

## Introduction

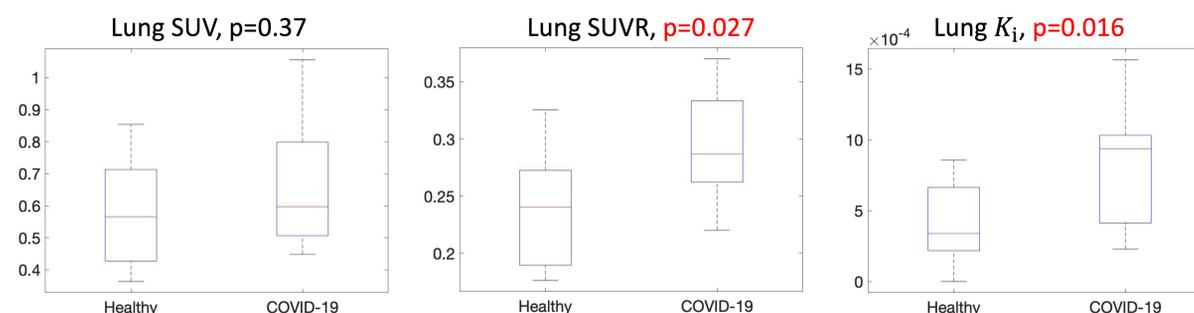
- COVID-19 can affect multiple organs and the prolonged impacts have not been thoroughly investigated.
- Total-body dynamic  $^{18}\text{F}$ -FDG PET, e.g., on the 2-m long uEXPLORER system, when combined with kinetic modeling, allows a quantitative evaluation of metabolism in the entire body.
- In this study, we investigate the metabolic changes in multiple organs of COVID-19 subjects in the early recovery period using total-body dynamic  $^{18}\text{F}$ -FDG PET and kinetic modeling.

## Methods

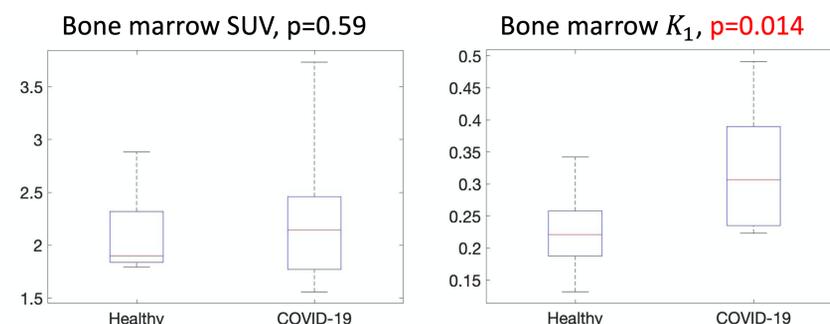
- The study enrolled thirteen healthy subjects and eight recovering COVID-19 patients who were within two months of confirmed diagnosis.
- Each subject had an  $^{18}\text{F}$ -FDG scan on the uEXPLORER system for one hour.
- Regions of interest (ROIs) were placed in multiple organs in the reconstructed total-body images to obtain parameters. The ROI-based PET parameters include the standardized uptake value (SUV), SUV ratio (SUVR) relative to blood,  $^{18}\text{F}$ -FDG rate constants  $K_1 \sim k_4$  by compartmental modeling, and net influx rate  $K_1 = K_1 k_3 / (k_2 + k_3)$ .
- T-tests were performed to examine differences between the two groups over the parameters.
- We further generated parametric images to confirm the ROI-based findings.

## Results

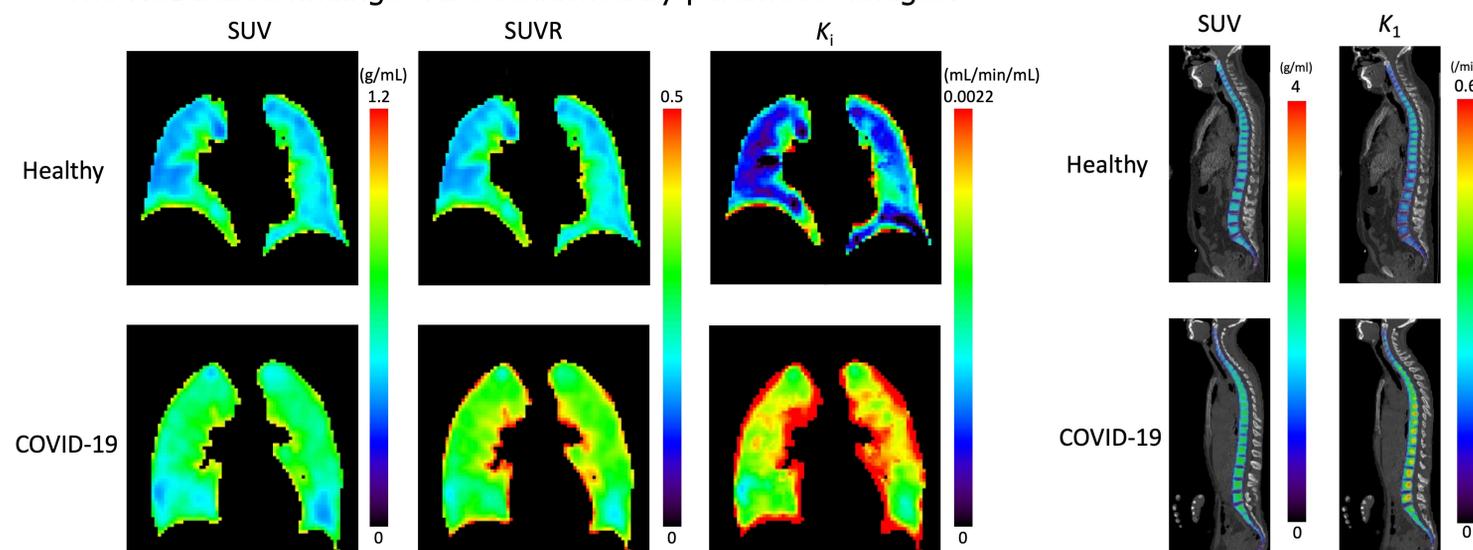
- We detected significant increases in lung SUVR (T-test p-value  $p = 0.027$ ) and  $K_1$  ( $p = 0.016$ ) in the COVID-19 group, while there is no significant difference in SUV (T-test p-value  $p = 0.37$ ) between the two groups.



- For bone marrow, there is a significantly increased  $^{18}\text{F}$ -FDG delivery rate  $K_1$  ( $p = 0.014$ ) in the COVID-19 group but no group difference in SUV ( $p = 0.59$ ).



- The ROI-based findings were confirmed by parametric images.



## Discussion

- The increases in  $^{18}\text{F}$ -FDG lung metabolism (represented by SUVR and  $K_1$ ) and bone marrow  $^{18}\text{F}$ -FDG delivery (represented by  $K_1$ ) may imply prolonged inflammation and immune response during the early recovery.

## Conclusions/Further Study

- We detected increased lung glucose metabolism and bone marrow glucose delivery of recovering COVID-19 patients, which suggests continued impacts in early recovery.
- Kinetic quantification enabled by total-body dynamic  $^{18}\text{F}$ -FDG PET provides a sensitive tool to monitor the metabolic changes in multiple organs.
- The study is still ongoing, and each of the COVID-19 recovering subjects will have a follow-up dynamic scan two months after the first scan.  $^{18}\text{F}$ -FDG kinetics will be further analyzed once the follow-up scan data are obtained.

## Acknowledgements

- This study is supported by NIH grants R01 CA206187, and R01 DK124803.
- UC Davis has a research agreement and a sales-based revenue sharing agreement with United Imaging Healthcare.