

Generative Adversarial Networks in Ophthalmology: What are these and how can they be used?

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Declaration of interest: Drs Lim, Wong and Ting are the co-inventors of a deep learning system for retinal diseases.

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Sources of Funding:

This work was supported by grants R01EY19474, R01 EY031331, R21 EY031883, and P30 EY10572 from the National Institutes of Health (Bethesda, MD), and by unrestricted departmental funding and a Career Development Award (JPC) from Research to Prevent Blindness (New York, NY).

Financial Disclosures:

J. Peter Campbell is a consultant to Boston AI.

ABSTRACT

Purpose of review

The development of deep learning (DL) systems requires a large amount of data, which may be limited by costs, protection of patient information and low prevalence of some conditions. Recent developments in artificial intelligence techniques have provided an innovative alternative to this challenge via the synthesis of biomedical images within a DL framework known as Generative Adversarial Networks (GANs). This paper aims to introduce how GANs can be deployed for image synthesis in ophthalmology, and to discuss the potential applications of GANs-produced images.

Recent findings

Image synthesis is the most relevant function of GANs to the medical field, and it has been widely used for generating “new” medical images of various modalities. In ophthalmology, GANs have mainly been utilized for augmenting classification and predictive tasks, by synthesizing fundus images and OCT images with and without pathologies such as age-related macular degeneration and diabetic retinopathy. Despite their ability to generate high-resolution images, the development of GANs remains data intensive, and there is a lack of consensus on how best to evaluate the outputs produced by GANs.

Summary

Although the problem of artificial biomedical data generation is of great interest, image synthesis by GANs represents an innovation with yet unclear relevance for ophthalmology.

Keywords (3 to 5):

Artificial intelligence, deep learning, generative adversarial networks, medical image synthesis, ophthalmology

INTRODUCTION

In the field of ophthalmology over the last few years, clinically applicable deep learning (DL) systems have been developed to detect different eye diseases, such as diabetic retinopathy (DR),[1-4] glaucoma,[3, 5] age-related macular degeneration(AMD),[6, 7] and retinopathy of prematurity (ROP) [8]. This has led to the real possibility that such DL systems may be implemented soon in appropriate clinical settings, such as in DR screening programs [9, 10].

Despite the substantial promise of DL, the development of a robust DL algorithm or system is data intensive, meaning that a large amount of data exhibiting representative variability (i.e., disease and normal) is required for the training and validation process [11]. The availability of such large datasets is often limited by the lack of corresponding clinical cohorts, the high costs of starting a primary data collection from baseline, and the need to protect the privacy of patients. Personal information of patients must be protected under rigorously controlled conditions and in accordance with the best research practices [12, 13]. Moreover, medical images are considered identifiable personal information and cannot be anonymized easily, and consent is difficult to obtain for large retrospective datasets [14-16]. In addition, annotated data of the more severe phenotypes of certain pathologies such as advanced glaucoma and neovascular or late AMD are often too uncommon in existing population studies to be useful for conducting DL analysis.

Recent developments in DL have provided an innovative alternative to these challenges, by using generative adversarial networks (GANs) to artificially create new images based on smaller real image datasets. There is significant potential to generate a large number of images required to train, develop, validate and test new DL algorithms and systems.

WHAT ARE GANS?

GANs are a special type of neural network model based on a game theoretic approach, with the objective being to find Nash equilibrium between two networks: a generator and a discriminator (Figure 1). The idea is to sample from a simple distribution, like Gaussian, and then learn to transform this noise to a targeted data distribution, using universal function approximators such as convolutional neural

networks (CNNs), by the adversarial training of generator and discriminator simultaneously. The generator model learns to capture the targeted data distribution, and the discriminator model estimates the probability that a sample comes from the targeted data distribution rather than the distribution generated by the generator. In other words, the task of generator is to generate natural looking images and the task of discriminator is to decide whether the image is fake or real. This can be thought of as a minimax two-player game, i.e. generator vs discriminator, where the performance of both the networks ideally improves over time iteratively. While the generator tries to fool the discriminator by generating images that appear as real as possible, the discriminator tries to not get fooled by the generator by improving its discriminative capability [17].

There are many ways to incorporate GANs in medical imaging tasks, such as segmentation,[18] classification,[19] detection,[20] registration,[21] image reconstruction[22] and image synthesis (Table 1)[23]. GANs have been used in research studies for generating medical images of various image modalities, including breast ultrasound,[24] mammograms,[25] computed tomography (CT),[26-29] magnetic resonance images (MRI),[30] cancer pathology images,[31] and contrast agent-free ischemic heart disease images[32]. Moreover, GANs have been shown to be capable of cross modality image synthesis, such as generating MRI based on ultrasound[33] or CT[34, 35]. This paper focuses on the image synthesis aspect of GANs in ophthalmology via introducing different types of GANs, summarizing GAN models reported in the literatures of ophthalmology (Table 2), discussing the outcome measures for GANs, and evaluating the advantages and disadvantages of GANs.

DIFFERENT TYPES OF GANS

There are many different formulations of GANs,[36] which might firstly be categorized according to their objectives. While GANs have been employed to generate sequential data such as text, their most common usage has been in imaging tasks, including video. Within image processing, GANs have been employed to perform texture synthesis, super-resolution, object detection and image synthesis, all of which could have potential applications in medicine. GANs can be further defined by their features, the more prominent of which may be their neural

architecture for generators and discriminators, their objective function, and their training procedure. Each of these components has undergone significant development since the introduction of GANs. For example, the conditioning of GANs referred as cGAN, on both the generator and discriminator would be demonstrated soon after their inception,[37] through the addition of a prior as input. Various objective functions have been proposed to address issues such as instability in training, some of the most prominent amongst of which include Wasserstein GAN,[38] which seeks to maintain a continuous distance when real and generated data distributions are disjoint, and LS-GAN,[39] which encourages the generated data distribution to be closer to the real distribution through the implementation of mean squared loss instead of log loss. PatchGAN[40] is proposed to run discriminators on patches or on images at different scales in order to improve the quality of image synthesis such as angiograph image synthesis. As for training procedure, individual learning rates for the discriminator and generator have been shown to converge to a local equilibrium [41].

Improving the quality of synthesized images at higher resolutions has been of particular interest for medical applications, which are often especially sensitive to subtle details within images. This has been enabled by innovations such as the progressive growing of the generator and discriminator with Progressive GAN (ProGAN),[42] orthogonal regularization with BigGAN,[43] and the extension of ProGAN with a number of incremental improvements such as a mapping network bilinear upsampling and block noise into StyleGAN [44]. It should be noted that many individual features of various GANs are possibly compatible, and may thus be combined into custom GAN architectures towards specific applications.

GAN FOR IMAGE SYNTHESIS IN OPHTHALMOLOGY

Fundus images

Using generated vessel trees as an intermediate stage, Costa et al. were among the first few groups to deploy adversarial learning for building an end-to-end system for synthesizing retinal fundus images. This system was trained on a small data set of 614 normal fundus images and tends to generate fixed outputs that are lack of diversity and pathological features [45]. Following this, various methods have been attempted to improve the quality and diversity of the synthetic fundus photos. Guibas

et al. proposed a two-stage GAN pipeline by first generating synthetic retinal blood vessel trees and then translating these masks to photorealistic images. The synthetic images were used to train a U-net segmentation network, achieving similar F1-score as the network trained on real images [46]. Using direct mapping from manual tubular structured annotation back to a raw image, the model developed by Zhao et al. can synthesize multiple images with diversity using a dataset as small as 10 training examples [47]. Yu et al. built a multiple-channels-multiple-landmarks pipeline using a combination of vessel tree, optic disc and optic cup images to generate color fundus photos, which produced superior images than single vessel-based approach [48].

Besides generating normal fundus images, GANs have been used to synthesize fundus photos of specific eye diseases, such as AMD, glaucoma, DR and ROP (Figure 2). Using over 100,000 color fundus photographs from the Age-Related Eye Disease Study (AREDS), Burlina et al. built two ProGAN models to synthesize referable (intermediate and/or advanced AMD) and non-referable AMD images respectively [49]. The outputs were realistic enough that two retinal specialists could not distinguish real images from the synthetic ones, with accuracy of 53.67% and 59.50% respectively. Furthermore, the DL system trained on synthetic data alone showed comparable diagnostic performance to the algorithm trained on real images in detecting referable AMD (AUC of 0.92 vs. AUC of 0.97). Diaz-Pinto et al. built a retinal image synthesizer and semi-supervised classifier for glaucoma detection using GANs on over 80,000 images. Their system was capable of generating realistic synthetic images with features of glaucoma and labelling glaucomatous images automatically with an AUC of 0.902 [50]. Furthermore, Zhou et al. report a DR fundus photo generator that can be directly deployed to train DR classifier, via modification with arbitrary grading and lesion information to synthesize high quality images [51]. Beers et al. trained a ProGAN model with 5,550 posterior pole retinal photographs of ROP, which could produce realistic fundus images of ROP. They evaluated the performance of a segmentation algorithm trained on synthetic images, reporting an AUC of 0.97 comparing to the segmentation maps from real images [52].

OCT

Utilizing over 100,000 OCT images from eyes with balanced distribution of urgent referrals (choroidal neovascularization, diabetic macular edema) and non-urgent referrals (drusen and normal eyes), Zheng et al. built a ProGAN model to synthesize OCT images that are realistic to retinal specialists. The DL framework trained on synthetic OCT images achieved an AUC of 0.905 in classifying urgent and nonurgent referrals, which was non-inferior to the performance of the model trained on real OCT images (AUC = 0.937) [53]. Apart from synthesizing OCT images from scratch, GAN has been used to enhance the image quality of OCT scans via denoising. Using noisy images and corresponding high-quality images from one normal eye, Ma et al. built an image-to-image cGAN, enabling the competition of the generator and the discriminator to learn the underlying structure of the retina layers and to reduce OCT speckle noise. Despite the small training dataset, their model was capable of generalizing to images with low signal-to-noise (LSTN) ratio from pathological eyes and from different OCT scanners [54]. Similarly, using a small set of OCT images with high signal-to-noise (HSTN) ratio and LSTN ratio from the same eye of 28 patients, Kande et al. equipped a GAN model with Siamese network to generate denoised spectra-domain OCT images that are closer to the ground truth images with HSTN ratio. The discriminator was designed to fool the generator to produce a denoised image via enabling extraction of the discriminative features from the HSNR patch and denoised patch by passing them through a twin network [55].

The image synthesis function of GANs could also be applied for predicting the post-treatment OCT images of patients receiving anti-vascular endothelial growth factor (anti-VEGF). Utilizing 476 pairs of pre- and post-therapeutic OCT images of patients with neovascular AMD (nAMD) who received anti-VEGF treatment, Liu et al. proposed an image-to-image GAN model to generate predicted post-therapeutic OCT images based on their pre-therapeutic ones. Their GAN model achieved a sensitivity of 84% and specificity of 86% in predicting the post-treatment macular classification (wet or dry macula) [56]. Following this, Lee et al. trained a cGAN model to predict post-treatment OCT images in patients with nAMD receiving anti-VEGF, using a larger dataset of 15,183 paired OCT B-scans from 241 patients [57]. This cGAN model was designed to predict the presence of four abnormal structures on post-treatment OCT, namely the intraretinal fluid, subretinal fluid, pigmented epithelial detachment, and subretinal hyperreflective material, with sensitivity and

specificity ranging from 21.2-88.2% and 94.6-95.1% respectively. The predictive performance was enhanced after adding fluorescein angiography and indocyanine green angiography images to the training datasets. As a result of the low sensitivity, the authors concluded that this model is not suitable to be used as a screening tool and further work with a dataset of more variations is warranted. Although GAN synthesized retinal images have overall consistent appearance, generating realistic images with pathological retinal lesions remains as a challenge [58].

Other image modalities

Recently, Tavakkoli et al. proposed a GAN model capable of producing fluorescein angiography (FA) from retinal fundus photographs, which was the first DL application to generate images from distinct modalities in ophthalmology [59]. Using pairs of FA and fundus images from 59 patients (30 with diabetic retinopathy, 29 normal) as the training dataset, they designed a multi-scale cGAN comprised of two generators and four discriminators. Their framework was able to produce FA images that are indistinguishable from real ones by three experts and are more accurate than the images produced by another two state-of-the-art cGAN models, as evidenced by significantly lower Fréchet Inception Distance and higher structural similarity measures. This cGAN technique may be a novel alternative to the invasive FA and the expensive OCT angiography with limited field of view. Furthermore, the ability to generate FA based on fundus photographs may improve the efficiency of tele-medicine, in particular during the COVID-19 pandemic when in-clinic examinations becomes challenging [59].

OUTCOME MEASURES FOR GANS

Since GANs are generative models, the evaluation of their outputs – usually images – is not as straightforward as for discriminative models like classifiers, where the predicted label can simply be compared against the ground truth, in a supervised context. For GANs, the evaluation of their outputs may intuitively be based on human judgment of their “realness”, which in turn can be broadly considered in two aspects. Firstly, fidelity, in the sense that the generated samples are visually indistinguishable from real samples of the intended class. Fidelity in turn generally depends on various characteristics of the generated images, such as overall quality (being in-focus, etc.), demonstrating plausible object texture and structure, and so forth. Secondly,

diversity, where the full range of variation of the intended class is generated. For example, if a GAN had been trained for cars, it would exhibit poor diversity if all images that it generated were of a particular car manufacturer or colour, despite the cars being otherwise realistic. When this happens, possibly due to the GAN's generator fixating on some particularly plausible output, the GAN is said to undergo mode collapse.

Many techniques have been proposed to evaluate GANs outputs, for both qualitative and quantitative measurements [60]. Qualitative methods may involve humans in inspecting and curating the generated images based on graders' subjective decisions. This may be particularly appropriate in the biomedical domain, since it may not be easily articulated as to why some generated samples are not physiologically plausible. Quantitative methods on the other hand tend not to involve direct human intervention, and the most natural test would perhaps be through classifying the generated samples with a discriminative model trained on real samples. A popular generalization of this basic idea is the Inception Score (IS), [61] which uses an Inception v3 model pre-trained on ImageNet, to classify a set of generated images. The IS was claimed by its developers to be highly correlated with human judgment. However, due to its reliance on the ImageNet dataset, it has been noted to be possibly inappropriate when applied to GANs trained on other datasets, being overly sensitive to network weights, and insensitive to prior distribution over labels [62, 63].

The Fréchet Inception Distance [41] is perhaps among the most commonly used outcome metrics for GANs, and involves embedding generated images into a feature space expressed by an embedding layer of the Inception v3 model, estimating the mean and covariance for the embeddings of the generated and real data, and computing the Fréchet distance between these two Gaussians. This improves upon IS by being able to quantify mode dropping/collapse, but is still unable to recognize overfitting, as when the GANs reproduce samples from the training data [62]. To address this, qualitative measures such as comparing generated samples to their nearest neighbours in the training data, might be considered [60].

Additionally, generated synthetic images may be evaluated against the GAN training dataset of real images, to ensure that the synthetic images are not merely minimally-adapted from the real data. Metrics such as the structural similarity index measure (SSIM) may be used to efficiently compare a sample of the generated data against the real images, in a pairwise manner [64].

ADVANTAGES AND DISADVANTAGES OF GANS

GANs have gained recognition due to their various advantages over previous generative models. While other unsupervised generative models share the advantage of not requiring initial human annotation, GANs differ from deep graphical models in that they do not require careful architectural design, and further do not rely on Markov chains for sampling, unlike generative autoencoders [17]. This simplifies and generalizes the modelling process, which may be convenient for researchers primarily interested in medical applications. Empirically, recent GAN architectures have been favoured over alternative generative models for the fidelity of their generated high-resolution images. Moreover, while GANs do not promise inference capabilities, techniques have been developed to explore their encoded latent space [65]. In the medical domain, GANs have been proposed to augment existing datasets and help preserve patient confidentiality through the generation of additional samples, especially of rarer conditions [66, 67].

Nevertheless, GANs do still possess certain disadvantages. Other than the abovementioned possibility of mode collapse, GANs do not explicitly represent the generator's distribution over the data, which is detrimental to model interpretability. However, this is also the case for other popular generative models. Also, GANs tend to be data intensive, in that the training data needs to sufficiently represent the desired underlying class. To give a concrete example, consider a GAN that is trained to generate "abnormal fundus images", but its training data are almost entirely composed of diabetic retinopathy samples, with a few maculopathy samples, and next to no glaucoma examples. In this case, it is very unlikely that the GAN will be able to generate an acceptable diversity of glaucomatous images. This might be addressed to an extent through human-in-the-loop training (Figure 3), where human guidance is introduced to select acceptable synthetic data generated by the GAN, during the GAN training process. These selected synthetic images can then be used

to augment both the training of the discriminator model, and the further fine-tuning of the GAN generator model itself.

CONCLUSION AND FUTURE DIRECTION

In conclusion, GANs offer a potential innovative solution to address a key challenge in the DL field for ophthalmology: that of limited access to large datasets. Of all the functions of GANs, image synthesis is the most relevant and explored by biomedical research. In ophthalmology, GANs are mainly utilized for synthesizing fundus images and OCT images with and without pathology such as AMD and DR. Although the unmet need of artificial biomedical data generation is of great interest, DL solutions such as GANs still face many challenges in the retinal image synthesis field. First, the development of GANs is data intensive, but how much data is considered sufficient to train GANs remains unknown, and the amount is likely to be task dependent. Second, although Inception Score and Fréchet Inception Distance have been commonly used for quantitative measurement of GANs outputs, current qualitative measurement for the outputs of GANs mainly relies on the subjective judgement of human. Therefore, an objective scale to evaluate the quality of synthetic images such as the realness may be proposed for comparison among different GAN models. Lastly, to test whether GANs could really be the solution to limited access to datasets, future research is warranted to evaluate if GANs generated images could augment the development of DL systems and to test the performance of synthetic image trained DL systems using independent datasets.

REFERENCE

1. Gulshan V, Peng L, Coram M, Stumpe MC, Wu D, Narayanaswamy A et al. Development and Validation of a Deep Learning Algorithm for Detection of Diabetic Retinopathy in Retinal Fundus Photographs. *JAMA*. 2016;316(22):2402-10. doi:10.1001/jama.2016.17216.
2. Abramoff MD, Lou Y, Erginay A, Clarida W, Amelon R, Folk JC et al. Improved Automated Detection of Diabetic Retinopathy on a Publicly Available Dataset Through Integration of Deep Learning. *Investigative Ophthalmology & Visual Science*. 2016;57(13):5200-6. doi:10.1167/iovs.16-19964.
3. Ting DSW, Cheung CY-L, Lim G, Tan GSW, Quang ND, Gan A et al. Development and Validation of a Deep Learning System for Diabetic Retinopathy and Related Eye Diseases Using Retinal Images From Multiethnic Populations With Diabetes. *JAMA*. 2017;318(22):2211-23. doi:10.1001/jama.2017.18152.
4. Gargeya R, Leng T. Automated Identification of Diabetic Retinopathy Using Deep Learning. *Ophthalmology*. 2017;124(7):962-9. doi:10.1016/j.ophtha.2017.02.008.
5. Hood DC, De Moraes CG. Efficacy of a Deep Learning System for Detecting Glaucomatous Optic Neuropathy Based on Color Fundus Photographs. *Ophthalmology*. 2018;125(8):1207-8. doi:10.1016/j.ophtha.2018.04.020.
6. Burlina PM, Joshi N, Pekala M, Pacheco KD, Freund DE, Bressler NM. Automated Grading of Age-Related Macular Degeneration From Color Fundus Images Using Deep Convolutional Neural Networks. *JAMA Ophthalmology*. 2017;135(11):1170-6. doi:10.1001/jamaophthalmol.2017.3782.
7. Grassmann F, Mengelkamp J, Brandl C, Harsch S, Zimmermann ME, Linkohr B et al. A Deep Learning Algorithm for Prediction of Age-Related Eye Disease Study Severity Scale for Age-Related Macular Degeneration from Color Fundus Photography. *Ophthalmology*. 2018;125(9):1410-20. doi:10.1016/j.ophtha.2018.02.037.
8. Brown JM, Campbell JP, Beers A, Chang K, Ostmo S, Chan RVP et al. Automated Diagnosis of Plus Disease in Retinopathy of Prematurity Using Deep Convolutional Neural Networks. *JAMA Ophthalmol*. 2018;136(7):803-10. doi:10.1001/jamaophthalmol.2018.1934.
9. Keel S, Lee PY, Scheetz J, Li Z, Kotowicz MA, Maclsaac RJ et al. Feasibility and patient acceptability of a novel artificial intelligence-based screening model for diabetic retinopathy at endocrinology outpatient services: a pilot study. *Scientific Reports*. 2018;8(1):4330. doi:10.1038/s41598-018-22612-2.
10. Bhuiyan A, Govindaiah A, Deobhakta A, Gupta M, Rosen R, Saleem S et al. Development and Validation of an Automated Diabetic Retinopathy Screening Tool for Primary Care Setting. *Diabetes Care*. 2020;43(10):e147-e8. doi:10.2337/dc19-2133.
11. Ting DSW, Lee AY, Wong TY. An Ophthalmologist's Guide to Deciphering Studies in Artificial Intelligence. *Ophthalmology*. 2019;126(11):1475-9. doi:10.1016/j.ophtha.2019.09.014.
12. The Health Insurance Portability and Accountability Act of 1996 (HIPAA), Security and Privacy 45 CFR Part 160.103 (2002) (1996).
13. HIPAA for Professionals. U.S. Department of Health & Human Services. <https://www.hhs.gov/hipaa/for-professionals/index.html>. Accessed 25 May 2021.
14. The L. Striking the right balance between privacy and public good. *The Lancet*. 2006;367(9507):275. doi:10.1016/S0140-6736(06)68043-4.

15. Your Rights Under HIPAA. HHS.gov. <https://www.hhs.gov/hipaa/for-individuals/guidance-materials-for-consumers/index.html>. Accessed 18 Mar 2021.
16. Informed consent for medical photographs. Dysmorphology Subcommittee of the Clinical Practice Committee, American College of Medical Genetics. *Genet Med*. 2000;2(6):353-5. doi:10.1097/00125817-200011000-00010.
17. Goodfellow IJ, Pouget-Abadie J, Mirza M, Xu B, Warde-Farley D, Ozair S et al., editors. *Generative Adversarial Nets* 2014. Cambridge, MA, USA: MIT Press.
18. Rezaei M, Yang H, Meinel C, editors. *Whole Heart and Great Vessel Segmentation with Context-aware of Generative Adversarial Networks* 2018; Berlin, Heidelberg: Springer Berlin Heidelberg.
19. Madani A, Moradi M, Karargyris A, Syeda-Mahmood T, editors. *Semi-supervised learning with generative adversarial networks for chest X-ray classification with ability of data domain adaptation*. 2018 IEEE 15th International Symposium on Biomedical Imaging (ISBI 2018); 2018 4-7 April 2018.
20. Alex V, Safwan K. P. M, Chennamsetty SS, Krishnamurthi G. *Generative adversarial networks for brain lesion detection*. *SPIE Medical Imaging*. SPIE; 2017.
21. Yan P, Xu S, Rastinehad A, Wood B, editors. *Adversarial Image Registration with Application for MR and TRUS Image Fusion*. *MLMI@MICCAI*; 2018.
22. Wolterink JM, Leiner T, Viergever MA, Išgum I. *Generative Adversarial Networks for Noise Reduction in Low-Dose CT*. *IEEE Trans Med Imaging*. 2017;36(12):2536-45. doi:10.1109/TMI.2017.2708987.
23. Yi X, Walia E, Babyn P. *Generative adversarial network in medical imaging: A review*. *Medical Image Analysis*. 2019;58:101552. doi:<https://doi.org/10.1016/j.media.2019.101552>.
24. Fujioka T, Mori M, Kubota K, Kikuchi Y, Katsuta L, Adachi M et al. *Breast Ultrasound Image Synthesis using Deep Convolutional Generative Adversarial Networks*. *Diagnostics (Basel)*. 2019;9(4). doi:10.3390/diagnostics9040176.
25. Guan S, Loew M. *Breast cancer detection using synthetic mammograms from generative adversarial networks in convolutional neural networks*. *J Med Imaging (Bellingham)*. 2019;6(3):031411. doi:10.1117/1.Jmi.6.3.031411.
26. Baydoun A, Xu KE, Heo JU, Yang H, Zhou F, Bethell LA et al. *Synthetic CT Generation of the Pelvis in Patients With Cervical Cancer: A Single Input Approach Using Generative Adversarial Network*. *IEEE Access*. 2021;9:17208-21. doi:10.1109/access.2021.3049781.
27. Jiang Y, Chen H, Loew M, Ko H. *COVID-19 CT Image Synthesis With a Conditional Generative Adversarial Network*. *IEEE J Biomed Health Inform*. 2021;25(2):441-52. doi:10.1109/jbhi.2020.3042523.
28. Liu Y, Meng L, Zhong J. *MAGAN: Mask Attention Generative Adversarial Network for Liver Tumor CT Image Synthesis*. *J Healthc Eng*. 2021;2021:6675259. doi:10.1155/2021/6675259.
29. Toda R, Teramoto A, Tsujimoto M, Toyama H, Imaizumi K, Saito K et al. *Synthetic CT image generation of shape-controlled lung cancer using semi-conditional InfoGAN and its applicability for type classification*. *Int J Comput Assist Radiol Surg*. 2021;16(2):241-51. doi:10.1007/s11548-021-02308-1.
30. Berm´dez C, Plassard A, Davis LT, Newton A, Resnick S, Landman B, editors. *Learning implicit brain MRI manifolds with deep learning*. *Medical Imaging*; 2018.
31. Levine AB, Peng J, Farnell D, Nursey M, Wang Y, Naso JR et al. *Synthesis of diagnostic quality cancer pathology images by generative adversarial networks*. *J Pathol*. 2020;252(2):178-88. doi:10.1002/path.5509.

32. Xu C, Xu L, Ohorodnyk P, Roth M, Chen B, Li S. Contrast agent-free synthesis and segmentation of ischemic heart disease images using progressive sequential causal GANs. *Med Image Anal.* 2020;62:101668. doi:10.1016/j.media.2020.101668.
33. Jiao J, Namburete ALL, Papageorghiou AT, Noble JA. Self-Supervised Ultrasound to MRI Fetal Brain Image Synthesis. *IEEE Trans Med Imaging.* 2020;39(12):4413-24. doi:10.1109/tmi.2020.3018560.
34. Jin C-B, Kim H, Liu M, Jung W, Joo S, Park E et al. Deep CT to MR Synthesis Using Paired and Unpaired Data. *Sensors (Basel).* 2019;19(10):2361. doi:10.3390/s19102361.
35. Lee JH, Han IH, Kim DH, Yu S, Lee IS, Song YS et al. Spine Computed Tomography to Magnetic Resonance Image Synthesis Using Generative Adversarial Networks : A Preliminary Study. *J Korean Neurosurg Soc.* 2020;63(3):386-96. doi:10.3340/jkns.2019.0084.
36. Gui J, Sun Z, Wen Y, Tao D, Ye J-p. A Review on Generative Adversarial Networks: Algorithms, Theory, and Applications. *ArXiv.* 2020;abs/2001.06937.
37. Mirza M, Osindero S. Conditional Generative Adversarial Nets. *ArXiv.* 2014;abs/1411.1784.
38. Arjovsky M, Chintala S, Bottou L. Wasserstein Generative Adversarial Networks. In: Doina P, Yee Whye T, editors. *Proceedings of the 34th International Conference on Machine Learning; Proceedings of Machine Learning Research: PMLR;* 2017. p. 214--23.
39. Mao X, Li Q, Xie H, Lau RYK, Wang Z, Smolley SP. Least Squares Generative Adversarial Networks. *2017 IEEE International Conference on Computer Vision (ICCV).* 2017:2813-21.
40. Li C, Wand M. Precomputed Real-Time Texture Synthesis with Markovian Generative Adversarial Networks. *ArXiv.* 2016;abs/1604.04382.
41. Heusel M, Ramsauer H, Unterthiner T, Nessler B, Hochreiter S, editors. *GANs Trained by a Two Time-Scale Update Rule Converge to a Local Nash Equilibrium.* NIPS; 2017.
42. Karras T, Aila T, Laine S, Lehtinen J. Progressive Growing of GANs for Improved Quality, Stability, and Variation. *ArXiv.* 2018;abs/1710.10196.
43. Brock A, Donahue J, Simonyan K. Large Scale GAN Training for High Fidelity Natural Image Synthesis. *ArXiv.* 2019;abs/1809.11096.
44. Karras T, Laine S, Aila T. A Style-Based Generator Architecture for Generative Adversarial Networks. *2019 IEEE/CVF Conference on Computer Vision and Pattern Recognition (CVPR).* 2019:4396-405.
45. Costa P, Galdran A, Meyer MI, Niemeijer M, Abramoff M, Mendonca AM et al. End-to-End Adversarial Retinal Image Synthesis. *IEEE Trans Med Imaging.* 2018;37(3):781-91. doi:10.1109/TMI.2017.2759102.
46. Guibas JT, Viridi TS, Li P. Synthetic Medical Images from Dual Generative Adversarial Networks. *ArXiv.* 2017;abs/1709.01872.
47. Zhao H, Li H, Maurer-Stroh S, Cheng L. Synthesizing retinal and neuronal images with generative adversarial nets. *Medical Image Analysis.* 2018;49:14-26. doi:10.1016/j.media.2018.07.001.
48. Yu Z, Xiang Q, Meng J, Kou C, Ren Q, Lu Y. Retinal image synthesis from multiple-landmarks input with generative adversarial networks. *Biomed Eng Online.* 2019;18(1):62. doi:10.1186/s12938-019-0682-x.
49. Burlina PM, Joshi N, Pacheco KD, Liu TYA, Bressler NM. Assessment of Deep Generative Models for High-Resolution Synthetic Retinal Image Generation of Age-Related Macular Degeneration. *JAMA Ophthalmology.* 2019;137(3):258-64. doi:10.1001/jamaophthalmol.2018.6156.

50. Diaz-Pinto A, Colomer A, Naranjo V, Morales S, Xu Y, Frangi AF. Retinal Image Synthesis and Semi-Supervised Learning for Glaucoma Assessment. *IEEE Trans Med Imaging*. 2019;38(9):2211-8. doi:10.1109/tmi.2019.2903434.
51. Zhou Y, Wang B, He X, Cui S, Shao L. DR-GAN: Conditional Generative Adversarial Network for Fine-Grained Lesion Synthesis on Diabetic Retinopathy Images. *IEEE J Biomed Health Inform*. 2020;Pp. doi:10.1109/jbhi.2020.3045475.
52. Beers A, Brown J, Chang K, Campbell J, Ostmo S, Chiang M et al. High-resolution medical image synthesis using progressively grown generative adversarial networks. 2018.
53. Zheng C, Xie X, Zhou K, Chen B, Chen J, Ye H et al. Assessment of Generative Adversarial Networks Model for Synthetic Optical Coherence Tomography Images of Retinal Disorders. *Transl Vis Sci Technol*. 2020;9(2):29. doi:10.1167/tvst.9.2.29.
54. Ma Y, Chen X, Zhu W, Cheng X, Xiang D, Shi F. Speckle noise reduction in optical coherence tomography images based on edge-sensitive cGAN. *Biomed Opt Express*. 2018;9(11):5129-46. doi:10.1364/boe.9.005129.
55. Kande NA, Dakhane R, Dukkipati A, Yalavarthy PK. SiameseGAN: A Generative Model for Denoising of Spectral Domain Optical Coherence Tomography Images. *IEEE Trans Med Imaging*. 2021;40(1):180-92. doi:10.1109/tmi.2020.3024097.
56. Liu Y, Yang J, Zhou Y, Wang W, Zhao J, Yu W et al. Prediction of OCT images of short-term response to anti-VEGF treatment for neovascular age-related macular degeneration using generative adversarial network. *Br J Ophthalmol*. 2020;104(12):1735-40. doi:10.1136/bjophthalmol-2019-315338.
57. Lee H, Kim S, Kim MA, Chung H, Kim HC. POST-TREATMENT PREDICTION OF OPTICAL COHERENCE TOMOGRAPHY USING A CONDITIONAL GENERATIVE ADVERSARIAL NETWORK IN AGE-RELATED MACULAR DEGENERATION. *Retina*. 2021;41(3):572-80. doi:10.1097/iae.0000000000002898.
58. Bellemo V, Burlina P, Yong L, Wong TY, Ting DSW, editors. *Generative Adversarial Networks (GANs) for Retinal Fundus Image Synthesis 2019*; Cham: Springer International Publishing.
59. Tavakkoli A, Kamran SA, Hossain KF, Zuckerbrod SL. A novel deep learning conditional generative adversarial network for producing angiography images from retinal fundus photographs. *Sci Rep*. 2020;10(1):21580. doi:10.1038/s41598-020-78696-2.
60. Borji A. Pros and cons of GAN evaluation measures. *Computer Vision and Image Understanding*. 2019;179:41-65. doi:<https://doi.org/10.1016/j.cviu.2018.10.009>.
61. Salimans T, Goodfellow I, Zaremba W, Cheung V, Radford A, Chen X, editors. *Improved Techniques for Training GANs*. NIPS; 2016.
62. Barratt ST, Sharma R. A Note on the Inception Score. *ArXiv*. 2018;abs/1801.01973.
63. Lucic M, Kurach K, Michalski M, Gelly S, Bousquet O, editors. *Are GANs Created Equal? A Large-Scale Study*. NeurIPS; 2018.
64. Zhou W, Bovik AC, Sheikh HR, Simoncelli EP. Image quality assessment: from error visibility to structural similarity. *IEEE Transactions on Image Processing*. 2004;13(4):600-12. doi:10.1109/TIP.2003.819861.
65. Abdal R, Qin Y, Wonka P. Image2StyleGAN: How to Embed Images Into the StyleGAN Latent Space? 2019 IEEE/CVF International Conference on Computer Vision (ICCV). 2019:4431-40.
66. Lim G, Thombre P, Lee ML, Hsu W, editors. *Generative Data Augmentation for Diabetic Retinopathy Classification*. 2020 IEEE 32nd International Conference on Tools with Artificial Intelligence (ICTAI); 2020 9-11 Nov. 2020.

67. Qasim AB, Ezhov I, Shit S, Schoppe O, Paetzold JC, Sekuboyina A et al. Red-GAN: Attacking class imbalance via conditioned generation. Yet another medical imaging perspective. In: Tal A, Ismail Ben A, Marleen de B, Maxime D, Herve L, Christopher P, editors. Proceedings of the Third Conference on Medical Imaging with Deep Learning; Proceedings of Machine Learning Research: PMLR; 2020. p. 655--68.

Functions of GAN	Methods	Examples
Segmentation	Use discriminator to count the adversarial losses	Conditional GAN for segmenting myocardium and blood pool simultaneously on patients with congenital heart disease[18]
Classification	Use generator and discriminator as a feature extractor, or use discriminator as a classifier	Use semi-supervised GAN for cardiac abnormality classification in chest X-rays[19]
Detection	Train discriminator to detect the lesions	Use GAN to detect brain lesions on MRI[20]
Registration	Train generator to produce transformation parameters or the transformed images	Use spatial transformation network to perform prostate MR to transrectal ultrasound image registration[21]
Image reconstruction	Use cGANs to tackle low spatial resolution, noise contamination and under-sampling	Use pix2pix framework for low dose CT denoising[22]

Table 1: Other applications of GANs in medical imaging besides image synthesis[23]

GAN models	Year	Target imaging modality	Type of GANs	Training datasets	Outputs
Costa et al.[45]	2018	Fundus images	Adversarial autoencoder, image-to-image translation	614 normal fundus images	An end-to-end retinal image synthesis system with corresponding vessel networks
Guibas et al.[46]	2017	Fundus images	Deep convolutional GAN	40 pairs of retinal fundus images and vessel segmentation masks	- F1 accuracy rating of 0.8877 and 0.8988 for synthetically trained and real image trained u-net respectively - Kullback-Leibler (KL) divergence score of 4.759 comparing the synthetic and real images
Zhao et al.[47]	2018	Fundus images	Tub-GAN	10 fundus images	Able to synthesize multiple images with diversity despite a small training set
Yu et al.[48]	2019	Fundus images	Cycle-GAN	101 fundus images (70 glaucomatous eyes and 31 normal eyes)	Higher structural similarity index and peak signal-to-noise ratio than single vessel tree approach

Burlina et al.[49]	2019	AMD fundus images of AMD	ProGAN	133,821 AREDS fundus images	- Accuracy of 53.67% and 59.50% in differentiating real from synthetic images - AUC of DLS trained on synthetic images alone: 0.9235		
Diaz-Pinto et al.[50]	2019	Glaucomatous fundus images	Deep convolutional GAN	86,926 fundus images	- t-SNE plots showing that the synthesized images are closer to the real images than Costa's method and semi-supervised DCGAN model - AUC of 0.9017 in detecting glaucomatous fundus images		
Zhou et al.[51]	2020	DR fundus images	DR-GAN	1,842 images	- Accuracy of 65.8% in differentiating real from synthetic images - Accuracy of 90.46% in DR grading by the classifying model augmented with synthetic images		
Beers et al.[52]	2018	Posterior pole retinal photographs of ROP	ProGAN	5,550 images	AUC of 0.97 comparing to the segmentation maps from real images		
Zheng et al.[53]	2020	OCT images	ProGAN	108,312 OCT images	- Accuracy of discriminating real versus synthetic OCT images: 59.50% and 53.67% - AUC of 0.905 in detecting urgent referrals by DL system trained on synthetic data		
Liu et al.[56]	2019	Post-treatment OCT images	Pix2pixHD	476 pairs of pre- and post-therapeutic OCT images	- sensitivity 84% and specificity 86% in predicting the post-treatment macular classification		
Lee et al.[57]	2021	Post-treatment OCT images	cGAN	15,183 paired OCT B-scans from 241 patients	Predicting	Sensitivity	Specificity
					Intraretinal fluid	33.3%	95.1%
					Subretinal fluid	21.2%	95.1%
					Pigmented epithelial detachment	70.4%	94.6%
Subretinal hyperreflective material	76.5%	94.1%					
Tavakoli et al.[59]	2020	Generate FA from fundus images	cGAN	Pairs of FA and fundus images from 59 patients	Fréchet Inception Distance: 43 Structural similarity measures: 0.67		

Table 2: the summary of studies using image synthesis function of GANs in ophthalmology

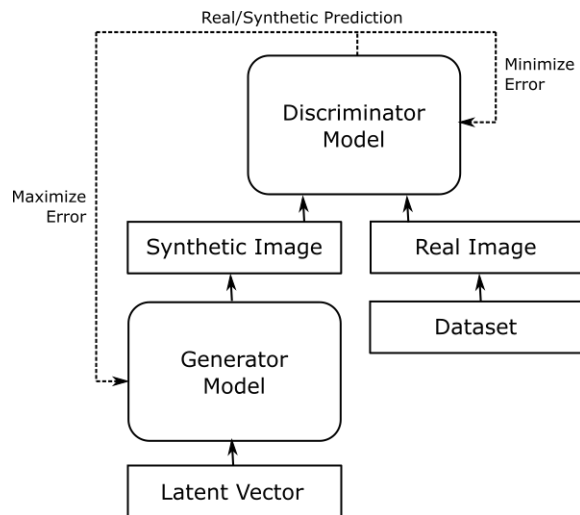


Figure 1: General architecture of a GAN. The generator model and discriminator model are in competition with each other, with the generator model's objective being to produce increasingly realistic synthetic images, and the discriminator model's objective being to distinguish these synthetic images from real images

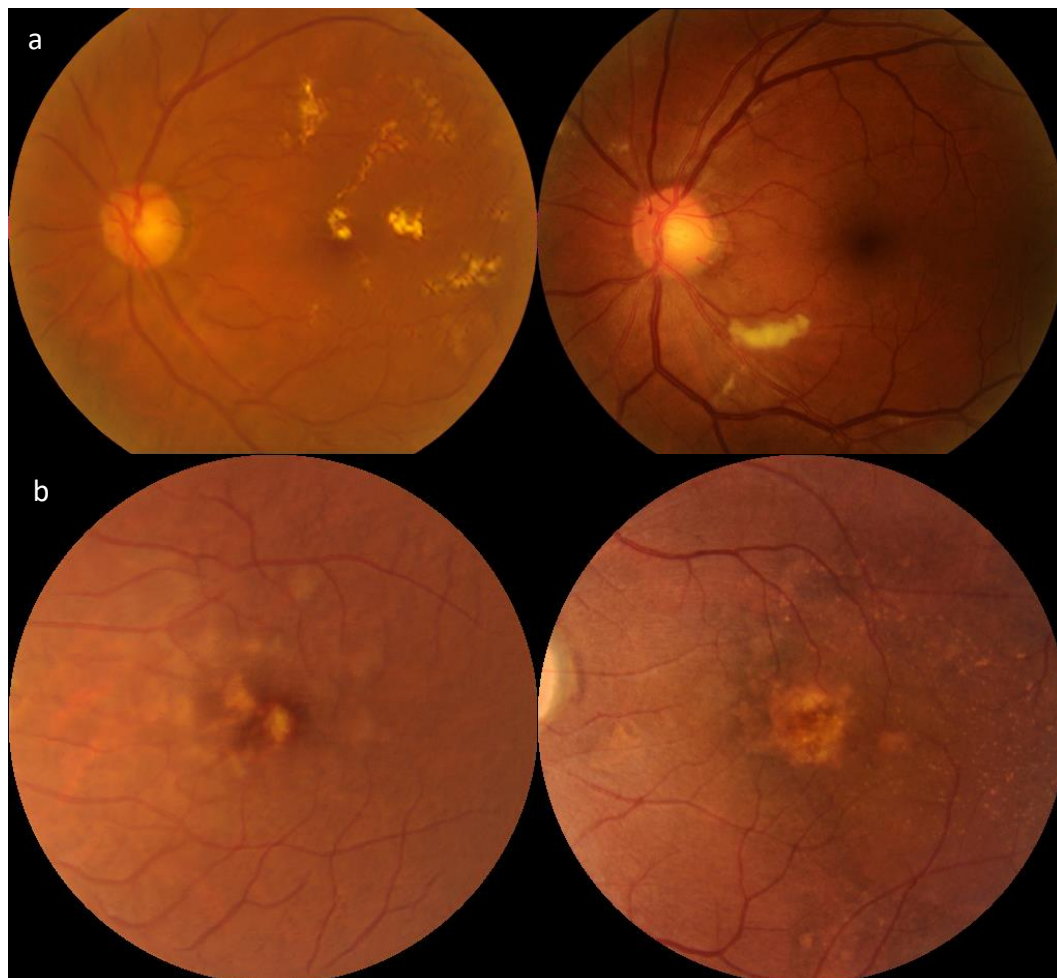


Figure 2: GAN synthesized images (left) compared to real images (right) of DR (a) and AMD (b). The AMD images were preprocessed with macular segmentation.

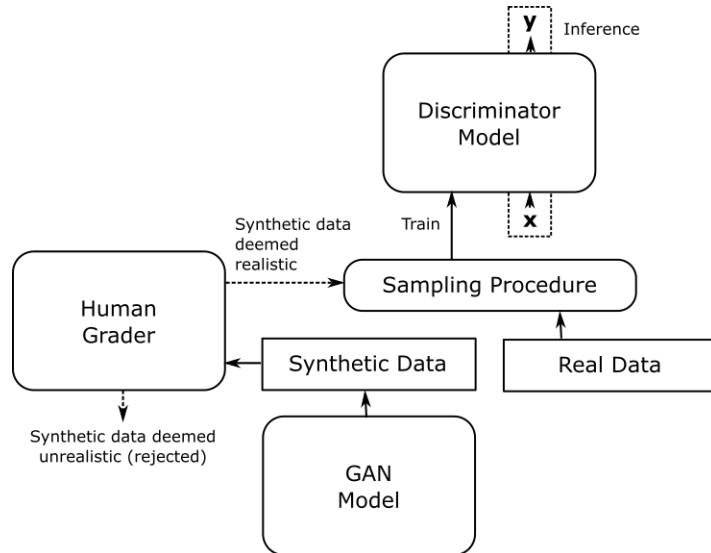


Figure 3: Human-in-the-loop training with a GAN. Human grader(s) arbitrate the generated synthetic data for realism, and the acceptable synthetic data is sampled together with real data to train an improved discriminator model