



Alone we are rare. Together we are strong.®

REQUEST FOR PROPOSALS

The National Organization for Rare Disorders (NORD)'s Rare Disease Research Grants Program

*Announces a research grant opportunity for one grant
up to \$30,000 US for*

Chromosome 6, Partial Trisomy 6q

DEADLINE FOR INITIAL APPLICATIONS: *October 12, 2025 (11:59 pm PT)*

NORD, with donations from the Partial Trisomy 6q community and Dylan's Rare Chromosome Dream Team, is accepting applications for one grant, up to \$30,000 US, for scientific and/or clinical research studies related to Chromosome 6, Partial Trisomy 6q.

Chromosome 6, partial trisomy 6q is an extremely rare chromosomal disorder in which a portion of the 6th chromosome (6q) is present three times (trisomy) rather than twice in cells of the body. Associated symptoms and findings vary in range and severity from case to case. However, many affected infants and children have slow physical development (growth delay) before and after birth, intellectual disabilities; malformation of the skull and facial (craniofacial) region; an unusually short, webbed neck; abnormal bending (flexion), extension, and fixation of certain (joint contractures) such as the fingers, wrists, elbows, knees and hips; and other physical abnormalities that may cause limitations of movement and abnormal postures. In rare cases, individuals with Chromosome 6, partial trisomy 6q may also have internal organ malformations. These may include heart (cardiac), intestinal, kidney (renal), and/or brain (cerebral) abnormalities. Chromosome 6, partial trisomy 6q appears to affect males and females equally.

In individuals with chromosome 6, partial trisomy 6q, all or a portion of the end (distal) region of the long arm (q) of chromosome 6 is present 3 times (trisomy) rather than twice in cells of the body. Reports indicate that, in those with the disorder, the duplicated portion of 6q2 has begun at various points between bands 6q21 to 6q26 and may extend to the end (or "terminal") of chromosome 6q (qter). In most cases, chromosome 6, partial trisomy 6q has been the result of a balanced chromosomal rearrangement in one of the parents. The parental chromosomal rearrangement has usually been a "balanced translocation".

Other names for chromosome 6, partial trisomy 6q include: 6q+ Syndrome, Partial; Chromosome 6, Trisomy 6q2; Distal Duplication 6q; Distal Trisomy 6q; Duplication 6q, Partial; Trisomy 6q, Partial; Trisomy 6q Syndrome, Partial

References

Asadi, Shahin & Ghazinejad, Neda & Ghayor, Shiva. (2023). A comprehensive review of Chromosome 6, Partial Trisomy 6q Syndrome. International Journal of Diabetes and Metabolism. 8. 291-295.

NORD REQUEST FOR PROPOSALS: Chromosome 6, Partial Trisomy 6q

Email: research-programs@rarediseases.org

Kulkarni, S. (2009, April 10). NORD Rare Disease Report: Chromosome 6, Partial Trisomy 6q.
<https://rarediseases.org/rare-diseases/chromosome-6-partial-trisomy-6q/>

Research Objectives

The NORD Rare Disease Research Grant Program was established in 1989 to encourage meritorious scientific and clinical studies designed to improve the diagnosis, understanding of underlying disease mechanisms, or therapy of specific rare diseases. Studies related to Chromosome 6, Partial Trisomy 6q will be considered. Grants will be awarded to qualified researchers to initiate small scientific research studies or clinical trials, the results of which could be used to obtain funding from the NIH, FDA, or other funding agencies, or to attract a corporate sponsor. Ideally, the proposed research should have the potential to lead to the ultimate development of a new or better therapy. Evaluation of proposals will include careful consideration of protocol design, objectiveness of parameters measured, and statistical evaluation proposed.

About NORD

The National Organization for Rare Disorders (NORD) is the leading independent advocacy organization representing all patients and families affected by rare diseases. NORD is committed to the identification, treatment and cure of the more than 7,000 rare diseases, of which approximately 90 percent are still without an FDA-approved treatment or therapy. Rare diseases affect 25-30 million Americans. More than half of those affected are children. NORD began as a small group of patient advocates that formed a coalition to unify and mobilize support to pass the Orphan Drug Act of 1983. For more than 35 years, NORD has led the way in voicing the needs of the rare disease community, driving supportive policies and education, advancing medical research and providing patient and family services for those who need them most. NORD is made strong together with over 300 disease-specific member organizations and their communities and collaborates with many other organizations on specific causes of importance to the rare disease patient community.

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Email: research-programs@rarediseases.org

APPLICATION PROCESS OVERVIEW

INITIAL APPLICATIONS

- Requests for proposals will be released by NORD.
- Letters of intent will be due **October 12, 2025**.
- Letters of intent will be reviewed by NORD reviewers.
- Requests for full proposals will be issued via email in late December 2025.

FULL PROPOSALS

- Full proposal invitations will be issued via email.
- Application requirements for full proposals will accompany these invitations.

AWARDING OF GRANT

- Award announcements will be made via email and posted on NORD's website in May 2026.
- Funding will begin after all necessary documents (e.g., IRB forms, patient consent forms, signed grant agreements) have been received by NORD.

FURTHER INFORMATION

- If the study involves human or animal subjects, copies of governance documents will be required from each site involved in the study before payment can be issued.
- Clinical drug trials must meet requirements established by the U.S. Food & Drug Administration (FDA).
- Duplicate/overlapping funds from any other private or public source are not to be used.
- All applications determined to have scientific merit will be considered, however before an award is issued compliance with international funding regulations must be confirmed when applicable.

INITIAL APPLICATION

Interested applicants should submit a completed application and letter of intent electronically to research-programs@rarediseases.org with "NORD Letter of Intent" as the subject line. Applicants can complete the requested information directly in the document provided below and/or merge any additional required documents into a single PDF file. Incomplete applications may not be considered. All applications must be received by **October 12, 2025 (11:59 pm PT)**.

REQUIRED ELEMENTS CHECKLIST	PAGE	✓
Application summary	4	
Letter of Intent (Maximum Length 2 Pages)	5	
Biographical sketch	6	
List of co-investigators, if applicable	7	
Budget outline	7	
OPTIONAL ELEMENTS	PAGE	✓
Reviewer information	8	

NORD REQUEST FOR PROPOSALS: Chromosome 6, Partial Trisomy 6q

Email: research-programs@rarediseases.org

Letter of support	8	
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APPLICATION SUMMARY

Deadline for Initial Applications: October 12, 2025 (11:59 pm PT)

PRINCIPAL INVESTIGATOR INFORMATION	
Name	
Position/Title	
Email	
Mailing Address	
Telephone	
PROPOSAL INFORMATION	
Project Title	
Project Term	2 YEARS
Funding Amount Requested (\$ US) <i>not to exceed \$30,000 US</i>	
Institution(s) where research will be conducted	
City, State/Province, Country of Institution(s)	
Will research involve human subjects?	☐ YES ☐ NO
Will research involve animals?	☐ YES ☐ NO
How did you hear about this RFP?	☐ NORD ☐ Rare Disease Organization ☐ Professional Organization ☐ Academic Organization ☐ Referral from Colleague ☐ Other (please specify)
Please be as specific as possible:	
Did you hear about this RFP via:	☐ NORD Website ☐ Google ☐ Social Media (Facebook, Twitter, LinkedIn, Instagram) ☐ Other (please specify) ☐ Email ☐ Medical/Research Publication ☐ Rare Action Network

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Please be as specific as possible:	
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LETTER OF INTENT

Please provide a Letter of Intent addressing the following elements of your research. Applicants may use the form below or attach a separate document to address the following details.

Letter of Intent (Not to exceed 2 pages)	
Summary Statement	
Statement of Need - What issue will the research address - Significance of the work - Why you have chosen this issue - Who will benefit from this research - Why this funding is essential	
Project Activity - Overview of research activities - Why this approach is novel - How the research builds upon and differs from previous research (if applicable)	
Credentials Why your institution/program is best equipped to do this research	
Closing	
Principal Investigator Signature REQUIRED	

BIOGRAPHICAL SKETCH

Please provide a biographical sketch and bibliography for the principal investigator. Applicant may use this form or the NIH Biosketch form. Please modify the form to include, when applicable, the following:

Name Position/Title				
Education/Training Begin with baccalaureate or other initial professional education and include postdoctoral training.	Institution and Location	Degree	Year(s)	Field of Study
Research and Professional Experience Concluding with present position, list in chronological order previous employment, experience, and honors. Include present membership on any advisory committee.				
Honors and Awards				

CO-INVESTIGATOR(S)

Name of Co-Investigator	
Position/Title	
Institution	
Email	

Name of Co-Investigator	
Position/Title	
Institution	
Email	

Name of Co-Investigator	
Position/Title	
Institution	
Email	

BUDGET

Please provide a brief budget outline describing how the funding will be used. Do not include PI salary, overhead, or indirect costs. Funding can be used to cover expenses such as staff salary, technical assistance, supplies, and small equipment.

REVIEWER INFORMATION (OPTIONAL)

Please list up to five areas of scientific/medical expertise needed to review this application (optional). Do not list names of individuals.

- 1.
- 2.
- 3.
- 4.
- 5.

Please list below any individuals who should not review this application (optional).

NAME	INSTITUTION	JUSTIFICATION

LETTER OF SUPPORT (OPTIONAL)

Please include a letter of support for your research from an individual advocate (e.g., patient, care partner) or patient advocacy group in this disease space (optional but encouraged).

To receive notification of future funding opportunities through NORD, sign up for NORD research news here: <https://rarediseases.org/communications-sign-up/>