Why are ultrasound (US) images of thyroid nodules suitable for deep learning (DL) analysis?

DL is a part of artificial intelligence (AI) systems, which is designed to have human’s way of thinking. DL has been applied in medical image analysis such as chest radiographs, retinal image, pathology, and US images, and showed comparable diagnostic performance with clinicians (1-4). To allow the DL to analyze images, we should input hundreds or thousands of images labelled with the answers, which is called training. Then the DL algorithm trains itself by extracting specific features from the images, and becomes able to predict the answer when a new test image is given. For instance, if one properly trains a DL algorithm with multiple images of cats and dogs labelled with the answers, the DL extracts the features of the two species and finally becomes to be able to differentiate them.

Neck US is a safe diagnostic imaging modality and the gold standard method for evaluation of thyroid nodules (5). Moreover, recent high-resolution US is easily utilized by clinicians for obtaining and interpreting sonographic neck images including thyroid nodules. The characteristics
of these nodules can be captured in one representative image making thyroid US suitable for DL analysis using convolutional neural network. This can be established by adding the features of thyroid nodules obtained by US to a DL algorithm in the form of a whole captured image or alternatively by cropping the image into multiple squares. The latter is generally preferred to avoid the influence of neighboring structures such as trachea, carotid artery, or muscles on the efficacy and accuracy of a DL algorithm.

**Previous studies using Computer-aided diagnosis (CAD) for thyroid nodule US images**

CAD system is based on classical machine learning. Unlike DL which itself determines and extracts key features from the images, CAD system requires that human should define the key features of a subject on which the prediction is based. In the CAD system for thyroid US, human should define the malignant features of the thyroid nodules such as irregular margin, taller-than-wide shape, markedly hypoechoic echogenicity, and presence of calcification to allow the CAD to predict malignant nodules. Then, the CAD quantifies each predetermined feature and eventually predicts if the given nodule is benign or malignant.

Choi et al. (6) used commercialized CAD system for the diagnosis of US images of thyroid nodules. They reported that CAD system showed a similar sensitivity as the experienced radiologist (90.7% vs. 88.4%) and lower specificity (74.6% vs. 94.9%). Similarly, Jeong et al. (7) used the same commercialized CAD system and reported that the CAD system had comparable sensitivity and lower specificity than experienced radiologist. The reason for the relatively high sensitivity and low specificity could be because the CAD system is more sensitive and consistent than the human in identifying the malignant features of thyroid nodules (Table 1).

<table>
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<th>Table 1 Ultrasonic Features suggestive of malignant thyroid nodules</th>
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<td>Hypoechoic nodule</td>
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<td>Taller than wide shape of the nodule</td>
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<tr>
<td>Irregular margin of the nodules</td>
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<td>Presence of microcalcifications</td>
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**Previous studies using DL for thyroid nodule US images**

Unlike CAD, DL does not require predetermination of malignant features but instead the final result of the thyroid biopsy or the specimen. After introducing these results in a DL algorithm, DL works to independently determine the different radiographic features (unbeknownst to the clinician) that are used to interpret future US images.

Interestingly, these features may not include the standard facts we use for US diagnosis (e.g., size, shape, etc.). Furthermore, we cannot know the features that DL uses for training or prediction, and thus the DL algorithm is referred to as a black box. DL algorithms generally outperform CAD systems, and the majority of recent studies use DL.

Ko et al. (8) used 439 US images for the training set and tested 150 images, and reported that DL algorithm showed comparable performance with radiologists (AUC 0.834 to 0.850 for DL and 0.805 to 0.860 for radiologist). Moreover, Song et al. (9) used 1,358 images from a training dataset and tested an algorithm with an internal (n=55) and external (n=100) test set. The sensitivity for the internal and external test set was 95.2% and 94.0% respectively. Buda et al. (10) used images from 1,278 and 99 nodules as a training and test set, respectively. They reported 87% sensitivity and 52% specificity in recommending further intervention, which was comparable with that recommended by radiologists.

How do we obtain enough images for successful DL algorithm training?

To train the DL algorithm using US images only, which is called training from scratch, a large amount of labeled US images are necessary because the diagnostic performance of a DL algorithm improves according to the size of the training dataset (11). However, the amount of collectable data is limited due to manpower and costs restrictions. In addition, it is unknown how many images are required for successful training. There are several methods however to address this limitation. One of the popular methods is transfer learning which saves time as it uses a pre-trained model. A pre-trained model is trained on a large benchmark dataset to solve a problem similar to the one that we want to solve. For instance, Inception is one of the most popular models, and pre-trained with the ImageNet database, which contains over 1.2 million images of commonly seen items in daily life. Using a pre-trained model is more efficient than training the whole layer of the DL algorithm despite the dataset not including medical or includes US images (12).

Another method is data augmentation. Data augmentation generates more images artificially by changing the ratio of
width to height, adding noise, changing colors, or using horizontal flip. It is reported to be helpful to achieve better DL performance (13). Although data augmentation is useful to increase the size of a training set and has been used in thyroid nodule image analysis (14), caution should be taken because it has a high potential to distort shape, margin, echogenicity, and calcification, which are essential elements for sonographic diagnosis of thyroid nodules (15).

**Limitations of DL analysis for thyroid nodule US images**

**Limitations in US image collection**

US is a convenient and reliable diagnostic tool when evaluating thyroid nodules. In general, one representative image contains enough information to delineate the nature of the nodule. Therefore, US fits well with the concept of DL. Regardless, there are high intra- and inter-reader variability in US image acquisition, and there is still a chance that the captured image of a thyroid nodule may not completely represent the lesion. For example, the features suggesting malignancy such as micro-calcification or irregular margin may not be adequately captured, or some features may look differently between axial and transverse images. This limitation might decrease the accuracy of US and subsequently the performance of DL.

**Indeterminate category**

American College of Radiology Thyroid Imaging Reporting And Data System (ACS TIRADS) is a risk stratification system evaluating thyroid nodules (16). The risk of malignancy is determined by five categories including composition, echogenicity, shape, margin, extra-thyroidal extension, and echogenic foci on US. Yet, ACS TIRADS is not confirmative, and FNAC is recommended for further evaluation of radiologically suspicious nodules. However, some FNAC results, which are reported as Bethesda categories, are still not confirmative (17). Category III/IV/V nodules can have diverse results such as benign, follicular thyroid carcinoma (FTC), variant type papillary thyroid carcinoma (PTC), or PTC on surgical pathology.

DL analysis of thyroid nodules are solely based on US findings. The accuracy of the DL is greatly influenced by the proportion of the nodules with indeterminate categories, which consists of 16% to 38% of the FNAC-tested nodules (18). FTC is usually diagnosed as indeterminate or benign category on FNAC, and has more benign feature on US than PTC. The more FTC is included in the dataset, the worse the diagnostic performance should be. Nonetheless, the researchers can increase the diagnostic performance of DL algorithm by excluding the indeterminate category as well as FTCs in the training or test set. However, the results cannot be applicable to real practice. This is why the number of indeterminate nodules on FNAC such as FTC used for training and testing must be mentioned in studies.

To illustrate this, a recently published paper reported that DL was trained with more than 300,000 thyroid US images, and showed relatively high specificity and sensitivity (14). However, of the 17,627 malignant nodules they used for training set, only 74 (0.4%) were FTCs, and 17,440 (98.9%) were PTC. Moreover, the number of FTC used for the test set was only 4 out of 1,194 nodules. Considering the incidence of FTC is one seventh of that of PTC (19), the results in this study should be interpreted carefully.

Likewise, variant types of PTC such as follicular variant PTC, which accounts for 12% to 30% of all PTC, also have less malignant US features compared to classical PTC (20,21). The more variant types of PTC are included in the training or test set, the worse the diagnostic performance of DL becomes. Therefore, care must be taken when interpreting study results in which most of the included malignant nodules are classic PTC (8).

Additionally, the gold standard in diagnosing a thyroid nodule is based on surgical pathology. However, some nodules are not removed by surgery, and the ultimate diagnosis is based on FNAC, with its inherent limitations mentioned above (9,22).

Nodules with indeterminate category on FNAC should undergo further evaluation with molecular testing or be removed surgically. Therefore, it could be more practical to train DL to discriminate thyroid nodules into groups that require surgery versus those that do not rather than into benign or malignant. Further research however is needed to address these concerns.

**Further practical considerations**

The results of DL analysis are presented as probability of benignity or malignancy ranging 0 to 1, not simply as benign or malignant. In general, benign or malignant results are presented based on a probability threshold of 0.5. However, the threshold can be arbitrarily adjusted as
needed, and the sensitivity and the specificity to predict malignancy change depending on the threshold. If the threshold is set to increase the specificity, the chance for misinterpreting true malignant nodules as benign, resulting in undertreatment of malignant nodules may increase.

Therefore, the threshold should be set to have high sensitivity in order that the false negative rate (predicting malignant as benign) can be minimized even if the false positive rate (predicting benign as malignant) is somewhat high. Considering more than 90% of the thyroid nodules which undergo FNAC turn out to be benign (23), and misdiagnosis of true benign as malignant would merely lead to FNAC, this scenario is desirable because DL can help patients to avoid unnecessary FNAC. Using DL as a decision support tool can be considered if an experienced clinician is not available.

Lastly, DL algorithms have been trained and tested using one 2D US image from each nodule. However, thyroid nodules are three-dimensional, and one representative US image may not completely reflect all pertinent features. DL analysis using multiple US images should be tried in the future to increase diagnostic capability.

Conclusions

Applying DL in the diagnosis of thyroid nodules is still in the developmental stage. There are several limitations related to efficient collection of US images, setting a proper threshold for predicting malignancy, and proper inclusion of indeterminate nodules into the dataset. Although currently developed DL cannot replace standard practice in the diagnosis of thyroid nodules, it might serve as an adjunctive tool to support the decision-making process for biopsy and surgery in the future.

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Footnote

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