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| **Dataset** | **Priority** | **Status** | **Next step** |
| CAERS | ??? | Initial schemas have been created and dataset has been loaded into elastic search | FDA to determine if dataset has changed and if there are non-technical hurdles to overcome |
| Additional recall data elements (device only) | 1 | CDRH supplied ppt slides to Iodine during Oct. 2014 visit | Iodine to review dataset in detail  FDA to schedule follow up meeting with business & technical owners to discuss detail requirements |
| Registration & Listing | 2 | - | Same as above |
| Classification | 3 | - | Same as above |
| CDRH Inspections | 4 | - | Same as above |
| 510K | 5 | Iodine team has reviewed and vetted the dataset | Iodine to determine compatibility with existing MAUDE datasets |
| PMA | 6 | - | TBD |
| MAUDE (hierarchy) | 7 | Base MAUDE launched in Aug. 2014 | FDA to request more information about this request via email |
| Post-approval studies | 8 | - | TBD |
| 522 Postmarket surveillance studies | 9 | - | TBD |

Are we missing missing any datasets that we should explore for this first investigational phase?