

# SENATE BILL 819

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By: **Senator Britt**

Introduced and read first time: February 16, 2007

Assigned to: Rules

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## A BILL ENTITLED

1 AN ACT concerning

2 **HIV and AIDS – Consent for Testing and Guidelines**

3 FOR the purpose of requiring a certain form of consent for HIV testing; authorizing  
4 substitute consent if an individual is unable to provide a certain form of consent  
5 for HIV testing; requiring the AIDS Administration to convene a certain  
6 workgroup to review certain Centers for Disease Control and Prevention  
7 recommendations relating to HIV and AIDS; requiring the AIDS  
8 Administration to report to the Governor and General Assembly on or before a  
9 certain date; defining certain terms; and generally relating to HIV and AIDS  
10 consent for testing and guidelines.

11 BY repealing and reenacting, with amendments,  
12 Article – Health – General  
13 Section 18–336  
14 Annotated Code of Maryland  
15 (2005 Replacement Volume and 2006 Supplement)

16 SECTION 1. BE IT ENACTED BY THE GENERAL ASSEMBLY OF  
17 MARYLAND, That the Laws of Maryland read as follows:

18 **Article – Health – General**

19 18–336.

20 (a) (1) In this section the following words have the meanings indicated.

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EXPLANATION: CAPITALS INDICATE MATTER ADDED TO EXISTING LAW.

[Brackets] indicate matter deleted from existing law.



1           (2) “HIV” means the human immunodeficiency virus that causes  
2 acquired immune deficiency syndrome.

3           (3) “Health care provider” means a physician, nurse, or designee of a  
4 health care facility.

5           (b) Except as provided in Title 11, Subtitle 1, Part II of the Criminal  
6 Procedure Article or § 18–338.3 of this subtitle, before obtaining a fluid or tissue  
7 sample from the body of an individual for the purpose of testing the fluid or tissue for  
8 the presence of HIV infection, a health care provider shall:

9           (1) Obtain [written informed] consent from the individual [on a  
10 uniform HIV informed consent form that the Department shall develop consistent with  
11 the requirements of the Department as established by regulations adopted by the  
12 Department]; and

13           (2) Provide the individual with pretest counseling, including:

14                   (i) Education about HIV infection and methods for preventing  
15 transmission;

16                   (ii) Information about a physician’s duty to warn; and

17                   (iii) Assistance in accessing health care available to an  
18 individual who tests positive for the HIV infection.

19           (c) Refusal to consent to the HIV antibody test or a positive test result may  
20 not be used as the sole basis by an institution or laboratory to deny services or  
21 treatment.

22           (d) If the individual is unable to give [informed] consent, substitute consent  
23 may be given under § 5–605 of this article.

24           (e) A physician or physician’s designee who obtains a positive result from an  
25 HIV antibody test conducted in accordance with the provisions of subsection (b) of this  
26 section shall:

27                   (1) Notify the individual from whom the fluid or tissue sample was  
28 obtained of the positive result;

29                   (2) Provide the individual with a copy of the Department’s publication  
30 describing available counseling services;

1           (3) Counsel the individual to inform all sexual and needle-sharing  
2 partners of the individual's positive HIV status;

3           (4) Offer to assist in notifying the individual's sexual and  
4 needle-sharing partners; and

5           (5) If necessary, take action appropriate to comply with § 18-337 of  
6 this subtitle.

7           [(f) The informed consent document shall be distinct and separate from all  
8 other consent forms.

9           (g) A patient identifying number obtained from an anonymous and  
10 confidential test site which is approved by the Department of Health and Mental  
11 Hygiene may be evidence of a patient's informed consent in lieu of a patient's  
12 signature.]

13           SECTION 2. AND BE IT FURTHER ENACTED, That:

14           (a) (1) The AIDS Administration shall convene a workgroup of  
15 stakeholders who serve the HIV and AIDS population to review Centers for Disease  
16 Control and Prevention guidelines relating to the HIV and AIDS epidemic.

17           (2) For the purpose of this section, "stakeholders" include:

18                   (i) Licensed physicians, hospitals, and other health care  
19 providers;

20                   (ii) Academic medical institutions; and

21                   (iii) HIV and AIDS advocacy organizations.

22           (b) The AIDS Administration shall report to the Governor and, in accordance  
23 with § 2-1246 of the State Government Article, the General Assembly on or before  
24 December 1, 2007, on changes made to HIV and AIDS related laws and regulations.

25           SECTION 3. AND BE IT FURTHER ENACTED, That this Act shall take effect  
26 June 1, 2007.