

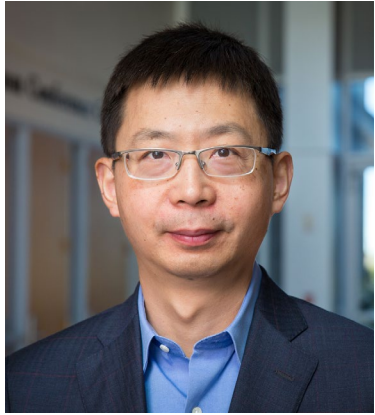
CCF Workshop Kidney (BUKMAP)

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Director, Kidney Translational Research Center



HuBMAP U54 KULMAP Team –Thank You



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Contact PI



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BUKMAP, Co-PI



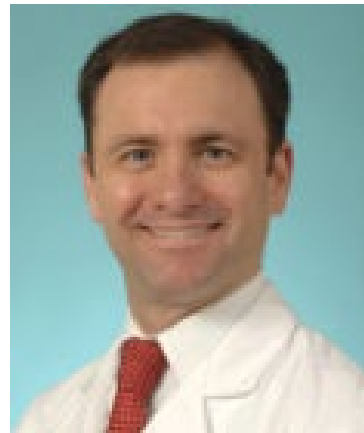
James Hagood@UNC,
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Blue Lake @UCSD
Co-I



Joe Gaut @ WashU
BUKMAP, Co-I



Gloria Pryhuber@Rochester,
LAPMAP; Co-I



Peter Kharchenko @ HMS
Data Core, Co-PI

Project overview

Hagood/Sun
@UCSD

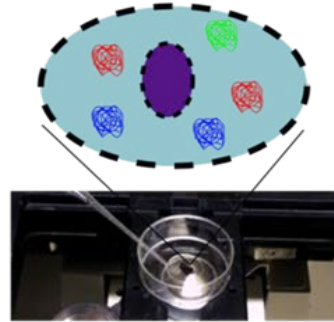
Zhang@UCSD

Kharchenko@HMS

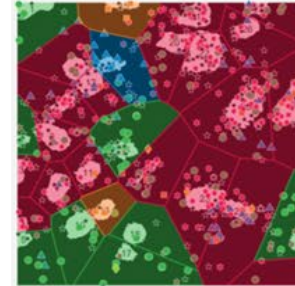
LAPMAP



Tissue blocks

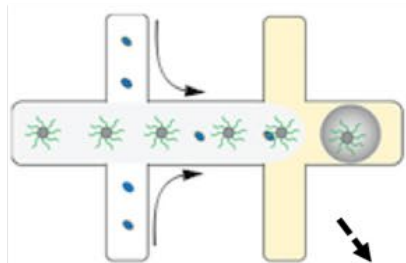
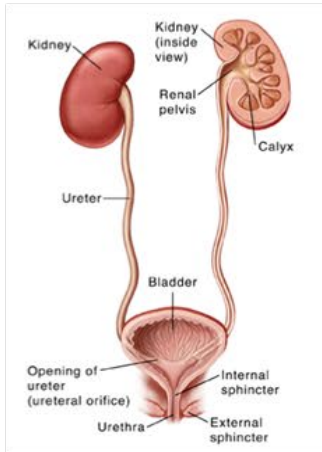


DART-FISH



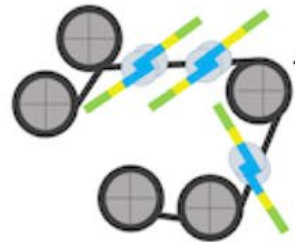
Jain@WashU

Nuclei

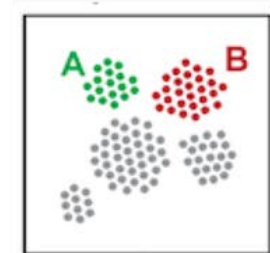


snDrop-Seq

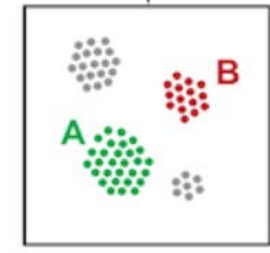
SNARE-Seq



scTHS-Seq



gene/accessible site mapping



Atlases

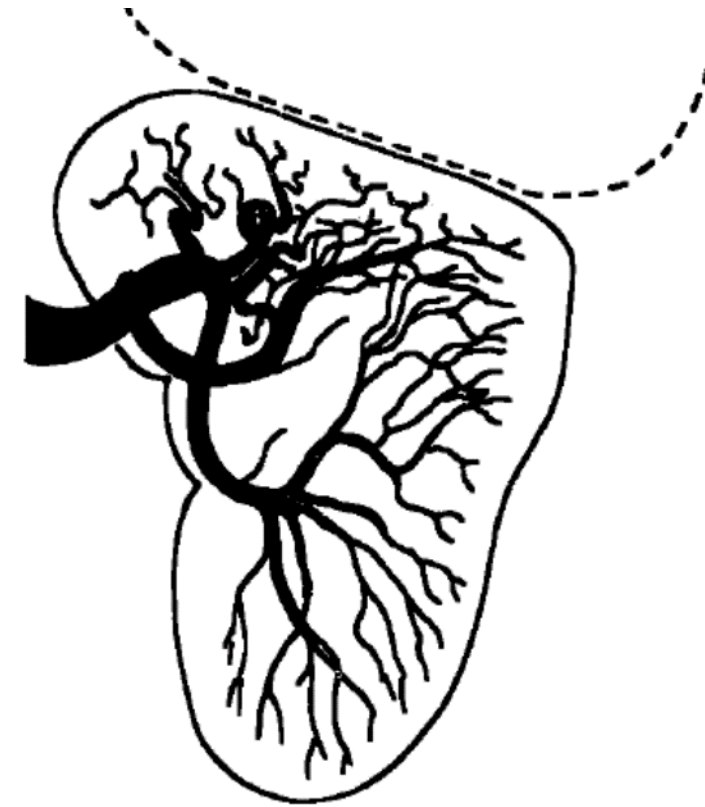
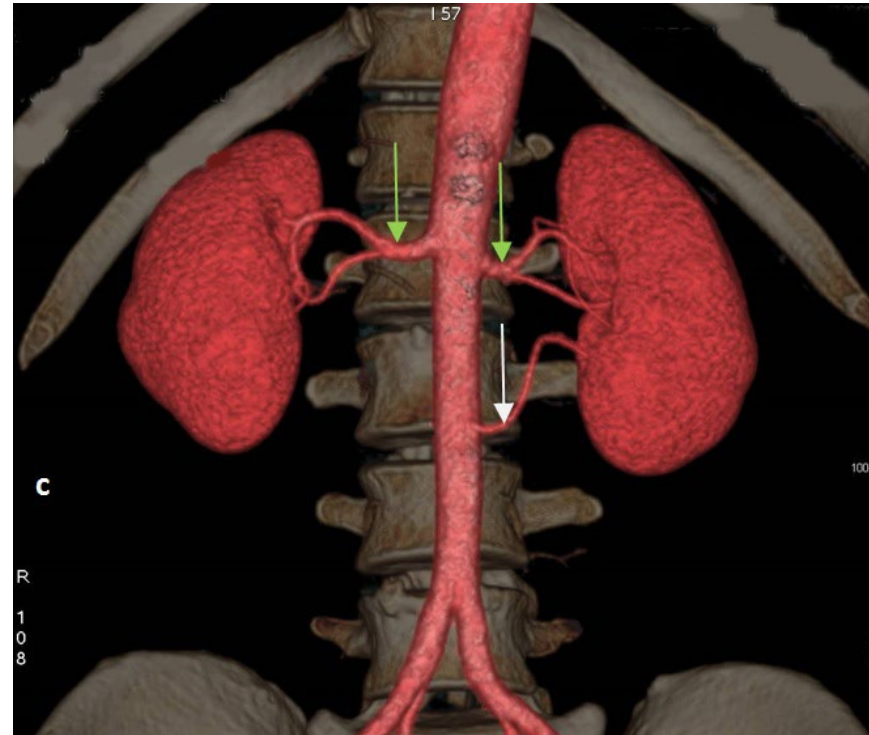
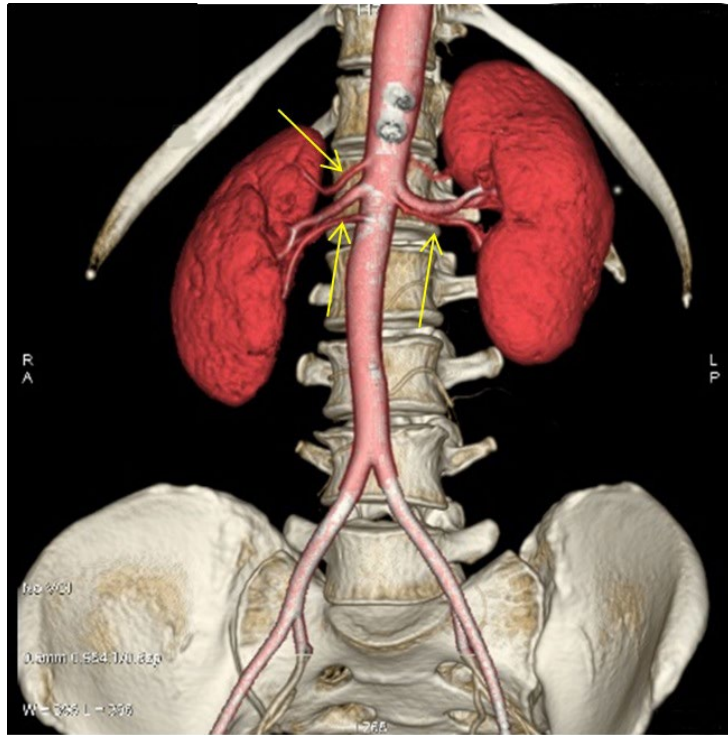
Organ spatial maps

- All cell types;
- Spatial distribution;
- Reference transcriptome;
- Reference chromatin accessibility map;
- Selected proteins

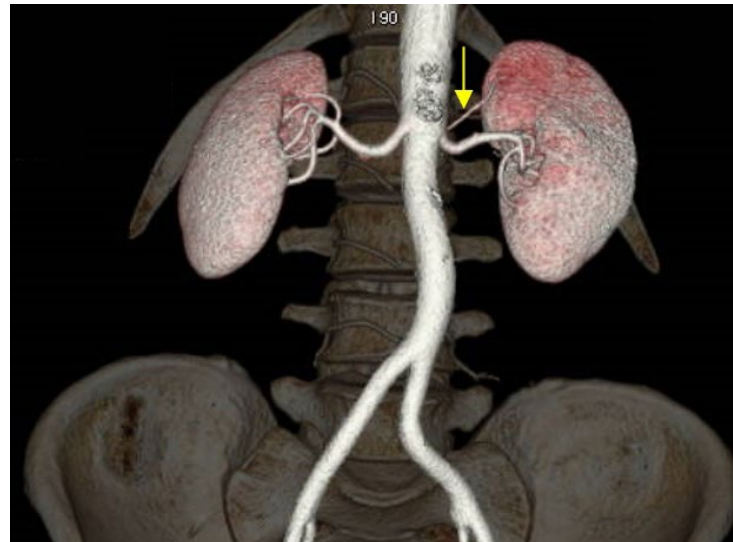
Overview

- Challenges in using common coordinates in kidney
- Approach to minimize challenges
- Glimpse of the workflow for BUKMAP

Variations in Kidney Vasculature: Intrinsic and Extrinsic Factors

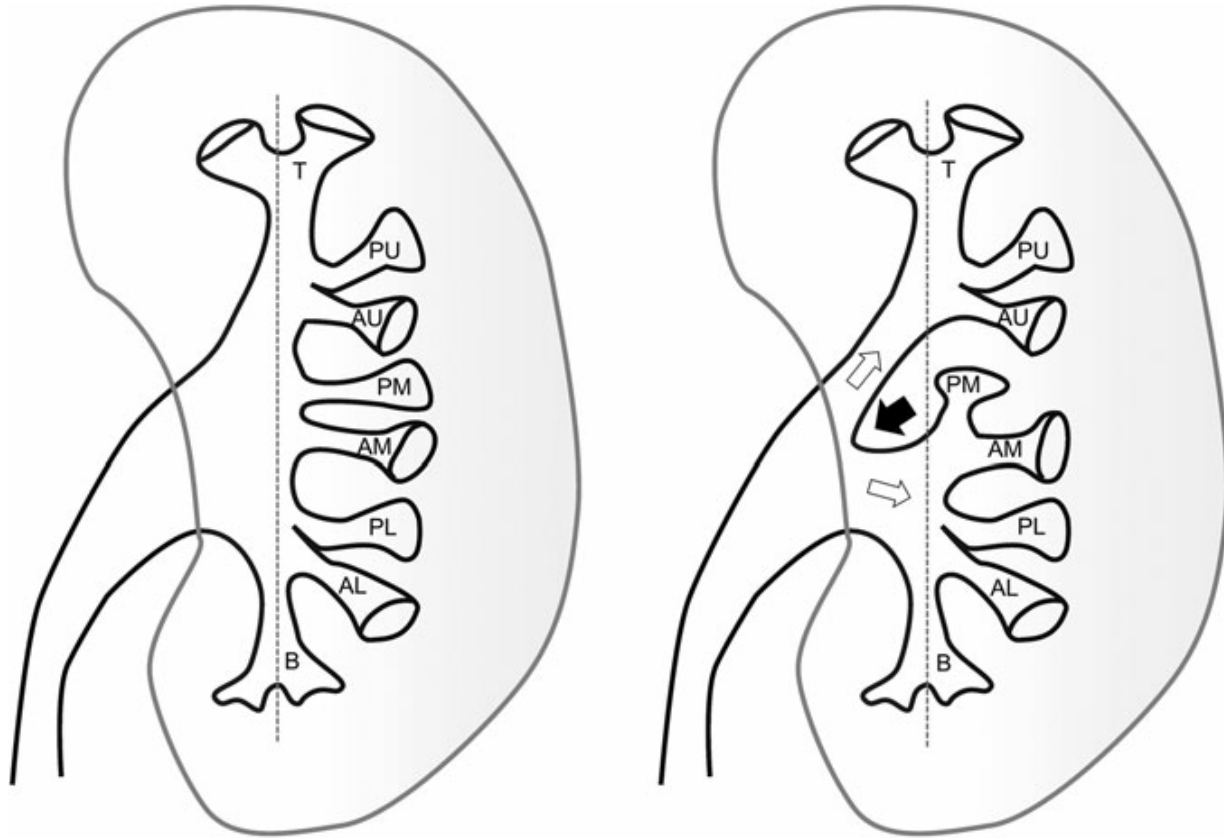


J. Frimann-Dahl (1961) Normal Variations of the Left Kidney, *Acta Radiologica*, 55:3, 207-216, DOI: 10.3109/00016926109174541



Munnusamy et al, Variations in Branching Pattern of Renal Artery in Kidney Donors Using CT Angiography, 2016, *J. Clin Diag Res*

Variations in Kidney Pelvicaliceal System: Intrinsic and Extrinsic Factors



Type I: Single pelvis (58%)

Type II: Divided pelvis (42%)

Proposal for a Simple Anatomical Classification of the Pelvicaliceal System for Endoscopic Surgery

Ryoji Takazawa, MD, PhD, Sachi Kitayama, MD, Yusuke Uchida, MD, Satoshi Yoshida, MD, Yusuke Kohno, MD, and Toshihiko Tsujii, MD, PhD, J of Endourology, 2018

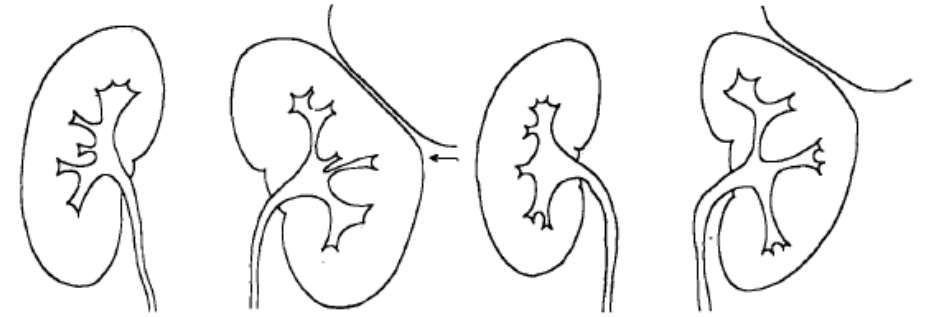


Fig. 8. Diagrammatic representations of typical normal variations of the calyces of the middle lobe of the left kidney.

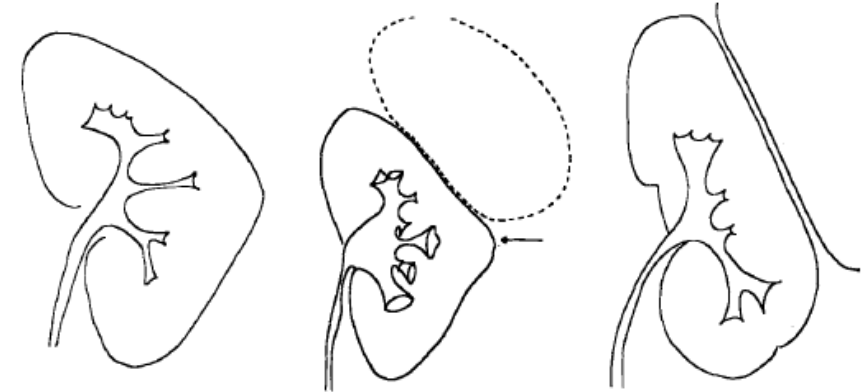


Fig. 9. Drawings showing influence of the spleen upon the renal pelvis.



Fig. 10. Normal variation of the upper, middle and lower lobe calyces.

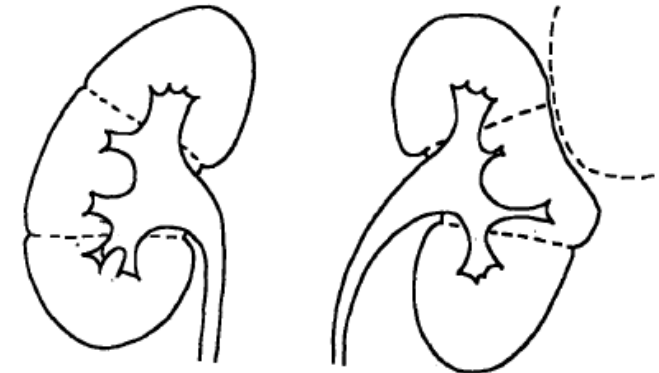
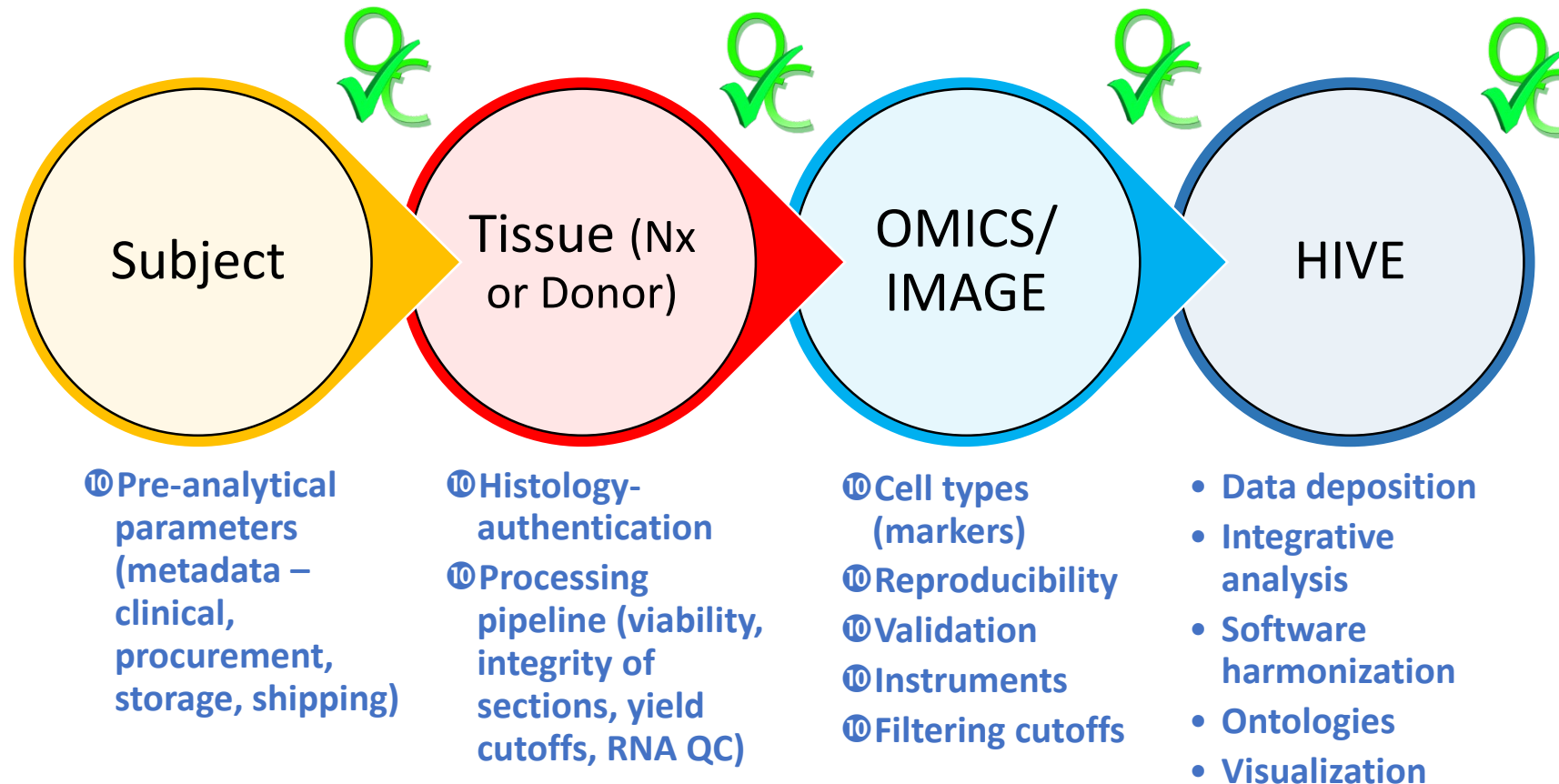
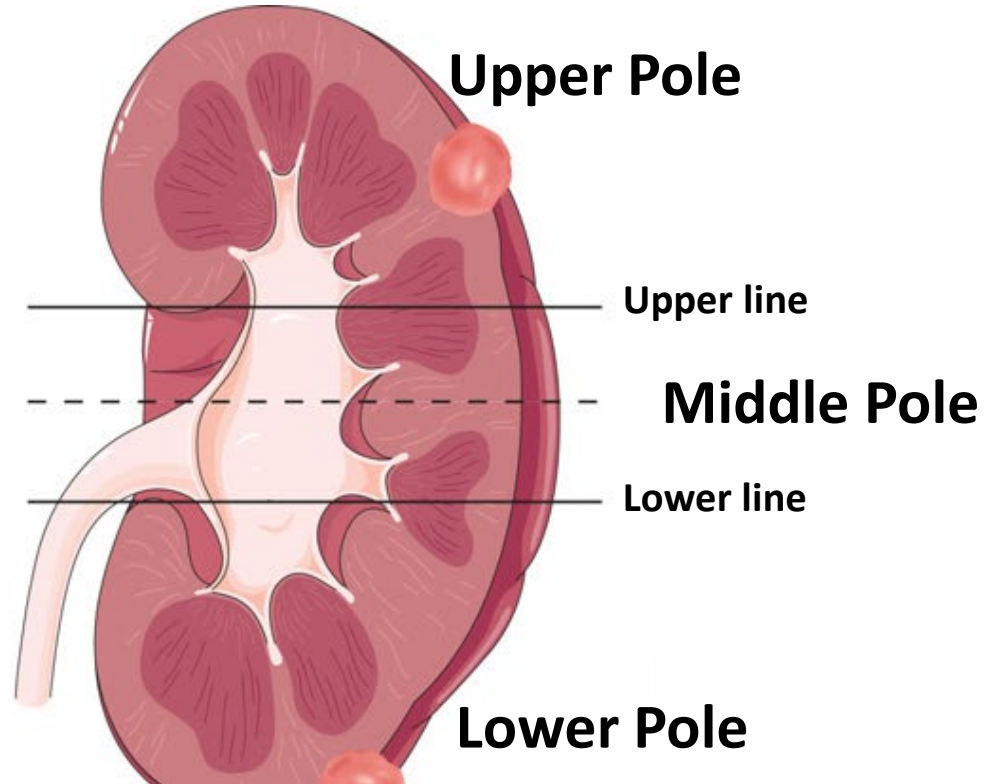


Fig. 11. Schematic drawing showing bulging of lateral contour of a lobulated left kidney.

Quality Control and Standardization = effective CCF

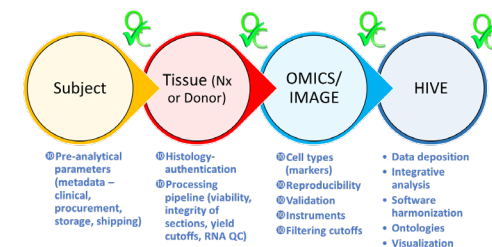


Gross Registration Using polar lines



The R.E.N.A.L. Nephrometry Score: A Comprehensive Standardized System for Quantitating Renal Tumor Size, Location and Depth

Alexander Kutikov and Robert G. Uzzo*, J. of Urology 2009

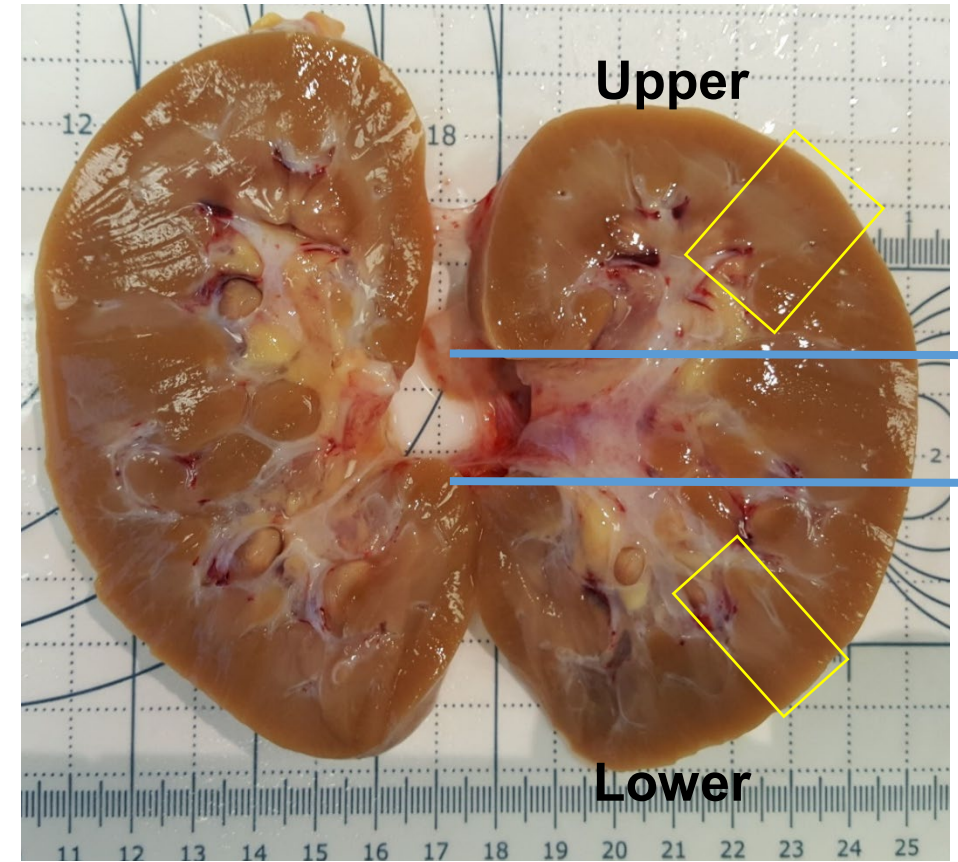
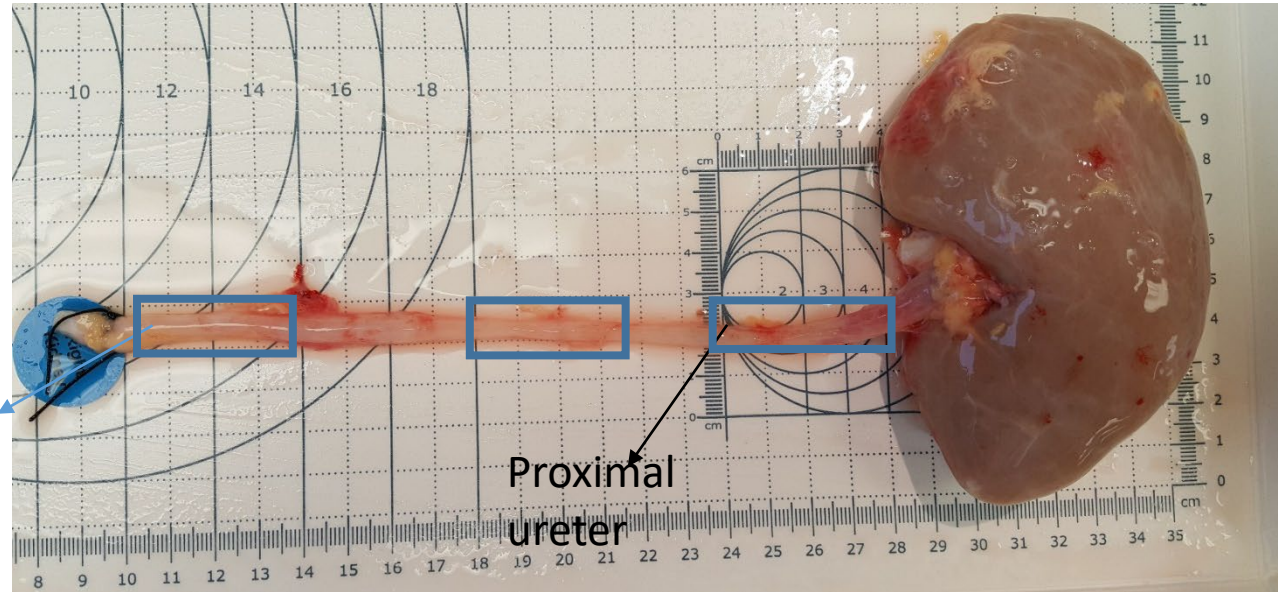


EXAMPLE

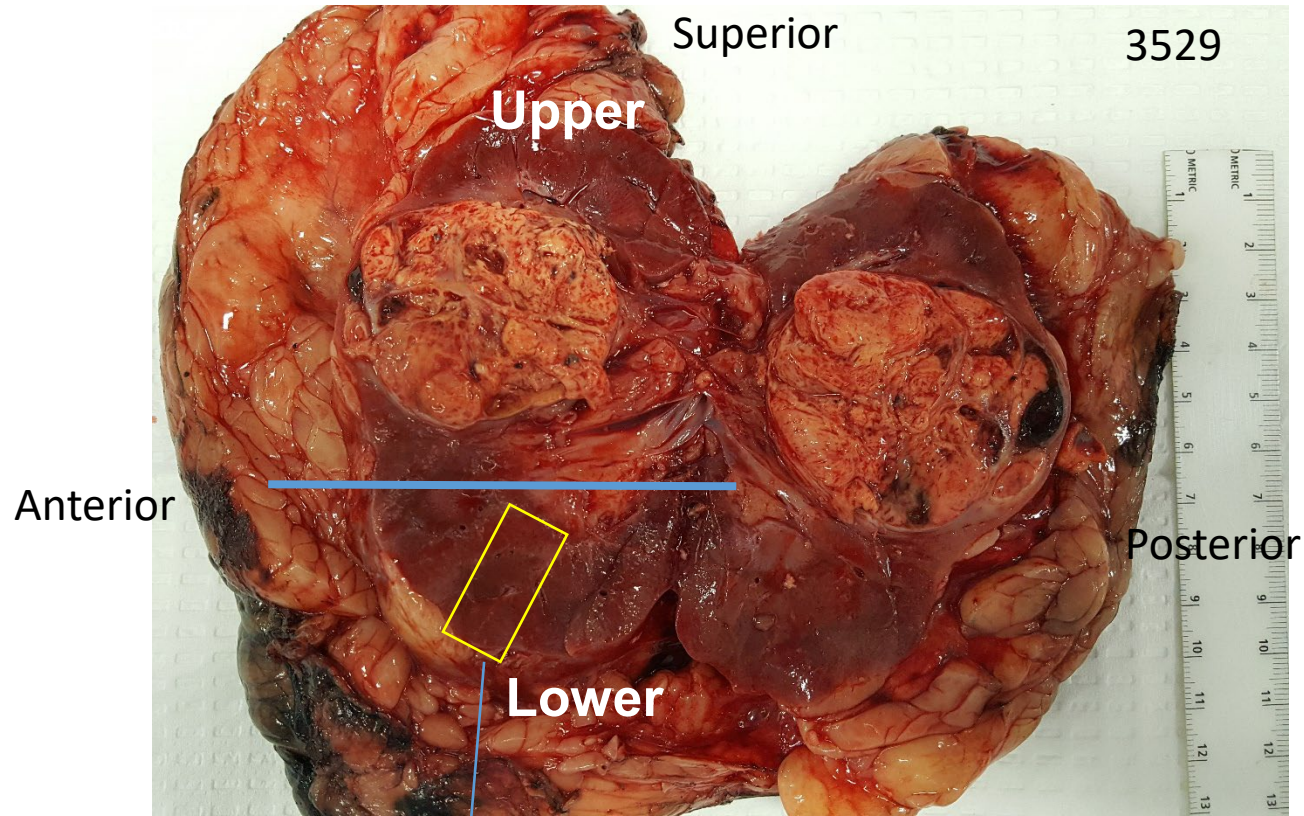
3505

Distal ureter

Proximal ureter



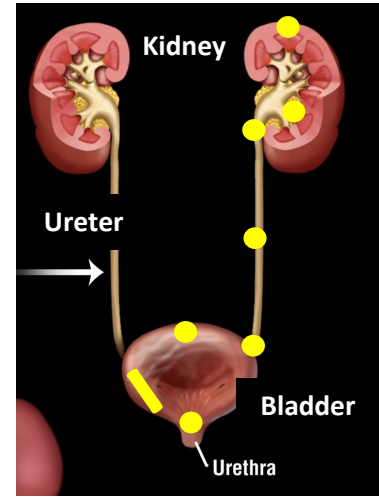
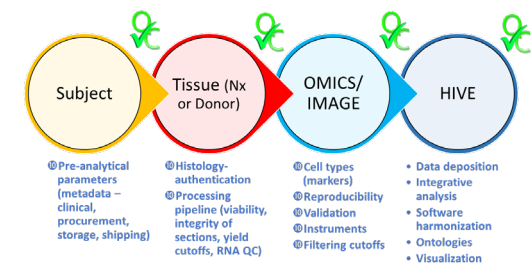
EXAMPLE



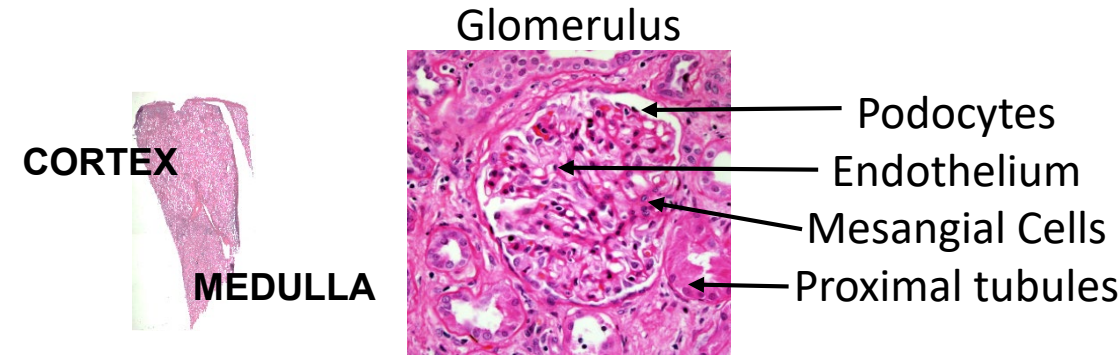
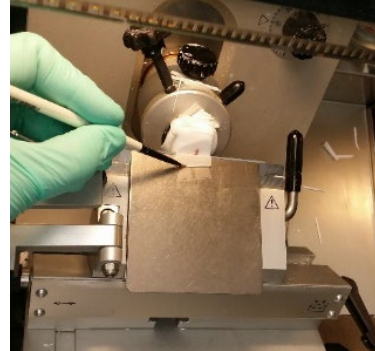
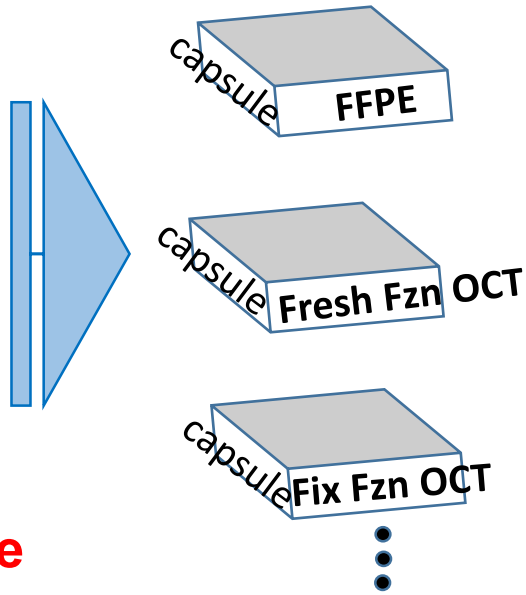
C-M1 3.8cm
from the
posterior

Depth 4.5cm.

Workflow



Gross reference



Histological reference

Chromatin accessibility (scTHS-Seq)

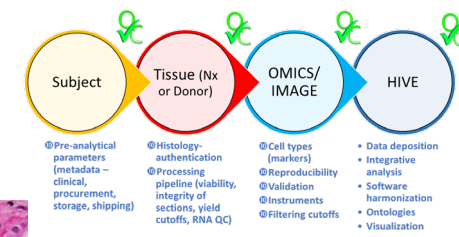
Single nucleus RNA seq (SPLIT-Seq)

3D RNA spatial mapping (DART-FISH)

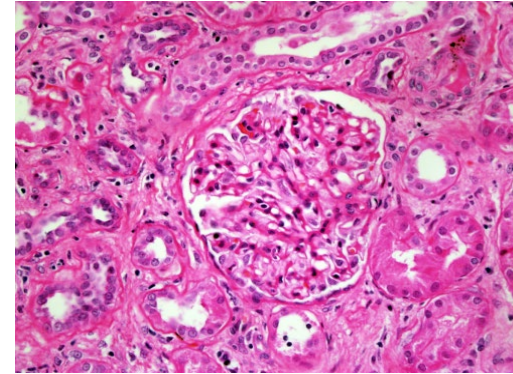
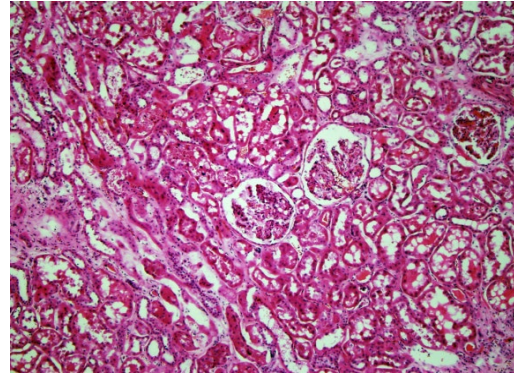
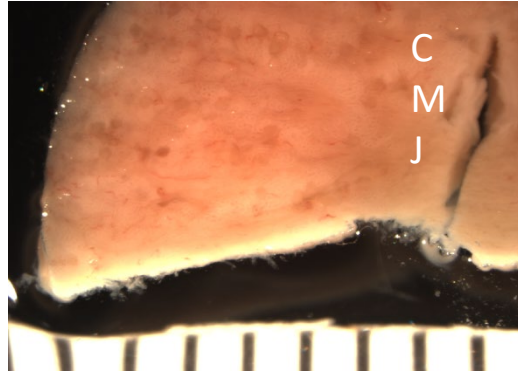
Cell type/state reference

Structure/region reference

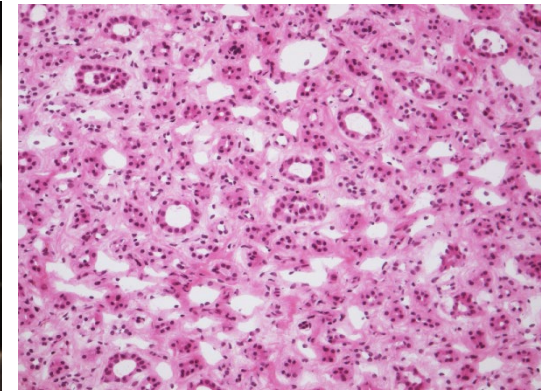
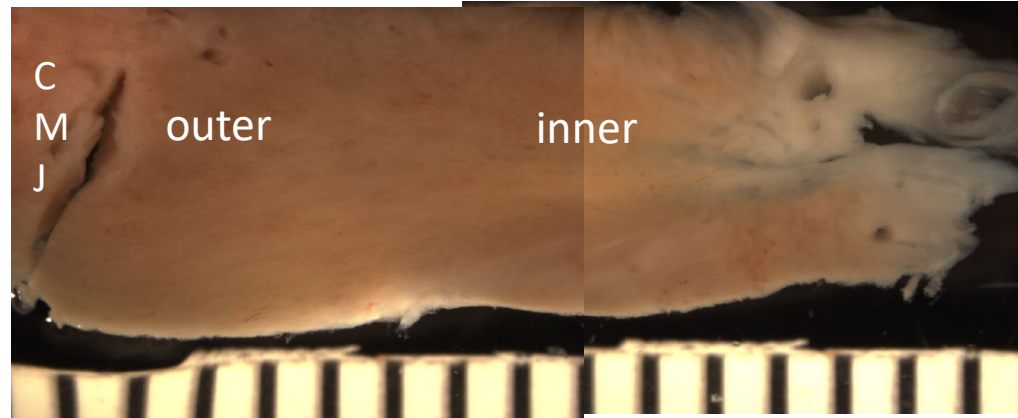
Structure - microscopic



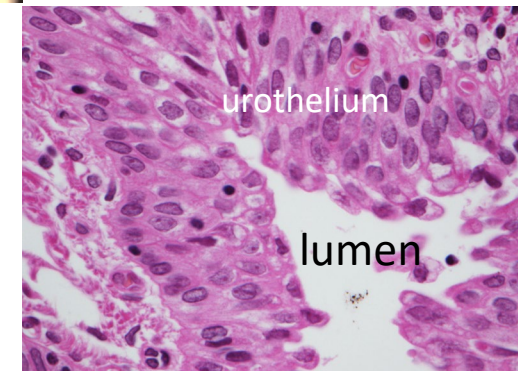
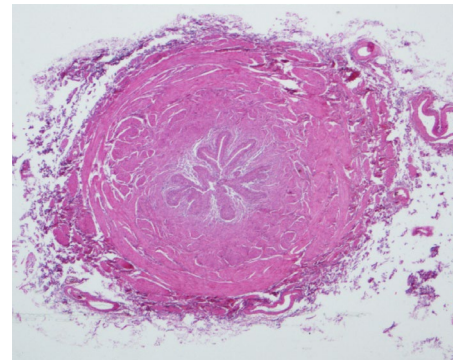
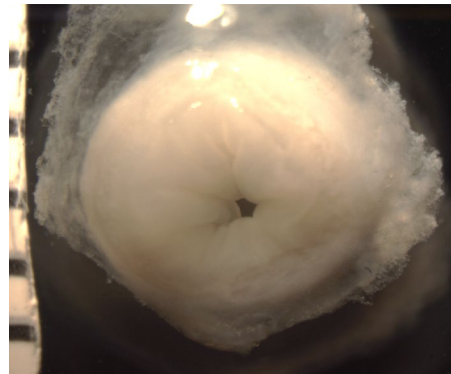
Cortex



Medulla

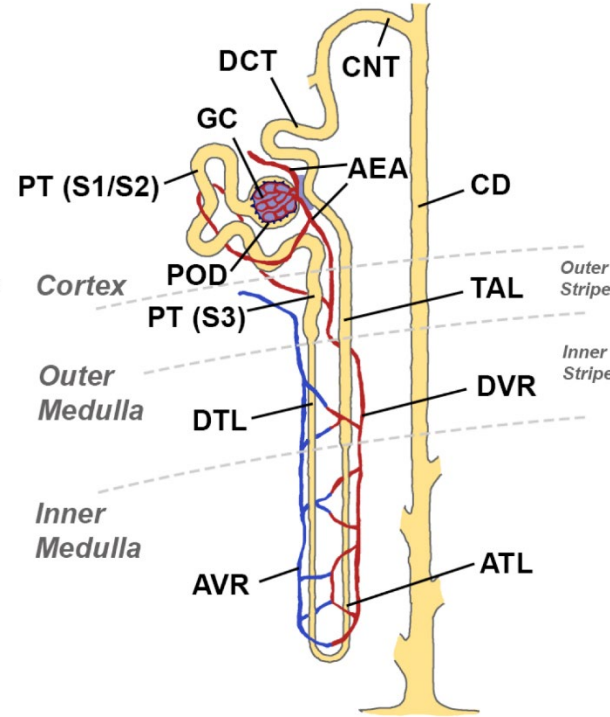
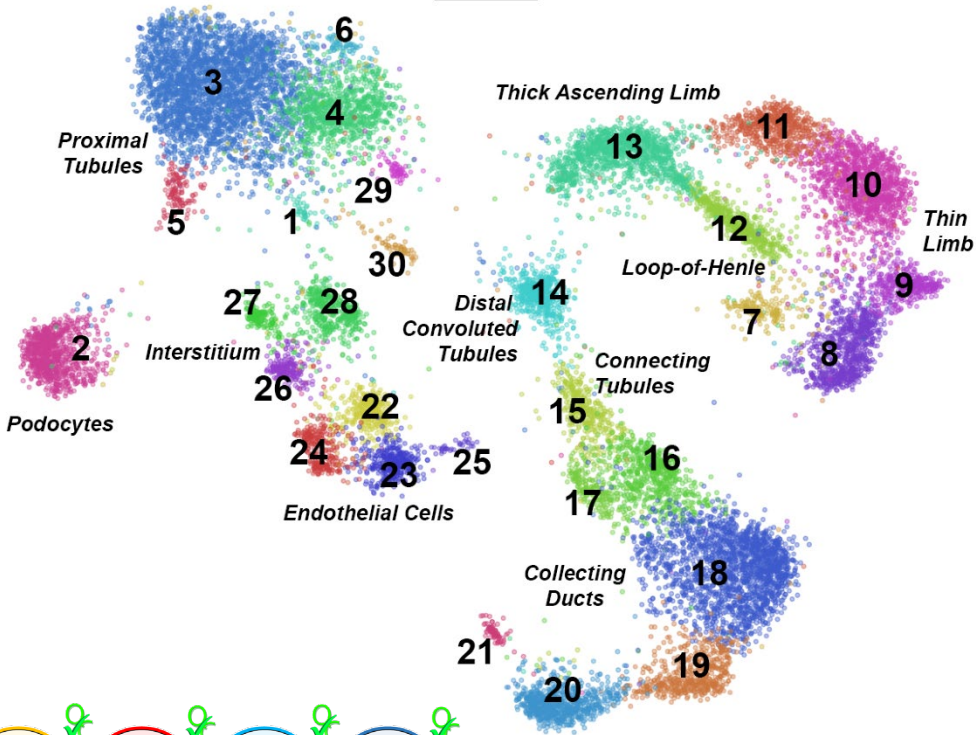


Ureter



snDrop-Seq: Clustering and Initial Annotation

UMAP



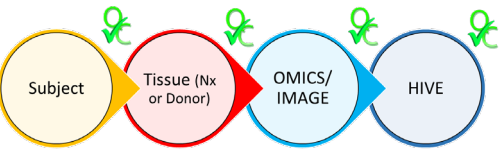
Cluster	Abbn	Annotation	# Nuc	Region Sampled
1	EPC	Epithelial Cells	55	Both
2	POD	Podocytes	859	Both
3	PT-1	Proximal Tubule Ep. Cells (S1/S2)	3238	Both
4	PT-2	Proximal Tubule Ep. Cells (S2)	920	Both
5	PT-3	PT Ep. Cells - <i>Inflam. Response</i>	124	Both
6	PT-4	PT Ep. Cells - Fibrinogen+ (S3)	86	Both
7	PT-5	Proximal Tubule Ep. Cells (S3)	219	Both
8	DTL	Descending Thin Limb	844	Both
9	ATL-1	Thin Ascending Limb	431	Both
10	ATL-2	Thin Ascending Limb	1325	Both
11	ATL-3	Thin Ascending Limb	623	Both
12	TAL-1	<i>Thick Ascending Limb - Surgery</i>	536	Both
13	TAL-2	Thick Ascending Limb	1217	Both
14	DCT	Distal Convoluted Tubule	568	Both
15	CNT	Connecting Tubule	395	Both
16	PC-1	Collecting System - PCs (Cortex)	663	Both
17	PC-2	<i>Collecting System - PCs - Stressed</i>	208	Both
18	PC-3	Collecting Duct - Principal Cells	1970	Both
19	IC-A2	Collecting Duct - Intercalated Cells	540	Both
20	IC-A1	Collecting Duct - Intercalated Cells	772	Both
21	IC-B	Collecting Duct - Intercalated Cells	84	Both
22	EC-1	Endothelial Cells - Glomerular Cap.	344	Both
23	EC-2	Endothelial Cells - AVR	366	Both
24	EC-3	Endothelial Cells - AEA & DVR	281	Both
25	EC-4	Endothelial Cells	42	Both
26	MC	Mesangial Cells	215	Both
27	vSMC/P	Vas. Sm. Muscle Cells/Pericytes	168	Both
28	INT	Interstitial	410	Both
29	Unk	Unknown - Novel PT (S2) Ep. Cells?	68	Both
30	IMM	Immune Cells - Macrophages	88	Both

Region Sampled

0 1

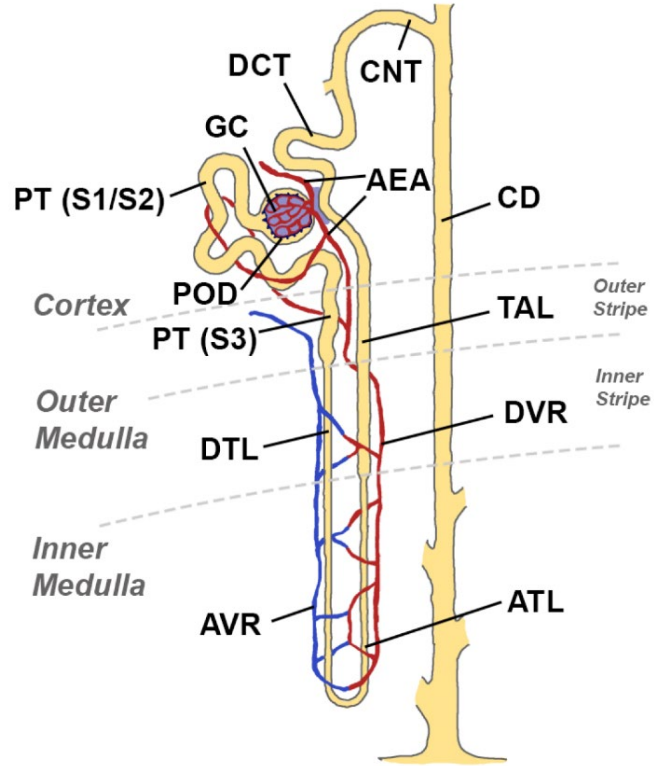
Cortex
Medulla
Both

Standardize segment names, cell types, markers

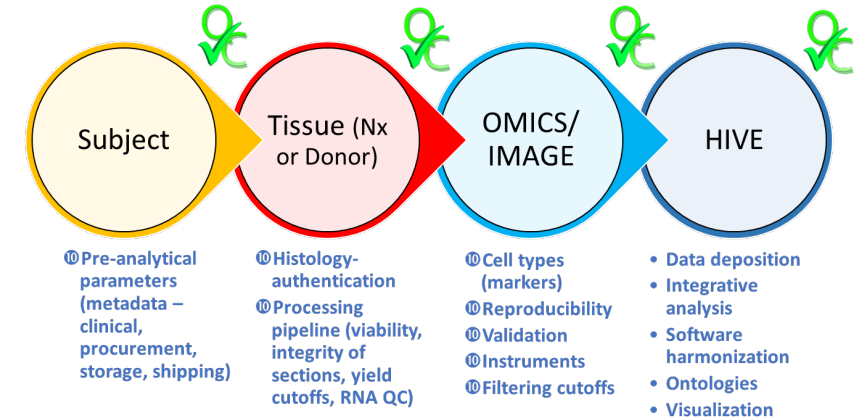


- Pre-analytical parameters (metadata - clinical, procurement, storage, shipping)
- Histology - authentication
- Processing pipeline (viability, integrity of sections, yield cutoffs, RNA QC)
- Cell types (markers)
- Reproducibility
- Validation
- Instruments
- Filtering cutoffs
- Data deposition
- Integrative analysis
- Software harmonization
- Ontologies
- Visualization

Harmonize – VUMC and BUKMAP



Metadata (many levels)
Anatomical coordinates
Structures
Regions
Cell types
Markers



There has been significant progress, but some more challenges