<table>
<thead>
<tr>
<th>No.</th>
<th>Question</th>
<th>Answer</th>
</tr>
</thead>
<tbody>
<tr>
<td>1.</td>
<td>How many years have you been a regulatory administrator/study coordinator?</td>
<td>[___________]</td>
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</tbody>
</table>
| 2.  | Of the multi-site, FDA-regulated studies you have coordinated, have any used a single IRB review process? | a. Yes  
b. Not yet and not planning → Skip to Q7  
c. Not yet but in the planning stages → End of participation |
|     | [DO NOT ASK: Pull from screening data.]                                  |                                                                        |
| 3.  | Now I will ask questions about your studies that used a single IRB process, including those that are only in the planning stages. Have any of these studies relied upon another IRB for ethics review? | a. Yes  
b. No  
c. Do not recall |
| 4.  | Has your institution's IRB ever served as the reviewing IRB for any of these studies? | a. Yes  
b. No  
c. Do not recall |
| 5.  | Were any these studies FDA-regulated?                                     | a. Yes  
b. No  
c. Do not recall |
| 6.  | In your role, how long have you been involved in the single IRB review process for multi-site studies? Please answer about all your studies—not just those that are FDA regulated. | a. Less than 1 year  
b. 1 to 2 years  
c. 3 to 4 years  
a. 5 or more years |
| 7.  | Can we re-contact you if we need to follow up on any of the information you provide during the interview? | a. Yes  
b. No |
| 8.  | Can we re-contact you to provide you with a summary of the study's findings? | a. Yes  
b. No |

Thank you. That concludes all my demographic questions. [Confirm interview scheduling]