BHD TUMOR CELL LINE AND RENAL CELL CARCINOMA LINE

SUMMARY (1024-CHARACTER LIMIT)
Scientists at the National Cancer Institute have developed a novel renal cell carcinoma (RCC) cell line designated UOK257, which was derived from the surgical kidney tissue of a patient with hereditary Birt-Hogg-Dube (BHD) syndrome and companion cell line UOK257-2 in which FLCN expression has been restored by lentivirus infection. The NCI Urologic Oncology Branch seeks parties interested in licensing or collaborative research to co-develop, evaluate, or commercialize kidney cancer tumor cell lines.

NIH REFERENCE NUMBER
E-131-2010

PRODUCT TYPE
- Research Materials

KEYWORDS
- Tumor Cell Line
- Renal Cell Line
- renal cell carcinoma
- RCC

COLLABORATION OPPORTUNITY
This invention is available for licensing and co-development.

CONTACT
- John D. Hewes
  NCI - National Cancer Institute
  240-276-5515
  John.Hewes@nih.gov

STATUS
Active

DESCRIPTION OF TECHNOLOGY
The NCI Urologic Oncology Branch is seeking statements of capability or interest from parties interested in collaborative research to further develop, evaluate, or commercialize kidney cancer tumor cell lines.

Scientists at the National Cancer Institute have developed a novel renal cell carcinoma (RCC) cell line designated UOK257, which was derived from the surgical kidney tissue of a patient with hereditary Birt-
Hogg-Dube’s (BHD) syndrome and companion cell line UOK257-2 in which FLCN expression has been restored by lentivirus infection. These cell lines harbor a germline mutation of FLCN gene (alias BHD) and displays loss of heterozygosity, can grow as xenograft in nude mice. Patients affected with BHD develop skin papules (fibrofolliculomas), lung cysts, spontaneous pneumothorax and an increased risk for bilateral multifocal renal tumors. Loss of both copies of the FLCN gene has been documented in BHD renal tumors; however, the molecular mechanisms by which inactivation of the encoded protein, folliculin, leads to the BHD phenotype are currently unknown. They have developed an important research tool for in vitro folliculin functional studies. The companion cell line will be extremely useful for comparative biochemical analyses of cell culture systems in which the FLCN gene is either expressed or inactivated, including identification of renal tumor biomarkers, alteration of biochemical pathways resulting from loss of FLCN function, tumorigenicity of FLCN null versus FLCN restored cells, preclinical therapeutic drug testing in xenograft animal models produced from injection of these cell lines, etc. UOK 257 and UOK257-2 are thus useful cell models for studying the underlying molecular derangements associated with mTOR pathways and other biogenesis pathways in human kidney cancer and for evaluating novel therapeutic approaches for this disease. UOK257 is also one of the 40-member renal cancer cell lines in the Tumor Cell Line Repository of the Urologic Oncology Branch (UOB), National Cancer Institute (NCI).

Topics of possible collaboration:

- For laboratory interests on the basis of metazoan tumor cell survival, including growth factor-regulated nutrient uptake; glucose or glutamine metabolism and epigenetic gene control; tumor cell bioenergetics and cell growth through AMPK and mTOR signaling pathways.
- In vitro and in vivo cell model for BHD cancer syndrome. It is voluble research tool for laboratory interested in identification of new BHD tumor antigens for immunotherapy.
- These paired cell lines for FLCN gene expression and function studies, including gene therapy, cytogenetics, gene mutation research, and examination abnormalities of interaction with other proteins that may contribute to BHD.
- The excellent in Vivo model for preclinical xenograin imaging, including stable transfection. Cells could be labeled with reagents for PET, Luciferase, Fluorescent, for transgenic mice, optical molecular imaging, etc. and provides useful platform for preclinical drug evaluations.

POTENTIAL COMMERCIAL APPLICATIONS

- In vitro and in vivo cell model for BHD cancer syndrome. Research tool for investigating the underlying molecular mechanisms contributing to advanced BHD, including the identification of new BHD tumor antigens for immunotherapy
- Research tool for studying genes transcription status of genes involved in BHD to reveal the genetic processes occurring in BHD tissues that may contribute to advanced disease
- Positive control cell line for FLCN gene expression and function studies, including cytogenetics, gene mutation research, and examination abnormalities of interaction with other proteins that may contribute to BHD
- Research tools for testing the activity of potential anti-cancer drugs against BHD, a disease which has
no effective treatment options; tool for searching tumor markers for diagnosis, prognosis and drug resistance

• Therapeutic drug testing for targeting BHD renal tumors, possible starting material for developing a cancer vaccine against BHD

COMPETITIVE ADVANTAGES

• Cell line is derived from a BHD patient
• Molecular and genetic features are well characterized

INVENTOR(S)

W. Marston Linehan (NCI)

DEVELOPMENT STAGE

• Basic (Target Identification)

PUBLICATIONS


Baba M, et al. Folliculin encoded by the BHD gene interacts with a binding protein, FNIP1, and AMPK, and is involved in AMPK and mTOR signaling. [PMID 17028174]

PATENT STATUS

• Research Material: NIH will not pursue patent prosecution for this technology