

Replay-free Sequential Fine-tuning of Medical VLMs

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Abstract

Catastrophic forgetting severely limits the adaptation of Vision Language Models (VLMs) for medical applications, where sequential learning on private data is often necessary. We propose that this issue stems from insufficient regularization and demonstrate that regularizing parameter updates during fine-tuning effectively mitigates forgetting without harming new task performance. To validate our method in a clinical context, we introduce **Medical-CL**, a new continual learning benchmark spanning pathology, cell microscopy, radiology, and surgery. Our streamlined, replay-free approach proves highly effective on this benchmark, offering a practical path toward building comprehensive, continually-learning medical VLMs and advancing the development of medical AI.

Keywords: Vision Language Models, Continual Learning, Catastrophic Forgetting.

Data and Code Availability We use publicly available datasets: MLLM-CL Benchmark, BSCCM, PitVis-2023, PathVQA, ROCOV2. We will make code and data publicly available.

Institutional Review Board (IRB) Our research does not require IRB approval.

1. Introduction

The emergence of Vision Language Models (VLMs) represents a significant milestone in artificial intelligence (Alayrac et al., 2022; Liu et al., 2023; Achiam et al., 2023). Building upon this foundational success, the field has increasingly focused on fine-tuning these powerful models for specialized medical applications, aiming to create robust medical VLMs.

However, the catastrophic forgetting phenomenon (Zhai et al., 2024; Shuttleworth et al., 2024) is an un-

avoidable issue for the development of medical VLM. This issue manifests as a severe degradation in a model’s performance on previously learned tasks after it has been fine-tuned for a new specialization (McCloskey and Cohen, 1989). Within the context of medical VLMs, fine-tuning on a specific dataset could inadvertently compromise its general reasoning abilities or diminish its acquired knowledge pertinent to other medical fields. Furthermore, given that a significant volume of medical data is private or sensitive and cannot be easily shared, sequential learning on different datasets is an unavoidable necessity for building comprehensive medical VLMs under current constraints.

Drawing inspiration from prior research into the loss landscapes of Large Language Models (Chen et al., 2025) and the significance of orthogonal subspaces during fine-tuning (Wang et al., 2023), we hypothesize that the forgetting phenomenon arises from insufficient regularization during the adaptation process. Based on reasonable assumptions derived from landscape and subspace theory, we present the rationale for the efficacy of *regularizing parameter updates*.

Our experimental results demonstrate that *regularization applied to parameter updates can effectively mitigate forgetting without compromising performance on the fine-tuning task*. This holds true across multiple sequential fine-tuning scenarios, including domain-continual learning (Zhao et al., 2025). Empirical evaluations reveal that our simple approach surpasses the performance of existing methods, particularly those that depend on a data replay buffer. This outcome highlights the strong, inherent capacity of Vision-Language Models for continual learning.

Subsequently, we extend our investigation to various medical domains, curating a novel medical continual learning benchmark, which we term **Medical-**

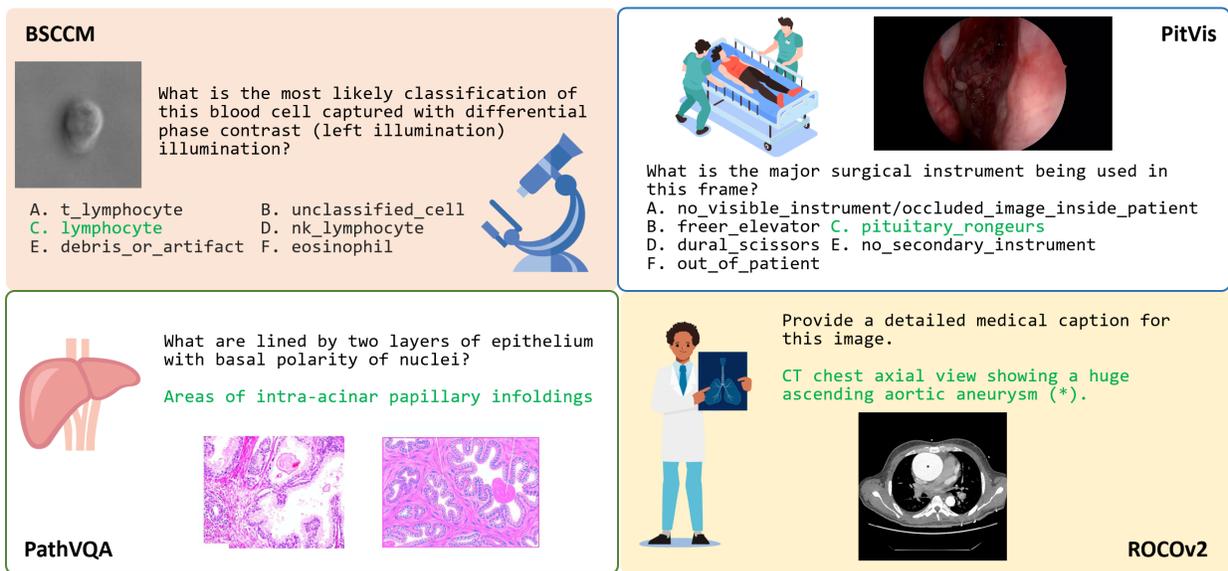


Figure 1: An Illustration of the Proposed Medical-CL Benchmark

75 **CL**, from a collection of public datasets (Pinkard
 76 et al., 2024; Das et al., 2025; He et al., 2020; Rückert
 77 et al., 2024). The **Medical-CL** benchmark in-
 78 corporates a variety of question formats, including
 79 multiple-choice, short-answer, and image caption-
 80 ing, to ensure a comprehensive evaluation. Adhering
 81 to a protocol similar to the **MLLM-CL** continual
 82 learning benchmark (Zhao et al., 2025), we confirm
 83 that our proposed method remains highly effective on
 84 this new benchmark.

85 In summary, this paper contributes a compre-
 86 hensive experimental investigation and correspond-
 87 ing theoretical analysis of the catastrophic forgetting
 88 phenomenon during VLM fine-tuning. Our work of-
 89 fers practical guidance for adapting these models to
 90 medical domains while preserving their previous ca-
 91 pabilities. This facilitates the sequential fine-tuning
 92 of VLMS on new medical data with minimal perfor-
 93 mance loss on previously learned tasks. We anticipate
 94 that this research will benefit practitioners in the field
 95 and advance the development of more robust and the-
 96 oretically grounded VLMS. Code, checkpoints and
 97 data are available through anonymous links in Ap-
 98 pendix A.

2. Challenge: Sequentially Fine-tuning of VLMS

2.1. MLLM-CL Benchmark for General Continual Learning

The challenge of general sequential fine-tuning is analogous to the continual learning problem. Accordingly, we adopt the MLLM-CL benchmark and its evaluation protocols as proposed in a recent study (Zhao et al., 2025). This sequential learning benchmark comprises five distinct domains: Remote Sensing (**RSVQA**), Medicine (**PathVQA**), Autonomous Driving (**DriveLM**), Science (**AI2D**, **SciVerse**, **MapQA**, **TQA**), and Finance (**StockQA**).

For simplicity, these tasks are denoted as **RS**, **Med**, **AD**, **Sci**, and **Fin**, respectively. In our experiments, we follow the sequential fine-tuning order established in the original MLLM-CL study: **RS** → **Med** → **AD** → **Sci** → **Fin**. Training Details are provided in Appendix C.

2.2. Evaluation Metrics

To ensure a fair and direct comparison, our evaluation protocol strictly adheres to the methodology outlined in MLLM-CL (Zhao et al., 2025). We report two primary metrics: *Last* and *Average*. The *Last* metric represents the average accuracy across all previously

Method	Last					Average				
	RS (%)	Med (%)	AD (%)	Sci (%)	Fin (%)	RS (%)	Med (%)	AD (%)	Sci (%)	Fin (%)
Zero-shot	32.29	28.28	15.59	35.55	62.56	-	-	-	-	-
<i>w/ replay buffer</i>										
LoRA	29.57	29.19	7.09	19.55	63.60	<u>80.87</u>	58.60	38.95	36.41	36.78
MoELoRA	40.23	23.58	5.19	18.35	74.89	80.00	56.91	34.69	31.70	31.36
O-LoRA	76.21	51.34	36.50	42.64	90.20	80.13	70.23	61.35	53.34	59.38
L2P	75.21	38.50	32.31	41.05	88.05	80.09	68.64	54.79	48.68	55.02
ModalPrompt	64.77	38.60	20.61	29.98	88.22	80.11	60.99	50.67	41.97	48.44
HiDe-LLaVA	75.36	39.23	37.17	45.02	81.89	81.51	62.37	49.37	50.61	55.73
MR-LoRA	79.87	62.71	<u>51.89</u>	52.48	89.69	80.82	72.19	<u>65.41</u>	62.52	67.31
IncLoRA (Ours)	77.43	<u>62.57</u>	52.00	52.48	<u>90.41</u>	78.30	71.93	65.38	62.12	66.98
SeqFull (Ours)	<u>78.94</u>	62.45	51.50	<u>52.08</u>	91.21	75.62	<u>72.16</u>	65.77	<u>62.32</u>	<u>67.24</u>
<i>w/o replay buffer</i>										
LoRA	26.75	25.76	0.79	18.69	70.44	<u>80.72</u>	59.68	40.51	18.64	28.49
MoELoRA	21.42	25.29	0.79	17.01	60.34	80.05	57.26	37.03	19.65	24.97
O-LoRA	62.68	35.17	16.93	34.44	92.16	80.22	67.56	51.51	44.28	48.28
L2P	63.82	34.63	22.96	38.58	92.98	80.02	68.86	51.57	45.12	50.59
ModalPrompt	65.99	37.35	23.27	37.61	87.60	80.11	59.66	46.86	42.97	50.36
HiDe-LLaVA	41.17	30.33	18.73	37.08	<u>92.21</u>	80.91	65.47	39.78	32.92	43.90
IncLoRA (Ours)	<u>77.20</u>	<u>58.97</u>	<u>51.43</u>	<u>47.44</u>	90.24	77.59	<u>71.59</u>	<u>64.40</u>	<u>60.22</u>	<u>65.06</u>
SeqFull (Ours)	79.10	61.22	52.36	50.52	91.29	77.06	72.75	66.09	62.49	67.44

Table 1: Results for domain continual learning in MLLM-CL benchmark. We highlight **the best result** and the second best result separately for *w/ replay buffer* and *w/o replay buffer*.

learned tasks after the model has completed training on the final task in the sequence. The *Average* metric captures performance throughout the entire training process, defined as the mean of the average accuracies calculated after each sequential task is learned.

2.3. Medical-CL Benchmark for Medical Continual Learning

To specifically evaluate sequential fine-tuning within the medical domain, we introduce a novel benchmark, termed **Medical-CL**. This benchmark is curated from several publicly available datasets, including BSCCM (Pinkard et al., 2024), PitVis (Das et al., 2025), PathVQA (He et al., 2020), and ROCov2 (Rückert et al., 2024).

These tasks are denoted as **Cell**, **Sur**, **Path**, and **Rad**, respectively. We follow MLLM-CL and use a randomized order of **Path** \rightarrow **Sur** \rightarrow **Cell** \rightarrow **Rad**. For **Cell**, **Sur** and **Path**, we use answer accuracy as the metric, while for **Rad**, we adapt BLEU (Papineni et al., 2002) and rescale it to 0-100 to fit the regular percentage accuracy. Training Details are provided in Appendix D.

As illustrated in Figure 1, each dataset was adapted for VLM evaluation. Our benchmark comprises four distinct medical datasets, spanning a

range of scenarios from cellular-level analysis and surgical procedures to high-level pathology and radiology interpretation. This diversity provides a robust framework for evaluating sequential learning capabilities in specialized medical contexts.

3. Analysis: Why the Forgetting is Happening?

Research indicates that pretrained Large Language Models (LLMs) have loss landscapes with wide, flat, and anisotropic basins, where model performance is stable (Chen et al., 2025; Xu et al., 2024). The pre-training on web-scale data creates a general foundational basin. The process of fine-tuning can be seen as optimizing a more specialized sub-basin within this foundational one to adapt the model to a specific target task.

For a pretrained model f_θ and a task \mathcal{T} , the loss landscape $\mathcal{L}_{f_\theta, \mathcal{T}}$ contains a task-specific sub-basin. Catastrophic forgetting occurs when subsequent parameter updates push the model outside of this specialized sub-basin. To analyze this, we can decompose the parameter space into two orthogonal subspaces: a **robust subspace** $\mathcal{R}_{f_\theta, \mathcal{T}}$, characterized by low loss curvature where parameter changes have

IncLoRA	Path	Sur	Cell	Rad	Average
Zero-shot	36.01	31.45	4.99	12.86	21.11
Path	68.85				68.84
Sur	63.83	53.43			58.63
Cell	59.93	51.62	84.34		65.30
Rad	53.58	47.12	75.58	18.12	48.57

Table 2: Performance of **IncLoRA** on the Medical-CL benchmark without a replay buffer.

SeqFull	Path	Sur	Cell	Rad	Average
Zero-shot	36.01	31.45	4.99	12.86	21.11
Path	61.17				61.17
Sur	59.85	58.03			58.94
Cell	59.24	56.47	82.09		65.94
Rad	54.54	57.54	80.27	16.20	52.09

Table 3: Performance of **SeqFull** on the Medical-CL benchmark without a replay buffer.

minimal impact, and a **sensitive subspace** $\mathcal{S}_{f_\theta, \mathcal{T}}$, defined by high loss curvature where performance is acutely sensitive to changes.

The key to preventing catastrophic forgetting is to regularize parameter updates to avoid disrupting the sensitive subspace $\mathcal{S}_{f_\theta, \mathcal{T}}$ of previously learned tasks. Common fine-tuning strategies achieve this through two primary forms of regularization:

- **Low Learning Rates** act as a *soft regularization*. By discouraging large steps, this approach biases the optimization process to remain within the robust subspace $\mathcal{R}_{f_\theta, \mathcal{T}}$, reducing the likelihood of venturing into the sensitive subspace.
- **Parameter-Efficient Fine-Tuning (PEFT)** methods like LoRA impose a *structural regularization*. They restrict all updates to a predefined, low-dimensional parameter subspace. This inherently limits the dimensionality of the sensitive subspace that can be altered, thereby preserving knowledge from prior tasks by design.

In essence, both strategies mitigate forgetting by constraining parameter updates, either in magnitude or direction, to protect the sensitive dimensions critical for retaining previously acquired knowledge.

4. Solution: Limiting the Parameter Update

4.1. Ablation Study on General MLLM-CL

As presented in Table 2, we conduct a comprehensive evaluation of our proposed methods, **IncLoRA** and **SeqFull**, on the general-purpose MLLM-CL benchmark. **IncLoRA** will reinitialize a new LoRA for each task and merge the LoRA weight into the model after learning each task. Then, the next task in stream will use this merged model as the new base model and repeat the process. **SeqFull** as its name,

will sequentially train all the LLM Backbone parameters without any trick.

In the setting where a replay buffer (details in Appendix B) is utilized, many contemporary methods employ sophisticated mechanisms to mitigate forgetting. As evidenced by the results, our simple, trick-free methods achieve performance that is highly comparable to the state-of-the-art. For instance, our **SeqFull** method achieves 78.94% on the **RS** task under the **Last** metric, closely trailing the 79.87% of the more complex **MR-LoRA**, while simultaneously outperforming it in the **Fin** domain.

The advantages of our methodology become even more pronounced in the more challenging and realistic scenario without a replay buffer, since medical data involves privacy and any possible leakage from replay buffer is unacceptable. In this setting, **IncLoRA** and **SeqFull** consistently outperforms all other competing methods, establishing new benchmarks across most domains.

4.2. Application on Medical-CL

To evaluate the effectiveness of our proposed methods in a specialized and privacy-sensitive domain, we applied **IncLoRA** and **SeqFull** to the Medical-CL benchmark, as shown in Table 2 and Table 3. The results demonstrate strong performance in this challenging, buffer-free setting.

Both methods significantly outperform the zero-shot baseline across all medical subdomains (Pathology, Surgery, Cell, and Radiology). For instance, after being trained on the full sequence of tasks, **SeqFull** achieves a final average score of 52.09, while **IncLoRA** achieves 48.57, both substantial improvements over the 21.11 baseline. Notably, both models show a strong ability to acquire new knowledge, with Cell performance reaching 84.34 for **IncLoRA** and 82.09 for **SeqFull**. These results highlight the viability of our simple yet effective continual learning

246 strategies for specialized applications where data pri-
247 vacy is paramount.

248 5. Discussion

249 Our findings have significant implications for medical
250 AI, offering a replay-free continual learning method
251 that addresses data privacy by enabling institutions
252 to fine-tune VLMS on local datasets. Future work
253 could focus on optimizing regularization strategies
254 for diverse medical modalities or assessing long-term
255 performance. These next steps are vital for develop-
256 ing theoretically grounded, adaptable, and scalable
257 VLMS that can safely learn across the vast landscape
258 of medical knowledge.

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Appendix A. Annoymous Links 339

We provide fully annoymous links to our supplement- 340
 ary materials here. 341

Code 342

[https://anonymous.4open.science/r/ 343
 replay-free-finetuning-medical-vlm-BB37 344](https://anonymous.4open.science/r/replay-free-finetuning-medical-vlm-BB37)

Model Checkpoints 345

[https://huggingface.co/ 346
 Replay-Free-Finetuning-Medical-VLM/ 347
 checkpoints 348](https://huggingface.co/Replay-Free-Finetuning-Medical-VLM/checkpoints)

Medical-CL Dataset 349

[https://huggingface.co/datasets/ 350
 Replay-Free-Finetuning-Medical-VLM/ 351
 Medical-CL 352](https://huggingface.co/datasets/Replay-Free-Finetuning-Medical-VLM/Medical-CL)

Appendix B. Fine-tuning Protocol 353

Base Model. For all the evaluations in this paper, 354
 we adapt LLaVA-7B as our base model for fine-tuning. 355
 The checkpoint could be downloaded from <https://huggingface.co/llava-hf/llava-1.5-7b-hf>. 356
 357

Fine-tuning Settings. For **IncLoRA** fine- 358
 tuning, we set the LLM backbone as the LoRA target 359
 and unfreeze the projector. For **SeqFull** fine-tuning, 360
 we all the paramters of LLM backbone and projector 361
 to be trainable. 362

Training Framework. Our experimental frame- 363
 work is built upon the LLaMA-Factory repository 364
 (Zheng et al., 2024). The training configurations ad- 365
 here to the official guidelines provided in the LLaVA 366
 model repositories (Liu et al., 2023). 367

Prompt Templates. For the LLaVA model, we 368
 utilized the corresponding system prompt templates 369
 provided within the LLaMA-Factory framework. For 370
 all the evaluations except the image captioning for 371
Radiology, we turn all the questions into multiple- 372
 choice and add format instruction in prompt to avoid 373
 the influence of the mismatch of output format. 374

Replay Buffer. We exactly follow the setting in 375
MLLM-CL Zhao et al. (2025), specifically, for each 376
 task, we collect a replay data buffer of size 20 samples. 377
 Then, for every downstream sequential fine-tuning, 378
 we directly hybrid the all the replay data into the 379
 training data. No over-sampling is implemented. 380

381 **Appendix C. MLLM-CL Fine-tuning**
 382 **Hyperparameters**

383 The training length for every task is aligned to
 384 MLLM-CL (Zhao et al., 2025) for fair comparison.

Config	Value
optimizer	AdamW
batch size	64
lr schedule	cosine decay
lr warmup ratio	0.1
base lr	8×10^{-5}
epoch for RS	1
epoch for Med	3
epoch for AD	1
epoch for Sci	2
epoch for Fin	1
LoRA rank	8

Table 4: Hyperparameters of **IncLoRA** in MLLM-CL Benchmark *w/o replay buffer*.

Config	Value
optimizer	AdamW
batch size	64
lr schedule	cosine decay
lr warmup ratio	0.1
base lr	8×10^{-5}
epoch for RS	1
epoch for Med	3
epoch for AD	1
epoch for Sci	2
epoch for Fin	1
LoRA rank	16

Table 5: Hyperparameters of **IncLoRA** in MLLM-CL Benchmark *w/ replay buffer*.

Config	Value
optimizer	AdamW
batch size	16
lr schedule	cosine decay
lr warmup ratio	0.1
base lr	1×10^{-6}
epoch for RS	1
epoch for Med	3
epoch for AD	1
epoch for Sci	2
epoch for Fin	1

Table 6: Hyperparameters of **SeqFull** in MLLM-CL Benchmark *w/o replay buffer*.

Config	Value
optimizer	AdamW
batch size	16
lr schedule	cosine decay
lr warmup ratio	0.1
base lr	1×10^{-6}
epoch for RS	1
epoch for Med	3
epoch for AD	1
epoch for Sci	2
epoch for Fin	1

Table 7: Hyperparameters of **SeqFull** in MLLM-CL Benchmark *w/ replay buffer*.

385 **Appendix D. Medical-CL Fine-tuning**
 386 **Hyperparameters**

387 All the experiment in this part is *w/o replay buffer*.

Config	Value
optimizer	AdamW
batch size	64
lr schedule	cosine decay
lr warmup ratio	0.1
base lr	1×10^{-6}
step for Path	2000
step for Sur	2000
step for Cell	2000
step for Rad	2000
LoRA rank	16

Table 8: Hyperparameters of **IncLoRA** in Medical-CL Benchmark.

Config	Value
optimizer	AdamW
batch size	16
lr schedule	cosine decay
lr warmup ratio	0.1
base lr	1×10^{-6}
step for Path	2000
step for Sur	2000
step for Cell	2000
step for Rad	2000

Table 9: Hyperparameters of **SeqFull** in Medical-CL Benchmark.