1	CDeep3M-Preview: Online segmentation
2	using the deep neural network model zoo
3	
4	Authors: Matthias G Haberl ¹ , Willy Wong ¹ , Sean Penticoff ¹ , Jihyeon Je ¹ ,
5	Matthew Madany ¹ , Adrian Borchardt ¹ , Daniela Boassa ¹ , Steven T Peltier ¹ , Mark
6	H Ellisman ¹
7	
8	¹ National Center for Microscopy and Imaging Research, School of Medicine, University of
9	California San Diego, Biomedical Science Building, 9500 Gilman Drive, La Jolla
10	
11	
12	Abstract
13	
14	Sharing deep neural networks and testing the performance of trained networks
15	typically involves a major initial commitment towards one algorithm, before knowing
16	how the network will perform on a different dataset. Here we release a free online tool,
17	CDeep3M-Preview, that allows end-users to rapidly test the performance of any of the
18	pre-trained neural network models hosted on the CIL-CDeep3M modelzoo. This
19	feature makes part of a set of complementary strategies we employ to facilitate
20	sharing, increase reproducibility and enable quicker insights into biology. Namely we:
21	(1) provide CDeep3M deep learning image segmentation software through cloud
22	applications (Colab and AWS) and containerized installations (Docker and Singularity)
23	(2) co-hosting trained deep neural networks with the relevant microscopy images and
24	(3) providing a CDeep3M-Preview feature, enabling quick tests of trained networks on

user provided test data or any of the publicly hosted large datasets. The CDeep3M modelzoo and the cellimagelibrary.org are open for contributions of both, trained

models as well as image datasets by the community and all services are free of charge.

28 **Main**

New deep neural networks are developed rapidly and a startling number of trained 29 models are available online for a wide range of image enhancement and analysis tasks 30 (see ¹ for a recent review). Since training new models is however expensive and 31 typically requires laboriously expert annotated training data, innovations in sharing 32 trained models effectively are critical to reduce time and cost in research. Model zoos 33 and GitHub repositories with different networks and/or trained models are currently the 34 35 most common way to share models, but are fairly disparate from the typical workflow or processing pipelines of biomedical labs. Passively hosted model zoos do not offer 36 37 an immediate entry point to evaluate the performance of a neural network or a trained 38 model, instead require to go first through complex installations - usually on high performance systems - before being able to know if the network will be useful for the 39 40 specific question at hand. The computations to test a deep neural network typically require installation of several requisites on a high-performance GPU-equipped system 41 42 and familiarization with the individual processing routines and configurations employed. Therefore, the use of model zoos has not been able to eliminate a major 43 44 time commitment required to reproduce results. Even recent developments of more user-friendly solutions for running deep neural networks for image segmentation²⁻⁴ on 45 local or cloud resources do require a time commitment of researchers at different levels 46 47 to: set up, configure and troubleshoot then familiarize themselves with software settings and testing parameters and their influence on performance. Further training 48 and testing different models on several small or individual very large imaging datasets 49 can be cumbersome. As a result, many cutting-edge developments for image analysis 50 with deep learning are still not used by a large portion of the biomedical imaging 51 52 community.

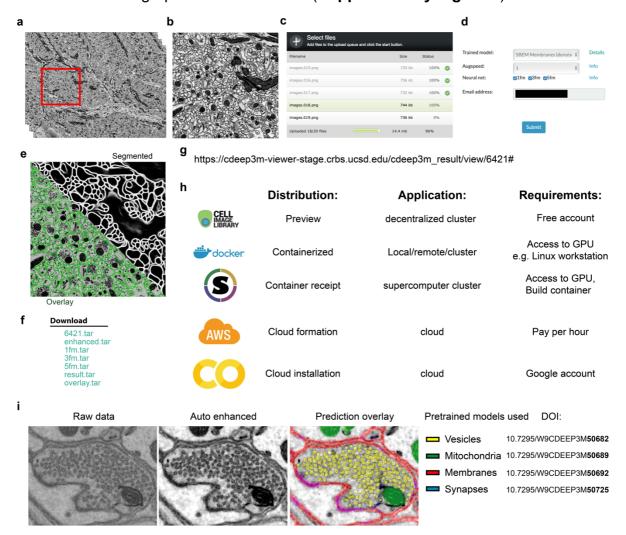
53 To facilitate sharing, increase reproducibility and enable quicker insights into biology we employ a set of innovative complementary strategies: (1) we recently 54 55 released a deep neural network platform, CDeep3M⁴, which circumvents installation 56 issues and hardware requirements for end-users. We now provide a docker container 57 of CDeep3M2 as well as a Google Colab installer with GUI (2) we are hosting CDeep3M pre-trained models in a public database (modelzoo), on cellimagelibrary.org 58 (CIL) that also hosts relevant large microscopy datasets⁵ and (3) we are now releasing 59 an online CDeep3M-Preview feature, offering instantaneous testing of any trained 60 61 neural network that is hosted on the CIL database. This allows users to 'test drive'

CDeep3M models within minutes on either their own data or on any region of interest 62 on a large number of publicly hosted imaging data to determine if a trained model of 63 interest performs well for their purpose and/or dataset (Figure 1a-g, Supplementary 64 Figure 1a-b). All results are displayed through a web interface, accessible to download 65 and can be shared with a link (Figure 1e-g, Supplementary Figure 1c). Users are 66 then guided through different options how to run the same model on a larger scale or 67 68 re-train the model with their own data using one of our distributions (Figure 1h). At the 69 same time the CDeep3M model uploader further provides users with a way to share 70 their trained models with the community in the modelzoo in a common place in a fully functional and testable state (Figure 2a). 71

72 Data processing with state-of-the-art deep neural networks requires high-end 73 hardware with GPUs with sufficient vRAM. Online processing using deep neural 74 networks for many users, as provided by CDeep3M-Preview, is limited by hardware 75 availability equipped with high-end GPUs. We therefore implemented a scheduling 76 system outsourcing the processing of the preview to the infrastructure of the Pacific 77 Research Pipeline (PRP), a decentralized computing cluster with GPUs and storage 78 nodes. The PRP cluster is managed by Kubernetes, using containerized applications to rapidly deploy computing jobs to available hardware. Our installation for the 79 CDeep3M-Preview is containerized as a Docker image on the PRP cluster, and is 80 streamed to available nodes allowing for near instantaneous start-up times (within 81 seconds). The imaging data and pre-trained models are sent from CIL and the 82 commands to initiate segmentation and the subsequent overlay with the segmentation 83 84 is submitted. Using a next-generation decentralized computer cluster, rather than running the backend processing on workstation/s, provides scalability at times of high 85 demand. In practice, this means that the end users can test whether a trained model 86 87 performs on their dataset in less than 5min total, without programming knowledge and 88 without any hardware requirements (Figure 1).

Once the user has identified a pretrained CDeep3M model and tested settings which perform well for a dataset of interest (**Figure 1e**), several routes are made available that provide quick implementations for applying the pre-trained network and the settings on the complete large-scale dataset. To this end we maintain several preconfigured installers for the use on local hardware, on supercomputer clusters or on cloud providers. Following pre-configured CDeep3M installers are provided: (i) Docker container (ii) Singularity recipe (iii) AWS cloudformation template (iv) Colab

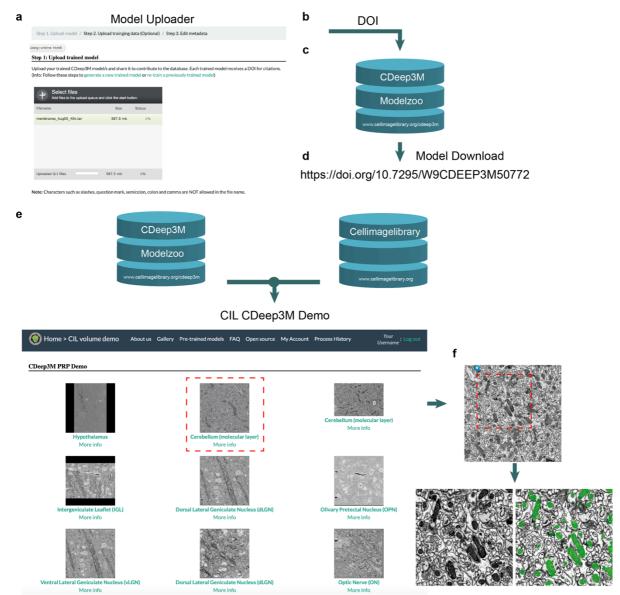
96 Notebook (**Figure 1h**). Detailed descriptions for configurations of each of those 97 solutions are available with the links provided. Without much effort or the obstacle of 98 requiring their own GPUs or funding for high-performance hardware the users can now 99 also take advantage of the free GPUs provided Google Colab, with the CDeep3M-100 colab installer and graphical user interface (**Supplementary Figure 2**).



101

102 Figure 1. Application of CDeep3M, using the preview function. (a-b) Selection of a region of interest 103 (ROI) from a large dataset to run CDeep3M-Preview. (c) Upload ROI data through web interface 104 (https://cdeep3m-stage.crbs.ucsd.edu/cdeep3m). (d) Chose trained model from CDeep3M modelzoo 105 and select parameters to perform preview. (e) Results are accessible through a web interface (here 106 shown segmented and overlay) and (f) data is accessible for download. (g) Through the web interface 107 the link can easily be shared with collaborators. (h) CDeep3M is available in multiple distributions to 108 facilitate access for many groups, for quick entry points (CDeep3M-Preview, Docker, AWS, Colab), 109 small scale tests (Preview, Colab) as well as large scale data science projects (Docker, Singularity, 110 AWS). The end-user can then either apply the trained model to the large dataset or re-train the model 111 with specific training data through one of those distributions. (i) Multiple pre-trained models are available 112 on the CDeep3M-modelzoo and were combined in this example (without re-training the network for this

- 113 dataset) to segment the cellular constituents of synapses. Auto-enhancement is performed in
- 114 CDeep3M2 providing generalizable models.



115

Figure 2. CDeep3M model uploader, database and demo function. (a) Community contributions to 116 117 the CDeep3M-modelzoo are facilitated through a web interface to upload and add metainformation to 118 trained models. (b) Each trained model receives a citable DOI and (c) is added to the CDeep3M 119 database, and becomes therefore accessible for the Preview function and (d) can be easily downloaded. 120 (e) The cellimagelibrary hosts many large-scale imaging datasets to which any of the trained networks 121 in the modelzoo can now be applied through the CDeep3M-demo function in an image browser. (f) 122 Example using a broadly trained CDeep3M model on a dataset for which it was no trained (before any 123 transfer learning is applied). Results can be easily shared through using the specific job ID and the web 124 interface. Results from (f) are at: https://cdeep3m-viewer.crbs.ucsd.edu/cdeep3m_result/view/6447 125

126 It is advantageous to co-host trained models and imaging data on the same 127 platform and maximize cross-linking between the two and facilitate testing across 128 several large datasets for generalizability. With the new extensions to the CDeep3M-

Modelzoo the users can upload their trained models to the CIL repository (Figure 2a-129 d), in order to share them or to apply them through the preview function on one of the 130 imaging datasets. In addition, metadata about the available trained models is stored in 131 the database, such as the targeted cell component, staining procedures, imaging 132 modality and voxel dimensions. When releasing a trained model on the CIL database 133 to the public, it will generate a citable Digital Object Identifier (DOI) (Figure 2b, 2d). 134 135 The DOI is a persistent identifier used to identify objects uniquely, standardized by the 136 International Organization for Standardization (ISO). To help other groups unlock 137 valuable large scale data we are providing the CDeep3M-Demo, facilitating to test pre-138 trained models on areas of interest on the large scale datasets available on 139 http://cellimagelibrary.org/ (CIL; Figure 2e). We co-host trained CDeep3M models on the CIL providing us with the infrastructure already in place for data storage, metadata 140 141 organization and large-scale image visualization. CIL is open-source software providing storage and user interfaces to deliver a publicly searchable database of 142 143 microscopy images and metadata to facilitate data sharing and reuse. The CIL data 144 input form allows end-users to submit images to the CIL data repository and annotate 145 the images with the ontology markup.

146

Together with the online preview function we release an upgrade to CDeep3M2, 147 which provides additional functionalities and reduced runtimes. The new version of 148 149 CDeep3M is backwards compatible, so that all previously trained models can still be applied and used for transfer learning with the new release. Importantly, we 150 incorporated enhanced image augmentation strategies, that can easily be configured, 151 to facilitate the training of more broadly tuned models. In the enhanced training 152 153 augmentation pipeline, the images are first processed through the sixteen rotations 154 and inversions (x/y and z) before each stack will go through a set of additional secondary and/or tertiary augmentations. The augmentations are performed as 155 156 follows: primary augmentations consisting of rotations and inversions (x: left/right, y: top/down, z: forward/reverse) are always performed, secondary augmentations 157 158 consisting of image filters (contrasting, sharpening, blurring, total variation denoising, 159 introduction of uniform noise, histogram equalization, skewing, elastic distortion; and 160 tertiary augmentations, resizing the images. Secondary and tertiary augmentation strengths are determined by the user with a scaling factor between 0 (no additional 161 162 augmentation) and 10 (strong augmentation). The combination of augmentation

strength can be customized for each dataset depending on the purpose of the training (fine tuning of model for one individual dataset or broad training for generalizable model). Furthermore, users can now easily provide multiple training datasets that will be used during training, which facilitates generating broadly applicable trained networks.

Applying the CDeep3M-Preview and Demo functions on the cellimagelibrary 168 169 serves us as an extreme test case scenario at unprecedented scale to test and improve 170 how well a trained deep neural network performs on previously unseen data 171 (generalizability). The available datasets are stored at various imaging conditions 172 (pixel- and voxelsize), ranging from 8bit unsigned integer to 32bit signed integer, with 173 different levels of noise, staining intensities and imaging conditions, acquired from 174 different tissue types. These constraints are very typical for different biological sample 175 preparation and imaging. Rather than performing training for each individual dataset, 176 we focused on training and providing more generalizing models, and stabilize their 177 performance through improved image pre-processing, which will reduce the 178 requirements to re-train the models. Mainly we automatized the following steps: image 179 conversion to 8bit, with simultaneous clipping of outlier pixels, readjustment of contrast 180 and a total variation denoising (Figure 1i). Overall, we note that these implementations improved the generalizability of trained models making them broadly applicable to 181 182 many more datasets, since this reduces most of the extreme variations in signal-to-183 noise levels and contrast between different SBEM datasets.

184 Altogether, the CDeep3M-Preview and new set of tools provided here gives biomedical researchers immediate access to experiment with AI for image 185 186 segmentation and the ability to test different trained models near-instantaneously. A similar guick entry approach has been taken recently with DeepCell, which allows 187 188 users to track cells in their own live cell imaging data with deep learning⁶. On the CDeep3M modelzoo a broad range of pre-trained models, trained on segmentation 189 190 tasks for electron microscopy, X-ray microCT and light microscopy data are readily available. Furthermore, by taking advantage of an emerging cyberinfrastructure of 191 192 decentralized compute cluster, the preview is scalable to account for the high demand 193 of many users. This approach can serve as an entry point for community members 194 with no experience in deep or machine learning to become familiar with the technology 195 and experiment with the effect of different parameter settings and will contribute to 196 democratize deep learning in the bioimaging community while allowing them to scale

their use case afterwards quickly to extremely large datasets through the CDeep3Mbuilt in processing pipelines.

199

200 **References**

- 1. Moen, E. et al. Deep learning for cellular image analysis. Nature Methods 16, 1233–
- 202 1246 (2019).
- 203 2. Falk, T. *et al.* U-Net: deep learning for cell counting, detection, and morphometry.
- 204 *Nature Methods* **16**, 67–70 (2019).
- 3. McQuin, C. et al. CellProfiler 3.0: Next-generation image processing for biology.
- 206 *PLOS Biology* **16**, e2005970 (2018).
- 4. Haberl, M. G. et al. CDeep3M—Plug-and-Play cloud-based deep learning for image
- segmentation. *Nature Methods* **15**, 677–680 (2018).
- 5. Orloff, D. N., Iwasa, J. H., Martone, M. E., Ellisman, M. H. & Kane, C. M. The cell:
- an image library-CCDB: a curated repository of microscopy data. *Nucleic Acids*
- 211 Research **41**, D1241–D1250 (2012).
- 6. Moen, E. et al. Accurate cell tracking and lineage construction in live-cell imaging
- experiments with deep learning. *bioRxiv* 803205 (2019) doi:10.1101/803205.
- 214

215 Acknowledgements

This research was funded by grants from the National Institutes of Health under award numbers 3R01GM082949, 3P41GM103412, 1RF1MH120685, 1R01AG065549 and 5R01DA038896. The CDeep3M-Preview and Demo are using PRP/Chase-CI, which is supported by its members institutions and the United States National Science Foundation through the NSF awards CNS-1456638, CNS-1730158, ACI-1540112, ACI-1541349, and OAC-1826967.

222

223 Author contributions

M.G.H. performed computational experiments. M.G.H. and J.J. wrote CDeep3M
 upgrade. W.W. implemented CDeep3M-Preview and CDeep3M-Demo interface on

CIL and model zoo uploader and database. S.P. implemented preview backend
scheduler. A.B. maintains singularity recipe and AWS cloud formation template.
M.G.H. wrote colab implementation with GUI. M.G.H. and M.M trained models hosted
on modelzoo. D.B. acquired SBEM data. M.G.H., S.T.P. and M.H.E. supervised
project. M.G.H. wrote manuscript with input from all authors.

231

232 Competing interests

- 233 The authors declare no competing interests.
- 234

235 Methods

CIL backend system. Under the CIL system, the metadata will be converted into 236 JSON format and stored in a NoSQL datastore. The CIL utilizes Elasticsearch as its 237 JSON search engine since it provides a distributed, multitenant-capable full-text 238 239 search engine with an HTTP web interface. When JSON data is imported into the 240 Elasticsearch datastore, all of the data fields are automatically indexed and immediately searchable using the built-in web-service. These built-in functions in 241 242 Elasticsearch are crucial for software development because it saves tremendous development time otherwise spent building data models and backend services. 243

244

245 Backend operations Scheduling system. At the core of the scheduling system for 246 the CDeep3M-Preview and Demo functions is the beanstalkd queue (https://beanstalkd.github.io/). Beanstalkd is a robust multichannel FIFO queuing 247 system. A single queue (or tube) contains the jobs in the order they are submitted. 248 249 Each job consists of the UID of the requestor, a description of the job to be run and its arguments, and an authentication token with a timestamp. A job can be in one of 4 250 251 states, ready, reserved, delayed, or buried. A costum written Perl based web API (stalker web) is used as an abstraction layer for the interaction with beanstalkd. A job 252 253 is submitted to the system via a client; the client does a few basic sanity checks against 254 the submitted job, secures a token and submits the job to the queue via stalker web 255 and if the submission is successful is returned the job id in the tube. Once in the tube 256 the job is in the ready state.

257 On the processing side there is a worker that periodically checks the tube for 258 ready jobs, if a job is found, it's parsed and if the worker has the capabilities needed 259 to run the job it verifies the token then reserves the job and then places it in the delayed state for an amount of time equal to the expected runtime (ERT). This removes the job from the ready state so no other workers will see the job. Since these are typically longer running jobs and are designed to run on geographically disparate hardware in an ad libitum fashion the delay is set to remove the need for the constant communication between worker and queue required by the reserved state.

The worker then sets the environment, downloads any data or models it needs, and proceeds to process the job as the UID of the submitter, collecting the output of the commands into a log. A separate watchdog process is started that will check every 90% of ERT if the worker process is still running and if it is, try to reset the delay on that job to ERT until successful or the parent process exits. Once the commands finish, the worker deletes the job from the queue and the log is published to a separate tube named with the job id of the original job.

The worker sends the output data back to the requestor via an API call. If the worker should die or communication between the worker and the queue goes down the delay time will eventually run out and the job will go back into ready state for another worker to pick up and process.

276

277 **PRP platform.** CDeep3M-preview is running as a Kubernetes Pod on the Pacific

278 Research Platform (PRP) and configured to use gpu-pods equipped with GPUs with

least 11GB vRAM. PRP is spanning over 20 universities and institutions, all

connected by dedicated optical light-paths at speeds of 10-100gb/s. A list of currently

available PRP resources can be found at: <u>https://ucsd-</u>

282 prp.gitlab.io/userdocs/running/gpu-pods/.

283

284 **CDeep3M2 data augmentations.** Data augmentations are used to avoid overfitting to 285 training data and intended to increase generalizability of trained models. To facilitate 286 regulating which data augmentation strategies are used we chose to separate 287 augmentation strategies into following three categories:

Primary augmentations refer to augmentations that only change the image orientations, such as data rotations and flipping in x, y and z directions. Since those leave the data unaltered they are always performed, on each training dataset, to generate 16 variations of the same data.

292 *Secondary augmentations* are data augmentations which alter the noise level, the 293 brightness or the contrast of the images. Secondary augmentations are performed if a

setting between 1 (weak) to 10 (strong) is chosen. Following operations are used in 294 295 secondary augmentations: increase and lowering of the image contrast; sharpening and gaussian blurring of the image stack; total variation using the Chambolle and/or 296 297 Bregman filter to denoise the image: adding of random binary gaussian noise: normalization of the image by performing histogram equalization; random skewing of 298 299 the image stack to four different directions (upper left, upper right, lower left, lower 300 right) while maintaining the image's aspect ratio; elastic distortions across the image 301 stack in which the first and the last images of the stack are distorted by a randomly 302 generated gaussian vector field while the images in between are distorted by the 303 interpolated field in between the two.

304 *Tertiary Augmentation* are performed if a setting between 1 (weak) to 10 (strong) is 305 chosen. During the tertiary augmentations images are resized, according to a strength 306 selected by the end user (values 1-10). Data is resized using upscaling alternating with 307 downscaling, to broaden the networks capabilities to recognize the same object at a 308 different pixelsize.

309

310 Code availability

All code is open access. The CDeep3M2 Docker container can be pulled directly from Docker-Hub at <u>https://hub.docker.com/r/ncmir/cdeep3m</u> or simply running 'docker pull ncmir/cdeep3m:latest'. The Google Colab Jupyter Notebooks with graphical userinterface (GUI) are available on GitHub at <u>https://github.com/haberlmatt/cdeep3m-</u> <u>colab</u>. The CDeep3M2 AWS cloudformation template is available <u>here</u>. CDeep3M2 source code is available on GitHub <u>https://github.com/CRBS/cdeep3m2</u>. The singularity image is available at: <u>http://cellimagelibrary.org/cdeep3m/singularity</u>.