Office of Sponsor and Regulatory Oversight

The newly established Office of Sponsor and Regulatory Oversight (OSRO) ensures CCRs regulatory compliance with sponsor obligations for Investigational New Drugs (IND) and Investigational Device Exemptions (IDE), a critically important role for the CCR clinical research program and its investigators. In addition, this office provides analytic support, leads the pharmacovigilance program, monitors clinical trials, and serves as the subject matter experts regarding FDA regulations.

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- **Policies**
- **SOPs**
- **Forms and Instructions**

- **IND/IDE Filing Status**
- **OSRO/SROS Contact Information**
- **Seminar Series Materials**
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