Joseph G. Gleeson, MD and his associates are conducting a research study to find out more about the inherited causes of brain diseases in childhood (genetic abnormalities of brain development such as mental retardation). Your child has been asked to participate because your child has a neurological disease thought to be due to a genetic defect. The goal of this study is to identify the specific genetic defect underlying the condition in your family. There will be approximately 800 new participants joining this study every year. To date there have been over 2700 individuals participating in this project.

If you agree to be in this study, the following will happen to you:

1. A blood sample (approximately 3 tablespoons) will be drawn from a vein in your child’s arm, or if your child is having surgery, from the IV line that is placed for anesthesia, or a saliva sample will be collected from your child and sent to Dr. Gleeson’s laboratory.
2. If your child is having surgery, the surgeon will save a small piece of tissue that would have been thrown away otherwise.
3. In order to localize the genetic cause of brain development in your child’s family, we also need to obtain DNA sample from each member of your family that might also carry the diseased gene. This determination will be made by Dr. Gleeson or one of his associates in advance, but may include your child’s parents, children, siblings, cousins and nieces/nephews. DNA will not be obtained from members of your family not at risk for carrying the diseased gene.
4. You will sign a release to allow your child’s medical records to be forwarded to Dr. Gleeson for review.
5. If you choose, you will be notified of results obtained through this study. We will not disclose non-maternity or non-paternity information.
6. You will not receive compensation for participating in this study, though you may be reimbursed for your travel expenses. Additionally, the necessary phlebotomy procedures or doctor visits will be of no cost to you.

Participation in this study may involve some added risks and discomforts. These include:

1. The blood draw may hurt slightly. Risks of drawing blood include possible pain, discomfort, and bruising at the puncture site, possible dizziness and fainting and possible infection. Prolonged bleeding is treated with pressure to the needle site, and bruising may leave the needle site temporarily discolored. If infection occurs, it will require medical attention.
2. There is a chance that participation in this study could cause psychological distress, economic and social harm. Some people involved in genetic studies have felt anxious about the possibility of carrying an altered gene that places them at risk or that can be passed on to children. If these feelings arise at any time during the study, you may contact us and we will arrange for you to speak with a genetic counselor.
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3. Social and economic harms which can be associated with the gathering of genetic information. We will do our best to keep all information confidential, but the testing may find an inherited defective gene that puts your child at risk for developing a genetic disease at some time in the future. This information, if disclosed to the wrong source, may adversely affect you economically and socially and adversely affect your child’s insurability and employability and cause you to be stigmatized. We will not make this information available to anyone unless you ask us to do so. All of the information we collect during the course of this study will be kept locked and secured and it will be released to no one unless you ask us to do so. In the course of collection of the clinical information we may need to review your medical records such as MRI scans, pathology reports, laboratory or progress reports etc. This information as well, will be kept confidential and not shared with anyone outside this project. We will not release this information to insurance companies, family members, work places or any other institutions. Even though, the risk of losing confidentiality via medical records cannot be fully eradicated, we take all the precautions to protect this information.

4. In order to help advance future patient screening strategies, a new condition in our research study might require that your child’s DNA sequence, diagnosis, and de-identified pedigree data be deposited into the NIH’s (National Institute of Health) dbGaP database. Your child’s personally identifiable information (PII) will not be shared on this database. Your child’s PII will remain confidential. Only de-identified genetic data would be deposited into the dbGaP database.

5. Issues of non-paternity and non-maternity. Non-paternity or non-maternity refers to the situations when as a result of genetic testing we are able to indicate that reported relationship between one or both parents with one or multiple children is inconsistent with the genetic make-up. In other words, based on the DNA analysis, we can demonstrate that the father or mother is not in fact biological parent of the child. Non-paternity or non-maternity will not be revealed to the family, will not be reported in findings nor released to anyone else. In these rare cases we exclude the individual or a family in question from our study.

After drawing your blood, or taking saliva or surgical tissue if applicable, a sample of your DNA will be kept with Dr. Gleeson indefinitely, and Dr. Gleeson, his associates or successors in these studies will be responsible for deciding how it will be used. In addition to Dr. Gleeson, your DNA may also be studied by Dr. William Dobyns at the University of Chicago; Dr. Elliot Sherr at the University of California, San Francisco; and Dr. Charles Schwartz at the Greenwood Genetic Center. A small amount of your DNA will be sent to a DNA bank for Dr. Gleeson, Dr. Dobyns, Dr. Sherr, Dr. Schwartz, and other future researchers collaborating on this project to use. The sample will not include your name or any other identifying information, but it will be sent with the name of the condition that we are studying in your family. Your sample may be used to validate new genetic mutations or to identify additional mutations in new genes involved in the condition in your family.
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Dr. Gleeson will be responsible for deciding how your child’s specimens will be used. The specimens collected from your child may also be used in additional research to be conducted by the University of California personnel collaborating in this research. These specimens and their derivatives may have significant therapeutic or commercial value. You consent to such uses.

If you decide later that you do not want the specimens collected from you to be used for future research, you may tell this to Dr. Gleeson, who will use his best efforts to stop any additional studies. However, in some cases, such as if your cells are grown up and are found to be generally useful, it may be impossible to locate and stop such future research once the materials have been widely shared with other researchers at the University of California.

If you are injured as a direct result of participation in this research, the University of California will provide any medical care you need to treat those injuries. The University will not provide any other form of compensation to you if you are injured. You may call the Human Research Protections Program at (858) 246-4777 for more information about this, to inquire about your rights as a research subject, or to report research-related problems.

This study may be of no direct benefit to you or members of your family. If as a result of participation in this study we obtain information that could significantly affect your health or well being, we will attempt to inform you of the existence of this information. You may then decide if you wish to know what we have learned. Dr. Gleeson hopes to be able to identify the gene responsible for the condition in your family, as well as develop improved diagnostic procedures and possibly new methods of treatment. In addition, this information may also be used to further our understanding of neurological disorders in other individuals and families.

Instances are known in which a subject in a research study has been required to furnish genetic information as a precondition in applying for health insurance and/or a job. Participation in this study does not mean that you have had genetic testing. Genetic testing means having a test performed and the results provided to you and your doctor. If you are interested in having genetic testing performed, you should consult your doctor, as some commercial tests are available. Your doctor can provide you with the necessary information to determine if such a test would be appropriate for you.

Dr. Gleeson and/or ____________________________ on Gleeson’s behalf has explained this study to you and answered your questions. If you have other questions or research-related problems, you may reach Dr. Gleeson at (858) 246-0547.

Your alternative option to participation in this study is not to participate. Your involvement in research is entirely voluntary. You may refuse to participate or withdraw at any time without jeopardy to the medical care you will receive at this or any institution.
Subject’s Name: __________________________________________

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Investigator’s Name: Dr. Joseph Gleeson

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Research records will be kept confidential to the extent provided by law.

You have received a copy of this consent document and a copy of “Experimental Subject’s Bill of Rights” to keep.

You agree to participate.

__________________________________________
Parent or Guardian’s Signature

__________________________
Witness

________
Date