Medical Imaging

Team VcLD

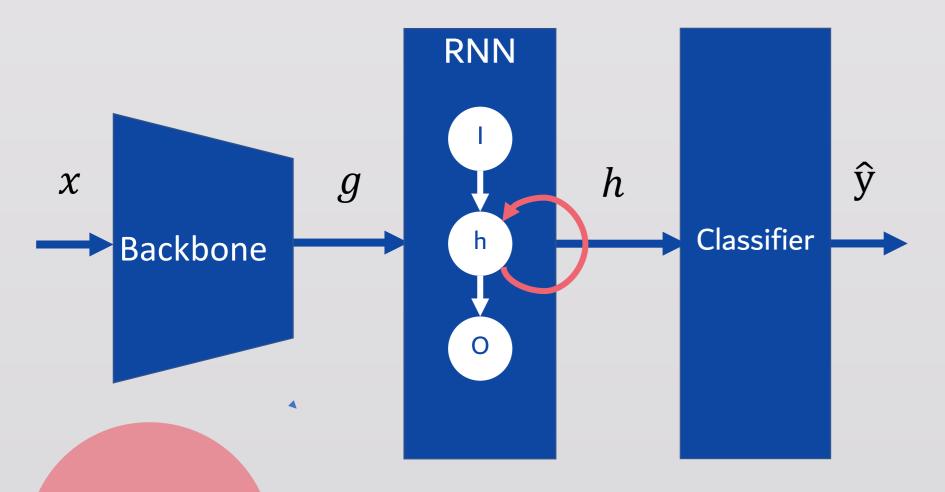
Data Preprocessing

- Random rotation
- Horizontal / Vertical flip
- Brightness adjustment
- MultiScaleCrop ... etc

Model Architecture

Backbone + RNN + Classifier

ResNet18 + BiGRU + FC



Hyperparameters

- Batch Size: 4 (patient_ids)
- Optimizer: AdamW
- $\gamma_{-}, \gamma_{+} = 4, 1$
- T = 8
- $\epsilon = 0.5$ $p_1, p_2 = 15, 0.15$
- Learning Rate: 0.001
- Learning Rate Schedule: Reduce the learning rate by half each time the validation loss is non-decreasing for three continuous epochs.
 Stop the training stage if learning rate is smaller than 1e-6.

Performance

F2-score evaluation:

Leaderboard	Public	Private
Single Model	0.78255	0.78312
Ensemble Model	0.79395	0.79353
Small dataset	0.72908	0.73567

Ablation Study

Model	F2-Score (Public)
MTM (remove diversity)	0.77197
MTM (remove CPC)	0.77362
MTM (remove masking strategy)	0.77411
MTM (remove AC)	0.77866
MTM (remove CC)	0.77889
MTM (remove LP)	0.78007
MTM	0.78255

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Methodology

- x: input images
- y:input labels
- *g* : output of ResNet18
- *h* : output of Bi-GRU
- \hat{y} : output of classifier
- res, rnn, cls: function of ResNet18, bi-GRU, and classifier.

Tasks:

1. Multi-label Classification: *Asy* is the asymmetric loss function.

$$L_1 = Asy(y, \hat{y})$$

$$Asy(y, \hat{y}) = -y(1 - \hat{y})^{\gamma_+} \log(\hat{y}) - (1 - y)\hat{y}^{\gamma_-} \log(1 - \hat{y})$$

2. Auxiliary Classification (AC): Remove the RNN layer and only use g to predict labels.

$$L_2 = Asy(y, cls(g))$$

3. Number of Label Prediction (LP): Predict the number of labels for each image. The objective function is cross entropy loss.

$$L_3 = CE(y, cls_2(g))$$

4. Class-Correlation Classification (CC): It uses the Euclidean distance between g and label embedding to predict the labels. The distance between a feature g and k-th label embedding should be small if the feature contains k-th label. (Asymmetric Loss)

$$d_k = f_d(g, c_k)$$

$$d = [d_1, d_2, ..., d_K] \text{ (K=5)}$$

$$q = -\operatorname{softmax}(d)$$

$$L_4 = Asy(y, q)$$

5. Contrastive Predictive Coding (CPC): This is an unsupervised learning task which is used to extract useful representations from high-dimensional data. This task aims to predict $g_{t+1}, g_{t+2}, ..., g_{t+T}$ from h_t (InfoNCE Loss)

$$L_5 = -E \left[log \frac{f_k(g_{t+k}, h_t)}{\sum_{g_i} f_k(g_j, h_t)} \right]$$

Loss functions:

- $L = \alpha L_1 + \beta L_2 + \gamma L_3 + \mu L_4 + \nu L_5$
- $\alpha = \beta = 1, \gamma = \mu = \nu = 0$ (Pretraining)
- $\alpha = \beta = \gamma = \mu = \nu = 1$ (Training MTM)

Initialization:

Pretrained model using MC and AC tasks only (L1 and L2)

Generalization:

Masking:

Choose p_1 % of the image positions randomly. If i-th image is chosen, we replace g_i with all zeros.

Diversity:

Sample a pair of images (z_1,z_2) from training data with following conditions.

i. z_1 and z_2 are adjacent to each other, with z_2 behind z_1

ii. z_2 must always have one more label than z_1

iii. z_1 and z_2 's label sequences must be identical except for the one label z_2 has that z_1 does not. (z_2 contains an additional label l which does not exist in z_1 , [0,1,1,0,0] and [0,1,1,0,1])

Therefore,
$$info(l) = z_2 - z_1$$

Procedures for constructing a new image sequence with image sequence $\{x_1, x_2, ..., x_m\}$ and labels $\{y_1, y_2, ..., y_m\}$:

- i. Let $n = m * p_2$
- ii. Choose n images from the sequence randomly.
- iii. Sample n pairs of images which satisfy the conditions above. Calculate $info(l_k)$ for each pairs of images and obtain $\{info(l_{k_1})_1, info(l_{k_2})_2, ..., info(l_{k_n})_n\}, k_1, k_2, ..., k_n \in \{1, ..., K\}$
- iv. Add $\inf o(l_{k_i})_i$ to its corresponding i-th chosen images in step ii.
- v. Add l_{k_i} to the labels of i-th chosen images if it doesn't exist already Similar to scheduled sampling strategy in NLP, for each patient, we define a probability ϵ to decide whether the model is trained by the original image sequence or the generated image sequence (by applying masking or adding additional labels as described above)