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| **Section I: Instructions** |
| **1.** Use this worksheet to document the information needed for OSF to request CRBO services. |
| **2.** Press the F1 key while in a field to show help and/or example text. |
| **Section II: General Information** |
| **1. Principal Investigator (PI) Full Name:**       |
| **2. PI Employer:**       |
| **3. Protocol Title:**       |
| **4. Protocol Number:**       |
| **5. NCT Number:**       |
| **6. IRB of Record:**       |
| **7. Study location(s), including all facilities and study related activities:**  |
|  | Location Name | Study Related Activity |  |
|  |  |  |  |
|  |
| **8. Study documents provided:** |
| [ ]  Final Protocol\* |
| [ ]  Draft Informed Consent Form (ICF)\* |
| [ ]  Draft Sponsor Budget\* |
| [ ]  Clinical Trial Agreement (CTA) Template or any Contractual/Grant Documents\* ***(NO PDF)*** |
| [ ]  Investigator’s Brochure/Device Manual\* |
| [ ]  Drug Study Documents: |
| [ ]  IND Exempt Determination |
| [ ]  FDA Approval Letter for IND |
| [ ]  Device Study Documents\*\*:  |
| [ ]  FDA Approval Letter for IDE |
| [ ]  CMS Approval Letter for IDEs (dated prior to 1/1/15) |
| [ ]  Other:  |
| **9. IND# / IDE#:**      -OR-[ ]  NA  |
| **10. Sponsor contact information:**  |
| Name & Title:       |
| Phone #:       |
| Email:       |
| **11. Any other important items PFS should know:**       |
| **12. Indicate who the payee will be:**       |
| **13. Indicate if negotiations should be delayed until site lead/MGR gives approval:**       |

\* – Required

\*\* – Investigator Initiated Studies – CA only