

Enhanced heterologous production of eicosapentaenoic acid in *Escherichia coli* cells that co-express eicosapentaenoic acid biosynthesis *pfa* genes and foreign DNA fragments including a high-performance catalase gene, *vktA*

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There were typing errors in Tables 1 and 2 of the original. The corrected Tables plus an updated reference are shown on the following page.

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Table 1 Fatty acid composition of *E. coli* DH5 α and its various transformants and recovered amount of EPA from cultures

Strains ^a	Fatty acid ^b (% total)					Content of EPA ($\mu\text{g ml}^{-1}$)
	16:0	16:1(9)	18:1(11)	EPA	Others ^c	
<i>E. coli</i> DH5 α	36.0 \pm 1.0	29.6 \pm 0.7	22.0 \pm 0.6	0	12.5 \pm 1.4	0
<i>E. coli</i> DH5 α (pEPA Δ 1)	35.6 \pm 0.9	26.9 \pm 1.5	21.8 \pm 0.9	2.5 \pm 0.2	13.2 \pm 2.7	1.7 \pm 0.1
<i>E. coli</i> DH5 α (pEPA Δ 1) (pGBM3)	38.6 \pm 1.8	28.2 \pm 0.6	20.8 \pm 0.3	3.2 \pm 1.7	9.2 \pm 1.1	1.5 \pm 1.3
<i>E. coli</i> DH5 α (pEPA Δ 1)[pGBM3::sal(<i>vktA</i>)]	35.9 \pm 3.1	18.5 \pm 0.4	22.9 \pm 1.9	12.3 \pm 0.7	10.3 \pm 0.8	7.3 \pm 1.2
<i>E. coli</i> DH5 α (pEPA Δ 1) [pGBM3::sal(Δ <i>vktA</i>)]	34.0 \pm 0.7	26.7 \pm 0.2	24.1 \pm 1.2	5.9 \pm 0.2	9.2 \pm 1.7	3.3 \pm 0.2

^a The cells were grown at 20 °C until the culture had an OD₆₆₀ of 1.0

^b Fatty acids are denoted as number of carbon atoms:number of double bond. The Δ -position of double bond is presented in parenthesis

^c Others include 12:0, 14:0, 18:0, and 3-hydroxyl 14:0

Table 2 Catalase activity of *E. coli* DH5 α and its various transformants

Strains ^a	Catalase activity (U mg protein ⁻¹)
<i>E. coli</i> DH5 α (pEPA Δ 1)	3.3
<i>E. coli</i> DH5 α (pEPA Δ 1)(pGBM3)	3.2
<i>E. coli</i> DH5 α (pEPA Δ 1)[pGBM3::sal(<i>vktA</i>)]	535
<i>E. coli</i> DH5 α (pEPA Δ 1)[pGBM3::sal(Δ <i>vktA</i>)]	1.7

^a Cells were washed three times with phosphate buffer (pH 7.5) by centrifugations at 3,000 \times *g* for 15 min, and they were then suspended in 0.2 ml of 60 mM potassium phosphate buffer (pH 7.0). Cells were disrupted by sonic oscillation using a Sonifier Cell Disruptor (model W185; Branson Ultrasonic Corp., Danbury, CT) for 40 s in an ice bath. Supernatants were removed after the centrifugation of cell lysates at 20,000 \times *g* for 60 min and were used as cell-free extracts

Reference

Okuyama H, Orikasa Y, Nishida T, Watanabe K, Morita N (2007) Bacterial genes responsible for the biosynthesis of eicosapentaenoic and docosahexaenoic acids and their heterologous expression. *Appl Environ Microbiol* 73:665–670