

Existing heart disease is undiagnosed in half of women who have a first heart attack.

Management of chest pains differ by sex and race. Men are more likely than women to receive definitive diagnoses of angina as opposed to vague chest pain. Women and blacks typically receive fewer cardiovascular medications than men and whites.

Lack of studies on women limits usefulness of research on coronary heart disease. Although CHD causes more than 250,000 deaths in women each year, much of the research on CHD in the last 20 years has either excluded women or included very few women. As a result, many of the tests and therapies used to treat women for CHD are based on studies conducted predominantly in men and may not be as effective in women.

Again, I want to thank Representative MILLENDER-McDONALD for her leadership, and I urge all of my colleagues to support H. Con. Res. 52.

Mr. PALLONE. Mr. Speaker, I yield back the balance of my time.

The SPEAKER pro tempore. The question is on the motion offered by the gentleman from New Jersey (Mr. PALLONE) that the House suspend the rules and agree to the concurrent resolution, H. Con. Res. 52.

The question was taken.

The SPEAKER pro tempore. In the opinion of the Chair, two-thirds being in the affirmative, the yeas have it.

Mr. PALLONE. Mr. Speaker, on that I demand the yeas and nays.

The yeas and nays were ordered.

The SPEAKER pro tempore. Pursuant to clause 8 of rule XX and the Chair's prior announcement, further proceedings on this question will be postponed.

EXPRESSING SENSE OF CONGRESS REGARDING NEED FOR ADDITIONAL RESEARCH INTO HYDROCEPHALUS

Mr. PALLONE. Mr. Speaker, I move to suspend the rules and agree to the concurrent resolution (H. Con. Res. 74) expressing the sense of the Congress regarding the need for additional research into the chronic neurological condition hydrocephalus, and for other purposes.

The Clerk read as follows:

H. CON. RES. 74

Whereas hydrocephalus is a serious neurological condition, characterized by the abnormal buildup of cerebrospinal fluids in the ventricles of the brain;

Whereas there is no known cure for hydrocephalus;

Whereas hydrocephalus affects an estimated one million Americans;

Whereas 1 or 2 in every 1000 babies are born with hydrocephalus;

Whereas over 375,000 older Americans have hydrocephalus, which often goes undetected or is misdiagnosed as dementia, Alzheimer's disease, or Parkinson's disease;

Whereas with appropriate diagnosis and treatment, people with hydrocephalus are able to live full and productive lives;

Whereas the standard treatment for hydrocephalus was developed in 1952, and carries multiple risks including shunt failure, infection, and overdrainage;

Whereas there are fewer than 10 centers in the United States specializing in the treatment of adults with normal pressure hydrocephalus;

Whereas each year, the people of the United States spend in excess of \$1 billion to treat hydrocephalus;

Whereas a September 2005 conference sponsored by 7 institutes of the National Institutes of Health—"Hydrocephalus: Myths, New Facts, Clear Directions"—resulted in efforts to initiate new, collaborative research and treatment efforts; and

Whereas the Hydrocephalus Association is one of the Nation's oldest and largest patient and research advocacy and support networks for individuals suffering from hydrocephalus: Now, therefore, be it

Resolved by the House of Representatives (the Senate concurring), That—

(1) the Congress commends the Director of the National Institutes of Health for working with leading scientists and researchers to organize the first-ever National Institutes of Health conference on hydrocephalus; and

(2) it is the sense of the Congress that—

(A) the Director of the National Institutes of Health should continue the current collaboration with respect to hydrocephalus among the National Eye Institute; the National Human Genome Research Institute; the National Institute of Biomedical Imaging and Bioengineering; the National Institute of Child Health and Human Development; the National Institute of Neurological Disorders and Stroke; the National Institute on Aging; and the Office of Rare Diseases;

(B) further research into the epidemiology, pathophysiology, disease burden, and improved treatment of hydrocephalus should be conducted or supported; and

(C) public awareness and professional education regarding hydrocephalus should increase through partnerships between the Federal Government and patient advocacy organizations, such as the Hydrocephalus Association.

The SPEAKER pro tempore. Pursuant to the rule, the gentleman from New Jersey (Mr. PALLONE) and the gentleman from Oklahoma (Mr. SULLIVAN) each will control 20 minutes.

The Chair recognizes the gentleman from New Jersey.

GENERAL LEAVE

Mr. PALLONE. Mr. Speaker, I ask unanimous consent that all Members may have 5 legislative days to revise and extend their remarks and include extraneous material on the bill that we are considering.

The SPEAKER pro tempore. Is there objection to the request of the gentleman from New Jersey?

There was no objection.

Mr. PALLONE. Mr. Speaker, I yield myself such time as I may consume.

I rise in support of H. Con. Res. 74, expressing the sense of the Congress regarding the need for additional research into the chronic neurological condition hydrocephalus.

Hydrocephalus simply means water on the brain. The term "hydrocephalus" defines a condition characterized by an excessive accumulation of fluid in the brain. This buildup of fluid inside the skull causes the brain to swell, infections of the nervous sys-

tem, lesions or tumors of the brain or spinal cord, and decreased mental function among other symptoms.

The causes of hydrocephalus are not all well understood. It may result from genetic inheritance or developmental disorders. Other possible causes include complications of premature birth, diseases or infections caught before birth, and injury before, during or after childbirth.

Hydrocephalus is believed to affect approximately one in every 500 children. At present, most of these cases are diagnosed prenatally, at the time of delivery, or in early childhood. Advances in diagnostic imaging technology allow more accurate diagnoses in individuals with atypical presentations, including adults with conditions such as normal pressure hydrocephalus.

The National Institute of Neurological Disorders and Stroke, a part of the National Institutes of Health, is the leading support of research on hydrocephalus within the Federal Government. NINDS works collaboratively with other institutes at NIH to further research on the influence of hydrocephalus on development and on the more general issue of the effect of early brain injury. The knowledge gained from this research will foster hope for new methods to treat and prevent developmental brain disorders such as hydrocephalus.

I would like to thank Congressman MIKE THOMPSON of California for his work to bring this resolution before us today, and I would urge my colleagues to support H. Con. Res. 74.

Mr. Speaker, at this time I reserve the balance of my time.

Mr. SULLIVAN. Mr. Speaker, I yield myself as much time as I may consume.

I stand here today in support of this resolution, House Concurrent Resolution 74, addressing the need for additional research into the chronic neurological condition hydrocephalus.

This disease, for which there is no cure, affects an estimated 1 million Americans. Often the symptoms of hydrocephalus are confused with those of dementia, Alzheimer's disease or Parkinson's disease. When the disease is properly identified, people with hydrocephalus are able to live full and productive lives.

The National Institutes of Health has responded to the needs of the hydrocephalus community by working with scientists and researchers to organize a conference in September of 2005 called "Hydrocephalus: Myths, New Facts, Clear Directions."

Demonstrating the need for collaborative research at the National Institutes of Health, seven institutes were able to work together and initiate new research and treatment efforts for hydrocephalus.

I thank Representative MIKE THOMPSON for his work in bringing awareness to this issue.

Mr. Speaker, I yield back the balance of my time.

Mr. PALLONE. Mr. Speaker, I am pleased to yield 5 minutes to the gentleman from California (Mr. THOMPSON), the sponsor of the House concurrent resolution.

Mr. THOMPSON of California. Mr. Speaker, I thank the gentleman for yielding.

I am here today to ask all of my colleagues to support this resolution. Hydrocephalus is an abnormal buildup of fluid in the brain, and it affects nearly 1 million Americans. Without treatment, hydrocephalus is fatal, but when treated, most people with this condition can lead full and productive lives.

I became aware of this condition through a constituent and a friend of mine, Cynthia Solomon, who has a family member who suffers from this condition. Cynthia struggled to find information about hydrocephalus and wanted to connect with other affected families. So she cofounded the first patient advocacy organization for this condition, the Hydrocephalus Association.

As anyone who has been affected by this condition can tell you, the symptoms are many and they vary from person to person. Excess fluid in the brain can cause head enlargement, blurred vision, seizures, learning disabilities and impaired physical development. In older adults, symptoms can mimic dementia, Alzheimer's disease and Parkinson's disease, often leading to a misdiagnosis and a delay in receiving critical and proper treatment.

Doctors do not yet understand the specific causes of hydrocephalus. However, the current treatment was developed back in 1952 and involves surgically inserting a shunt into the brain. This carries serious risk of shunt failure, infection and obstructions. Overdrainage is also a threat. This can trigger a vertical collapse, causing blood vessels to tear and possibly resulting in a subdural hematoma.

Improvement in this treatment is long overdue, and with additional research, we can make it happen.

The National Institutes of Health recognizes this need and recently organized their first ever conference on hydrocephalus. This has resulted in efforts to initiate new collaborative research projects and an expansion of their focus on the development of new treatments.

This resolution commends the NIH for their action and encourages them to continue their collaborative efforts. It also calls for additional research into this serious condition.

However, we cannot depend solely on Federal efforts to expand awareness about hydrocephalus. I commend the Hydrocephalus Association and other groups for their commitment to patient advocacy and public education. Partnerships between these groups, health care providers and the government will bring us closer to our common goal: improved treatment of this condition.

I would like to say a special thanks to Dory Kranz, who is the current di-

rector of the Hydrocephalus Association, for her help in putting this resolution together and her ongoing work in this regard.

I ask my colleagues for their support of this resolution so we can further research into this very serious and important condition and we can bring about improved treatment to those individuals who are affected by this very, very serious and debilitating condition.

Ms. PELOSI. Mr. Speaker, I rise in strong support of H. Con. Res. 74 which has been introduced by my colleague from California, Congressman MIKE THOMPSON, and co-sponsored by Democratic and Republican Members alike.

H. Con. Res. 74 encourages additional support for research into the prevention and treatment of the neurological condition hydrocephalus. It is a chronic medical condition that, like other conditions affecting a relatively small number of people, receives inadequate attention and resources, which delays research that could achieve great breakthroughs. Passage of H. Con. Res. 74 will demonstrate the support of the Congress for aggressive research to find improved methods for detecting and treating hydrocephalus not only among children, but within the increasingly large number to adults who are affected by late onset of the condition.

I am especially proud that the Hydrocephalus Association is headquartered in my congressional district in San Francisco, and that the couple whose pioneering efforts have encouraged and supported so many people with hydrocephalus and their families are San Franciscans—Emily and Russell Fudge, as is the Association's Executive Director, Dory Kranz.

Under their leadership, together with the board composed of leading physicians and researchers, parents and people with hydrocephalus, the Hydrocephalus Association has raised public awareness of this condition and the enormous impact it has on over one million Americans. Because of the medical advances and the advocacy efforts promoted by the Association, most of these children and adults are able to lead full and productive lives and make enormous contributions to our society.

These successes have inadvertently complicated the efforts to advance research, diagnosis and treatment. The typical surgical treatment—the insertion of a shunt to carry away excessive cerebral fluid from the brain—was developed over 50 years ago. Because shunting has alleviated many of the more grave aspects of pre-shunt hydrocephalus, many believe it represents a cure. But it does not. Shunt surgery and the frequent repairs, which are well known to those with hydrocephalus and their families, are not only serious operations, but cost a billion dollars a year, much of which might well be averted with development of advanced treatment strategies.

Promoting additional research through increased federal support is the goal of this resolution. Those advances will benefit not only those with hydrocephalus, but will help to reduce excessive costs in our health care system, and allow hundreds of thousands of people with hydrocephalus to live even fuller lives freed from the anxieties and costs associated with shunt failure and related complications.

Seven of the institutes of the National Institutes of Health—including the Office of Rare

Diseases—sponsored a major national conference in September 2005 on "Hydrocephalus: Myths, New Facts, Clear Directions" which has encouraged aggressive action in the areas of research and treatment. Now it is time for the Congress to join the campaign to expand our understanding of the causes and modernize the treatment of hydrocephalus. I call upon my colleagues to support H. Con. Res. 74 to encourage our nation's leading medical institutions and researchers to expand their focus on achieving breakthrough research in the diagnosis and treatment of hydrocephalus.

Mr. WAXMAN. Mr. Speaker, I rise in strong support of H. Con. Res. 74. This resolution will encourage research into Hydrocephalus, a chronic and often devastating neurological condition.

Hydrocephalus affects an estimated one million Americans—which classifies it as a rare disease. And, unfortunately, like so many other rare diseases, insufficient resources have been directed toward it. Individuals with this disease are forced to undergo "shunting," a highly invasive surgical procedure that carries with it serious safety risks. This procedure also takes a heavy toll on our entire health-care system, costing an average of \$35,000 per procedure.

We can avoid paying this price. With more research and focus on this disease, better treatment—and perhaps even a cure—is within our reach. Patients can be spared the trauma of brain surgery and American citizens can avoid paying more than a billion dollars each year for this treatment.

The NIH has already taken some positive steps toward this goal. By initiating a collaborative effort among 7 NIH institutes and sponsoring a major national conference, the NIH has begun the work that must be done. Now we need to send a strong statement that we want this work to continue.

Cures for rare diseases like Hydrocephalus will never be found unless we increase our effort and follow the scientific promise. We can start with this vote today.

Mr. GEORGE MILLER of California. Mr. Speaker, I rise today to express my support for H. Con. Res. 74 sponsored by Congressman MIKE THOMPSON. Passage of this bill will express federal support for Hydrocephalus research.

Hydrocephalus, a chronic neurological condition that causes cerebrospinal fluid to build up in the brain cavity instead of being reabsorbed into the body, is a disease that affects over one million Americans. This disease can cause head enlargement and blurred vision, learning disabilities and impaired physical development and is fatal if untreated.

Like many other diseases that affect a relatively small portion of our population, Hydrocephalus research lacks proper funding. It is deplorable that the current standard treatment, which requires the insertion of a shunt into the brain to drain out the fluid, was designed in 1952. Shunts are extremely prone to infections and frequently require repair through major surgery.

Modern medicine can do better. I am certain that with federal support for additional research we can develop a better treatment, if not a cure, for those suffering from Hydrocephalus and help them live healthier, fuller lives.

I applaud my colleague, Mr. THOMPSON, for his efforts in this area and I encourage my colleagues to support this resolution.

Mr. PALLONE. Mr. Speaker, I have no additional requests for time, and I yield back the balance of my time.

The SPEAKER pro tempore. The question is on the motion offered by the gentleman from New Jersey (Mr. PALLONE) that the House suspend the rules and agree to the concurrent resolution, H. Con. Res. 74.

The question was taken; and (two-thirds being in the affirmative) the rules were suspended and the concurrent resolution was agreed to.

A motion to reconsider was laid on the table.

RECESS

The SPEAKER pro tempore. Pursuant to clause 12(a) of rule I, the Chair declares the House in recess until approximately 6:30 p.m. today.

Accordingly (at 5 o'clock and 14 minutes p.m.), the House stood in recess until approximately 6:30 p.m.

□ 1830

AFTER RECESS

The recess having expired, the House was called to order by the Speaker pro tempore (Mr. ALTMIRE) at 6 o'clock and 30 minutes p.m.

ANNOUNCEMENT BY THE SPEAKER PRO TEMPORE

The SPEAKER pro tempore. Pursuant to clause 8 of rule XX, proceedings will resume on motions to suspend the rules previously postponed.

Votes will be taken in the following order:

H. Con. Res 47, by the yeas and nays; H.R. 755, by the yeas and nays;

H.R. 884, by the yeas and nays.

The vote on H. Con. Res 52 will be taken tomorrow.

The first electronic vote will be conducted as a 15-minute vote. Remaining electronic votes will be conducted as 5-minute votes.

SUPPORTING THE GOALS AND IDEALS OF A NATIONAL MEDAL OF HONOR DAY

The SPEAKER pro tempore. The pending business is the question of suspending the rules and agreeing to the concurrent resolution, H. Con. Res. 47.

The Clerk read the title of the concurrent resolution.

The SPEAKER pro tempore. The question is on the motion offered by the gentleman from Missouri (Mr. SKELTON) that the House suspend the rules and agree to the concurrent resolution, H. Con. Res. 47, on which the yeas and nays are ordered.

The vote was taken by electronic device, and there were—yeas 411, nays 0, not voting 22, as follows:

[Roll No. 103]

YEAS—411

Abercrombie	Delahunt	Jordan
Ackerman	DeLauro	Kagen
Aderholt	Dent	Kanjorski
Akin	Diaz-Balart, L.	Kaptur
Alexander	Diaz-Balart, M.	Keller
Allen	Dicks	Kennedy
Altmire	Dingell	Kildee
Andrews	Doggett	Kilpatrick
Arcuri	Donnelly	Kind
Baca	Doolittle	King (IA)
Bachmann	Doyle	King (NY)
Bachus	Drake	Kirk
Baird	Dreier	Klein (FL)
Baker	Duncan	Kline (MN)
Baldwin	Edwards	Knollenberg
Barrett (SC)	Ehlers	Kucinich
Barrow	Ellison	Kuhl (NY)
Bartlett (MD)	Ellsworth	LaHood
Barton (TX)	Emerson	Lamborn
Bean	Engel	Lampson
Becerra	English (PA)	Langevin
Berkley	Eshoo	Lantos
Berman	Etheridge	Larsen (WA)
Berry	Everett	Larson (CT)
Biggert	Fallin	Latham
Bilbray	Farr	LaTourette
Bilirakis	Fattah	Lee
Bishop (GA)	Feeney	Levin
Bishop (NY)	Ferguson	Lewis (GA)
Bishop (UT)	Filner	Lewis (KY)
Blackburn	Flake	Linder
Blumenauer	Forbes	Lipinski
Blunt	Fortenberry	LoBiondo
Boehner	Fossella	Loeb
Bonner	Fox	Lofgren, Zoe
Bono	Frank (MA)	Lowey
Boozman	Franks (AZ)	Lucas
Boren	Frelinghuysen	Lungren, Daniel
Boswell	Galleghy	E.
Boucher	Garrett (NJ)	Lynch
Boustany	Gerlach	Mack
Boyd (FL)	Giffords	Mahoney (FL)
Boyd (KS)	Gilchrest	Maloney (NY)
Brady (PA)	Gillibrand	Manzullo
Braley (IA)	Gillmor	Marchant
Brown (SC)	Gingrey	Markey
Brown-Waite,	Gohmert	Marshall
Ginny	Gonzalez	Matheson
Buchanan	Goode	Matsui
Burgess	Goodlatte	McCarthy (CA)
Burton (IN)	Gordon	McCarthy (NY)
Butterfield	Granger	McCaul (TX)
Buyer	Graves	McCollum (MN)
Calvert	Green, Al	McCotter
Camp (MI)	Green, Gene	McCrery
Campbell (CA)	Grijalva	McDermott
Cannon	Gutierrez	McGovern
Cantor	Hall (NY)	McHenry
Capito	Hall (TX)	McHugh
Capps	Hare	McIntyre
Capuano	Harman	McKeon
Cardoza	Hastings (FL)	McMorris
Carnahan	Hayes	Rodgers
Carney	Heller	McNerney
Carson	Hensarling	McNulty
Carter	Herger	Meehan
Castle	Herseth	Meek (FL)
Castor	Higgins	Melancon
Chabot	Hill	Mica
Chandler	Hinche	Michaud
Clarke	Hinojosa	Millender-
Cleaver	Hirono	McDonald
Clyburn	Hobson	Miller (FL)
Coble	Hodes	Miller (MI)
Cohen	Hoekstra	Miller (NC)
Cole (OK)	Holden	Miller, Gary
Conaway	Holt	Miller, George
Conyers	Honda	Mitchell
Cooper	Hooley	Mollohan
Costa	Hoyer	Moore (KS)
Costello	Hulshof	Moore (WI)
Courtney	Inglis (SC)	Moran (VA)
Cramer	Inslee	Murphy (CT)
Crenshaw	Israel	Murphy, Patrick
Crowley	Issa	Murphy, Tim
Cuellar	Jackson (IL)	Murtha
Culberson	Jackson-Lee	Musgrave
Cummings	(TX)	Myrick
Davis (AL)	Jefferson	Nadler
Davis (CA)	Jindal	Napolitano
Davis (IL)	Johnson (GA)	Neal (MA)
Davis (KY)	Johnson (IL)	Neugebauer
Davis, David	Johnson, E. B.	Nunes
Davis, Lincoln	Johnson, Sam	Oberstar
Deal (GA)	Jones (NC)	Obey
DeGette	Jones (OH)	Olver

Ortiz	Salazar	Thompson (CA)
Pallone	Sali	Thompson (MS)
Pascarella	Sánchez, Linda	Thornberry
Pastor	T.	Tiahrt
Paul	Sanchez, Loretta	Tiberi
Payne	Sarbanes	Tierney
Pearce	Saxton	Towns
Pence	Schakowsky	Turner
Perlmutter	Schiff	Udall (CO)
Peterson (MN)	Schmidt	Udall (NM)
Peterson (PA)	Schwartz	Upton
Petri	Scott (GA)	Van Hollen
Pickering	Scott (VA)	Velázquez
Pitts	Sensenbrenner	Visclosky
Platts	Serrano	Walberg
Poe	Sessions	Walden (OR)
Pomeroy	Sestak	Walsh (NY)
Porter	Shadegg	Walz (MN)
Price (GA)	Shays	Wamp
Price (NC)	Shea-Porter	Wasserman
Pryce (OH)	Sherman	Wasserman
Putnam	Shimkus	Schultz
Radanovich	Shuler	Waters
Rahall	Shuster	Watson
Ramstad	Simpson	Watt
Rangel	Sires	Waxman
Regula	Skelton	Weiner
Rehberg	Slaughter	Welch (VT)
Reichert	Smith (NE)	Weldon (FL)
Renzi	Smith (NJ)	Weller
Reyes	Smith (WA)	Westmoreland
Reynolds	Snyder	Whitfield
Rodriguez	Solis	Wicker
Rogers (AL)	Souder	Wilson (NM)
Rogers (KY)	Spratt	Wilson (OH)
Rogers (MI)	Stearns	Wilson (SC)
Rohrabacher	Stupak	Wolf
Ros-Lehtinen	Sullivan	Woolsey
Roskam	Sutton	Wu
Roybal-Allard	Tancredo	Wynn
Royce	Tanner	Yarmuth
Ruppersberger	Tauscher	Young (AK)
Ryan (OH)	Taylor	Young (FL)
Ryan (WI)	Terry	

NOT VOTING—22

Brady (TX)	Hastert	Rothman
Brown, Corrine	Hastings (WA)	Rush
Clay	Hunter	Smith (TX)
Cubin	Kingston	Space
Davis, Jo Ann	Lewis (CA)	Stark
Davis, Tom	Meeks (NY)	Wexler
DeFazio	Moran (KS)	
Emanuel	Ross	

□ 1903

Mr. REICHERT and Mr. FLAKE changed their vote from “nay” to “yea.”

So (two-thirds being in the affirmative) the rules were suspended and the concurrent resolution was agreed to.

The result of the vote was announced as above recorded.

A motion to reconsider was laid on the table.

PROMOTING TRANSPARENCY IN FINANCIAL REPORTING ACT OF 2007

The SPEAKER pro tempore. The pending business is the question of suspending the rules and passing the bill, H.R. 755.

The Clerk read the title of the bill.

The SPEAKER pro tempore. The question is on the motion offered by the gentleman from Georgia (Mr. SCOTT) that the House suspend the rules and pass the bill, H.R. 755, on which the yeas and nays are ordered.

This will be a 5-minute vote.

The vote was taken by electronic device, and there were—yeas 412, nays 0, not voting 21, as follows: