

The potential of radiomics with PET/CT: study of correlations with the metabolic profile and its discriminative power

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Introduction

The treatment of lung cancer is still a challenging research, partly due to late-stage diagnosis of patients. This leads directly to a high mortality rate, so an **early diagnosis** is a crucial step in the treatment of patients with lung cancer. This research focusses on one of the most common types of lung cancer, more specifically, **early-stage non-small cell lung cancer (NSCLC)**. All patients included in this study underwent a **lobectomy**.

This study examines the **discriminative** potential of combining specific metabolic biomarkers from blood plasma (**metabolomics**) with features out of medical images (**radiomics**). This way metabolomics and radiomics might be at the base to develop a **more patient specific treatment plan** for lung cancer patients.

Research questions

Goals:

1. Combining **metabolomics and radiomics** datasets from NSCLC patients, to unravel the underlying **correlations**.
2. Generate a **discriminative model** between malignant and non-malignant lung lesions, and between adenocarcinoma and squamous cell carcinoma based on the **radiomics** features.

Materials and method

Metabolomics and radiomics:

- 39 patients
- **ACCURATE** tool (creating volumes of interest (VOIs)) and **RADIOMICS** tool (extracting radiomics features)
- 483 radiomics parameters
- 238 metabolic parameters obtained with proton nuclear magnetic resonance ($^1\text{H-NMR}$)
- **Correlation test**



Figure 1: The $^1\text{H-NMR}$ spectroscopic analysis [1]

Radiomics:

- 85 patients
- **ACCURATE** tool (creating VOIs) and **RADIOMICS** tool (extracting radiomics features)
- 483 radiomics parameters out of PET-CT images
- **Correlation test**
- **Logistic regression** to generate discriminative model



Figure 2: The Biograph Horizon detector from Siemens [2]

Results

Table 1: Summary of the total amount of correlated radiomics features to metabolic variables related to plasma glucose and glycerol for all three segmentation methods

	↑ Glucose	↓ Glycerol
First segmentation method (PET/CT)	12 features	8 features
Second segmentation method (PET/CT)	16 features	12 features
Third segmentation method (PET)	17 features	15 features

Table 2: Summary of the sensitivity and specificity of the generated models to discriminate between malignant and non-malignant lung nodules, and between adenocarcinoma and squamous cell carcinoma

	Sensitivity	Specificity
Malignant/non-malignant	0.9474	0.0000
Adenocarcinoma/Squamous cell carcinoma	0.4000	0.9167

Conclusion

- High glucose uptake in tumor cells
- Supporting role gluconeogenesis in normal cells
- **6** features positively correlated to **increased** plasma glucose
- **13** features negatively correlated to **decreased** plasma glycerol
- **Discriminative models:**
- Malignant/non-malignant and adenocarcinoma/squamous cell carcinoma
- **Fragile** due to small patient cohort

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[1]: ProLUNG study

[2]: Siemens Healthineers