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Standardized Evaluation of Subcutaneous Glucose Monitoring Systems Under Routine Environmental Conditions

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Summary

The purpose of this study is to assess accuracy and reliability of the SPIDIMAN CGM sensor system with regard to values as measured by Super GL and compare these results with similar evaluations of the Medtronic MiniMed 640G system, the Abbott FreeStyle Libre Flash Sensor and the DexCom G4 Platinum Sensor in patients with type 1 Diabetes.

Description

The study is a single-center open-label study in patients with type 1 diabetes treated with continuous subcutaneous insulin infusion (CSI) or multiple daily injections (MDI) therapy. The study will include a total of 12 subjects.

Study Design

Intervention Model: Single Group Assignment, Masking: Open Label, Primary Purpose: Basic Science

Conditions

Diabetes Mellitus, Type 1

Intervention

Glucose Sensor

Status

Active, not recruiting

Source

Medical University of Graz

Results (where available)

[View Results](#)

Links

- Source: <http://clinicaltrials.gov/show/NCT02614768>
- ClinicalTrials.gov processed this data on November 25, 2015

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