The result of the vote was announced as above recorded.

A motion to reconsider was laid on the table.

□ 1900

HONORING THE MEMORY OF TONY CASTILLO

(Mr. COFFMAN asked and was given permission to address the House for 1 minute and to revise and extend his remarks.)

Mr. COFFMAN. Mr. Speaker, I rise today to honor the memory of Tony Castillo, a longtime friend of mine from my hometown of Aurora, Colorado, who recently passed away.

Tony and I had a friendship that started in our youth. We had so much in common. Tony and I both came from military families. Our late fathers had both married war brides in the aftermath of World War II. They were career enlisted soldiers who were transferred to Fitzsimons Army Medical Center for their last assignment in 1964 when we were both just 9 years old. Our military families both came from previous assignments in Europe.

While I followed in my father's footsteps and joined the military, Tony stayed in Aurora and eventually married the love of his life, Nita Adkins of Pueblo, Colorado. Tony and Nita raised two extraordinary children, Ben and Jess, in a loving family that has been inseparable.

Tony was an extraordinary example of a great friend, a loving husband, a devoted father, and he will always be remembered and missed by all who knew him.

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NATIONAL FOSTER CARE MONTH

(Mr. LANGEVIN asked and was given permission to address the House for 1 minute and to revise and extend his remarks.)

Mr. LANGEVIN. Mr. Speaker, as cochair of the Congressional Caucus on

Foster Youth, I rise today in recognition of May as National Foster Care Month.

Growing up, my parents welcomed many foster children into our family, providing them with a stable and nurturing environment.

One of the biggest challenges for youth in foster care is finding their forever family. Even youth who age out of foster care still need a place to call home after they turn 18, and whether through reunification, adoption, or an individual plan, every foster youth deserves a permanent home.

Mr. Speaker, these children belong to all of us, and we are all responsible for their well-being. I urge my colleagues to join me in recognizing May as National Foster Care Month.

RECOGNIZING MEMBERS OF PENN STATE'S RED CELL ANALYTICS LAB

(Mr. THOMPSON of Pennsylvania asked and was given permission to address the House for 1 minute and to revise and extend his remarks.)

Mr. THOMPSON of Pennsylvania. Mr. Speaker, I rise today to recognize members of Penn State University's Red Cell Analytics Lab, a unique organization intended to educate students on how to combat threats in an age of ever-evolving technology.

Last weekend, I visited with members of the lab, and I listened in awe to how they are working with local law enforcement to keep students, staff, and spectators safe at events such as Penn State football games and the university's annual dance marathon, or THON

Quite simply, these students have become experts at recognizing threats and security holes—essential skills in today's world where governments, technology firms, and corporations must stay one step ahead of those who seek to do us harm. I was happy to learn that members of the Red Cell Analytics Lab move on to great careers, working for companies in our financial and technology sectors, along with the Federal Government.

It is unfortunate that these talents are necessary in today's world, but I am glad they are being cultivated at Penn State University.

THE HEROIN EPIDEMIC

(Mr. RYAN of Ohio asked and was given permission to address the House for 1 minute and to revise and extend his remarks.)

Mr. RYAN of Ohio. Mr. Speaker, this week, we are going to take up a package of bills that I support dealing with the massive heroin epidemic plaguing places like Ohio, but all over the country.

In my congressional district, we saw, a few months back, over 22 overdoses. In Trumbull County alone, 7 deaths; in Summit County, city of Akron, 56 people died of a heroin overdose, compared

to just 40 in 2013. Now, in Ohio, these drug overdoses have been the leading cause of accidental deaths in our State, bypassing car accidents.

This House this week is going to do the responsible thing and pass key legislation and hopefully combine with the Senate legislation. But there is a missing piece on the recovery side. We need to make sure that, as we address this issue, it is comprehensive: it is about addiction and it is about treatment, but it also must be about recovery.

So I commend my colleagues on both sides of the aisle for these important pieces of legislation that are coming forth. I will support them, but knowing that we have got a lot more work to do and that the bottom line is we need money behind these programs.

The President put in his budget over \$1 billion. We need to make sure that the resources get down to the local community so that we can help stem the tide of this epidemic in our country.

SUPPORTING TSC GLOBAL AWARENESS DAY AND MAX LUCCA

(Ms. ROS-LEHTINEN asked and was given permission to address the House for 1 minute and to revise and extend her remarks.)

Ms. ROS-LEHTINEN. Mr. Speaker, today I rise to lend my voice in support of TSC Global Awareness Day on May

Tuberous sclerosis complex, or TSC, is a rare genetic disease that causes uncontrolled tumor growth across the face, body, and organs of affected individuals. Unfortunately, there is no cure for TSC.

South Florida's own Max Lucca—look at that sweet young face—was diagnosed with TSC when he was only 2 weeks old. Now he is 8, and Max Lucca has thrived because of the constant love and care provided by his parents, Vanessa and Max. Max Lucca enjoys baseball—as you can see—and hanging out with his best buddy.

So please think of this sweet young boy, Max Lucca, and others living with TSC on May 15 for TSC Global Awareness Day, and let's work together to find new treatments and a cure for this tragic disease.

GIVING HOPE TO DUCHENNE MUSCULAR DYSTROPHY PATIENTS

(Mr. PAULSEN asked and was given permission to address the House for 1 minute and to revise and extend his remarks.)

Mr. PAULSEN. Mr. Speaker, I rise today to speak about a genetic disorder, Duchenne muscular dystrophy, that has caused heartbreak for families across the country. The genetic mutation, found typically in boys, weakens muscles and leaves most sufferers confined to wheelchairs. Due to lung or heart failure, most afflicted with Duchenne pass away before their 25th birthday.