

HR 3205

Patient Safety and Quality Improvement Act of 2005

Congress: 109 (2005–2007, Ended)

Chamber: House Policy Area: Health Introduced: Jul 12, 2005

Current Status: Placed on the Union Calendar, Calendar No. 117.

Latest Action: Placed on the Union Calendar, Calendar No. 117. (Jul 27, 2005) **Official Text:** https://www.congress.gov/bill/109th-congress/house-bill/3205

Sponsor

Name: Rep. Bilirakis, Michael [R-FL-9]

Party: Republican • State: FL • Chamber: House

Cosponsors (18 total)

Cosponsor	Party / State	Role	Date Joined
Rep. Brown, Sherrod [D-OH-13]	D · OH		Jul 12, 2005
Rep. Deal, Nathan [R-GA-10]	R · GA		Jul 12, 2005
Rep. Waxman, Henry A. [D-CA-30]	D · CA		Jul 12, 2005
Rep. Murphy, Tim [R-PA-18]	$R \cdot PA$		Jul 14, 2005
Rep. Baird, Brian [D-WA-3]	D · WA		Jul 18, 2005
Rep. Eshoo, Anna G. [D-CA-14]	D · CA		Jul 18, 2005
Rep. Whitfield, Ed [R-KY-1]	$R \cdot KY$		Jul 18, 2005
Rep. Bishop, Sanford D., Jr. [D-GA-2]	D · GA		Jul 21, 2005
Rep. Bono, Mary [R-CA-45]	$R \cdot CA$		Jul 22, 2005
Rep. Burgess, Michael C. [R-TX-26]	$R \cdot TX$		Jul 22, 2005
Rep. Norwood, Charles W. [R-GA-9]	$R \cdot GA$		Jul 22, 2005
Rep. Upton, Fred [R-MI-6]	$R \cdot MI$		Jul 22, 2005
Rep. Walden, Greg [R-OR-2]	$R \cdot OR$		Jul 22, 2005
Rep. Drake, Thelma D. [R-VA-2]	$R \cdot VA$		Jul 25, 2005
Rep. Emanuel, Rahm [D-IL-5]	D·IL		Jul 25, 2005
Rep. Green, Gene [D-TX-29]	$D \cdot TX$		Jul 25, 2005
Rep. Pickering, Charles W. "Chip" [R-MS-3]	$R \cdot MS$		Jul 25, 2005
Rep. McDermott, Jim [D-WA-7]	D · WA		Jul 26, 2005

Committee Activity

Committee	Chamber	Activity	Date
Energy and Commerce Committee	House	Reported by	Jul 14, 2005

Subjects & Policy Tags

Policy Area:

Health

Related Bills

Bill	Relationship	Last Action
109 S 544	Related bill	Aug 5, 2005: Referred to the Subcommittee on Health.

Patient Safety and Quality Improvement Act of 2005 - Amends the Public Health Service Act to designate patient safety work product as privileged and not subject to: (1) a subpoena or discovery in a civil, criminal, or administrative disciplinary proceeding against a provider; (2) disclosure under the Freedom of Information Act (FOIA) or a similar law; (3) admission as evidence in any civil, criminal, or administrative proceeding; or (4) admission in a professional disciplinary proceeding. Defines "patient safety work product" as any data, reports, records, memoranda, analysis, or written or oral statements which: (1) are assembled or developed by a provider for reporting to a patient safety organization (PSO); (2) are developed by a PSO for patient safety activities and which could result in improved patient safety or health care quality or outcomes; or (3) identify or constitute the deliberations or analysis of, or identify the fact of reporting pursuant to, a patient safety evaluation system.

Designates such work product as confidential, but permits: (1) disclosure for use in a criminal proceeding after a court determines that such work product contains evidence of a criminal act and is material to the proceeding and not reasonably available from any other source; (2) disclosure to the extent required to seek redress for violations through a civil action; and (3) disclosure if authorized by each provider identified in such work product.

Excludes certain disclosures from confidentiality protection, including: (1) disclosures to carry out patient safety activities; (2) disclosures of nonidentifiable work product; (3) disclosures otherwise allowed under Health Insurance Portability and Accountability Act of 1996 (HIPAA) confidentiality regulations to entities carrying out sanctioned research, evaluation, or demonstration products; (4) disclosures by a provider to the Food and Drug Administration (FDA); (5) voluntary disclosures by a provider to the provider's accrediting body; (6) disclosures necessary for business operations consistent with the goals of this Act; (7) disclosures to law enforcement authorities relating to the commission of a crime if the person making the disclosure reasonably believes the disclosure is necessary; and (8) disclosures that do not include materials that assess the quality of care of an identifiable provider or pertain to actions or failures to act by an identifiable provider by a person other than a PSO. Excludes from privilege any voluntary disclosure of nonidentifiable work product.

Instructs that privilege and confidentiality of work product continues for disclosures under this Act, with certain exceptions. Prevents a PSO from being compelled to disclose information collected or developed unless such information is identified, is not work product, and is not reasonably available from another source.

Prohibits an accrediting body from: (1) taking an accrediting action against a provider based on the provider's good faith participation in the collection, development, reporting, or maintenance of work product in accordance with this Act; and (2) requiring a provider to reveal communications with any PSO. Prevents a provider from taking an adverse employment action against an individual for the good faith reporting of information.

Sets forth penalties for the disclosure of identifiable work product in knowing or reckless violation of this Act. Allows an individual to bring a civil action to enjoin any act or practice that violates this Act.

Declares that this Act shall not: (1) limit greater privileges or confidentiality protections under federal, state, or local laws; (2) limit, alter, or affect the legal requirements pertaining to information that is not privileged or confidential; (3) alter or affect the implementation of HIPAA confidentiality regulations; (4) limit the authority of any provider, PSO, or other entity to require greater confidentiality or delegate authority to make disclosures; (5) preempt or affect any state law requiring a provider to report information that is not work product; (6) limit, alter, or affect any requirement for reporting safety information to the FDA; or (7) prohibit any person from conducting additional analysis.

Requires the Secretary to report to Congress on effective strategies for reducing medical errors and increasing patient safety.

Requires the Secretary to facilitate the creation of and maintain a network of patient safety databases that: (1) provide an interactive evidence-based management resource for providers, PSOs, and other entities; and (2) have the capacity to accept, aggregate across the network, and analyze voluntarily reported nonidentifiable work product. Requires the Secretary to assess the feasibility of providing for a single point of access to the network for qualified researchers for information aggregated across the network and, if feasible, provide for implementation. Allows the Secretary to determine common formats for reporting to the databases that are consistent with the administration simplification provisions of the Social Security Act. Requires that information reported to the databases be used to analyze national and regional statistics and be made available to the public.

Requires PSOs to submit an initial certification to the Secretary and renew their certification every three years. Sets forth criteria for certification.

Requires the Secretary to: (1) determine if the initial and subsequent certifications meet the requirements of this Act; and (2) compile and maintain a listing of certified PSOs. Makes provisions for work product if a PSO's certification is expired or revoked.

Allows the Secretary, acting through the Director of the Agency for Healthcare Research and Quality (AHRQ), to provide technical assistance to PSOs.

Requires the Comptroller General to study the effectiveness of this Act.

Actions Timeline

- Jul 27, 2005: Reported (Amended) by the Committee on Energy and Commerce. H. Rept. 109-197.
- Jul 27, 2005: Reported (Amended) by the Committee on Energy and Commerce. H. Rept. 109-197.
- Jul 27, 2005: Placed on the Union Calendar, Calendar No. 117.
- Jul 20, 2005: Committee Consideration and Mark-up Session Held.
- Jul 20, 2005: Ordered to be Reported (Amended) by Voice Vote.
- Jul 14, 2005: Referred to the Subcommittee on Health.
- Jul 14, 2005: Subcommittee Consideration and Mark-up Session Held.
- Jul 14, 2005: Forwarded by Subcommittee to Full Committee by Voice Vote.
- Jul 12, 2005: Introduced in House
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- Jul 12, 2005: Referred to the House Committee on Energy and Commerce.