

August 10, 2007

Lynda Cedar  
Study Director  
Atlantic Life Sciences, Inc.  
2110, boulevard Decarie, Suite 102  
Montreal, Quebec. H4A-3J3

Dear Ms. Cedar,

**Re: Successful Completion of Study**

On behalf of the team at \_\_\_\_\_ kindly allow me to take this opportunity to thank you and your staff for all of your hard work and effort in helping \_\_\_\_\_ meet the deficiency response timelines of the above named project.

Needless to say, this was not your atypical "first project" as we were bound by very tight timelines from the beginning, but overall, the project was handled successfully and in a very efficient manner. The ability for all groups (Sponsor, Clinic, and bioanalytical site) to work together, under pressure, to meet the timelines stipulated by the regulatory agency was truly appreciated.

My overall impressions of the clinic activities carried out during my monitoring visit on June 4, 2007, and performed in accordance with GCP requirements, the protocol, pertinent SOPs, and federal law, were good. During my June 4<sup>th</sup> visit, I was very pleased to see that the observations made during our May-2007 Pre-study Clinic Site Audit were either revised or implemented. Of particular note were the revisions requested to your clinic SOPs and training records. The source documents reviewed were also well developed allowing sufficient detail to ensure that the study would be executed properly and always under Good Clinical Practices (GCPs).

Most sincerely,