

already seeing its impacts across the country. But it also provides economic opportunity, if we invest in R&D to develop new renewable sources and efficiency technologies. This bill would jeopardize American innovation in this critical area.

Today's bill also meddles in decision-making at our federal research institutions, decreasing funding at certain directorates at the National Science Foundation and imposing new requirements in the grant-making process. Our science agencies have a robust review process in place to fund the most critical research. Politics should have no part in that process.

Unlike earlier America COMPETES bills that were built on broad consensus, HR 1806 is opposed by the vast majority of our nation's scientific community. I urge my colleagues to listen to these scientists and bring forward a bill that invests in American innovation and supports the cutting-edge research necessary to maintain our leadership in the world.

SUPPORTING THE PEOPLE OF NEPAL

HON. TED POE

OF TEXAS

IN THE HOUSE OF REPRESENTATIVES

Thursday, May 21, 2015

Mr. POE of Texas. Mr. Speaker, after the massive earthquake shook Nepal, Eric Jean and Della Hoffman were stranded on a remote trail with a group of other backpackers. Friends of theirs from their time in college at Rice University, which is in my district, contacted my office for help. We worked with the State Department to set them free. Five days after the earthquake, Special Forces came to the village and rescued them.

Ms. Hoffman would later recall, "I don't think we even knew what was happening until some of the villagers ran out of the hut and then, immediately after that, the boulders just started coming down from both sides of the canyon, including into the village and on top of the houses."

Six U.S. Marines and two Nepalese soldiers sacrificed their lives attempting to save others just like Eric and Della when their helicopter crashed.

I stand with my colleagues in support of House Resolution 235, and urge the administration to work with the Nepal government and the international community to deliver aid quickly, easily, and with long-term rebuilding in mind.

And that's just the way it is.

HONORING BUENA VISTA WINERY'S WINE TOOL MUSEUM

HON. MIKE THOMPSON

OF CALIFORNIA

IN THE HOUSE OF REPRESENTATIVES

Thursday, May 21, 2015

Mr. THOMPSON of California. Mr. Speaker, I rise today to recognize Buena Vista Winery on the occasion of the opening of their new, first-of-its-kind Wine Tool Museum. The oldest premium winery in the United States, Buena Vista has been in operation since 1857 when a Hungarian immigrant, Count Agoston Haraszthy, established the vineyards and built

the winery. Over the intervening years, the winery has had a colorful history, passing through many hands before ultimately being purchased by Jean-Charles Boisset in 2011. Boisset immediately hatched a plan to turn the property around that eventually culminated in the creation of the Wine Tool Museum, which officially opened to the public on March 24, 2015.

The Boisset family already had an impressive wine tool collection when Jean-Charles's sister encountered a man in Burgundy looking for a buyer for his enormous thirty thousand item collection of wine tools. With the combination of Boisset family implements and the new acquisitions, the Museum's collection encompasses items as diverse as antique plows and blades, secateurs, and wine harvest baskets.

Along with its impressive collection of tools, the Museum guides visitors through history with an educational film that traces winemaking in the region from the early days of Haraszthy through the plague of phylloxera. The first of its kind museum tells the story of California's wine community. People will not only be able see, but also learn about, the tools that brought wine from the vine to the bottle a century and a half ago.

Mr. Speaker, it is fitting and proper that we honor Buena Vista Winery at this time. Its commitment to not only preserving viticultural history, but demonstrating the evolution of the profession, will help increase awareness and appreciation for California and Sonoma's long history of winemaking.

RECOGNIZING THE SERVICE OF DEPUTY SHERIFF JOSE ALVARADO

HON. JOHN KATKO

OF NEW YORK

IN THE HOUSE OF REPRESENTATIVES

Thursday, May 21, 2015

Mr. KATKO. Mr. Speaker, I rise today to recognize the service of José "Joey" Alvarado to the community of Wayne County, New York. Deputy Sheriff Alvarado has given 30 years of dedicated service to the Wayne County Sheriff's Office. In his role as Deputy Sheriff, he has played an essential role in safeguarding the rights and freedoms of the residents of his county.

Throughout his 30 years of service to the Wayne County Sheriff's Office, Deputy Sheriff Joey Alvarado has consistently performed with professionalism and dedication, working towards the goal of making his community a safer place.

Since 1985, Deputy Sheriff Joey Alvarado has served as a Jailor/Dispatcher, Correction Officer, and Deputy Sheriff. During his tenure as Deputy Sheriff, he was also assigned to the Sheriff's Office Special Investigation Unit. He has made an outstanding contribution through his career as Deputy Sheriff to the quality of life for all Wayne County residents.

I commend Deputy Sheriff Alvarado's sacrifice and contribution to the Wayne County community and wish him the very best in his retirement.

HONORING PENNSYLVANIA CYSTIC FIBROSIS, INC. (PACFI)

HON. TOM MARINO

OF PENNSYLVANIA

IN THE HOUSE OF REPRESENTATIVES

Thursday, May 21, 2015

Mr. MARINO. Mr. Speaker, I rise today in order to congratulate Pennsylvania Cystic Fibrosis, Inc. (PACFI) for 30 years of work in raising Cystic Fibrosis awareness and funding Cystic Fibrosis research efforts. PACFI is an independent, non-profit, all volunteer organization that provides crucial services and much needed support for Pennsylvania families affected by Cystic Fibrosis.

PACFI was founded on October 2, 1985. The PACFI organization is unique in that they do not have a paid staff and operate solely with volunteers. This allows PACFI to use 100% of donations they receive to provide benefits for Pennsylvania families such as paying for emergency and other medical expenses.

PACFI also works to fund Cystic Fibrosis research around the country. To date, they have raised more than \$565,000 for institutions and universities that are on the leading edge of Cystic Fibrosis research. These donations are helping to discover better treatment options and will hopefully lead to a cure.

PACFI is doing excellent work in Pennsylvania on one of the most common and fatal genetic diseases. Cystic Fibrosis affects approximately 1 in 2000 people and their life expectancy is only 35 years. I commend PACFI for doing great work in the field of Cystic Fibrosis research and supporting families that need assistance with the costs of Cystic Fibrosis treatment.

CONGRATULATING SOFIA VICTORIA DE LA PENA ON FOUNDING THE FIT KIDS DAY NON-PROFIT

HON. MARIO DIAZ-BALART

OF FLORIDA

IN THE HOUSE OF REPRESENTATIVES

Thursday, May 21, 2015

Mr. DIAZ-BALART. Mr. Speaker, I rise today to congratulate Sofia Victoria de la Pena on her work founding Fit Kids Day, and establishing the group in the South Florida community.

Sofia Victoria was a ninth grader at Carrollton School of the Sacred Heart in Coconut Grove, FL when she recognized an increased need for health programs for children. She wanted to add to the work being done to combat childhood obesity, and came up with a plan. Her idea was an entire day focused on fitness, which became known as Fit Kids Day.

To get the event started, Sofia Victoria reached out to leading students at other schools to be ambassadors for the program. The students walked throughout their neighborhoods, to spread awareness for the event and attract community support. Local businesses chipped in to help with the first event, providing services and food to the participants, which were offered free of charge. Since the first Fit Kids Day, the program has expanded, and multiple cities have organized their own events. In these cities, mayors or city managers organize a day of activities that are popular in their area.

Fit Kids Day caught on as an event, and is now a nonprofit organization. The Presidential Fitness Counsel has since talked to Sofia Victoria about using the Fit Kids Day model for their organization. The event was created by kids, for kids, and offers many leadership opportunities. In addition, the program has helped introduce fitness ideas and plans to less fortunate communities.

Mr. Speaker, I am honored to congratulate Sofia Victoria de la Pena on her accomplishment, and I ask my colleagues to join me in recognizing her outstanding achievement. It is an honor to know a family that continues to work hard to improve their community every day.

**HONORING DAVIESS COUNTY, KY,
ON CELEBRATING ITS BICENTEN-
NIAL**

HON. BRETT GUTHRIE

OF KENTUCKY

IN THE HOUSE OF REPRESENTATIVES

Thursday, May 21, 2015

Mr. GUTHRIE. Mr. Speaker, I rise today to congratulate Daviess County, Kentucky, on celebrating its bicentennial.

Daviess County came into its existence on June 1, 1815, through an Act of the Kentucky General Assembly. But on May 30, 2015, Daviess County will begin to celebrate its 200th birthday.

Daviess County lies in the Western Kentucky Coalfield region and is also an oil producer. Bounded by the Ohio River, it serves as an important trade and transportation artery—making it a major manufacturing center, transportation hub and U.S. Customs Port of Entry on the Ohio River. It is also a leader in health care, medical research and pharmaceutical experimentation.

The keystone celebration, on May 30th, includes an early morning 5 k run/walk at the Mount Saint Joseph Motherhouse of the Ursuline Sisters. Activities for people of all ages will take place around the Courthouse Square, and the day will end with a reception and the opening of a special Bicentennial Art Exhibit at the Owensboro Museum of Fine Art.

Other activities will include honoring African Americans from Daviess County who served in the Union Army during the Civil War with the dedication of an historic highway marker on the Courthouse lawn. There will also be a series of Bicentennial-related programs at the Daviess County Public Library throughout the summer.

A new history of Daviess County, Kentucky, *Celebrating Our Heritage*, has been published and highlights many aspects of daily life in the county. Among these topics is Daviess County's rich history in the agriculture industry—citing corn, soybean and tobacco producers.

Daviess County is the home to some of Kentucky's great colleges and universities, including: Brescia University, Kentucky Wesleyan College, a campus of the Kentucky Community and Technical College System and a branch of Western Kentucky University.

It is important to also highlight the leadership in Daviess County and all their efforts to make this a successful and thriving community. Thank you for making Daviess County what it is today.

I encourage everyone in Kentucky's Second District to join in the festivities to celebrate the

rich history and traditions of Daviess County. I congratulate all who live and serve the county and look forward to taking part in some of these celebrations myself. Here's to many more years of success.

**ADVANCING RESEARCH FOR
HYDROCEPHALUS PATIENTS**

HON. CHRISTOPHER H. SMITH

OF NEW JERSEY

IN THE HOUSE OF REPRESENTATIVES

Thursday, May 21, 2015

Mr. SMITH of New Jersey. Mr. Speaker, I rise today to call attention to legislation I introduced last week, the Advancing Research for Hydrocephalus Act, and urge my colleagues to cosponsor this important bill. My new legislation—which is supported by the national Hydrocephalus Association—will facilitate better research into this devastating condition by requiring the collection of demographic information on the hydrocephalus community.

Hydrocephalus, which is defined as an abnormal accumulation of cerebrospinal fluid (CSF) within cavities in the brain, can cause brain damage, vision issues, and extreme pain for those affected.

One of those affected, Adrienne D'Oria, a 22 year old from my Congressional District, has suffered from hydrocephalus since she was 10 months old. In addition to the excruciating pain, complications from shunt malfunctions, dozens of brain surgeries and hundreds of hospital visits have essentially eliminated any chance of a normal childhood. Hydrocephalus continues to limit her options for the future:

All of my friends, everyone I went to school with is graduating and starting the next stage in their life. I can't do that," she said recently. "I had to withdraw from so many classes because of hospital admissions and all the surgeries. Even though I've been out of high school for four years I only have the credits of a freshman. My friends are graduating and I'm stuck in limbo. I can't control it.

Unfortunately for Adrienne and thousands like her, the most common treatment for hydrocephalus remains a surgically-inserted shunt. Shunts drain the fluid from the brain through the neck and into other parts of the body. They frequently become blocked, malfunction, or cause infection. In almost half of all cases in children, the shunt fails within the first two years. When they do, patients must immediately locate a medical facility and a neurosurgeon who can correct the problem. This precarious situation is a constant source of fear for those who suffer from hydrocephalus and their families. In fact, hydrocephalus is the most common reason for brain surgery in children.

The scientific and medical communities not only have very few resources that can help them in understanding this condition, they are not even aware of the true impact of this disorder. Without better data and research, they cannot develop more effective treatments.

Mr. Speaker, there are some estimates that this condition affects roughly one million Americans. Yet given that hydrocephalus can occur either congenitally or be acquired, oftentimes through infection or traumatic brain injury, reporting of hydrocephalus has been inconsistent. Currently no mechanism exists to

identify and track persons with hydrocephalus who develop the condition after birth. As a result, we do not have a good grasp on the demographics of hydrocephalus patients.

My bill provides a remedy. The Advancing Research for Hydrocephalus Act will establish a National Hydrocephalus Surveillance System (NHSS) to collect information on the incidence and prevalence of hydrocephalus among a range of demographics, including changes in epidemiology over time. This surveillance system would provide a wealth of data for researchers. Better surveillance will facilitate better research and lead to better outcomes, treatment and care for the infants, children, and adults experiencing the agonizing pain of hydrocephalus.

So I urge my colleagues to support my legislation to help provide assistance and raise the quality of life for individuals, like Adrienne, who are suffering from this condition.

**IN RECOGNITION OF EAST HAR-
LEM COUNCIL FOR HUMAN SER-
VICES, CELEBRATING 50 YEARS OF
SERVICE**

HON. CHARLES B. RANGEL

OF NEW YORK

IN THE HOUSE OF REPRESENTATIVES

Thursday, May 21, 2015

Mr. RANGEL. Mr. Speaker I rise today to give recognition to the East Harlem Council for Human Services, which is celebrating 50 years of serving the local community. The work done by the Council has been vital to our community and has changed the lives of so many for decades. Groups like the Council and institutions like the Boriken Care Center are worth fighting for, and I will continue to make sure there are adequate resources available for them to continue to thrive and serve their community.

I am proud to honor the East Harlem Council for Human Services which was incorporated in 1965 though grass-roots efforts of local East Harlem residents committed to addressing unmet needs in their community. The Council is a multi-service not-for-profit which coordinates an array of affordable and no cost services to more than 10,000 individuals each year without regard for an individual's ability to pay for services. The Board of Directors and more than almost 200 staff members are united in a commitment to the Council's mission of providing the highest quality of comprehensive, community-based, fully bilingual services to the East Harlem community.

The Council is the largest grass-roots, multi-service not-for-profit in our East Harlem community. By continuing its strong history of diverse community-based leadership, and commitment to the self-determination of this institution and the East Harlem community at large, the Council has positioned itself for continued growth. The Council continuously renews its commitment to the residents of El Barrio to ensure that the Boriken Neighborhood Health Center and its sister programs will continue to provide affordable quality comprehensive services in East Harlem for many more decades.

There's no finer work than fighting to bring health and essential wellness to those struggling to make ends meet in underserved communities. Everyone deserves access to quality