May 16, 2018

Conformis, Inc.
Emmanuel Nyakako
Sr. Vice President, Quality and Regulatory
600 Technology Park Drive, 4th Floor
Billerica, Massachusetts 01821

Re: K180906
Trade/Device Name: Conformis iTotal Cruciate Retaining (CR) Knee Replacement System
Regulation Number: 21 CFR 888.3560
Regulation Name: Knee Joint Patellofemorotibial Polymer/Metal/Polymer Semi-Constrained Cemented Prosthesis
Regulatory Class: Class II
Product Code: JWH, OOG, OIY
Dated: April 4, 2018
Received: April 6, 2018

Dear Emmanuel Nyakako:

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration. Please note: CDRH does not evaluate information related to contract liability warranties. We remind you, however, that device labeling must be truthful and not misleading.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); medical device reporting (reporting of medical device-related adverse events) (21 CFR 803); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR Part 807.97). For questions regarding the reporting of adverse events under the MDR regulation (21 CFR Part 803), please go to http://www.fda.gov/MedicalDevices/Safety/ReportaProblem/default.htm for the CDRH's Office of Surveillance and Biometrics/Division of Postmarket Surveillance.

For comprehensive regulatory information about medical devices and radiation-emitting products, including information about labeling regulations, please see Device Advice (https://www.fda.gov/MedicalDevices/DeviceRegulationandGuidance/) and CDRH Learn (http://www.fda.gov/Training/CDRHLearn). Additionally, you may contact the Division of Industry and Consumer Education (DICE) to ask a question about a specific regulatory topic. See the DICE website (http://www.fda.gov/DICE) for more information or contact DICE by email (DICE@fda.hhs.gov) or phone (1-800-638-2041 or 301-796-7100).

Sincerely,

Mark N. Melkerson -S

Mark N. Melkerson
Director
Division of Orthopedic Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

DEPARTMENT OF HEALTH AND HUMAN SERVICES
Food and Drug Administration

Indications for Use

Form Approved: OMB No. 0910-0120
Expiration Date: 06/30/2020
See PRA Statement below.

510(k) Number *(if known)*
K180906

Device Name

Conformis iTotal Cruciate Retaining (CR) Knee Replacement System

Indications for Use *(Describe)*

The iTotal Cruciate Retaining (CR) Knee Replacement System (KRS) is intended for use as a total knee replacement in patients with knee joint pain and disability whose conditions cannot be solely addressed by the use of a prosthetic device that treats only one or two of the three knee compartments, such as a unicondylar, patellofemoral or bicompartmental prosthesis.

The Indications for Use include:

• Painful joint disease due to osteoarthritis, traumatic arthritis, rheumatoid arthritis or osteonecrosis of the knee.

• Post traumatic loss of joint function.

• Moderate varus, valgus or flexion deformity in which the ligamentous structures can be returned to adequate function and stability.

• Failed osteotomies, hemiarthroplasties, and unicondylar, patellofemoral or bicompartmental implants.

• Revision procedures provided that anatomic landmarks necessary for alignment and positioning of the implant are identifiable on patient imaging scans.

This implant is intended for cemented use only.

Type of Use *(Select one or both, as applicable)*

☒ Prescription Use (Part 21 CFR 801 Subpart D) ☐ Over-The-Counter Use (21 CFR 801 Subpart C)

CONTINUE ON A SEPARATE PAGE IF NEEDED.

510(K) SUMMARY (PAGE 1 OF 5)

Submitter's Name and Address:	Conformis, Inc. 600 Technology Park Dr. Billerica, MA 01821
Establishment Registration Number:	3009844603 and 3004153240
Date 510(k) Summary was Prepared:	April 4, 2018
Contact Person:	Emmanuel O. Nyakako, Sr. Vice President, Quality and Regulatory
Telephone Number:	(781) 345-9164
Fax Number:	(781) 345-0147
Name of the Device(s):	Conformis® iTotal® Cruciate Retaining Knee Replacement System (iTotal CR KRS)
Common Name(s):	Knee Replacement System
Regulatory Status and Regulation Number:	Class II 21 CFR 888.3560
Classification Name:	Knee joint patellofemorotibial polymer/metal/polymer semi-constrained cemented prosthesis
Device Classification:	Product Codes: JWH: Knee joint patellofemorotibial polymer/metal/polymer semi-constrained cemented prosthesis. OOG: Knee Arthroplasty Implantation System. Intended to be used to assist in the implantation of a specific knee arthroplasty device or a set of specific knee arthroplasty devices. Indicated to include guiding alignment, making or establishing cuts, selecting, sizing, attaching, positioning or orienting implant components. OIY: Prosthesis, knee, patellofemorotibial, semi-constrained, cemented, polymer + additive/metal/polymer + additive. This generic type of device includes prostheses that have a femoral implant made of alloys, such as cobalt-chromium-molybdenum, and a tibial component(s) and/or a retropatellar resurfacing component made of ultra-high molecular weight polyethylene plus an additive, such as α-tocopherol.

510(K) SUMMARY (PAGE 2 OF 5)

Indications for Use:

The iTotal Cruciate Retaining (CR) Knee Replacement System (KRS) is intended for use as a total knee replacement in patients with knee joint pain and disability whose conditions cannot be solely addressed by the use of a prosthetic device that treats only one or two of the three knee compartments, such as a unicondylar, patellofemoral or bicompartmental prosthesis.

The Indications for Use include:

- Painful joint disease due to osteoarthritis, traumatic arthritis, rheumatoid arthritis or osteonecrosis of the knee.
- Post traumatic loss of joint function.
- Moderate varus, valgus or flexion deformity in which the ligamentous structures can be returned to adequate function and stability.
- Failed osteotomies, hemiarthroplasties, and unicondylar, patellofemoral or bicompartmental implants.
- Revision procedures provided that anatomic landmarks necessary for alignment and positioning of the implant are identifiable on patient imaging scans.

This implant is intended for cemented use only.

Identification of the Legally Marketed Device(s) (Predicate Device(s)):

Conformis iTotal CR Knee Replacement System (iTotal CR KRS)
Device Class**:** II
Product Code: JWH, OOG, OIY
Regulation Number: 21 CFR 888.3560
510(k) Number: K161366

510(K) SUMMARY (PAGE 3 OF 5)

Device Description:

The iTotal CR Knee Replacement System (hereafter referred to as the "iTotal CR KRS") is a patient-specific, tricompartmental, faceted posterior cruciate ligament (PCL) retaining knee replacement system. The iTotal CR KRS is a semi-constrained, cemented knee implant which consists of a femoral, tibial, and patellar component.

Using patient imaging and a combination of proprietary and off-the-shelf software, a patient-specific implant is designed that best meets the geometric and anatomic requirements of the specific patient. The femoral implant is manufactured from cobalt chromium molybdenum (CoCrMo) alloy. The tibial component includes a metal tray manufactured from CoCrMo alloy and either one or two polyethylene inserts. The polyethylene inserts may be manufactured from either UHMWPE or iPoly® XE (a highly cross-linked vitamin E infused polyethylene) The patellar component may also be manufactured from either UHMWPE or iPoly® XE.

For user convenience, and similar to the predicate iTotal CR KRS, single-use, patient-specific ancillary orthopedic manual surgical instruments designed for use with the proposed iTotal CR KRS are provided to assist in the positioning of the total knee replacement components intraoperatively and in guiding the cutting of bone. In addition, reusable orthopedic manual surgical instruments are provided separately.

The function and general design features of the patient-specific implants and ancillary instruments remain similar to those described in the predicate 510(k), **K161366**.

510(K) SUMMARY (PAGE 4 OF 5)

Comparison Summary of Technological Characteristics and Modifications Proposed:

The proposed iTotal CR KRS and predicate iTotal CR KRS have the same technological characteristics. There have been no changes to the proposed iTotal CR KRS device with respect to design, materials or methods of manufacture, packaging, or sterilization.

Modifications, which represent improving manufacturing efficiencies, are primarily focused on increasing automation of the CAD manufacturing processes via the proprietary software modules as shown in **Table 8-1** below.

Table 8-1: Comparison Between the Proposed and Predicate Device

Characteristic	Predicate iTotal CR KRS (K161366)	Proposed iTotal CR KRS
Generation of Femoral Implant	iTotalWorks 5.1 and Fem Addin or Manual CAD Process	iTotalWorks 6.0 & iTotalFem CR 1.0 or Manual CAD Process
Generation of Femoral iJigs	iTotalWorks 5.1 & iTotal FemJigs 3.0 or Manual CAD Process	iTotalWorks 6.0 and iTotalFem CR 1.0 or Manual CAD Process
Generation of Tibial Implants and iJigs	iTotalTib CR 4.3 or Manual CAD Process	iTotalTib CR 5.0 or Manual CAD Process
Generation of Patient-Specific Surgical Plan (iView)	iTotal CR iView 3.0 or Manual CAD Process	iTotal CR iView 4.0 or Manual CAD Process

510(K) SUMMARY (PAGE 5 OF 5)

Substantial Equivalence: The determination of substantial equivalence for this device was based on a detailed device description. Non-clinical laboratory testing was performed demonstrating that the device can be considered substantially equivalent to the predicate device for its intended use. The following non-clinical laboratory testing was performed:

Software verification and validation testing of proprietary software

Conclusion: Based on the testing conducted, it is concluded that the proposed iTotal Cruciate Retaining Knee Replacement System is substantially equivalent to the predicate iTotal Cruciate Retaining Knee Replacement System (**K161366**, cleared June 14, 2016).