

Genetic obesity disorders: Quantitating visceral fat as a predictor of cardiometabolic risk

Lara Musa^{1,2}, Raj Attariwala³, Qing-San Xiang⁴ and William Gibson^{1,2}

¹ Department of Medical Genetics, BC Children's Hospital, Vancouver B.C. ² Diabetes Research Program, Child and Family Research Institute, Vancouver B.C. ³ AIM Medical Imaging, Vancouver B.C. ⁴ Department of Radiology, BC Children's and St. Paul's Hospital, Vancouver B.C.

OBJECTIVES

- Develop a method to quantitate total visceral and total subcutaneous fat mass
- Determine the absolute amount and proportion accounted for by each layer

BACKGROUND

Body Composition

The human body can be modeled as a three compartment system: bone mass, fat mass and lean body mass (muscle, heart, liver and other organs). We know that an increase in a person's body fat is often associated with increased risk for severe diseases, but the *distribution* of body fat also affects one's risk level.

Fat may be classified into two categories: the subcutaneous and the visceral layer. Visceral fat breakdown leads to the release of low-density lipoproteins (LDL) and triglycerides into the bloodstream. Thus, the amount of visceral fat correlates strongly with serious diseases, namely hypertension, diabetes and coronary heart disease.

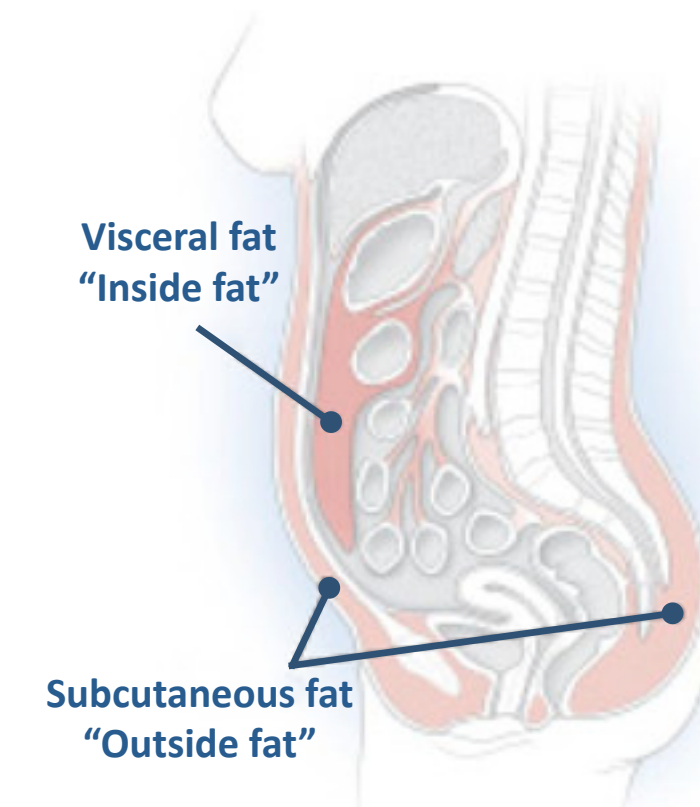
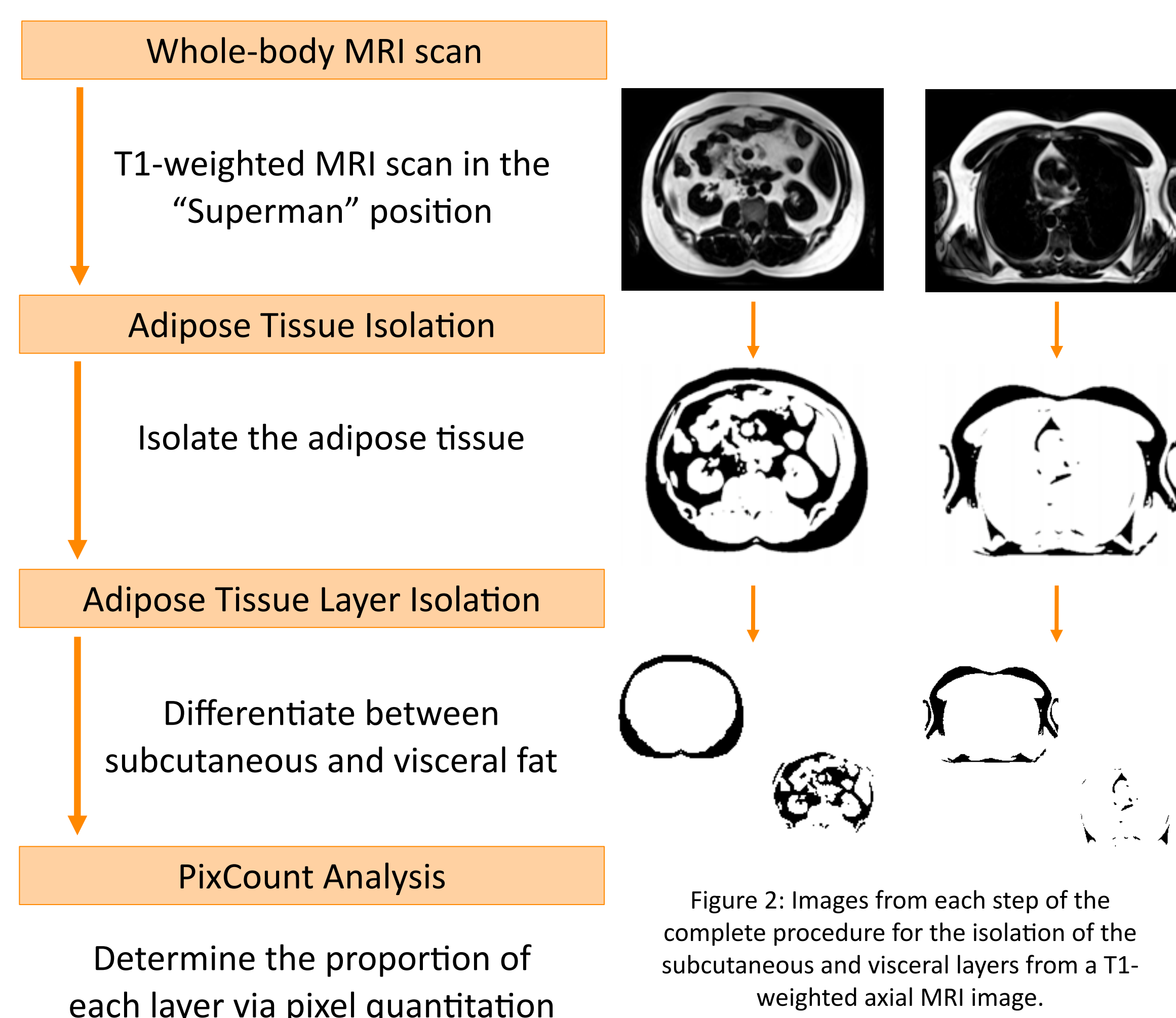


Figure 1: The distribution and accumulation of subcutaneous and visceral adipose tissue (Gather Health 2007).

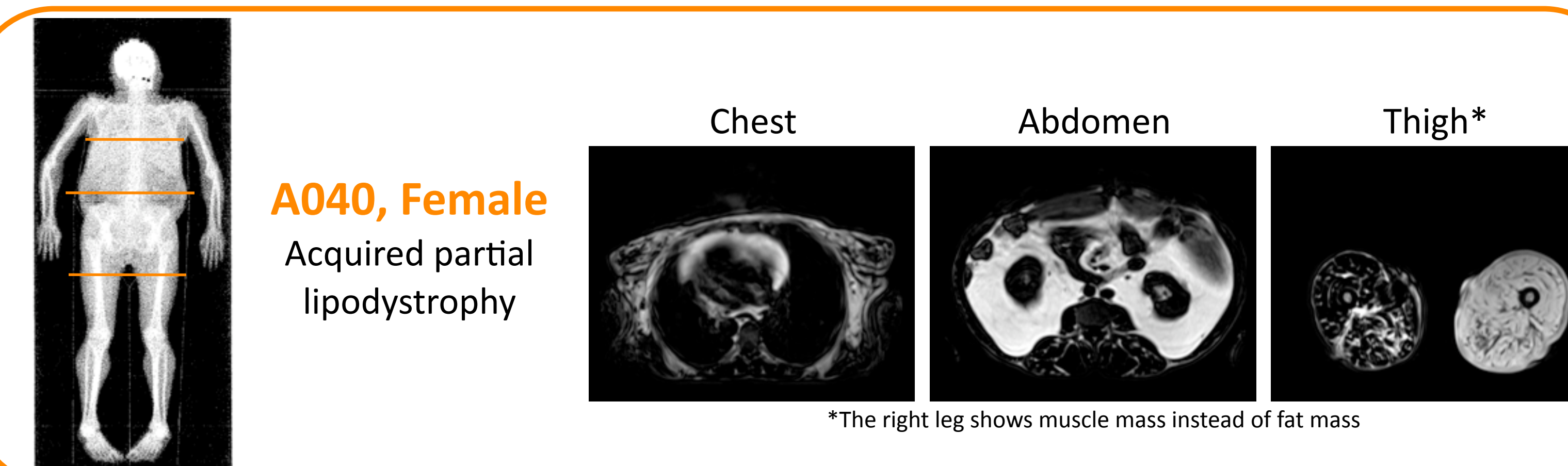
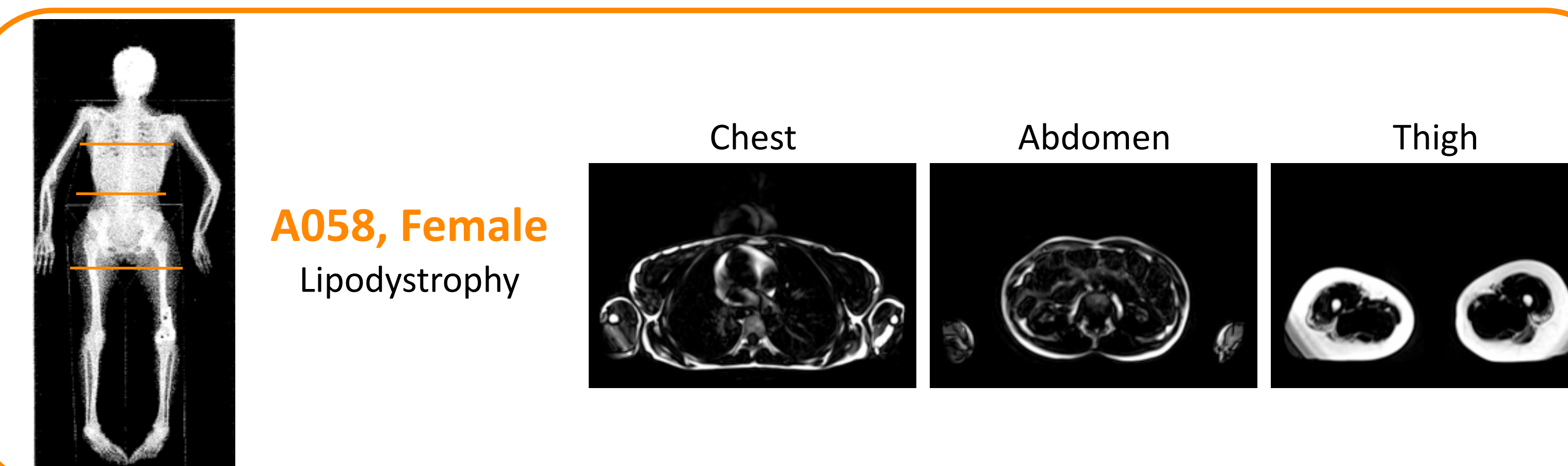
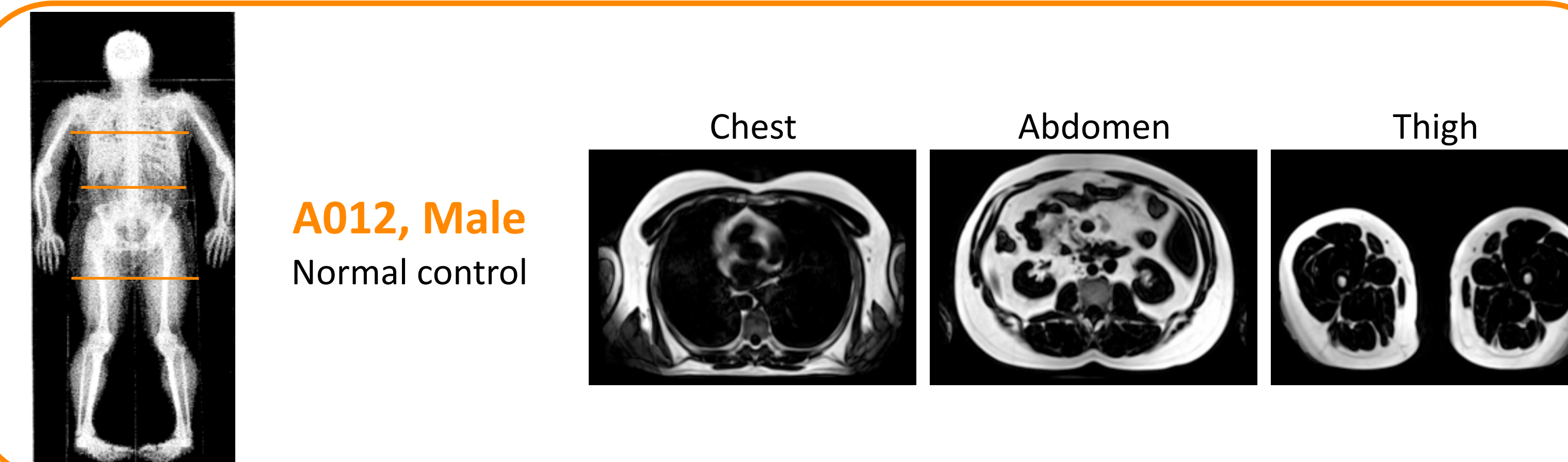
Methods for measuring body composition

Many different techniques are used to measure body composition, in particular body fat percentage. However, these techniques are limited since they do not reliably distinguish visceral from subcutaneous fat. Whole-body MRI captures data on both layers, which must first be separated with computational methods.

METHODS



STUDY SUBJECTS



DISCUSSION & CONCLUSION

- Fat distribution varies between individuals
- Visceral fat percentage cannot be predicted from the total body fat mass among individuals with genetic obesity disorders
- Pixel quantitation of MRI images provides a potential means to segregate fat mass into two compartments

FUTURE DIRECTIONS & APPLICATIONS

Future Directions

- Use of multiple coronal slices, instead of axial slices, may provide the same amount of data with less processing time

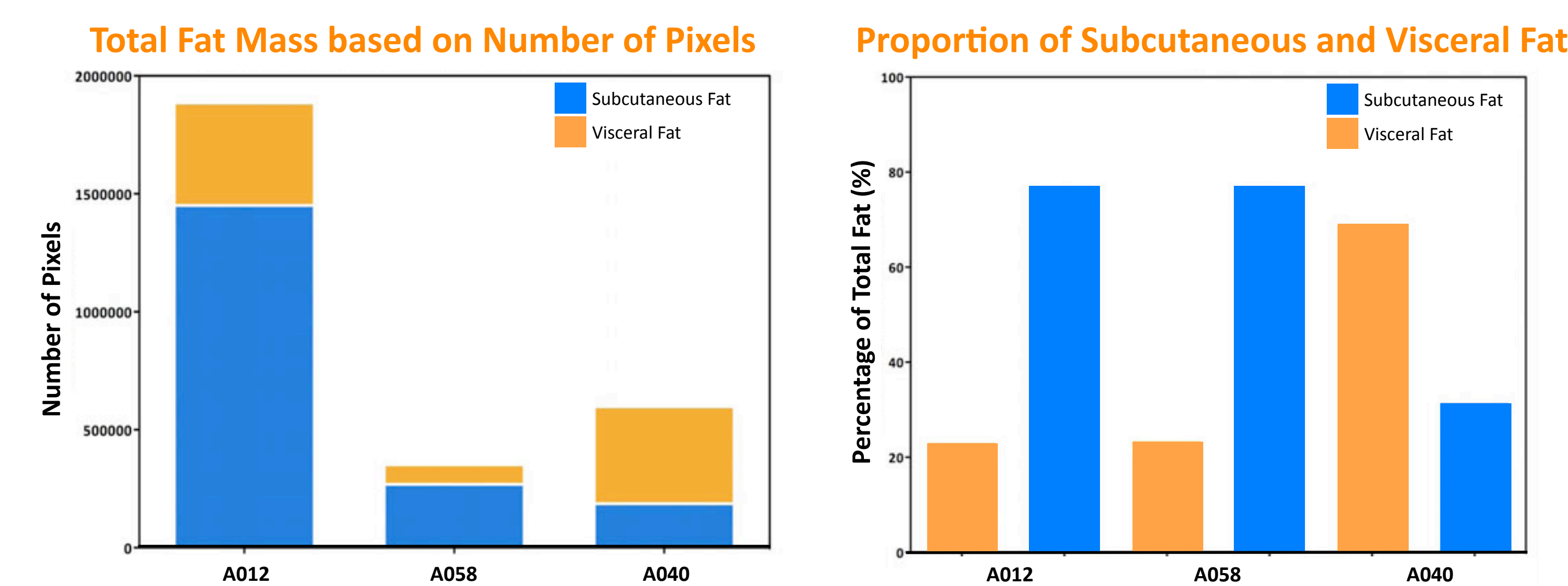
Applications

- Genotype-phenotype correlation**
Identify novel determinants of visceral fat accumulation and the partitioning of fat into the subcutaneous and visceral depots.
- Rare genetic obesity disorders**
Improve understanding of how individuals with rare genetic obesity disorders, such as Prader-Willi syndrome, are affected by visceral adiposity.



Figure 3: An example of a T1-weighted whole body coronal slice image

RESULTS



Total Fat Mass

- A012, male normal control, is shown to have the greatest amount of total fat mass while both females have significantly lower levels of total fat.

Fat Distribution

- Men have a greater proportion of their total fat as visceral fat than do women. A058, female with lipodystrophy, has the same proportion of visceral fat as the male control which is different from the norm for a female.
- A040 has a significant decrease in her subcutaneous adipose tissue, but appears to retain normal amounts of visceral fat.

ACKNOWLEDGEMENTS

Thank you to Dr. Raj Attariwala, Wayne Picker and other staff members from AIM Medical Imaging for conducting the whole-body MRI scans. Special thanks to Dr. Qing-San Xiang for developing the PixCount and DicomSorter programs used for this study.

Dr. Gibson gratefully acknowledges the salary support provided by the Child and Family Research Institute and the Canadian Institutes of Health Research Clinician Scientist programs. This work is supported by the British Columbia Prader-Willi Syndrome Association student research award and CIHR.



Contact Information: Lara Musa at lmusa@cfri.ca
604-875-2000 ext 6783