

SPECIAL 510(k): Device Modification
ODE Review Memorandum

To: THE FILE

RE: DOCUMENT NUMBER k060256

This 510(k) submission contains information/data on modifications made to the SUBMITTER'S own Class II, Class III or Reserved Class I devices requiring 510(k). The following items are present and acceptable (delete/add items as necessary) for the UNICEL DXC 600I SYNCHRON ACCESS CLINICAL SYSTEM:

1. The name and 510(k) number of BECKMAN COULTER, INC.'s previously cleared devices, Synchron LXi 725 and Unicel DxC 600 and 800 clinical systems, cleared under k023049 and k042291 respectively. (For a preamendments device, a statement to this effect has been provided.)
2. Submitter'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.
3. 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 replacement of the LX20 Pro analyzer component with a Unicel DxC 600 analyzer with appropriate DxC modifications. Updated barcode reader, peri-pump, syringe drive and module on the Closed Tube Aliquoter. Updates for the Access 2 analyzer component based on the current stand-alone Access 2 analyzer. Software updates to all modules and general overall cosmetics.**
4. **Comparison Information** (similarities and differences) to applicant's legally marketed predicate device including, labeling, intended use, and physical characteristics.
5. 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
 - c) A declaration of conformity with design controls. The declaration of conformity should include:
 - i) A statement signed by the individual responsible, that, as required by the risk analysis, all verification and validation activities were performed by the designated individual(s) and the results demonstrated that the predetermined acceptance criteria were met, and
 - ii) A statement signed by the individual responsible, that the manufacturing facility is in conformance with design control procedure requirements as specified in 21 CFR 820.30 and the records are available for review.
6. A **Truthful and Accurate Statement, a 510(k) Summary or Statement and the Indications for Use Enclosure (and Class III Summary for Class III devices).**

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 submitter'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 submitter 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.

Comments – Validation Protocols were adequate to address the identified causes of hazards identified in the Risk analysis (FTA & FMEA)