Subject: Re: Biological Agents Registry Form: Pin
From: Christopher Pin <cpin@uwo.ca>
Date: 7/6/2012 5:29 PM
To: Jennifer Stanley <jstanle2@uwo.ca>

Hi Jennifer,

Sorry to not respond to this sooner. I will work on the corrections and send next week. HOWEVER...

I am a little surprised at why I should separate the LRTGT and my laboratory into two different documents. I oversee, and therefore am responsible, for both. The people in the LRTGT often work in my laboratory space using the same procedures and and space, and vice versa. As I am sure you are aware, the process involved to complete a BARF requires many hours of work. Since the committee is requesting that I put this additional time into completing a second document, I think it would be appropriate to provide a reasonable explanation as to why this is necessary. Is there some regulation that is involved? Are there rules that govern this process? I know that the committee's time is precious as is mine, so I think its fair to ask for a detailed explanation.

Thanks for your understanding.

Chris

On 6/26/12 7:06 PM, "Jennifer Stanley" <jstanle2@uwo.ca> wrote:

> re: Biological Agents Registry Form
>
> Hello there
>
> Your form was recently reviewed at the Biohazards Subcommittee meeting. Please address these issues and re-send.
>
> This researcher's personal research should be separated from research done in the Transgenic Facility, LRTGT (i.e. the work his/her grant does not fund). If Tamoxifen is being used in the Transgenic Facility, it should be used in accordance with the latest animal care protocol (to be stated in the summary). Section 5.0 should reflect E1A. The appendix includes some oncogenes. If these oncogenes are not being used this should be indicated. The Transgenic Facility (LRTGT) should complete a separate form.
>
> Regards
> Jennifer

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