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Chamberlin, John

<http://hdl.handle.net/2027.42/78190>
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Biobanking: Honoring Donors’ Rights
Biobank Basics

• Collection of biological material and data
• Information available for research
• Have existed since the 1800s
• Registries and surveys prompted recent attention
Questions of Morality

• Consent procedures
• Confidentiality
• Research results
• Incidental findings
Consent Procedures

• Blanket consent
  • No restrictions on researchers
  • European preference and emerging trend
• Tailored consent
  • Choices about research types
  • American preference
Consent Procedures

- Blanket consent disadvantages
  - The more general, the less informed
  - Only broad descriptions about the purpose are available
Tailored consent advantages

- Types of research
- Uses of data
- International sharing
- Commercial partners
- Link data to personal and medical information
- Can re-contact donors
Research Results

• Normally published long after the study began
• Published in science journals
• Not always published
  • No value
  • Misleading information
Incidental Findings

• Individual research participant
• Potential health or reproductive importance
• Discovered in the course of conducting research
• Beyond the aims of study
Reduced level of social benefit in return for honoring rights of donors
Biobanking: Honoring Donors’ Rights

Questions and Conversation