
Flash Invariant Point Attention

Andrew Liu, Axel Elaldi, Nicholas T Franklin, Nathan Russell
Gurinder S Atwal, Yih-En Andrew Ban, Olivia Viessmann

Flagship Pioneering
Cambridge, MA, United States

[anliu,aelaldi,nfranklin,nrussell,matwal,aban,oviessmann]@flagshippioneering.com

Abstract

Invariant Point Attention (IPA) is a key algorithm for geometry-aware modeling in structural biology, central to many protein and RNA models. However, its quadratic complexity limits the input sequence length. We introduce FlashIPA, a factorized reformulation of IPA that leverages hardware-efficient FlashAttention to achieve linear scaling in GPU memory and wall-clock time with sequence length. FlashIPA matches or exceeds standard IPA performance while substantially reducing computational costs. FlashIPA extends training to previously unattainable lengths, and we demonstrate this by re-training generative models without length restrictions and generating structures of thousands of residues. FlashIPA is available at https://github.com/flagshippioneering/flash_ipa.

1 Introduction

Invariant Point Attention, IPA, is a geometry-aware attention operation that has been the workhorse of generative structural design models for proteins and RNA, initially popularized by AlphaFold2 [1] and AlphaFold-Multimer [2], and subsequently widely adopted across structural biology modeling. Amongst them are structure prediction models like OpenFold and ESMFold for proteins [3, 4], RhoFold for RNA [5], generative protein backbone models such as FrameDiff [6], FrameFlow [7, 8], the FoldFlow family [9, 10], FrameDiPT [11], Proteus [12], FADiff [13], Genie [14], IgDiff [15], GAFL [16], P2DFlow [17], RNA generative models such as RNA-FrameFlow [18], and scoring models like lociPARSE [19]. A list of models that rely on IPA for their structure modeling is provided in Appendix A.1. The advantage of IPA is its roto-translational ($SE(3)$) invariant representation of the molecular structure, enforcing the idea that rotations and translations of a molecule results in an equivalent structure prediction. This inductive geometric bias accelerates training and improves performance in limited data settings, as is the case for structurally resolved biomolecules. IPA’s quadratic scaling ($O(L^2)$) limits its scalability, rapidly exhausting GPU memory when modeling longer biomolecules. As of May 2025, 42% of structures within the PDB have more than 512 residues, and 33% with more than 756 and 23% with more than 1024 (see Appendix A.2). Most trainings across the literature reduce their data to chains and cropped structures below 512 residues. Despite this, they still commonly run on costly multi-GPU setups with trainings that span from days up to a month [5, 6].

Such engineering compromises, namely truncating biologically relevant lengths, limiting dataset sizes, and relying on expensive computational infrastructure, restrict the progress of the field. In this work, we propose a recasting of the original IPA algorithm to a simple attention form to leverage off-the-shelf I/O reduction methods like FlashAttention, which replace the quadratic $O(L^2)$ with an effective linear $O(L)$ scaling behavior. We show empirically that FlashIPA exceeds the validation performance of IPA in benchmarking models and datasets. We then demonstrate the memory and

compute time efficiency by retraining benchmarking models more efficiently and without length restrictions on the data.

We provide FlashIPA as an importable uv package at https://anonymous.4open.science/r/flash_ipa-07CE/ with an API similar to existing repositories using IPA to facilitate drop-in usage.

2 Preliminaries

Introduced by AlphaFold2 [1], IPA is a specialized attention mechanism designed to preserve 3D geometric relationships directly in transformer-based models for structural biology applications. IPA enforces invariance under 3D rotations and translations by utilizing a learnable local coordinate frame representation for each residue (point) within proteins and RNAs.

Formally, a function $f : \mathcal{X} \rightarrow \mathcal{Y}$ is *invariant* to transformations in the abstract group G if $f(g \cdot x) = f(x)$ for all $g \in G$. This differs from *equivariance* to G , where the transformations of the group commute with f , or $f(g \cdot x) = g \cdot f(x)$ for all $g \in G$. Here, we are concerned with the transformations T (referred to as "frames") in the special Euclidean group $SE(3)$, a continuous group of rigid transformations that includes rotations and translations, but not reflections. Intuitively, the output of an $SE(3)$ -invariant function does not change under rotations and translations, which is desirable for biomolecular structures, that do not change function under those transformations.

2.1 Frame representations for proteins and RNA

Meaningful frames for biomolecules are commonly defined on the backbone structures. In proteins, each residue frame is typically determined by three of the four atoms from the backbone, namely the alpha carbon (C_α), nitrogen (N), and carbon (C), with C_α at the origin. The frame transformation T maps the standard positions of these backbone atoms into their actual positions in global coordinates: $[N, C, C_\alpha, O] = T \cdot [N', C', C'_\alpha, O']$. The frame has a rotation and translation component $T = (R, t)$. The rotation component of the transformation, R , is computed using Gram–Schmidt orthonormalization on vectors formed by $C_\alpha - N$ and $C_\alpha - C$ bonds, while the position of the oxygen (O) is captured by the torsion angle around the $C_\alpha - C$ bond.

For RNA, the backbone structure involves 13 atoms per nucleotide, leading to more complex flexibility. RNA-FrameFlow [18], define RNA frames using the atoms $C3'$, $C4'$, and $O4'$, identified as the least variable backbone positions. The rest of the RNA backbone geometry can then be parameterized efficiently through a set of eight torsion angles.

2.2 IPA algorithm

The IPA algorithm is a 1D sequence-to-sequence transformation. It transforms an input sequence representation s_i , local frames T_i , and pair representations z_{ij} (with sequence indices $i, j \in 1, \dots, L$) into an output sequence \tilde{s}_i . Each attention head, indexed by $h \in 1, \dots, H$, involves linear transformations of the input sequence to produce scalar queries, keys, values ($\mathbf{q}_i^h, \mathbf{k}_i^h, \mathbf{v}_i^h \in \mathbb{R}^c$) and Euclidean (geometric) counterparts ($\tilde{\mathbf{q}}_i^{hp}, \tilde{\mathbf{k}}_i^{hp}, \tilde{\mathbf{v}}_i^{hp} \in \mathbb{R}^3$). Additionally, the pair representation $\mathbf{z}_{ij} \in \mathbb{R}^{L \times L \times d_z}$ contributes to a bias term b_{ij}^h through a linear projection:

$$\text{Single rep: } \mathbf{q}_i^h, \tilde{\mathbf{q}}_i^{hp}, \mathbf{k}_i^h, \tilde{\mathbf{k}}_i^{hp}, \mathbf{v}_i^h, \tilde{\mathbf{v}}_i^{hp} \leftarrow \text{Linear}(\mathbf{s}_i) \quad (1)$$

$$\text{Pair rep. (bias term): } b_{ij}^h \leftarrow \text{Linear}(\mathbf{z}_{ij}) \quad (2)$$

$$\text{IPA update: } a_{ij}^h = \text{softmax}_j \left(\underbrace{w_L \left(\frac{1}{\sqrt{c}} \mathbf{q}_i^h \mathbf{k}_j^h + b_{ij}^h - \frac{\gamma^h w_C}{2} \sum_p \left\| T_i \circ \tilde{\mathbf{q}}_i^{hp} - T_j \circ \tilde{\mathbf{k}}_j^{hp} \right\|_2^2 \right)}_{O(L^2)} \right) \quad (3)$$

$$\text{Attention aggregation: } \tilde{\mathbf{o}}_i^h = \sum_j a_{ij}^h \mathbf{z}_{ij}, \quad \mathbf{o}_i^h = \sum_j a_{ij}^h \mathbf{v}_j^h, \quad \tilde{\mathbf{o}}_i^{hp} = T_i^{-1} \circ \sum_j a_{ij}^h \left(T_j \circ \tilde{\mathbf{v}}_j^{hp} \right) \quad (4)$$

$$\text{Concat, project, return: } \tilde{s}_i = \text{Linear} \left(\text{concat}_{h,p} \left(\tilde{\mathbf{o}}_i^h, \mathbf{o}_i^h, \tilde{\mathbf{o}}_i^{hp}, \left\| \tilde{\mathbf{o}}_i^{hp} \right\|_2 \right) \right) \quad (5)$$

The proof of SE(3) invariance of the IPA transformation can be found in the appendix of AlphaFold2 [1]. This form of IPA incurs $O(L^2)$ I/O and memory complexity due to the explicit materialization of the quadratic attention matrix in equation 3.

2.3 FlashAttention and I/O reduction

The quadratic scaling of the attention operation is a well-known challenge in deep learning. Operations are generally I/O-bound on the GPU, with performance primarily limited by data transfers between the GPU’s high-bandwidth memory (HBM) and its static random access memory (SRAM). I/O reduction is typically achieved through kernel fusion, which reduces memory traffic by combining multiple operations into a single CUDA kernel. FlashAttention [20, 21] achieves kernel fusion via an online and tiled computation of the softmax, which, instead of materializing the full quadratic $\mathbf{M} = \text{softmax}(\mathbf{q}\mathbf{k}^\top)$ matrix, performs an equivalent computation by accumulating partial contributions to the output $\mathbf{Y} = \mathbf{M}\mathbf{V}$ one tile at a time. This technique improves scalability and performance for transformer-based models. We provide the Flash-Attention pseudo-code in appendix A.3. Building on FlashAttention-1, FlashAttention-2 further reduced the number of non-matmul FLOPs, increased parallelism across thread blocks, and distributed work between warps to reduce communication through shared memory [21].

2.4 FlashIPA: combining geometry-awareness and efficiency

Our efficiency gains come from expressing the entire IPA update in a form that is amenable to FlashAttention (i.e. a factorized version of form $\mathbf{q}^\top \mathbf{k}$), which then computes the attention update without materializing any quadratic attention matrix. To do so, we rewrite the softmax argument in the IPA update (eq. (3)) as a single inner product. We also need to parameterize the pair representation (bias term eq. (2)) $\mathbf{z}_{ij} \in \mathbb{R}^{L \times L \times d_z}$ in a factorized form $\mathbf{z}_{ij} = \mathbf{z}_i^1 \mathbf{z}_j^2$, where $\mathbf{z}_i^1, \mathbf{z}_j^2 \in \mathbb{R}^{L \times r \times d_z}$. Here, r can be interpreted as the "rank" of the factorization, and is the dimension on which we perform the contraction. We then expand the sum of squared norms in the third term of eq. (3) and collect terms, resulting in an equivalent update rule, given in Algorithm 1. Note that factorizing \mathbf{z}_{ij} allows not only for combining the softmax terms, but also allows us to compute the pair representation part of the attention as $\tilde{\mathbf{o}}_i^h = \sum_j a_{ij} \mathbf{z}_{ij} = \mathbf{z}_i^1 \mathbf{z}_j^2 \left(\sum_j a_{ij} \mathbf{z}_j^2 \right)$, avoiding materializing any quadratic object. As a result, the attention components from eq. (1) are lifted to $\hat{\mathbf{q}}_i^h, \hat{\mathbf{k}}_j^h \in \mathbb{R}^{c+5N_{\text{query}}+rd_z}, \hat{\mathbf{v}}_i^h \in \mathbb{R}^{c+3N_{\text{value}}+rd_z}$. Regular attention is then computed on the lifted components.

With the new update in the form of regular attention, we then leverage FlashAttention to perform the computation. This leads to significant speedups in wall-clock time and since the online computation of FlashAttention avoids materializing the quadratic attention matrix, we only need to store the queries, keys, and values in the GPU’s HBM, incurring only $O(L)$ memory.

2.5 Factorizing the pair representation

General forms of pair representations \mathbf{z}_{ij} typically involve quadratic memory complexity and are not inherently decomposable. However, common implementations of IPA often compute these quadratic representations from components that intrinsically scale linearly, implying redundancy in the resulting quadratic tensor \mathbf{z}_{ij} (eq. (2)). This suggests that pair representations can be efficiently approximated through low-rank factorization in latent space. In particular, common pairwise tensors in structural biology modeling, such as distance matrices and contact maps, are either low-rank or sparse due to the smooth, local nature of physical interactions like electrostatics and steric constraints. This motivates a simplified factorization strategy for initializing and updating \mathbf{z}_{ij} . Instead of explicitly constructing and storing a full quadratic tensor of shape (B, L, L, d) , we represent each feature (e.g., positional encodings, distograms, diffusion time embeddings in generative models, a.s.o.) using two lower-dimensional tensors (pseudo-factors) each of shape (B, L, r, d) , obtained via linear projections. We show empirically that this parametrization, despite relaxing certain inductive biases, recovers downstream performance with substantially lower memory consumption.

Algorithm 1 FlashIPA with factorized pair representations

Given input sequence $\mathbf{s} \in \mathbb{R}^{L \times d_{in}}$, factorized pair representation $\mathbf{z}_{ij} = \mathbf{z}_i^1 \top \mathbf{z}_j^2$ (where $\mathbf{z}^1, \mathbf{z}^2 \in \mathbb{R}^{L \times r \times d_z}$ and $\mathbf{z} \in \mathbb{R}^{L \times L \times d_z}$), frames $T_i = (R_i, \mathbf{t}_i), \forall i, j \in \{1, \dots, L\}$

- 1: $\mathbf{q}_i^h, \bar{\mathbf{q}}_i^{hp}, \mathbf{k}_i^h, \bar{\mathbf{k}}_i^{hp}, \mathbf{v}_i^h, \bar{\mathbf{v}}_i^{hp} \leftarrow \text{Linear}(\mathbf{s}_i)$
 - 2: $\mathbf{b}_i^{h1} \leftarrow \text{Linear}(\mathbf{z}_i^1)$
 - 3: $\mathbf{b}_i^{h2} \leftarrow \text{Linear}(\mathbf{z}_i^2)$
 - 4: $\hat{\mathbf{q}}_i^h \leftarrow \text{concat} \left(\mathbf{q}_i^h, \left\{ T_i \circ \bar{\mathbf{q}}_i^{hp} \right\}_{p=1}^{N_{\text{Query}}}, \left\{ \left\| T_i \circ \bar{\mathbf{q}}_i^{hp} \right\|_2^2 \right\}_{p=1}^{N_{\text{Query}}}, \mathbf{1}_{N_{\text{Query}}}, \mathbf{b}_i^{h1} \right)$
 - 5: $\hat{\mathbf{k}}_i^h \leftarrow \text{concat} \left(\frac{w_L}{\sqrt{c}} \mathbf{k}_i^h, \left\{ \gamma^h w_L w_C T_i \circ \bar{\mathbf{k}}_i^{hp} \right\}_{p=1}^{N_{\text{Query}}}, \frac{-\gamma^h w_L w_C}{2} \mathbf{1}_{N_{\text{Query}}}, \left\{ \frac{-\gamma^h w_L w_C}{2} \left\| T_i \circ \bar{\mathbf{k}}_i^{hp} \right\|_2^2 \right\}_{p=1}^{N_{\text{Query}}}, \mathbf{b}_i^{h2} \right)$
 - 6: $\hat{\mathbf{v}}_i^h \leftarrow \text{concat} \left(\mathbf{v}_i^h, \left\{ T_i \circ \bar{\mathbf{v}}_i^{hp} \right\}_{p=1}^{N_{\text{Values}}}, \mathbf{z}_i^2 \right)$
 - 7: $\hat{\mathbf{o}}_i^h \leftarrow \text{FlashAttention}(\hat{\mathbf{Q}}^h, \hat{\mathbf{K}}^h, \hat{\mathbf{V}}^h)_i$
 - 8: $\mathbf{o}_i^h, \bar{\mathbf{o}}_i^{hp}, \bar{\mathbf{o}}_i^h \leftarrow \text{split}(\hat{\mathbf{o}}_i^h)$
 - 9: $\tilde{\mathbf{o}}_i^h \leftarrow \mathbf{z}_i^1 \top \bar{\mathbf{o}}_i^h$
 - 10: $\tilde{\mathbf{s}}_i \leftarrow \text{Linear} \left(\text{concat}_{h,p} \left(\bar{\mathbf{o}}_i^h, \mathbf{o}_i^h, \bar{\mathbf{o}}_i^{hp}, \left\| \bar{\mathbf{o}}_i^{hp} \right\|_2 \right) \right)$
 - 11: Return $\tilde{\mathbf{s}}$
-

In many structure models \mathbf{z}_{ij} is updated multiple times by a non-linear EdgeTransition module that is not part of IPA. Because of that each factor is usually highly non-linear with respect to the input, despite the low-dimensional contraction. Thus, we still retain substantial representation power with this factorization. The benefits of highly nonlinear low-dimensional factors is well motivated in the literature: Instead of large head dimensions and linear factors, i.e. $q, k = \text{Linear}(X)$, as is the case for transformers, many state space models, such as Mamba [22] use factors that are non-linear transformations of the input (e.g. $q, k = \text{Swish}(\text{Conv1d}(X))$), and much smaller contraction dimensions (often $r = 16$).

We also note that most implementations of IPA rely on pair representations \mathbf{z}_{ij} that depend on distograms computed from pairwise distances. To simultaneously ensure invariance and avoid computing activations on the full distogram, we only keep the distances of the k nearest neighbors (B, L, k, n_{bins}) instead of (B, L, L, n_{bins}) . In practice, we compute these using a full $L \times L$ distance matrix during pre-processing, which does not impact the runtime complexity of FlashIPA. We keep track of neighbor identities by applying positional encodings on their indices. Emphasizing nearest neighbor distances is also motivated by the fact that local geometry (e.g. chemical bonds) is more important than global geometry when it comes to validity of biomolecular structures. We surmise that in future work it may be beneficial to compress all $\frac{L(L-1)}{2}$ distances into factors without materializing the pairwise matrix, in a manner similar to FlashAttention.

3 Experiments

3.1 FlashIPA is SE(3) invariant and scales linearly in memory and wall-clock time

We tested the SE(3)-invariance by applying random roto-translations to input Gaussian point clouds and measuring the output deviation after applying a single layer of the original IPA and FlashIPA. Original IPA output deviation was $< 10^{-6}$ and FlashIPA $< 10^{-3}$. We evaluate the scaling behavior in GPU memory and wall-clock time by passing single-sample batches of increasing lengths L through original and FlashIPA, see Fig. 1. We use a polynomial fit (green and red dotted lines), and find an approximate GPU memory scaling for FlashIPA of $y[\text{MB}] = -7 \cdot 10^{-12} \cdot L^2 + 7.5 \cdot 10^{-2} \cdot L$ versus original IPA $y[\text{MB}] = 2.4 \times 10^{-3} \cdot L^2 + 1.4 \cdot 10^{-2} \cdot L$. The computational complexity of attention still remains $O(L^2)$ due to the softmax operation. FlashAttention reduces the memory complexity to $O(L)$. Practically I/O, rather than computation, often dominates runtime on GPU hardware, and this is reflected in the observed linear wall-clock scaling in Fig. 2A.

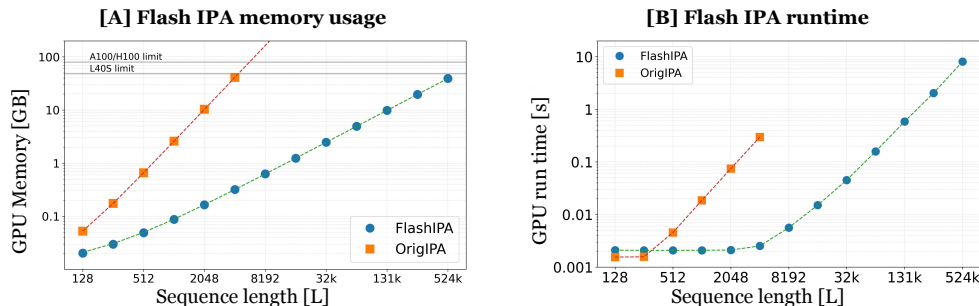


Figure 1: Scaling as a function of input sequence length on a single-sample batch forward pass. [A] GPU memory usage in GB. Original IPA scaled approximately quadratically with sequence length ($y \text{ [MB]} = 2.4 \times 10^{-3} \cdot L^2 + 1.4 \cdot 10^{-2} \cdot L$), FlashIPA follows a linear trend ($y \text{ [MB]} = -7 \cdot 10^{-12} \cdot L^2 + 7.5 \cdot 10^{-2} \cdot L$). [B] Wall-clock time in seconds.

3.2 Integration test with external repositories

We selected two recent model approaches where code and data were available for retraining of the original model and allowed for FlashIPA insertion. We chose FoldFlow [10] for proteins and RNA-FrameFlow [18] for RNAs, both are flow-matching generative backbone models. All experiments were run on L40S GPU instances with 48 GB HBM memory. FlashIPA hyperparameters were matched to the IPA parameters chosen by the original authors (embedding sizes, hidden dimensions, number of heads, etc.). For the factorization of the pair representation we tested rank $\in \{1, 2\}$, and found rank 2 and pair-wise distograms with $k = 20$ to match and partially surpass the loss convergence of original IPA.

3.3 FlashIPA improves performance and extends to larger proteins

We retrained the FoldFlow Base model in its original form with IPA and with FlashIPA. We reran the PDB data pre-processing pipeline by the authors, which yielded a total of 40,492 single-chain protein monomers for training. The original training used a maximum length cut-off of 512 residues, which results in a 10% reduction of the training data to 36,600 structures. We match the training strategy of the original authors and train on this reduced dataset on 4 GPUs with DDP. We kept model and train parameters identical to the published values of the authors, however for the FlashIPA version we had to make two adjustments: The original repository runs 4 blocks of IPA with a hidden dimension of 256. FlashAttention becomes incompatible with DDP at that dimension, so we reduced FlashIPA hidden dimension to 128 and used 5 blocks (instead of 4) to match model parameter sizes (17.4M versus 17.1M, theirs versus ours). To keep memory consumption at an efficient constant level, the original implementation heuristically chose an effective batch size according to the quadratic rule $\text{eff}_{bs} = \max \left[\text{round} \left(\frac{500.000 \times n_{GPUs}}{N^2} \right), 1 \right]$, which kept GPU memory at approximately 90% throughout training. For FlashIPA we were able to achieve a linear effective batch size $\text{eff}_{bs} = \max \left[\text{round} \left(\frac{20.000 \times n_{GPUs}}{N} \right), 1 \right]$ that resulted in similar memory consumption. For example IPA only has a batch size of 1 for a protein of length 512, whereas FlashIPA can batch together 39 samples. We found that FlashIPA converged slightly faster than IPA as expected with larger batch size, but we also compared partial loss terms, and found that particularly the number of steric clashes reduced faster for FlashIPA (we provide loss curves in the Appendix A.4). This suggests that our local k-nearest neighbour distogram emphasis is more efficient at capturing local geometry. To demonstrate the immediate enablement of linear scaling we ran one additional training with FlashIPA on the entire monomer dataset without length restrictions. The largest chain in the dataset is 8.8k residues. We follow the original authors' validation test that generates structures at varying lengths, subsequently inverse folds with Protein-MPNN [23] and forward-folds with ESMFold [4]. Generated and forward-folded structures are then aligned via the Kabsch algorithm and the resulting self-consistency RMSD (sc-RMDS) is a proxy for generation fidelity. We perform this test for all three models: original IPA and FlashIPA trained up to length 512, and FlashIPA on all data. Similar to the original paper we sampled 50 structures per length from 100 to 500, and sample 8 inverse folded sequences to forward fold per structure. All models are evaluated on checkpoint number 200,000. Fig. 2 A) shows that the sc-RMSD is lower for FlashIPA compared to the original IPA

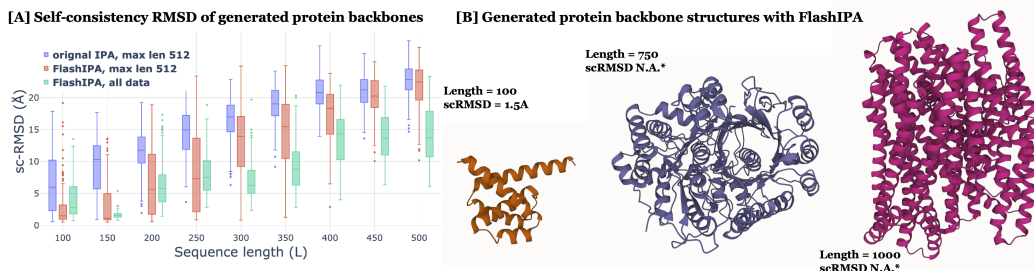


Figure 2: FoldFlow self-consistency validation after 200k optimization steps. A) The sc-RMSD of the FlashIPA (red, green) models is consistent or better than the original IPA model (blue). Extending training to larger structures with FlashIPA further improves sc-RMSD. B) Three exemplar generated backbone structures with FoldFlow FlashIPA. *ESMFold gets out of memory for lengths beyond 500+ residues and sc-RMSD could not be assessed.

trained on the same length-restricted data. Sc-RMSD is further improved when FlashIPA training is extended to the full dataset, as expected. We emphasize that FlashIPA is not designed to improve generation quality per se, but to enable training on larger and more complex structures. We suspect that the observed sc-RMSD values reflect accumulated model errors across three stages of validation: FoldFlow generation, Protein-MPNN inverse folding, and ESMFold forward folding. We could not assess sc-RMSF for larger structures, as the ESMFold also depends on IPA and ran out of memory. Overall we conclude that FlashIPA is more performant and more efficient than IPA. Fig. 2 B) shows example structures generated with the full data trained FlashIPA version.

3.4 FlashIPA trains more efficiently and extends to larger RNAs

We retrained the RNA-FrameFlow model using the code provided by the original authors. We ingested the same BGSU version 3.382 of the RNASolo2 dataset [24], comprising a total of 14,995 structures (see Appendix A.5). First we re-trained RNA-FrameFlow with original and FlashIPA on all structures within 40 to 150 residue (6,030 structures total), as proposed by the authors. We kept all hyperparameters consistent with the original model. We matched the authors’ training setup and models were trained on 4 GPUs with DDP and an effective batch size of 4×28 . 4 GPUs are necessary for effective training with IPA, but with FlashIPA it becomes feasible to run a comparable training on a single GPU instance. To demonstrate the cost efficiency we also performed an additional FlashIPA training run on the same data with a batch size of 512 on single GPU. All models were trained for a fixed compute time of 20 hours, which resulted in approximately 156k iterations with original IPA, roughly 230k with FlashIPA on 4GPUs and 194k with FlashIPA on a single GPU. During training we found models to converge comparably with overlapping loss curves, see Appendix A.6. For an apples-to-apples comparison we validated the models using identical validity, novelty, and diversity metrics as defined in the original paper, which uses gRNAde [25] for inverse-folding of the generated structure and RhoFold [26] for forward-folding. For validity, we report the average self-consistency template modeling score (sc-TM). To do so, we generated 50 structures per length ranging from 40 to 150 residues, in increments of 10. The results are presented in Table 1 and Fig. 4. We observe comparable scores between all models. In particular, the single-GPU training matches performance at a quarter of the compute cost than the original.

To highlight the advantages of using FlashIPA in the RNA-FrameFlow model, we analyzed the GPU runtime required to generate RNA backbones based on RNA sequence length and batch size. The results are in Fig. 3. We observed improvements with FlashIPA compared to the original IPA model, especially for longer sequences and larger batch sizes. For example it takes 30 times longer to generate an RNA sequence of length 2048 with the original IPA than with FlashIPA.

We also re-trained RNA-FrameFlow with FlashIPA on the full RNASolo dataset without maximum length restrictions (the longest structures being 4,417 nucleotides) and filtering out structures shorter than 40 nucleotides. We trained the model on 4 GPUs for 48 hours, with batch size of 4×28 , resulting in approximately 310k iterations. In Fig. 4, we present samples of large generated RNA structures with lengths of 2000 and 4000 nucleotides. In comparison, the original RNA-FrameFlow can not be

Model	Validity (\uparrow) (scTM \pm std)	Diversity (\uparrow) (qTM cluster)	Novelty (\downarrow) (pdbTM \pm std)	Checkpoint (#steps)	Cost (# GPU hours)
RNA-FrameFlow	0.42 ± 0.21	0.15	0.81 ± 0.10	156k	80
. + FlashIPA (ours)	0.38 ± 0.20	0.14	0.82 ± 0.10	230k	80
. + FlashIPA single GPU ** (ours)	0.41 ± 0.21	0.08	0.77 ± 0.09	194k	20
. + FlashIPA ** + All data (ours)	0.36 ± 0.14	0.14	0.74 ± 0.08	310k	192

Table 1: Average validity, diversity, and novelty scores for 600 generated RNAs of length ≤ 150 with RNA-FrameFlow, with and without FlashIPA. FlashIPA provides competitive results at a fraction of the cost of an original IPA layer. ** indicates that batch size has been increased to match GPU memory capacities.

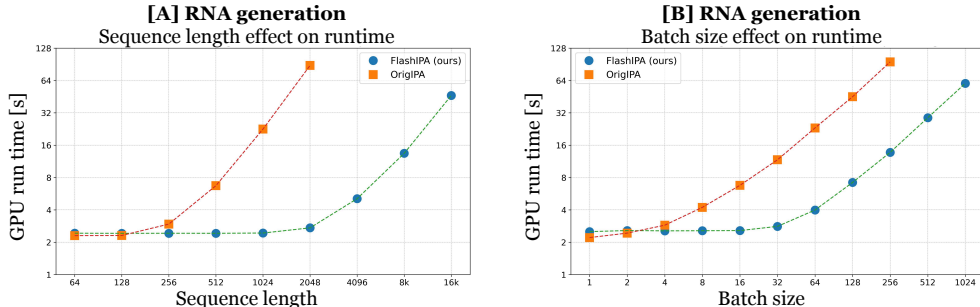


Figure 3: Scaling of FlashIPA versus IPA for RNA generation using RNA-FrameFlow model, with a number of diffusion timestep $N_T = 50$. [A] Impact of the generated sequence length on the generation runtime, using a batch size of 1. [B] Impact of the generated batch size on the generation runtime, for generated sequence of length 128.

trained or generate such long sequences. The inverse-forward-folding consistency test cannot be run for structures of thousands of residues as the validation models (gRNAde [25] for inverse-folding and RhoFold [26] for forward-folding) run out of memory at such lengths. However, we qualitatively observe that the generated structures resemble RNA, we don't necessary suspect those to be valid, as the amount of training data at such length is still comparably small. We note that sc-TM scores vary across sequence lengths, with some fluctuations (e.g., at 120 residues). This pattern is consistent with the original RNA-FrameFlow paper, which attributes it to a length imbalance in the training distribution and biases introduced by the RhoFold structure predictor used during evaluation (see [18] Appendix A.3).

4 Discussion

FlashIPA provides memory and wall-clock efficient $SE(3)$ invariant training on biomolecules and allows for training on structures of thousands of residues. This approach opens up the possibility of multi-chain complex modeling in situations constrained by the original IPA module. Our work permit usage of IPA in contexts where it may have been ruled out due to its scaling limitations, including representations beyond polymeric backbone that include more atomic detail.

Efficiency and scalability to long contexts has often been a neglected aspect of geometric deep learning research and many modules have quadratic to cubic complexity. An example is triangular attention used in popular models like AF-3[27], Chai-1[28] and Boltz-1 [29]. Recently, TriFast [30] was released, which uses fused triangular attention kernels to reduce I/O complexity from cubic to quadratic. We assume that a factorized version might even achieve a linear scaling. Most efficient, sub-quadratic models tend not to incorporate geometric inductive biases. Our work can be seen as a first step at bridging this divide, enabling models that respect invariances, are cost-effective, and can scale to larger or more fine-grained biomolecular systems.

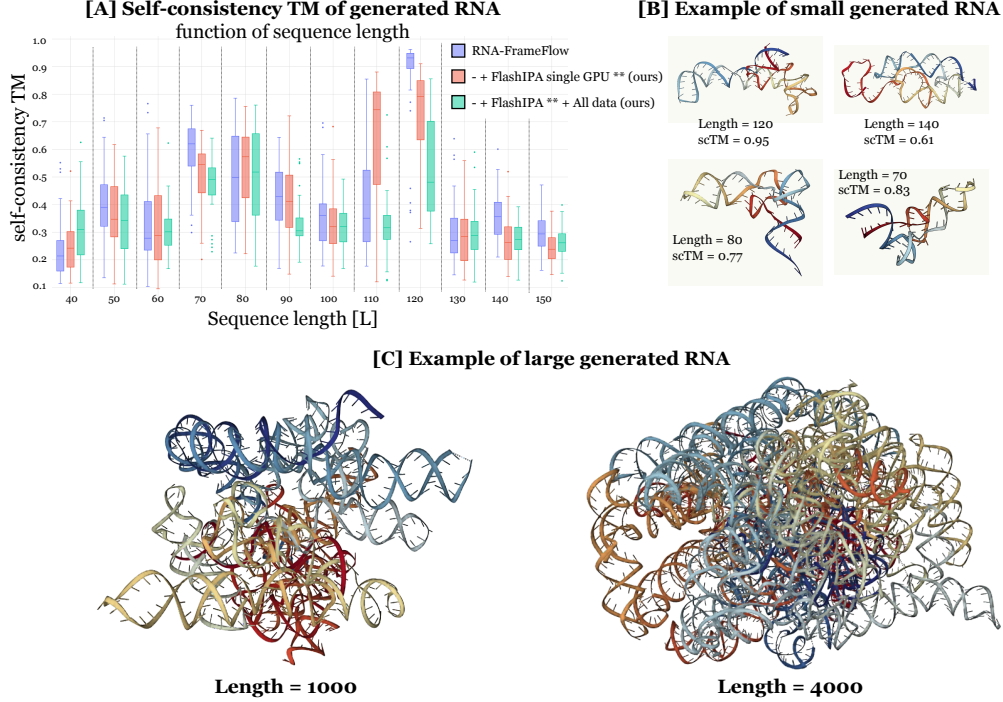


Figure 4: RNA-FrameFlow generated RNAs results. **[A]** Comparison of the scTM score depending on the generated RNA sequence length and the IPA module used. The RNA-FrameFlow and RNA-FrameFlow + FlashIPA models are both trained on only short sequences ≤ 150 , while the All data model has been trained without maximum sequence length limit. We don’t observe a significant difference between the different trained RNA-FrameFlow models. **[B]** Example of short generated RNA structures with our FlashIPA RNA-FrameFlow model trained on short sequences. **[C]** Example of long generated RNA structured with our FlashIPA RNA-FrameFlow model trained on the full dataset.

4.1 Limitations

FlashIPA relies on factorization of the pair representation, which ultimately is an approximation. Here we did not find a decrease in performance, but this may not be guaranteed for more general applications to other forms of more dense pair representations. Exploring alternative schemes specialized for certain pair representations (e.g. cross-concatenations, distograms, pair index differences) may further improve downstream performance.

Despite substantial memory and I/O savings, current FlashAttention implementations (e.g., FlashAttention2) impose a maximum head dimension of 256, due to the kernel’s internal design, which expects stacked query/key/value vectors. Since our method relies on augmenting the attention components, this means that we must have $\max(c + 5N_{query} + rd_z, c + 3N_{value} + rd_z) \leq 256$. Here, we achieved competitive results using head dimensions around 200, so this was not a practical limitation. However, the Triton implementation of FlashAttention2 based on AMD CDNA (MI200, MI300) and RDNA GPUs allows for extending the head dimension to arbitrary size. We expect these improvements to apply more broadly to a wider class of GPU architectures in the months to come. Unlocking larger head dimension would allow us to increase our factorization rank, thereby better approximating dense pair representations and closing potential gaps against quadratic IPA.

Furthermore, we note that despite achieving $O(L)$ in I/O and memory, the underlying compute cost is still $O(L^2)$ due to the softmax. Removing the softmax and using linear attention variants such as Mamba would allow us to achieve $O(L)$ cost in both compute and memory, and is thus a promising avenue of future work.

References

- [1] John Jumper, Richard Evans, Alexander Pritzel, Tim Green, Michael Figurnov, Olaf Ronneberger, Kathryn Tunyasuvunakool, Russ Bates, Augustin Žídek, Anna Potapenko, et al. Highly accurate protein structure prediction with alphafold. *nature*, 596(7873):583–589, 2021.
- [2] Richard Evans, Michael O’Neill, Alexander Pritzel, Natasha Antropova, Andrew Senior, Tim Green, Augustin Žídek, Russ Bates, Sam Blackwell, Jason Yim, et al. Protein complex prediction with alphafold-multimer. *bioRxiv*, pages 2021–10, 2021.
- [3] Gustaf Ahdritz, Nazim Bouatta, Christina Floristean, Sachin Kadyan, Qinghui Xia, William Gerecke, Timothy J O’Donnell, Daniel Berenberg, Ian Fisk, Niccolò Zanichelli, et al. Openfold: Retraining alphafold2 yields new insights into its learning mechanisms and capacity for generalization. *Nature Methods*, 21(8):1514–1524, 2024.
- [4] Zeming Lin, Halil Akin, Roshan Rao, Brian Hie, Zhongkai Zhu, Wenting Lu, Allan dos Santos Costa, Maryam Fazel-Zarandi, Tom Sercu, Sal Candido, et al. Language models of protein sequences at the scale of evolution enable accurate structure prediction. *BioRxiv*, 2022:500902, 2022.
- [5] Tao Shen, Zhihang Hu, Siqi Sun, Di Liu, Felix Wong, Jiuming Wang, Jiayang Chen, Yixuan Wang, Liang Hong, Jin Xiao, et al. Accurate rna 3d structure prediction using a language model-based deep learning approach. *Nature Methods*, pages 1–12, 2024.
- [6] Jason Yim, Brian L Trippe, Valentin De Bortoli, Emile Mathieu, Arnaud Doucet, Regina Barzilay, and Tommi Jaakkola. Se (3) diffusion model with application to protein backbone generation. *arXiv preprint arXiv:2302.02277*, 2023.
- [7] Jason Yim, Andrew Campbell, Emile Mathieu, Andrew Y. K. Foong, Michael Gastegger, Jose Jimenez-Luna, Sarah Lewis, Victor Garcia Satorras, Bastiaan S. Veeling, Frank Noe, Regina Barzilay, and Tommi Jaakkola. Improved motif-scaffolding with SE(3) flow matching. *Transactions on Machine Learning Research*, 2024.
- [8] Jason Yim, Andrew Campbell, Andrew YK Foong, Michael Gastegger, José Jiménez-Luna, Sarah Lewis, Victor Garcia Satorras, Bastiaan S Veeling, Regina Barzilay, Tommi Jaakkola, et al. Fast protein backbone generation with se (3) flow matching. *arXiv preprint arXiv:2310.05297*, 2023.
- [9] Guillaume Huguet, James Vuckovic, Kilian Fatras, Eric Thibodeau-Laufer, Pablo Lemos, Riashat Islam, Cheng-Hao Liu, Jarrid Rector-Brooks, Tara Akhound-Sadegh, Michael Bronstein, et al. Sequence-augmented se (3)-flow matching for conditional protein backbone generation. *Advances in neural information processing systems*, 2024.
- [10] Joey Bose, Tara Akhound-Sadegh, Guillaume Huguet, Kilian FATRAS, Jarrid Rector-Brooks, Cheng-Hao Liu, Andrei Cristian Nica, Maksym Korablyov, Michael M Bronstein, and Alexander Tong. Se (3)-stochastic flow matching for protein backbone generation. In *The Twelfth International Conference on Learning Representations*, 2024.
- [11] Cheng Zhang, Adam Leach, Thomas Makkink, Miguel Arbesú, Ibtissem Kadri, Daniel Luo, Liron Mizrahi, Sabrine Krichen, Maren Lang, Andrey Tovchigrechko, Nicolas Lopez Carranza, Uğur Şahin, Karim Beguir, Michael Rooney, and Yunguan Fu. Framedipt: Se(3) diffusion model for protein structure inpainting. *bioRxiv*, 2023.
- [12] Chentong Wang, Yannan Qu, Zhangzhi Peng, Yukai Wang, Hongli Zhu, Dachuan Chen, and Longxing Cao. Proteus: exploring protein structure generation for enhanced designability and efficiency. *bioRxiv*, pages 2024–02, 2024.
- [13] Ke Liu, Weian Mao, Shuaike Shen, Xiaoran Jiao, Zheng Sun, Hao Chen, and Chunhua Shen. Floating anchor diffusion model for multi-motif scaffolding. 2024.
- [14] Yeqing Lin and Mohammed AlQuraishi. Generating novel, designable, and diverse protein structures by equivariantly diffusing oriented residue clouds. *arXiv preprint arXiv:2301.12485*, 2023.

- [15] Daniel Cutting, Frédéric A Dreyer, David Errington, Constantin Schneider, and Charlotte M Deane. De novo antibody design with se (3) diffusion. *Journal of Computational Biology*, 2024.
- [16] Simon Wagner, Leif Seute, Vsevolod Viliuga, Nicolas Wolf, Frauke Gräter, and Jan Stühmer. Generating highly designable proteins with geometric algebra flow matching. In *Thirty-eighth Conference on Neural Information Processing Systems*, 2024.
- [17] Yaowei Jin, Qi Huang, Ziyang Song, Mingyue Zheng, Dan Teng, and Qian Shi. P2dflow: A protein ensemble generative model with se(3) flow matching. *Journal of Chemical Theory and Computation*, 2025.
- [18] Rishabh Anand, Chaitanya K. Joshi, Alex Morehead, Arian R. Jamasb, Charles Harris, Simon Mathis, Kieran Didi, Bryan Hooi, and Pietro Liò. Rna-frameflow: Flow matching for de novo 3d rna backbone design. *arXiv preprint arXiv:2406.13839*, 2024.
- [19] Sumit Tarafder and Debswapna Bhattacharya. lociparse: a locality-aware invariant point attention model for scoring rna 3d structures. *Journal of Chemical Information and Modeling*, 64(22):8655–8664, 2024.
- [20] Tri Dao, Dan Fu, Stefano Ermon, Atri Rudra, and Christopher Ré. Flashattention: Fast and memory-efficient exact attention with io-awareness. *Advances in neural information processing systems*, 35:16344–16359, 2022.
- [21] Tri Dao. Flashattention-2: Faster attention with better parallelism and work partitioning. *International Conference on Learning Representations*, 2024.
- [22] Albert Gu and Tri Dao. Mamba: Linear-time sequence modeling with selective state spaces. *arXiv preprint arXiv:2312.00752*, 2023.
- [23] Justas Dauparas, Ivan Anishchenko, Nathaniel Bennett, Hua Bai, Robert J Ragotte, Lukas F Milles, Basile IM Wicky, Alexis Courbet, Rob J de Haas, Neville Bethel, et al. Robust deep learning-based protein sequence design using proteinmpnn. *Science*, 378(6615):49–56, 2022.
- [24] Bartosz Adamczyk, Maciej Antczak, and Marta Szachniuk. Rnasolo: a repository of cleaned pdb-derived rna 3d structures. *Bioinformatics*, 38(14):3668–3670, 2022.
- [25] Chaitanya K Joshi, Arian R Jamasb, Ramon Viñas, Charles Harris, Simon V Mathis, Alex Morehead, Rishabh Anand, and Pietro Liò. gRNade: Geometric deep learning for 3d rna inverse design. In *International Conference on Learning Representations (ICLR)*, 2025.
- [26] Tao Shen, Zhihang Hu, Zhangzhi Peng, Jiayang Chen, Peng Xiong, Liang Hong, Liangzhen Zheng, Yixuan Wang, Irwin King, Sheng Wang, et al. E2efold-3d: End-to-end deep learning method for accurate de novo rna 3d structure prediction. *arXiv preprint arXiv:2207.01586*, 2022.
- [27] Josh Abramson, Jonas Adler, Jack Dunger, Richard Evans, Tim Green, Alexander Pritzel, Olaf Ronneberger, Lindsay Willmore, Andrew J Ballard, Joshua Bambrick, et al. Accurate structure prediction of biomolecular interactions with alphafold 3. *Nature*, 630(8016):493–500, 2024.
- [28] Chai Discovery team, Jacques Boitreaud, Jack Dent, Matthew McPartlon, Joshua Meier, Vinicius Reis, Alex Rogozhonikov, and Kevin Wu. Chai-1: Decoding the molecular interactions of life. *BioRxiv*, pages 2024–10, 2024.
- [29] Jeremy Wohlwend, Gabriele Corso, Saro Passaro, Mateo Reveiz, Ken Leidal, Wojtek Swiderski, Tally Portnoi, Itamar Chinn, Jacob Silterra, Tommi Jaakkola, et al. Boltz-1: Democratizing biomolecular interaction modeling. *bioRxiv*, pages 2024–11, 2024.
- [30] Liam Atkinson. Trifast. <https://github.com/latkins/trifast>, 2025.
- [31] Andrew Campbell, Jason Yim, Regina Barzilay, Tom Rainforth, and Tommi Jaakkola. Generative flows on discrete state-spaces: Enabling multimodal flows with applications to protein co-design. *arXiv preprint arXiv:2402.04997*, 2024.
- [32] Aleksa Gordic. Eli5: Flashattention. <https://gordicaleksa.medium.com/eli5-flash-attention-5c44017022ad>, 2023.

A Appendix

A.1 List of biomolecular design models with IPA structure modules

Table 2: A list of models that are based on IPA structure modeling. Models were identified via GitHub advanced search, searching for commonalities in code implementations.

	Model name	Modeling approach	Training data	Train length cut-offs
Protein	OpenFold [3]	protein structure prediction	130k single protein chains (PDB)	<256, fine-tuning <384,<512,<5120
	FrameDiff [6]	diffusion for backbone generation	20k monomers (PDB)	60-512
	FrameDiPT [11]	diffusion for backbone inpainting	32k monomers (PDB)	60-512
	Proteus [12]	diffusion for backbone generation	51k single chains from oligo- and monomers (PDB)	60-512
	FADiff [13]	diffusion for multi-motif backbone generation	Identical to FrameDiff	60-512
	Genie [14]	diffusion for backbone generation	26k monomers (PDB)	60-512
	IgDiff [15]	FrameDiff finetuning for antibody backbone generation	Synthetic antibody structures from folding	Not reported
	FrameFlow [8, 7]	flow-matching for backbone generation and motif scaffolding	Identical to FrameDiff	60-512
	FoldFlow [10, 9]	Stochastic flow matching for backbone generation	22.2k monomers (PDB)	60-512
	GAFL [16]	Geometric algebra flow matching for backbone generation	Identical to FrameDiff	60-512
	Multiflow [31]	Discrete and continuous flow matching for joint sequence-backbone generation	18.7k monomers (PDB)	60-384
	P2DFlow [17]	flow matching for backbone ensemble generation	100 MD simulation trajectories from ATLAS	Not reported
RNA	Rho-Fold [5]	RNA structure prediction	5.5k chains (PDB)	<1024
	RNA-FrameFlow [18]	flow matching for backbone generation	5.3k original structures + 1.1k cropped structures (PDB)	40-150
	lociPARSE [19]	Locality-aware IPA for structure scoring	52k synthetic structures from 1.4k sequence targets	<200

A.2 Residue counts of macromolecules in the PDB

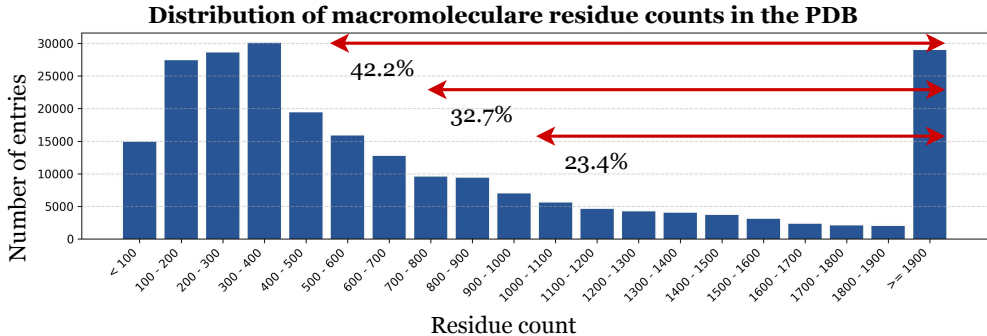


Figure 5: Distribution of protein residue counts of the proteins resolved in the PDB (Figure reproduced and adapted from the data at RCSB PDB Statistics: Sequence length distribution; <https://www.rcsb.org/stats/distribution-residue-count>; accessed 6 May 2025).

A.3 FlashAttention Algorithm

Pseudo-code for the FlashAttention kernel, an online tiled computation of the softmax, which, instead of materializing the full quadratic $M = \text{softmax}(\mathbf{qk}^T)$ matrix, performs an equivalent computation by accumulating partial contributions to the output $\mathbf{Y} = \mathbf{MV}$ one tile at a time [20, 21]. A good introduction with line-by-line explanations can also be found here [32].

Algorithm 2 Flash Attention-2 via online softmax (Dao 2024)

Given $\mathbf{Q}, \mathbf{K}, \mathbf{V} \in \mathbb{R}^{L \times d}$, T_c , T_r row blocks of size B_r , T_c column blocks of size B_c .

```
1: for  $i = 1$  to  $T_r$  do
2:   Load  $\mathbf{Q}_i$  from HBM to SRAM
3:    $\mathbf{O}_0 = \mathbf{0}_{B_r \times d}$ ,  $l_0 = \mathbf{0}_{B_r}$ ,  $m_0 = -\infty_{B_r}$ 
4:   for  $j = 1$  to  $T_c$  do
5:     Load  $\mathbf{K}_j, \mathbf{V}_j$  from HBM to SRAM.
6:     On chip, compute  $\mathbf{S}_i^{(j)} = \mathbf{Q}_i \mathbf{K}_j^T$ 
7:     On chip, compute  $m_i^{(j)} = \max(m_i^{(j-1)}, \text{rowmax}(\mathbf{S}_i^{(j)}))$ ,  $\tilde{\mathbf{P}}_i^{(j)} = \exp(\mathbf{S}_i^{(j)} - m_i^{(j)})$ ,
        $l_i^{(j)} = e^{m_i^{(j-1)} - m_i^{(j)}} l_i^{(j-1)} + \text{rowsum}(\tilde{\mathbf{P}}_i^{(j)})$ 
8:     On chip, compute  $\mathbf{O}_i^{(j)} = \text{diag}(e^{m_i^{(j-1)} - m_i^{(j)}})^{-1} \mathbf{O}_i^{(j-1)} + \tilde{\mathbf{P}}_i^{(j)} \mathbf{V}_j$ 
9:   end for
10:  On chip, compute  $\mathbf{O}_i = \text{diag}(l_i^{(T_c)})^{-1} \mathbf{O}_i^{(T_c)}$ 
11:  Write  $\mathbf{O}_i$  to HBM as  $i$ th block of  $\mathbf{O}$ 
12: end for
13: Return  $\mathbf{O}$ 
```

A.4 FoldFlow training convergence

We provide loss curves for the training of the FoldFlow base model with original and FlashIPA below. FlashIPA converged slightly faster for the same number of optimization steps, which is expected from the bigger effective batch size, but we also found in particular local loss terms, such as the steric clash loss to decrease noticeably more efficient.

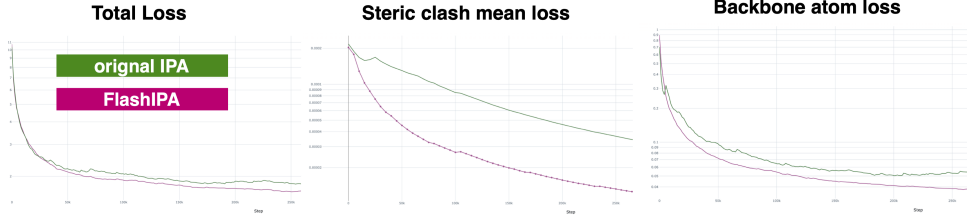


Figure 6: Loss behaviour for FoldFlow model training.

A.5 Nucleotide residue counts of RNA structures in the RNASolo2 dataset.

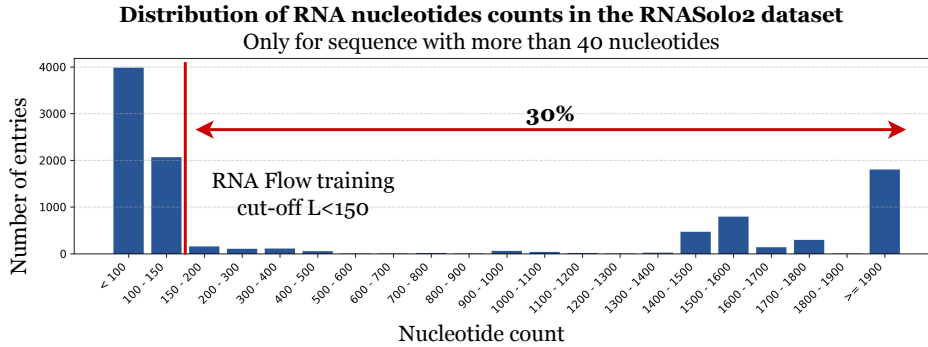


Figure 7: Distribution of nucleotide residue counts of the RNA in the RNASolo2 dataset, filtering out the short structures of length < 40 nucleotides. The training cut-off of RNA Flow discards all sequences of length > 150 , accounting for 30% of the dataset.

A.6 RNA-FrameFlow training convergence

We provide loss curves for the training of the RNA-FrameFlow base model with original and FlashIPA below. At similar hyperparameters and hardware, RNA-FrameFlow with and without FlashIPA behaves similarly.

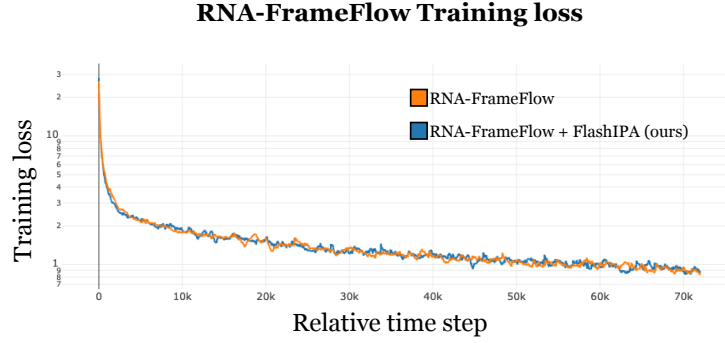


Figure 8: Loss behaviour for RNA-FrameFlow model training.

NeurIPS Paper Checklist

1. Claims

Question: Do the main claims made in the abstract and introduction accurately reflect the paper’s contributions and scope?

Answer: [\[Yes\]](#)

Justification: The abstract and introduction summarize the scaling laws and downstream performance that we demonstrate in the main text. We provide experiments and benchmarks to validate the claims. We perform invariance tests and run a scaling study to demonstrate our claim of linear scaling with FlashIPA. We apply the proposed method to popular models in generative biomolecular design, namely a protein and an RNA backbone generative model, to demonstrate the impact.

2. Limitations

Question: Does the paper discuss the limitations of the work performed by the authors?

Answer: [\[Yes\]](#)

Justification: We clearly state any assumptions, for example we clearly state that the factorization is approximate, we provide experiments on the downstream performance to assess if there are any effects of the approximations made. We include a comprehensive Limitations section where we address limitations of FlashIPA, including restrictions from FlashAttention and opportunities for improvement.

3. Theory assumptions and proofs

Question: For each theoretical result, does the paper provide the full set of assumptions and a complete (and correct) proof?

Answer: [\[Yes\]](#)

Justification: We provide a thorough walk-through our proposed algorithm and state all assumptions. The proof of invariance is referred to from the AlphaFold2 paper, which provides a full proof in their appendix.

4. Experimental result reproducibility

Question: Does the paper fully disclose all the information needed to reproduce the main experimental results of the paper to the extent that it affects the main claims and/or conclusions of the paper (regardless of whether the code and data are provided or not)?

Answer: [\[Yes\]](#)

Justification: We describe our experiments, including hyperparameters and hardware, as well as the data used for training. We provide the FlashIPA code via an anonymized github link. The repository provides a test script as well.

5. Open access to data and code

Question: Does the paper provide open access to the data and code, with sufficient instructions to faithfully reproduce the main experimental results, as described in supplemental material?

Answer: [\[Yes\]](#)

Justification: We provide an anonymised link to a github repository with the FlashIPA code and a test script. The data and code we used for re-training is referenced, as it is adopted from other author’s repositories. The data we used for training is provided by other sources, which we reference as well.

6. Experimental setting/details

Question: Does the paper specify all the training and test details (e.g., data splits, hyperparameters, how they were chosen, type of optimizer, etc.) necessary to understand the results?

Answer: [\[Yes\]](#)

Justification: Training hyperparameters and hardware are described in the experiments section. In many cases we adopted hyperparameter choices from external authors’ models, and reference their publication.

7. **Experiment statistical significance**

Question: Does the paper report error bars suitably and correctly defined or other appropriate information about the statistical significance of the experiments?

Answer: [Yes]

Justification: We compare invariance errors between IPA and FlashIPA. For all other results, there are no statistical error bars as it does not apply to the nature of the measure.

8. **Experiments compute resources**

Question: For each experiment, does the paper provide sufficient information on the computer resources (type of compute workers, memory, time of execution) needed to reproduce the experiments?

Answer: [Yes]

Justification: We state the instance type and GPU specifications and not how many GPUs were used for how many hours/days.

9. **Code of ethics**

Question: Does the research conducted in the paper conform, in every respect, with the NeurIPS Code of Ethics <https://neurips.cc/public/EthicsGuidelines>?

Answer: [Yes]

Justification: No human subjects were used in the study. Training was performed on publicly available academic datasets. The entire manuscript is anonymised, with no link or reference to the authors. The github link is anonymised too.

10. **Broader impacts**

Question: Does the paper discuss both potential positive societal impacts and negative societal impacts of the work performed?

Answer: [NA] .

There are no anticipated negative or positive societal impacts.

11. **Safeguards**

Question: Does the paper describe safeguards that have been put in place for responsible release of data or models that have a high risk for misuse (e.g., pretrained language models, image generators, or scraped datasets)?

Answer: [NA] .

There is no risk of misuse to our knowledge.

12. **Licenses for existing assets**

Question: Are the creators or original owners of assets (e.g., code, data, models), used in the paper, properly credited and are the license and terms of use explicitly mentioned and properly respected?

Answer: [Yes]

Justification: All references are cited throughout the text. We did not use any work with restricted licenses.

13. **New assets**

Question: Are new assets introduced in the paper well documented and is the documentation provided alongside the assets?

Answer: [NA] .

No new assest are introduced.

14. **Crowdsourcing and research with human subjects**

Question: For crowdsourcing experiments and research with human subjects, does the paper include the full text of instructions given to participants and screenshots, if applicable, as well as details about compensation (if any)?

Answer: [NA] .

No crowdsourcing with human subjects was used.

15. Institutional review board (IRB) approvals or equivalent for research with human subjects

Question: Does the paper describe potential risks incurred by study participants, whether such risks were disclosed to the subjects, and whether Institutional Review Board (IRB) approvals (or an equivalent approval/review based on the requirements of your country or institution) were obtained?

Answer:[NA] .

No human subjects are involved in this research.

16. Declaration of LLM usage

Question: Does the paper describe the usage of LLMs if it is an important, original, or non-standard component of the core methods in this research? Note that if the LLM is used only for writing, editing, or formatting purposes and does not impact the core methodology, scientific rigorousness, or originality of the research, declaration is not required.

Answer: [NA]

Justification: LLMs only used for editing.