

# Core Facilities

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## SEARCHABLE CORE LISTING

### Core Facilities

## Core Facilities

*Please remember to recognize the contributions of core facility personnel to your research project when appropriate. ABRF Recommended Guidelines for Authorship on Manuscripts can be found on the [Core Facilities Policies and Guidelines Resources](#) page of this website.*

Below is an alphabetical listing of our core facilities.

Each core facility's name links to their individual website. Clicking on the '[Profile/Services](#)' in each listing brings up a brief profile of the core, as well as a link to their equipment list if applicable.

### ***This is a searchable database.***

You can search the listing either by broad **categories**, specific **keywords** such as 'sequencing' or 'RNA', by **director name**, or by type of **equipment**, such as 'PCR'.

Please click on the sentence below 'Search for a Core Facility by Category or Keyword' to bring up the search function. Contact the core facility for more information and to discuss your project.

### [Search for a Core Facility by Category or Keyword](#)

#### [Animal Clinical Chemistry and Gene Expression Laboratories](#)

Contact: [Hyung-Suk Kim, Director](#)

**Facility Overview:** [Profile/Services](#) [Click on Profile/Services for overview](#)

#### [Animal Histopathology Core Facility \(Mice\)](#)

Contact: [Dawud Hilliard, Director](#)

**Facility Overview:** [Profile/Services](#) [Click on Profile/Services for overview](#)

# UNC Cores of Pathology Interest

- Animal Histopathology Core\*
- Translational Pathology Laboratory\*
- Center for Gastrointestinal Biology and Disease (CGIBD): Histology Core
- Histology Research Core
- Tissue Procurement Core
- Microscopy Services Laboratory\*
- Biomedical Research Imaging Center (BRIC): Small Animal Imaging
- Injury Biomarker Core\*
- Animal Clinical Chemistry and Gene Expression Laboratories\*
- Flow Cytometry Core

# Animal Histopathology Core

## UNC, Lineberger Comprehensive Cancer Center

- Faculty Director
  - Stephanie Montgomery (2014)
- Facility Director
  - Dawud Hilliard (2014)
- 3 Full time histotechnologists
  - Lily Wai – Histotechnologist (2009)
  - Traci Raley – Histotechnologist (2013)
  - Amanda Brown – Histotechnologist (2015)



# Animal Histopathology Core

UNC, Lineberger Comprehensive Cancer Center

## Comprehensive histopathology services

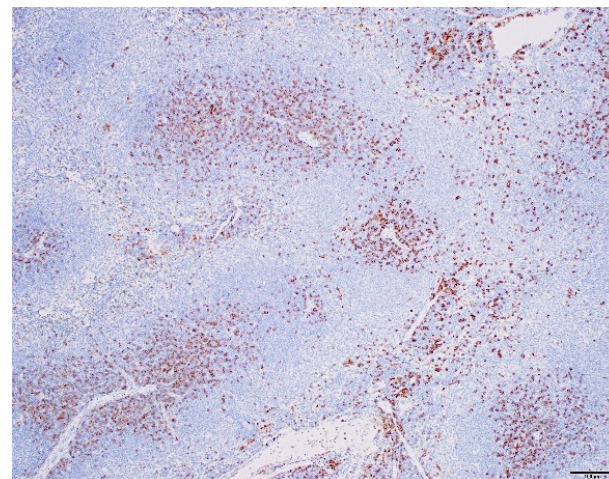
- Tissue processing and embedding
- Histology sectioning (paraffin or frozen)
- Routine and Special Stains (35+)
- Immunohistochemistry & Immunofluorescent stains
  - Multicolor, multiplex IHC or IF, automated
  - 45+ protocols available (FFPE, Mouse)
  - Investigator supplied antibody
- TUNEL staining
- Pathology consultation



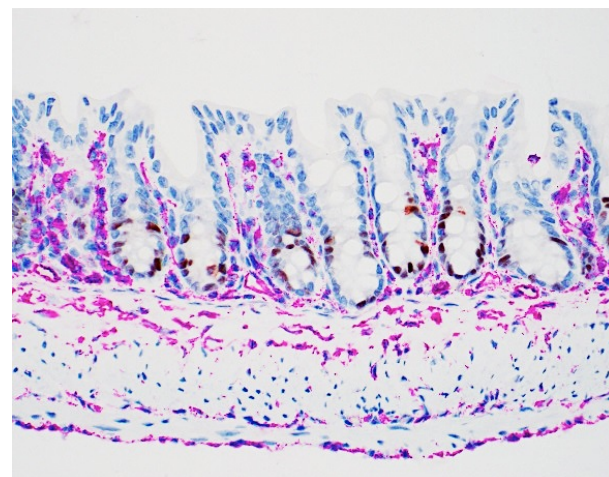
# Animal Histopathology Core

## UNC, Lineberger Comprehensive Cancer Center

- Alpha Fap
- Arginase 1
- AT8
- B220 (CD45R)
- Beta-Catenin
- BrdU
- Calbindin
- Cleaved Caspase-3
- CD11b
- CD26
- CD206 (mannose)
- CD3
- CD31
- **CD4**
- CD45
- CD47
- CD49b(Alpha 2)
- **CD8**
- CD95 (Fas)
- Epcam
- Fas(C18C12)
- F4/80
- FoxP3
- HT7
- iNOS
- GFP
- Glut 2
- Ly6G (Gr1)
- Ki-67
- MCI
- Myeloperoxidase
- Phospho Akt
- Phospho-Histone H2A.X
- PTEN
- RFP (tdTomato, ds Red)
- S100
- SARS N antigen
- Stat 6
- Survivin
- Tau 12
- Vimentin



CD8 IHC, mouse spleen



Ki-67 & vimentin, mouse colon





# Animal Histopathology Core

## UNC, Lineberger Comprehensive Cancer Center

### Histology Pricing

#### AHP UNC Investigator

#### Histo Lab NCSU

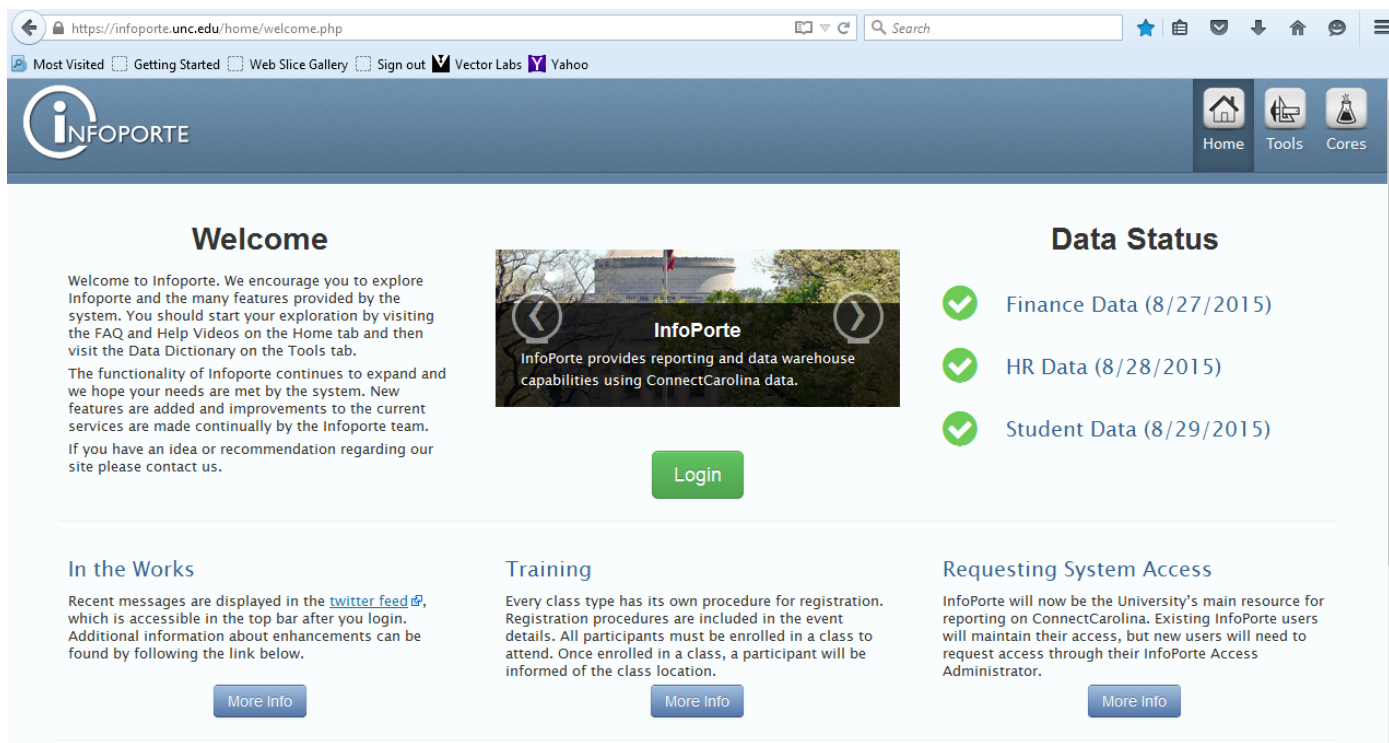
Process & Embed	\$3.16	
Section 1 slide	\$2.92	\$3.90
H&E stain	\$3.30	\$4.60
<b>1 H&amp;E slide from fixed tissue</b>	<b>\$9.38</b>	<b>\$8.30</b>
Special Stain - Level 1	\$3.60	\$14.40
Special Stain - Level 2	\$4.92	
Special Stain - Level 3	\$8.55	\$15.40
<b>IHC single color</b>	<b>\$20.54</b>	<b>\$32.30(27.10)</b>
IF single color	\$19.98	
Dual color IHC	\$34	
Tech time (Special req)	\$43.74/hr	
Frozen section	Tech rate+dispos	\$8.00/Tissue

# Animal Histopathology Core

## UNC, Lineberger Comprehensive Cancer Center

### Submitting an order

- First-time user: contact us to create an account
- Submit an online request
- Drop off or ship specimens



The screenshot shows the InfoPorte website interface. At the top is a navigation bar with the InfoPorte logo, a search bar, and links for Home, Tools, and Cores. Below the navigation bar is a 'Welcome' section with a detailed message about the system's functionality and a 'Login' button. To the right of the welcome message is a 'Data Status' section showing three items: Finance Data (8/27/2015), HR Data (8/28/2015), and Student Data (8/29/2015), each with a green checkmark. Below the welcome message are three sections: 'In the Works', 'Training', and 'Requesting System Access', each with a 'More Info' button.

<https://infoporte.unc.edu/home/welcome.php>

Most Visited Getting Started Web Slice Gallery Sign out Vector Labs Yahoo

**INFOPORTE**

Home Tools Cores

### Welcome

Welcome to Infoporte. We encourage you to explore Infoporte and the many features provided by the system. You should start your exploration by visiting the FAQ and Help Videos on the Home tab and then visit the Data Dictionary on the Tools tab.

The functionality of Infoporte continues to expand and we hope your needs are met by the system. New features are added and improvements to the current services are made continually by the Infoporte team.

If you have an idea or recommendation regarding our site please contact us.

**InfoPorte**

InfoPorte provides reporting and data warehouse capabilities using ConnectCarolina data.

Login

### Data Status

- ✓ Finance Data (8/27/2015)
- ✓ HR Data (8/28/2015)
- ✓ Student Data (8/29/2015)

### In the Works

Recent messages are displayed in the [twitter feed](#), which is accessible in the top bar after you login. Additional information about enhancements can be found by following the link below.

More Info

### Training

Every class type has its own procedure for registration. Registration procedures are included in the event details. All participants must be enrolled in a class to attend. Once enrolled in a class, a participant will be informed of the class location.

More Info

### Requesting System Access

InfoPorte will now be the University's main resource for reporting on ConnectCarolina. Existing InfoPorte users will maintain their access, but new users will need to request access through their InfoPorte Access Administrator.

More Info

# Animal Histopathology Core

UNC, Lineberger Comprehensive Cancer Center

## LCCC Animal Histopathology Core

(919) 966-3653

[lcccanimalistopathology@med.unc.edu](mailto:lcccanimalistopathology@med.unc.edu)

333 S. Columbia, 426 MacNider Hall

University of North Carolina

Chapel Hill, NC 27599

Stephanie A. Montgomery, PhD, DVM, Dipl. ACVP,

[stephanie\\_montgomery@med.unc.edu](mailto:stephanie_montgomery@med.unc.edu)

Dawud Hilliard, MPM, HTL (ASCP) QIHC,

[dawud\\_hilliard@med.unc.edu](mailto:dawud_hilliard@med.unc.edu)



# Microscopy Services Laboratory

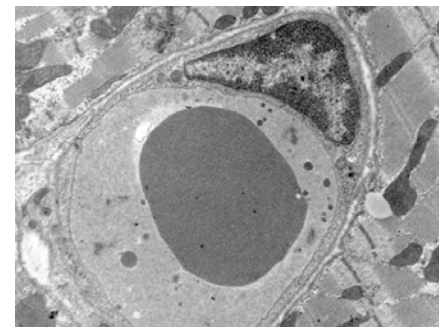
## UNC, Lineberger Comprehensive Cancer Center

Dr. Ariel, PhD, Director

(919) 966-2413, [pablo\\_ariel@med.unc.edu](mailto:pablo_ariel@med.unc.edu)

- Electron microscopy
  - TEM, Immuno-TEM, negative staining EM, SEM
- Light Microscopy
  - Zeiss LSM 710 Spectral Confocal Laser Scanning Microscope
  - Zeiss LSM 700 Confocal Laser Scanning Microscope: 4 excitation lines, 2 fluorescent and 1 transmitted channel
  - Olympus IX70 Inverted wide field, fluorescence, Live stage housed in an incubator for long term live cell imaging.
  - Leica M420 Macroscope
- Morphometry

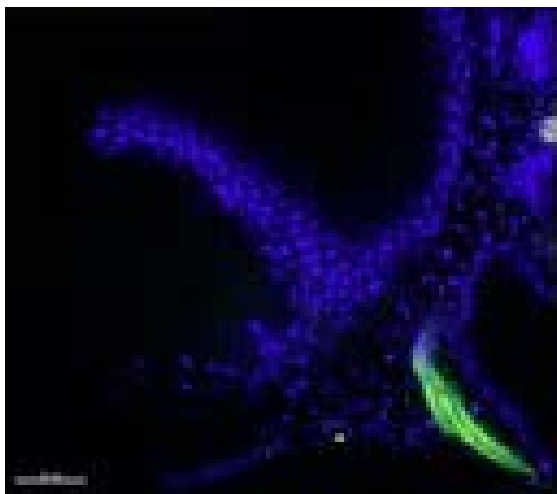
TEM



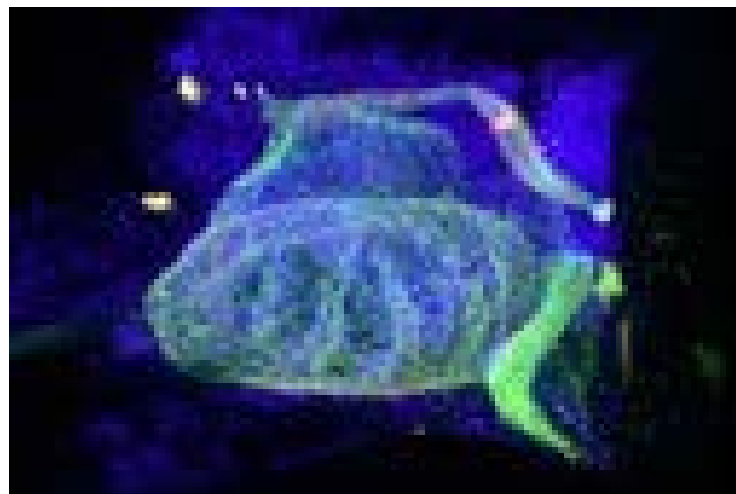
# Microscopy Services Laboratory

## UNC, Lineberger Comprehensive Cancer Center

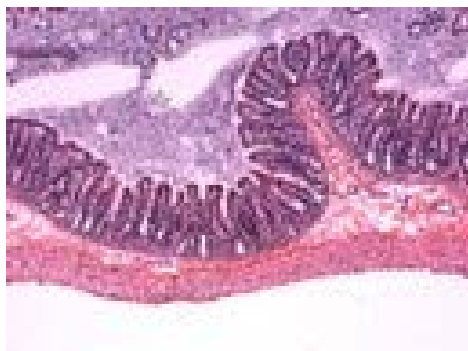
3D Recon Frog Heart



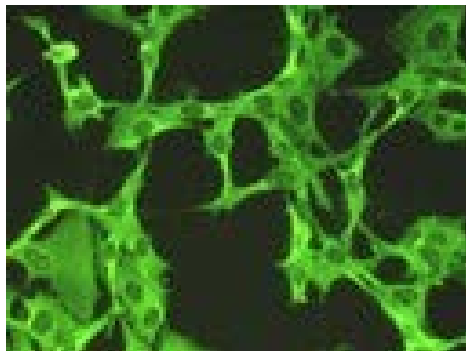
Z-Stack Frog Heart



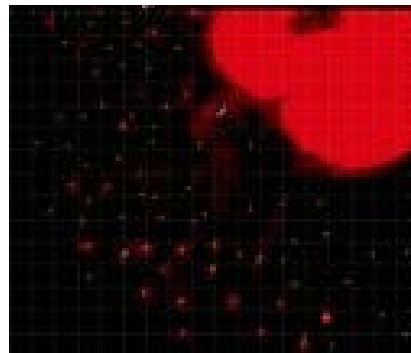
BF



IF



Live Cell

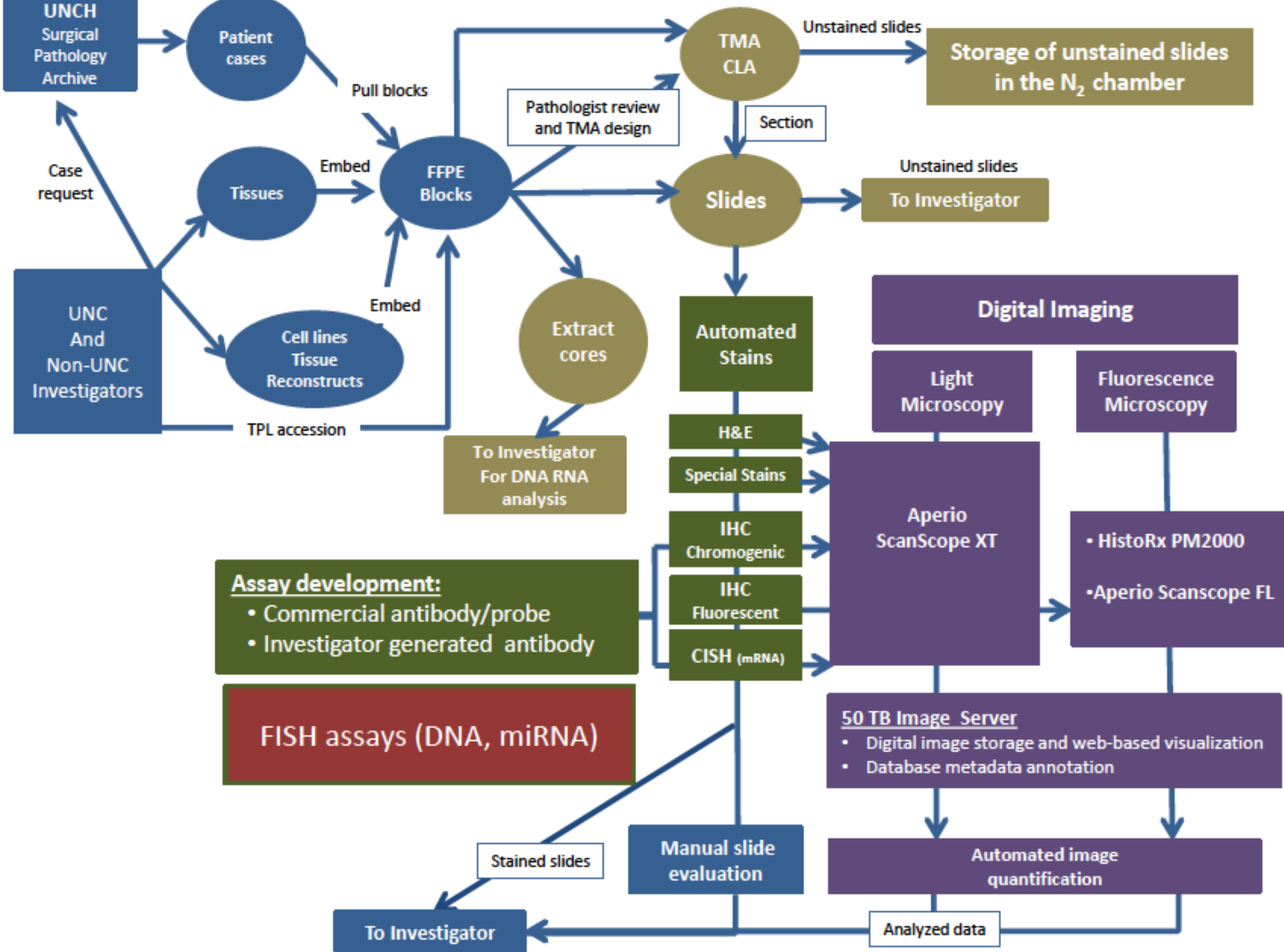


Macro



# **Histology and Digital Pathology services at UNC Translational Pathology Laboratory**

Nana N. Feinberg  
Translational Pathology Laboratory  
UNC-CH  
Facility Director

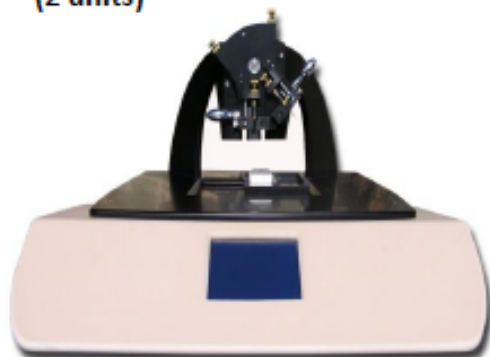


# TMA Arrayer

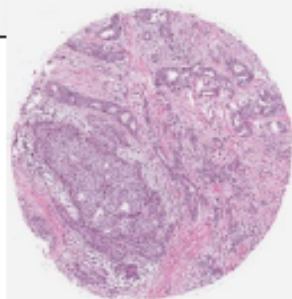
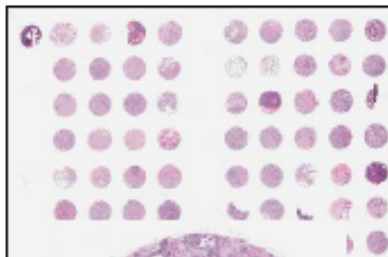
Pathology Devices

Semi-automated Tissue Arrayer

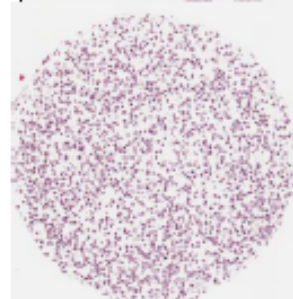
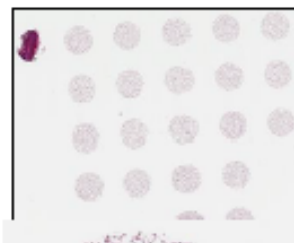
(2 units)



**Tissue (human/mouse)**



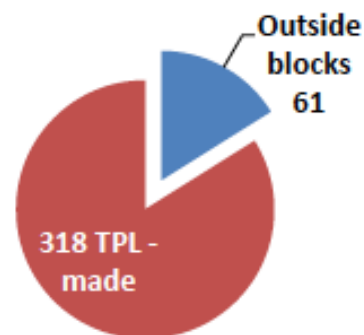
**Cell lines**



**Tissue Reconstructs**



Nitrogen Cabinets for storage of unstained slides





# Major slide staining equipment (Leica)



Bond max



Bond III (2 units)

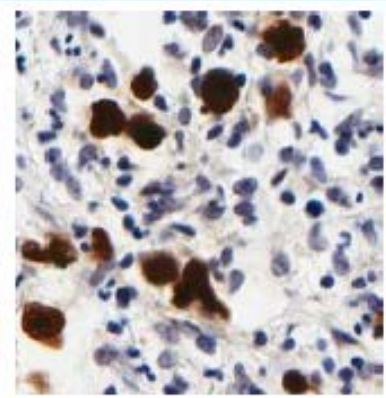
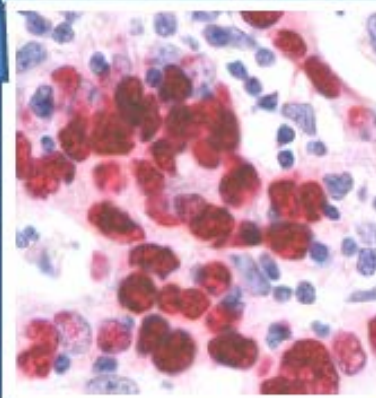
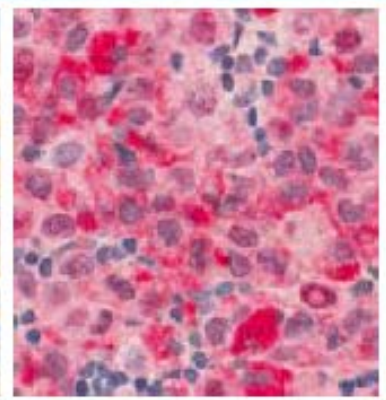
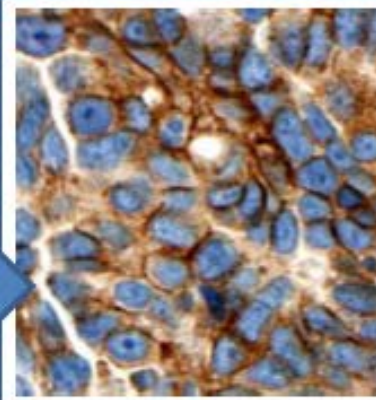


Bond Rx

IHC

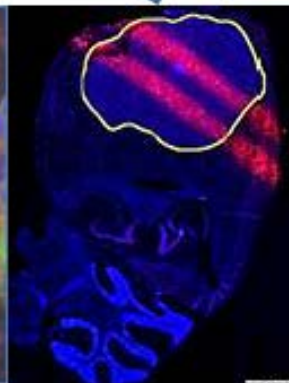
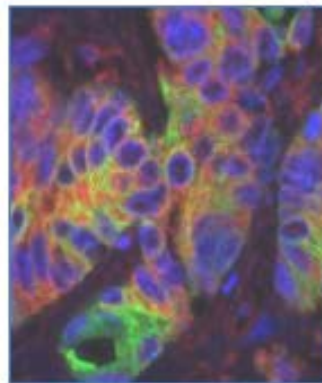
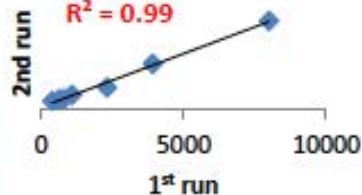
ISH

IF

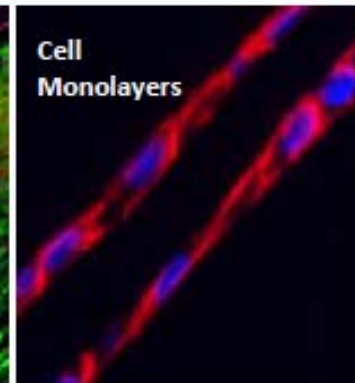


P-Chk2 assay  
reproducibility on  
irradiated NHF1 cells

$R^2 = 0.99$

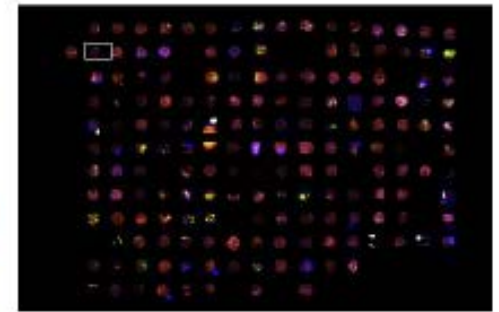
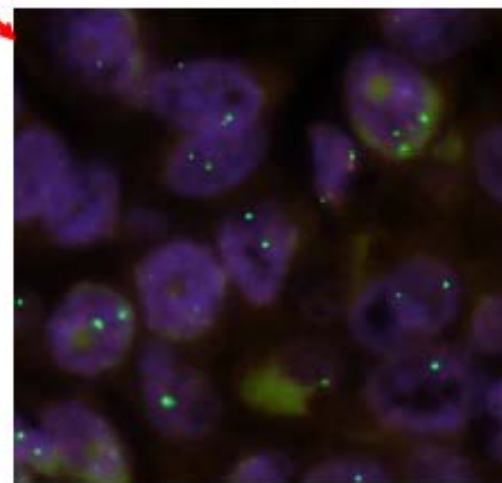
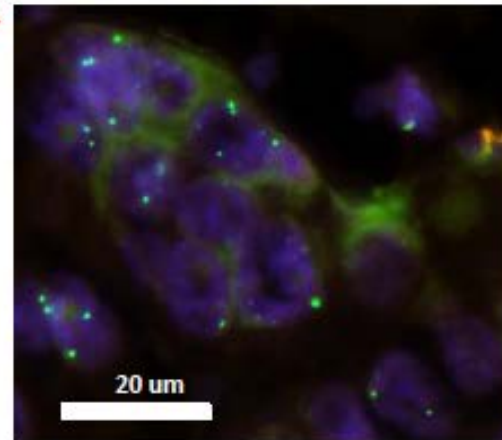
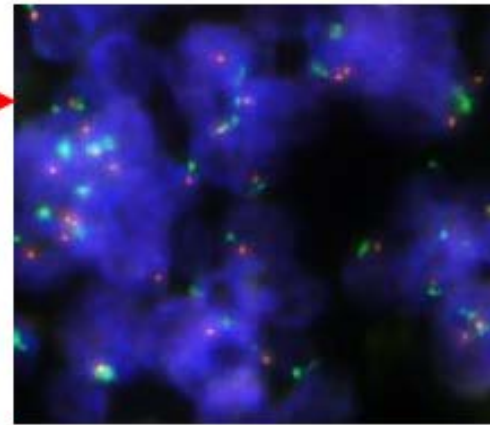
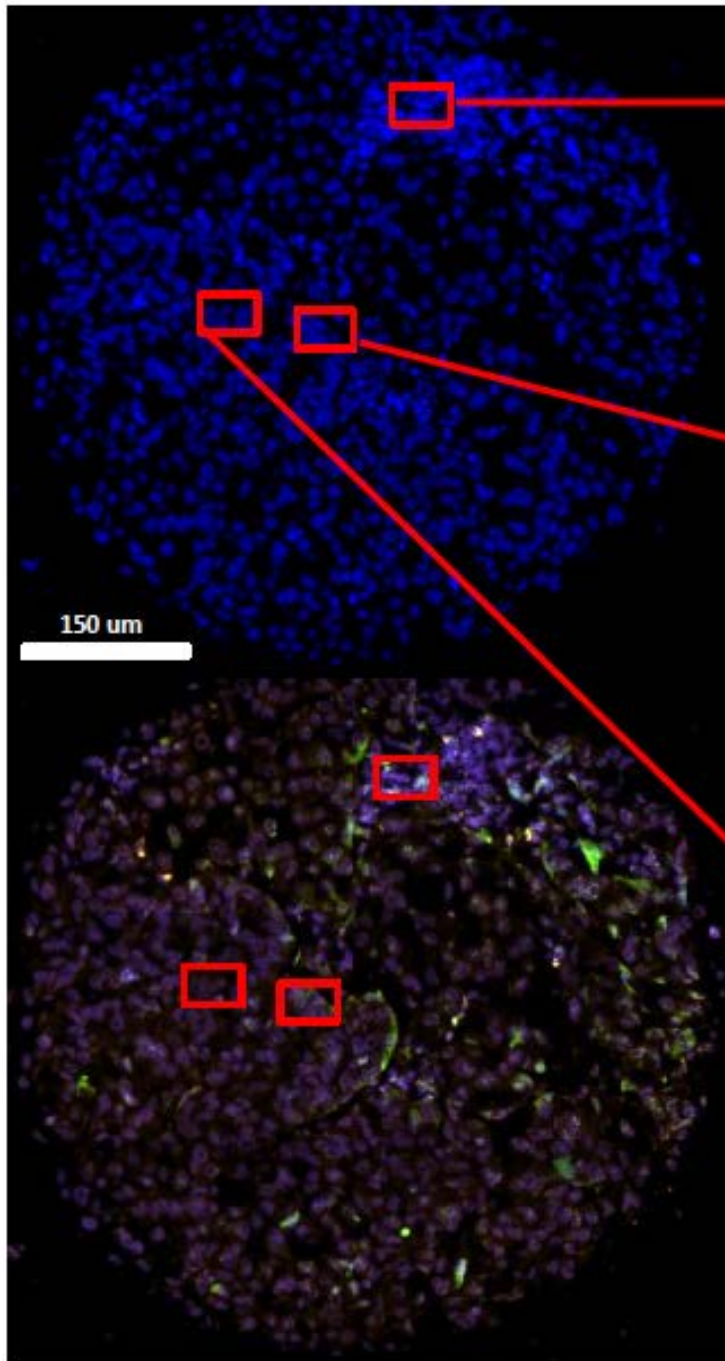


Tissue  
Reconstruct



Cell  
Monolayers





CDKN2A<sup>Red</sup>  
CEP9<sup>Green</sup>

# CDKN2A FISH

TMA 100A1  
section 21  
ARIOL Scan

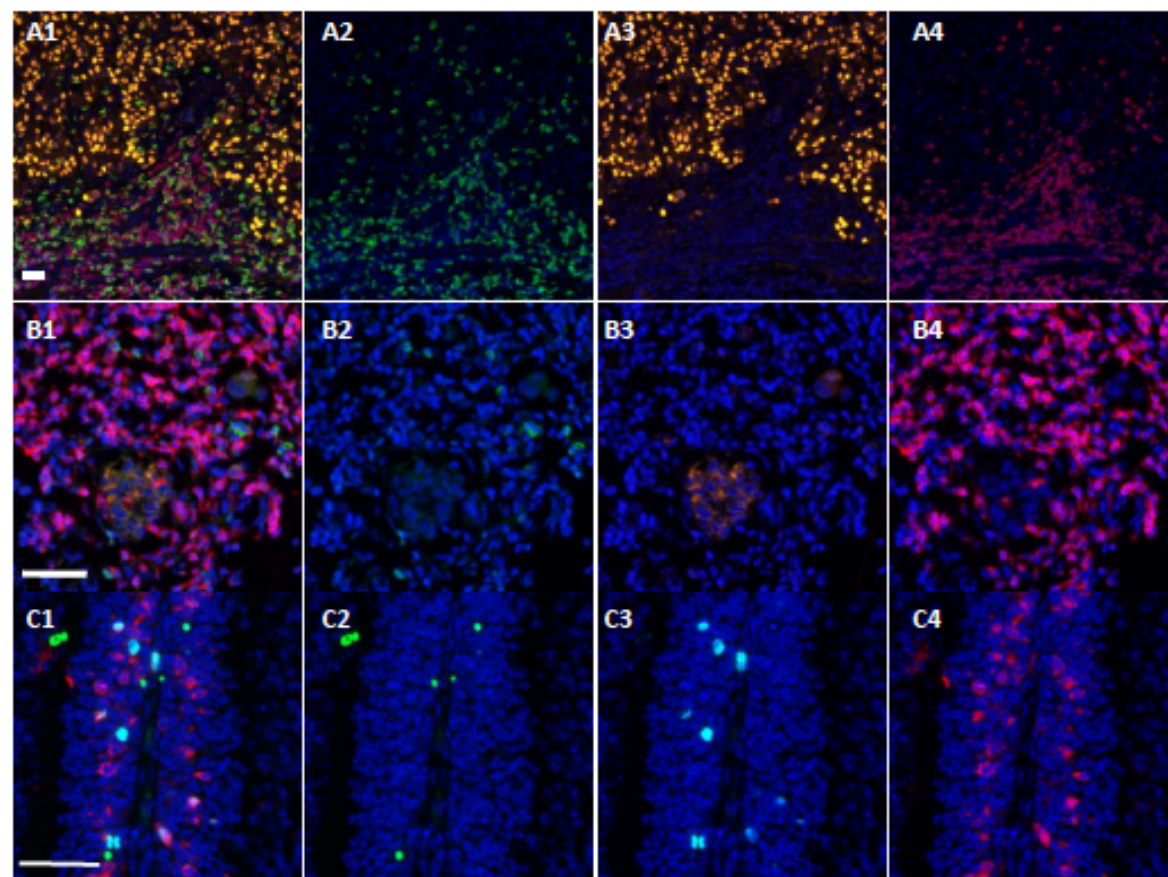
### 3-plex IF images done by TPL

**Panel A.** (A1) Human Melanoma FFPE tissue stained with Sox10 in orange (Cy3), CD3 in red (Cy5), CD8 in green (Alexa488), and Hoechst 33258 nuclear counterstain in blue. (A2-A4) the same image shown in dual channels: (A2) CD8 and nuclear, (A3) Sox10 and nuclear, (A4) CD3 and nuclear.

**Panel B.** (B1) Breast carcinoma FFPE tissue stained with Palladin in green (FITC), Pan-Cytokerstin (CK) in orange (Cy3), CD3 in red (Cy5), and Hoechst 33258 nuclear counterstain in blue. (B2-B4) the same image shown in dual channels: (B2) Palladin and nuclear, (B3) CK and nuclear, (B4) CD3 and nuclear.

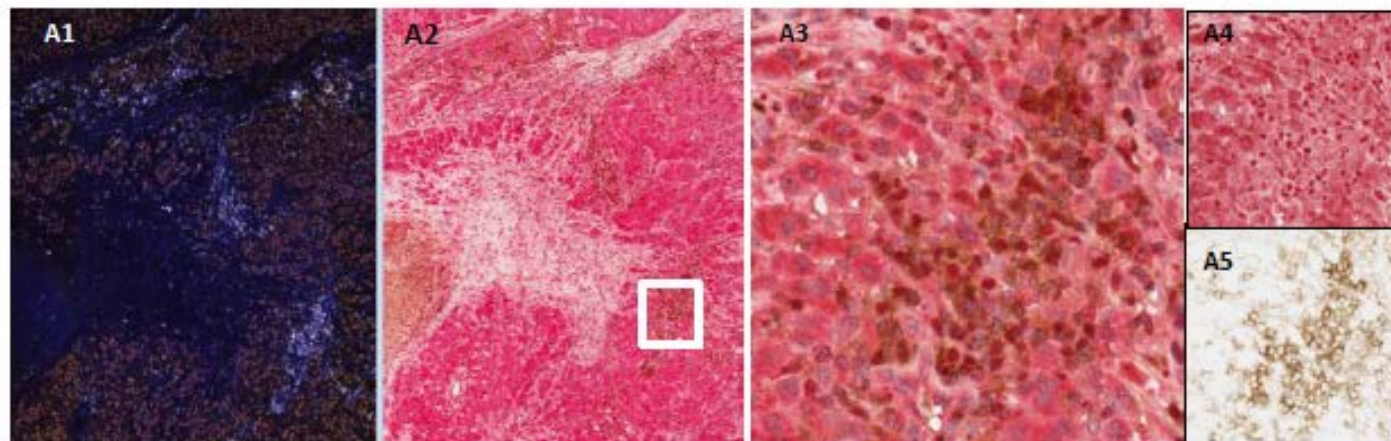
**Panel C.** (C1) FFPE Mouse brain stained with  $\gamma$ H2AX in green (Alexa488), phosphoH3 (pH3) in turquoise (Cy3), Cyclin B1 in red (Cy5), and Hoechst 33258 nuclear counterstain in blue. (C2-C4) the same image shown in dual channels: (C2)  $\gamma$ H2AX and nuclear, (C3) pH3 and nuclear, (C4) Cyclin B1 and nuclear.

The scale bars on all panels equal to 50  $\mu$ m.



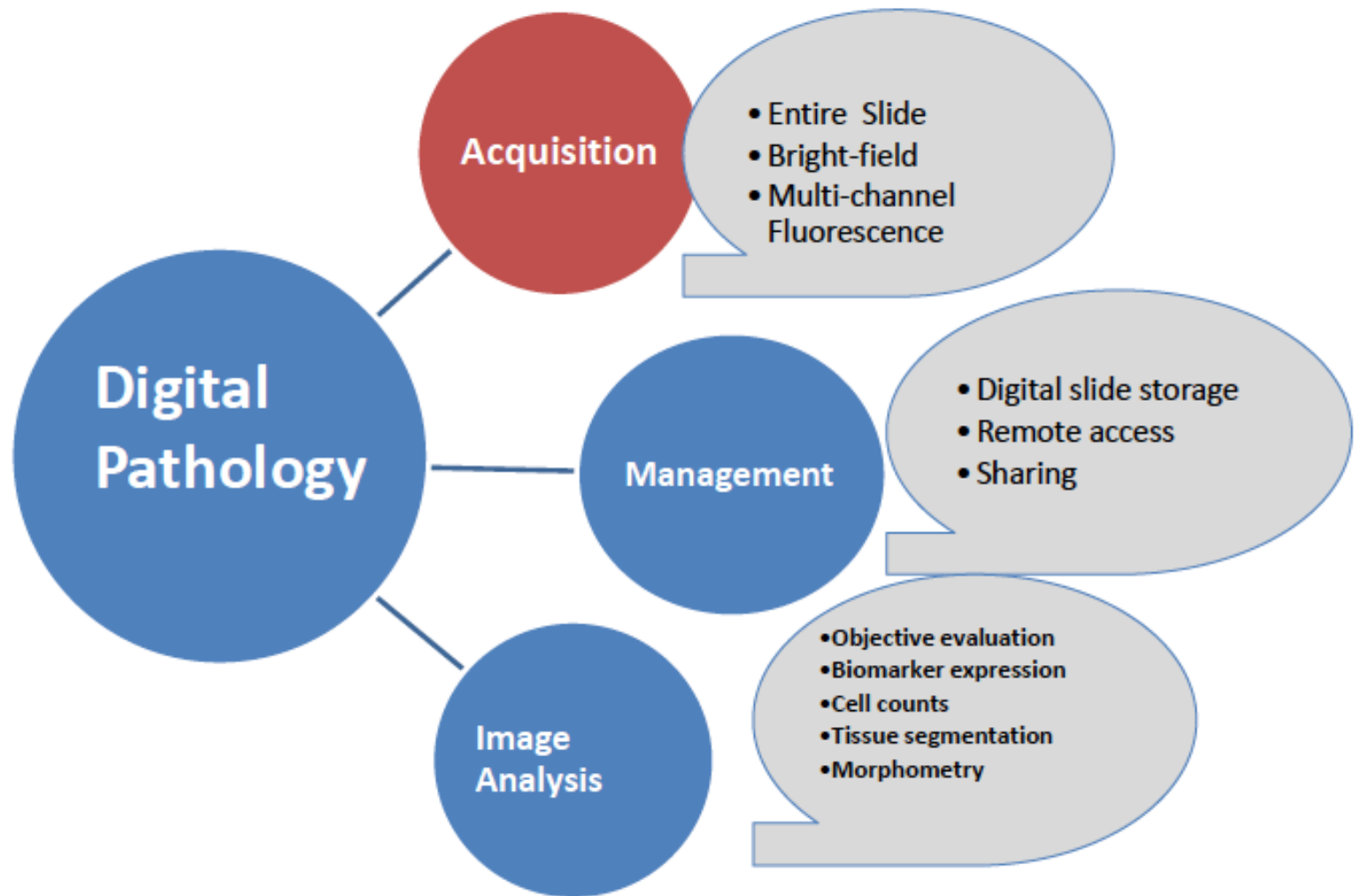


## 5-plex (IF+IHC)



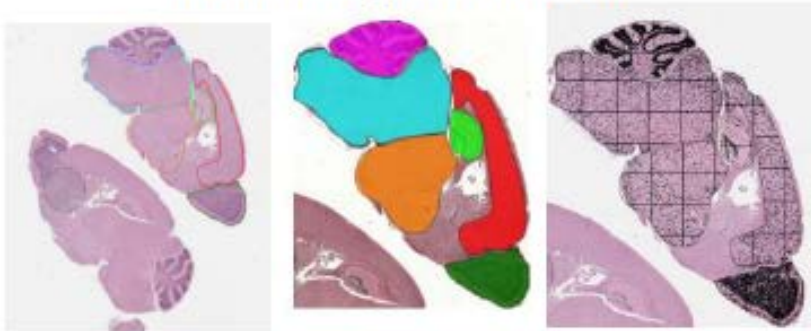
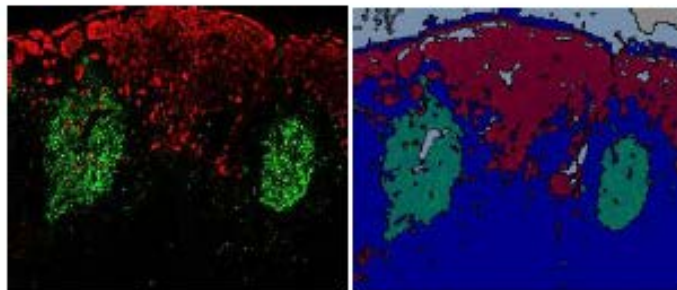
**A1.** Melanoma FFPE tissue (section #10) stained with Sox10 (cy3), CD3 (cy5) and CD8 (Alexa488). Sox10 is shown in orange, CD3 and CD8 in white. **A2-A5.** The same slide (section #10) shown in A1 was stained with CD45 IHC (brown) and PD-L1 (red), scanned and analyzed using Aperio color deconvolution algorithm. (A2) shows CD45-PD-L1/brown-red dual IHC image. (A3) is enlarged from the white box on A2. (A4, A5) show A3 image of the deconvolved red (PD-L1) and brown (CD45) channels respectively by the Aperio color deconvolution algorithm.

# Digital Pathology:

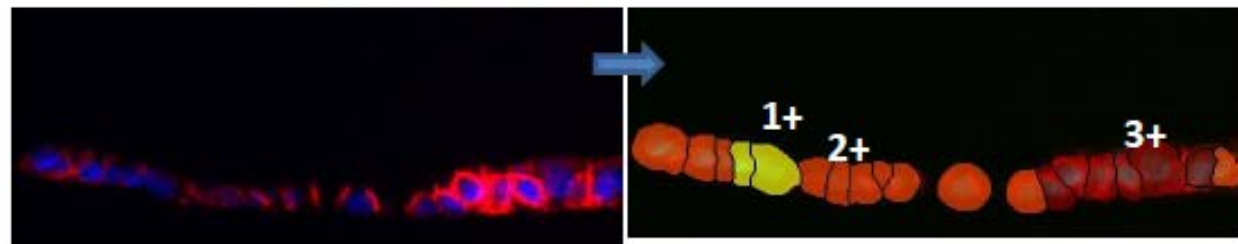
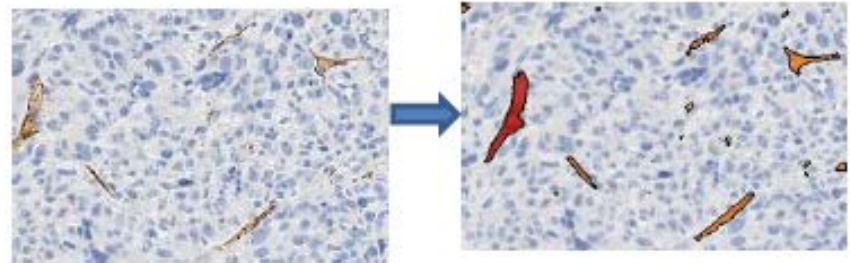
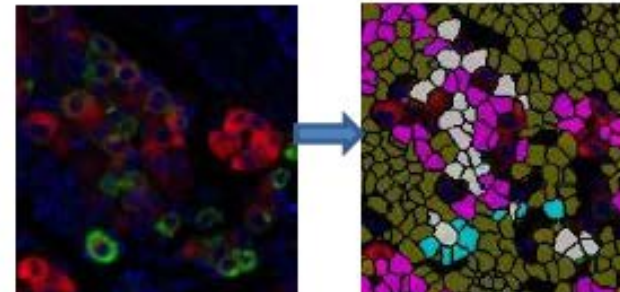


# Definiens Tissue Studio

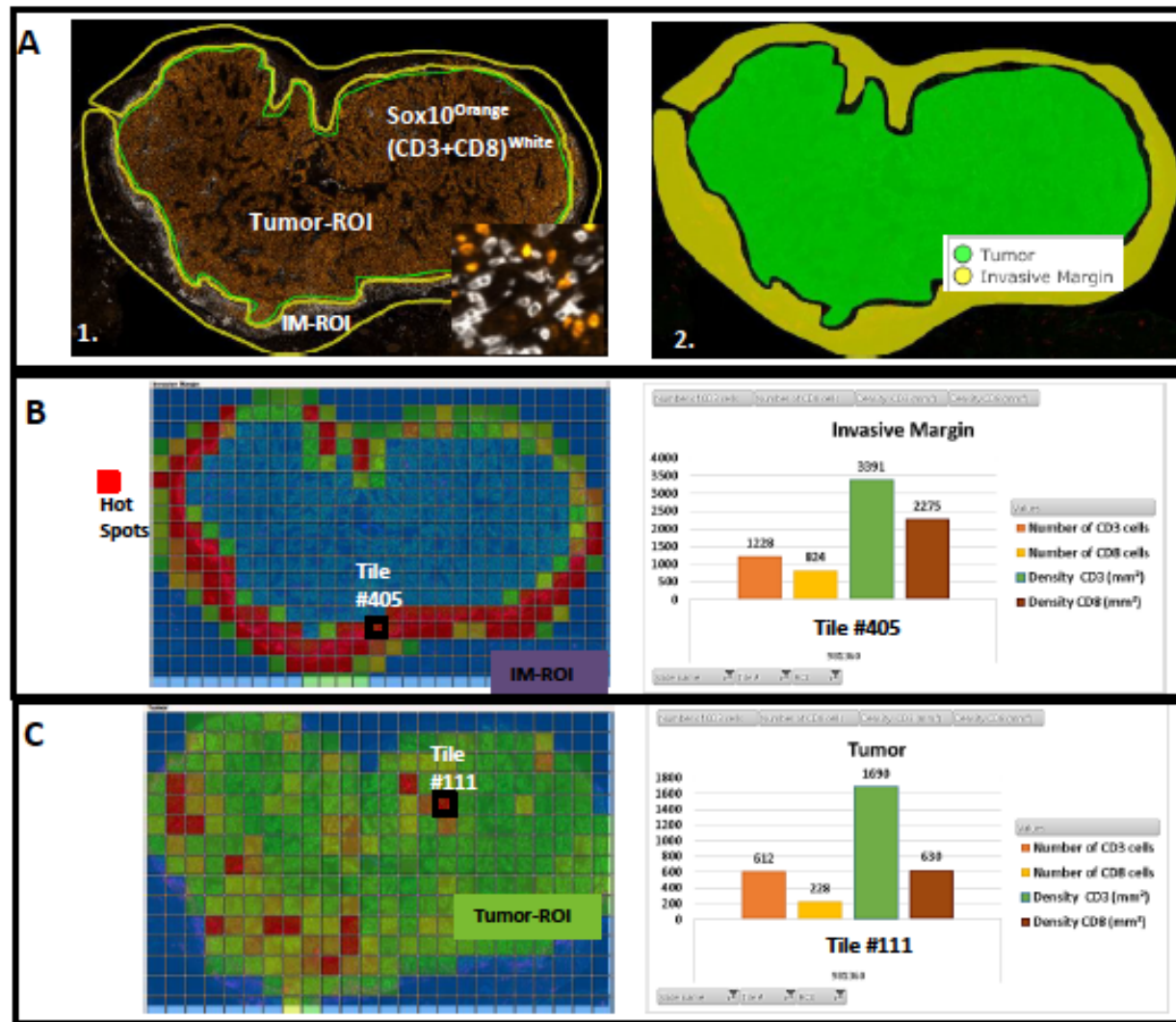
## Tissue segmentation

[illegible]

Algorithms:  
Nuclear, Co-localization,  
Microvessel, Membrane,



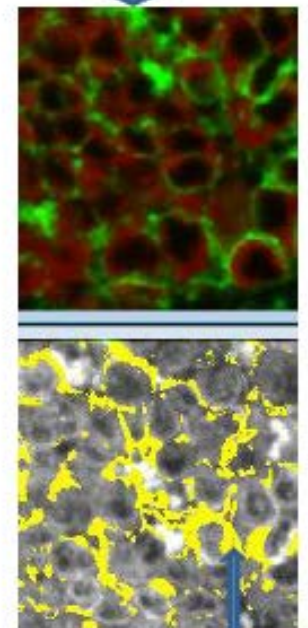
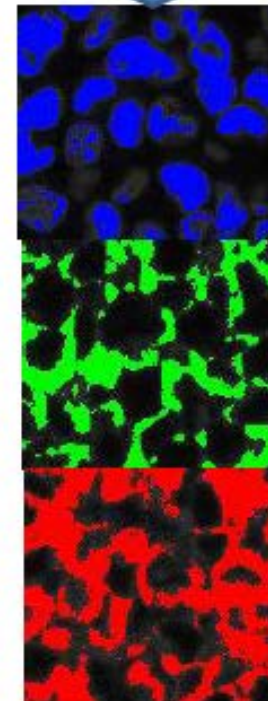
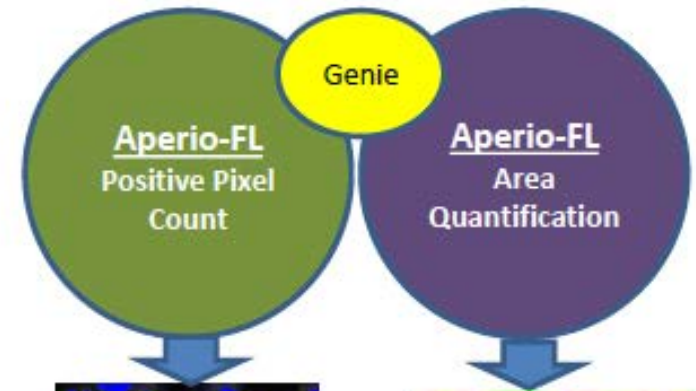
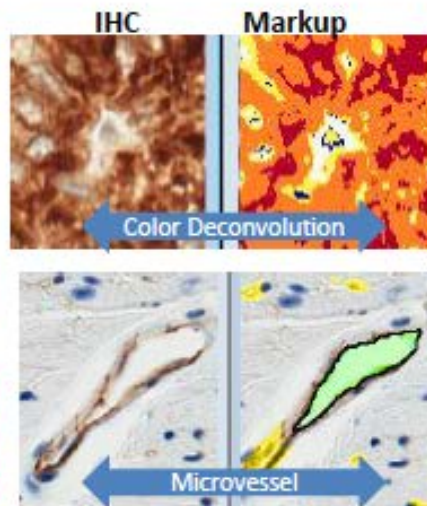
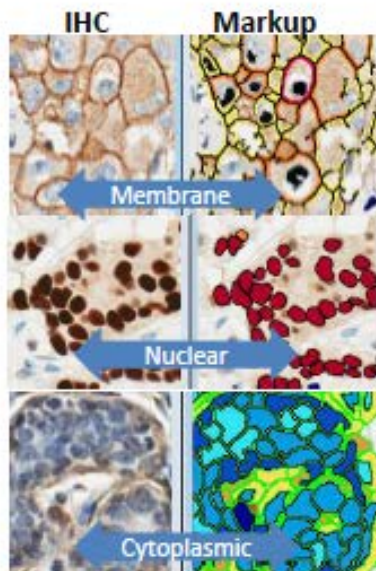
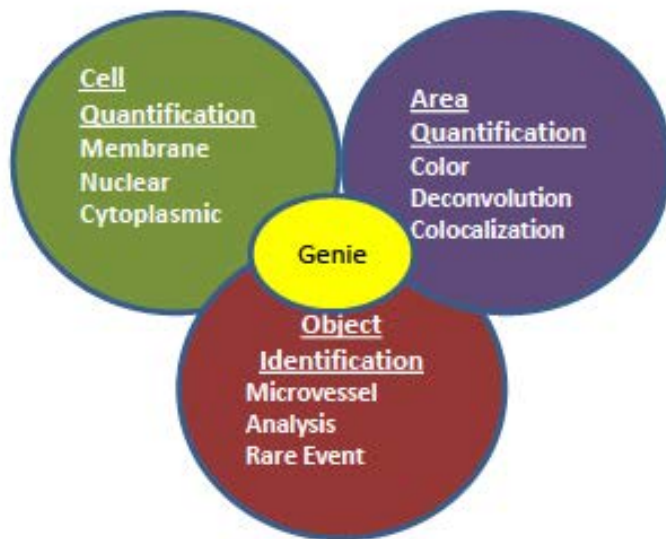




- A. (1) Manually defined Tumor and IM ROIs in the Aperio Imagscope. Sox10 is shown in orange and CD3+CD8 cells in white. Sox10 positivity was used to circle tumor ROI and an adjacent 0.5-1mm wide invasive margin (IM). (2) Tumor and IM defined by Definiens Tissue Studio
- B. using imported ROIs from Aperio. B-C. Demonstrates Tissue Studio analysis: left panels show CD3 heat map per ROI. The tiles with high density of CD3 are shown in red. The charts on the right display number and density/mm<sup>2</sup> of CD3 and CD8 cells in the black-outlined tile on the heat map.



# Aperio image analysis algorithms

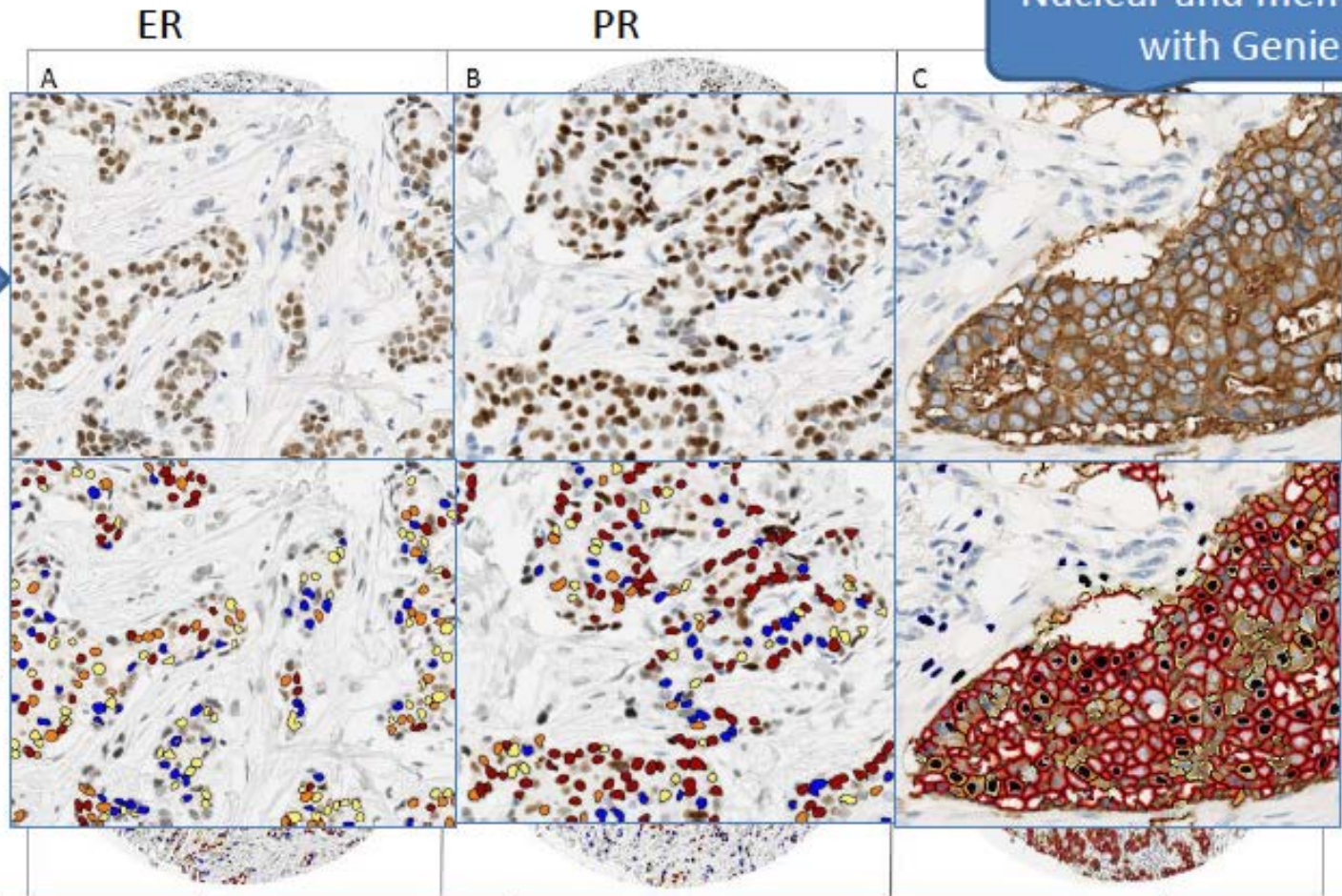


Co-localized area



# Validation of Aperio Algorithms

Nuclear and membrane  
with Genie



IHC

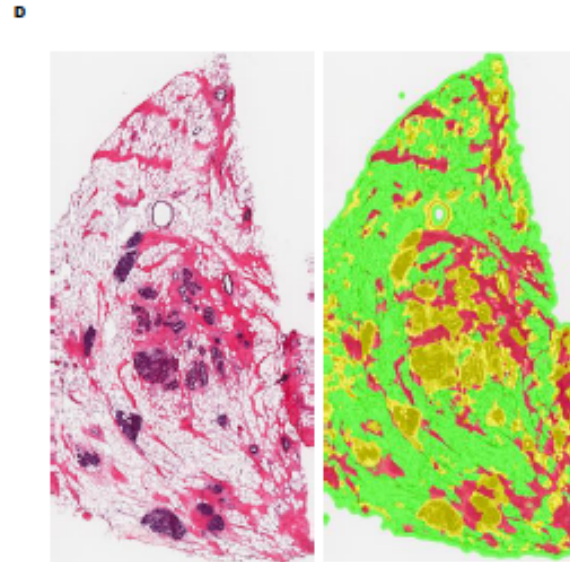
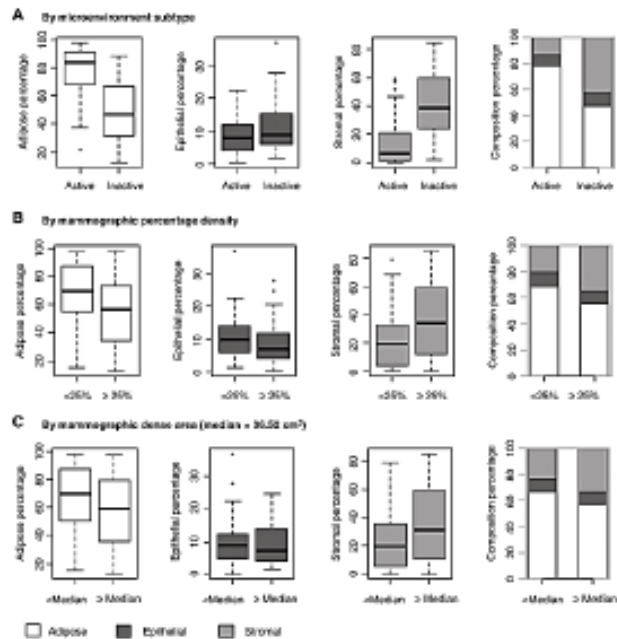


Markups

Marker	N	Observed agreement	Kappa (95% CI)	Sensitivity	Specificity
ER	242	89%	0.80 (0.73-0.86)	86%	93%
PR	280	91%	0.83 (0.78-0.89)	96%	85%
HER2	309	93%	0.79 (0.71-0.87)	98%	94%

# Breast tissue composition distribution by the Active/Inactive subtype and mammographic density.

Genie  
Algorithm



Adipose Tissue	
Epithelium	
Stroma	
Glass	

Sun X, Gierach GL, Sandhu R, Williams T, **Midkiff BR**, Lissowska J, Wesolowska E, Boyd NF, Johnson NB, Figueroa JD, Sherman ME, Troester MA. Relationship of mammographic density and gene expression: analysis of normal breast tissue surrounding breast cancer. Clin Cancer Res. 2013;19(18):4972-82.

# Animal Clinical Chemistry and Gene Expression Laboratories

UNC School of Medicine, Dept Pathology and Lab Med

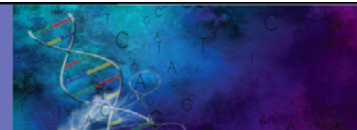
Dr. Kim PhD, Director

(919) 966-3539 [hskim@med.unc.edu](mailto:hskim@med.unc.edu)

- Hematological tests - 25ul EDTA-whole blood per sample
  - **WBC:** Lymphocytes, Monocytes, Granulocytes
  - **RBC:** HGB, HCT, MCV, MCH, MCHC, RDW
  - **PLT:** MPV
- Clinical Chemistries - 10ul plasma, serum, or urine/test +10ul
  - **Ions:** Ammonia, Blood Urea Nitrogen, Ca, Cl, CO<sub>2</sub>, Fe, Total Iron-Binding Capacity, K, Lithium, Mg, Na, and Phosphorus
  - **Protein/Organics:** Albumin, Bilirubin, Creatinine, Cholesterol, HDL-Cholesterol, Glucose, Total Protein, C-Reactive Protein, Urine Protein, Cerebral Spinal Fluid Protein, Triglycerides, Uric Acid, and Urine Creatinine
  - **Enzymes:** Alanine Aminotransferase, Acid Phosphatase, Alkaline Phosphatase, Amylase, Aspartate Aminotransferase, Cholinesterase, Creatinine Kinase MB, GGT, Lactate, LDH, and Lipase



# The University of North Carolina Organ Injury Biomarker Core



We are excited to announce the launch of The Hamner-UNC Organ Injury Biomarker Core\*, a full service core designed to assist investigators in study design, measurement, and interpretation of recently discovered biomarkers of organ injury. The traditional biomarkers utilized to detect the presence of hepatic, renal and/or cardiac injury are suboptimal, lacking sensitivity and specificity. The new biomarkers not only provide enhanced sensitivity and specificity for injury detection but can additionally inform on mechanism of injury. We currently are focusing on liver, kidney, and cardiac injury biomarkers in clinical and non clinical species; however, we are willing to expand our menu of assays to include other organs based on the needs of the investigators we serve. Please contact us.

## Advantages of working with our core:

- In-house expertise in biomarker measurement/interpretation
- Access to *in silico* modeling that incorporates release and clearance kinetics to improve study design and data interpretation (liver only at present)
- Collaborations with external experts in the fields of liver, kidney, and cardiac injury

## Multiple technologies available including:

- Standard clinical chemistry analyzer
- Meso Scale Discovery (MSD)
- Lumindex
- Firefly Bioworks

## Services offered:

- **Study design assistance** (recommendations for biofluid sampling times, biofluid collection/processing, and pertinent biomarkers to assess)
- **Biomarker measurements** (dependent on species; including but not limited to)
  - o **Hepatic:** microRNA profiling, miR-122, HMGB1 (total), K18 (full length and cleaved), GLDH, total bile acids, ALT, AST, ALP, total bilirubin, direct bilirubin, LDH, and albumin
  - o **Renal:** KIM-1, clusterin, albumin, total protein, cystatin c, RPA-1,  $\beta$ 2-microglobulin, osteopontin, NGAL, GST $\alpha$ , GST $\mu$ , sCr, and BUN
  - o **Cardiac:** microRNA profiling, cTnT, cTnI, hFABP, BNP, NT-proBNP, CK-MB, LDH, AST
- **Data Analyses/Interpretation**

## MicroRNA Profiling - Now Available

MicroRNAs are small, noncoding RNA species that post-transcriptionally regulate gene expression. These species make ideal biomarkers because they are highly stable in biofluids, such as blood and urine, and can be released into circulation following organ injury. Utilizing minimal sample volume, we can now profile targeted panels (68-plex) of microRNAs in a high throughput format.

MicroRNA panels are available for the following areas of interest:  
Immunology, Cardiology, Liver Toxicology, Neurology, and Oncology

\*Supported by the UNC TrACS Institute and CTSA grant 1UL1TR001111.



UNC  
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THE UNIVERSITY  
of NORTH CAROLINA  
at CHAPEL HILL

## Rapid funding opportunity available now



For a limited time, we are accepting one-page proposals to initiate collaborations in which biomarker measurements, analyses, and interpretations will be conducted by our core for little to no cost to investigators (up to a maximum of \$2000 in consumables supplied). We are seeking projects in which liver, renal, or cardiac injury are suspected in clinical or nonclinical studies but have not yet been explored with the offered biomarkers. Please contact us for more details or to submit a proposal. Funding decision will be made within two weeks of submission.

For more information, to submit a study proposal, or to request a consultation,

contact: Rachel Church, Ph.D., Director

[rchurch@unc.edu](mailto:rchurch@unc.edu) • 919-226-3145

# Organ Injury Biomarker Core

Hamner-UNC / Eshelman School of Pharmacy

Dr. Rachel Church, PhD, Director

(919) 226-3145, [rchurch@unc.edu](mailto:rchurch@unc.edu)

- Study design, data analysis/interpretation, different species
- Biomarker measurements

**Hepatic:** MicroRNA, miR-122, HMGB1, K18, GLDH, Bile acids, ALT, AST, ALP, Bilirubin, LDH, albumin

**Renal:** KIM-1, clusterin, albumin, total protein, cystatin c, RPA-1, beta2-microglobulin, osteopontin, NGAL, GSTalpha, sCr, BUN

**Cardiac:** MicroRNA, cTnT, cTnl, hFABP, BNP, NT-proBNP, CK-MP, LDH, AST

**MicroRNA profiling** - 68 plex technology, Firefly, biofluids

- Technology - *small sample size, multiplex*

Firefly, Meso Scale Discovery (MSD), Luminex, chemistry analyzer