DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
10903 New Hampshire Avenue
Document Control Center – WO66-G609
Silver Spring, MD 20993-0002

January 14, 2016

GENOSS Co., Ltd.
c/o Mr. Dachan Kwon
iCT America, Inc.
180 Sylvan Avenue, 2nd Floor
Englewood Cliffs, New Jersey 07632

Re: K15842
Trade/Device Name: rainbow™ Shine
Regulation Number: 21 CFR 872.6660
Regulation Name: Porcelain powder for clinical use
Regulatory Class: II
Product Code: EIH
Dated: December 7, 2015
Received: December 9, 2015

Dear Mr. Kwon:

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.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Division of Industry and Consumer Education at its toll-free number (800) 638-2041 or (301) 796-7100 or at its Internet address http://www.fda.gov/MedicalDevices/ResourcesforYou/Industry/default.htm. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR 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.

You may obtain other general information on your responsibilities under the Act from the Division of Industry and Consumer Education at its toll-free number (800) 638-2041 or (301) 796-7100 or at its Internet address http://www.fda.gov/MedicalDevices/ResourcesforYou/Industry/default.htm.

Sincerely yours,

for Erin I. Keith, M.S.
Director
Division of Anesthesiology, General Hospital,
Respiratory, Infection Control and Dental Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

GENOSS

Indication for use

510(k) Number: K151842

Device Name: rainbow™ Shine

Indication for use:

rainbow™ Shine is used in the manufacture of a dental core through milling by machine (MAD/MAM or CAD/CAM) followed by sintering.

Prescription Use ___√___ **AND/OR** **Over-The-Counter Use** _____

(Part 21 CFR 801 Subpart D) **(21 CFR 801 Subpart C)**

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Concurrence of CDRH, Office of Device Evaluation (ODE)

510(K) Number: **K151842**

510(k) Summary

12/07/2015

1. Company

	Submitter
Name	GENOSS Co., Ltd.
Address	1F, Gyeonggi R&DB Center / 226, 2F, GSBC, 105 Gwanggyo-ro, Yeongtong-gu, Suwon-si, Gyeonggi-do, Korea
Phone/Fax	+82-31-888-5100/ +82-31-888-5105
Contact person	Byungsun Kim / RA bskim@genoss.com
Summary Date	12/07/2015

2. Device Name

Proprietary name	:	rainbow™ Shine
Regulation number	:	21 CFR 872.6660
Classification name	:	Porcelain powder for clinical use
Product code	:	EIH
Device class	:	Class II

3. Predicated Device

K092513 RAINBOW BLOCK

4. Description

rainbow™ Shine is a partially sintered dental ceramic made out of colored ZrO_2(Y-TZP). rainbow™ Shine is milled into cores of artificial teeth and then is finally sintered in a furnace to harden the ZrO_2. Then, the core is layered with porcelain to make a finished tooth.

5. Indication for use

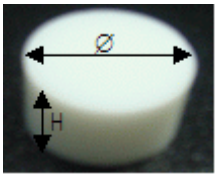
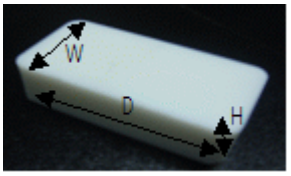
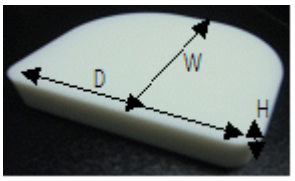
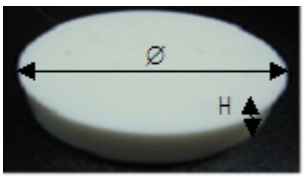
rainbow™ Shine is used in the manufacture of a dental core through milling by machine (MAD/ MAM or CAD/CAM) followed by sintering.

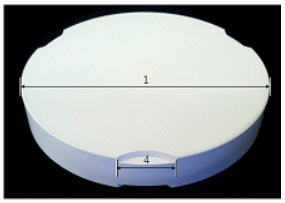
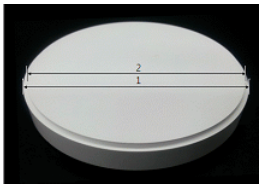
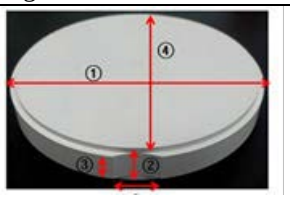
6. Technological Characteristics

The following comparison table of the technological characteristics of the subject device and the predicate devices outlines and provides the similarities and the substantial equivalency of the rainbow™ Shine and the predicate.

Comparison of Characteristics

Device name	**rainbow™ Shine**	**RAINBOW BLOCK**	**Comparison**
Manufacturer	Genoss Co., Ltd.	Genoss Co., Ltd.	Same
510(k) Number	New Device	K092513	-
Materials	Tosoh Powder	Tosoh Powder	Same
Form	Preformed block	Preformed block	Same
Type, class of dental ceramic	Type II - Class 5	Type II - Class 6	The subject and predicate devices satisfy dental ceramic classification requirement by ISO 6872:2008
Sterilization	Non-sterile	Non-sterile	Same
Indication for use	rainbow™ Shine is used in the manufacture of a dental core through milling by machine (MAD/ MAM or CAD/CAM) followed by sintering.	Rainbow Block is used in the manufacture of a dental core through milling by machine (MAD/MAM or CAD/CAM) followed by sintering.	Same
Use	Prescription	Prescription	Same
Technical characteristics			
Bending Strength (Flexural strength)	706MPa	1144MPa	Bending strength is higher than required by ISO 6872:2008 for Class 5 dental ceramics. (> 500 Mpa)
Sintering Density (g/cm^3)	6	6.00~6.04	Same

Radioactivity (Bq/g)		< 1.0 Bq/g	< 1.0 Bq/g	Same
Chemical Solubility ($\mu g/cm^2$)		3	14.47	Chemical solubility satisfies requirement by ISO 6872:2008 for Class 5 dental ceramics. (< 2000 $\mu g/cm^2$)
Biocompatibility		Biocompatible	Biocompatible	Same
Shape	**Single**	 Diameter 25.3mm Height 12, 16, 22mm	 Diameter 25mm Height 12, 16, 22mm	Similar The subject and predicate devices are available in numerous shapes and sizes in order to be compatible with multiple milling machines.
	2~9 bridge	 Diameter 44.5~75.5mm Width 25.5, 25, 26, 38mm Height 12, 16, 22mm	 Diameter 33~75.5mm Width 25~38mm Height 12mm	
	10~16 bridge	 Diameter 89, 95mm Width 55, 75mm Height 12, 16, 22mm	 Diameter 80~95mm Width 55, 75mm Height 12mm	
	Disk	 Diameter 87, 95, 98, 100mm Height 10~28mm	 Diameter 87, 95, 98,100mm Width 10~26mm Height 12, 16, 22mm	

Shape	**Step block**	 Diameter 95mm Height 10~28mm	 Diameter 95, 98mm Height 10~28mm	
		 Diameter 98mm Height 10~28mm	 Diameter 95, 98mm Height 10~28mm	
		 Diameter 98mm Height 10~28mm	–	

There are two minor differences that are worth discussing:

1) The subject and predicate device have slight difference in materials. The ZrO_2 and Y_2O_3 are major materials of general zirconia blank. The differences in the composition between the subject and predicate device relate to the percentages of Y_2O_3 and they have effect on the crystal structure and consequently on the translucency of the device. SiO_2, Fe_2O_3, Na_2O and Er_2O_3 are trace elements used as colorants, but they have no material effects on the biocompatibility of the device. The non-clinical tests(bench test and biocompatibility test) demonstrate the Technical characteristics.

2) The differences of technological characteristics(Bending strength, chemical solubility) are within what is expected of this type of device. The rainbow Shine has a lower bending strength than the predicate device, due to the fact that the material is completely stabilized with Y2O3 in order to achieve higher translucency. However, the performance test results satisfy the requirement by ISO 6872:2008

7. Summary of non-clinical testing

Non-clinical device testing was conducted to confirm the performance of the subject device. Bench testing was conducted in accordance with the FDA recognized consensus standard(Recognition number : 4-178: ISO 6872 Third edition 2008-09-01, dentistry - ceramic materials) Bench tests for performance comparison of the subject device and the predicate device includes the following testing:

- Bending Strength
- Linear thermal expansion coefficient
- Chemical Solubility

Technical characteristics of both devices satisfy the requirements by ISO 6872:2008. Slight differences between the subject and predicate devices do not raise any new issues of safety and effectiveness.

Biocompatibility testing was conducted on the device pursuant to the ISO 10993-1:2009 Biological evaluation of medical device - Part 1: Evaluation and testing within a risk management process.

- Cytotoxicity test(ISO 10993-5)
- Irritation or intracutaneous reactivity(ISO 10993-10)
- Sensitization(ISO 10993-10)
- Acute systemic toxicity(ISO 10993-11)
- Genotoxocity(ISO 10993-03)

The result of biocompatibility testing demonstrated that no issue of biocompatibility arises.

8. Conclusion

Based on the information provided in this premarket notification of GENOSS Co., Ltd. Concludes that rainbow™ Shine is substantially equivalent to the predicate device.