

Deep Learning and Radiomics in Triple-Negative Breast Cancer: Predicting Long-Term Prognosis and Clinical Outcomes

Chen Cheng^{1,*}, Yan Wang^{2,3,*}, Jine Zhao^{4,*}, Di Wu¹, Hong Li⁵, Hongyan Zhao¹

¹Department of Ultrasound, Lianyungang Traditional Chinese Medicine Hospital, Lianyungang, 222004, People's Republic of China; ²Department of Ultrasound, Lianyungang Municipal Oriental Hospital, Lianyungang, 222046, People's Republic of China; ³Department of Ultrasound, Xuzhou Medical University Affiliated Hospital, Lianyungang, Jiangsu, 222061, People's Republic of China; ⁴Department of Ultrasound, Donghai County People's Hospital, Lianyungang, Jiangsu, 222300, People's Republic of China; ⁵Department of Ultrasound, the First People's Hospital of Lianyungang, Lianyungang, Jiangsu, 222061, People's Republic of China

*These authors contributed equally to this work

Correspondence: Hongyan Zhao, Department of Ultrasound, Lianyungang Traditional Chinese Medicine Hospital, No. 160, Chaoyang Middle Road, Haizhou District, Lianyungang, 222004, People's Republic of China, Tel +86 0518-85574003, Email zhaohongyanzhy01@126.com

Abstract: Triple-negative breast cancer (TNBC) is a unique breast cancer subtype characterized by the lack of estrogen receptor (ER), progesterone receptor (PR), and human epidermal growth factor receptor 2 (HER2) expression in tumor cells. TNBC represents about 15% to 20% of all breast cancers and is aggressive and highly malignant. Currently, TNBC diagnosis primarily depends on pathological examination, while treatment efficacy is assessed through imaging, biomarker detection, pathological evaluation, and clinical symptom improvement. Among these, biomarker detection and pathological assessments are invasive, time-intensive procedures that may be difficult for patients with severe comorbidities and high complication risks. Thus, there is an urgent need for new, supportive tools in TNBC diagnosis and treatment. Deep learning and radiomics techniques represent advanced machine learning methodologies and are also emerging outcomes in the medical-engineering field in recent years. They are extensions of conventional imaging diagnostic methods and have demonstrated tremendous potential in image segmentation, reconstruction, recognition, and classification. These techniques hold certain application prospects for the diagnosis of TNBC, assessment of treatment response, and long-term prognosis prediction. This article reviews recent progress in the application of deep learning, ultrasound, MRI, and radiomics for TNBC diagnosis and treatment, based on research from both domestic and international scholars.

Keywords: deep learning, MRI, radiomics, triple negative breast cancer, ultrasound

Breast cancer is among the most prevalent cancers affecting women worldwide, posing a significant threat to women's health. According to the World Health Organization (WHO), it is the most frequently diagnosed cancer globally and ranks fifth in cancer-related mortality.¹ Triple-negative breast cancer (TNBC) represents a distinct subtype, accounting for about 10 to 20% of all breast cancer cases.² TNBC is defined by the absence of ER, PR, and HER2 expression in tumor cells, making these patients unresponsive to hormone therapy and HER2-targeted treatments.³ Consequently, chemotherapy remains the primary treatment option.⁴ Numerous studies have indicated that TNBC generally carries a higher recurrence rate and poorer prognosis compared to other breast cancer subtypes.⁵⁻⁷ Furthermore, TNBC often occurs in younger women and is associated with a higher prevalence of BRCA1 gene mutations.⁸ Some researchers suggest that this may partly explain the higher incidence of TNBC in certain populations, such as Ashkenazi Jews.^{9,10}

With the rapid advancement of artificial intelligence (AI), interest is surging in its medical applications to aid clinical decision-making. AI-driven techniques like deep learning and radiomics, which combine medical imaging with digital analysis, are increasingly being utilized to extract disease-specific information that is difficult to assess visually. In recent years, these methods have become significant research focal points.¹¹ In this context, quantitative analysis of multimodal,

multidimensional data is essential for evaluating the spatiotemporal characteristics of various tissues, organs, and their surrounding microenvironments.¹² This article reviews recent research in deep learning and radiomics, focusing on their applications in the differential diagnosis of TNBC, assessing neoadjuvant chemotherapy (NAC) efficacy, and predicting long-term prognosis post-NAC. The objective is to examine the clinical potential and future applications of these technologies.

The literature retrieval strategy employed in this study extensively covered multiple authoritative databases, including PubMed, Web of Science, and China National Knowledge Infrastructure (CNKI), to ensure the comprehensiveness and diversity of the retrieval results. During the retrieval process, we meticulously selected a series of keywords that could be used independently or in combination to maximize the capture of relevant information. These keywords encompassed “ultrasound”, “MRI”, “radiomics”, “deep learning”, “triple-negative breast cancer/breast cancer”, and “neoadjuvant chemotherapy.” Our retrieval focused on literature published between 2016 and 2023, reflecting the latest advancements and research findings in this field in recent years.

Overview of Deep Learning and Radiomics

Radiomics involves extracting numerous features from medical imaging data to quantify tumor heterogeneity using advanced image processing and analysis techniques.¹³ These features include tumor shape characteristics, first-order statistical properties, and texture features, all of which are not discernible through conventional imaging observation. Radiomics enhances accuracy in disease detection, diagnosis, classification, and prognostic evaluation.¹⁴ By analyzing radiological images of tumors and surrounding tissues, it reveals subtle biological processes that provide insight into tumor behavior and treatment response.¹⁵ The radiomics workflow typically includes the following steps:¹⁶ ① Image Acquisition: High-quality medical images are obtained using modalities such as CT, MRI, and PET. ② Image Preprocessing: This stage involves standardizing images, reducing noise, and applying corrections to improve analysis quality. ③ Region of Interest (ROI) Selection and Segmentation: This step requires precisely delineating the area for analysis, usually focusing on the tumor region. ④ Feature Extraction: Quantitative features are extracted from the ROIs, capturing details of shape, intensity, and texture. ⑤ Feature Analysis: Statistical and machine learning techniques are applied to these features, facilitating disease characterization and aiding in predictions of treatment effectiveness and patient prognosis.

Deep Learning (DL) is a subset of machine learning that employs a multi-layer architecture of artificial neural networks, mimicking the information processing capabilities of the human brain.¹⁷ This enables the model to make predictions and decisions by learning intricate patterns and features from extensive datasets. DL not only improves the accuracy and efficiency of feature extraction but also reveals potential image features that may be overlooked during conventional observation. With technological advancements, deep learning has become a pivotal force in advancing medical imaging, providing robust support for accurate diagnoses and personalized tumor treatments.^{18,19}

In this framework, deep learning acquires features by conducting end-to-end training on coarse regions containing the targets.^{20,21} Upon completing the training, depth features can either be integrated with semantic features for radiomics analysis or used directly for model predictions. In medical applications, convolutional neural network architectures—such as ResNet, GoogLeNet, and DenseNet—are frequently employed for tasks involving image and video processing. Over the past decade, deep learning has accelerated the development and integration of artificial intelligence technologies in healthcare. However, challenges persist, including the demand for substantial data and computational resources, along with concerns regarding model interpretability and transparency that necessitate urgent attention.²²

Application and Comparison of Deep Learning and Radiomics in the Diagnosis and Treatment of TNBC

Recent research has established that deep learning and radiomics hold significant promise in predicting the biological characteristics of TNBC.²³ These methodologies aid clinicians in accurately identifying the tumor type and stage, choosing the most suitable treatment options, monitoring treatment effectiveness, evaluating patient survival probabilities, and delivering more personalized treatment recommendations. Given that ultrasound is the most widely utilized

method for breast examination and MRI provides the highest resolution imaging in clinical settings, this article reviews the implementation of artificial intelligence technologies in these two imaging modalities.

Application of Deep Learning and Radiomics in the Diagnosis of TNBC

The current application of deep learning and radiomics in diagnosing and classifying TNBC has showed significant diagnostic efficacy.^{24–37} For detailed information, refer to [Table 1](#). In a study by Sha et al, involving 1223 patients, the diagnostic value of MRI radiomics for TNBC was evaluated, revealing a combined sensitivity of 0.72 and specificity of 0.91, with an area under the curve (AUC) of 0.88. Ma et al (2021) explored the use of quantitative features extracted from dynamic contrast-enhanced MRI (DCE-MRI) images of three-dimensional tumor volumes to differentiate TNBC from non-TNBC.^{24,25} The findings indicated that a radiomics model based on 15 features achieved optimal performance,

Table 1 Deep Learning and Radiomics in the Diagnosis and Management of TNBC

Clinical Events	Test methods	Accuracy	Sensitivity	Specificity	AUC/C-index	Author (Publication Time)
Diagnosis of TNBC	MRI, Radiomics	—	0.72	0.91	0.88	Sha YS et al, 2022 ²⁴
Identify TNBC from NTNBC	DCE-MRI, Radiomics	—	—	—	AUC was 0.741 for cross-validation and 0.867 for independent testing cohorts	Mingming Ma et al, 2021 ²⁵
Molecular typing of breast cancer	Conventional ultrasound combined with contrast-enhanced ultrasound	0.854	—	—	—	Gong X et al, 2023 ²⁶
Identify TNBC from NTNBC	Ultrasound deep learning (Resnet50)	0.889	0.875	0.9	0.9	Ye, H et al, 2021 ²⁷
Diagnosis of TNBC	MRI deep learning	—	—	—	0.944	Yin HL et al, 2023 ²⁸
Predicting pCR status after NAC in patients with TNBC	DWI + DCE-MRI, deep learning	—	—	—	The AUC was 0.97 ± 0.04 in the training group and 0.82 ± 0.10 in the internal validation group, while it was 0.86 ± 0.03 in the independent validation group and 0.83 ± 0.02 in the prospective blinded trial group AUC=0.94(95% CI:0.91–0.97)	Zhou Z et al, 2023 ²⁹
Predicting pCR status after NAC in advanced breast cancer	Ultrasound, deep learning	—	—	—	—	Jiang M et al, 2021 ³¹
Predicting response to NAC therapy in patients with TNBC	Radiomics	—	—	—	—	—
Predicting recurrence within 3 years after NAC in patients with TNBC	MRI, Radiomics	—	—	—	AUC = 0.78 and 0.72 in training and testing cohorts The radiomics model, which integrated MRI features obtained before and after neoadjuvant chemotherapy (NAC), achieved an AUC of 0.963 in the test group and 0.933 in the validation group	Hwang KP et al, 2023 ³⁰ Mingming Ma et al, 2022 ³⁴

(Continued)

Table 1 (Continued).

Clinical Events	Test methods	Accuracy	Sensitivity	Specificity	AUC/C-index	Author (Publication Time)
Predicting the likelihood of long-term recurrence and metastasis following neoadjuvant therapy in TNBC	DCE-MRI, Radiomics	87.5% in training group and 82.9% in validation group	—	—	AUC (0.917 for training group and 0.859 for validation group)	Xia B et al, 2021 ³⁶
Predicting disease-free survival in triple-negative breast cancer	Ultrasound, Radiomics	—	—	—	The C-index of the model was 0.75 (95% CI = 0.72 to 0.78) in the training set and 0.73 (95% CI = 0.71 to 0.75) in the validation set	Yu F et al, 2021 ³⁷

with AUCs of 0.741 during cross-validation and 0.867 in an independent test cohort. Gong et al combined conventional ultrasound with contrast-enhanced ultrasound radiomics to predict breast cancer molecular subtypes, showing that the accuracy of the combined ultrasound model surpassed that of the conventional model (85.4% vs 81.3%, $p < 0.01$).²⁶ These results highlight the substantial potential of MRI and ultrasound radiomics in the differential diagnosis and classification of TNBC.

Ye et al conducted a retrospective analysis of 934 ultrasound images of breast malignant tumors, which included 110 cases of TNBC and 824 cases of non-triple-negative breast cancer (NTNBC).²⁷ They employed a ResNet50 deep convolutional neural network for the analysis, resulting in an AUC of 0.900 for differentiating TNBC from NTNBC. The model showed an accuracy of 88.89%, with a sensitivity of 87.5% and specificity of 90.00%. Yin et al assessed the effectiveness of a deep learning model using multiparametric MRI to enhance radiologists' accuracy in the differential diagnosis of TNBC.²⁸ Their results showed that the deep learning approach performed exceptionally well in the validation group, achieving an AUC of 0.944. With the integration of artificial intelligence, the diagnostic AUC improved from 0.833 to 0.885 and from 0.823 to 0.876 for the two primary radiologists, respectively. For the two senior radiologists, the AUC increased from 0.901 and 0.950 to 0.925 and 0.975, respectively. These findings suggest that imaging combined with deep learning offers an automated and improved diagnostic process, making them promising noninvasive clinical tools for diagnosing TNBC. The incorporation of deep learning and radiomics techniques can substantially enhance diagnostic efficiency, potentially leading to earlier and more accurate detection and classification of TNBC.

To thoroughly evaluate the effectiveness of deep learning in ultrasound and MRI, alongside the application of radiomics in diagnosing TNBC, it is clear that the efficacy of deep learning and radiomics in MRI applications generally exceeds that of ultrasound diagnostics. Nevertheless, variations in study populations and the diagnostic criteria utilized across different research efforts highlight the need for further standardization in comparative analyses of the diagnostic efficacy of these two methodologies.

Deep Learning and Radiomics in TNBC Therapy Evaluation

Accurate evaluation of the impact of NAC for breast cancer is essential for determining appropriate surgical approaches and guiding the extent of breast cancer resection. Currently, the assessment of treatment efficacy in TNBC predominantly relies on biomarker detection, imaging studies, and clinical observations. Numerous studies suggest that deep learning and imaging-omics techniques play a significant role in optimizing treatment options and monitoring efficacy in TNBC patients. Zhou et al explored the effectiveness of applying deep learning to DCE-MRI and diffusion-weighted imaging (DWI) to predict pathological complete response (pCR) status in patients with TNBC.²⁹ Their study utilized images from 130 patients with TNBC to develop the models, achieving impressive results with an AUC of 0.97 ± 0.04 in the training group and 0.82 ± 0.10 in the internal validation group. In an independent validation cohort of 32 cases, the model

maintained a robust AUC of 0.86 ± 0.03 . Additionally, in another prospective blinded trial involving 48 patients, the model yielded an AUC of 0.83 ± 0.02 . These findings indicate that multiparametric MRI-based deep learning can accurately predict pCR or non-pCR in patients with TNBC at an early stage. Hwang et al examined the utility of a radiomics model based on quantitative atlases derived from MRI to predict the response to NAC in patients with TNBC.³⁰ The multivariate radiomics model developed during mid-treatment achieved AUC values of 0.78 in the training cohort and 0.72 in the testing cohort. Although these results indicated moderate predictive power, they suggest that MRI-based radiological features obtained during mid-treatment may be valuable in identifying early responders to NAC in patients with TNBC.

Recently, numerous studies have employed multimodal methods to construct predictive models for pCR status in patients with TNBC. Jiang et al developed and validated a deep learning radiomic nomogram (DLRN) that utilized ultrasound and clinical data to assess pCR status in patients with locally advanced breast cancer following NAC.³¹ The findings indicated that the DLRN accurately predicted pCR status, achieving an AUC of 0.94 in the validation cohort, with a 95% confidence interval (CI) of 0.91 to 0.97. Furthermore, the model's calibration was proven to be effective.

Hacking et al and Jimenez et al incorporated MRI radiomics alongside pathological sections and tumor-infiltrating lymphocyte (TIL) levels to predict pCR after neoadjuvant systemic therapy.^{32,33} Their studies exhibited promising results for the clinical application of these approaches in accurately assessing treatment response.

In summary, deep learning and radiomics enable the automatic extraction of a wide range of quantitative features from medical images in a non-invasive manner, enhancing the evaluation of treatment efficacy in TNBC. The integration of fusion models, which incorporate multi-learning fusion or various machine learning techniques, can further refine the prediction of patients with TNBC responses to NAC. This approach helps tailor treatment decisions by identifying patients who may not benefit from standard care, enabling the implementation of more appropriate treatment regimens and ultimately improving treatment effectiveness and patient survival.

Application of Deep Learning and Radiomics in Predicting the Long-Term Prognosis of TNBC

In recent years, some researchers have begun to explore the application value of radiomics in predicting the long-term prognosis of TNBC, conducting preliminary investigations in this area. Ma et al (2022) developed a radiomics prediction model for systemic recurrence after NAC in TNBC based on automated segmented MRI.³⁴ The findings indicated that the AUC for the clinical model was 0.747 in the training set and 0.737 in the validation group for predicting systemic recurrence. In contrast, the AUCs for the radiomics model, which incorporated MRI features before NAC, after NAC, and both before and after NAC, were significantly higher, achieving values of 0.879, 0.91, and 0.963 in the training set, and 0.814, 0.802, and 0.933 in the validation set, respectively. These results show that the radiomics model based on a combination of MRI features before and after NAC is effective in predicting whether patients with TNBC will experience systemic recurrence within three years after treatment. This model could aid in the non-invasive identification of high-risk patients with TNBC at the risk of relapse after NAC, thus enhancing follow-up and treatment strategies and ultimately improving their prognosis.

Jiang et al focused on predicting TNBC prognosis by extracting imaging features related to peri-tumoral heterogeneity using DCE-MRI.³⁵ This model successfully predicted recurrence-free survival ($P = 0.01$) and overall survival ($P = 0.004$) in patients with TNBC. Bingqing also utilized radiomics features extracted from DCE-MRI prior to treatment, incorporating time-domain characteristics to construct a radiomics model aimed at predicting long-term recurrence and metastasis risk following neoadjuvant therapy.³⁶ The optimal model achieved an AUC of 0.917 in the training group and 0.859 in the validation group, with accuracy rates of 87.5% in the training group and 82.9% in the validation group. This study concluded that the DCE-MRI machine learning radiomics model, enhanced with time-domain characteristics, is valuable for predicting the long-term prognosis of patients with TNBC undergoing NAC.

While there are few radiomics studies in the field of ultrasound, Yu et al conducted a multicenter application study utilizing ultrasound radiomics features to predict disease-free survival (DFS) in TNBC.³⁷ This study identified ten significant radiomics characteristics to construct omics signatures, which were found to be independent risk factors for

predicting DFS in both the training and validation sets ($p < 0.05$). The resulting clinic-radiomics model, which included axillary lymph node stage, Ki-67 index, and radiomics signatures, achieved a concordance index (C-index) of 0.75 (95% CI: 0.72 to 0.78) in the training set and 0.73 (95% CI: 0.71 to 0.75) in the validation set, showing moderate predictive efficacy.

After a thorough review of the existing domestic and international literature, we observed a limited number of studies focusing on the application of deep learning in predicting long-term prognosis for TNBC. This scarcity may be attributed to the complex and heterogeneous biological characteristics of TNBC, which complicate the extraction of imaging features that are closely linked to long-term outcomes. Despite significant advancements in deep learning for medical imaging analysis, its application in prognostic predictions for specific cancers remains largely exploratory. To enhance the validity and reliability of these models, more comprehensive clinical data and long-term follow-up studies are essential. Therefore, the current lack of research not only highlights the existing challenges in this domain but also suggests potential avenues for future investigations.

Comparative Analysis of Diagnostic and Predictive Efficacy of Deep Learning and Radiomics

There is currently no consensus on whether deep learning or radiomics demonstrates superior diagnostic or predictive power. The predictive accuracy of these two approaches is influenced by various factors, including individual patient differences, the type and volume of available data, and the methodologies used for model training. In certain contexts, models that utilize high-precision radiomic features and undergo iterative dimensionality reduction may provide better interpretability and performance.

In recent years, multimodal strategies that combine radiomics with deep learning have shown promise in enhancing predictive accuracy. The DLRN capitalizes on the strengths of both methods, improving the accuracy of prognostic predictions through the synergistic integration of automatically extracted features and radiomic characteristics derived from deep learning.^{38,39}

Comparative Analysis of the Diagnostic and Predictive Efficacy of Deep Learning and Radiomics in HER-2 Positive Vs HER-2 Negative Breast Cancer

As another subtype of breast cancer with poor prognosis, HER-2 positive breast cancer is characterized by over-expression or amplification of the HER-2 receptor on the surface of tumor cells. Compared to other subtypes such as ER+/PR+/HER-2- and ER+/PR-/HER-2-, HER-2 positive breast cancer exhibits a higher risk of recurrence and metastasis. Therefore, imaging-assisted diagnosis and monitoring are particularly important in the management of this type of cancer. Existing studies have used deep learning algorithms to analyze breast MRI images to differentiate between HER-2 positive and HER-2 negative breast cancer. The results show that deep learning models achieve an accuracy rate of about 80% in identifying HER-2 positive breast cancer.^{40,41} A literature search from 2019 to 2024 reveals that there are still very few studies on the evaluation of the efficacy of neoadjuvant chemotherapy for HER-2 positive breast cancer using deep learning and radiomics. Kim SY's team⁴² developed a CNN deep learning model based on post-NAC dynamic contrast-enhanced MRI and clinical data to identify pathological complete remission in HER2 positive and triple-negative breast cancer. The study shows that the model established after cropping lesion images has certain diagnostic potential, but further optimization and external validation are needed to improve accuracy. It is evident that machine learning techniques have significant application potential not only in the clinical diagnosis and treatment of TNBC but also in the diagnosis and prognostic evaluation of other breast cancer subtypes. However, further development and exploration are still required.

Conclusion

Deep learning and radiomics are rapidly advancing in the research on diagnosis and treatment of triple-negative breast cancer (TNBC), demonstrating immense potential, particularly in playing a crucial role in assessing tumor heterogeneity. These advanced technologies can provide more comprehensive and detailed tumor information, effectively compensating

for the limitations of traditional biopsy sampling. By deeply analyzing the relationship between radiomic features and tumor biological behavior, and developing more sophisticated machine learning algorithms, researchers can more accurately detect and analyze tumor heterogeneity, providing robust support for clinical diagnosis and treatment decisions. However, there remains a notable lack of research focused on predicting long-term prognosis. Meanwhile, deep learning and radiomics also face the following challenges in the diagnosis and treatment of TNBC.

Key challenges include the application of fusion methods, such as the DLRN, in diagnosing and treating TNBC. Additionally, there is a pressing need for the comprehensive acquisition and analysis of multimodal imaging data, particularly the integration of ultrasound and MRI using artificial intelligence. Furthermore, challenges related to artificial intelligence data acquisition and processing, the lack of interpretability of algorithms, and the limitations of clinical applications remain significant hurdles.

To address these challenges, it is essential to promote interdisciplinary collaboration, integrate multimodal data, and drive technological advancements. Such concerted efforts can systematically tackle the obstacles currently faced, unlocking the full potential of deep learning and radiomics and enabling their transformative impact on the precision diagnosis and treatment of TNBC.

Abbreviations

MRI, Magnetic Resonance Imaging; US, Ultrasound; TNBC, Triple-Negative Breast Cancer; ER, Estrogen Receptor; PR, Progesterone Receptor; HER-2, Human Epidermal Growth Factor-2; NTNBC, Non-Triple-Negative Breast Cancer; NAC, Neoadjuvant Chemotherapy; ROI, Region Of Interest; CNN, Convolutional Neural Network; ROC, Receiver Operating Characteristic; AUC, Area Under The Curve; DCA, Decision Curve Analysis; DLRN, Deep Learning Radiomics Nomogram; pCR, Pathological Complete Remission; npCR, Non-Pathological Complete Remission; DL, Deep Learning; AI, Artificial Intelligence.

Data Sharing Statement

All data generated or analysed during this study are included in this article. Further enquiries can be directed to the corresponding author.

Ethics Approval and Consent to Participate

The study is a literature review and does not require ethics.

Consent for Publication

Not applicable

Acknowledgments

We would like to acknowledge the hard and dedicated work of all the staff that implemented the intervention and evaluation components of the study.

Funding

2023 Lianyungang City “521 Project” funded Young talent project (LYG06521202387) 2022 Lianyungang Health Youth Science and Technology Project (QN202204). 2024 The Project Supported by the Science Foundation of Kangda College, Nanjing Medical University (KD2024KYJJ082). 2024 Lianyungang City Cancer Prevention and Control Science and Technology Development Plan Project (QN202426).

Disclosure

The authors declare that they have no competing interests.

References

1. Mridha MF, Hamid MA, Monowar MM, et al. A comprehensive survey on deep-learning-based breast cancer diagnosis. *Cancers (Basel)*. 2021;13(23):6116. PMID: 34885225; PMCID: PMC8656730.doi:10.3390/cancers13236116
2. Sung H, Ferlay J, Siegel RL, et al. Global Cancer Statistics 2020: GLOBOCAN estimates of incidence and mortality worldwide for 36 cancers in 185 countries. *CA Cancer J Clin*. 2021;71(3):209–249. PMID: 33538338.doi:10.3322/caac.21660
3. Keenan TE, Tolaney SM. Role of immunotherapy in triple-negative breast cancer. *J Natl Compr Canc Netw*. 2020;18(4):479–489. PMID: 32259782.doi:10.6004/jnccn.2020.7554
4. Ge J, Zuo W, Chen Y, Shao Z, Yu K. The advance of adjuvant treatment for triple-negative breast cancer. *Cancer Biol Med*. 2021;19(2):187–201. PMID: 34448553; PMCID: PMC8832962.doi:10.20892/j.issn.2095-3941.2020.0752
5. Dent R, Trudeau M, Pritchard KI, et al. Triple-negative breast cancer: clinical features and patterns of recurrence. *Clin Cancer Res*. 2007;13(15 Pt 1):4429–4434. PMID: 17671126.doi:10.1158/1078-0432.CCR-06-3045
6. Nunnery SE, Mayer IA, Balko JM. Triple-negative breast cancer: breast tumors with an identity crisis. *Cancer J*. 2021;27(1):2–7. PMID: 33475287; PMCID: PMC8109153.doi:10.1097/PPO.0000000000000494
7. Yagata H, Kajiyama Y, Yamauchi H. Current strategy for triple-negative breast cancer: appropriate combination of surgery, radiation, and chemotherapy. *Breast Cancer*. 2011;18(3):165–173. Erratum in: *Breast Cancer*. 2012;19(4):369. PMID: 21290263.doi:10.1007/s12282-011-0254-9
8. Foulkes WD, Smith IE, Reis-Filho JS. Triple-negative breast cancer. *N Engl J Med*. 2010;363(20):1938–1948. PMID: 21067385.doi:10.1056/NEJMra1001389
9. Greenup R, Buchanan A, Lorizio W, et al. Prevalence of BRCA mutations among women with triple-negative breast cancer (TNBC) in a genetic counseling cohort. *Ann Surg Oncol*. 2013;20(10):3254–3258. PMID: 23975317.doi:10.1245/s10434-013-3205-1
10. Lee E, McKean-Cowdin R, Ma H, et al. Characteristics of triple-negative breast cancer in patients with a BRCA1 mutation: results from a population-based study of young women. *J Clin Oncol*. 2011;29(33):4373–4380. PMID: 22010008; PMCID: PMC3221522.doi:10.1200/JCO.2010.33.6446
11. Mireştean CC, Volovăţ C, Iancu RI, Iancu DPT. Radiomics in triple negative breast cancer: new horizons in an aggressive subtype of the disease. *J Clin Med*. 2022;11(3):616. PMID: 35160069; PMCID: PMC8836903.doi:10.3390/jcm11030616
12. Corredor G, Bharadwaj S, Pathak T, Viswanathan VS, Toro P, Madabhushi A. A review of ai-based radiomics and computational pathology approaches in triple-negative breast cancer: current applications and perspectives. *Clin Breast Cancer*. 2023;23(8):800–812. PMID: 37380569; PMCID: PMC10733554.doi:10.1016/j.clbc.2023.06.004
13. Mayerhoefer ME, Materka A, Langs G, et al. Introduction to Radiomics. *J Nucl Med*. 2020;61(4):488–495. PMID: 32060219; PMCID: PMC9374044.doi:10.2967/jnumed.118.222893
14. Lambin P, Leijenaar RTH, Deist TM, et al. Radiomics: the bridge between medical imaging and personalized medicine. *Nat Rev Clin Oncol*. 2017;14(12):749–762. PMID: 28975929.doi:10.1038/nrclinonc.2017.141
15. Aerts HJ. The potential of radiomic-based phenotyping in precision medicine: a review. *JAMA Oncol*. 2016;2(12):1636–1642. PMID: 27541161.doi:10.1001/jamaoncol.2016.2631
16. van Timmeren JE, Cester D, Tanadini-Lang S, Alkadhi H, Baessler B. Radiomics in medical imaging—“how-to” guide and critical reflection. *Insights Imag*. 2020;11(1):91. PMID: 32785796; PMCID: PMC7423816.doi:10.1186/s13244-020-00887-2
17. Chan HP, Samala RK, Hadjiiski LM, Zhou C. Deep learning in medical image analysis. *Adv Exp Med Biol*. 2020;1213:3–21. PMID: 32030660; PMCID: PMC7442218.doi:10.1007/978-3-030-33128-3_1
18. Esteva A, Robicquet A, Ramsundar B, et al. A guide to deep learning in healthcare. *Nat Med*. 2019;25(1):24–29. PMID: 30617335.doi:10.1038/s41591-018-0316-z
19. Currie G, Hawk KE, Rohren E, Vial A, Klein R. Machine learning and deep learning in medical imaging: intelligent imaging. *J Med Imaging Radiat Sci*. 2019;50(4):477–487. PMID: 31601480.doi:10.1016/j.jmir.2019.09.005
20. Zhang X, Zhang Y, Zhang G, et al. Deep learning with radiomics for disease diagnosis and treatment: challenges and potential. *Front Oncol*. 2022;12:773840. PMID: 35251962; PMCID: PMC8891653.doi:10.3389/fonc.2022.773840
21. Choudhary K, DeCost B, Chen C, et al. Recent advances and applications of deep learning methods in materials science. *npj Comput Mater*. 2022;8:59.
22. Tran KA, Kondrashova O, Bradley A, Williams ED, Pearson JV, Waddell N. Deep learning in cancer diagnosis, prognosis and treatment selection. *Genome Med*. 2021;13(1):152. PMID: 34579788; PMCID: PMC8477474.doi:10.1186/s13073-021-00968-x
23. Li J, Fang Z, Zhou J, et al. The association between molecular biomarkers and ultrasonographic radiomics features for triple negative invasive breast carcinoma. *Chin J Ultrasonography*. 2019;28(2):137–143.
24. Sha YS, Chen JF. MRI-based radiomics for the diagnosis of triple-negative breast cancer: a meta-analysis. *Clin Radiol*. 2022;77(9):655–663. PMID: 35641339.doi:10.1016/j.crad.2022.04.015
25. Ma M, Gan L, Jiang Y, et al. Radiomics analysis based on automatic image segmentation of DCE-MRI for predicting triple-negative and nontriple-negative breast cancer. *Comput Math Methods Med*. 2021;2021:2140465. PMID: 34422088; PMCID: PMC8371618.doi:10.1155/2021/2140465
26. Gong X, Li Q, Gu L, et al. Conventional ultrasound and contrast-enhanced ultrasound radiomics in breast cancer and molecular subtype diagnosis. *Front Oncol*. 2023;13:1158736. PMID: 37287927; PMCID: PMC10242104.doi:10.3389/fonc.2023.1158736
27. Ye H, Hang J, Zhang M, et al. Automatic identification of triple negative breast cancer in ultrasonography using a deep convolutional neural network. *Sci Rep*. 2021;11(1):20474. PMID: 34650065; PMCID: PMC8517009.doi:10.1038/s41598-021-00018-x
28. Yin HL, Jiang Y, Xu Z, Jia HH, Lin GW. Combined diagnosis of multiparametric MRI-based deep learning models facilitates differentiating triple-negative breast cancer from fibroadenoma magnetic resonance BI-RADS 4 lesions. *J Cancer Res Clin Oncol*. 2023;149(6):2575–2584. PMID: 35771263.doi:10.1007/s00432-022-04142-7
29. Zhou Z, Adrada BE, Candelaria RP, et al. Prediction of pathologic complete response to neoadjuvant systemic therapy in triple negative breast cancer using deep learning on multiparametric MRI. *Sci Rep*. 2023;13(1):1171. PMID: 36670144; PMCID: PMC9859781.doi:10.1038/s41598-023-27518-2

30. Hwang KP, Elshafeey NA, Kotrotsou A, et al. A radiomics model based on synthetic MRI acquisition for predicting neoadjuvant systemic treatment response in triple-negative breast cancer. *Radiol Imag Cancer*. 2023;5(4):e230009. PMID: 37505106; PMCID: PMC10413296.doi:10.1148/rycan.230009
31. Jiang M, Li CL, Luo XM, et al. Ultrasound-based deep learning radiomics in the assessment of pathological complete response to neoadjuvant chemotherapy in locally advanced breast cancer. *Eur J Cancer*. 2021;147:95–105. PMID: 33639324.doi:10.1016/j.ejca.2021.01.028
32. Hacking SM, Windsor G, Cooper R, Jiao Z, Lourenco A, Wang Y. A novel approach correlating pathologic complete response with digital pathology and radiomics in triple-negative breast cancer. *Breast Cancer*. 2024;31(3):529–535. PMID: 38351366.doi:10.1007/s12282-024-01544-y
33. Jimenez JE, Abdelhafez A, Mittendorf EA, et al. A model combining pretreatment MRI radiomic features and tumor-infiltrating lymphocytes to predict response to neoadjuvant systemic therapy in triple-negative breast cancer. *Eur J Radiol*. 2022;149:110220. PMID: 35193025.doi:10.1016/j.ejrad.2022.110220
34. Ma M, Gan L, Liu Y, et al. Radiomics features based on automatic segmented MRI images: prognostic biomarkers for triple-negative breast cancer treated with neoadjuvant chemotherapy. *Eur J Radiol*. 2022;146:110095. PMID: 34890936.doi:10.1016/j.ejrad.2021.110095
35. Jiang L, You C, Xiao Y, et al. Radiogenomic analysis reveals tumor heterogeneity of triple-negative breast cancer. *Cell Rep Med*. 2022;3(7):100694. PMID: 35858585; PMCID: PMC9381418.doi:10.1016/j.xcrm.2022.100694
36. Xia B, Li C, Qian Z, et al. Radiomics based on machine learning in predicting the long-term prognosis for triple-negative breast cancer after neoadjuvant chemotherapy. *Chin J Radiol*. 2021;55(10):1059–1064.
37. Yu F, Wang J, Deng J, et al. Radiomics features on ultrasound imaging for the prediction of disease-free survival in triple negative breast cancer: A multi-institutional study. *Chin J Ultrasonography*. 2021;30(6):519–525.
38. Li Q, Zhang X, Sun Y. [Application of imaging omics and deep learning in the evaluation of lymph node status in rectal cancer]. *Chin J Radiol*. 2023;57(2):231–234.
39. Lin T, Liu A. Advances in medical image-based radiomics and deep learning in patients with hepatocellular carcinoma. *Chin J Med Imag*. 2022;30(4):401–405.
40. Xu Z, Yang Q, Li M, et al. Predicting HER2 status in breast cancer on ultrasound images using deep learning method. *Front Oncol*. 2022;12:829041. doi:10.3389/fonc.2022.829041
41. Chen ZH, Zha HL, Yao Q, et al. Predicting pathological characteristics of HER2-positive breast cancer from ultrasound images: a deep ensemble approach. *J Imaging Inform Med*. 2024. PMID: 39187701. doi:10.1007/s10278-024-01229-0
42. Kim SY, Lee J, Cho N, Kim YG. Deep-learning based discrimination of pathologic complete response using MRI in HER2-positive and triple-negative breast cancer. *Sci Rep*. 2024;14(1):23065. doi:10.1038/s41598-024-74276-w