

This MICCAI paper is the Open Access version, provided by the MICCAI Society. It is identical to the accepted version, except for the format and this watermark; the final published version is available on SpringerLink.

Paired Image Generation with Diffusion-Guided Diffusion Models

Haoxuan Zhang $^{1,2\star},$ Wenju Cui $^{1,2\star},$ Yuzhu Cao $^{1,2},$ Tao Tan 3, Jie Liu 4, Yunsong Peng $^{1,2,5\boxtimes},$ and Jian Zheng $^{1,2\boxtimes}$

- School of Biomedical Engineering (Suzhou), Division of Life Sciences and Medicine, University of Science and Technology of China, Hefei Anhui, China
- ² Suzhou Institute of Biomedical Engineering and Technology, Chinese Academy of Science, Suzhou Jiangsu, China
 - ³ Faculty of Applied Sciences, Macao Polytechnic University, Macao, China
- ⁴ Department of Radiology, Suzhou Municipal Hospital, Suzhou Jiangsu, China
- ⁵ Key Laboratory of Advanced Medical Imaging and Intelligent Computing of Guizhou Province, Guizhou Provincial People's Hospital, Guiyang Guizhou, China

Abstract. The segmentation of mass lesions in digital breast tomosynthesis (DBT) images is very significant for the early screening of breast cancer. However, the high-density breast tissue often leads to high concealment of the mass lesions, which makes manual annotation difficult and time-consuming. As a result, there is a lack of annotated data for model training. Diffusion models are commonly used for data augmentation, but the existing methods face two challenges. First, due to the high concealment of lesions, it is difficult for the model to learn the features of the lesion area. This leads to the low generation quality of the lesion areas, thus limiting the quality of the generated images. Second, existing methods can only generate images and cannot generate corresponding annotations, which restricts the usability of the generated images in supervised training. In this work, we propose a paired image generation method. The method does not require external conditions and can achieve the generation of paired images by training an extra diffusion guider for the conditional diffusion model. During the experimental phase, we generated paired DBT slices and mass lesion masks. Then, we incorporated them into the supervised training process of the mass lesion segmentation task. The experimental results show that our method can improve the generation quality without external conditions. Moreover, it contributes to alleviating the shortage of annotated data, thus enhancing the performance of downstream tasks.

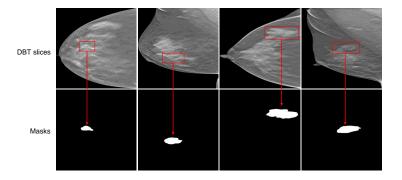
Keywords: Paired image generation Diffusion-guided diffusion models \cdot DBT mass segmentation

1 Introduction

The segmentation of masses in Digital Breast Tomosynthesis (DBT) images is of great significance for the early screening of breast cancer [12]. However, the

^{★:} Equal contribution; ⊠: pys@mail.ustc.edu.cn, zhengj@sibet.ac.cn

dense fibroglandular tissue structure in mammograms often results in highly concealed mass lesions, as illustrated in Fig. 1 [17]. This inherent concealment poses significant challenges for radiologists in manual annotation, leading to labor-intensive workflows and a severe scarcity of high-quality annotated datasets for training deep learning models. To mitigate this data limitation, diffusion models have emerged as a promising solution for synthetic data augmentation in medical imaging [8,16,2]. The generative framework of diffusion models can be broadly categorized into two paradigms. Unconditional generation methods, such as DDPM [5] and DDIM [15], synthesize realistic images from pure noise by progressively denoising random Gaussian distributions through Markov chains. Conditional generation methods, on the other hand, achieve controllable generation by adding guidance signals to the denoising process of the network. These guidance signals can be in the form of classes [1,4], texts [13,11], images [18,10,9], or features [6].



 ${f Fig.\,1.}$ Hidden mass lesions in DBT slices.

However, the application of the above methods in the medical field has two problems. First, for unconditional generation methods, due to the concealment of the lesions, it is difficult for the model to learn the features of the lesion areas. This results in a relatively low generation quality of the lesion areas, thereby restricting the overall quality of the generated images [9]. Second, existing methods only generate images and cannot generate the corresponding annotations. Therefore, conditional generation methods are often used to obtain annotated images, where the annotations are input to guide the generation process. However, the conditions rely on manual input, which limits the diversity of annotations. As a result, the diversity of the generated images is restricted, and the generated images cannot efficiently improve the performance of the model through supervised training in downstream tasks. To tackle these two problems, we propose a Paired Image Generation (PIG) method based on diffusion models. This method does not require external conditional inputs. Instead, it leverages the paired relationship between images as the guiding signal within the generation process. Specifically, we model the generation process of paired images first. Through mathematical derivations, we prove that the unconditional generation process of paired images can be equivalent to two diffusion processes guided mutually. Thus, similar to the classifier-guided diffusion model approach [1], we can achieve the generation of paired images by training an additional diffusion guider for the conditional diffusion model. In the experimental stage, we generated paired DBT slices and mass lesion masks, and added the images generated by different methods to the training of the mass segmentation model. The experimental results show that compared with other generation methods, the images generated by our method have a 15.66 improvement in the FID metric and a 2.24% improvement in the Dice metric. This indicates that our method contributes to enhancing the generation quality and alleviating the shortage of annotated data.

2 Methodology

2.1 Revisiting Diffusion Models

We adopt the diffusion models as the backbone of our proposed PIG method. The diffusion process of the diffusion models consists of the forward process and the reverse process [5,15,9].

The Forward Process. In the forward process, Gaussian noise is gradually added to the image until the image becomes pure noise. The forward process of the diffusion models is Markovian, and the probability distribution q of this process satisfies Eq. (1), where x_0 is the clean image, x_t is the noised image at the t-th time-step, T is the number of time-steps, \mathcal{N} denotes the Gaussian distribution, $\beta_t \in (0,1)$ is the predefined noise schedule and I is an identity matrix of the same shape as x_0 . Eq. (2) represents the single-step noise addition process, and the t-step noise addition process can be obtained through recursion as shown in Eq. (3), where $\bar{\alpha}_t = \prod_{i=1}^t (1 - \beta_i) \to 0$.

$$q(x_{1:T}|x_0) = \prod_{t=1}^{T} q(x_t|x_{t-1}), q(x_t|x_{t-1}) = \mathcal{N}(x_t; \sqrt{1-\beta_t}x_{t-1}, \beta_t I)$$
 (1)

$$x_t = \sqrt{1 - \beta_t} x_{t-1} + \sqrt{\beta_t} \epsilon_t, \epsilon_t \sim \mathcal{N}(0, I)$$
 (2)

$$x_t = \sqrt{\bar{\alpha}_t} x_0 + \sqrt{1 - \bar{\alpha}_t} \epsilon, \epsilon \sim \mathcal{N}(0, I)$$
(3)

The Reverse Process. The reverse process generates images from pure noise by gradually denoising through a trained model θ . Due to the concealment of lesions, in order to improve the generation quality of the lesion part, the mask of the lesion is often input as an external condition c, thereby guiding the model to pay more attention to the generation of the lesion area. The probability distribution p_{θ} of this process satisfies Eq. (4) and Eq. (5), where $p(x_T) = \mathcal{N}(x_T; 0, I)$, $\mu_{\theta}(x_t, c, t) = \sqrt{\bar{\alpha}_{t-1}}x_0 + \sqrt{1 - \bar{\alpha}_{t-1} - \sigma_t^2} \cdot \epsilon_{\theta}(x_t, c, t)$ and σ_t is the predefined

4 H. Zhang et al.

noise variance. Specifically, the model predicts the noise $\epsilon_{\theta}(x_t, c, t)$ and obtains the predicted clean image \hat{x}_0 at the t-th time-step using Eq. (6). Subsequently, x_{t-1} is obtained through Eq. (7) for the next denoising step.

$$p_{\theta}(x_{0:T}|c) = p(x_T) \prod_{t=1}^{T} p_{\theta}(x_{t-1}|x_t, c)$$
(4)

$$p_{\theta}(x_{t-1}|x_t, c) = \mathcal{N}(x_{t-1}; \mu_{\theta}(x_t, c, t), \sigma_t^2 I)$$
(5)

$$\hat{x}_0 = \frac{x_t - \sqrt{1 - \bar{\alpha}_t} \epsilon_{\theta}(x_t, c, t)}{\sqrt{\bar{\alpha}_t}} \tag{6}$$

$$x_{t-1} = \sqrt{\bar{\alpha}_{t-1}}\hat{x}_0 + \sqrt{1 - \bar{\alpha}_{t-1} - \sigma_t^2} \cdot \epsilon_{\theta}(x_t, c, t) + \sigma_t \epsilon_t, \epsilon_t \sim \mathcal{N}(0, I)$$
 (7)

Limitations. Since the condition c relies on external input, manually creating these image conditions is time-consuming, and their diversity is limited. Therefore, we aim to generate c simultaneously when generating images to obtain more diverse lesion annotations. In addition, another advantage of this approach is that it can introduce guiding signals into the image generation process without external condition input, thereby enhancing the generation quality. To achieve this goal, we propose a Paired Image Generation (PIG) method.

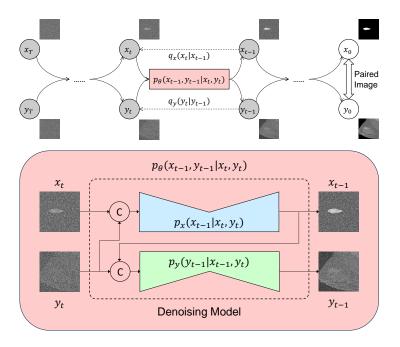


Fig. 2. Graphical model for paired image generation process.

2.2 Paired Image Generation

In this section, we conduct the mathematical modeling for the PIG method based on the diffusion models. The PIG method combines two diffusion processes, and its graphical model is illustrated in Fig. 2.

The Forward Process. During the forward process, we adopt two Markovian noise addition processes similar to Eq. (1)-(3) as shown in Eq. (8)-(10), where (x_0, y_0) represents the clean paired images and $\epsilon_1, \epsilon_2 \sim \mathcal{N}(0, I)$.

$$q(x_{1:T}, y_{1:T}|x_0, y_0) = \prod_{t=1}^{T} q_x(x_t|x_{t-1})q_y(y_t|y_{t-1})$$
(8)

$$q_x(x_t|x_{t-1}) = \mathcal{N}(x_t; \sqrt{1-\beta_t}x_{t-1}, \beta_t I), x_t = \sqrt{\bar{\alpha}_t}x_0 + \sqrt{1-\bar{\alpha}_t}\epsilon_1$$
 (9)

$$q_{y}(y_{t}|y_{t-1}) = \mathcal{N}(y_{t}; \sqrt{1-\beta_{t}}y_{t-1}, \beta_{t}I), y_{t} = \sqrt{\bar{\alpha}_{t}}y_{0} + \sqrt{1-\bar{\alpha}_{t}}\epsilon_{2}$$
 (10)

The Reverse Process. The probability distribution p_{θ} of the joint denoising process satisfies Eq. (11), which can be implemented with two diffusion models x and y to fit p_x and p_y respectively as proved in Eq. (12). In Eq. (11)-(12), $P(\cdot)$ is the abbreviation of probability distribution, $P(x_T) = \mathcal{N}(x_T; 0, I)$ and $P(y_T) = \mathcal{N}(y_T; 0, I)$. Specifically, we can train an additional diffusion guider as model x. It takes y_t as the guiding signal to generate x_{t-1} for guiding model y. Moreover, according to Eq. (6)-(7), in the generation process of x_{t-1} , \hat{x}_0 will be predicted first. Therefore, the conditional diffusion model $p_y(y_{t-1}|x_0, y_t)$ (such as Maisi [2], a generative foundation model pretrained on large-scale unannotated datasets) can be used as model y to avoid the influence of the known noise ϵ_x . During the generation process, this diffusion-guided diffusion model architecture can generate guiding signals for each other, thereby improving the image quality. The training and sampling processes are shown in Alg 1-2, where we set $\sigma_t = 0$.

$$p_{\theta}(x_{0:T}, y_{0:T}) = P(x_T)P(y_T) \prod_{t=1}^{T} p_{\theta}(x_{t-1}, y_{t-1}|x_t, y_t)$$
(11)

Proposition 1. The joint denoising process p_{θ} of the paired images can be implemented with two conditional denoising processes p_x and p_y .

Proof. According to the conditional probability formula and the Bayesian probability formula, the derivation process is as follows.

$$p_{\theta}(x_{t-1}, y_{t-1}|x_{t}, y_{t})$$

$$=p_{x}(x_{t-1}|x_{t}, y_{t}) \cdot P(y_{t-1}|x_{t-1}, x_{t}, y_{t})$$

$$=p_{x}(x_{t-1}|x_{t}, y_{t}) \cdot \frac{p_{y}(y_{t-1}|x_{t-1}, y_{t}) \cdot P(x_{t}|x_{t-1}, y_{t-1}, y_{t})}{P(x_{t}|x_{t-1}, y_{t})}$$

$$=p_{x}(x_{t-1}|x_{t}, y_{t}) \cdot \frac{p_{y}(y_{t-1}|x_{t-1}, y_{t}) \cdot q_{x}(x_{t}|x_{t-1})}{q_{x}(x_{t}|x_{t-1})}$$

$$=p_{x}(x_{t-1}|x_{t}, y_{t}) \cdot p_{y}(y_{t-1}|x_{t-1}, y_{t})$$
(12)

Algorithm 1 The training process of the diffusion guider.

Input: dataset distribution $p(x_0, y_0)$, noise schedule β_t , number of timesteps T Output: trained $model_x$

```
1: repeat
         sample (x_0, y_0) \sim p(x_0, y_0), \epsilon_1 \sim \mathcal{N}(0, I), \epsilon_2 \sim \mathcal{N}(0, I), t \sim Uniform(\{1, ..., T\})
2:
3:
         \bar{\alpha}_t = \prod_{i=1}^t (1 - \beta_i)
         \bar{\alpha}_t = \prod_{i=1}^{r} (1 - \beta_i) 
 x_t = \sqrt{\bar{\alpha}_t} x_0 + \sqrt{1 - \bar{\alpha}_t} \epsilon_1, y_t = \sqrt{\bar{\alpha}_t} y_0 + \sqrt{1 - \bar{\alpha}_t} \epsilon_2
4:
                                                                                                                               ⊳ Eq. (9)-(10)
5:
         \epsilon_x = model_x(x_t, y_t, t)
                                                                                                                           \triangleright p_x(x_{t-1}|x_t,y_t)
6:
         l_x = (\epsilon_1 - \epsilon_x).square().mean()
7:
         update x with \nabla_x l_x
8: until converged
9: return model_x
```

Algorithm 2 The sampling process of paired images.

Input: noise schedule β_t , number of timesteps T, trained $model_x$, trained $model_y$

```
Output: generated images \hat{x}_0, \hat{y}_0
   1: sample x_T \sim \mathcal{N}(0, I), y_T \sim \mathcal{N}(0, I)
   2: for t=T,T-1,...,1 do
                       \bar{\alpha}_t = \prod_{i=1}^t (1 - \beta_i)
                     \hat{x}_{0} = \frac{x_{t} - \sqrt{1 - \bar{\alpha}_{t}} \epsilon_{x}}{\sqrt{\bar{\alpha}_{t}}}, x_{t-1} = \sqrt{\bar{\alpha}_{t-1}} \hat{x}_{0} + \sqrt{1 - \bar{\alpha}_{t-1}} \cdot \epsilon_{x} \qquad \qquad \triangleright \text{Eq. (6)-(7)}
\epsilon_{y} = model_{y}(\hat{x}_{0}, y_{t}, t) \qquad \qquad \triangleright p_{y}(y_{t-1}|\hat{x}_{0}, y_{t}) = p_{y}(y_{t-1}|\frac{x_{t-1} - \sqrt{1 - \bar{\alpha}_{t-1}} \epsilon_{x}}{\sqrt{\bar{\alpha}_{t-1}}}, y_{t})
\hat{y}_{0} = \frac{y_{t} - \sqrt{1 - \bar{\alpha}_{t}} \epsilon_{y}}{\sqrt{\bar{\alpha}_{t}}}, y_{t-1} = \sqrt{\bar{\alpha}_{t-1}} \hat{y}_{0} + \sqrt{1 - \bar{\alpha}_{t-1}} \cdot \epsilon_{y} \qquad \qquad \triangleright \text{Eq. (6)-(7)}
\mathbf{nd for}
    8: end for
   9: return \hat{x}_0, \hat{y}_0
```

3 Experiments

We conducted a paired image generation experiment taking the mass lesion masks (x_0) and the DBT slices (y_0) as paired images. Then, we applied the generated paired images in the supervised training of the DBT mass lesion segmentation task to verify the effectiveness of the generated images.

3.1 **Datasets and Experimental Setup**

DBTMassSeg. This private DBT mass segmentation dataset is sourced from Suzhou Municipal Hospital and Guizhou Provincial People's Hospital⁶. The dataset contains data of 367 patients and the data of each patient includes images from the CC and MLO perspectives and the corresponding mass lesion masks, which are manually annotated by two experienced radiologists. There are a total of 8,723 slices containing mass lesions. In the preprocessing stage, we cropped the blank parts and resized the images to 512×512 .

⁶ The data used in this study were approved by the Ethics Committee of Suzhou Municipal Hospital (No. 2024320) and Guizhou Provincial People's Hospital (No. 2024328). Written informed consent was waived as the data were fully anonymized.

Experimental Setup. We linearly scaled the images to [-1,1] and set the noise schedule β_t to constants increasing linearly from $\beta_1 = 10^{-4}$ to $\beta_T = 0.02$ [5]. The maximum time step T was set to 1024. During the sampling process, we adopted the uniform step sampling method of DDIM [15], with the number of sampling steps set to 256. We used U-Net [14] as the network architecture of the two diffusion models. We set the number of the input channels to 2 and introduced the guiding signal through concatenation in the channel dimension [9,7].

3.2 Results

Focused Generation of Lesion Regions. The unconditional paired image generation process is shown in Fig. 3. It can be seen that during the generation process of the lesion masks \hat{x}_0 and the DBT slices \hat{y}_0 , the shape of the lesion is generated based on the lesion masks at the early stage, and the overall texture is refined in the subsequent process. It indicates that the network is more focused on the generation of lesion regions according to the diffusion guidance.

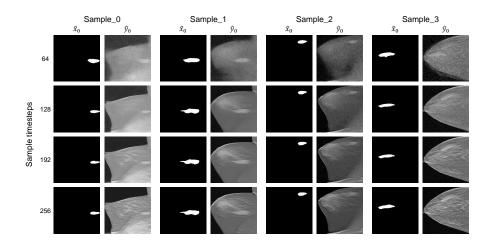


Fig. 3. The progressive generation process of PIG.

Comparative Experiments with Unconditional Diffusion Models. Since the proposed PIG method is a generation approach that does not require external conditions, we conducted comparative experiments with the unconditional diffusion models, including DDPM [5] and DDIM [15]. Specifically, we set the number of sampling steps to 1024 for DDPM and 256 for DDIM, and employed each method to generate 2048 DBT slices respectively. We adopted the Frechet Inception Distance (FID) [3] metric to evaluate the generation quality.

The experimental results presented in Table 1 show that during the image generation process, PIG generates guiding signals simultaneously, which ensures

Table 1. The FID metric of different methods

Method	DDPM [5]	DDIM [15]	PIG
FID↓	31.84	31.54	15.88

that the process requires no additional input conditions and enhances the image generation quality.

Comparative Experiments with Conditional Diffusion Models. Conditional diffusion models can be guided by inputting lesion masks for image generation. Therefore, we evaluated the quality of the generated images through a lesion segmentation task. We used U-Net [14] as the segmentation model.

First, we conducted five-fold cross-validation on the DBTMassSeg dataset. Subsequently, we selected LDM [13], SegGuidedDif [9] and ControlNet [18] as comparative methods and used the existing masks in the training set as guidance to generate 2048 DBT slices respectively. Meanwhile, our proposed PIG method generated 1024, 2048 and 3072 paired images successively. Then we added the images generated by different methods to the supervised training process respectively. We used the Dice, IoU, Precision, and Recall metrics to evaluate the performance of the segmentation task.

Table 2. The experimental results of the mass segmentation task.

Dataset	Dice(%)↑	IoU(%)↑	Precision(%)↑	Recall(%)↑
DBTMassSeg	52.36	41.60	56.14	56.81
$+LDM_{2048} [13]$	56.91	45.88	60.92	60.99
+SegGuidedDif_2048 [9]	58.30	46.99	60.40	64.76
$+ControlNet_2048$ [18]	57.38	46.09	61.53	61.78
$+\mathrm{PIG}_1024$	56.52	45.39	59.86	61.70
$+\mathrm{PIG}_2048$	60.54	49.10	62.92	66.01
$+\mathrm{PIG}_3072$	59.14	47.92	60.65	65.66

The average results of the five-fold cross-validation experiment are presented in Table 2. Here, "+PIG_2048" denotes the addition of 2048 annotated images generated by the PIG method to the original DBTMassSeg dataset and the notations of the other datasets follow the same principle. As shown in the results, since PIG can generate more diverse lesion masks, when generating the same number of DBT slices, the images generated by PIG can help the network learn the features of lesion areas better, thereby enhancing the network's generalization and performance. Additionally, in the experiment, all the generation methods exhibit similar trends. Specifically, the performance peaks after generating about 2048 images, and further increasing the proportion of generated images leads to performance fluctuations. For PIG, the performance dips to its lowest point after generating about 3072 images, which outperforms the highest values achieved by other methods. This indicates the high quality and stability of PIG.

4 Conclusion

We propose an innovative unconditional paired image generation method. First, through mathematical derivations, we prove that the generation process of paired images can be achieved by two diffusion processes guided mutually. Therefore, by training an additional diffusion guider for the conditional diffusion model, we can introduce the guiding signal into the diffusion process without external conditions. The experimental results show that our generation method can improve the generation quality and effectively alleviate the shortage of annotated data.

Acknowledgments. This study was partly supported by National Natural Science Foundation of China (No. 62371449 and No. 82302286), Jiangsu Provincial Key Research and Development Program Social Development Project (No. BE2022720), Guizhou Provincial Basic Research Program (Natural Science) (zk[2025]-513), and Guizhou Provincial Health Commission Science Technology Fund Project (gzwkj2024-474).

Disclosure of Interests. The authors have no competing interests to declare that are relevant to the content of this article.

References

- Dhariwal, P., Nichol, A.: Diffusion models beat gans on image synthesis. In: Ranzato, M., Beygelzimer, A., Dauphin, Y., Liang, P., Vaughan, J.W. (eds.) Advances in Neural Information Processing Systems. vol. 34, pp. 8780–8794. Curran Associates, Inc. (2021), https://proceedings.neurips.cc/paper_files/paper/2021/file/49ad23d1ec9fa4bd8d77d02681df5cfa-Paper.pdf
- Guo, P., Zhao, C., Yang, D., Xu, Z., Nath, V., Tang, Y., Simon, B., Belue, M.J., Harmon, S., Turkbey, B.I., Xu, D.: Maisi: Medical ai for synthetic imaging. ArXiv abs/2409.11169 (2024), https://api.semanticscholar.org/CorpusID:272694143
- Heusel, M., Ramsauer, H., Unterthiner, T., Nessler, B., Hochreiter, S.: Gans trained by a two time-scale update rule converge to a local nash equilibrium. In: Proceedings of the 31st International Conference on Neural Information Processing Systems. p. 6629–6640. NIPS'17, Curran Associates Inc., Red Hook, NY, USA (2017)
- Ho, J.: Classifier-free diffusion guidance. ArXiv abs/2207.12598 (2022), https://api.semanticscholar.org/CorpusID:249145348
- 5. Ho, J., Jain, A., Abbeel, P.: Denoising diffusion probabilistic models. In: Larochelle, H., Ranzato, M., Hadsell, R., Balcan, M., Lin, H. (eds.) Advances in Neural Information Processing Systems. vol. 33, pp. 6840–6851. Curran Associates, Inc. (2020), https://proceedings.neurips.cc/paper_files/paper/2020/file/4c5bcfec8584af0d967f1ab10179ca4b-Paper.pdf
- Hong, S., Lee, G., Jang, W., Kim, S.: Improving Sample Quality of Diffusion Models Using Self-Attention Guidance . In: 2023 IEEE/CVF International Conference on Computer Vision (ICCV). pp. 7428–7437. IEEE Computer Society, Los Alamitos, CA, USA (Oct 2023). https://doi.org/10.1109/ICCV51070.2023.00686, https://doi.ieeecomputersociety.org/10.1109/ICCV51070.2023.00686
- Ji, Y., Chen, Z., Xie, E., Hong, L., Liu, X., Liu, Z., Lu, T., Li, Z., Luo, P.: Ddp: Diffusion model for dense visual prediction. In: Proceedings of the IEEE/CVF International Conference on Computer Vision (ICCV). pp. 21741–21752 (October 2023)

- Kazerouni, A., Aghdam, E.K., Heidari, M., Azad, R., Fayyaz, M., Hacihaliloglu, I., Merhof, D.: Diffusion models in medical imaging: A comprehensive survey. Medical Image Analysis 88, 102846 (2023). https://doi.org/https://doi.org/10.1016/j.media.2023.102846, https://www.sciencedirect.com/science/article/pii/S1361841523001068
- Konz, N., Chen, Y., Dong, H., Mazurowski, M.A.: Anatomically-controllable medical image generation with segmentation-guided diffusion models. In: Linguraru, M.G., Dou, Q., Feragen, A., Giannarou, S., Glocker, B., Lekadir, K., Schnabel, J.A. (eds.) Medical Image Computing and Computer Assisted Intervention MICCAI 2024. pp. 88–98. Springer Nature Switzerland, Cham (2024)
- Liu, X., Park, D.H., Azadi, S., Zhang, G., Chopikyan, A., Hu, Y., Shi, H., Rohrbach, A., Darrell, T.: More control for free! image synthesis with semantic diffusion guidance. In: Proceedings of the IEEE/CVF Winter Conference on Applications of Computer Vision (2023)
- Nichol, A., Dhariwal, P., Ramesh, A., Shyam, P., Mishkin, P., McGrew, B., Sutskever, I., Chen, M.: Glide: Towards photorealistic image generation and editing with text-guided diffusion models. In: International Conference on Machine Learning (2021), https://api.semanticscholar.org/CorpusID:245335086
- Rafferty, E.A., Durand, M.A., Conant, E.F., Copit, D.S., Friedewald, S.M., Plecha, D.M., Miller, D.P.: Breast cancer screening using tomosynthesis and digital mammography in dense and nondense breasts. JAMA 315(16), 1784–1786 (04 2016). https://doi.org/10.1001/jama.2016.1708, https://doi.org/10.1001/jama.2016.1708
- Rombach, R., Blattmann, A., Lorenz, D., Esser, P., Ommer, B.: High-resolution image synthesis with latent diffusion models. In: 2022 IEEE/CVF Conference on Computer Vision and Pattern Recognition (CVPR). pp. 10674–10685 (2022). https://doi.org/10.1109/CVPR52688.2022.01042
- Ronneberger, O., Fischer, P., Brox, T.: U-net: Convolutional networks for biomedical image segmentation. In: Navab, N., Hornegger, J., Wells, W.M., Frangi, A.F. (eds.) Medical Image Computing and Computer-Assisted Intervention MICCAI 2015. pp. 234–241. Springer International Publishing, Cham (2015)
- 15. Song, J., Meng, C., Ermon, S.: Denoising diffusion implicit models. arXiv:2010.02502 (October 2020), https://arxiv.org/abs/2010.02502
- 16. Wang, J., Wang, K., Yu, Y., Lu, Y., Xiao, W., Sun, Z., Liu, F., Zou, Z., Gao, Y., Yang, L., Zhou, H.Y., Miao, H., Zhao, W., Huang, L., Zeng, L., Guo, R., Chong, I., Deng, B.J., Cheng, L., Chen, X., Luo, J., Zhu, M., Baptista-Hon, D.T., Monteiro, O., Li, M., Ke, Y., Li, J., Zeng, S., Guan, T., Zeng, J., Xue, K., Oermann, E.K., Luo, H., Yin, Y., Zhang, K., Qu, J.: Self-improving generative foundation model for synthetic medical image generation and clinical applications. Nature medicine (2024), https://api.semanticscholar.org/CorpusID:274655051
- 17. Xiong, X., Wang, C., Li, W., Li, G.: Semi- and weakly-supervised learning for mammogram mass segmentation with limited annotations. 2024 IEEE International Symposium on Biomedical Imaging (ISBI) pp. 1–5 (2024), https://api.semanticscholar.org/CorpusID:268385465
- 18. Zhang, L., Rao, A., Agrawala, M.: Adding conditional control to text-to-image diffusion models. In: Proceedings of the IEEE/CVF International Conference on Computer Vision (ICCV). pp. 3836–3847 (October 2023)