

Comprehensive

165

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Cancer Center

Aims

- The CAP-accredited BRHS SR's primary aims are to:
- Provide tissue analysis, including histological, immunohistochemical, immunofluorescence staining, routine and special histologic technique, and tissue microarray
- Consent, bank, and distribute biospecimens and provide consultation to CINJ investigators to optimize the collection of primary and correlative data from clinical trials

CIPT

Research Program Support (2018–2022)



GICG, CMI, CIPT

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GICG



Cancer Res. 2021







Leading Personnel & Roles



Gregory Riedlinger, MD, PhD Interim Director



Kelly Walton Histopathology Manager



Zhongren (David) Zhou, MD *Co-Director*



Joseph Rosenberg Biorepository Manager



Kathleen Dwyer Program Administrator

Services & Innovation





Services & Innovation

New

- Consenting patients to CINJ banking protocol and other research studies – Cooperman Barnabas and other RWJBH hospitals
- Online Biospecimen query dashboard
- Processing, embedding, cutting or snap-freezing tissue – New Sakura tissue processor, embedding center, and automated coverslipper

A Main

Personnel

- New CK4600 automated tissue microarrayer
- Multiplex IHC

Primary Site	SPECIMEN SITE			
Select all	Select all			
(Blank)	ABDOMINAL			
ABDOMINAL WALL	ADRENAL			
ADRENAL	ANUS			
AMPULLA	APPENDIX			
AMPULLA OF VATER	ASSORTED TISSUE T			
ANUS	BILE DUCT			
APPENDIX	BLADDER			
BLADDER	BRAIN			





Emphasis & Directions

Continuing

Utilization & Management

- Targeted collection & distribution of biospecimens for IRB approved research studies
- Immunohistochemistry & Immunofluorescence
- Routine and Special histology staining
- Preparation of tissue slides for Laser Capture Microdissection
- Sample management system uses OnCore Biospecimen Module



Parkin Ubiquitinates Phosphoglycerate Dehydrogenase (PHGDH) to Suppress Serine Synthesis and Tumor Progression



The BRHS SR used a special TMA composed of 200 primary breast tumors and performed IHC to show that Parkin inhibits tumorigenesis through negative regulation of PHGDH. Additional TMAs for lung and breast tumors were obtained by US Biomax. Representative images of IHC staining of Parkin and PHGDH in human breast cancer specimens and lung cancer specimens

IMPACT

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RUTGERS HEALTH

PHGDH is frequently overexpressed in human cancer, including breast and lung cancers. This overexpression activates serine synthesis to promote cancer progression and is associated with poor prognosis in cancer patients

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The authors present evidence that through ubiquitination and degradation of PHGDH, Parkin suppresses serine synthesis, which contributes greatly to the tumor-suppressive function of Parkin



Personnel

N=approximately 24-30



Phase 1 Trial of Pembrolizumab Administered Concurrently with Chemoradiotherapy for Locally Advanced Non-Small Cell Lung Cancer

Subject must have inoperable non-small cell lung cancer to be treated with definitive chemoradiation

Subject must have measurable disease based on RECIST 1.1 subjects with inoperable non-small cell lung cancer

IMPACT

(MK-3475) beginning at 200 mg Q3W starting 2-6 weeks after chemoradiation with subsequent dose escalation and advancing the timing to during chemoradiation

Pembrolizumab

Monitor for safety

Aisner, Jabbour ⊠ (GICG)

Lin (BSR)

JAMA Oncol, 2020 6(6):848-855 The BRHS SR provided PD-L1 IHC 22C3 pharmDx service for this clinical trial. The authors were able to demonstrate feasibility and tolerance of the Incorporation of pembrolizumab with concurrent chemoradiotherapy in the definitive management of locally advanced NSCLC with a the 12-month progression free survival was 69.7%







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mTOR Regulates Aerobic Glycolysis Through NEAT1 and Nuclear Paraspeckle-mediated Mechanism in Hepatocellular Carcinoma



The BRHS SR performed H&E staining and optimized anti-HA IHC staining for the investigator. Mouse liver tumor tissues were stained by IHC positively for the HCC marker Arginase, but negatively for the cholangiocarcinoma marker CK19. Neat1_2 was stained using RNAscope

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IMPACT

Previous human clinical trials of rapalogs in advanced HCC failed to achieve desired endpoints and better patient survival outcomes

Our observations reveal that NEAT1 expression/paraspeckle biogenesis is a key determinant for the success of mTORC1targeted cancer therapy in liver cancer





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FILIP1L Loss is a Driver of Aggressive Mucinous Colorectal Adenocarcinoma and Mediates Cytokinesis Defects through PFDN1

D Е F H&E FILIP1L PAS E Zhou (BRHS SR co-author) Nov 1; 81(21): 5523-5539

IMPACT

The authors have shown that Filamin A interacting protein 1-like (FILIP1L) is a versatile tumor suppressor in many types of cancer

Here they show that FILIP1L increases xenograft growth in vivo, drives colonic epithelial hyperplasia in mice, and increases mucin secretion and mitotic defects in mucinous colorectal adenocarcinoma. an aggressive subtype of colorectal cancer with poor prognosis

The BRHS provided FFPE of normal colon, serrated polyps, mucinous adenocarcinoma, and nonmucinous adenocarcinoma, IHC and independent pathologist staining under blinded conditions for FILIP1L. Colons were fixed and stained with H&E (D), FILIP1L (E), and PAS (F)



Verzi, Libutti ☑ (GICG)

Cancer Res, 2021

Services & Innovation

Emphasis & Directions

Utilization & Management

Services & Innovation





Emphasis & Future Directions

A Main

CINJ Priority Cancer Collections by Ethnicity

(Jan 2018 - Dec 2022)

Obtain CINJ vouchers to boost collections and utilization of samples in BIPOC populations

Race	Breast	Cervix	Colon	Lung	Melanoma	Prostate	Rectum	Totals
Asian	5	0	35	4	2	31	0	77
Black or African American	9	0	48	4	0	39	10	110
Unknown	0	0	3	1	0	4	0	8
White	62	6	202	151	41	195	0	657
Totals	76	6	288	160	43	269	10	852

IIT Support Emphasis



Cancer Immunotherapy PI: Hinrichs. CT83 IHC assay is used to identify suitable candidates for targeted cellular therapy protocol by testing their tumor tissue for expression of KK-LC-1 antigen

Future Directions

 Recruit a Chief of Oncologic Pathology who will also serve as the new BRHS director

Personnel

 Add a pediatric tumor biobank with its own consent protocol in the next few months

Utilization & Management

Emphasis & Directions

 We also expect to integrate additional RWJBH hospitals

RUTGERS Cancer Institute of New Jersey RUTGERS HEALTH

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Supporting Information

