March 24, 2022

The Honorable Chuck Schumer Majority Leader U.S. Senate 322 Hart Senate Office Building Washington, DC 20510

The Honorable Mitch McConnell Minority Leader U.S. Senate 317 Russell Senate Office Building Washington, DC 20510 The Honorable Nancy Pelosi Speaker U.S. House of Representatives 1236 Longworth House Office Building Washington, DC 20515

The Honorable Kevin McCarthy Minority Leader U.S. House of Representatives 2468 Rayburn House Office Building Washington, DC 20515

Dear Majority Leader Schumer, Minority Leader McConnell, Speaker Pelosi, and Minority Leader McCarthy:

On behalf of a broad spectrum of patients, advocacy organizations, cancer centers and healthcare professionals, we are writing today to express our support for the *Reducing Hereditary Cancer Act (H.R. 4110/S.B. 3656)*, legislation that would ensure Medicare beneficiary access to genetic testing for hereditary cancer risk, increased screening and risk reducing interventions, when medically necessary and appropriate.

Under existing Medicare guidelines, only a person with "signs, symptoms, complaints, or personal histories of disease" meets the criteria for coverage of medical services. Recognizing the value of cancer prevention and early detection, in recent years Congress has passed legislation allowing for coverage of certain cancer screenings (e.g., mammograms, colonoscopies and PSA tests) for the "average risk" population.

There has been tremendous progress in cancer prevention, detection, and treatment over the past quarter century. Research shows that inherited genetic mutations play a major role in approximately 10% of cancers, including breast, ovarian, endometrial, prostate, pancreatic and colorectal.² Major cancer organizations, genetics, and medical professional societies including the National Comprehensive Cancer Network (NCCN), American Society of Clinical Oncology (ASCO) and others have established guidelines for the assessment and management of hereditary cancer risk.

The U.S. Preventive Services Task Force recognizes the significance of genetics in cancer risk. In 2013, the Task Force published recommendations for *Risk Assessment, Genetic Counseling, and Genetic Testing for BRCA-Related Cancer*. For those with an inherited mutation, the USPSTF notes that management consists of "a variety of interventions to lower future cancer risk. This includes intensive screening, risk-reducing medications, and risk-reducing mastectomy and salpingo-oophorectomy."³

Medicare covers genetic testing only for beneficiaries *already diagnosed with cancer* (regardless of family cancer history or a known genetic mutation in the family). Most private insurers cover genetic counseling and testing for appropriate individuals, including those without a cancer diagnosis, as well as people with a cancer diagnosis. It is crucial that Medicare beneficiaries have access to the same cancer screening and preventive measures as their counterparts with private insurance.

If someone without cancer learns they have an inherited mutation that increases cancer risk (e.g., BRCA1 or BRCA2) prior to Medicare eligibility—or pays out of pocket for genetic testing and finds that they carry an inherited mutation while on Medicare—the individual cannot access the recommended high-risk cancer screenings. Medicare is not permitted to cover these screenings, despite the fact that they are proven to detect cancer earlier, when it is less invasive, less costly, and more easily treated.

Similarly, Medicare is barred from covering potentially lifesaving, risk-reducing procedures, such as bilateral salpingo-oophorectomy (removal ovaries and fallopian tubes). This surgery is crucial for women at high risk of ovarian cancer because there is no reliable screening or early detection; more than 75% of affected women are diagnosed with advanced-stage disease (Stage III or IV).⁴ Up to 25% of ovarian cancers are attributable to an inherited genetic mutation. The median age for diagnosis of ovarian cancer in the U.S. is 63 years, meaning almost half of all persons with ovarian cancer are Medicare beneficiaries. Furthermore, those aged 65 or older with ovarian cancer have significantly worse cancer-related survival than younger patients.⁵

Medicare's inability to cover these potentially life-saving tests and interventions exacerbates health disparities. Access to screening and risk-reducing interventions enables early detection and reduces risk for individuals who carry an inherited mutation. We must prioritize screening, early detection, and prevention in Medicare but to do this requires Congressional action. The National Cancer Institute (NCI) predicts nearly 10,000 excess deaths in the U.S. from breast and colorectal cancer alone over the next 10 years because of pandemic-related delays in cancer screening and treatment.⁶ Enactment of the *Reducing Hereditary Cancer Act of 2021 (H.R. 4110/S.B. 3656)* will improve access to critical screening and preventive care, and may mitigate some of the predicted cancer-related deaths over the next decade and beyond.

We encourage your support of this lifesaving legislation that will ultimately save Medicare tens of millions of dollars and thank you for your time and consideration today. Please contact <u>Lisa Schlager</u> at FORCE or <u>Alyssa Schatz</u> at NCCN with any questions.

Sincerely,

Patient Advocacy Organizations

AliveAndKickn
Alliance for Aging Research
Alliance for Patient Access
American Cancer Society Cancer Action Network
Breast Cancer Action
Brem Foundation to Defeat Breast Cancer

Bright Pink

Cancer ABCs

Cancer Care

Cancer Support Community

Colon Cancer Alliance for Research & Education for Lynch Syndrome

Colon Cancer Coalition

Community Oncology Alliance Patient Advocacy Network (CPAN)

DenseBreast-info, Inc.

Disability Rights Legal Center

Dr. Susan Love Foundation for Breast Cancer Research

Fairview Health Services

Fight Colorectal Cancer

Florida Breast Cancer Foundation

FORCE - Facing Our Risk of Cancer Empowered

Genetic Alliance

GI Cancers Alliance

HealthyWomen

Hereditary Colon Cancer Foundation

HIS Breast Cancer Awareness

ICAN, International Cancer Advocacy Network

ICARE

Hope For Stomach Cancer

The Jewish Federations of North America

Kamie K Preston Hereditary Cancer Foundation

Let's Win! Pancreatic Cancer

Living Beyond Breast Cancer

Living LFS

Lynch Syndrome International

Male Breast Cancer Coalition

National Alliance Against Disparities in Patient Health

National Coalition for Cancer Survivorship

National Ovarian Cancer Coalition

National Patient Advocate Foundation

NothingPink

Ovarian Cancer Project

Ovarian Cancer Research Alliance

Pancreatic Cancer Action Network

Prevent Cancer Foundation

Prostate Cancer Foundation

Raymond Foundation, Inc.

Research Advocacy Network

SHARE Cancer Support

Sharsheret | The Jewish Breast & Ovarian Cancer Community

Stupid Cancer, Inc.

Susan G. Komen

Thelma D. Jones Breast Cancer Fund

Tigerlily Foundation

Triage Cancer

Us TOO International, Inc.

Young Survival Coalition

ZERO - The End of Prostate Cancer

Academia/Professional Societies/Medical Institutions

Abramson Cancer Center, Penn Medicine

Academy of Oncology Nurse and Patient Navigators (AONN)

Advocate Aurora Health

American College of Medical Genetics and Genomics

American College of Obstetricians and Gynecologists (ACOG)

American Urological Association

The American Society of Breast Surgeons

Association for Clinical Oncology

Association for Molecular Pathology

Association of American Cancer Institutes

Association of Community Cancer Centers (ACCC)

Basser Center for BRCA, Penn Medicine

Center for Genomic Interpretation

Community Oncology Alliance (COA)

Consortium for Science, Policy & Outcomes, Arizona State University

Fairview Health Services

Florida Association of Genetic Counselors

Fox Chase Cancer Center

Georgetown Lombardi Comprehensive Cancer Center

Huntsman Cancer Institute at the University of Utah

IL Society of Genetic Professionals

Intermountain Healthcare

International Society of Nurses in Genetics

JScreen

The Lynch Syndrome Screening Network

MHealth Fairview

Michigan Cancer Genetics Alliance

Moffitt Cancer Center

National Association for Nurse Practitioners in Women's Health

National Cancer Registrars Association

National Comprehensive Cancer Network (NCCN)

National Society of Genetic Counselors

Northwestern University Feinberg School of Medicine

Oncology Nursing Society

Palo Alto Medical Foundation

Society of Gynecologic Oncology

Swedish Cancer Institute

UC Santa Cruz Genomics Institute

University of Miami

University of Rochester Medical Center

US Oncology Network

William C. Bernstein Familial Cancer Registry, University of Minnesota

¹ American Society of Clinical Oncology, Genetic Testing Coverage & Reimbursement

² National Cancer Institute, <u>The Genetics of Cancer</u>, Accessed March 22, 2021

³ JAMA | US Preventive Services Task Force | RECOMMENDATION STATEMENT, <u>Risk Assessment, Genetic Counseling, and Genetic Testing for BRCA-Related Cancer</u>, August 20, 2019

⁴ Am Fam Physician. 2016 Jun 1;93(11):937-944

⁵ American Cancer Society, Ovarian Cancer Risk Factors, Accessed March 19, 2021

⁶ Sharpless, N. | COVID-19 and Cancer. Science. June 19, 2020