


Providing Newborn Screening Specimens for Research: Legal Issues Faced by State Health Departments Webinar Series

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States that store and provide residual dried blood spots (DBS) for secondary uses, or are considering storage and secondary uses, may face an array of legal issues. This webinar series is intended to cover these legal issues – and related ethical and policy issues – concerning secondary uses of DBS and associated data.

This webinar series is intended for public health attorneys, newborn screening programs and laboratories, institutional review boards, HIPAA privacy boards, privacy officers, researchers, and others who have interest in legal, and ethical and policy issues related to secondary uses of residual DBS.

Date	Topics	Registration Link
April 26, 2012 at 1:00pm EST	Introduction	Click here
May 31, 2012 at 1:00pm EST	Ownership and control of specimens	Click Here
June 28, 2012 at 1:00pm EST	Protection of privacy	Click Here
July 26, 2012 at 1:00pm EST	Protection of human research subjects	Click Here
August 30, 2012 at 1:00pm EST	Ownership and control of research results	Click Here
September 27, 2012 at 1:00pm EST	Summary of core issues	Click Here

Continuing education credits may be available, please contact ismotrich@acmg.net if interested in finding out more!



The Network
for Public Health Law



Jointly presented by the Newborn Screening Translational Network (NBSTRN), The Network for Public Health Law, the National Newborn Screening and Genetics Resource Center (NNSGRC), and the Association for Public Health Laboratories (APHL)