## **PRODUCTION OF PALM-BASED TOCOTRIENOLS-ENHANCED FRACTION (TEF)**

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ocotrienols as well as tocopherols are homologues of vitamin E. Vitamin E is a fatsoluble natural antioxidant and present in small amount in vegetable oils and fats. The importance of vitamin E, an essential vitamin, for keeping the good health in animals and human has long been recognized. Deficiency of vitamin E in animals would lead to complications, such as foetal resorption and muscular dystrophy. Although  $\alpha$ -tocopherol is the most active form of vitamin E in terms of foetal resorption tests, it is interesting to note that progress on research over the last few years indicates that tocotrienols form of vitamin E are superior than  $\alpha$ -tocopherol in activities such as hypocholesterolemic and anti-cancer effects. In order to harness the full benefits of tocotrienols, there is a need for tocotrienols preparation with minimal amount of tocopherols in it. Almost all current tocotrienols preparations in the market contain tocopherols, for instance Palm Vitamin E, a tocotrienols-rich fraction (TRF) is a mixture of tocotrienols and tocopherols at a typical ratio of 75 to 25. A process for preparing tocotrienols-enhanced fraction (TEF) products with a ratio of tocotrienols to tocopherols more than 90:10 is presented.

### **PRODUCTION OF TEF**

The current sources for extraction of tocotrienols are from agro-based industries including palm oil and rice bran oil. TRF preparations are available in the market, for example at 20%, 50% and 70% total vitamin E content, with a typical ratio of tocotrienols to tocopherols at 75:25. We propose a technology for increasing the content of tocotrienols to more than 80% and improving the ratio of tocotrienols to tocopherols to 90:10 or more. An example is the production of MPOB-T3Mix88, a TEF from palm fatty acid distillate (Figure 1). The process involves the preparation of palm-based TRF and subsequently extraction of tocopherol resulting in an increase of tocotrienols content from 57.3% to 88.0% and the ratio of tocotrienols to tocopherols from 76:24 to 97:3 (Table 1). Another product, MPOB-T3 Mix 50

Vitamin E Concentration 0.4%-0.8% Palm fatty acid distillate **ESTERIFICATION** Methyl esters 0.4%-0.8% DISTILLATION ➔ Distilled Methyl esters Distillation 5%-10% residue CONCENTRATION Tocotrienols-60%-80% **Rich Fraction** (TRF) TOCOPHEROL **ISOLATION** >80% Tocotrienols-**Enhanced Fraction** (TEF)

Figure 1. Production of palm-based tocotrienolsenhanced fraction (TEF).





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Tocotrienols Preparation	α-Tocopherol (%)	α-Tocotrienol (%)	γ-Tocotrienol (%)	δ-Tocotrienol (%)	Total (%)
Palm Vitamin E (TRF)	17.8	17.1	23.0	17.2	75.1
MPOB-T3 Mix88 (TEF)	2.6	25.2	37.9	24.9	90.6
MPOB-T3 Mix50	1.5	14.6	21.6	14.8	52.5

# TABLE 1. TOCOPHEROLS AND TOCOTRIENOLS CONTENT OF<br/>TOCOTRIENOLS PREPARATIONS

Notes: TRF = tocotrienols-rich fraction.

TEF = tocotrienols-enhanced fraction.

(*Figure 2*) with tocotrienols content 51% is prepared by blending MPOB-T3Mix88 with RBD palm olein accordingly. The product (MPOB-T3 Mix50) with a minimal tocotrienols content of 50%, can be easily promoted for various applications.



Figure 2. Palm-based tocotrienols (MPOB-T3Mix50).

### POTENTIAL BENEFITS OF TOCOTRIENOLS

Both tocopherols and tocotrienols are known for their antioxidant activities in foods and biological systems (Ab Gapor *et al.*, 1989; Minhajuddin *et al.*, 2005).  $\alpha$ -Tocotrienol showed better antioxidant activity than  $\alpha$ -tocopherol in the order of 40-60 times against lipid peroxidation (Serbinova *et al.*, 1991). In a human study, O'Byrne *et al.* (2000) reported that  $\alpha$ -tocotrienol may be potent in decreasing low-density lipoprotein (LDL) oxidizability. Tocotrienols have been shown to be a potential hypocholesterolemic agent (Qureshi et al., 1995; Hasselwander et al., 2002). Tocotrienols have also been shown to have anti-cancer activities (Kato et al., 1985; Komiyama et al., 1989; Wan Ngah et al., 1999; Nesaretnam et al., 2000). The superiority of tocotrienols over tocopherols in anti-cancer effects have been confirmed (Guthrie et al., 1997; Yu et al., 1999; McIntyre et al., 2000; Yu et al., 2003). Tocotrienols have been reported to show anti-angiogenic activity, suggesting potential usefulness as a therapeutic agent for tumour angiogenesis (Miyazawa et al., 2004).  $\alpha$ -Tocotrienol but not tocopherols has been shown to be a potent neuroprotection agent (Sen et al., 2000; Osakada et al., 2004). Anderson et al. (2003) reported that tocotrienols induce IKBKAP expression, thus suggesting a possible therapy for familial dysautonomia.

#### CONCLUSION

A substantial amount of evidence indicates that tocotrienols are more effective than tocopherols in health promoting activities including hypocholesterolemic, anti-cancer and antioxidant properties. Technology for producing TEF is available for further development. The products are expected to have applications in the high end nutraceuticals, functional foods and cosmetics industries.

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#### REFERENCES

AB GAPOR MD TOP; ONG, A S H; KATO, A; WATANABE, H and KAWADA, T (1989). Antioxidant activities of palm vitamin E with special reference to tocotrienols. *Elaeis*, *1*(1): 63-67.

ANDERSON, S L; QUI, J and RUBIN, B Y (2003). Tocotrienols induce IKBKAP expression: a possible therapy for familial dysautonomia. *Biochem Biophys Res Comm*, 306: 303-309.

GUTHRIE, N; GAPOR, A; CHAMBERS, A F and CARROLL, K K (1997). Inhibition of proliferation of estrogen receptor-negative MDA-MB-435 and -positive MCF-7 human breast cancer cells by palm oil tocotrienols and tamoxifen, alone and in combination. *J. Nutr.*, *127*(*3*): 544S-548S.

HASSELWANDER, O; KRAMER, K; HOPPE, P P; OBERFRANK, U; BALDENIUS, K; SCHRODER, H; KAUFMANN, W; BAHNEMANN, R and NOWAKOWSKY, B (2002). Effects of feeding various tocotrienol sources on plasma lipids and aortic atherosclerotic in cholesterol-fed rabbits. *Fd. Res. Intern.*, *35*: 245-251.

KATO, A; YAMAOKA, M; TANAKA, A; KOMIYAMA, K and UMEZAWA, I (1985). Physiological effect of tocotrienol. *J. Japan Oil Chem. Soc.* (YUKUGAKU), 34: 375-376.

KOMIYAMA, K; IIZUKA, K; YAMAOKA, M; WATANABE, H; TSUCHIYA, N and UMEZAWA, I (1989). Studies on the biological activity of tocotrienols. *Chem. Phar. Bull.*, *37*(5): 1369-1371.

MCINTYRE, B S, BRISKI, K P; TIRMENSTEIN, M A; FARISS, M W; GAPOR, A and SYLVESTER, P W (2000). Antiproliferative and apoptotic effects of tocopherols and tocotrienols on normal mouse mammary epithelial cells. *Lipids*, *35*(2): 171-180.

MINHAJUDDIN, M; BEG, Z H and IQBAL, J (2005). Hypolipidemic and antioxidant properties of tocotrirnol rich fraction isolated from rice bran oil in experimentally induced hyperlipidemic rats. *Fd Chem Toxico*, 43: 747-753.

MIYAZAWA, T; INOKUCHI, H; HIROKANE, H; TSUZUKI, T; NAKAGAWA, K and IGARASHI, M (2004). Anti-angiogenic potential of tocotrienol *in vitro*. *Biochemistry* (*Moscow*), 69(1): 67-69.

NESARETNAM, K; DORAISAMY, S and DARBRE, P D (2000). Tocotrienols inhibit growth of ZR-75-1

breast cancer cells. Internat. J. Fd. Sci. Nut., 5: S95-S103.

O'BYRNE, D; GRUNDY, S; PACKER, L; DEVARAJ, S; BALDENIUS, K; HOPPE, P P; KRAEMER, K; JIALAL, I and TRABER, M G (2000). Studies of LDL oxidation following  $\alpha$ -,  $\gamma$ - and  $\delta$ -tocotrienol acetate supplementation of hypercholesterolemic humans. *Free Radical Biology & Medicine*, 29(9): 834-845.

OSAKADA, F; HASHINO, A; KUME, T; KATSUKI, H; KANEKO, S and AKAIKE, A (2004). α-Tocotrienol provides the most potent neuroprotection among vitamin E analogs on cultured striatal neurons. *Neuropharmacology*, 447: 904-915.

QURESHI, A A; BRADLOW, B A; BRACE, L; MANHANDLE, J; PETERSON, D M; PEARCE, B C; WRIGHT, J K; GAPOR, A and ELSON, C E (1995). Response of hypercholesterolemic subjects to administration of tocotrienols. *Lipids*, *30* (12): 1171-1177.

SEN, C K; KHANNA, S; ROY, S and PACKER, L (2000). Molecular basis of vitamin E action. *J. Bio. Chem.*, 275(17): 13049-13055.

SERBINOVA, E; KAGAN, V; HAN, D and PACKER, L (1991). Free radical recycling and intramembrane mobility in the antioxidant properties of alpha-tocopherol and alpha-tocotrienol. *Free Radical Biology* & *Medicine*, 10: 263-275.

YU, W; SIMMONS-MENCHACHA, M; GAPOR, A; SANDERS, B G and KLINE, K (1999). Induction of apoptosis in human breast cancer cells by tocopherols and tocotrienols. *Nutrition and Cancer*, *33*(*1*): 26-32.

YU, F L; GAPOR, A and BENDER, W (2003). Evidence for the preventive effect of the polyunsaturated phytol side chain in tocotrienols on 17b-estradiol epoxidation. Paper presented at PIPOC 2003 International Palm Oil Congress  $\beta$ - Food Technology and Nutrition Conference. 24-28 August 2003. Putrajaya Marriott Hotel.

WAN NGAH, W Z; AZIAN, A L; NARIMAH, A H H; PERMEEN, Y; NASARUDDIN, A; AB GAPOR, M T; MAZLAN, Z and KHALID, B A K (1999). Palmvitee supplementation of chronic hepatits B patients and *in vitro* studies on cancer cell lines. *Proc. of the 1999 PORIM International Palm Oil International Congress - Nutrition Congress*. p. 233-240. For more information kindly contact:

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