**Project Description**
Phase 2 – Electronic Messaging

Phase 1 - Develop a GS1 standard and/or guideline that would detail the best practice approach to implementation of GS1 standards for Electronic Communications in the pharmaceutical clinical trial supply chain. This could include EDI and/or EPCIS

**Deliverable/Objective**
Develop a GS1 standard and/or guideline that would detail the best practice approach to implementation of GS1 standards for Electronic Communications in the pharmaceutical clinical trial supply chain

**Completed last 30 days**
- Implementation Guideline passed eBallot
- Implementation Guideline Ratified

**Next steps**
- CT EDI SME is developing the mapping of the CT business messages with XML messages
- Once complete, a new Work Request will be submitted

**Risks and Issues**

**Project approved**
April ‘19
- Seated group

**Requirements**
October ‘19
- Com Rev & eBallot

**Draft standard**
February ‘20
- Com Rev & eBallot
- IP review
- BCS and MB

**Publication**
* February ‘20
- GS1 website

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<table>
<thead>
<tr>
<th>Company participation</th>
<th>Stakeholders</th>
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<tbody>
<tr>
<td><strong>Actual roster</strong></td>
<td><strong>LT Sponsor</strong> Marianne Timmons</td>
</tr>
<tr>
<td>Manufacturer</td>
<td>15</td>
</tr>
<tr>
<td>Hospital</td>
<td>4</td>
</tr>
<tr>
<td>MO's</td>
<td>18</td>
</tr>
<tr>
<td>Other</td>
<td>14</td>
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<td><strong>Required roster</strong></td>
<td><strong>Chairs</strong> Hans von Steiger – Pfizer, Sylvain Alberola – SANOFI / Pierre Fernandez Barbera – SANOFI / Olivia Chauvel - CH Victor Dupouy</td>
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<tr>
<td><strong>Minimum votes</strong></td>
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</tr>
<tr>
<td>Manufacturer</td>
<td>4</td>
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<tr>
<td>Hospital</td>
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<tr>
<td>MO's</td>
<td>4</td>
</tr>
<tr>
<td>Other</td>
<td>2</td>
</tr>
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**Legend**
- G: On schedule
- Y: Minor Risk/≈10% behind schedule
- R: Significant risk/10%+ behind schedule
- C: Completes

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