_{1}+1}$$
(18)

$$=\tau\left(\sum_{i=1}^{t_1}\tau^{t_1-i}I_i - \left(U_{\rm th}\sum_{i=1}^{t_1-1}\tau^{t_1-1-i}s_i + v_{\rm th}\right) + v_{\rm th}\right) - s_{t_1}U_{\rm th} + I_{t_1+1} \qquad (19)$$

$$=\tau \sum_{i=1}^{t_1} \tau^{t_1-i} I_i + I_{t_1+1} - U_{\text{th}} \sum_{i=1}^{t_1-1} \tau^{t_1-1-i} s_i - s_{t_1} U_{\text{th}}$$
(20)

$$=\sum_{i=1}^{t_1+1} \tau^{t_1+1-i} I_i - \left(U_{\rm th} \sum_{i=1}^{t_1} \tau^{t_1-i} s_i + v_{\rm th} \right) + v_{\rm th}$$
(21)

$$=k_{t_1+1} - m_{t_1+1} + v_{\rm th} \tag{22}$$

Therefore, Eqs. (7-8) hold for $t = t_1 + 1 \in [2, L]$. According to the Mathematical Induction Theory (Bather, 1994), we can conclude that Eqs. (7-8) hold for all the $t \in [1, L]$, so that Assertion 3.1 holds. Q.E.D.

A.2 THE PROOF OF ASSERTION 3.2

We assume that the dimension of input signal I is L for the refractory LIF neurons with the soft reset. Firstly, according to Eq. (7) and Eq. (8), we have

$$k_1 = I_1, \quad m_1 = v_{\rm th}.$$
 (23)

Hence we can definitely obtain $s_1 = H_s (k_1 - m_1)$ in the first iteration of PMBC. Then, without loss of generality, for the (a + 1)-th iteration of PMBC, we assume that the first *a* spiking signals (i.e. s_1, s_2, \dots, s_a) have been obtained. Note that m_{a+1} is only related to the first *a* spiking signals:

$$k_{a+1} = \sum_{i=1}^{a+1} p_{a+1-i}^{a+1} y_i, \quad m_{a+1} = U_{\text{th}} \sum_{i=1}^{a} q_{a-i}^{a+1} s_i + v_{\text{th}},$$
(24)

where $p^{a+1} = (\tau^0, \tau^1, \dots, \tau^a)$, $q^{a+1} = (0, \tau^0, \tau^1, \dots, \tau^{a-1})$. Therefore, in the *b*-th iteration of PMBC, at least one new spiking signal (s_{a+1}) can be definitely obtained by $s_{a+1} = H_s(k_{a+1} - m_{a+1})$, thus the maximum number of iterations of PMBC is no more than *L*. Q.E.D. 810 In fact, due to the parallel computing mechanism of PMBC, the actual number of iterations is much 811 smaller than L. As shown in Figure 2(b), after only 5 iterations, the explicit spiking state exceeds 812 99%, whereas processing a 1024-dimensional input sequence serially would require 1024 iterations, 813 highlighting the efficiency of PMBC. Detailed experimental proofs and analysis are shown in Sec-814 tion B.1. Based on the proof process for Assertion 3.1, we can conclude that the distribution of k_t is closely related to y_t , where y_t is the output obtained from the previous SSM block, calcu-815 lated through layer normalization. Therefore, if SPikE-SSM initializes $v_{th} = 1$ during training, the 816 first iteration of PMBC can effectively determine that most of the spiking signals are zero. This 817 significantly reduces the number of required iterations and accelerates training efficiency. 818

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A.3 THE PROOF OF ASSERTION 3.3

Similarly to the proof of Assertion 3.1, we utilize the Mathematical Induction Theory (Bather, 1994) to proof Assertion 3.3. As the presupposition, we set $s_0 = 0$, $u_0 = 0$, and $R_0 = 0$, which is reasonable since there's no signal at t = 0 with the time step of the spiking signal s, membrane voltage u and input current I signal all considered as 1 to L, that is s_t , u_t , and I_t , $t = 1, \dots, L$. On this basis, according to Eqs. (11-12) ($u_t = \tau u_{t-1} + I_t - R_t U_{\text{th}}$, $R_t = \tau_r R_{t-1} + s_{t-1}$, $s_t = H(u_t - v_{\text{th}})$), we can transform the problem into proving the following equation first.

$$u_t = k_t - m_t + v_{\text{th}}, \quad s_t = H_s \left(k_t - m_t \right),$$
(25)

where
$$k_t = \sum_{i=1}^{t} \tau^{t-i} I_i$$
, $m_t = U_{\text{th}} \sum_{i=1}^{t-1} \tau^{t-1-i} R_{i+1} + v_{\text{th}}$, $R_{t+1} = \tau_r R_t + s_t$, (26)

Firstly, consider t = 1, we can obtain:

$$u_1 = \tau u_0 - R_1 U_{\rm th} + I_1 \xrightarrow[R_1 = \tau_r R_0 + s_0]{} k_1 = I_1, \quad m_1 = v_{\rm th},$$
(27)

which is congruent to Eq. (26) with t = 1. Then, to verify that Eqs. (25-26) hold for all $t \in [1, L]$, we assume that Eqs. (25-26) hold for $t = t_1 \in [1, L-1]$, that is

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$$u_{t_1} = k_{t_1} - m_{t_1} + v_{th}, \quad s_{t_1} = H_s \left(k_{t_1} - m_{t_1} \right), \tag{28}$$

where
$$k_{t_1} = \sum_{i=1}^{t_1} \tau^{t_1 - i} I_i$$
, $m_{t_1} = U_{\text{th}} \sum_{i=1}^{t_1 - 1} \tau^{t_1 - 1 - i} R_{i+1} + v_{\text{th}}$, $R_{t_1 + 1} = \tau_r R_{t_1} + s_t$, (29)

based on this we have:

$$u_{t_1+1} = \tau u_{t_1} - R_{t_1+1}U_{\text{th}} + I_{t_1+1}, \quad s_{t_1+1} = H_s \left(u_{t_1+1} - v_{\text{th}} \right).$$
(30)

According to substituting the u_{t_1} in Eq. (28) into Eq. (30), we have:

$$u_{t_{1}+1} = \tau \left(k_{t_{1}} - m_{t_{1}} + v_{th}\right) - R_{t_{1}+1}U_{th} + I_{t_{1}+1}$$
(31)

$$=\tau\left(\sum_{i=1}^{t_1}\tau^{t_1-i}I_i - \left(U_{\rm th}\sum_{i=1}^{t_1-1}\tau^{t_1-1-i}R_{i+1} + v_{\rm th}\right) + v_{\rm th}\right) - R_{t_1+1}U_{\rm th} + I_{t_1+1} \quad (32)$$

$$=\tau \sum_{i=1}^{t_1} \tau^{t_1-i} I_i + I_{t_1+1} - U_{\text{th}} \sum_{i=1}^{t_1-1} \tau^{t_1-1-i} R_{i+1} - R_{t_1+1} U_{\text{th}}$$
(33)

$$=\sum_{i=1}^{t_1+1} \tau^{t_1+1-i} I_i - \left(U_{\rm th} \sum_{i=1}^{t_1} \tau^{t_1-i} R_{i+1} + v_{\rm th} \right) + v_{\rm th}$$
(34)

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$$=k_{t_1+1} - m_{t_1+1} + v_{\text{th}}$$
(35)

Therefore, Eqs. (25-26) hold for $t = t_1 + 1 \in [2, L]$. According to the Mathematical Induction Theory (Bather, 1994), we can conclude that Eqs. (25-26) hold for all the $t \in [1, L]$. Subsequently, the parallel calculation form of the membrane potential of the LIF neuron with soft reset and refractory =

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⁸⁶⁴ period can be expressed as:

$$u_t = \sum_{i=1}^t \tau^{t-i} I_i - U_{\text{th}} \sum_{i=1}^t \tau^{t-i} R_i$$
(36)

$$=\sum_{i=1}^{t} \tau^{t-i} I_i - U_{\text{th}} \sum_{i=1}^{t} \tau^{t-i} (\sum_{j=1}^{i-1} \tau_r^{i-1-j} s_j)$$
(37)

$$=\sum_{i=1}^{t} \tau^{t-i} I_i - U_{\text{th}} \sum_{i=1}^{t-1} \sum_{j=0}^{t-1-i} \tau^j \tau_r^{t-1-i-j} s_i$$
(38)

$$=\sum_{i=1}^{t} \tau^{t-i} I_i - U_{\rm th} \sum_{i=1}^{t-1} \sum_{j=0}^{t-1-i} (\tau/\tau_r)^j \tau_r^{t-1-i} s_i \tag{39}$$

$$s_t = H_s(u_t - v_{\text{th}}), \quad k_t = \sum_{i=1}^{\tau} \tau^{t-i} I_i$$
 (40)

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$$m_t = U_{\rm th} \sum_{i=1}^{t-1} \sum_{j=0}^{t-1-i} (\tau/\tau_r)^j \tau_r^{t-1-i} s_i + v_{\rm th}$$
(41)

Q.E.D.

Therefore, Assertion 3.3 holds.

Apparently, Eqs. (11-12) degrade to the version of SPikE-SSM-RS without the refractory period when $\tau_r = 0$.

B MORE DETAILS RELATED TO SPIKE-SSM

B.1 DETAILED DESCRIPTION AND ANALYSIS FOR FIGURE 2

To clearly demonstrate the implementation process and effectiveness of PMBC, we present the evolution of the boundary and convergence process in Figure 2.

(1) In Figure 2(a), we illustrate the boundary's evolution of a particular neuron chosen at random 894 during training as PMBC iterations increase over different time steps. In the top part of Figure 2(a), 895 before the first iteration, with all spiking signals s_i setting to 1, $m_t^{up}(0)$ increases slowly as the time 896 step t progresses, with growth gradually leveling off until stabilizing. This is consistent with Eq. (8), 897 as $0 < \tau < 1$ causes τ^t to approach 0 over time, $m_t^{up}(0)$ approaches $U_{\rm th}/(1-\tau) + v_{\rm th}$ when t is large. Conversely, with all spiking signals s_i setting to 0, $m_t^{low}(0)$ remains v_{th} across all time steps. 899 After the first iteration of PMBC, partial binary spiking signals s_i are explicitly obtained, leading 900 to two outcomes: in some time steps, $m_t^{low}(1)$ apparently increases compared to $m_t^{low}(0)$, while 901 in others, $m_t^{up}(1)$ apparently decreases compared to $m_t^{up}(0)$. It is observed that spiking states that 902 are quickly identified are often following several k_t that are too large or too small in succession. In the bottom part of Figure 2(a), inherited from the convergence process of m_t^{up} and m_t^{low} in 903 the top part, the results after 5 iterations of PMBC are shown, where $m_t^{low}(5)$ and $m_t^{up}(5)$ become 904 almost identical. Eventually, nearly all binary spiking signals s_i are explicitly determined based on 905 $s_t = H_s(k_t - m_t)$, by comparing k_t and m_t in parallel and efficiently, omitting the serial computing 906 of membrane potential u_t . 907

(2) In Figure 2(b), we show the convergence curve of the explicit spiking state as PMBC iterations increase. The explicit spiking state refers to the proportion of spiking signals that are explicitly determined across all time steps. From Figure 2(b), it is evident that PMBC can resolve most of the spikes in just a few iterations, significantly fewer than the original serial computation method used for LIF neurons (Eq. (4)). After only 5 iterations, the explicit spiking state exceeds 99%, whereas processing a 1024-dimensional input sequence serially would require 1024 iterations, highlighting the efficiency of PMBC.

915 (3) In Figure 2(c), we provide an intuitive comparison between the PMBC method based on parallel 916 computing and the traditional serial computation method. For a sequence of length L, the serial 917 method requires L iterations, whereas the PMBC approach processes all L tokens simultaneously in 918 parallel, requiring only M iterations, with M much smaller than L.

Alg	orithm 2 The Optimization Process of PMBC for the Refractory LIF Neuron
In	put: Parameters $\tau, \tau_r, U_{\text{th}}$ and v_{th} ; Input signal $I \in \mathbb{R}^{1 \times L}$; Maximum of iterations M .
0ι	itput: Spiking signals $s \in \mathbb{R}^{1 \times L}$.
1:	Define $p = (\tau^0, \tau^1, \cdots, \tau^{L-1}), q : q_t = \tau_r^t \cdot \sum_{j=0}^t (\tau/\tau_r)^j; k = iFFT (FFT (I) \cdot FFT (p)).$
2:	Initialize $s^{up} = (1, \dots, 1) \in \mathbb{R}^{1 \times L}$ and $s^{low} = (0, \dots, 0) \in \mathbb{R}^{1 \times L}$.
3:	Repeat up to <i>M</i> times:
4:	$m^{up} = U_{\rm th} \cdot {\rm iFFT} \left({\rm FFT} \left(q \right) \cdot {\rm FFT} \left(s^{up} \right) \right) + v_{\rm th};$
5:	$m^{low} = U_{th} \cdot iFFT(FFT(q) \cdot FFT(s^{low})) + v_{th};$
6.	If $k_{\pm} > m_{\pm}^{up}$ then $s_{\pm}^{low} = 1$. If $k_{\pm} < m_{\pm}^{low}$ then $s_{\pm}^{up} = 0$.
7.	Until convergence of spike rate $\frac{1}{2}\sum_{t}s^{low}$
· · ·	$\mathbf{D}_{\mathbf{d}_{i}} = \mathbf{D}_{\mathbf{d}_{i}} = \mathbf{D}_{\mathbf{d}$
8:	Return $s \equiv s^{100}$.

In fact, the parallel PMBC method makes explicitly training parametric LIF neurons more efficient and feasible, especially in long sequence scenarios. In contrast, the SDN-based approach in SpikingSSM (Shen et al., 2024) is limited to effectively training only the threshold voltage (v_{th}) , while it's unclear whether other hyper-parameters can be trained end-to-end. This ambiguity highlights the advantage of PMBC in handling more comprehensive parameter optimization with trainable temporal dynamics.

⁹³⁸ The experimental setup for this part is described as follows: ⁹³⁹

For Figure 2(a): Both the soft reset mechanism and refractory period are considered in the LIF neuron of SPikE-SSM block trained with PMBC. The parameters τ , τ_r , U_{th} , and v_{th} are fixed as 0.3, 0.4, 1, and 0.5 respectively. The number of iterations in PMBC is set to 10 during training. The curve in Figure 2(a) shows the results with different iteration numbers in PMBC during inference.

For Figure 2(b): Only the soft reset mechanism is considered in the LIF neuron of the SPikE-SSM block trained with PMBC. The parameters τ , U_{th} , and v_{th} are fixed as 0.1, 1, and 3 respectively. The number of iterations in PMBC is set to 10 during training. After all the iterations of PMBC, the still fuzzy spiking signals are set to 1 by default, i.e. the spiking signals for these time steps are set to fire by default. The curve in Figure 2(b) shows the explicit spiking state with different iteration numbers in PMBC during inference.

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B.2 THE OPTIMIZATION PROCESS OF PMBC FOR THE REFRACTORY LIF NEURON

The pseudo-code of the optimization process of PMBC for the Refractory LIF Neuron is summarized in Algorithm 2, which only differs from the LIF neuron with soft reset in the representation of m_t in Algorithm 1.

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B.3 SURROGATE GRADIENT IN SPIKE-SSM

Since the Heaviside function H_s in Eq. (13) is non-differentiable at x = 0, several surrogate gradient (SG) methods are proposed to enable training through gradient descent. Common SG functions are differentiable at all points and possess non-zero derivatives near the threshold, allowing them to approximate the original discontinuous gradient of the spiking activation function, such as the rectangular function (Zheng et al., 2021) and the triangular function (Bellec et al., 2018). In SPikE-SSM, the piecewise quadratic surrogate spiking function g(x) is utilized with $\alpha = 1$. g(x) and its gradient g'(x) are defined as:

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$$\begin{array}{ll} \mathbf{968} \\ \mathbf{969} \\ \mathbf{970} \end{array} \qquad g(x) = \begin{cases} 0, & \text{if } x < -\frac{1}{\alpha} \\ -\frac{1}{2}\alpha^2 |x|x + \alpha x + \frac{1}{2}, & \text{if } |x| \le \frac{1}{\alpha} \\ 1, & \text{if } x > \frac{1}{\alpha} \end{cases}, \quad g'(x) = \begin{cases} 0, & \text{if } |x| > \frac{1}{\alpha} \\ -\alpha^2 |x| + \alpha, & \text{if } |x| \le \frac{1}{\alpha} \end{cases}$$

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 $\frac{\frac{1}{\alpha}}{\frac{1}{2}}$. (42)



Figure 5: Comparison on sCIFAR10 between SPikE-SSM-SRR with fixed τ and τ_r and that with trainable τ and τ_r . U_{th} and U_{th} are both set to 1 in SPikE-SSM-SRR.

Table 7: The hyper-parameters of our experiments on these datasets. **H** denotes the model dimension, **N** denotes the state dimension, **LR** denotes learning rate, **WD** denotes weight decay and **BS** denotes the batch size. **BN** and **LN** refer to Batch Normalization and Layer Normalization.

Dataset	Depth	Н	Ν	Norm	pNorm	Dropout	LR	BS	Epochs	WD	$(\Delta_{min}, \Delta_{max})$
sMNIST	2	128	64	LN	False	0.1	0.01	64	25	0.01	(0.001,0.1)
psMNIST	4	128	64	LN	False	0.1	0.01	64	60	0.01	(0.001, 0.1)
sCIFAR10	4	128	64	LN	False	0.1	0.01	64	100	0.01	(0.001, 0.1)
ListOps	8	128	64	BN	False	0	0.01	50	40	0.05	(0.001, 0.1)
Text	6	256	64	BN	True	0	0.01	16	32	0.01	(0.001, 0.1)
Retrieval	6	256	64	BN	True	0	0.01	64	20	0.01	(0.001, 0.1)
Image	6	512	64	LN	False	0.1	0.01	50	200	0.01	(0.001, 0.1)
Pathfinder	6	256	64	BN	True	0	0.004	64	200	0.01	(0.001, 0.1)
Path-X	6	256	64	BN	True	0	0.0005	32	50	0.01	(0.0001, 0.01)
WT-103	16	1024	64	LN	True	0.1	0.0005	1	200	0.01	(0.001,0.1)

C MORE DETAILS RELATED TO EXPERIMENTS

1001 C.1 DETAILS OF DATASETS

1003 C.1.1 SEQUENTIAL VISION DATASETS

The MNIST dataset (Deng, 2012) is a classic benchmark in machine learning, featuring 70,000 grayscale images of handwritten digits (0-9), with 60,000 samples for training and 10,000 for testing, each image sized at 28×28 pixels. It has long been a cornerstone for evaluating image classi-fication models due to its simplicity and widespread availability. The sequential MNIST (sMNIST) dataset (Le et al., 2015) transforms these 2D images into sequences of 784 elements by flattening the pixel grid into a 1D sequence. This transformation poses a more complex challenge, as models must process and retain information over longer time steps to accurately classify the digit. To further increase the difficulty, the permuted sequential MNIST (psMNIST) (Le et al., 2015) applies a fixed random permutation to the pixel sequences, effectively scrambling their spatial order and disrupting any inherent structure. This variant demands even greater computational ability from models, as they must learn to extract meaningful features from a sequence with no obvious temporal or spatial coherence, making psMNIST a far more challenging task compared to the original sMNIST. The sCIFAR10 dataset is a sequential version of the CIFAR-10 dataset (Krizhevsky et al., 2009), where the original static images are processed in a temporal manner, typically by feeding the image pixels row-by-row or in a pre-defined sequence.

1019 C.1.2 LRA BENCHMARK

1021The LRA benchmark (Tay et al., 2020) was specifically designed to evaluate the performance of se-1022quence models in long-context scenarios, where capturing dependencies across extended sequences1023is crucial. It consists of six diverse tasks, with sequence lengths ranging from 1K to 16K steps,1024covering multiple modalities including visual data, mathematical expressions, and natural language.1025These tasks are carefully curated to challenge models on various aspects of long-context compre-
hensions, such as text classification, document retrieval, image recognition, the pathfinder problem,

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Table 8: Ablation studies of SPikE-SSM-SR (v_{th} and U_{th} are trainable) with different τ . Acc, SpkR and FzR denote test accuracy, spiking rate and fuzzy rate respectively.

Iter Nums	τ	0.2	0.15	0.1
5	$\begin{vmatrix} Acc (\%) \uparrow \\ SpkR (\%) \downarrow \\ FzR (\%) \downarrow \end{vmatrix}$	84.91 10.66 1.88	84.93 10.69 1.29	84.93 9.91 1.30
20	$\begin{vmatrix} Acc (\%) \uparrow \\ SpkR (\%) \downarrow \\ FzR (\%) \downarrow \end{vmatrix}$	85.24 10.03 0.48	84.37 9.72 0.36	85.12 9.98 0.37

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Table 9: Ablation studies of SPikE-SSM-Full with different τ and τ_r . Acc, SpkR and FzR denote test accuracy, spiking rate and fuzzy rate respectively. Fuzzy rate is defined as the mean proportion of final unidentified spiking signals s_i in all neurons after finite iterations of PMBC during inference.

au			0.2			0.15			0.1		
Iter Nums	$ au_r$	0.6	0.75	0.9	0.6	0.75	0.9	0.6	0.75	0.9	
5	$\begin{vmatrix} Acc (\%) \uparrow \\ SpkR (\%) \downarrow \\ FzR (\%) \downarrow \end{vmatrix}$	84.97 10.27 2.54	84.60 10.23 3.09	83.32 10.21 6.25	84.13 9.93 2.03	84.34 9.94 2.27	84.75 9.94 5.74	85.17 9.46 2.00	85.04 9.63 2.39	84.37 9.61 5.82	
20	$\begin{vmatrix} Acc (\%) \uparrow \\ SpkR (\%) \downarrow \\ FzR (\%) \downarrow \end{vmatrix}$	84.64 10.28 1.07	84.56 10.55 1.64	83.32 10.73 3.05	84.25 10.41 0.94	84.17 10.39 1.21	83.85 10.22 3.1	84.66 10.10 0.95	84.47 10.00 1.18	83.56 10.33 2.87	

and list operations (ListOps), making it a comprehensive testbed for assessing a model's ability to 1053 process and reason over extended input sequences. 1054

1056 C.1.3 WIKITEXT-103

1057 The WikiText-103 dataset (Merity et al., 2016) is a large-scale corpus containing over 100 million to-1058 kens extracted from Wikipedia articles that have been rated as Good or Featured. Spanning a broad 1059 spectrum of topics and domains, it offers a rich variety of linguistic patterns and structures. Unlike many other datasets, WikiText-103 consists of full-length articles rather than isolated snippets, 1061 making it particularly well-suited for models designed to capture long-term dependencies across 1062 extended contexts. Due to its depth and diversity, it has become a pivotal benchmark for word-level language modeling, providing a robust testing ground for evaluating models' capacity to understand 1064 and generate coherent text over lengthy sequences.

C.2 DETAILS OF EXPERIMENTAL SETTINGS 1067

1068 Table 7 describes the particular training details of experiments on different tasks, where LRA bench-1069 marks consist of six tasks, including ListOps, Text, Retrieval, Image, Pathfinder and Path-X. Depth 1070 denotes the number of SPikE-SSM blocks, pNorm denotes the pre-norm, and WT - 103 means 1071 WikiText-103 dataset.

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- C.3 MORE EXPERIMENTAL RESULTS 1074
- 1075 C.3.1 PARAMETER SENSITIVITY ANALYSIS

In the blocks of the proposed model, we compare the performances of SPikE-SSM-SSR with fixed 1077 τ and τ_r and that with trainable τ and τ_r . The results are shown in Figure 5, which shows that 1078 SPikE-SSM with fixed τ and τ_r can achieve lower sparsity and higher accuracy. Therefore, we τ and 1079 τ_r are set fixed in our method. Then we investigate the impacts of different τ and τ_r on our model.

Table 10: The impact of different fire modes for the fuzzy spiking signals on SPikE-SSM-SR with fixed $U_{\rm th} = 1$ and $V_{\rm th} = 1$.

Criterion	No Reset	Fire Mode 1	Fire Mode 2	Fire Mode 3	Fire Mode 4
Accuracy (%) ↑	85.31	86.11	85.64	85.68	85.55
Spiking Rate (%) ↓	12.27	13.07	12.19	12.19	11.94

Table 11: The impact of different fire modes for the fuzzy spiking signals on SPikE-SSM-Full with refractory period and trainable $U_{\rm th}$ and $V_{\rm th}$.

Criterion	No Reset	Fire Mode 1	Fire Mode 2	Fire Mode 3	Fire Mode 4
Accuracy (%) ↑	85.45	85.34	84.80	84.84	85.23
Spiking Rate (%)↓	14.81	10.16	9.97	10.87	9.75

1097 The results for SPikE-SSM-Full and SPikE-SSM-SR (v_{th} and U_{th} are trainable) are shown in Table 8 1098 and Table 9 respectively. Note that there are two hyper-parameters (τ and τ_r) in SPikE-SSM-Full, 1099 and only one hyper-parameter (τ) in SPikE-SSM-SR. As previously stated by the default setting, the 1000 fuzzy spiking signals s_f are all set to False during training (i.e. $s_f = 0$).

From Table 8, we can observe that: (1) In SPikE-SSM-SR with only one hyper-parameter τ , re-1101 gardless of the number of PMBC iterations, the change in τ has minimal impact on the model's 1102 overall accuracy and sparsity, indicating that the performance of SPikE-SSM-SR is not sensitive to 1103 the parameter τ , which demonstrates the robustness of our proposed method. (2) As the number 1104 of PMBC iterations increases, the Fuzzy Rate significantly decreases, as more iterations in PMBC 1105 allow for more spiking signals to be identified. (3) Under the same number of PMBC iterations, the 1106 Fuzzy Rate decreases significantly as the hyper-parameter τ becomes smaller. This indicates that a 1107 smaller τ helps improve the computational efficiency of PMBC and the accuracy of the model. 1108

From Table 9, we can observe that: (1) In SPikE-SSM-SR with two hyper-parameters τ and τ_r , under the same number of PMBC iterations and τ_r , the change in τ has minimal impact on the model's overall accuracy, but smaller τ leads to smaller Spiking Rate and Fuzzy Rate. This indicates that a smaller τ helps reduce the sparsity and improve the accuracy of the model. (2) Under the same number of PMBC iterations and τ , the change in τ_r has minimal impact on the model's overall accuracy and sparsity, demonstrating the robustness of our method.

Therefore, the hyper-parameters τ and τ_r are set to 0.1 and 0.9 respectively in the full SPikE-SSMby default, which can achieve a more balanced performance between the sparsity and accuracy.

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1118 C.3.2 THE IMPACT OF FUZZY FIRE MODES IN PMBC

1119 Due to the parallel computation of PMBC, most of the spiking signals can be definitely obtained in 1120 5-10 iterations of Algorithm 1 and Algorithm 2, with only a few spiking signals are still fuzzy. To 1121 improve computing efficiency, we adopt several fire modes to address these fuzzy spiking signals to 1122 reduce the iterations of PMBC, including:

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• No Reset: Train the SPikE-SSM without reset mechanism (so $U_{\rm th}$ and refractory period are not applicable).

- Fire Mode 1: The fuzzy spiking signals s_f are all set to True (i.e. $s_f = 1$).
- Fire Mode 2: The fuzzy spiking signals s_f are all set to False (i.e. $s_f = 0$).
- Fire Mode 3: The fuzzy spiking signals s_f randomly are determined by the mean spiking rate value of definite piking signals s_d .
- Fire Mode 4: The fuzzy spiking signals s_f are determined by their current corresponding upper and lower bounds. For example, after all the iterations of PMBC, if s_i is still fuzzy in time step t = i, and its upper and lower bounds are m_i^{up} and m_i^{low} . If $k_i > (m_i^{up} + m_i^{low})/2$, then $s_i = 1$, or else $s_i = 0$.

1135	Table 12: The impact of different numbers of iterations in PMBC on the accuracy and spiking rate
1136	performances of our method.

Number of Iterations	1	2	5	10	30
Accuracy (%) ↑	84.46	84.74	84.74	85.02	84.99
Spiking Rate (%) \downarrow	13.32	12.90	12.66	12.11	12.32
Fuzzy Rate (%) \downarrow	8.80	6.77	4.48	2.62	0.82

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Table 13: The impact of different numbers of iterations in PMBC on the speed and time cost of our 1144 method. "serial computing" means serially training SPikE-SSM without PMBC. 1145

1146 1147	Number of Iterations	1	2	5	10	30	50	serial computing
1148	Speed (iters/s) ↑	25.32	21.51	14.63	9.50	3.97	2.42	1.21
1149	Time (ms/iters) \downarrow	39.49	46.49	68.35	105.26	251.89	413.22	826.45

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To investigate the impact of different fire modes of fuzzy spiking signals for PMBC, we trained the 1152 SPikE-SSM-SR and SPikE-SSM-Full with different fire modes on dataset sCifar10 with 10 PMBC 1153 iterations and 100 epochs, and the performances are shown in Table 10 and Table 11 respectively. 1154

1155 According to Table 10, we can observe that SPikE-SSM-SR with Fire Mode 2, Fire Mode 3 and Fire 1156 Mode 4 can all achieve both higher accuracy and lower spiking rate than that of SPikE-SSM without 1157 the reset mechanism, which verifies the effectiveness of reset mechanism in the refractory LIF neuron model. Although SPikE-SSM-SR with Fire Mode 1 achieves the highest spiking rate, it also 1158 achieves the highest accuracy among the five versions, which can be attributed to the fact that too 1159 sparse signals may lead to the loss of important information during training. 1160

1161 According to Table 11, we can observe that SPikE-SSM-Full with Fire Mode 1, Fire Mode 2, Fire 1162 Mode 3, and Fire Mode 4 can all achieve lower spiking rates than that of SPikE-SSM without the 1163 reset mechanism. Notably, compared with Table 10 and Table 11, we can observe that SPikE-SSM-1164 Full with both trainable U_{th} and v_{th} can achieve lower spiking rate than SPikE-SSM-SR with the same Fire Modes in all the four different Fire Modes, which indicates that the trainable $U_{\rm th}$ and $V_{\rm th}$ 1165 can more effectively model the temporal dynamics with biological interpretability. 1166

1167 In our method, we choose Fire Mode 2 as the default setting since it can further reduce the spiking 1168 rate with all the fuzzy spiking signals set to False ($s_f = 0$), and it can also achieve a more balanced 1169 performance between accuracy and spiking rate.

1170

1171 C.3.3 THE IMPACT OF DIFFERENT ITERATIONS IN PMBC 1172

In this section, we investigate the impact of different numbers of iterations in PMBC on the accuracy 1173 and sparsity performances of SPikE-SSM. We trained the SPikE-SSM-SR with fixed $U_{th} = v_{th} = 1$ 1174 and 4 layers on dataset sCIFAR10 with 100 epochs and $\tau = 0.2$. This experiment is conducted on a 1175 V-100 GPU. The results with different iterations in PMBC are shown in Table 12. 1176

1177 According to Table 12, we can observe that: (1) As the number of PMBC iterations increases, 1178 the model's accuracy gradually improves, while the fuzzy rate decreases. This is because more 1179 spiking signals are explicitly calculated with additional iterations, enhancing the model's robustness and leading to a steady rise in accuracy. (2) As the number of PMBC iterations increases, the 1180 overall spiking rate gradually decreases, indicating that more distinct spiking signals can enhance 1181 the model's sparsity. (3) The SPikE-SSM-SR with 10 iterations of PMBC achieves both higher 1182 accuracy and a lower spiking rate compared to the version with 30 iterations, despite having a 1183 significantly higher fuzzy rate. This suggests that more iterations of PMBC are not always better, as 1184 fuzzy spiking signals may function similarly to dropout. 1185

In addition, we investigate the impact of different and more granular numbers of iterations in PMBC 1186 on the inference speed and the cost times of SPikE-SSM. The experimental settings are the same as 1187 Table 12. Note that the sequence length L of sCIFAR10 is 1024. The experimental results are shown in Table 13, from which we can observe that: (1) As the number of iterations in PMBC increases, the model's inference speed progressively decreases, with each iteration requiring more time to com-plete. This slowdown becomes more noticeable as iteration counts grow. (2) The fastest inference speed is achieved when only a single iteration is performed. However, even when multiple itera-tions are conducted in parallel within PMBC, the inference speed remains considerably faster than the traditional sequential iteration method that doesn't use PMBC. These findings further highlight the efficiency of our proposed PMBC-based training approach, demonstrating that it significantly accelerates the model's inference while maintaining robust performance.

In our method, we set the default number of iterations of PMBC as 3, which can achieve a more balanced performance between the high-efficiency inference and stable accuracy.