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## Overall protocol for MicroPOTS LCMS top down proteomics of kidney tissue sections

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Mowei Zhou<sup>1</sup>, James M Fulcher<sup>1</sup>, Yen-Chen Liao<sup>1</sup>, Ljiljana.PasaTolic<sup>1</sup>

<sup>1</sup>Pacific Northwest National Laboratory

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Mowei Zhou

Pacific Northwest National Laboratory

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**We use this protocol and it's working**

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## Abstract

This is the overall workflow for LCMS top down proteomics of kidney functional units from tissue sections using the MicroPOTS platform. The expected outcomes are proteoform identification and quantitation values from selected tissue functional units.

## Troubleshooting



## Tissue collection

- 1 The tissue sections were prepared and shipped from Vanderbilt-TMC following the protocol below:

### Protocol

NAME

### Cryostat Sectioning of Tissues for 3D Multimodal Molecular Imaging

CREATED BY

Jamie Allen

Preview

## Sample preparation

- 2 Functional units (glomerulus, medullary, tubule) were dissected and collected into the microPOTS platform using the method below:

### Protocol

NAME

### Laser Capture Microdissection of Tissue Functional Units for microPOTS Top-Down Proteomics

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James M Fulcher

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## Data Acquisition

- 3 The samples were analyzed by LCMS top down proteomics as described below:



## Protocol

NAME

### Top Down Proteomics Data Collection for Microdissected Kidney Tissue Functional Units

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## Data Analysis

- 4 LCMS datasets were analyzed for proteoform identification and quantitation. The final results are reported.

## Protocol

NAME

### Proteoform Identification and Quantitation with TopPIC and TDPortal for Human Tissues

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