



U.S. Food and Drug Administration  
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Ofer Cohen  
Quality Assurance Manager  
COLORCHIP (ISRAEL) LTD.  
TAVOR BUILDING 1, P.O. BOX 158  
NEW INDUSTRIAL PARK  
YOKNEAM, ISRAEL 2069203

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This is to acknowledge receipt of your August 29, 2018, document, which was filed pursuant to the regulations for the administration and enforcement of the Radiation Control for Health and Safety Act of 1968 (title 21, code of Federal Regulations, Subchapter J) as they pertain to Annual Report requirements.

Your document has been assigned an Accession Number of 1831697-000, and has been classified as a(n) Annual Report (pursuant to Part 1002, Subpart B of the Regulation referenced above).

Further, the submittal has been assigned an informal subject title of "This submission is a(n) Annual Report. These Data Measurement, Transmit, Control Laser Products cover the period from July 01, 2017 to June 30, 2018."

This acknowledgement does not constitute approval of the document. You will be contacted if any questions or comments arise concerning your document.

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If you have any questions, please contact the Director of the Division of Radiological Health, or the branch chief of your respective product area, as listed on the CDRH Management Directory, under the Office of In Vitro Diagnostics and Radiological Health, Division of Radiological Health.  
<http://www.fda.gov/AboutFDA/CentersOffices/OfficeofMedicalProductsandTobacco/CDRH/CDRHOffices/ucm127854.htm>

Please include a primary (and optional secondary) contact email address in all submissions (and/or cover letters) to facilitate electronic correspondence.

Sincerely yours,

Division of Radiological Health  
Office of In Vitro Diagnostics and Radiological Health  
Center for Devices and Radiological Health