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Institutional review boards, investigators and compliance officials have a new forum to review issues of importance in human subject regulations, courtesy of the HHS Office for Human Research Protections (OHRP). And the agency is soliciting ideas for future discussions.

On Sept. 7, OHRP hosted what it is calling its “inaugural workshop,” titled “Meeting New Challenges in Informed Consent in Clinical Research.” The workshop ran more than seven hours.

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