# MedDRA & WHODrug User Group Meeting

**February 6, 2018 – Novotel Bengaluru Techpark Hotel, Bangalore, India**

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<thead>
<tr>
<th>Time</th>
<th>Session</th>
<th>Speakers</th>
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<tr>
<td>8:00-9:00</td>
<td>Registration</td>
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<tr>
<td>9:00-9:30</td>
<td>Opening &amp; introduction</td>
<td>Damon Fahimi, UMC and Pat Revelle, MedDRA MSSO</td>
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<td>9:30-10:00</td>
<td>Keynote: Coding – What’s the point?</td>
<td>Birgitta Grundmark, UMC</td>
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<td>10:00-11:15</td>
<td>Report from UMC</td>
<td>Anna Mattsson, Damon Fahimi and Malin Fladvad, UMC</td>
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<td>11:15-11:45</td>
<td>Coffee &amp; tea break</td>
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<td>11:45-13:00</td>
<td>Report from MedDRA MSSO</td>
<td>Pat Revelle, MedDRA MSSO</td>
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<td>13:00-14:00</td>
<td>Lunch</td>
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<td>14:00-15:30</td>
<td>Application of MedDRA and WHODrug to a clinical trial – an interactive session</td>
<td>Anna Baumgarten, UMC and Jane Knight, MedDRA MSSO</td>
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<td>15:30-16:00</td>
<td>Coffee &amp; tea break</td>
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<td>16:00-16:25</td>
<td>Implementation of the WHODrug B3-format</td>
<td>Dhanya Poulose, IQVIA</td>
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<td>16:25-16:50</td>
<td>Overview and importance of common terminology criteria for adverse events (CTCAE) in oncology trials</td>
<td>Amreen Taj and Rajiv Rai, Chiltern</td>
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<td>16:50-18:05</td>
<td>MedDRA &amp; WHODrug Panel discussion</td>
<td>Moderator: Shawn Davis Raj, eClinical Solutions India</td>
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<td>Panelists: Amit Kulkarni, Novartis Healthcare</td>
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<td>Manoj Shanmugasundaram, IQVIA</td>
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<td>Ranganath Nagarjuna, Chiltern</td>
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<td>Shubhadeep Sinha, Hetero Drugs</td>
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<td>18:05-18:30</td>
<td>Open Q&amp;A session, wrap-up and closing remarks</td>
<td>All presenters</td>
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<td>18:30-19:15</td>
<td>Drink reception</td>
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<td>19:15</td>
<td>Networking dinner</td>
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This event is co-hosted by the MedDRA MSSO and the Uppsala Monitoring Centre (UMC).

*For any questions, please contact: [WHODrug@who-umc.org](mailto:WHODrug@who-umc.org) or [mssohelp@meddra.org](mailto:mssohelp@meddra.org)