

Home-Monitoring in ICD patients: MONITOR-ICD

Summary:

With the medical benefit of implantable cardioverter-defibrillators (ICD) for the reduction of sudden cardiac death being proven, the cost-effectiveness of the therapy is playing a role due to limited resources in the health system. With modern ICDs remote monitoring (home-monitoring, HM) is possible. Given these premises, the aim of the monitor-ICD study is to investigate whether the follow-up costs of the home monitoring group are lower than those of the conventional follow-up. Furthermore, we aim to analyse whether the monitor-ICD patients have a clinical relevant benefit due to a faster medical treatment and if subsequent events and clusters of events can be prevented in this group.

MONITOR-ICD is a multi centre randomized controlled trial. With a recruitment period of 24 months, 416 patients are planned to be included in the study, either getting an ICD with home-monitoring or an ICD with conventional follow-up. Follow-ups will take place at 1, 3, 6, 12, 18 and 24 months after study start. For the home-monitoring patients, follow-ups at 3, 6 and 18 months can be discarded due to online retrieval of the data. Besides the health economic and clinical endpoints, overall and disease specific quality of life will be measured.

Principal investigator:

Zabel, MD, Universität Göttingen

Willich, MD, MPH, MBA

Project coordinator:

Müller-Riemenschneider, MD, MSc

Reinhold, MPH

Research associate:

Data management:

Stasun

Project coordination office:

Wagner

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