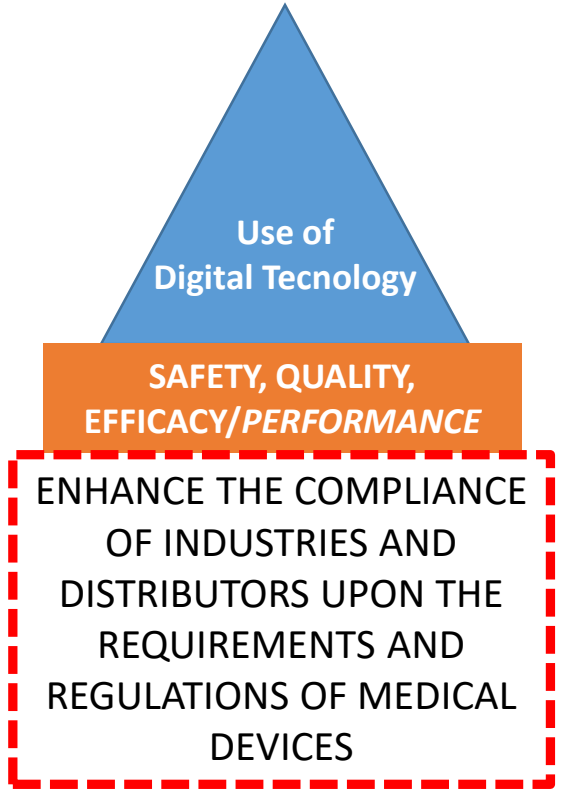
KEMENTERIAN KESEHATAN REPUBLIK INDONESIA

Implementation of Medical Device Control During The New Normal Era

GERMAS Gerakan Masyarakat Hidup Sehat

***Pre market Control* in the New Normal Era:**

- Optimize the use of digital information technology in the process of licensing (facility and product license).
- Focus on new medical device innovation.
- Adjustment the requirement in the new normal condition (*clinical evaluation*, Certificate of Free Sales, Letter of Authorization and supported documents).
- Easiness and flexibility of consultation/assistance through virtual system.



***Post market Control* in the New Normal Era :**

- Conducting remote audit to production and distribution facilities.
- The use of information and technology to conduct Monitoring/vigilance.
- Optimize the use of electronic based reporting (*e-report* and *e-watch*)
- Conducting medical device product Sampling through the online courier services.