

**Fundamentals of Oncology Treatment Clinical Trials**  
**October 24-25, 2006**  
**Lister Hill Center Auditorium**  
**First floor, Building 38A**  
**8600 Rockville Pike**  
**Bethesda, MD**

**Agenda**

**Tuesday, October 24, 2006**

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| Moderator:          | <b><i>Georgie Cusack, R.N., M.S., A.O.C.N.<sup>®</sup></i></b>   |
| 7:30 am – 8:00 am   | <b>Sign-in Day 1</b>   |
| 8:00 am – 8:10 am   | <b>Welcome and Introduction</b>  |
| 8:10 am – 9:00 am   | <b>Clinical Trials Update: Phase 0 Trials</b><br><b><i>James Doroshov, M.D., F.A.C.P.</i></b>  |
| 9:00 – 10:00 am     | <b>Protection of Human Subjects</b><br><b><i>Elizabeth Ness, R.N., M.S.</i></b>  |
| 10:00 am – 10:15 am | <b>BREAK</b>   |
| 10:15 am – 11:15 am | <b>Clinical Trial Design and Protocol Development</b><br><b><i>Elizabeth Ness, R.N., M.S.</i></b>                                      |
| 11:15 am – 12:00 pm | <b>Managing Data in Clinical Research</b><br><b><i>Allison Wise, B.S., R.H.I.A.</i></b>  |
| 12:00 pm – 1:00 pm  | <b>LUNCH</b> ( <i>On your own</i> )  |
| 1:00 pm – 1:45 pm   | <b>Documentation in Clinical Research</b><br><b><i>Susan Booher, R.N., M.S.</i></b>  |
| 1:45 pm – 2:30 pm   | <b>IND Clinical Trials and the Drug Development Process</b><br><b><i>Maureen Edgerly, R.N., M.A., O.C.N.<sup>®</sup>, C.C.R.C.</i></b> |
| 2:30 pm – 2:45 pm   | <b>BREAK</b>   |
| 2:45 pm – 3:30 pm   | <b>Roles &amp; Responsibilities: The Sponsor</b><br><b><i>Maureen Edgerly, R.N., M.A., O.C.N.<sup>®</sup>, C.C.R.C.</i></b>            |
| 3:30 pm – 4:15 pm   | <b>Roles &amp; Responsibilities: The Research Team</b><br><b><i>Rebecca Babb, R.N., B.S.N., O.C.N.<sup>®</sup></i></b>                 |
| 4:15 pm – 4:30 pm   | <b>Questions</b><br><b>Day 1 Evaluation</b>  |

**Wednesday, October 25, 2006**

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| Moderator:          | <b>Georgie Cusack, R.N., M.S., A.O.C.N.<sup>®</sup></b>   |
| 7:30 am – 8:00 am   | <b>Sign-in Day 2</b>  |
| 8:00 am – 9:00 am   | <b>Informed Consent Process</b><br><b>Georgie Cusack, R.N., M.S., A.O.C.N.<sup>®</sup></b>                |
| 9:00 am – 9:30 am   | <b>RECIST: Applying the Rules</b><br><b>Elizabeth Ness, R.N., M.S.</b>                                    |
| 9:30 am – 10:15 am  | BREAK   |
| 10:15 am – 11:45 am | <b>Adverse Events and IND Safety Reports</b><br><b>Elizabeth Ness, R.N., M.S.</b>                         |
| 11:45 am – 12:45 pm | LUNCH ( <i>On your own</i> )  |
| 12:45 pm – 1:15 pm  | <b>Regulatory Binder</b><br><b>Michelle Eugeni, R.N., B.A., B.S.N.</b>                                    |
| 1:15 pm – 2:15 pm   | <b>Monitoring of Sponsored Treatment Clinical Trials</b><br><b>Geoffrey D. Seidel, R.N., B.S.N., M.S.</b> |
| 2:15 pm – 2:30 pm   | BREAK   |
| 2:30 pm – 3:30 pm   | <b>Patient Recruitment</b><br><b>Susan McMullen, R.N., B.S.</b>   |
| 3:30 pm – 4:15 pm   | <b>Clinical Trial Jeopardy</b>  |
| 4:15 pm – 4:30 pm   | <b>Questions</b><br><b>Day 2 Evaluation</b><br><b>Overall Workshop Course Evaluation</b>                  |