NCIC Clinical Trials Group
2003 Expanded Fall Meeting of Committees
Clinical Research Associates Committee

Clinical Research Associates Committee Meeting

Delta Chelsea Hotel, Toronto, ON
Friday, October 17, 2003
8:00 am – 12:00 pm

Room: Churchill Ballroom

Chair: Kay Cranston

8:15 - 8:30 **Welcome and Steering Committee Report**
Kay Cranston, Chair CRA Committee

8:30 - 8:40 **CRA Disease Site Reps**
Phyllis Bettello, Chair CRA Disease Site Representatives

8:40 – 9:10 **Quality Assurance – Central Office**
Bev Koski, Quality Assurance Coordinator, NCIC Clinical Trials Group

9:10-9:30 **Quality Assurance – CRA Committee – Tools for Data Collection**
Nancy Page, Chair CRA Quality Assurance Subcommittee

9:30 – 9:50 **Ethics, Regulatory and Intergroup (ERI) Update—Central Office**
Melanie Walker, ERI Coordinator, NCIC Clinical Trials Group

9:50-10:05 **BREAK**

10:05-11:10 **CRA ERI Subcommittee Educational Session:**
The Process of Obtaining Informed Consent
CRA ERI Subcommittee

11:10-11:20 **CTSU Update**
Anita Nelson, Protocol Coordinator

11:20 – 11:45 **CTCAE v3.0 - Common Terminology Criteria for Adverse Events**
Monica Bacon, Study Coordinator, NCIC Clinical Trials Group

12:00 – 1:30 **CRA Disease Site & Information Fair / Data Management & Pharmacy Poster Session**
Buffet Lunch Provided – Mountbatten Salon B

1:30 – 4:30 **NCIC CTG Central Office Trial Workshops** *For rooms and schedule- see reverse*

MY.10…….. Nancy Paul
MA.27…….. Cathy Elliott
MA.21…….. Tina Liinamaa
MA.22…….. Paula Richardson
CO.17…….. Ann-Marie Sargeant
BR.19……..Nadine Magoski
BR.20…….. Marina Djurfeldt
AdEERS ……Jean Powers & Ann Setser
RECIST (Response Evaluation Criteria in Solid Tumours)..... Katherine Hann
CONSENT Form Development…… ERI