



# NEW JERSEY CENTER FOR TOURETTE SYNDROME AND ASSOCIATED DISORDERS, INC.

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## Be an important part of Tourette Syndrome Genetics research by donating DNA in 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 New Jersey 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 the Yale University Child Study Center, will be at the Cell and DNA Repository at Rutgers University all day on **Thursday and Friday, April 9th and 10th**, 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 e-mail Dr. Gary Heiman, a team leader at the repository, at [familystudy@biology.rutgers.edu](mailto:familystudy@biology.rutgers.edu). For more information on the program, please visit [www.rucdr.org](http://www.rucdr.org).

