

THE GENERAL ASSEMBLY OF PENNSYLVANIA

SENATE BILL

No. 761 Session of 2019

INTRODUCED BY GORDNER, JUNE 12, 2019

REFERRED TO BANKING AND INSURANCE, JUNE 12, 2019

AN ACT

1 Amending the act of March 20, 2002 (P.L.154, No.13), entitled  
 2 "An act reforming the law on medical professional liability;  
 3 providing for patient safety and reporting; establishing the  
 4 Patient Safety Authority and the Patient Safety Trust Fund;  
 5 abrogating regulations; providing for medical professional  
 6 liability informed consent, damages, expert qualifications,  
 7 limitations of actions and medical records; establishing the  
 8 Interbranch Commission on Venue; providing for medical  
 9 professional liability insurance; establishing the Medical  
 10 Care Availability and Reduction of Error Fund; providing for  
 11 medical professional liability claims; establishing the Joint  
 12 Underwriting Association; regulating medical professional  
 13 liability insurance; providing for medical licensure  
 14 regulation; providing for administration; imposing penalties;  
 15 and making repeals," in medical professional liability,  
 16 further providing for informed consent.

17 The General Assembly of the Commonwealth of Pennsylvania  
 18 hereby enacts as follows:

19 Section 1. Section 504 of the act of March 20, 2002  
 20 (P.L.154, No.13), known as the Medical Care Availability and  
 21 Reduction of Error (Mcare) Act, is amended to read:

22 Section 504. Informed consent.

23 (a) Duty of physicians.--Except in emergencies, a physician  
 24 owes a duty, which may be fulfilled by a physician or by a  
 25 qualified practitioner under subsection (b), to a patient to

1 obtain the informed consent of the patient or the patient's  
2 authorized representative prior to conducting the following  
3 procedures:

- 4 (1) Performing surgery, including the related  
5 administration of anesthesia.
- 6 (2) Administering radiation or chemotherapy.
- 7 (3) Administering a blood transfusion.
- 8 (4) Inserting a surgical device or appliance.
- 9 (5) Administering an experimental medication, using an  
10 experimental device or using an approved medication or device  
11 in an experimental manner.

12 (b) [Description of procedure] Requirements to obtain  
13 informed consent.--Consent is informed if the patient or the  
14 patient's authorized representative has been given a description  
15 of a procedure set forth in subsection (a) and the risks and  
16 alternatives that a reasonably prudent patient would require to  
17 make an informed decision as to that procedure. [The physician]  
18 A physician may delegate the task of obtaining the informed  
19 consent of the patient or the patient's authorized  
20 representative to a qualified practitioner for a procedure under  
21 subsection (a) performed by a physician or performed by a  
22 qualified practitioner. If claims for failure to obtain informed  
23 consent are alleged, the physician or qualified practitioner  
24 shall be entitled to present evidence of the description of that  
25 procedure and those risks and alternatives that a physician or  
26 qualified practitioner, acting in accordance with accepted  
27 medical standards of medical practice, would provide.

28 (b.1) Consent from another qualified practitioner.--A  
29 physician or qualified practitioner performing a procedure under  
30 subsection (a) may rely on information provided by another

1 qualified practitioner to obtain the informed consent of the  
2 patient or the patient's authorized representative.

3 (b.2) Evidence.--Information provided by another qualified  
4 practitioner under subsection (b.1) shall be competent evidence  
5 in a proceeding in which it is alleged that a physician or  
6 qualified practitioner performing a procedure under subsection  
7 (a) failed to obtain informed consent.

8 (b.3) Construction.--Nothing under this section shall be  
9 construed to require a physician to delegate the authority to  
10 obtain informed consent to a qualified practitioner.

11 (c) Expert testimony.--Expert testimony is required to  
12 determine whether the procedure constituted the type of  
13 procedure set forth in subsection (a) and to identify the risks  
14 of that procedure, the alternatives to that procedure and the  
15 risks of these alternatives.

16 (d) Liability.--

17 (1) [A physician is liable] Liability under this section  
18 for failure to obtain the informed consent only may be  
19 established if the patient proves that receiving such  
20 information would have been a substantial factor in the  
21 patient's decision whether to undergo a procedure set forth  
22 in subsection (a).

23 (2) [A physician may be held liable] Liability may be  
24 established under this section for failure to seek a  
25 patient's informed consent if the physician or qualified  
26 practitioner knowingly misrepresents to the patient [his or  
27 her] the professional credentials, training or experience[.]  
28 of the physician or qualified practitioner who performs the  
29 procedure.

30 (e) Human research exception.--The requirements under this

1 section shall be deemed satisfied if informed consent is  
2 obtained for human research conducted pursuant to approval by an  
3 institutional review board or similar entity in accordance with  
4 21 CFR Pt. 50 (relating to protection of human subjects), 45 CFR  
5 Pt. 46 (relating to protection of human subjects) and any other  
6 applicable Federal laws and regulations.

7 (f) Applicability--A physician or qualified practitioner  
8 performing a procedure under subsection (a) shall not be  
9 required to obtain a separate or new informed consent from the  
10 patient or the patient's authorized representative, provided  
11 that informed consent was already obtained by a physician or  
12 another qualified practitioner with respect to the procedure.

13 (g) Definition.--As used in this section, the term  
14 "qualified practitioner" means a:

15 (1) "Physician assistant" as defined in section 2 of the  
16 act of December 20, 1985 (P.L.457, No.112), known as the  
17 Medical Practice Act of 1985, or section 2 of the act of  
18 October 5, 1978 (P.L.1109, No.261), known as the Osteopathic  
19 Medical Practice Act;

20 (2) "Certified registered nurse practitioner" as defined  
21 in section 2(12) of the act of May 22, 1951 (P.L.317, No.69),  
22 known as The Professional Nursing Law;

23 (3) "Midwife or nurse-midwife" as defined in section 2  
24 of the Medical Practice Act of 1985; and

25 (4) Registered nurse under section 3 of The Professional  
26 Nursing Law who is authorized under the registered nurse's  
27 scope of practice to perform the procedure as delegated by  
28 the physician or a registered nurse authorized to administer  
29 anesthesia under 49 Pa. Code § 21.17 (relating to anesthesia)  
30 or a successor statute or regulation.

1 The term shall include another physician and a physician  
2 participating in a medical residency or fellowship training  
3 program. A qualified practitioner shall have knowledge of the  
4 patient's condition and the procedure enumerated under  
5 subsection (a) to be conducted on the patient and shall be  
6 acting under the supervision of, at the direction of, or in  
7 collaboration or cooperation with, the physician.

8       Section 2. The amendment of section 504 of the act shall  
9 apply to all pending litigation. The term "pending litigation"  
10 means any action in which a final order has not yet been entered  
11 prior to the effective date of this section.

12       Section 3. This act shall take effect immediately.