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ExcelVite Passes US FDA Inspection

June 13th, 2017 – New Jersey, USA – ExcelVite, the first and only PIC/S GMP-certified palm phytonutrients producer is pleased to announce the successful completion of its quality audit / inspection by the United States' Food and Drug Administration (US FDA).

The US FDA inspection was conducted over three days at its production site in Malaysia. ExcelVite's production plant and Quality Management System (QMS) were inspected according to FDA's Code of Federal Regulation – Title 21 Current Good Manufacturing Practices (21 CFR-cGMP). There was no non-compliance within ExcelVite's systems and hence, no Non-Conformance was issued at the end of the inspection.

"As part of the company's commitment to ensure that our QMS is compliant with current good manufacturing practice (cGMP), we are proud to know that our existing system meets the quality criteria and requirements of the US FDA. It is of utmost importance to ensure that the entire system is adhered to and implemented at all times in order to ensure product quality and safety, which is our commitment to customers," says Ms. YT Chan, Quality Assurance Manager of ExcelVite.

"Being a dedicated and largest manufacturer and exporter of palm phytonutrients especially EVNol™ natural full spectrum tocotrienol complex and EVTene™ natural mixed-carotene complex, ExcelVite is committed to the highest level of product quality, safety and manufacturing standards. In addition to enforcing cGMP PIC/S Guide (Pharmaceutical Inspection Co-Operation Scheme's Guide to Good Manufacturing Practice for Medicinal Products), ExcelVite is proud to have achieved this important milestone in the FDA audit. It represents and underscores our compliance with the 21 CFR-cGMP Guidelines that allows for continuous export to the United States," says WH Leong, CEO of ExcelVite.

"I want to congratulate and thank Team EV for their effort in upholding the most stringent production and quality systems in its daily operation that have led to the success in this important FDA inspection. The commitment and conscientiousness shown are commendable and highly appreciated. The company is proud to have such a team and we look forward to implementing continuous improvement efforts to maintain and sustain our leadership position in the market," added Mr. Leong.

This is the second time that the US FDA has inspected ExcelVite's manufacturing facility located at Chemor, Perak, Malaysia. The first inspection had been conducted in 2012.

About ExcelVite

ExcelVite Sdn. Bhd., incorporated in Malaysia in 2013, is the leading and largest producer of natural full spectrum tocotrienol / tocopherol complex (EVNol™, and EVNol SupraBio™), natural mixed carotenoids complex (EVTene™), phytosterol complex (EVRol™), red palm oil concentrate (EVSpectra™) in the world via a patented technology.

ExcelVite is the only tocotrienol producer that operates in accordance to GMP (PIC/S) Guide to Good Manufacturing Practice for Medicinal Products. Its laboratory is accredited with ISO/IEC 17025 accreditation.

EVNol SupraBio™ is a patented (US Patent No. 6,596,306) self-emulsifying palm tocotrienol complex that ensures optimal tocotrienols oral absorption.

ExcelVite manufactures and markets its products under the tradenames: EVNol™, EVNol SupraBio™, EVTene™, EVRol™, and EVSpectra™. These branded ingredients are Non-GMO, Kosher and Halal certified. ExcelVite supports the production of certified sustainable palm oil (CSPO) through the GreenPalm Program.

Websites: www.excelvite.com and www.tocotrienol.org

Contacts:

WH Leong, CEO	Tel: +61 (03) 9801 3881	Email: info@excelvite.com
Bryan See, Regional Product Manager	Tel: +1 (732) 906 1901	Email: klsee@excelvite.com
CheeYen Lau, Nutritionist	Tel: +605 2014 192	Email: cylau@excelvite.com