## Business purpose
- This group will develop the appropriate solution required by the European Commission to develop a Master UDI-DI for implementation of a new level of identification of eyewear products and other devices as part of the UDI requirements based on the European Medical Device Regulation (MDR).

## Key issues and risks
- EU COM delegated act has not been finalised. The team is moving forward with the current understanding of the potential language in the delegated act which could impact the draft GSCN.
- If Spectacles are required in the GSCN this could delay completion date of Q1 2024

## Accomplishments this period
- Conducted a Poll to choose one MUDI Option for Made to Order contacts
- Drafted first GSCN
- On the 16 November call it was raised that EUCOM could potentially include Spectacles into the Regulation

## Upcoming period activities
- Continue to work on GSCN and work to make it flexible enough that spectacles can be incorporated if needed
# New EU requirements for medical devices identification MSWG

## Overall Status

<table>
<thead>
<tr>
<th>#</th>
<th>Standard/guideline Milestones</th>
<th>Due Date</th>
<th>Notes</th>
<th>#</th>
<th>Collateral milestones</th>
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<td>1</td>
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## Stakeholders

- **Sponsor**: Marianne Timmons
- **SDL**: Greg Rowe
- **CE/Sol Liaison**: Geraldine Lissalde Bonnet
- **SME**: Neil Piper
- **AG Liaison**: John Terwilliger - Abbott
- **Chairs**: Scott Durland - J&J, Lionel Tussau - atrify, Sylvia Reingardt – GS1 Germany

## Project Scope, Resources and Timeline Change Management

- IESC approval date: 9/21

## Change Management

<table>
<thead>
<tr>
<th>New risks identified</th>
<th>Yes/no</th>
<th>Change request included</th>
<th>Yes/no</th>
<th>Legend</th>
<th>Complete</th>
<th>Not Started</th>
<th>On target</th>
<th>Risk of being late</th>
<th>Change needed (not recoverable)</th>
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