



NJ Center for Tourette Syndrome
AND ASSOCIATED DISORDERS, INC.



Be an important part of Tourette Syndrome genetics research by donating DNA in March/April at Rutgers University



Would you like to see more people working on a cure for Tourette Syndrome? How about research on medications specifically for TS? Now is your opportunity to join the NJ Center for Tourette Syndrome & Associated Disorders (NJCTS) efforts and personally be part of making that happen.

Dr. Robert King, a professor of child psychiatry at the Yale University Medical School and medical director of the TS/OCD Clinic at Yale University Child Study Center, will be at the NJCTS Cell and DNA Sharing Repository at Rutgers University all day on **Thursday and**

Friday, March 31 & April 1 2016 to interview individuals and families interested in participating in the National Institute of Mental Health NJCTS Genetics Sharing Program.

For more information or to schedule an appointment with Dr. King, please email Dr. Gary Heiman, a team leader at the repository, at familystudy@biology.rutgers.edu. For more information on the program, please visit www.rucdr.org.



The NJ Center for Tourette Syndrome & Associated Disorders, Inc. (NJCTS) is the nation's first Center of Excellence for Tourette Syndrome (TS). Through partnerships and collaborations, NJCTS provides a continuum of services, support and education for families; outreach and training for medical and educational professionals; and advocacy for collaborative research for better treatments and a cure for TS.

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