



December 4, 2024

The Honorable Chuck Schumer
Majority Leader
United States Senate
322 Hart Senate Office Building
Washington, D.C. 20510

The Honorable Mike Johnson
Speaker of the House
United States House of Representatives
568 Cannon House Office Building
Washington, D.C. 20515

The Honorable Mitch McConnell
Minority Leader
United States Senate
317 Russell Senate Office Building
Washington, D.C. 20510

The Honorable Hakeem Jeffries
Minority Leader
United States House of Representatives
2433 Rayburn House Office Building
Washington, D.C. 20515

Dear Majority Leader Schumer, and Minority Leader McConnell, Speaker Johnson, Minority Leader Jeffries,

As you work to finalize legislation for consideration before the end of the 118th Congress, the undersigned 213 organizations urge you to pass or include within any larger bill the provisions of the Creating Hope Reauthorization Act (H.R. 7384/S. 4583), which would reauthorize the highly effective Rare Pediatric Disease Priority Review Voucher (PRV) program for at least five years. Reauthorizing the Rare Pediatric Disease PRV program has broad bipartisan support, including approval by the full House of Representatives in September as part of the amended Give Kids a Chance Act (H.R. 3433). The current authorization is set to expire on December 20th, and a timely, clean and long-term reauthorization is critical to maintaining this important incentive which has effectively spurred drug development to help children living with rare diseases.

Since its creation by Congress in 2012, the Rare Pediatric Disease PRV program has helped bring to market therapies for children affected by 39 rare diseases.¹ Without treatment, many of these diseases lead to death or debilitating illness before the children reach adulthood, and only three of these rare diseases had a safe and effective FDA-approved therapy on the market before the program began.² Additionally, more than half of all Rare Pediatric Disease PRV designations occurred in the last four years,³ showing the program is fostering robust drug development where significant unmet therapeutic needs currently exist.

Developing drugs for rare pediatric diseases often presents unique challenges, including small patient populations and difficulties conducting pediatric clinical trials. The incentive established by the Rare Pediatric Disease PRV program is simple and according to the Congressional Budget Office (CBO) score⁴

¹ See: https://rarediseases.org/wp-content/uploads/2024/05/NORD_PRV-white-paper_FINAL.pdf

² Ibid.

³ Mease, C., Miller, K. L., Fermaglich, L. J., Best, J., Liu, G., & Torjusen, E. (2024). Analysis of the first ten years of FDA's rare pediatric disease priority review voucher program: designations, diseases, and drug development. *Orphanet Journal of Rare Diseases*. https://link.springer.com/epdf/10.1186/s13023-024-03097-x03097x?sharing_token=tVsdxcxCuGoLKGG18G02G_BpE1tBhCbncw3BuzI2ROyCDnBKl_41BmSn3a_5qrzjgrLXsufvRX0wtQEnALK9Za3v_5zjNTa3quYxLJ0LC4dnFV94TbHqovQ6Vq5sRWu7_u2v1C7h16jaeLChSswkyx4eSqy_KycTNielqfGSM

⁴ https://www.cbo.gov/system/files/2024-09/suspensions_week_of_9_23_2024_1.pdf

would not have any effect on direct federal spending: if a product manufacturer develops an FDA-approved therapy to treat a rare pediatric disease, the company earns a transferable priority review voucher – and the right to a more expeditious FDA review timeline – that can be used for a subsequent product application or can be sold and transferred to another company. If the Rare Pediatric Disease PRV program is not reauthorized by Congress, a key incentive that has effectively helped bring treatments and cures to kids and their families will end. We cannot afford to let this happen.

As noted, a five-year extension of the Rare Pediatric Disease PRV program passed unanimously by the House of Representatives and has garnered significant bipartisan support. It is also widely supported by the rare disease patient community, with nearly [200 patient organizations](#) signing on to a letter of support for the bill this summer. We urge that you build upon the work done to date and pass a timely, clean and long-term reauthorization to ensure promising science can be translated into treatments and hope for children and families affected by rare diseases.

Thank you for considering this request and please don't hesitate to reach out to Jamie Sullivan at the EveryLife Foundation for Rare Diseases, at jsullivan@everylifefoundation.org and Hayley Mason, Policy Analyst with the National Organization for Rare Disorders, at HMason@rarediseases.org with any questions.

Sincerely,

EveryLife Foundation for Rare Diseases
National Organization for Rare Disorders
Acromegaly Community Inc.
Adrenal Insufficiency United
Advocates for Medically Fragile Kids NC
Aimed Alliance
Aislinn's Wish Foundation
Alagille Syndrome Alliance
Alpha-1 Foundation
Alport Syndrome Foundation
AMDA
American Kidney Fund
Angelman Syndrome Foundation
APBDRFoundation
Aplastic Anemia and MDS International
Foundation
ASXL Rare Research Endowment
Autism Science Foundation
Avery's Hope
BARE Inc
Barth Syndrome Foundation
BDSRA Foundation
BPAN WARRIORS
Bubba's Light
CA Action Link for Rare Diseases (Cal Rare)
CACNA1A Foundation

Canavan Foundation
CDH International
Center for Innovation & Value Research
Charcot Marie Tooth Research Foundation
Child Neurology Foundation
Chondrosarcoma CS Foundation, Inc.
Coalition to Cure Calpain 3
Coalition to Cure CHD2
COMBINEDBrain
Congenital Hyperinsulinism International
Conquer MG
Cooley's Anemia Foundation
CSNK2A1 FOUNDATION
CTNNB1 Connect and Cure
Cure CMD
CURE GABA-A
Cure GM1 Foundation
Cure KCNH1 Foundation
Cure Lowe Foundation
Cure Mito Foundation
Cure Sanfilippo Foundation
Cure SMA
CureARS
CURED Nfp (Campaign Urging Research for
Eosinophilic Diseases)
CureLGMD2i

CureSHANK
 Cyclic Vomiting Syndrome Association
 Cystic Fibrosis Research Institute
 Dana's Angels Research Trust
 debra of America
 Dion Foundation for Children with Rare Diseases
 Dravet Syndrome Foundation
 Dreamsickle Kids Foundation, Inc.
 Dup15q Alliance
 EB Research Partnership
 End AxD
 Endosalpingiosis Foundation INC
 Eosinophilic & Rare Disease Cooperative
 Epilepsy Foundation of America
 Fabry Support & Information Group
 FAM177A1 RESEARCH FUND
 Familial Dysautonomia Foundation
 Family Heart Foundation
 FD/MAS Alliance
 Fighting H.A.R.D. Foundation
 flok Health
 Fondazione Telethon
 Foundation for Angelman Syndrome
 Therapeutics (FAST)
 Foundation for Prader-Willi Research
 Foundation to Fight H-abc
 FRAXA Research Foundation
 Friedreich's Ataxia Research Alliance (FARA)
 GABA-A Alliance
 Galactosemia Foundation
 Gaucher Community Alliance
 GBS|CIDP Foundation International
 Gene Giraffe Project
 Global Genes
 Global Liver Institute
 Glut1 Deficiency Foundation
 Haystack Project
 HCU Network America
 HD-CARE - Huntington's Disease Community
 Advocacy & Education
 Hemophilia Foundation of Southern California
 Hermansky-Pudlak Syndrome Network
 Hope for Hypothalamic Hamartomas
 Hope in Focus
 Huntington's Disease Society of America
 Hydrocephalus Association
 HypoPARAthyroidism Association
 Immune Deficiency Foundation
 INADcure Foundation
 Indo US Organization for Rare Diseases (IndoUSrare)
 International Fibrodysplasia Ossificans Progressiva
 (FOP) Association
 International Foundation for CDKL5 Research
 International Rett Syndrome Foundation
 International Waldenstrom's Macroglobulinemia
 Foundation
 Jack McGovern Coats' Disease Foundation
 Jansen's Foundation
 Jordan's Guardian Angels
 Juju and Friends CLN2 Warrior Foundation
 Kabuki Syndrome Foundation
 KCNQ2 Cure Alliance
 Koolen-de Vries Syndrome Foundation
 Krabbe Connect
 Krishnan Family Foundation
 Lambert Eaton LEMS Family Association
 Lennox-Gastaut Syndrome (LGS) Foundation
 Leukodystrophy Newborn Screening Action Network
 LGMD Awareness Foundation, Inc
 LGMD2D Foundation
 Li-Fraumeni Syndrome Association (LFS Association)
 Little Hercules Foundation
 Lung Transplant Foundation
 MECP2 Duplication Foundation
 Mellie J Foundation
 Mission: Cure
 Mississippi Metabolics Foundation
 MitoAction
 MLD Foundation
 MSUD Family Support Group
 MTM-CNM Family Connection
 Muscular Dystrophy Association
 Myasthenia Gravis Association
 Myositis Support and Understanding
 N=1 Collaborative
 National Alliance for Caregiving
 National Alliance for PANS/PANDAS Action
 National Ataxia Foundation
 National Eosinophilia Myalgia Syndrome Network
 National Fragile X Foundation

National Health Council
National Kidney Foundation
National MPS Society
National Tay-Sachs & Allied Diseases Association
NBIA Disorders Association
Necrotizing Enterocolitis (NEC) Society
Neev Kolte & Brave Ronil Foundation
NephCure
Noah's Hope - Hope4Bridget
NTM Info & Research, Inc.
NW Rare Disease Coalition
Ogden CARES
Organic Acidemia Association
Parent Project Muscular Dystrophy
Partnership to Fight Chronic Disease
Pathways for Rare and Orphan Solutions
Petronille Healthy Society
PMD Foundation
Pompe Alliance
Project Alive
PTEN Hamartoma Tumor Syndrome Foundation
PWSA | USA - Prader-Willi Syndrome Association
Rare New England
Rare Trait Hope Fund
RareRising
Raymond A. Wood Foundation
Rett Syndrome Research Trust
Sanfilippo Children's Foundation
SANFILIPPO SUD
SATB2 Gene Foundation
SCAD Alliance
SHINE Syndrome Foundation
Shwachman-Diamond Syndrome Alliance Inc
Sickle cell association of Kentuckiana
Sisters Hope Foundation
Sleep Consortium
SMS Research Foundation
Spina Bifida Association
Stronger Than Sarcoidosis

Superior Mesenteric Artery Syndrome Research
Awareness and Support
Supporters of Families with Sickle Cell Disease, Inc.
SynGAP Research Fund, DBA cureSYNGAP1
Taylor's Tale
Team Telomere
The Association for Frontotemporal Degeneration
The Bluefield Project to Cure FTD
The Bonnell Foundation: Living with cystic fibrosis
The Children's Medical Research Foundation, Inc.
The DDX3X Foundation
The E.WE Foundation
The Global Foundation for Peroxisomal Disorders
THE KAT6 FOUNDATION INC
The LAM Foundation
The Little Legs Big Heart Foundation
The Louisa Adelynn Johnson Fund for Complex Disease
The MED13L Foundation Inc.
The Mended Hearts, Inc.
The National Adrenal Diseases Foundation
The National PKU Alliance
The Oxalosis and Hyperoxaluria Foundation
The RYR-1 Foundation
The Akari Foundation
Tough Genes
TSC Alliance
U.R. Our Hope
Undiagnosed Diseases Network Foundation
United Mitochondrial Disease Foundation
United MSD Foundation
United Ostomy Associations of America, Inc.
United Porphyrins Association
Uriel E. Owens Sickle Cell Disease Association
of the Midwest
Vasculitis Foundation
Wake Up Narcolepsy, Inc.
Wilson Disease Association
Wisconsin Rare Disease Alliance
Wyldey Nation Foundation
ZTTK SON-Shine Foundation

Cc: The Honorable Bernie Sanders, Chairman, Senate Committee on Health, Education,
Labor and Pensions

The Honorable Bill Cassidy, Ranking Member, Senate Committee on Health, Education,
Labor and Pensions

The Honorable Cathy McMorris Rodgers, Chair, House Committee on Energy and
Commerce

The Honorable Frank Pallone, Ranking Member, House Committee on Energy and
Commerce

The Honorable Robert Casey, Lead Sponsor, Creating Hope Reauthorization Act

The Honorable Markwayne Mullin, Lead Cosponsor, Creating Hope Reauthorization Act

The Honorable Sherrod Brown, Lead Cosponsor, Creating Hope Reauthorization Act

The Honorable Susan Collins, Lead Cosponsor, Creating Hope Reauthorization Act

The Honorable Michael McCaul, Lead Sponsor, Creating Hope Reauthorization Act

The Honorable Anna Eshoo, Lead Cosponsor, Creating Hope Reauthorization Act

The Honorable Gus Bilirakis, Lead Cosponsor, Creating Hope Reauthorization Act

The Honorable Nanette Barragan, Lead Cosponsor, Creating Hope Reauthorization Act

The Honorable Lori Trahan, Lead Cosponsor, Creating Hope Reauthorization Act

The Honorable Michael Burgess, Lead Cosponsor, Creating Hope Reauthorization Act