<table>
<thead>
<tr>
<th>Question Text</th>
<th>Author Response</th>
<th>Custom Submission Question ID</th>
</tr>
</thead>
<tbody>
<tr>
<td>Please complete the following questions in accordance with the <a href="http://dx.doi.org/10.2106/JBJS.19.00981">International Committee of Medical Journal Editors’ recommendations on data sharing in clinical trials (guidelines and examples are available here)</a>.</td>
<td>yes</td>
<td>54</td>
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<tr>
<td>Will individual participant data be available (including data dictionaries)?</td>
<td>All of the individual participant data that underlie the results reported in this article and all collected data, after deidentification (text, figures, tables).</td>
<td>55</td>
</tr>
<tr>
<td>What data in particular will be shared? (Examples include all individual participant data after deidentification, only participant data that underlies the results, or not available.)</td>
<td>study protocol, ethical Votum, statistical Analysis plan, ICF, clinical study report, analytic code</td>
<td>56</td>
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<tr>
<td>What other documents will be available?</td>
<td>anyone who wishes to Access the data</td>
<td>58</td>
</tr>
<tr>
<td>When will data be available (start and end dates)?</td>
<td>immediately after publication, no end date</td>
<td>57</td>
</tr>
<tr>
<td>With whom? (Examples include anyone who wishes to access the data, researchers who provide a proposal, or not applicable.)</td>
<td>proposals may be submitted up to 36 months after publication. After 36 months data will be available in our University’s data warehouse but without Investigator support.</td>
<td>59</td>
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</tbody>
</table>