

## SRES 161

A resolution recognizing June 2009 as the first National Hereditary Hemorrhagic Telangiectasia (HHT) month, established to increase awareness of HHT, which is a complex genetic blood vessel disorder that affects approximately 70,000 people in the United States.

**Congress:** 111 (2009–2011, Ended)

**Chamber:** Senate

**Policy Area:** Health

**Introduced:** May 21, 2009

**Current Status:** Submitted in the Senate, considered, and agreed to without amendment and with a preamble by Unanimous

**Latest Action:** Submitted in the Senate, considered, and agreed to without amendment and with a preamble by Unanimous Consent. (consideration: CR S5885; text as passed Senate: CR S5885; text of measure as introduced: CR S5877) (May 21, 2009)

**Official Text:** <https://www.congress.gov/bill/111th-congress/senate-resolution/161>

### Sponsor

**Name:** Sen. Johnson, Tim [D-SD]

**Party:** Democratic • **State:** SD • **Chamber:** Senate

### Cosponsors (7 total)

Cosponsor	Party / State	Role	Date Joined
Sen. Bennett, Robert F. [R-UT]	R · UT		Jun 23, 2009
Sen. Bingaman, Jeff [D-NM]	D · NM		Jun 23, 2009
Sen. Chambliss, Saxby [R-GA]	R · GA		Jun 23, 2009
Sen. Kerry, John F. [D-MA]	D · MA		Jun 23, 2009
Sen. Warner, Mark R. [D-VA]	D · VA		Jun 23, 2009
Sen. Isakson, Johnny [R-GA]	R · GA		Jul 14, 2009
Sen. Mikulski, Barbara A. [D-MD]	D · MD		Sep 8, 2009

### Committee Activity

*No committee referrals or activity are recorded for this bill.*

### Subjects & Policy Tags

**Policy Area:**

Health

## Related Bills

Bill	Relationship	Last Action
<a href="#">111 SRES 508</a>	Related bill	<b>May 26, 2010:</b> Resolution agreed to in Senate without amendment and with a preamble by Unanimous Consent. (consideration: CR S4469-4470; text as passed Senate: CR S4470)
<a href="#">111 HRES 536</a>	Related bill	<b>Jun 12, 2009:</b> Referred to the Subcommittee on Health.
<a href="#">111 SRES 141</a>	Related bill	<b>May 11, 2009:</b> Referred to the Committee on Health, Education, Labor, and Pensions. (text of measure as introduced: CR S5332)

## Summary (as of May 21, 2009)

Expresses support for the designation of June 2009 as National Hereditary Hemorrhagic Telangiectasia (HHT) month.

Recognizes: (1) the need to pursue research to find better treatments and a cure for HHT; (2) the HHT Foundation International as the only U.S. advocacy organization working to find a cure for HHT while saving the lives and improving the well-being of individuals and families affected; (3) the importance of comprehensive care centers in providing complete care and treatment for HHT patients; (4) that stroke, lung, and brain hemorrhages can be prevented through early diagnosis, screening, and treatment of HHT; (5) that severe hemorrhages in the nose and gastrointestinal tract can be controlled through intervention and that heart failure can be managed through proper diagnosis and treatments of HHT; and (6) that a leading medical and academic institution estimated that \$6.6 billion of one-time health care costs can be saved through aggressive management of HHT in the at-risk population.

Acknowledges the need to identify the approximately 90% of the HHT population that has not yet been diagnosed and that is at risk for death or disability due to sudden rupture of the blood vessels in major organs in the body.

## Actions Timeline

- **May 21, 2009:** Introduced in Senate
- **May 21, 2009:** Passed/agreed to in Senate: Submitted in the Senate, considered, and agreed to without amendment and with a preamble by Unanimous Consent.(consideration: CR S5885; text as passed Senate: CR S5885; text of measure as introduced: CR S5877)
- **May 21, 2009:** Submitted in the Senate, considered, and agreed to without amendment and with a preamble by Unanimous Consent. (consideration: CR S5885; text as passed Senate: CR S5885; text of measure as introduced: CR S5877)