The medical devices act (MDA) says that – in order to protect the patient – the use of medical devices has to be documented persistently. If you employ MedITEX MaSter (Material and Sterile Goods Management), there is a high-performance software to take over the annoying liabilities of the manual process documentation. And in addition you will always have the overview of your medical equipment stock.

**Your application for MDA – direct interface to the IVF EMR MedITEX IVF**

Thus at all times a medical practice has to provide evidence of all data referring to a patient including appointments, suppliers, manufacturers and products to the point of the exact batch number of the used preparation. Besides security data sheets and operating manuals have to be immediately available for each product. From the fertility database management system MedITEX IVF the patients can be opened easily in MedITEX MaSter and the documentation can start immediately. All relevant patient data will be overtaken from MedITEX IVF.

More information ...