

## SPECIAL 510(K): DEVICE MODIFICATION OIR DECISION SUMMARY

**510(k) Number:** k181273

This 510(k) submission contains information/data on modifications made to the applicant's own class II or class I devices requiring 510(k). The following items are present and acceptable:

---

The name and 510(k) number of the applicant's previously cleared device. CareSens N Premier Blood Glucose Monitoring System, and CareSens N Premier BT Blood Glucose Monitoring System (k170614) (For a preamendments device, a statement to this effect has been provided.)

1. Applicant's statement that the **INDICATION/INTENDED USE** of the modified device as described in its labeling **HAS NOT CHANGED** along with the proposed labeling which includes instructions for use, package labeling, and, if available, advertisements or promotional materials (labeling changes are permitted as long as they do not affect the intended use).
2. A description of the device **MODIFICATION(S)**, including clearly labeled diagrams, engineering drawings, photographs, user's and/or service manuals in sufficient detail to demonstrate that the **FUNDAMENTAL SCIENTIFIC TECHNOLOGY** of the modified device **has not changed**.

### **This change was for:**

The CareSens N Premier BT Blood Glucose Monitoring System was modified as follows to create the GLUCOCARD Shine Connex Blood Glucose Monitoring System:

- Change in meter material, from Polycarbonate to Acryl.
- Change in LCD display from a negative to a positive display.
- Change in data transfer port from a phone jack to a Micro USB port.

The CareSens N Premier Blood Glucose Monitoring System was modified as follows to create the GLUCOCARD Shine Express Blood Glucose Monitoring System:

- Addition of a talking feature.
- Change in meter material, from Polycarbonate to Acryl.
- Change in LCD display from a negative to a positive display.
- Change in data transfer port from a phone jack to a Micro USB port.
- Change in Power source from two 3.0V Lithium batteries (CR2032) to two 1.5 V AAA alkaline batteries.

3. **Comparison Information** (similarities and differences) to applicant's legally marketed predicate device including, labeling, intended use, physical characteristics, and specifications.
4. A **Design Control Activities Summary** which includes:
  - a) Identification of Risk Analysis method(s) used to assess the impact of the modification on the device and its components, and the results of the analysis
  - b) Based on the Risk Analysis, an identification of the verification and/or validation activities required, including methods or tests used and acceptance criteria to be applied

The labeling for this modified subject device has been reviewed to verify that the indication/intended use for the device is unaffected by the modification. In addition, the applicant's description of the

particular modification(s) and the comparative information between the modified and unmodified devices demonstrate that the fundamental scientific technology has not changed. The applicant has provided the design control information as specified in The New 510(k) Paradigm and on this basis, I recommend the device be determined substantially equivalent to the previously cleared (or their preamendment) device.

GLUCOCARD Shine Connex Blood Glucose Monitoring System and GLUCOCARD Shine Express Blood Glucose Monitoring System are intended for single patient home use. Disinfection efficacy studies were performed on the materials comprising the meter and lancing device by an outside commercial laboratory testing services demonstrating complete inactivation of hepatitis B Virus (HBV) with Clorox Germicidal Wipe (EPA Reg. No: 67619-12). The sponsor also conducted robustness studies and demonstrated that there was no change in performance or in the external materials of the meter after 260 cleaning and disinfection cycles representing 5 years of single patient use. Labeling was reviewed for adequate instructions for the validated cleaning and disinfection procedures and found to be acceptable.