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Enhanced Temporal Correlation for Universal Lesion Detection

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ABSTRACT

Universal lesion detection (ULD) methods for computed tomography (CT) images play a vital role in the modern clinical medicine and intelligent automation. It is well known that single 2D CT slices lack spatial-temporal characteristics and contextual information compared to 3D CT blocks. However, 3D CT blocks necessitate significantly higher hardware resources during the learning phase. Therefore, efficiently exploiting temporal correlation and spatial-temporal features of 2D CT slices is crucial for ULD tasks. In this paper, we propose a ULD network with the enhanced temporal correlation for this purpose, named TCE-Net. The designed TCE module is applied to enrich the discriminate feature representation of multiple sequential CT slices. Besides, we employ multi-scale feature maps to facilitate the localization and detection of lesions in various sizes. Extensive experiments are conducted on the DeepLesion benchmark demonstrate that this method achieves 66.84% and 78.18% for FS@0.5 and FS@1.0, respectively, outperforming compared state-of-the-art methods.

KEYWORDS

Universal lesion detection; computational biology; medical computing; deep learning; enhanced temporal correlation

1 Introduction

Computed tomography (CT), a widely used medical aid in diagnosis, scans and creates images of body parts from multiple directions and cross-sections. As the number of CT images in the medical system explosively grows, the limited number of professional radiologists makes it an urgent task to research medical diagnostic applications. With the continuous development of deep learning, which integrating with medical treatments to improve medical efficiency and solve the problem of uneven distribution of medical resources. At the same time, excellent lesion detection models have been proposed, including methods for single organs, such as lung nodule detection [1–10], lymph nodes [11–13], and liver tumors [14–39]. However, these methods cannot meet the clinical demand for detecting multiple lesioned organs at the same time. Therefore, universal lesion detection has gradually gained the attention of researchers, while many promising achievements have been obtained, such as [1–3,5,15,16], etc.



In recent years, Yan et al. [4] proposed a large dataset DeepLesion [4], which had greatly contributed to the development of universal lesion detection models. Based on the DeepLesion, some researchers [15–23] have proposed valid detection methods and have shown better metrics, respectively. Among the existing ULD methods, the accuracy is generally improved by the following aspects: anchor-free mechanism [24,25] or enhancing feature representation [1,16,20,26,27], optimizing anchor mechanism [15,28,29], and domain knowledge augment [30]. We focus on the idea of enhancing feature representation, which mainly include: (1) adding attention mechanism [1,26,27,31–34], the context information inter slices and the spatial information intra slices can be concentrated in the most prominent regions to enrich the feature representation; (2) introducing deformable convolution in the network [20], which can increase the receptive field of feature extraction; (3) using pseudo-3D [34] to process 3D CT input data reduces the number of network parameters to some extent. Although these schemes are effective, there is still room for improvement in terms of detection properties and efficiency.

We show the effectiveness of several methods for detecting single-organ lesions on a multi-organ lesion detection task in Fig. 1. We can see that a universal lesion detection network has more potential and values. Inspired by camouflaged object detection in video [17], enhancing the temporal correlation of multiple sequential slices contributes to the capture of lesions between slices by the network. Accordingly, by enhancing the temporal correlation, the feature representation of slices can be improved for improving the detection accuracy. Through the above observation, we find that the existing universal lesion detection methods neglect to focus on the correlation between CT slices, which can provide richer features for complex lesion detection tasks. Therefore, we propose an enhanced temporal correlation detection network (i.e., TCE-Net), in which the temporal correlation enhancement (TCE) module is introduced to extract temporal correlations of multiple sequential CT slices. In addition, multi-scale feature maps of sequential slices are extracted using a pre-trained Feature Pyramid Network (FPN) [31] for detecting lesions of different sizes.

The contributions of this work are threefold:

- (i) An end-to-end network called TCE-Net is proposed for universal lesion detection on CT images.
- (ii) A temporal correlation enhancement module was designed for lesion detection by extracting the temporal correlation inter multiple sequential CT slices.
- (iii) The experimental outcomes demonstrate that the developed framework performs favourably against state-of-the-art approaches on the DeepLesion benchmarks.

The remainder of this article are organized as follows. Section 2 introduces related work. The exploited TCE-Net for universal lesion detection is described in details in Section 3. Then, experimental results are provided in Section 4. Finally, we draw a conclusion of this paper in Section 5.

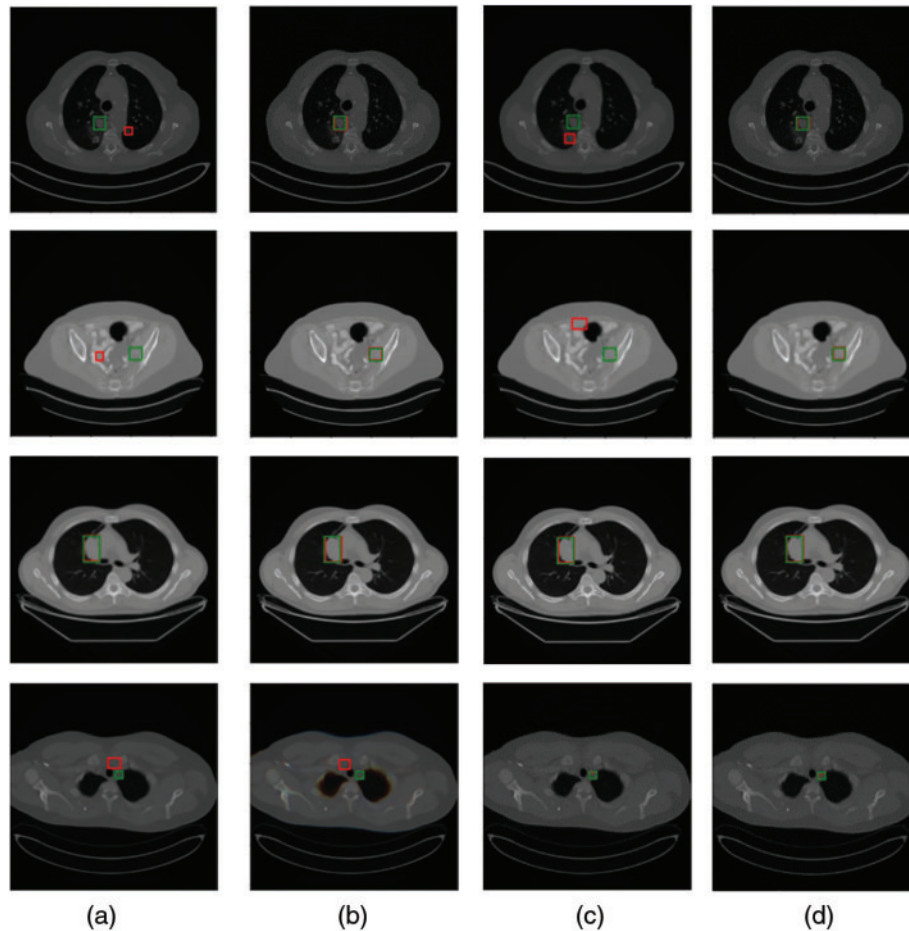


Figure 1: Comparison of visualization results of several single organ detection networks with ours. Green and red boxes correspond to ground-truths in the test set and predicted true positives. (a) The method proposed in [37]; (b) a modified SSD model by [38]; (c) MSCR [39]; (d) ours

2 Related Work

A Universal Lesion Detection (ULD) is a task that has great potential for development in object detection and is in a high clinical demand in clinical [40–51]. In recent years, the ULD task has received great attention, and related research works have been proposed one after another. Since 3D convolutional neural networks are very high memory consuming and expensive to train, and it is also a complex and difficult task to label 3D boxes for 3D CT dataset, Yan et al. [5] proposed a 3D context enhanced region-based CNN, called 3DCE, incorporating 3D context into a 2D CNN network to extract features and using a 2D detection network to make the final prediction. Shao et al. [1] proposed a Multi-Scale Booster, called MSB, which is used different dilation rates to capture the lesion responses from each pyramid feature map, respectively. In [27], with the aim of enriching feature representations and improving feature discrimination, they introduced a cross-slice context attention that selectively aggregated information from different slices and an intra-slice spatial attention to focus feature learning on the most salient regions. Meanwhile, pseudo-3D has also been applied to lesion detection, such as MP3D FPN proposed by Zhang et al. [16], which can efficiently extract 3D context

for universal lesion detection in CT slices. As in [1], the MVP-Net introduced by Li et al. [3] also extracts multi-scale features, with the difference that the MVP-Net used three FPNs corresponding to three window widths simultaneously to suit different detection parts. In addition to the above methods, there are also approaches based on domain knowledge enhancement, such as Sheoran et al. [30]. Meanwhile, refining the anchor mechanism is also a common mindset [15,24,25,28,29].

Currently, enhancing feature representation is still the majority route. Yet existing universal lesion detection models ignore the hidden temporal correlations between sequential slices which facilitate the catching of lesions when enhancing feature representation. With this motivation as a beginning point, we explore TCE-Net, where the TCE module focuses on extracting temporal feature descriptions, and has achieved impressive outcomes. More details are illustrated in Sections 3 and 4.

3 The Proposed Method

This section introduces a framework for universal lesion detection based on CT slices, named the TCE-Net method. Fig. 2 shows the framework of our proposed method, which will be detailed in the following subsections. First of all, an overall description of our approach is given in Section 3.1. In this framework, we develop the temporal correlation enhancement (TCE) module (see Section 3.2), whose structure is shown in Fig. 3. Then, Faster R-CNN [21] is used as a detector to demonstrate the effectiveness of our module. As for the total loss of the training phase, a detailed description is given in Section 3.3.

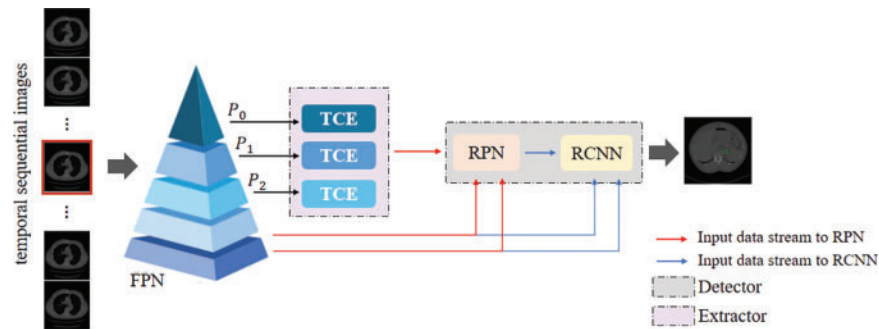


Figure 2: The framework of the proposed TCE-Net. TCE module learns the temporal correlation representation from features of sequential slices

3.1 TCE-Net

Given a set of sequential CT slices from a single subject, we learn temporal correlation and spatial-temporal features from multiple sequential slices to improve the characterization of the input data for accurate lesion detection. During the training stage, we match the predicted bounding box with the ground truth using the total loss. At the inference time, the overall network generates the final prediction directly.

Here, assume that the location of the target slice is t and the locations of the neighboring slices are $(t - 2\tau, t)$ and $(t, t + 2\tau)$, so the slices in the range of $[t - 2\tau, t + 2\tau]$ are defined as a sequence of input slices, which are fed to TCE-Net.

Firstly, the pre-trained FPN is cited in [31], including bottom-up, top-down and lateral connection components and is employed to extract multi-scale feature maps from the input data to prepare for the detection of lesions of different sizes. We define the feature maps of the output of each layer of the

pyramid from top to bottom as P_i , where i denotes the i -th layer. The channel number of P_i is set to 128.

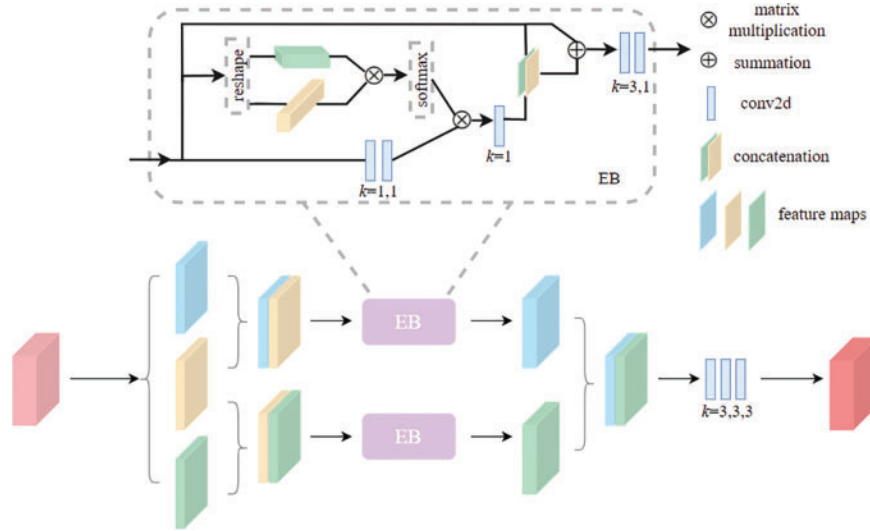


Figure 3: The pipeline of the TCE module, where the Enhanced Block (EB) helps to obtain correlation relationship between neighbouring slices group. The k in the figure is the convolution kernel size. Black arrows indicate the direction of data flow

Subsequently, the P_0, P_1, P_2 are further processed by the TCE module to improve the ability of detection network to track lesions by boosting the inter-slice correlation description. Mathematically, this process can be formulated as follows:

$$T_i = C(P_i), \tag{1}$$

where $C(\cdot)$ denotes the processing of the TCE module, and T_i represents hidden temporal correlation of feature representation at different size. Here, i takes 0, 1, 2. The enhanced modules in layers 0 to 2 share weights in the network.

Finally, the multi-scale enhanced temporal correlation features obtained by the TCE module cascade shallow features P_3 and P_4 are passed to the RPN in [21] and the detector Faster R-CNN for classification and regression in turn for lesion detection.

$$R = D(T_i, P_3, P_4), \tag{2}$$

where $D(\cdot)$ indicates the detector, and R represents the detected results.

3.2 TCE Module

In this section, we give detailed descriptions about the proposed TCE module for temporal correlation enhancement.

As shown in Fig. 3, in the TCE module, to aggregate correlations between multiple neighboring slices and reinforce the temporal correlation representation between the whole sequential slices, the study first divides the obtained feature maps of multiple slices G into N groups (N is set to 3 in this paper). Arithmetically, we define the features of N groups in G :

$$G = \{\zeta_1, \zeta_2, \dots, \zeta_{N-1}, \zeta_N\}. \tag{3}$$

In Eq. (3), ζ_j denotes the j -th group in the G. At the next step, Enhanced Block (EB) is employed to enhance the temporal correlation between two adjacent groups.

$$F_{j \sim j+1} = EB(\zeta_j, \zeta_{j+1}), \quad (4)$$

where, $j \geq 1 \cap j + 1 \leq N, j \in \mathbb{N}$. The final obtained multiple temporal correlation characterizations of adjacent groups F from EBs which contain the enhanced correlation are integrated, and then use three convolution layers to achieve the aim of further fusion.

Enhanced Block For learning temporal correlation representations from adjacent slices, we adopt an architecture, called Enhanced Block, i.e., EB.

The study begins by extracting the context information in the $x \times y$ -plane and the correlation information in the z -axis of the input feature map x entered into the Enhanced Block, to obtain temporal correlation and spatial-temporal feature between adjacent slices, defined as $\phi(x)$ and $\varphi(x)$, respectively, and then proceed to perform a matrix multiplication operation \otimes to fuse the representations. To enhance the discriminative representation of correlation information, we use softmax to acquire the correspondent probability distribution of fused representations y_1 , and multiply it with the deeper feature of original input y_2 . This can be expressed as follows:

$$y_1 = softmax(\phi(x) \otimes \varphi(x)), \quad (5)$$

$$y_2 = Conv(x), \quad (6)$$

$$y_0 = y_1 \otimes y_2, \quad (7)$$

where $Conv$ is the convolutional operation with 3×3 kernel. After that, the result y_0 passes through a convolution layer and cascades with the input x and a residual operation in order, to protect against the loss of feature information. The above steps are expressed by the formulation as

$$y = x + cat(Conv(y_0), x), \quad (8)$$

where the $cat(\cdot)$ and y is cascade operation and output separately. Finally, the output result is obtained by two 3×3 convolutions. The resolutions of features remain constant throughout the whole procedure to maintain the representations contained.

3.3 Total Loos

To train the TCE-Net, we use lesion classification loss and bounding-box regression loss in Faster R-CNN.

$$L(\{p_i\}, \{t_i\}) = \frac{1}{N_{cls}} \sum L_{cls}(p_i, p_i^*) + \lambda \frac{1}{N_{reg}} \sum p_i^* L_{reg}(t_i, t_i^*). \quad (9)$$

where p_i is the predicted classification probability of anchor $[i]$. The $p_i^* = 1$ when anchor $[i]$ is a positive sample and $p_i^* = 0$ when anchor $[i]$ is a negative sample. t_i indicates the parameterized coordinates of the predicted bounding box. t_i^* represents ground truth. λ is the weight balance parameter that is set to 10. N_{cls} means mini-batch size, N_{reg} is the number of anchor location. The L_{reg} use smooth L1 loss, and the L_{cls} adopt cross entropy. They are optimized jointly.

4 Experiments

4.1 Datasets and Evaluation Metrics

DeepLesion [4], is the largest multi-category, lesion-level annotated dataset of clinical medical CT images ever published by the NIH Clinical Center, which collects a total of 10594 CT scans before and after 4427 individual anonymous patients, including 32735 instances of labeled lesions. DeepLesion

divides all data into a 70% training set, a 15% validation set, and a 15% test set. The study uses DeepLesion’s training set to train the model and tests the model in the test set to obtain the average sensitivity under different false positives, serving as an evaluation metric to compare performance with other models.

We train the model on an NVIDIA GeForce RTX 3090 GPU, using the FPN with ResNet-50 that pre-trained on ImageNet [23] in our framework. We set five anchor scales (16, 32, 64, 128, 256) and three anchor ratios (0.5, 1.0, 2.0) in RPN while the threshold value of IOU is set to 0.7, in order to balance learning speed and learning effectiveness of model. The CT slice is resized to 800×800 pixels.

We adopt the stochastic gradient descent (SGD) optimizer with momentum 0.9 and weight decay 0.0001 to train the model for 14 epochs. The base learning rate is set at 0.002, reduced by a factor of 10 after the 11th and 13th epochs.

4.2 Comparison with Other Methods

We compare the proposed TCE-Net with some state-of-the-art methods including Faster R-CNN [21], 3DCE [5], ULDOR [19], GPN [18], MVP-Net [3], LENS [22] and Deformable Dilated Faster R-CNN [20]. To be fair comparison, we have re-implemented 3DCE using the FPN backbone. We perform the evaluation by computing the sensitivity values at different false positives rates and mean average precision (mAP) as illustrated in Table 1.

Table 1: On the DeepLesion test set, the proposed method is compared with the state-of-the-art method. Lesion detection sensitivity values are now reported at different false positive (FP) rates. The number after “,” denotes the number of slices

Methods	FP@							mAP
	0.5	1	2	3	4	8	16	
Faster R-CNN21	0.57	0.67	0.76	–	0.82	0.86	0.89	–
3DCE5, 3	0.522	0.6297	0.7351	0.7881	0.8263	0.8919	0.9294	0.3365
3DCE, 9	0.5482	0.6547	0.7459	0.7897	0.8166	0.8719	0.9151	0.2988
3DCE, 27	0.5967	0.704	0.7836	0.8247	0.8469	0.8921	0.9235	0.3349
GPN18	0.36	0.5	0.62	–	0.72	0.76	0.79	–
ULDOR19	0.5286	0.648	0.7484	–	0.8438	0.8717	0.918	–
MVP-Net3, 3	0.5704	0.6893	0.796	0.8394	0.8634	0.9161	0.9442	0.3529
MVP-Net, 9	0.6224	0.7492	0.8308	0.8669	0.8884	0.9261	0.9491	0.4219
LENS22	0.403	0.5	0.596	–	0.695	0.78	–	–
Deformable dilated faster R-CNN20	0.427	0.531	0.65	–	0.775	0.775	–	–
Ours, 3	0.6055	0.7288	0.8164	0.8534	0.8793	0.9208	0.9497	0.3921
Ours, 7	0.625	0.7522	0.8396	0.8789	0.8984	0.9296	0.9532	0.4026
Ours, 9	0.6684	0.7818	0.8585	0.8872	0.9072	0.9365	0.9536	0.4241

4.2.1 Quantitative Results

From Table 1 and Fig. 4, we can see that GPN uses lesion detection under a single-scale feature map, and the performance is significantly poorer than that of multi-scale feature-based lesion detection

models like MVP-Net and ours, which illustrates the essential need for multi-scale feature extraction. Our model outperforms 3DCE 3 slices by 9.91% and MVP-Net 3 slices by 3.95% for FPs@4.0 for the same 3 slices case. More importantly, our proposed TCE-Net 9 slices achieves a performance boost of over 7.78% for FPs@1.0 and 4.6% for FPs@0.5 compared to 3DCE 27 slices and MVP-Net 9 slices, respectively. Meanwhile, the indicator mAP reaches its greatest at TCE-Net 9 slices.

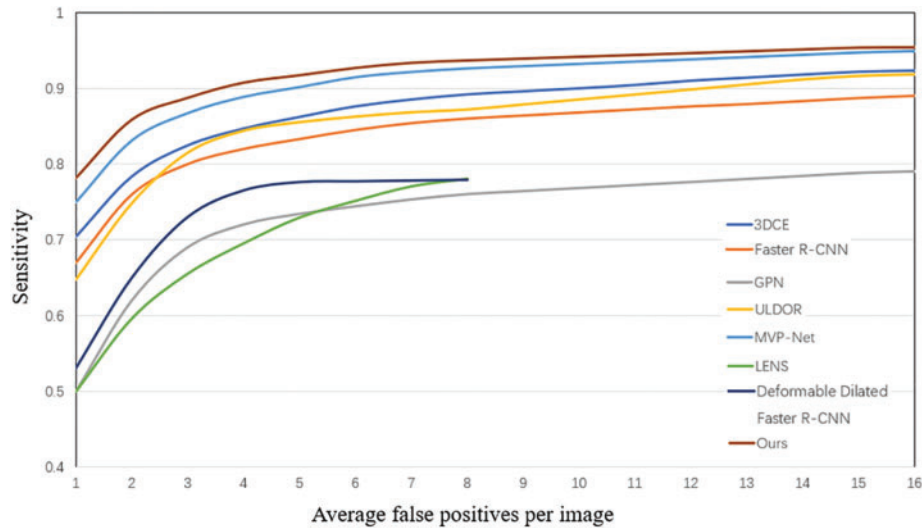


Figure 4: FROC curves of various methods on the test set

4.2.2 Qualitative Results

Predicted bounding boxes generated from several compared models are illustrated in Fig. 5. Our model shows superior capability to predict both the lesion location and size when compared with other methods. It is evident that the proposed method enhances the performance in detecting very small-sized or larger lesions. However, the insufficient training data for certain organ lesions resulted in inadequate generalization for the detection of these lesions as can be seen in Fig. 6.

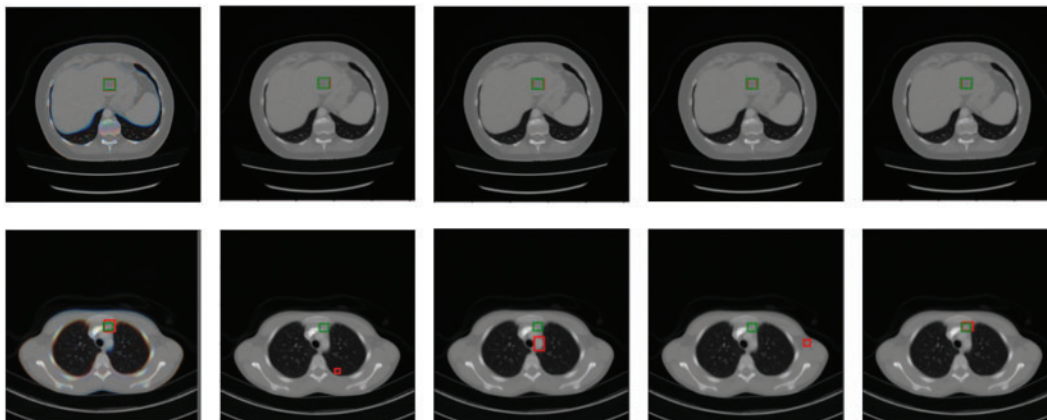


Figure 5: (Continued)

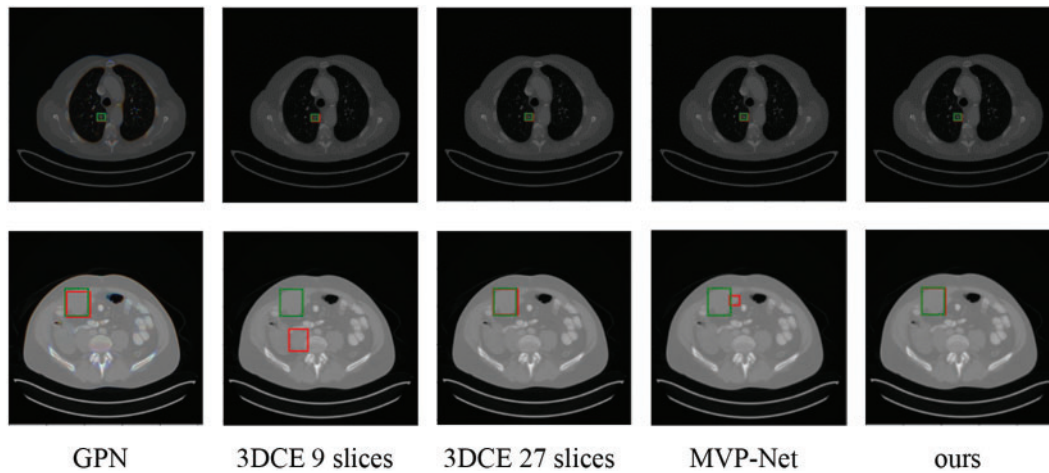


Figure 5: Qualitative comparison of several existing universal lesion detection models and ours on the testing set of DeepLesion. Green and red boxes correspond to ground-truths in the test set and predicted true positives

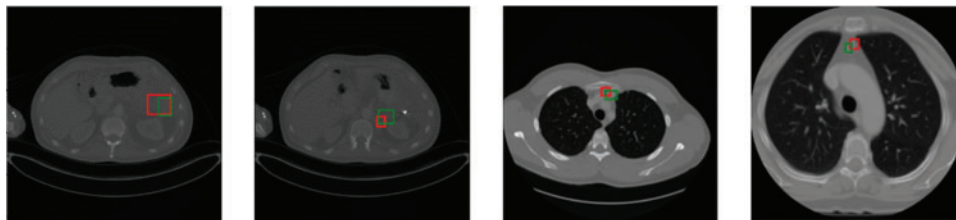


Figure 6: Visualization of inaccurate lesion detection results. Green and red boxes correspond to ground-truths in the test set and predicted true positives

4.3 Ablation Experiment

We perform a number of ablations to probe into our TCE-Net. As shown in Table 2, it shows that the use of the temporal correlation enhancement module has improved the metric by 1.52% compared to the baseline FPN, on the basis of which the feature information is increased by adding the number of input slices to 7 and 9 slices, respectively. And it can be seen that the accuracy of the model was improved by 3.86% and 7.41%, respectively, and the best results are achieved when using the combination of TCE and input 9 slices.

Table 2: Performance of our proposed method for doing ablation experiments on the DeepLesion dataset. The number after “,” denotes the number of slices

Methods	FP@						
	0.5	1	2	3	4	8	16
FPN	0.5943	0.7136	0.8101	0.8498	0.8748	0.9194	0.9497
FPN+TCE, 3	0.6055	0.7288	0.8164	0.8534	0.8793	0.9208	0.9497
FPN+TCE, 7	0.625	0.7522	0.8396	0.8789	0.8984	0.9296	0.9532

(Continued)

Table 2 (continued)

Methods	FP@						
	0.5	1	2	3	4	8	16
FPN+TCE, 9	0.6684	0.7818	0.8585	0.8872	0.9072	0.9365	0.9536

5 Conclusions

This work investigates the effectiveness of the enhanced temporal correlation detection network (TCE-Net) for universal lesion detection (ULD) in CT scans. The temporal correlation enhancement module has been proposed to enhance the effective fusion of multi-slice features through temporal correlation between multiple adjacent slices. We validate the effectiveness of our method with various experimental and analytic results. It shows that the proposed method brings a performance boost in the lesion detection tasks in CT scans. Universal lesion detection is a challenging task, partly because lesions in some organs have minimal differences from adjacent tissues. Therefore, false positives are high for detection in these organs. Meanwhile, the imbalance of data samples from various types of organs also limits the generalization ability of TCE-Net. In the future enriched organ CT data will be needed to train TCE-Net to improve the performance.

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Availability of Data and Materials: Preprocessed data are available on request from the first author.

Conflicts of Interest: The authors declare that they have no conflicts of interest to report regarding the present study.

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