What is your primary role at your organization?

- Medical Coder: 13%
- Manager, Medical Coding: 55%
- Medical Reviewer: 0%
- Biostatistician: 0%
- System Developer: 3%
- Data Manager: 11%
- Other: 18%
What function(s) are responsible for MedDRA coding at your organization?

- Central Coding Group Codes both Pharmacovigilance and Clinical Trial data: 36%
- Separate Central Coding Group; one codes Pharmacovigilance, and one codes Clinical Trial data: 31%
- Decentralized coding for Pharmacovigilance, Centralized for Clinical Trials: 6%
- Centralized for Pharmacovigilance, Decentralized for Clinical Trials: 3%
- Medical Coding Activities Fully outsourced: 6%
- I do all of the coding: 6%
- Other: 14%
Does your organization utilize terminologies other than MedDRA for coding?

- ICD-9: 8%
- ICD-10: 11%
- SNOMED: 8%
- Other: 0%
- No Way! MedDRA is the bomb!: 74%
Has your organization participated in any coding experiments using machine learning or artificial intelligence?

- 4: Robots taking over ... ackkkk!
- 7: Yes
- 26: No
- 2: Don't Know
Any other thoughts on this MedDRA User Group Meeting or future user group discussion topics that you'd like to share?

- Electronic medical records-status
- its really helpful and very well organized. thank you for hosting such very informative meetings.
- Efficiency in data review
- Good meeting. The AI discussion / presentations was very interesting.
- More Update and follow up for machine learning
- Is the location of the Spring 2019 meeting already decided?
- Something on signal detection
- What do organizations mean when data is versioned to the next MedDRA version?
- MedDRA and SNOMED mapping progress status
Any other thoughts on this MedDRA User Group Meeting or future user group discussion topics that you'd like to share?

Updates on the “requirements” in the FDA guidance docs as to what truly is required

Do any of your companies use vendors who manage non-SMQ Baskets?

Can the slides presented today be emailed to the attendees?

How are those in the device space using MedDRA

Customized MedDRA query

From MSSO- what goes into denying a change request

Would like to know of the companies attending the actual percentage that already code to WHODrug primary ATC based on indication.

How to handle compound terms for clinical trials when they are able to be added via a change request.

Can we have a list of participants & the company they are from?
Any other thoughts on this MedDRA User Group Meeting or future user group discussion topics that you'd like to share?

- Have the MSSO and UMC host an annual conference.
- Excellent meeting, topics, discussion, venue, food & preparation. Thank you!
Roche

Genentech

Thank You!