Researchers from the Greehey Children's Cancer Research Institute at UT Health San Antonio co-authored a paper published Sept. 26 in *Science Signaling* that looks at the reliability of a common research tool to study RAS cancer mutations.

RAS mutations are implicated in several cancers, including lung, colorectal and pancreatic ductal carcinoma. The three RAS genes are the most frequently mutated oncogenes in cancer, implicated in more than a quarter of all human cancers.

In pediatric cancers, mutations that activate the RAS pathway are frequently found in embryonal rhabdomyosarcoma (35 percent of tumors) and recurrent neuroblastoma (60 percent).

The *Science Signaling* article says that although there is intense interest in studying the development of anti-cancer strategies targeting RAS proteins, the field is held back by poor reliability of anti-RAS antibodies to recognize the members of the RAS family of proteins.

These antibodies are a critical research tool, but lack of reliability limits useful—and replicable—study results, the authors noted.

Angelina V. Vaseva, Ph.D., and Peter J. Houghton, Ph.D., director of the Greehey Institute, are the co-authors from UT Health San Antonio. Dr. Vaseva is a visiting scientist from the laboratory of Dr. Channing Der, the senior author on the study performed at the University of North Carolina at Chapel Hill, and done in collaboration with the National Cancer Institute RAS Initiative. Dr. Vaseva conducts research in the Houghton laboratory related to therapeutics of RAS-mutant childhood cancers.

The team from multiple institutions evaluated 22 commercially available anti-RAS antibodies for their capability to recognize RAS subtypes and mutated RAS.

The collaborators identified antibodies that selectively recognize each of the four human RAS proteins in human cancer cells.

This capability is only in Western blot assays, one of the most commonly used research analyses, and not in other research analyses used by laboratories worldwide.


Provided by University of Texas Health Science Center at San Antonio

This document is subject to copyright. Apart from any fair dealing for the purpose of private study or research, no part may be reproduced without the written permission. The content is provided for information purposes only.