November 15, 2021

The Honorable Patty Murray Chairwoman Senate Committee on Health Education, Labor and Pensions 428 Dirksen Senate Office Building Washington, DC 20510

The Honorable Richard Neal Chairman House Committee on Ways and Means 1102 Longworth House Office Building Washington, DC 20515

The Honorable Frank Pallone, Jr.
Chairman
House Energy and Commerce Committee
2125 Rayburn House Office Building
Washington, DC 20515

The Honorable Richard Burr Ranking Member Senate Committee on Health Education, Labor and Pensions 428 Dirksen Senate Office Building Washington, DC 20510

The Honorable Kevin Brady Ranking Member House Committee on Ways and Means 1102 Longworth House Office Building Washington, DC 20515

The Honorable Cathy McMorris Rodgers Ranking Member House Energy and Commerce Committee 2125 Rayburn House Office Building Washington, DC 20515

Dear Chairs Murray, Neal, and Pallone and Ranking Members Burr, Brady, and McMorris Rodgers:

The undersigned organizations represent millions of patients, health care providers, and advocates for health equity who are deeply committed to improving our science base through the adequate representation of underrepresented communities in vital medical research. We strongly support passage of the Diversifying Investigations Via Equitable Research Studies for Everyone (DIVERSE) Trials Act (H.R. 5030/S. 2706) and urge you to prioritize this bill in your Committees to ensure it is enacted into law this year. The DIVERSE Trials Act would make it easier for *all* patients to participate in clinical trials while removing barriers that are known to keep certain racial and ethnic groups, older adults, rural residents, and those with limited incomes from being appropriately represented. This legislation has earned bipartisan, bicameral support, takes a practical approach that will lead to near-term improvements, and entails little cost to taxpayers.

The three provisions of the DIVERSE Trials Act will result in more efficient, inclusive, and accessible research, which will benefit the health of all Americans:

First, the legislation would permit individuals to receive financial support for the non-medical costs associated with their participation in clinical trials, by creating a statutory safe harbor for clinical trial sponsors to use in reimbursing such costs. Today, clinical trial sponsors generally supply without charge the drug or technology being investigated while insurers generally cover the other treatment costs associated with a trial. Often, however, patients must pay non-medical costs associated with participating in a clinical trial—such as transportation to a trial site, lodging in another city, meals, and additional childcare. For many people, these costs are difficult to bear and may be the difference between saying "yes" or "no" to a clinical trial that could improve their own care and the care of others suffering from the same disease. For example, a recent study showed that patients making less than

\$50,000 a year are 30 percent less likely to enroll in a cancer clinical trial. Financial barriers are a major barrier for communities underrepresented in clinical trials resulting in trials that do not adequately represent the patient populations who are intended to use the treatments being studied. By clarifying that trial sponsors can financially support patients' non-medical costs, the DIVERSE Trials Act can change this reality for the better.

Second, the DIVERSE Trials Act would allow trial sponsors to provide individuals with the technology necessary for them to participate remotely in clinical trials. Clinical trials almost always require participants to report their condition, symptoms, side effects, or other data on a regular basis and/or to have certain health indicators monitored by web-enabled technology. Some trials of new treatments consist entirely or substantially of such monitoring and so lend themselves to more comfortable and efficient remote participation. For this to work, however, participants must have access to technologies, such as portable computers, tablets, and software applications. This legislation would take the commonsense step of allowing sponsors to provide such technology to participants in a clinical trial. The benefits of this provision include eliminating a financial barrier to participation in clinical trials that are increasingly incorporating web-enabled technology and fostering the conduct of trials that utilize remote participation and include larger cohorts of underrepresented groups.

Finally, the legislation would require the Department of Health and Human Services (HHS) to issue guidance on decentralized clinical trials. While the COVID-19 pandemic accelerated the need for clinical trials in which patients could participate at or close to home, researchers have long endeavored to make clinical trials more convenient for participants. The decentralized approach could open the door to clinical trials for a much broader array of participants, such as those who live hours from a trial site or do not have the ability to make repeated visits to the trial site due to work or caregiving schedules. A significant hinderance to the conduct of decentralized trials has been the absence of clear guidance on the design, conduct, oversight, and other requirements of this research approach. Issuing guidance on the conduct of decentralized clinical trials is an urgent priority for patients, advocates, and health care researchers alike.

We stand at the cusp of a health care revolution—as breakthroughs in genetics, immunotherapy, systems biology, precision medicine, and other specialties lead to more effective treatments and even potential cures for many of the most intractable diseases. Clinical trials are the means by which such breakthroughs are shown to be effective and safe. Making clinical trials more efficient, inclusive, and accessible—as the DIVERSE Trials Act would do—means bringing the growing benefits of medicine and science closer to every American.

Please prioritize the DIVERSE Trials Act (H.R. 5030/ S. 2706) this year to ensure more patients—regardless of their race, ethnicity, economic means, or geographic location—have a chance to enroll in clinical trials.

Sincerely,

Abramson Cancer Center at Penn Medicine Academy of Oncology Nurse & Patient Navigators (AONN) Active Surveillance Patients international ADAP Advocacy Association

¹ Unger, J. M., Gralow, J. R., Albain, K. S., Ramsey, S. D., & Hershman, D. L. (2016). Patient Income Level and Cancer Clinical Trial Participation: A Prospective Survey Study. *JAMA Oncology*, 2(1), 137–139. https://doi.org/10.1001/jamaoncol.2015.3924

Adult Congenital Heart Association

Aimed Alliance

ALK Positive Inc.

Alliance for Aging Research

ALPHA-1 FOUNDATION

American Association for Cancer Research

American Cancer Society Cancer Action Network

American Heart Association

American Liver Foundation

American Lung Association

American Muslim Health Professionals (AMHP)

American Society for Radiation Oncology

American Urological Association

Answer Cancer Foundation

Arthritis Foundation

Asian & Pacific Islander American Health Form

Association for Clinical Oncology

Association of American Cancer Institutes

Association of Black Cardiologists

Association of Community Cancer Centers (ACCC)

Association of Oncology Social Work

Association of Women in Rheumatology

Asthma and Allergy Foundation of America

Autoimmune Association

BlackDoctor, Inc

Breastcancer.org

Cancer Action Coalition of Virginia

Cancer and Careers

Cancer Support Community

CANCER101

Cancer Care

Caregiver Action Network

Children's Cancer Cause

Cholangiocarcinoma Foundation

CLL Society

Color of Crohn's & Chronic Illness

Colorectal Cancer Alliance

Community Access National Network (CANN)

Crohn's & Colitis Foundation

Deadliest Cancers Coalition

Debbie's Dream Foundation: Curing Stomach Cancer

Depression and Bipolar Support Alliance

Digestive Disease National Coalition

Digestive Health Physicians Association (DHPA)

Diverse Health Hub

Epilepsy Foundation

EveryLife Foundation for Rare Diseases

Fight Colorectal Cancer

FORCE: Facing Our Risk of Cancer Empowered

Friends of Cancer Research

Global Healthy Living Foundation

Global Liver Institute

GO2 Foundation for Lung Cancer

GRASP

Head and Neck Cancer Alliance

HealthTree Foundation

Healthy Americas Foundation

HealthyWomen

Heart Valve Voice US

Hemophilia Federation of America

Hep B United

Hepatitis B Foundation

Hispanic Health Network

HISPANIC HEALTHCARE TASK FORCE

HIV + Hepatitis Policy Institute

Hope for Stomach Cancer

ICAN, International Cancer Advocacy Network

Infinite Strength, Inc.

Infusion Access Foundation (IAF)

International Foundation for Autoimmune & Autoinflammatory Arthritis (AiArthritis)

JDRF

KidneyCAN

KRAS Kickers

Latino Commission on AIDS

Lazarex Cancer Foundation

LUNGevity Foundation

Lupus and Allied Diseases Association, Inc.

Lupus Foundation of America

Male Breast Cancer Coalition

Malecare Cancer Support

Metastatic Breast Cancer Network

Moffitt Cancer Center

Multiple Sclerosis Association of America

Muscular Dystrophy Association

Musella Foundation For Brain Tumor Research & Information, Inc.

National Alliance for Hispanic Health

National Alliance of State Prostate Cancer Coalitions

National Association of Nutrition and Aging Services Programs

National Black Nurses Association

National Brain Tumor Society

National Cancer Registrars Association

National Comprehensive Cancer Network

National Eczema Association

National Health Council

National Hispanic Cancer Awareness Day

National Hispanic Council on Aging

National Hispanic Medical Association

National Infusion Center Association (NICA)

National LGBT Cancer Project

National Marrow Donor Program/Be The Match

National Medical Association

National Minority Quality Forum

National MS Society

National Patient Advocate Foundation

National Rural Health Association

National Urban League

NCODA

Oncology Nursing Society

Oral Cancer Foundation

Ovarian Cancer Research Alliance

Pancreatic Cancer Action Network

Patient Advocates In Research (PAIR)

Patient Empowerment Network

Patients Rising

Pennsylvania Prostate Cancer Coalition (PPCC)

Phelan-McDermid Syndrome Foundation

PlusInc

Preparedness and Treatment Equity Coalition

Prevent Blindness

Preventive Cardiovascular Nurses Association

Prostate Cancer International

Prostate Conditions Education Council

Qmetrics Technologies

Research Advocacy Network

RetireSafe

Research!America

Scleroderma Research Foundation

Skin of Color Society

Society for Women's Health Research

Solve M.E.

Spondylitis Association of America

Stand Up To Cancer

Susan G. Komen

THANC (Thyroid, Head and Neck Cancer) Foundation

The EndBrainCancer Initiative (EBCI) (formerly the Chris Elliott Fund)

The Gerontological Society of America

The Kennedy Forum

The Leukemia & Lymphoma Society

The Mended Hearts, Inc

The Multiple Myeloma Research Foundation (MMRF)

The Prostate Health Education Network PHEN

Tigerlily Foundation

Triage Cancer

TSC Alliance

Twisted Pink
U.S. Pain Foundation
UsAgainstAlzheimer's
USC Norris Comprehensive Cancer Center
Vaccinate Your Family
WomenHeart: The National Coalition for Women with Heart Disease
Young Survival Coalition
ZERO - The End of Prostate Cancer

CC: The Honorable Robert Menendez
The Honorable Tim Scott
The Honorable Raul Ruiz
The Honorable Larry Bucshon