

## PROTOCOLS FOR WESTERN IRB REVIEW CHECKLIST

<b>PI:</b>
<b>Title:</b>
<b>Sponsor:</b>
<b>Sponsor Protocol Number:</b>
<b>NUM_OPPHI:</b>

- MSC Submission Form (signed by the PI and the Dept. Director)
- WIRB\_Initial Review Submission Form
- FDA 1572 Form (Signed by Investigator)
- Research Protocol
- Informed Consents
  - English
  - Spanish
  - Information required about [ClinicalTrials.gov](http://ClinicalTrials.gov) (Informed Consents)
- Diaries and Questionnaires
  - English
  - Spanish
- Investigator's Brochure
- Log of Research Personnel (Including name, Institution of Affiliation, role, etc.)
- Evidence that the protocol was submitted to the Institutional Biosafety Committee (IBC) and CITI Program\_Certificate of Biosafety
- Evidence protocol processing with Office of Contracts [(Lcda. Rosa I. Martínez Addarich -Letter or Email). Recuerde que una vez tenga la aprobación del IRB que evalúa el protocolo debe

llevar copia de la carta a la Oficina de Contratos para continuar el proceso y presentarlo para la firma del Rector.]

Other:

Revised by:	
_____	_____
Name	Sign
Date: _____	