The Republican House has been in session for 110 days. I am deeply disappointed that the Republican leadership has chosen to ignore important bills that I have supported, such as H.R. 1664—the Investing in America: A Penny for Progress Act. H.R. 1664 would shore up the Highway Trust Fund through new Invest in America bonds and a modest increase in the fuel excise tax. Another bill, H.R. 1265—the Rebuilding America's Airport Infrastructure Act, would eliminate the cap on passenger facility charges, allowing local airports the ability to raise additional funds in order to build up more airport infrastructure such as new terminals and runways. H.R. 2510—the Water Quality Protection and Job Creation Act of 2017 would help bolster financing for new water and wastewater infrastructure projects all across the United States. These are real bills already introduced in Congress that we could consider today.

Mr. Speaker, we cannot afford to wait. There are a number of solutions already before us that this chamber could consider. It will take a display of political will by the Republican Party to consider these practical solutions to our nation's infrastructure woes. The American people are demanding that we act swiftly on these policies so that we can focus on what is most important—the efficient movement of the people, goods, and services which drive our economy forward. The time to act is now. I strongly urge my colleagues to support a comprehensive plan to shore up our nation's infrastructure.

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

The SPEAKER pro tempore (Mr. DUNN). Members are reminded to refrain from engaging in personalities toward the President.

PARALYSIS RESOURCE CENTER

The SPEAKER pro tempore. Under the Speaker's announced policy of January 3, 2017, the Chair recognizes the gentleman from Florida (Mr. DEUTCH) for 30 minutes.

Mr. DEUTCH. Mr. Speaker, I rise today to share the important work of the Paralysis Resource Center at the Christopher and Dana Reeve Foundation.

Christopher Reeve put a face on spinal cord injury and had an unrelenting drive to pursue the best research in the world. It was his vision, it was his passion, and it was his brilliance that attracted young scientists to take on the cause and advance the field of spinal cord research.

And while his wife, Dana, was universally known as the model for caregiving, her real legacy is the creation of the foundation's Paralysis Resource Center that has reached so many of those living with paralysis and their families with life-changing resources.

The President's budget eliminates all funding for the Paralysis Resource Center and suggests that the program is duplicative. Fortunately, my colleagues in the House appreciate the unique nature of the work that the PRC does.

The PRC consists of a variety of services, communities, and programs, including:

Information specialists trained to help anyone, from newly paralyzed individuals and their family members to persons who have lived with disabilities for quite some time, by providing individualized support and information;

The Peer and Family Support Program, which builds peer-to-peer connections to help individuals find support and resources among the communities that best understand the day-to-day realities and long-term challenges of individuals living with paralysis;

The Quality of Life Grants Program, which has awarded over 2,900 grants, totaling more than $22 million, to broaden the impact of nonprofits across the country that foster community engagement and involvement while promoting health and wellness for individuals living with paralysis; and

The Military and Veterans Program, the MVP, which supports the unique needs of current servicemembers and veterans regardless of when they served or how their injury was obtained.

I urge my colleagues to reach out to the Paralysis Resource Center to learn of the important work that they are doing for vulnerable constituents in their districts. If you do, you can see that the PRC is so much more than a line item in a budget. It is more than dollars and cents. It is more than just another program.

The value of these programs can only be appreciated when you learn the stories of the thousands of Americans that the PRC has helped with their services. Tonight, I would like to tell some of those stories. It is through these stories that we can see the amazing work of the Paralysis Resource Center, the lasting legacy of Christopher and Dana Reeve, and the bright future of so many who have been served through the PRC's programs.

You will notice that, throughout these stories, you hear the same names pop up again and again. That is because the PRC builds community and it builds connections.

When a spinal cord injury leaves Americans with no place else to turn, the PRC steps in to help guide the way; and, in return, many of those who have benefited from PRC turn back and look to help others with spinal cord injuries, to support their journey of recovery and rehabilitation.

Mr. Speaker, in order to move forward and to tell these stories, I would like to start by yielding to the gentleman from Rhode Island (Mr. LANGEVIN), my friend, who will share some thoughts with us now.

Mr. LANGEVIN. Mr. Speaker, I want to begin by thanking the gentleman from Florida (Mr. DEUTCH), my colleague, for yielding. It is an honor to join him this evening to discuss the cause and mission that is very close to my heart.

As many of my colleagues know, I was injured back in 1980, as a young police cadet, when I was accidentally shot and paralyzed in the police locker room when a police officer's gun accidentally discharged. So, the cause of finding a cure and better treatments for spinal cord injuries and supporting family members as caregivers of people with spinal cord injuries has, again, been a passion of mine and mission for many years.

In late 2001, it was my privilege to join Christopher and Dana Reeve, two very dear friends of mine whom I got to know very well over the years, in announcing the creation of the Paralysis Resource Center, or the PRC, which was officially established in 2002 as part of the Christopher and Dana Reeve Foundation. These were two remarkable individuals, as I said, I got to know and who became friends of mine over the years.

I can remember back when Christopher Reeve was paralyzed in his horse riding accident and how, instead of being quiet and introverted and just dealing with his recovery, he made it his mission to call attention to the cause of spinal cord injuries and the need for further research to be done and support services to be provided. He was an amazing inspiration to me because he gave voice and was a face of spinal cord injuries and the work that needed to be done to find a cure.

Because of his work and that of the Christopher and Dana Reeve Foundation, we are getting closer and closer every day to finding a cure for spinal cord injuries; and we are getting closer and closer every day to providing support for better quality of life for people with disabilities and paralysis and further support for the families of those who care for people with disabilities, especially those with spinal cord injuries.

In 2009, I was so proud to support the enactment of the Christopher and Dana Reeve Paralysis Act, which formally authorized some of the programs that have led to the success of the PRC. This center has grown to include a network of information specialists, a number of veterans programs, a Peer and Family Support Program, a substantial Quality of Life Grants Program, and a neurorecovery rehabilitation network.

It really is an inclusive center. It conveys the message that people dealing with spinal cord injuries or nerve-related conditions are not alone, that there are people and organizations around them who care and are willing to give the advice and counsel support that they need. So whether you are living with paralysis or you are the caregiver of a loved one who is paralyzed, the PRC provides resources and information to help serve the specific needs of those affected by spinal cord injuries.

We know there are more than 43 million caregivers in the U.S. providing support to loved ones with chronic disabling conditions on a variety of disabilities, whether it is a sick child or a sick parent or, in particular, those living with spinal cord injuries and paralysis.

Reports estimate that the annual economic value, if we had to put a price tag on this uncompensated care of family caregivers, to be somewhere approximately at the level of $470 billion annually—$470 billion annually, if we had to put a dollar figure on this uncompensated care. That is more than the total Medicaid spending was in 2013. That is when this study was done and those statistics were made public.

The benefits of family caregiving, of course, are plentiful and it is what families do for one another. If you have a sick child or a sick parent or a loved one, families so often step up to the plate and they provide that uncompensated care. Again, it is what families do. But, at the same time, it can also take an emotional, mental, and physical toll.

□ 2030

As a result, respite is the most frequently requested support service among family caregivers. Respite is that little bit of assistance where you have someone coming in to give the caregiver a break during the course of a week. Maybe it is an hour or two a day or a few hours during the course of a week that can make all the difference in allowing the caregiver themselves to go to the doctor or go do their grocery shopping, the things that are important to the rest of the family, or bringing a child to a baseball game or soccer practice; again, just those little everyday things that maybe we often take for granted that a family caregiver may not be able to do but for the fact that they have respite coming in. As a result, respite is, again, the most frequently requested support services among family caregivers.

Supported by the Christopher and Dana Reeve Foundation, the Lifespan Respite Care Program, which I helped to create back in 1996 with then-Congressman Mike Ferguson, a Republican from Pennsylvania, it was a bipartisan effort in creating this program, which operates now in 35 States and the District of Columbia, complements the PRC's efforts to help paralyzed individuals achieve a high quality of life by ensuring that caregivers who help them also receive relief and care, again, and support themselves.

Access to respite services has been shown to promote caregiver health and well-being, promote family stability, reduce the likelihood of abuse and neglect, and delay or even avoid the need for admission to costlier, long-term institutional settings, resulting in significant savings for the healthcare system and ultimately taxpayers, but ultimately helping the loved one who needs the care receive better care, because the loved one providing the care gets the support and the break that they need.

So beyond supporting access to respite for caregivers, we know the PRC's services and programs have touched the lives of people across the country, including those residing in the Ocean State, where I am from in Rhode Island.

For instance, Sarah Galli of Rhode Island spoke about her brother, Jeff, who was injured in 1998, and I would like to share her impactful words on the importance of the PRC now. She wrote:

''July 4, 1998, was the day that my family changed forever. My healthy, strong, athletic 17-year-old brother dove into the shallow end of a swimming pool and was rendered a high-level quadriplegic.

''Each Independence Day that followed has been a reminder of how quickly and irrevocably Jeff had his physical independence taken from him. It is also a reminder of how grateful I am for the support that has enveloped my family with care.

''I can't speak for my brother's experiences as a quad. That is his story to tell. But here is mine. In the years following his accident, I have counted on the Paralysis Resource Center and the Reeve Foundation for support. I have referred friends and strangers to them who require information and advocacy.

''And as a theater major in college, I started a student cabaret, Born for Broadway, that expanded an annual gala in the years since, raising awareness and funds for paralysis organizations, including Reeve. And I have also picked up the phone on several occasions to call the PRC and cry. Sometimes you just need to know that you are not alone.

''A friend of mine, Reeve supporter Jeff Ruben, wrote a song about a pal who lives with paralysis, using the lyrics, 'I believe in miracles.'

''I don't believe in miracles, but I do believe in Jeff. And I believe in Reeve. Do you?''

Mr. Speaker, we know the PRC's network of support remains strong to this day. When another Rhode Islander, 21-year-old Jack Calbi, was injured in a mountain bike accident this past April and left paralyzed from the chest down, the PRC immediately reached out to Jack's family to discuss the array of resources available.

Nearly 6 million Americans live with paralysis, depend on robust services and supports, respite for caregivers that love them, and a continued investment in the science and innovation that will lead to the treatments and cures of tomorrow.

The PRC has become a beacon of hope for so many. And as my dear friend Christopher Reeve said, ''Once you choose hope, anything is possible.''

Christopher and Dana Reeve were friends of mine. I miss them dearly. They truly did provide hope to me and millions of others, whether affected by paralysis or some other nerve disorder or disabilities in general, or the families that are affected by this condition as well. They made a difference, as has the PRC. I am grateful for their work and I am grateful for all the efforts to keep the PRC going strong so that they can continue to help those living with paralysis, their families, and for those who may be affected in the future.

And together I know that one day, because of all the efforts that are going on through amazing research right now, as we get closer step by step every day to finding a cure for spinal cord injuries, that families will know that they are not alone, that we are in this together, and that PRC and the Christopher and Dana Reeve Foundation are helping us to one day get one step closer to finding the cures and those treatments that we so desperately look for.

Mr. Speaker, I thank my colleague from Florida for raising attention to this issue and for yielding to me.

Mr. DEUTCH. Mr. Speaker, I thank my friend from Rhode Island, Representative LANGEVIN. I thank him for his leadership here in the House on these important issues.

But just as you spoke about your friends Christopher and Dana Reeve and talked specifically about how Christopher Reeve was not quiet and introverted, that he was an inspiration to you, I think it is fair to say, and I am comfortable actually speaking in this one instance on behalf of my colleagues here in the House in saying that just as Christopher Reeve was an incredible leader and an inspiration to you, so, too, Representative LANGEVIN, are you an inspiration to all of us in the House. I thank you for participating tonight.

There are so many other stories to tell. Representative LANGEVIN shared some. I would like to share just a few others.

One is the story of Eric LeGrand. On October 16, 2010, Rutgers University football star Eric LeGrand sustained a spinal cord injury at his C3 and C4 vertebrae.

After a kickoff in the fourth quarter against Army, Eric sprinted down the field and he made the tackle, but after the collision, he stayed down on the field at MetLife Stadium. Eric was paralyzed from the neck down, and doctors gave him a zero to 5 percent chance of regaining neurologic function.

From the moment he was injured, the Paralysis Resource Center was by Eric's side with a wealth of services and support. To conquer the immediate challenges of paralysis, his mother, Karen LeGrand, spoke regularly with the PRC's information specialists, who provided a roadmap of the resources that helped Eric transition to the Kessler Institute for Rehabilitation in West Orange, New Jersey. At that point, Eric was already exceeding expectations for his recovery and was weaned off of his ventilator and his feeding tube.

To help his family understand the long-term realities of a spinal cord injury, Eric and Karen were put in touch with Alan Brown, a dear friend of mine I will speak more about in a bit. Alan is a certified peer mentor and staff member of the Reeve Foundation, and he offered advice from his own personal experience living with quadriplegia for over 25 years.

Eric was also enrolled in the Reeve Foundation's NeuroRecovery Network, the NRN. The NRN is a growing network of rehabilitation centers that develop and provide cutting-edge therapies, like Locomotor Training.

Thanks to the NRN therapies and Eric's determination, he recovered the ability to move his neck and his shoulders. He has also improved trunk, cardiovascular, and pulmonary control.

His quality of life, well-being, and health have dramatically improved through the efforts of the PRC.

To pay forward his gratitude and help others, Eric founded Team LeGrand of the Christopher and Dana Reeve Foundation to speed the development of treatments for spinal cord injury. Since its inception in September 2013, Team LeGrand has raised close to $1 million.

Karen LeGrand was also inspired by her experience with the PRC, and became a certified peer mentor to offer guidance and support to parents who have children living with paralysis.

Elizabeth Forst is someone else whose story we should focus on. An avid traveler and diver, Elizabeth ''E.B.'' Forst's life changed in 2014 when a dive into a swimming pool resulted in a C4–C5 spinal cord injury.

In the aftermath of her injury, Elizabeth's family struggled with how to move forward and to ensure that Elizabeth could still live life to the fullest.

Her older brother, Tracy, reached out to the Reeve Foundation to tap into the constellation of programs and resources offered by the Paralysis Resource Center. He spoke with information specialists to get a better understanding of his sister's condition and how his family could help her adapt physically, emotionally, and socially to living with a spinal cord injury.

Tracy also regularly referred to the Reeve Foundation's website to educate himself on life after paralysis from both Elizabeth's perspective and his role as her family member and caregiver.

Thankful for the support she received from the Paralysis Resource Center, Elizabeth became a trained and certified peer mentor to help fellow paralysis community members find beauty and joy after a life-changing injury.

She has also resumed her passion for diving and traveling, as well as blogging for the Reeve Foundation to share her tips for traveling with a disability. E.B. has said she still has a lot of travel destinations on her list, and the PRC is surrounding her with support to make sure she can continue to travel the world.

Denna Laing is someone else we should think about tonight. On December 31, 2015, Denna Laing, a standout hockey player for the NWHL Boston Pride, was playing in the Women's Winter Classic at Gillette Stadium when she crashed headfirst into the boards and sustained a spinal cord injury at her C5 vertebra.

While she was in the hospital, the Reeve Foundation mobilized a network of support through the Paralysis Resource Center to help her family understand the short-term and the long-term challenges of living with paralysis.

They spoke with information specialists who provided tailored assistance on transitioning Denna from the hospital to acute care, and then adjusting to life back in her community.

Denna and her family connected with certified peer mentors like Alan Brown, who offered support and guidance based on their own personal experiences.

She is currently building up her strength and working to improve her health by participating in a comprehensive therapy regime at Journey Forward in Canton, Massachusetts, which is part of the Reeve Foundation's NeuroRecovery Network.

□ 2045

Denna recently tackled the Boston Marathon with help from former NHL player Bobby Carpenter, who pushed her racing wheelchair to the finish line.

Known for her vibrant personality, she credits the support she received from the Reeve Foundation and paralysis community as the reason that she keeps smiling. In her words: ''I know the best of my life has yet to come.''

J.D. Bruning and Kelly Lamb are from Arkansas. After graduating from the University of Arkansas, J.D. was enjoying the warm weather with his friends when they decided to take a float trip on a local river. Near the end of the float trip, J.D. dove into what he thought was deep water, but it was too shallow, and he sustained a high-level spinal cord injury.

Since that life-changing moment over 4 years ago, J.D. and his family have tapped into the wealth of resources provided by the Reeve Foundation Paralysis Resource Center to help J.D. live a full and active life in his community.

Given the scarcity of resources for the paralysis community in Arkansas, the PRC was critical in establishing the best possible care for J.D. and helping to navigate the path forward for his family.

His girlfriend, Kelly Lamb, was so inspired by the Reeve Foundation's programs and mission that she ran both the Chicago and New York City Marathons in 2016, under the Team Reeve banner, and raised over $20,000 to accelerate innovative research.

Kelly has cited the Paralysis Resource Center as a lifeline for thousands of families impacted by paralysis in the State of Arkansas and across the Nation.

Sabrina Cohen is from Florida. Born and raised in Miami Beach, Sabrina sustained a C3–5 spinal cord injury in 1992, when a teenager driver, who promised to give Sabrina a ride to a party, started drag racing and, unfortunately, slammed into a tree. She was 14 years old, and her life was changed forever.

When Sabrina was first injured, there was no centralized resource to learn how to live with paralysis or connect with fellow community members.

When the Reeve Foundation Paralysis Resource Center was founded in 2002, it became and since remains a lifesaving resource for 5.4 million Americans living with paralysis from spinal cord injury, ALS, MS, spina bifida, stroke, and cerebral palsy.

However, physical barriers still existed in Sabrina's community as there was no beach in the Miami Beach area that was accessible to wheelchair users. Through the Sabrina Cohen Foundation, she applied for a Reeve Foundation Quality of Life Grant to fund her dream of a fully accessible beach and playground for the disabled, including seniors, veterans, and children with special needs.

The Quality of Life Grants Program is funded through the Paralysis Resource Center and awards financial support to nonprofits that foster greater health, quality of life, community involvement, and independence for individuals living with paralysis and their caregivers.

She made the most of a Quality of Life Grant to establish Miami Beach's first-ever dedicated public beach with improved wheelchair access and outdoor adaptive fitness and watersport activities.

Finally, I would like to talk about two other people, both also from Florida, and both good friends of mine.

Alan Brown, I mentioned earlier. He is a resident of Aventura, Florida, who was injured in 1988, at the age of 20, when his neck was crushed by an ocean wave while on vacation in the Caribbean. The accident caused a C5–6 spinal cord injury and left Alan a quadriplegic with no movement below his chest.

For nearly 30 years since the accident, Alan and his family have, through the Alan T. Brown Foundation to Cure Paralysis, provided support, education, and advocacy for the paralysis community.

Initially, the focus for the foundation was on finding a cure for paralysis, a magic bullet that would help everyone living with paralysis to walk again. But after years in a wheelchair, and the difficult challenges to overall health and emotional well-being that paralysis presents, he and his family decided to overhaul the mission of the foundation.

Today, they focus on improving the lives of people living with paralysis through peer mentoring, outreach, and fundraising to help others with basic needs like wheelchairs and ongoing medical care. And now, Alan and his foundation have partnered with the Reeve Foundation to extend the reach and impact of his message to the paralysis community.

Alan travels the country on behalf of the Reeve Foundation to advocate, to increase awareness, and to make an impact on the quality of life of other people living with paralysis and their families.

He currently serves as manager of public impact at the Christopher & Dana Reeve Foundation, and in this role, he focuses on extending the reach of the foundation's impact on the greater public, creating new opportunities for partnership with peer organizations, increasing the visibility of PRC and other Reeve Foundation programs, bolstering fundraising efforts, and developing the national Peer & Family Support Program.

I have been with Alan and been inspired by him on a regular basis, close up, and the work that he does in the paralysis community and for those who are experiencing paralysis firsthand. The work that he does at those early moments is extraordinary.

Finally, Shawn Friedkin, who is from my community in south Florida. In March of 1992, Shawn was injured in a motor vehicle accident 2 weeks before his 27th birthday. He was married with one child, and he spent much of the early years following his injury working to increase medical research funding for spinal cord injuries.

For his efforts, he was invited to the White House to meet with Hillary Clinton, then-First Lady, as part of her healthcare initiative.

Shawn testified before Congress on the need for an increase in research funding. And based on his desire to help others in the community, Shawn left his career in finance, where he worked as a hedge fund manager, to create Stand Among Friends in 1997.

Stand Among Friends, based in Boca Raton, Florida, is a cross-disability advocacy organization. It is focused on helping people with disabilities find employment through vocational training and adaptive technology. It has provided services to more than 4,000 people and helped over 1,000 people find employment. Shawn has grown the services and mission to become one of the top disability service providers in the State of Florida.

And each year, Shawn and the Stand Among Friends organization hosts the ‘‘emb(race).’’ The emb(race) is a day of diversity awareness that includes a half marathon, a 5K, and a family fun walk; and it is, indeed, a way for all of us to embrace one another.

Shawn, as I said, is a dear friend and an inspiration in his community and to all who know him.

Mr. Speaker, in closing, I would like to thank all of those who make the Paralysis Resource Center at the Christopher & Dana Reeve Foundation what it is today.

As you can see from these stories, the PRC comes in at a family's most vulnerable moment. These are the most serious injuries and illnesses one can imagine. They are life-changing, and they come in with an immediate response to help guide families who are hurting to help them create a positive vision of a new life and build a community of support for Americans living with paralysis.

I would like to leave with the words of Christopher Reeve. He said: ‘‘I think a hero is an ordinary individual who finds strength to persevere and endure in spite of overwhelming obstacles.’’

Tonight, Mr. Speaker, I would urge my colleagues to join me in supporting the Paralysis Resource Center in their work to help empower more American heroes in the fight against spinal cord injuries.

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

LEAVE OF ABSENCE

By unanimous consent, leave of absence was granted to:

Mr. POE of Texas (at the request of Mr. McCARTHY) for today on account of personal reasons.

Mr. CUMMINGS (at the request of Ms. PELOSI) for today and the balance of the week.

Mrs. NAPOLITANO (at the request of Ms. PELOSI) for today and the balance of the week, on account of attending to her husband's health situation in California.

ADJOURNMENT

Mr. DEUTCH. Mr. Speaker, I move that the House do now adjourn.

The motion was agreed to; accordingly (at 8 o'clock and 54 minutes p.m.), under its previous order, the House adjourned until tomorrow, Tuesday, July 25, 2017, at 10 a.m. for morning-hour debate.

EXECUTIVE COMMUNICATIONS, ETC.

Under clause 2 of rule XIV, executive communications were taken from the Speaker's table and referred as follows:

2048. A letter from the Deputy Assistant Secretary, Office of Legislative Affairs, Department of the Treasury, transmitting the Report to Congress from the Chairman of the National Advisory Council on International Monetary and Financial Policies, pursuant to 22 U.S.C. 262r(a); Public Law 95-118, Sec. 1701(a) (as amended by Public Law 105-277, Sec. 583); (112 Stat. 2681-202); to the Committee on Financial Services.

2049. A letter from the Secretary, Department of Education, transmitting the Department's final rule — Teacher Preparation Issues [Docket ID: ED-2014-OPE-0057] (RIN: 1840-AD07) received July 19, 2017, pursuant to 5 U.S.C. 801(a)(1)(A); Public Law 104-121, Sec. 251; (110 Stat. 868); to the Committee on Education and the Workforce.

2050. A letter from the Deputy Assistant Secretary, Bureau of Legislative Affairs, Department of State, transmitting the Department's Country Reports on Terrorism 2016, pursuant to Sec. 140 of the Foreign Relations Authorization Act for FY 1988 and 1989, as Amended (22 U.S.C. 2656f); to the Committee on Foreign Affairs.

2051. A letter from the Architect of the Capitol, transmitting the semiannual report of disbursements for the operations of the Architect of the Capitol for the period of January 1, 2017, through June 30, 2017, pursuant to 2 U.S.C. 1868a(a); Public Law 113-76, div. I, title I, Sec. 1301(a); (128 Stat. 428) (H. Doc. No. 115—56); to the Committee on House Administration and ordered to be printed.

2052. A letter from the General Counsel, National Credit Union Administration, transmitting the Administration's final rule — Civil Monetary Penalty Inflation Adjustment (RIN: 3133-AE67) received July 19, 2017, pursuant to 5 U.S.C. 801(a)(1)(A); Public Law 104-121, Sec. 251; (110 Stat. 868); to the Committee on the Judiciary.

2053. A letter from the Management and Program Analyst, FAA, Department of Transportation, transmitting the Department's final rule — Airworthiness Directives; Rolls-Royce plc Turbofan Engines [Docket No.: FAA-2017-0187; Directorate Identifier 2017-NE-08-AD; Amendment 39-18893; AD 2017-10-19] (RIN: 2120-AA64) received July 19, 2017, pursuant to 5 U.S.C. 801(a)(1)(A); Public Law 104-121, Sec. 251; (110 Stat. 868); to the Committee on Transportation and Infrastructure.

2054. A letter from the Management and Program Analyst, FAA, Department of Transportation, transmitting the Department's final rule — Airworthiness Directives; Bombardier, Inc., Airplanes [Docket No.: FAA-2017-0125; Directorate Identifier 2016-NM-193-AD; Amendment 39-18946; AD 2017-14-02] (RIN: 2120-AA64) received July 19, 2017, pursuant to 5 U.S.C. 801(a)(1)(A); Public Law 104-121, Sec. 251; (110 Stat. 868); to the Committee on Transportation and Infrastructure.

2055. A letter from the Management and Program Analyst, FAA, Department of Transportation, transmitting the Department's final rule — Airworthiness Directives; Airbus Airplanes [Docket No.: FAA-2016-3984; Directorate Identifier 2014-NM-119-AD; Amendment 39-18945; AD 2017-14-01] (RIN: 2120-AA64) received July 19, 2017, pursuant to 5 U.S.C. 801(a)(1)(A); Public Law 104-121, Sec. 251; (110 Stat. 868); to the Committee on Transportation and Infrastructure.

2056. A letter from the Management and Program Analyst, FAA, Department of Transportation, transmitting the Department's final rule — Airworthiness Directives; Dassault Aviation Airplanes [Docket No.: FAA-2016-9504; Directorate Identifier 2016-NM-107-AD; Amendment 39-18932; AD 2017-12-02] (RIN: 2120-AA64) received July 19, 2017, pursuant to 5 U.S.C. 801(a)(1)(A); Public Law 104-121, Sec. 251; (110 Stat. 868); to the Committee on Transportation and Infrastructure.

2057. A letter from the Management and Program Analyst, FAA, Department of Transportation, transmitting the Department's final rule — Airworthiness Directives; The Boeing Company Airplanes [Docket No.: FAA-2016-9391; Directorate Identifier 2016-NM-129-AD; Amendment 39-18931; AD 2017-13-01] (RIN: 2120-AA64) received July 19, 2017, pursuant to 5 U.S.C. 801(a)(1)(A); Public Law 104-121, Sec. 251; (110 Stat. 868); to the Committee on Transportation and Infrastructure.

2058. A letter from the Management and Program Analyst, FAA, Department of Transportation, transmitting the Department's final rule — Airworthiness Directives; DG Flugzeugbau GmbH Gliders [Docket No.: FAA-2017-0343; Directorate Identifier 2017-CE-005-AD; Amendment 39-18936; AD 2017-13-06] (RIN: 2120-AA64) received July 19, 2017, pursuant to 5 U.S.C. 801(a)(1)(A); Public Law 104-121, Sec. 251; (110 Stat. 868); to the Committee on Transportation and Infrastructure.

2059. A letter from the Management and Program Analyst, FAA, Department of