# RAPID GUDID Workgroup

9/14/16

# RAPID Progress

- Principle: Capturing DI at point of use in EHRs is future goal but registries and other downstream users can use it NOW.
- Public availability of GUDID will allow:
  - Model-specific outcomes searches
  - Lower cost for device data entry
  - Auto-population and improved accuracy of device data in registries
- GUDID WG Team: Manufacturers, Registry Owners, Vendors, FDA, Clinicians
- Major Learnings:
  - Understanding how data is used is crucial to development and improvement in quality of GUDID
  - Common issues in UDI adoption across data sources & device types
  - GUDID data ***CAN*** be integrated to improve registry data

# RAPID GUDID WG Accomplishments

- Identification of Core GUDID data elements for use by registries

| | | |
|---|---|---|
| Primary Device Identifier | GMDN term | Clinically Relevant Size Type |
| Secondary Device Identifier | SNOMED CD/Term | Clinically Relevant Size Value |
| Company Name | Device Description | Clinically Relevant Size Unit of Measure |
| Brand | Version or Model | Catalog Number |

- Understand importance of Global Medical Device Nomenclature (GMDN) to meet peripheral vascular device categorization goals
- Evaluate gaps in UDI data for registry purposes

# RAPID GUDID Outcomes and Next Steps

- Incorporated DI data into Vascular Quality Initiative Registry
- Plans for collecting DI data in EHR for auto-populating into VQI registry
- Identification of need for MDEpiNet and/or Learning UDI Community Workgroups on cross-pilot issues:
  - Additional UDI Data (AUDI)
  - Clinically Relevant Size
  - Catalog Number
  - GMDN/SNOMED

# Contact Information


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