Especially in medical engineering and diagnostics the regulatory requirements increased. For that reason we expanded our product portfolio for technical approval and the preparation of the necessary documents.

- Risk analyses according to DIN EN ISO 14971:2012
- Laser safety according to DIN EN 60825
- Electrical safety according to DIN EN 60601
- Premeasurement for EMC and ESD

Documentation and monitoring according to the requirements of specific medical approvals:

- CEmed annex II, section 3 of the directive 93/42/EWG
- IVD directive 98/79/EC
- FDA