

115TH CONGRESS
2D SESSION

S. 2652

AN ACT

To award a Congressional Gold Medal to Stephen Michael
Gleason.

1 *Be it enacted by the Senate and House of Representa-*
2 *tives of the United States of America in Congress assembled,*

1 **SECTION 1. SHORT TITLE.**

2 This Act may be cited as the “Stephen Michael Glea-
3 son Congressional Gold Medal Act”.

4 **SEC. 2. FINDINGS.**

5 The Congress finds the following:

6 (1) Stephen “Steve” Gleason was born March
7 19, 1977, in Spokane, Washington to Mike and Gail
8 Gleason.

9 (2) Steve attended Gonzaga Preparatory School
10 for high school where he excelled as both a football
11 and baseball player.

12 (3) In 1995, Steve enrolled at Washington
13 State University where he was a 2-sport athlete for
14 the baseball and football teams and helped the Cou-
15 gars football team advance to the 1997 Rose Bowl.

16 (4) In 2000, Steve signed a professional football
17 contract with the Indianapolis Colts of the National
18 Football League as an undrafted free agent but
19 later joined the New Orleans Saints in November of
20 that same season.

21 (5) Steve would go on to play 7 more seasons
22 as a member of the New Orleans Saints.

23 (6) Steve will always be remembered for his
24 blocked punt on September 25, 2006, against the
25 Atlanta Falcons, the night the Louisiana Superdome

1 reopened for the first time after Hurricane Katrina
2 in a game the Saints would win 23 to 3.

3 (7) In January, 2011 Steve was diagnosed with
4 amyotrophic lateral sclerosis or ALS, considered a
5 terminal neuro-muscular disease.

6 (8) Following his diagnosis, Steve, with the lov-
7 ing support of his wife, Michel, began a mission to
8 show that patients can not only live but thrive after
9 a diagnosis of ALS and established The Gleason Ini-
10 tiative Foundation also known simply as “Team
11 Gleason”.

12 (9) At the time of his diagnosis, however, Steve
13 said there will be “No White Flags”, which has be-
14 come the mantra of Team Gleason.

15 (10) The Gleason Initiative Foundation helps
16 provide individuals with neuromuscular diseases or
17 injuries with leading edge technology, equipment and
18 services, raises global awareness about ALS to find
19 solutions and an end to the disease, and has helped
20 hundreds of people with ALS experience life adven-
21 tures they never thought possible after their diag-
22 nosis.

23 (11) Steve’s story and mission have been told
24 by the NFL Network, ESPN, HBO, ABC, CBS,
25 CNN, and many local media outlets, as well as in a

1 2016 documentary titled “Gleason”, which was her-
2 alded at the Sundance Film Festival and premiered
3 across the country with Variety calling the produc-
4 tion “an emotional powerhouse”. The documentary
5 won several awards, including the 2016 Washington,
6 D.C. Area Film Critics Association Award for Best
7 Documentary.

8 (12) Steve was named one of two Sports
9 Illustrated’s Inspirations of the Year in 2014, has
10 been a keynote speaker for Microsoft and at two
11 United Nations sponsored Social Innovation Sum-
12 mits, and received the 2015 George S. Halas Cour-
13 age Award, given to a NFL player, coach or staff
14 member who overcomes the most adversity to suc-
15 ceed.

16 (13) Steve helped advocate for the Steve Glea-
17 son Act of 2015 (Public Law 114-40; 129 Stat.
18 441), and the Steve Gleason Enduring Voices Act of
19 2017, H.R. 2465, 115th Congress (2017), which
20 permanently ensures people living with diseases such
21 as ALS have access to speech generating devices re-
22 gardless of their setting, whether at home or a
23 healthcare institution.

24 (14) In 2014, Steve and Team Gleason hosted
25 a global summit to bring together researchers, pa-

1 tients, caregivers, and all ALS stakeholders to create
2 a plan to ultimately end ALS. That summit resulted
3 in the single largest coordinated and collaborative
4 ALS research project in the world, Answer ALS,
5 which brings together nearly two dozen research in-
6 stitutions, 1,000 patients and 20,000,000,000,000
7 data points that are important to the project and
8 that will define the unknown pathways that will lead
9 to treatments or finally a cure.

10 (15) In 2015, Steve and Microsoft worked to-
11 gether to create a method for people who are com-
12 pletely paralyzed to navigate their power wheelchairs
13 with their eyes. Today, Steve, Microsoft and all
14 wheelchair manufacturers are working collabora-
15 tively to make it widely available to all who need
16 this technology. In addition, Microsoft has also made
17 eye tracking technology part of all Windows 10
18 products across the globe.

19 (16) In 2011, 10 months after his diagnosis,
20 Steve and Michel made their most significant accom-
21 plishment, becoming parents to their son Rivers.

22 (17) Steve and Michel Gleason continue to fight
23 to find a solution for ALS so they can share many
24 years together and as parents to Rivers.

1 SEC. 3. CONGRESSIONAL GOLD MEDAL.

2 (a) AWARD AUTHORIZED.—The Speaker of the
3 House of Representatives and the President pro tempore
4 of the Senate shall make appropriate arrangements for the
5 award, on behalf of the Congress, of a single gold medal
6 of appropriate design to Stephen Michael Gleason.

7 (b) DESIGN AND STRIKING.—For the purposes of the
8 award referred to in subsection (a), the Secretary of the
9 Treasury (hereafter in this Act referred to as the “Sec-
10 retary”) shall strike the gold medal with suitable emblems,
11 devices, and inscriptions, to be determined by the Sec-
12 retary.

13 SEC. 4. DUPLICATE MEDALS.

14 Under such regulations as the Secretary may pre-
15 scribe, the Secretary may strike and sell duplicates in
16 bronze of the gold medal struck under section 3, at a price
17 sufficient to cover the costs of the medals, including labor,
18 materials, dies, use of machinery, and overhead expenses.

1 **SEC. 5. STATUS OF MEDALS.**

2 Medals struck pursuant to this Act are national med-
3 als for purposes of chapter 51 of title 31, United States
4 Code.

Passed the Senate June 14, 2018.

Attest:

Secretary.

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