A APPENDIX

{findings or impression}

You are a professional radiologist. Please determine if the anatomy ({anatomy}) is mentioned in this CT image report. Please answer directly with "Yes" or "No".

Figure 6: Prompt used to judge if an anatomy is mentioned in the "Findings" or "Impression" section of a clinical report.

{findings or impression}

You are a professional radiologist. Please extract the descriptive information about the specific anatomy ({anatomy}) from this CT image diagnostic report. Please follow these guidelines:

Precise extraction: Extract the descriptive information directly related to {*anatomy*} from the report. **Specify anatomical details:** If the report mentions specific areas, parts, or anatomical details of {*anatomy*}, make sure to include this information in the description. This could include affected areas, normal structures, or any notable features.

Concise and clear: Directly extract the report content, avoiding unnecessary explanations or background. **Format requirement:** Please provide the information in the format "*{anatomy}*: descriptive information", ensuring to use *{anatomy}* as the unified prefix for the item. Even if the anatomy has multiple independent parts or multiple lateral characteristics, it should be treated and described as a whole, returning only one comprehensive piece of information about that anatomy.

Figure 7: Prompt used to extract anatomy-specific description from the "Findings" or "Impression" section of a clinical report. Notably, we do not obtain the descriptions for all anatomies in a single query; rather, we strategically query the LLM for each anatomy individually. This approach significantly simplifies the complexity of description extraction and greatly enhances the quality of the extracted descriptions.

A.1 DETAILS ABOUT THE TEXT CLASSIFIER

We utilize the annotated validation and test sets of MedVL-CT69K to develop a text classifier that identifies 54 abnormalities in the generated radiology reports. To achieve this, we first merge these two sets and then re-split them into new training and validation sets using a 2:1 ratio. Afterwards, we train the classifier, which consists of a BERT-base encoder and a classification head, using the reports and corresponding disease labels form the training set. A binary cross-entropy loss is used to supervise the model training. Tab. [7] shows the precision, recall, and F1 scores of the text classifier across 54 abnormalities on the validation set. Notably, the model achieves an impressive average F1 score of 0.95. This high performance substantiates its reliability as a tool for assessing the diagnostic accuracy of report generation models.

A.2 IMPLEMENTATION DETAILS

For the abdominal MedVL-CT69K dataset, we reformat all CT scans so that the first axis points from inferior to superior, the second from posterior to anterior, and the third from left to right. We then resample the in-plane axial images to 1mm resolution and the out-of-plane slice thickness to 5mm spacing using trilinear interpolation. We map the Hounsfield unit range -300:400 to the range 0:1, clipping values that fall outside of this range. We use ViT-base Dosovitskiy et al. (2020), initialized with MAE ImageNet-1K pre-trained weights He et al. (2022), as the image encoder. The patch size is set to 16, 16, 32 along the axial, coronal, and sagittal axes, respectively. A pre-trained BERT-base Devlin et al. (2018) model is used as the text encoder. We train fVLM with an Adam optimizer. The learning rate linearly increases to 1e-4 in the first epoch and then decreases to 1e-6 with a cosine decay scheduler. The model undergoes training for 20 epochs on 4 A100 GPUs, with a batch size of 48. During model training, we apply RandomCrop and RandomFlip on the fly. The cropping size is set to 96, 256, and 384 along the axial, coronal, and sagittal axes, respectively. Notably,



Figure 8: Percentage of normal samples for each anatomy.



Figure 9: Performance comparison between our method and three radiologists. "n_pos" denotes the number of positive samples of each abnormality.

we observe that if a completely random cropping strategy is used, larger anatomies are more likely to be incomplete after cropping and consequently excluded from the loss calculation. This would introduce a data bias and potentially compromise the model's performance. To address this issue, we employ a uniform sampling strategy to randomly select an anatomy that must be completely included in the cropped image region. For the chest CT-RATE dataset, we apply the same image pre-processing as CT-CLIP Hamamci et al. (2024) to ensure a fair comparison with the competitors. In our co-teaching approach, we iteratively train two fVLMs, alternating between them after each iteration. We initiate a burn-in stage of 5 epochs to allow both models to establish a baseline level of performance. After that, we leverage each model to generate soft labels for its counterpart.

A.3 READER STUDY

To further validate our method's efficacy, we conduct a reader study to compare our approach with three board-certified radiologists. For this experiment, we randomly select 100 patients from the test set of MedVL-CT69K. Fig. 9 shows the results. Although our method has demonstrated sig-

Table 5: Performance of fVLM when using different models to correct contrastive labels.

	Baseline	Model self	Momentum	CoT
AUC	78.7	73.3	78.8	79.8
ACC	75.3	71.4	75.5	75.9







Figure 11: Visual activation maps of our model in diagnosing multiple diseases.



Figure 12: T-SNE visualization of visual embeddings for various abnormalities.

nificant improvements over previous approaches, there remains a noticeable performance gap compared to professional radiologists overall. However, for some diseases such as liver cirrhosis and splenomegaly, our method achieves comparable diagnostic accuracy to radiologists.

A.4 FURTHER ABLATION ANALYSIS

In Tab. 5, we compare the performance of fVLM when employing different models to correct contrastive learning labels during pre-training. It can be seen that utilizing the training model itself for label correction leads to a significant performance degradation, which could be attributed to the error accumulation issue. Moreover, the proposed CoT strategy yields greater performance gains compared to the momentum model. To explore this, we measure the difference between training model and label correction model by calculating the Euclidean distance of their parameters, as illustrated in Fig. 10. It can be observed that the momentum model, updated through exponential moving average, exhibit minimal discrepancy with the training model. This suggests they may produce similar predictions, potentially leading to error accumulation in the label correction process. In contrast, the iteratively trained models in our proposed CoT framework exhibit considerable distinctness, leading to diverse predictions and reducing the risk of error accumulation.

A.5 VISUALIZATION

We qualitatively assess the alignment efficacy of our proposed method through visualization in Fig. [1]. The heatmaps illustrates the correlation between anatomy-specific visual tokens and the textual embedding of abnormality. We observe high activation in specific affected areas for both localized lesions (*e.g.*, bladder stone) and diffuse abnormalities (*e.g.*, fatty liver). The results demonstrate the model's capacity to precisely localize pathological changes across a spectrum of conditions. Fig. [12] illustrates the distribution of visual embedding for a diverse array of abnormalities. In contrast to CLIP, our method exhibits more compact embedding clusters among positive cases of each abnormality. These findings demonstrate the improved semantic understanding and diagnostic interpretability of our fVLM.

Anatomical System	Anatomy	Grouped Anatomy	
, , , , , , , , , , , , , , , , , , ,	Face	Face	
	Brain	Brain	
	Esophagus	Esophagus	
	Trachea	Trachea	
	Lung upper lobe left		
	Lung lower lobe left		
	Lung upper lobe right	Lung	
	Lung middle lobe right	8	
	Lung lower lobe right		
	Heart myocardium		
	Heart atrium left		
	Heart atrium right	Heart	
	Heart ventricle left	ficart	
Organs	Heart ventricle right		
Organs	Adrenal gland right		
	Adrenal gland left	Adrenal gland	
	Kidnov right		
	Kidney light	Kidney	
		<u> </u>	
	Stomach	Stomach	
	Liver	Liver	
	Gall bladder	Gall bladder	
	Pancreas	Pancreas	
	Spleen	Spleen	
	Colon	Colon	
	Small bowel	Small bowel	
	Duodenum	Sinui Sover	
	Urinary bladder	Urinary bladder	
	Aorta	Aorta	
	Inferior vena cava	Inferior vena cava	
	Portal vein and splenic vein	Portal vein and splenic vei	
Vacala	Pulmonary artery	Pulmonary artery	
vessels	Iliac artery left	Ilio o ortorry	
	Iliac artery right	mac artery	
	Iliac vena left	TI .	
	Iliac vena right	lliac vena	
	Vertebrae L1-L4	Lumbar vertebrae	
	Vertebrae T1-T12	Thoracic vertebrae	
	Vertebrae C1-C7	Cervical vertebrae	
	Rib left 1-12		
	Rib right 1-12	Rib	
	Humerus left		
	Humerus right	Humerus	
	Scapula left		
Bones	Scapula right	Scapula	
Dones	Clavicula left		
	Clavicula right	Clavicula	
	Eawletia light		
	Femur right	Femur	
	Hip loft		
	Lip right	Hip	
	Soortum	Saamum	
		Sacruin	
	Gluteus maximus left		
	Gluteus maximus right	Gluteus	
	Gluteus medius left		
	Gluteus medius right		
Muscles	Gluteus minimus left		
	Gluteus minimus right		
	Iliopsoas left	Ilionsoas	
	Iliopsoas right	mopsoas	
	Autochthon left	Autochthon	
	Autochthon right	Autochuloli	

Table 6: Anatomy grouping.

Anatomical organ	Abnormality	Precision	Recall	F1-score
A due nel ele n d	Thickening	1.00	0.97	0.99
Adrenal gland	Nodule	1.00	0.96	0.98
Pladdar	Diverticulum	0.97	0.94	0.96
Diauuei	Stones	1.00	1.00	1.00
	Gas	0.84	0.79	0.81
	Effusion	0.81	0.71	0.75
	Obstruction	0.86	1.00	0.92
Colon	Diverticulum	0.97	1.00	0.98
Cololi	Colorectal Cancer	0.97	0.95	0.96
	Rectal Cancer	1.00	0.95	0.97
	Appendicitis	1.00	1.00	1.00
	Appendicolith	0.89	0.96	0.92
Fsonhagus	Hiatal Hernia	0.74	1.00	0.85
Lsophagus	Varicose Veins	1.00	1.00	1.00
	Cholecystitis	0.99	1.00	0.99
Gallbladder	Gallstone	1.00	1.00	1.00
	Adenomyomatosis	0.92	0.92	0.92
Heart	Cardiomegaly	1.00	0.95	0.97
Incurt	Pericardial Effusion	1.00	1.00	1.00
	Atrophy	0.97	0.88	0.92
Kidney	Cyst	0.97	0.96	0.97
Runey	Hydronephrosis	0.88	1.00	0.94
	Calculi	0.99	0.98	0.99
	Steatosis	0.99	1.00	0.99
	Glisson's Capsule Effusion	0.85	0.89	0.87
	Metastase	0.90	0.95	0.92
Liver	Intrahepatic Bile Duct Dilatation	0.96	0.97	0.97
	Cancer	1.00	1.00	1.00
	Cyst	0.99	0.99	0.99
	Abscess	0.91	0.95	0.93
	Cirrhosis	1.00	1.00	1.00
	Atelectasis	0.96	0.98	0.97
T	Bronchiectasis	0.97	0.9	0.93
Lung	Emphysema	1.00	0.96	0.98
	Pneumonia	0.98	0.96	0.97
	Pleural effusion	0.98	1.00	0.99
	Pancreatic cancer	1.00	0.89	0.94
Denenan	Atrophy	1.00	0.82	0.90
Pancreas	Pancreatius Demonstria durat dilatation	1.00	1.00	1.00
	Pancreatic duct dilatation	0.98	0.91	0.93
	Hupertension	0.97	0.87	0.92
Portal vein	Thrombosis	0.74	0.91	0.93
	Gas	0.74	0.74	0.74
Small Intestine	Gas	0.89	0.93	0.91
	Obstruction	0.89	0.92	0.91
	Diverticulum	0.93	1.00	0.95
	Intussusception	0.97	0.90	0.90
	Hemangioma	1.00	0.97	0.92
Snleen	Infarction	0.95	0.97	0.07
Spicen	Splenomegalv	1.00	0.99	1.00
	Gastric wall thickening	0.96	0.96	0.96
Stomach	Stomach cancer	1.00	0.90	0.90
Sacrum	Osteitis	0.97	1.00	0.99
			0.05	0.95
Avelage		0.25	0.95	0.95

Anatomy	Anatomy count	Abnormality	Abnormality count
Adrenal gland	63015	Thickening	3037
Autenai gianu	03913	Nodule	3687
Dladdar	62192	Diverticulum	283
Diadder	02182	Stones	109
		Gas	2173
		Effusion	975
		Obstruction	436
Cala	(2054	Diverticulum	1623
Colon	62054	Colorectal Cancer	817
		Rectal Cancer	858
		Appendicitis	1623
		Appendicolith	1119
	2/2/	Hiatal Hernia	184
Esophagus	2636	Varicose Veins	609
		Cholecystitis	3935
Gallbladder	63407	Gallstone	5500
Gundhudder	00107	Adenomyomatosis	1246
		Cardiomegaly	316
Heart	3701	Dericardial Effusion	1067
		Atrophy	021
		Cust	27010
Kidney	63618	Uyst	1140
		Calculi	5256
		Calcul	3330
	63690	Stealosis	4872
		Glisson's Capsule Effusion	915
		Metastase	2403
Liver		Intrahepatic Bile Duct Dilatation	6093
		Cancer	888
		Cyst	21/10
		Abscess	239
		Cirrhosis	1772
	6598	Atelectasis	1988
_		Bronchiectasis	781
Lung		Emphysema	190
		Pneumonia	1463
		Pleural effusion	4665
		Pancreatic cancer	933
	63627	Atrophy	942
Pancreas		Pancreatitis	1035
		Pancreatic duct dilatation	2697
		Steatosis	846
Doutol voin	62055	Hypertension	1149
Portal vein	03833	Thrombosis	760
	62419	Gas	2906
Small Intestine		Effusion	2326
		Obstruction	1174
		Diverticulum	2352
		Intussusception	168
		Hemangioma	718
Spleen	63749	Infarction	374
Spreen	55717	Splenomegalv	1732
		Gastric wall thickening	2871
Stomach	63682	Stomach cancer	1064
Sacrum	62055	Osteiti	246
Sacium	02055	Usicili	240

Table 8: The distribution of 54 tested abnormalities in the train set. We employ the well-developed text classifier to automatically extract abnormality labels from radiology reports for each sample.

Anatomy	Anatomy count	Abnormality	Abnormality count	
A dranal gland	1140	Thickening	62	
Autenai gianu	1149	Nodule	79	
		Gas	29	
		Effusion	13	
Colon	1127	Obstruction	5	
COIOII	1127	Colorectal Cancer	10	
		Rectal Cancer	16	
		Appendicitis	5	
		Cholecystitis	73	
Gallbladder	1054	Gallstone	127	
		Adenomyomatosis	35	
		Atrophy	16	
Kidnay	11/18	Cyst	492	
Klulley	1140	Hydronephrosis	14	
		Calculi	104	
	1146	Steatosis	97	
		Glisson's Capsule Effusion	20	
		Metastase	40	
Liver		Intrahepatic Bile Duct Dilatation	132	
		Cancer	10	
		Cyst	381	
		Cirrhosis	30	
		Pancreatic cancer	5	
		Atrophy	16	
Pancreas	1149	Pancreatitis	26	
		Pancreatic duct dilatation	49	
Dortal vain	1150	Hypertension	18	
r ontar veni	1150	Thrombosis	10	
	1131	Gas	34	
Small Intestine		Effusion	25	
		Obstruction	8	
		Hemangioma	12	
Spleen	1140	Infarction	8	
		Splenomegaly	35	
Stomach	1150	Gastric wall thickening	61	
Stomach	1150	Stomach cancer	20	

Table 9: The distribution of 36 annotated abnormalities in the validation set.

Anatomy	Anatomy count	Abnormality	Abnormality count
Adrenal gland	3418	Thickening	96
	0.10	Nodule	87
Bladder	3243	Diverticulum	21
		Stones	28
		Gas	129
		Chatmation	30
		Diverticulum	104
Colon	3213	Colorectal Cancer	104
		Rectal Cancer	90 73
		Appendicitis	19
		Appendicolith	74
		Hiatal Hernia	10
Esophagus	105	Varicose Veins	78
		Cholecystitis	246
Gallbladder	3134	Gallstone	355
Guildiadei	5151	Adenomyomatosis	60
		Cardiomegaly	20
Heart	234	Pericardial Effusion	77
		Atrophy	37
		Cvst	1646
Kidney	3313	Hydronephrosis	87
		Calculi	408
		Steatosis	263
	3281	Glisson's Capsule Effusion	68
		Metastase	122
T '		Intrahepatic Bile Duct Dilatation	264
Liver		Cancer	61
		Cyst	1264
		Abscess	12
		Cirrhosis	188
		Atelectasis	70
	126	Bronchiectasis	18
Lung		Emphysema	10
		Pneumonia	72
		Pleural effusion	94
		Pancreatic cancer	29
_	3328	Atrophy	37
Pancreas		Pancreatitis	77
		Pancreatic duct dilatation	94
		Steatosis	45
Portal vein	3410	Hypertension	54
		I hrombosis	33
	3248	Gas	188
Small Intestine		Effusion	142
		Discretion	01
		Intussussention	115
		Hemoneticme	10
Splaan	3350	Information	4/
spieen	3332	Splanomagely	252
		Gastric wall thickening	206
Stomach	3373	Stomach cancer	117
Sacrum	3242	Ostoiti	17
Sacium	5272	U U U U U U U U U U U U U U U U U U U	1/

Table 10: The distribution of 54 annotated abnormalities in the test set.

Anatomy	Abnormality	AUC	ACC	Spec	Sens
Adrenal gland	Thickening	64.6	62.1	64.8	59.4
Autenai gianu	Nodule	66.8	64.8	63.0	66.7
Bladder	Diverticulum	85.9	77.8	74.6	81.0
Diaduei	Stones	82.0	75.7	80.0	71.4
	Gas Accumulation	88.7	80.8	78.6	82.9
	Effusion	87.6	80.4	78.7	82.0
	Obstruction	99.5	98.6	97.2	100
Colon	Diverticulum	71.7	68.7	65.3	72.1
Colon	Colorectal Cancer	72.6	64.6	65.6	63.5
	Rectal Cancer	85.4	77.0	82.8	71.2
	Appendicitis	74.9	71.9	70.2	73.7
	Appendicolith	65.7	63.1	62.6	63.5
Fsonhagus	Hiatal Hernia	97.7	92.6	95.1	90
Lsophagus	Varicose Veins	98.8	97.9	97.1	98.7
	Cholecystitis	67.3	62.7	61.1	64.2
Gallbladder	Gallstone	64.6	61.8	58.3	65.4
	Adenomyomatosis	62.6	61.0	57.0	65.0
Heart	Cardiomegaly	95.1	88.7	87.3	90.0
Incart	Pericardial Effusion	76.5	74.4	64.4	84.4
	Atrophy	96.0	91.5	91.1	91.9
Kidney	Cyst	68.6	63.1	64.3	61.8
Ridiley	Hydronephrosis	75.7	69.5	78.1	60.9
	Calculi	57.9	56.6	56.4	56.9
	Steatosis	93.3	85.1	84.6	85.6
	Glisson's Capsule Effusion	86.5	78.9	82.7	75.0
	Metastase	78.8	71.6	70.3	73.0
Liver	Intrahepatic Bile Duct Dilatation	76.8	70.8	68.2	73.5
Liver	Cancer	84.9	79.1	77.8	80.3
	Cyst	62.9	59.4	61.4	57.3
	Abscess	81.8	79.5	75.7	83.3
	Cirrhosis	94.7	88.5	87.5	89.4
	Atelectasis	94.8	89.4	89.4	89.3
.	Bronchiectasis	81.6	74.1	76.0	72.2
Lung	Emphysema	75.0	69.3	68.6	70.0
	Pneumonia	72.8	69.0	79.8	58.2
	Pleural Effusion	86.7	81.3	80.5	82.0
	Pancreatic Cancer	87.0	79.7	80.2	79.3
D	Atrophy	86.4	77.3	76.3	78.4
Pancreas	Pancreatitis	91.0	87.0	93.3	81.8
	Pancreatic Duct Dilatation	11.2	70.9	/5.8	00.0 72.2
	Steatosis	84.9	/5./	/8.1	/3.3
Portal vein	Hypertension	96.8	92.2	95.5	88.9
		90.0	91.8	90.8	92.7
	Gas Accumulation	84.1	71.2	84.7	69.7 76.1
	Enusion	81.5	/4.2	12.4	/0.1
Small Intestine	Divertion	95.2	90.5	92.0	88.3 60.0
	Intussusception	75.0	07.1	78.0	09.0 66 7
	Homenzieme	62 0	60.6	70.0	617
Spleen	Information	03.8	00.0 86 0	39.3 00 4	01./ 01.0
	splanomagely	09.1	00.2 01.0	90.0 82 0	01.0 85 0
	Gastric Well Thickening	92.4	65 7	62.5	68.0
Stomach	Costria Concer	09.0	72.0	02.3	00.9 70.0
Cooming		10.1	12.0	13.1	10.9
Sacrum	Ostelli	07.3	03.0	03.3	00.2
Average		81.3	76.2	76.5	75.8

Table 11: Detailed zero-shot performance of our method on each abnormality.