

118TH CONGRESS  
2D SESSION

# H. RES. 1304

Expressing support for the designation of June 19, 2024, as “World Sickle Cell Awareness Day” in order to increase public awareness across the United States and global community about sickle cell disease and the continued need for empirical research, early detection screenings, novel effective treatments leading to a cure, and preventative care programs with respect to complications from sickle cell anemia and conditions relating to sickle cell disease.

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## IN THE HOUSE OF REPRESENTATIVES

JUNE 18, 2024

Mr. DAVIS of Illinois (for himself, Mr. BURGESS, Ms. LEE of California, Mr. CARTER of Georgia, Mr. BLUMENAUER, Mr. THANEDAR, Ms. WILD, Mr. JACKSON of Illinois, Mr. JOHNSON of Georgia, Ms. CROCKETT, Ms. ADAMS, Mrs. RAMIREZ, Mr. DAVIS of North Carolina, Mr. CARSON, Ms. JACKSON LEE, Mr. GRIJALVA, Ms. NORTON, and Mr. GARAMENDI) submitted the following resolution; which was referred to the Committee on Energy and Commerce

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## RESOLUTION

Expressing support for the designation of June 19, 2024, as “World Sickle Cell Awareness Day” in order to increase public awareness across the United States and global community about sickle cell disease and the continued need for empirical research, early detection screenings, novel effective treatments leading to a cure, and preventative care programs with respect to complications from sickle cell anemia and conditions relating to sickle cell disease.

Whereas sickle cell disease (referred to in this preamble as “SCD”) is a group of inherited red blood cell disorders, a genetic condition present at birth, and a major health problem in the United States and worldwide;

Whereas the 2024 theme of World Sickle Cell Awareness Day, “Hope Through Progress: Advancing Sickle Cell Care Globally”, is an immediate call to bring voices together to improve the health and quality of life for individuals living with SCD and their families;

Whereas, in 1972, Dr. Charles Whitten cofounded the Sickle Cell Disease Association of America to improve research, education, and health care for SCD patients and which is now headquartered in Hanover, Maryland;

Whereas, in 1972, Congress passed the National Sickle Cell Anemia Control Act (Public Law 92–294; 86 Stat. 136), which, for the first time, provided authority to establish education, information, screening, testing, counseling, research, and treatment programs for SCD;

Whereas sickle cell trait (referred to in this preamble as “SCT”) is a gene mutation that causes a single misspelling in the DNA instructions for hemoglobin, a protein that aids in carrying oxygen in the blood, and can result in chronic complications, including anemia, stroke, infections, organ failure, tissue damage, intense periods of pain referred to as vaso-occlusive crises, and even premature death in individuals living with SCD;

Whereas SCT occurs when an individual inherits 1 copy of the sickle cell gene from 1 parent, and, although most individuals who have SCT live normal lives, when both parents have SCT, there is a 25-percent chance that any of their children will have SCD;

Whereas there are an estimated 1,000,000 to 3,000,000 individuals with SCT in the United States, with many unaware of their status;

Whereas an estimated 100,000 individuals have SCD in the United States, with 1 out of every 365 African-American births and 1 out of every 16,300 Hispanic-American births resulting in SCD, and nearly 1 out of 13 African-American babies are born with SCT;

Whereas SCD affects millions of individuals throughout the world, especially individuals of genetic descent from certain countries in sub-Saharan Africa, South and Central America, the Caribbean, South Asia, the Middle East, and the Mediterranean basin;

Whereas the variance relating to the prevalence of SCT ranges greatly by region and demography, with overall rates as high as 40 percent in parts of sub-Saharan Africa and among newborns in parts of India;

Whereas, in many countries that are poor in resources, 90 percent of children with SCD do not live to see adulthood;

Whereas, approximately 1,000 children in Africa are born with SCD each day, more than one half of whom will die before their fifth birthday;

Whereas the high prevalence of SCD in the central and western regions of India results in approximately 20 percent of babies diagnosed with SCD in parts of the western region dying before the age of 2;

Whereas, in 2006, the World Health Assembly passed a resolution, adopted by the United Nations in 2009, recognizing SCD as a public health priority with a call to action for each country to implement measures to tackle

the disease, and in 2010, the World Health Assembly passed a resolution relating to preventing and managing birth defects, including SCD;

Whereas screening newborns for SCD is a crucial first step for families to obtain a timely diagnosis, to obtain comprehensive care, and to decrease the mortality rate for children with respect to SCD;

Whereas approved treatments for SCD are limited, with the Food and Drug Administration approving only 4 SCD therapies since 2017, but, as of the date of adoption of this resolution, there are more than 40 SCD therapies in development;

Whereas there is an immediate need for lifesaving therapeutics that can improve the duration and quality of life for individuals with SCD;

Whereas, in 2020, the National Academies of Sciences, Engineering, and Medicine developed a comprehensive strategic plan and blueprint for action to address SCD, which highlights the need to develop new innovative therapies and to address barriers to the equitable access of approved treatments;

Whereas, in 2020, the Department of Health and Human Services, in partnership with the American Society of Hematology and the SickleInAfrica Consortium, and in collaboration with the World Health Organization, hosted a webinar for a joint effort to strengthen efforts to combat SCD during the COVID–19 pandemic and beyond;

Whereas the late Kwaku Ohene-Frempong, M.D., Professor Emeritus of Pediatrics at the Perelman School of Medicine at the University of Pennsylvania, an American Society of Hematology member who founded and served as a

member of the Global Sickle Cell Disease Network, was a leader in advancing the body of knowledge in SCD research, public health, and medicine and is recognized as immeasurably benefitting thousands of children worldwide;

Whereas there are emerging genetic therapy technologies, including 2 therapies approved by the Food and Drug Administration in December of 2023, that can modify a patient’s own hematopoietic stem cells to enable them to generate healthy red blood cells to prevent sickle cell crises;

Whereas hematopoietic stem cell transplantation (commonly known as “HSCT”) is currently the only cure for SCD, and while advancements in treatment for complications associated with SCD have been made, more research is needed to find widely available and accessible treatments and cures to help individuals with SCD; and

Whereas, although June 19, 2024, has been designated as “World Sickle Cell Awareness Day” to increase public awareness across the United States and global community about SCD, there remains a continued need for empirical research, early detection screenings, novel effective treatments leading to a cure, and preventative care programs with respect to complications from sickle cell anemia and conditions relating to SCD: Now, therefore, be it

1        *Resolved*, That the House of Representatives—

2                (1) supports the goals and ideals of World Sick-  
3        le Cell Awareness Day;

4                (2) commits to ensuring equitable access to new  
5        sickle cell disease (referred to in this resolution as

1 “SCD”) treatments by shining the light among all  
2 economic, racial, and ethnic groups to improve  
3 health outcomes for individuals living with SCD;

4 (3) calls on the Department of Health and  
5 Human Services to create global policy solutions  
6 aimed at providing support for the global community  
7 with respect to SCD and, in partnership with local  
8 governments, the domestic resources needed to pro-  
9 vide access to newborn screening programs, thera-  
10 peutic interventions, and support services with re-  
11 spect to SCD;

12 (4) supports eliminating barriers to equitable  
13 access to innovative SCD therapies, including cell,  
14 gene, and gene-editing therapies in the Medicare and  
15 Medicaid systems for the most vulnerable patients;

16 (5) encourages the people of the United States  
17 and the world to hold appropriate programs, events,  
18 and activities on World Sickle Cell Awareness Day  
19 to raise public awareness of SCD traits, preventa-  
20 tive-care programs, treatments, and other patient  
21 services for those suffering from SCD, complications  
22 from SCD, and conditions relating to SCD;

23 (6) encourages the President to form a Sickle  
24 Cell Disease Interagency Group, which should in-  
25 clude the Department of Health and Human Serv-

1 ices, the Department of Veterans Affairs, the Na-  
2 tional Institutes of Health, the Food and Drug Ad-  
3 ministration, and the Centers for Medicare & Med-  
4 icaid Services, to work toward policies that will sup-  
5 port equitable and appropriate access to innovative  
6 SCD therapies; and

7 (7) with respect to the policies described in  
8 paragraph (6), urges the interagency group de-  
9 scribed in that paragraph to consider options that  
10 not only address access to potential future curative  
11 treatments for SCD, but also address the bias that  
12 the population most affected by SCD continues to  
13 face within the United States and global health care  
14 systems.

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