

| | Certification Bodies | | | Accreditation bodies | Authority (MOH) |
|--|---|---|-------------------------------|--|------------------------------------|
| | Notified Body | Registrar | Auditing Organization | | |
| What they do | QMS Audits Product testing CE Marking | QMS Audits Product testing | QMS Audits Product testing | Accredit Certification Bodies | Oversee Certification Bodies |
| Issue Device Registration Certificate? | Yes, For CE only | No. This is done by Health Canada | No | No | Yes, Except EU |
| Countries | EU | Canda | Five Countries | N/A | Local Market |
| Conduct MDSAP Audits | No | No | Yes | No | No |
| Examples | TUV-BSI - Intertek-DEKRA-SGS | | | ANSI National Accreditation Board (ANAB) | |