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Opportunities to enhance alternative sources of long-chain *n*-3 fatty acids within the diet

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Health benefits or advocated health benefits of long-chain (LC) *n*-3 PUFA are better known by medical doctors as well as by consumers, so that consumption increases. In addition, the development of aquaculture requires more fishmeal and fish oil. Humanisation of care of companion animals is also associated with addition of LC *n*-3 PUFA in pet foods. The risk of the increased demand for LC *n*-3 PUFA is the excess harvesting of natural sources, especially of marine origin (oily fishes, krill). In order to improve sustainability, alternative sources of LC *n*-3 PUFA have been developed. These alternative sources are: (a) terrestrial plants naturally or genetically enriched in stearidonic acid (SDA), which bypasses the first limiting step of (i.e. $\Delta 6$ desaturase) of the biosynthesis of LC *n*-3 PUFA; (b) single-cell oils rich in LC *n*-3 PUFA (microalgae, *Escherichia coli*) and krill. Currently, plants rich in SDA are expensive, metabolic engineering is unfavourably accepted by consumers in many countries, cultivation of microalgae is very expensive even though their ability (for some of them) to synthesise biofuels could induce a decrease in industrial costs, and Antarctic krill harvest must be restricted. Thus, it is difficult to predict their real development in the future.

n-3 Fatty acids: EPA: DHA: Nutrition: PUFA

According to a Packaged Facts™ report⁽¹⁾, the number of consumers who are seeking out high-*n*-3 products has increased dramatically over the past few years. In the US market, the percentage of adults who take fish oil supplements has jumped from 8% in 2006 to 17% in 2011; global consumer spending on *n*-3 products (excluding fish) was about $\$13 \times 10^9$ in 2011 and will reach $\$34.7 \times 10^9$ by 2016. Asia–Pacific is accounting for 43% of sales.

According to a Transparency Market Research™ report⁽²⁾, the demand for fish oil was 1035 kilotonnes in 2011 and is expected to reach 1130 kilotonnes in 2018. Aquaculture (salmon and trout) accounted for over 70% of the consumption of oil in 2011 primarily in Chile and Peru. However, direct human consumption of fish oil has been increasing over the past 5 years,

due to marketer's influence and due to an increase in awareness of health benefits of long-chain (LC) *n*-3 PUFA, such as prevention of type-2 diabetes (Asian, Eskimos), prevention of insulin-resistance^(3,4), decrease in plasma TAG⁽⁵⁾, anti-inflammatory effect⁽⁶⁾, the defect of brain and retina development (only if deficiency in LC *n*-3 PUFA), primary cardiovascular prevention⁽⁵⁾, anti-stress and anti-adrenergic effect^(7,8) and decrease in liver steatosis⁽⁹⁾. Because of these health benefits and of some deleterious effects of deficiency, many national and international recommendations have been proposed about the optimal intake of LC *n*-3 PUFA for the general population. They vary considerably depending on the country and on the type of illness to prevent and/or to treat (Table 1) (see website: www.omega3dressings.com/Dietary_Recommendation.html).

Abbreviations: LC, long-chain; SCO, single-cell oils; SDA, stearidonic acid.

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Table 1. Selected suggested long-chain *n*-3 (EPA+DHA) intakes for adults available from various agencies and bodies (from⁽¹⁰⁾)

Authority/Group	Intake (mg/d)
Omega Workshop, Adelaide, Australia, 2002	300–400
SACN/COT, UK, 2004	450
National Heart Foundation, 2008	500
American Dietetic Association and Dietitians of Canada, 2007	500
FAO/WHO Expert Consultation, 2008	250–2000*
American Heart Association, 2002	
CHD sufferers	1000
Those seeking to reduce TAG (blood fats)	2000–4000
Australia and New Zealand (suggested dietary targets), 2006	
Female	430
Male	610
French AFSSA, 2010	
General Nutrition	500
Metabolic disease risk reduction	500
CVD risk reduction	500–750
Breast and colon cancer risk reduction	500
Neuropsychiatric risk reduction	>200–300
AMD risk reduction	500
European Food Safety Agency	250
Ministry of Health, Labor and Welfare, Japan, 2005–2010	2000–2600

SACN/COT, Scientific Advisory Committee on Nutrition/Committee on Toxicity of Chemicals in Food, Consumer Products and the Environment; AFSSA, Agence française de sécurité sanitaire des aliments; AMD, age-related macular degeneration.

*For secondary prevention of CHD.

Table 2. Species of fish and countries where they are caught for fishmeal and oil production (from⁽¹²⁾)

Species	Country
Anchovy	Peru, Chile, South Africa, Namibia, Mexico, Morocco
Jack mackerel	Peru, Chile
Capelin	Norway, Iceland, Russian Federation
Menhaden	USA, Atlantic and Gulag of Mexico
Blue whiting	Norway, UK, Russian Federation, Ireland
Sand eel	Denmark, Norway, Faeroe Islands
Norway pout	Denmark, Norway, Faeroe Islands
Sprat	Denmark, Russian Federation

A calculation can be easily made: choosing a daily 500 mg requirement of EPA+DHA, a global population reaching 8×10^9 by 2025, assuming fish contains 2–5 mg/100 g oil and an LC *n*-3 PUFA content of 450 mg/150 g flesh, the current global fish harvest (93 metric tonnes per annum) will not supply this requirement. If the UK recommendation of intake of two meals including 140 g fish per week were followed⁽¹¹⁾, then annual per capita consumption would have to rise to 23.3 kg. This translates into an additional production of 40 million tonnes for 2008, rising to 82 million tonnes in 2050. Per capita fish consumption per year has increased hugely from the 1960s: 9.9 kg in the 1960s, 11.5 kg in the 1970s, 12.6 kg in the 1980s, 14.4 kg in the 1990s, 17.0 kg in 2007 (China 26.7 kg; 14.6 kg without China) and 17.1 kg in 2008. Fish and fish trimmings caught for fishmeal and fish oil productions are reported in Tables 2 and 3.

Myers & Worm⁽¹³⁾ have published in *Nature* a frequently cited paper where they conclude that

‘Industrialised fisheries typically reduced community biomass by 80 % within 15 years of exploitation. We estimate that large predatory fish biomass today is only about 10 % of pre-industrial levels. We conclude that some species compensation was evident, but often reversed within a decade or less, probably because of changes in targeting or bycatch.’

For all these reasons, concerns have been reached about the risk of depletion of oily fish communities (the most used source of LC *n*-3 PUFA). Thus, over the past few years, alternative sources of LC *n*-3 PUFA to oily fish have been developed.

Alternative sources of long-chain *n*-3 PUFA

In order to be acceptable for use in human subjects, alternative sources and final products must be safe, green and healthy. The first alternative to oily fish capture is aquaculture. However, aquaculture uses mainly fish oil and fishmeal⁽¹⁴⁾. Strobel *et al.*⁽¹⁵⁾ reported that, in Germany, fat content in fillets from cultured salmon was 12.3 v. 2.07 wt % in the wild counterpart, which resulted in 4-fold greater absolute amount in total LC *n*-3 PUFA (2.40 (sd 0.78) v. 0.53 (sd 0.21)% of fresh weight). Nichols *et al.*⁽¹⁶⁾ reported that in Australia the average content in LC *n*-3 PUFA of wild fish was 350 mg/150 g wet weight as compared with a range of 1200–3700 mg/150 g in farmed fish. This requires fishmeal and fish oil.

In order to increase sustainability of sources of LC *n*-3 PUFA, fish oil and fishmeal use can be limited to the last weeks of growth (finishing diets). However,

Table 3. Fish trimmings used for fishmeal and oil production (from⁽¹²⁾).

Species	Country
Catfish	USA, Vietnam
<i>Tuna</i> sp.	Thailand, Japan, USA, Australia, South Korea, China, France, Ecuador, Maldives Islands and others
Salmon (wild)	Norway, USA–Alaska (wild), UK, Ireland, Canada, Chile, Japan
Sardine/Pilchard	Peru, Chile, South Africa, Namibia, Japan, Spain, Mexico
White fish sp.	UK, USA–Alaska, Canada, Chile
Dogfish	Canada, USA
Pollock	USA–Alaska
Horse mackerel	Ireland, Norway, Denmark, Spain
Atlantic herring	Iceland, Norway, Denmark, UK, Faeroe Islands, Sweden, Ireland, Canada
Mackerel sp.	UK, Peru, Chile, South Africa
Hoki (Blue grenadier)	Australia, New Zealand

other sources should be used or developed for aquaculture or as a direct source of LC *n*-3 PUFA as food ingredient or as pharmaceutical. These alternatives are: (a) terrestrial or GM plant-based lipids; (b) single-cell oil (SCO); (c) krill.

Terrestrial plant based-lipids and transgenic oil seeds

Terrestrial plants, such as canola, sunflower, soya, flax, olive, palm or linseed contain linoleic acid (18:2*n*-6), oleic acid (18:1*n*-9) or α -linolenic acid (18:3*n*-3) have been used to feed farmed fish, but their flesh content in LC *n*-3 PUFA was lower than that of their wild counterparts. Consequently, they are now used only during the initial phase of growth and fish oils or fishmeal rich in LC *n*-3 PUFA are used as finishing diet over the final 10 weeks.

Stearidonic acid (SDA, 18:4*n*-3) is more efficiently converted to EPA than 18:3*n*-3 because it does not require the first rate-limiting step: Δ 6-desaturase. Foods rich in SDA are hemp oil, seeds from Boraginaceae family of plants including borage, blackcurrant seed oil (2–4% SDA) and *Echium plantagineum* seed oil (echium oil, 12–14% SDA)^(17,18). A new plant oil extracted from the seeds of *Buglossoides arvensis* (Ahiflower™ oil) is a rich source of SDA (20% SDA). Lemke *et al.*⁽¹⁹⁾ compared in healthy subjects either 1.5 g/d high-oleic sunflower ethyl ester oil capsules plus foods containing 7 g/d high-oleic sunflower oil (control) or 1.5 g/d EPA oil ethyl ester capsules plus foods containing 7 g/d high-oleic sunflower oil (EPA), or 1.5 g/d high-oleic sunflower ethyl ester oil capsules plus foods containing 7 g/d SDA soyabean oil for 12 weeks. The content in EPA of erythrocyte membranes was 0.50 ± 0.03, 2.17 ± 0.21 and 0.85 ± 0.05% for control, EPA and SDA, respectively. This demonstrates the ability of SDA to be converted into EPA in human subjects but the content in EPA was far below this obtained following EPA oil supplement.

A SDA-rich oil (*E. plantagineum* L., Boraginaceae; 14% SDA), given over 6 weeks to Atlantic salmon parr (freshwater phase) led to a similar accumulation of LC *n*-3 PUFA to fish oil⁽²⁰⁾. As a comparison, canola oil rich in 18:3*n*-3 did not lead to accumulation of EPA and DHA. Thus, SDA-rich aquafeeds might replace

fish oil in freshwater aquaculture. However, this occurred over the period before smolting, which has been shown to coincide with a period of peak of LC *n*-3 PUFA production. Conversely, when given to seawater Atlantic salmon, SDA-rich oil did not produce the same level of EPA and DHA as a fish oil diet⁽²¹⁾. Thus, further assessment of SDA-rich oil is needed. Furthermore, it is currently more expensive than fish oil.

A mixed vegetable oil (mix of rapeseed, palm and degummed linseed oils in a 3.7:2:1 ratio) resulted after the finishing diet period, in LC *n*-3 PUFA flesh content similar to that obtained with a 100% fish oil fed salmon without any difference in growth, mortality and sensory characteristics⁽²²⁾.

Transgenic plants have been created to produce LC *n*-3 PUFA owing to the characterisation of LC PUFA biosynthetic pathways and the availability of many genes encoding these pathways (see details in Venegas-Calerón *et al.*⁽²³⁾). Briefly, the (aerobic) Δ 6-pathway is widely spread in eukaryotes; the alternative pathway in algae and protists is the Δ 8-pathway (18:3*n*-3 is first elongated to 20:3 Δ 11,14,17 eicosatrienoic acid followed by a Δ 8-desaturation). A third (anaerobic) pathway (present in a few lower marine organisms) requires neither desaturases nor elongases, but a processive polyketide synthase-like reaction from malonyl-CoA (Fig. 1).

These transgenic plants can synthesise EPA and DHA, but most of them also accumulate intermediates of the *n*-6 pathway. Recently, Petrie *et al.*⁽¹⁴⁾ described the transgenic DHA production by *Arabidopsis thaliana* with a DHA content as high as 15% (higher than the 12% usually found in classical fish oils) and a high *n*-3: *n*-6 ratio (8:1 to 16:1 depending on the native 18:3*n*-3 and 18:2*n*-6 were included or not). However, the EPA content of this transgenic seed was only 1.8% (18% in classical fish oils). The same authors⁽¹⁴⁾ calculated that the application of this technology in crop species would result in 1 ha of a *Brassica napus* (canola) crop containing 12% DHA in seed oil to produce as much DHA as approximately 10 000 fish. Ruiz-Lopez *et al.*⁽²⁴⁾ have also developed a transgenic approach in the seed oil of the crop *Camelina sativa*. They obtained some seeds containing EPA levels up to 31% (mean 24%) and other

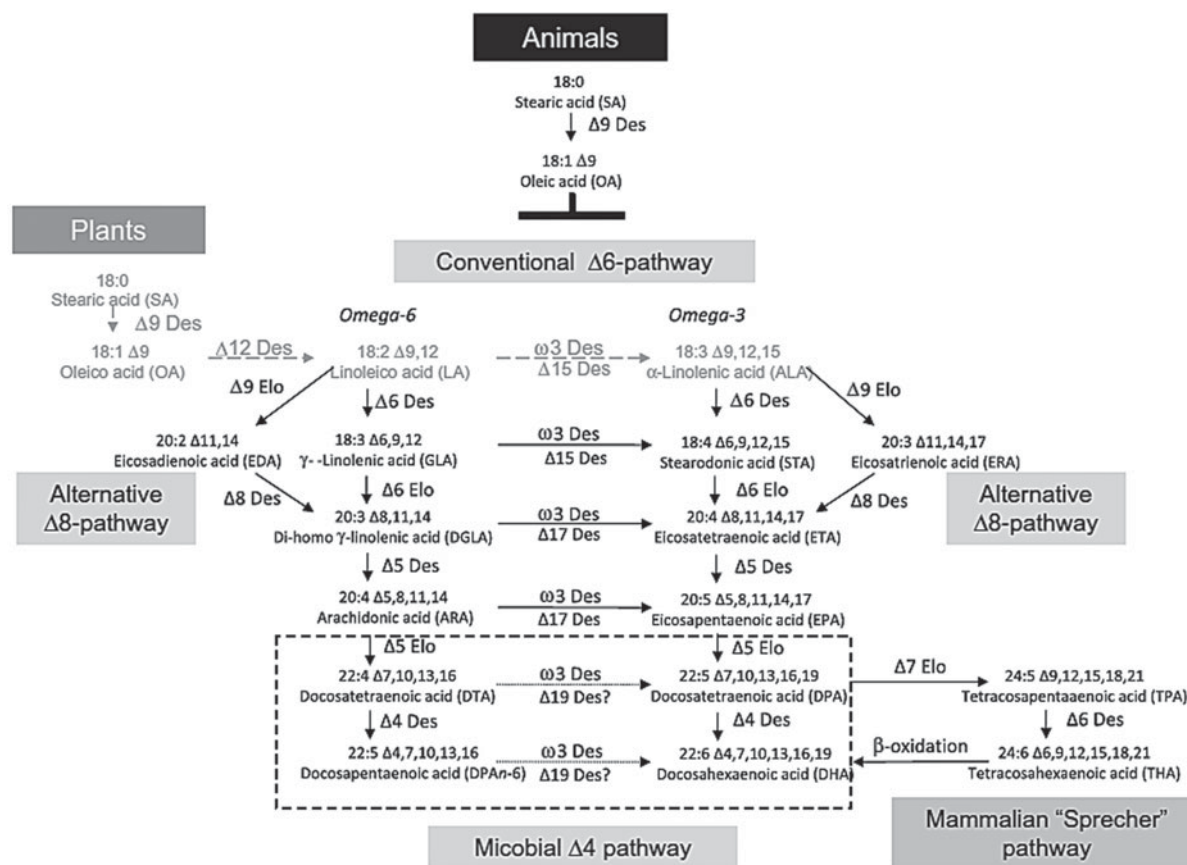


Fig. 1. Aerobic very long chain PUFA biosynthetic pathways. The various routes for synthesis of arachidonic acid, EPA and DHA are shown, as mediated by the consecutive action of desaturases and elongases. The predominant $\Delta 6$ -pathway is shown, as is the alternative $\Delta 8$ -pathway. Two routes for DHA synthesis are shown, microbial $\Delta 4$ -pathway and mammalian ‘Sprecher’ pathway. Des, desaturase; Elo, elongase (from ⁽²³⁾).

seeds accumulating up to 12% EPA and 14% DHA (mean 11% EPA and 8% DHA).

In Europe, GM organisms are not well accepted by the consumers. Thus, a more acceptable alternative could be to feed farmed fish with these transgenic plants to enhance EPA and DHA generation instead of fishmeal or fish oil.

Single-cell oil

SCO, extracted from micro-organisms grown under heterotrophic conditions, can also be rich in LC *n*-3 oils. Examples of micro-organisms are microalgae^(25–27) including thraustochytrids^(28,29) and *Escherichia coli*⁽³⁰⁾.

Microalgae

The ability of microalgae to be an alternative source of LC *n*-3PUFA has been recently reviewed⁽³¹⁾. In the marine food chain, microalgae (phytoplankton) are rich in EPA and DHA. They are consumed by zooplankton itself consumed by fish, so that the content in EPA and DHA is particularly high in marine oily fish. Microalgae offer a non-polluted resource for LC *n*-3 PUFA production, but for human consumption,

microalgae and their products must be non-pathogenic and non-toxic, genetically stable, and present high growth and product formation rates⁽³²⁾. Ideally, the selected strains should be able to accumulate high amounts of TAG rich in EPA and/or DHA (up to 50% biomass dry weight, including 30–70% DHA).

LC *n*-3 PUFA biosynthesis from α -linolenic acid by the elongase–desaturase pathway is common in microalgae. In some species, biosynthesis of EPA involves elongation of α -linolenic acid, followed by a $\Delta 8$ and $\Delta 5$ desaturation; DHA is obtained through EPA elongation into docosapentaenoic acid and subsequent desaturation by $\Delta 4$ desaturase, or through the anaerobic polyketide synthase pathway, as it has been suggested for thraustochytrids⁽²³⁾.

Photoautotrophic microalgae

Photoautotrophic LC *n*-3 PUFA producing microalgae are mainly marine species. The EPA and DHA content and the EPA:DHA ratio vary from species to species (Tables 4–6). Large variation in EPA and DHA content within one species is due to the culture conditions, the growth phase of the microalgae at the time of harvest and the extraction technique used to determine the

Table 4. Examples of marine microalgae species characterised by EPA production (adapted from⁽²⁶⁾)

	EPA content (% TFA)	EPA content (% DW)
<i>Nannochloropsis</i> sp. Hibberd	38–39	2–3
	15–18	5–6
	11–22	3–6
	15–27	4
	5–27	2–4
	30–35	3–4
	35–39	4
<i>Phaeodactylum tricomutum</i> Bohlin	31	5
	40–57	1–4
	28	3
	30–32	3
	38–42	4–5
<i>Nitzschia laevis</i> Hustedt	25–33	3–4
	11–16	2–3
<i>Porphyridium cruentum</i> Nägeli	25	3
	41	–
<i>Odontella aurita</i> Agardh	26	–
<i>Pavlova lutheri</i> Green	18–23	–
	22–29	–
<i>Cyclotella cryptica</i> Lewin and Guillard	17–23	1
<i>Cylindrotheca</i> sp. Rabenhorst	24–25	–

TFA, total fatty acids; DW, dry weight.

lipid content and fatty acid profile. Photoautotrophic technologies for marine microalgae cultures in open pond systems cultivation can utilise non-arable lands and water resources considered unsuitable for agriculture. These processes also have a more favourable net energy ratio, relative to heterotrophic systems, but specific growth rates in these systems are relatively low, harvesting may be costly, and the risks of salinity fluctuation and contamination can lead to variable quality and quantity of the final product. Photobioreactors allow better growth efficacy and productivity, better control over environmental parameters, significant reduction of contamination risks, and allow higher biomass concentrations, but high volume to surface ratio is currently limited due to light penetration restrictions. Lastly, they are very expensive. Thus, the use of these systems is currently justified only for production of high-value products.

Photosynthetic microalgae tend to produce higher levels of EPA than heterotrophs. *Nannochloropsis* Hibberd, *Phaeodactylum* Bohlin, *Nitzschia* Hassall and *Porphyridium* Nägeli can present a high percentage of EPA in total fatty acids, but its total amount in the biomass is low (Table 4). Photoautotrophic species contain lipids involved in the photosynthetic metabolism and, unlike various DHA-rich microalgae, the fatty acid profiles of EPA producers usually show other LC-PUFA, such as DHA and/or arachidonic acid (Table 5).

Phototrophic species, such as the eustigmatophyte *Nannochloropsis*, have long been used by the aquaculture industry with the aim to supply larval fish with *n*-3 LC-PUFA through the enrichment of live feeds

(e.g. rotifers; for review see Martins *et al.*⁽²⁶⁾). Gog *et al.*⁽³⁴⁾ described a *Nannochloropsis oculata* with a fatty acid profile containing 49% EPA, while most commonly *Nannochloropsis* strains show lower levels, in the range of 11–39%. Although *Nannochloropsis* accumulates TAG in response to environmental stressors, EPA deposition may be low within this lipid class, which is instead associated with shorter fatty acids. Deposition of EPA in *Nannochloropsis* seems to be favoured by low salinity, whereas total lipids appear to increase in higher salinity waters. The cultivation of *Nannochloropsis* sp. in ultra-dense cultures, with valuable EPA production, using frequent replacement of nutrient medium, has been described. In fact, the genus *Nannochloropsis* is currently the source of marketed oils due to its potential to produce high EPA lipids with very low DHA and arachidonic acid content, which is advantageous for the manufacture of dietary supplements (Table 7).

Heterotrophic microalgae

Heterotrophic microalgae produce EPA and/or DHA mainly as TAG and phospholipids. Heterotrophic DHA, predominantly obtained with *Schizochytrium* species, is already commercially available for human consumption, animal feed and aquaculture. Heterotrophic cultivation without light, using inexpensive mineral medium supplemented with a carbon source is feasible provided fermentation and culture conditions are optimised to obtain the highest content in EPA and/or DHA of each strain. Overall, application of photosynthetic species for the commercial production of oils requires advances in physiological and genetic engineering studies to enhance growth performances and lipid deposition, optimisation of fatty acid composition, biotechnological improvements regarding light capture and contamination, and lowering of costs involved in biomass production and harvesting⁽³³⁾. Several marine heterotrophic microalgae are considered as good DHA sources (Table 5). These species represent the most preeminent alternative industrial sources of oils rich in DHA (Table 7), with approved use in human foods, especially for application in infant formulas, since they are considered to be non-pathogenic and non-toxicogenic. Ward and Singh⁽³²⁾ reported that some *Schizochytrium* strains might produce levels as high as 94% DHA of total *n*-3 fatty acids. *Schizochytrium* sp. possess a number of traits that are advantageous for the industry, including high lipid content, elevated DHA production and presentation in the TAG form, as well as good growth in culture under elevated cell concentrations. In *Schizochytrium* strains, DHA may reach 30–35% of total fatty acids mainly in TAG fraction (see Martins *et al.*⁽²⁶⁾ for details).

Thraustochytrids have also been used in the grow-out phase of Atlantic salmon, either as whole-cell biomass or the extracted SCO. Complete replacement of fish oil by thraustochytrid oil in the diets resulted in a comparable growth and health associated with a higher concentration of DHA in the flesh. The large quantity of oil required throughout the life history of salmonids and marine

Table 5. Examples of marine microalgae species characterised by DHA production (adapted from⁽²⁶⁾)

Species	DHA content (% TFA)	DHA content (% DW)
<i>Schizochytrium mangrovei</i> Raghuk	31–41	12–21
<i>Schizochytrium limacinum</i> Honda and Yokochi	25–35	5–15
	–	15–19
<i>Schizochytrium</i> sp. (HX-308) Goldstein and Belsky	40–56	11–20
<i>Schizochytrium</i> sp. Goldstein and Belsky	45–52	20–24
	28	4
<i>Thraustochytrium</i> sp. Sparrow	23–24	16–17
<i>Thraustochytrium aureum</i> Goldstein	32–37	6–7
<i>Thraustochytrium striatum</i> Schneider	37	2
<i>Ulkenia</i> sp. Gaertner	10–23	5
<i>Aurantiochytrium</i> sp. Yokoyama and Honda	40	18
<i>Cryptothecodinium cohnii</i> Javornicky	19–34	2–4
	63	6
	53–57	5–6

TFA, total fatty acids; DW, dry weight.

Table 6. DHA and EPA content in various fish species (from⁽³³⁾)

Type	DHA (g/100g)	EPA (g/100g)	DHA+EPA (g/100g)	Ratio (DHA:EPA)
Tuna	1.141	0.363	1.504	3.1:1.0
Anchovy	0.911	0.538	1.449	1.7:1.0
Salmon				
Atlantic, farmed	1.457	0.690	2.147	2.1:1.0
Atlantic, wild	1.429	0.411	1.840	3.5:1.0
Chinook	0.727	1.010	1.737	1.0:1.4
Sockeye	0.700	0.530	1.230	1.3:1.0
Mackerel, Atlantic	0.699	0.504	1.203	1.4:1.0
Menhaden	1.19	1.84	3.03	1.0:1.54
Herring, Atlantic	1.105	0.909	2.014	1.2:1.0
Trout				
Rainbow, farmed	0.820	0.334	1.154	2.5:1.0
Rainbow, wild	0.520	0.468	0.988	1.1:1.0
Halibut	0.374	0.091	0.465	4.1:1.0
Cod	0.154	0.004	0.158	38.5:1.0
Haddock	0.162	0.076	0.238	2.1:1.0
Catfish				
Channel, farmed	0.128	0.049	0.177	2.6:1.0
Channel, wild	0.137	0.100	0.237	1.4:1.0
Swordfish	0.681	0.087	0.768	7.8:1.0
Grouper	0.213	0.035	0.248	6.1:1.0
Shrimp	0.144	0.171	0.315	1.0:1.2

fish makes such the use of SCO cost-prohibitive as compared to LC *n*-3 PUFA obtained from plants. A potentially cost-effective approach is to use SCO in diets during the final 6–12 weeks of fish growth. A reduction of production costs of SCO is awaited because of the increasing interest in the industrial biofuels production from microalgae⁽³⁵⁾.

Transgenic microalgae

Metabolic engineering of microalgae is, as in terrestrial plants, a way to enhance LC *n*-3 PUFA^(36,37). *Phaeodactylum tricorutum* is a unicellular diatom, which accumulates EPA (up to ~35%), but only trace levels of DHA. Hamilton *et al.*⁽³⁶⁾ have overexpressed,

in *P. tricorutum*, the Δ 6-desaturase and Δ 5-elongase, which significantly increased the DHA levels.

Krill

Antarctic krill (*Euphausia superba*) are the small, bug-eyed shrimp-like crustaceans that are the central diet for whales, penguins, seals and seabirds. Krill contain LC *n*-3 PUFA mainly as phospholipids⁽³⁸⁾ but also as TAG⁽³¹⁾. *E. superba* contains the highest concentration of EPA and DHA compared to other krill species⁽³⁹⁾. The potential health benefits⁽⁴⁰⁾ from these attributes are being marketed in various forms of krill oil products. Krill is considered as the single largest ‘underutilised’ commercial marine resource remaining. However, as

Table 7. Levels of EPA and/or DHA in commercially available oils derived from marine microalgae cultures (adapted from⁽²⁶⁾)

Company and commercial product designation	% EPA or DHA	Microbial sources	Techniques
Aurora Algae A2 EPA Pure™	65 % EPA (regular) 95 % EPA (pharma)	Undisclosed	Phototrophic, open-pond
Qualitas Health EicoOil™	25–30 % EPA	<i>Nannochloropsis oculata</i> Hibberd	Phototrophic, open-pond
Algae Biosciences AlgaeBio Omega-3 Origins™	20 % EPA; 20 % DHA	Undisclosed	Oil blend from two marine strains
DSM-NP life's DHA™	40–45 % DHA	<i>Cryptocodinium cohnii</i> Javornicky	Heterotrophic fermentation
DSM-NP life's DHA plus EPA™	10 % EPA; 22.5 % DHA	Schizochytrium sp. Goldstein and Belsky	Heterotrophic fermentation
Lonza DHAid™	35–40 % DHA	<i>Ulkenia</i> sp. Gaertner	Heterotrophic fermentation
Source – Omega Source Oil™	35–40 % DHA	<i>Schizochytrium</i> sp. Goldstein and Belsky	Heterotrophic fermentation
GCI Nutrients DHA Algae 35 % oil	35 % DHA	<i>Cryptocodinium cohnii</i> Javornicky	Heterotrophic fermentation

factory ships are deployed and climate is changing, krill and Antarctic food web that depend on it could be threatened. The Southern Ocean krill biomass is estimated at up to 700 million metric tonnes⁽⁴¹⁾. The annual krill harvest is still well within the very strict limits set by the Convention on the Conservation of Antarctic Marine Living Resources (twenty-four member countries and the EU), which regulates fishing in the Southern Ocean (www.ccamlr.org). The total annual catch level of the current Convention on the Conservation of Antarctic Marine Living Resources is 5 610 000 tonnes/year on sector 48 and 3 085 000 tonnes on sector 58. There is also a precautionary limit of 620 000 tonnes limit in area 48 (the so-called 'trigger level') and 452 000 for area 58-4-2. Once this limit is achieved, Convention on the Conservation of Antarctic Marine Living Resources might close the krill fishery until a procedure for the division of the overall catch limit into smaller management units.

As written by Naylor *et al.*⁽⁴²⁾ 'Krill is at the base of the Southern Ocean food web and is also particularly sensitive to environmental variables, including climate change. In some regions, considerable overlap exists between the krill fishery and the foraging ranges of land-based predators, such as penguins, which cannot move readily to new feeding areas. Local krill significantly reduce the food resources of other predators, such as seals and whales. Unfortunately, existing data on krill abundance and population variables are not sufficient to establish precautionary management of the krill fishery and its effect on the Antarctic ecosystem. Considerable care will thus be needed in setting local catch limits for krill harvest to protect key predators and other animals in the Southern Ocean ecosystem, and the Commission for the Conservation of Antarctic Marine Living Resources is trying to develop data-driven procedures to achieve this.'

Conclusion and perspectives

The ratio of wild to farmed fish has hugely decreased over the past decade to fall below one. This could be

good news for sustainability of natural resources. However, aquaculture is now the main user of fishmeal and fish oil. Progress in aquaculture has led to the conclusion that the use of fishmeal rich in LC *n*-3 PUFA could be used only as a finishing diet over the final 10 weeks of fish growth without altering growth, taste or content in LC *n*-3 PUFA as compared to wild fish counterparts. Efforts have been made to develop alternative sources of LC *n*-3 PUFA. Crops rich in SDA have a greater interest on their own in human subjects or given to animals. However, cost is very high. Genetic engineering has increased the ability of terrestrial plants to produce more SDA or EPA and/or DHA, but they are poorly generally accepted by the consumers. A more acceptable way (with some psychological limits anyway) could be their use as feed in animals secondarily consumed by human subjects. SCO rich in LC *n*-3 PUFA show great promise for production of oils and for reducing dependence on wild fisheries. Their high cost of production makes them infeasible for feeding of farmed fish. Their ability to produce bio-fuels should help to decrease the costs in the future. Metabolic engineering of SCO (microalgae, *E. coli*) is also a way to generate LC *n*-3 PUFA. Do not forget that human insulin was for many years produced from *E. coli* strains. The better delineation of the real daily needs of LC *n*-3 PUFA, as well as their real health benefits and the specific effect of EPA *v.* DHA will bring a real boost in the production of high-quality high-grade LC *n*-3 supplements or of enriched foods of good nutritional value. The choice of the best source of LC *n*-3 PUFA should be determined by price, availability, formulation, taste and sustainability.

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Conflicts of interest

None.

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Authorship

J. D. wrote the present paper; N. G. performed the literature search, gathered the articles, reviewed the manuscript and made the necessary corrections.

References

- Packaged Facts (2011) Omega-3: Global Product Trends and Opportunities. <http://www.packagedfacts.com/Omega-Global-Product-6385341/>
- Transparency Market Research (2013) Fish Oil Market for Aquaculture, Direct Human Consumption, Hydrogenation and Industrial Applications - Global Industry Analysis, Size, Share, Growth, Trends and Forecast, 2012–2018. <http://www.transparencymarketresearch.com/fish-oil.html>
- Delarue J, LeFoll C, Corporeau C *et al.* (2004) *n*-3 long chain polyunsaturated fatty acids: a nutritional tool to prevent insulin resistance associated to type 2 diabetes and obesity? *Reprod Nutr* **44**, 289–299.
- Zhang M, Picard-Deland E & Marette A (2013) Fish and marine omega-3 polyunsaturated fatty acid consumption and incidence of type 2 diabetes: a systematic review and meta-analysis. *Int J Endocrinol*. Available at: <http://dx.doi.org/10.1155/2013/501015>.
- Mozaffarian D & Wu JH (2012) (*n*-3) fatty acids and cardiovascular health: are effects of EPA and DHA shared or complementary? *J Nutr* **142**, 614S–625S.
- Calder PC (2011) Fatty acids and inflammation: the cutting edge between food and pharma. *Eur J Pharmacol* **668**, Suppl. 1, S50–S58.
- Delarue J, Matzinger O, Binnert C *et al.* (2003) Fish oil prevents the adrenal activation elicited by mental stress in healthy men. *Diab Metab* **29**, 289–295.
- Delarue J, Guillodo MP, Guillerm S *et al.* (2008) Fish oil attenuates adrenergic overactivity without altering glucose metabolism during an oral glucose load in haemodialysis patients. *Br J Nutr* **99**, 1041–1047.
- Di Minno MN, Russolillo A, Lupoli R, *et al.* (2012) Omega-3 fatty acids for the treatment of non-alcoholic fatty liver disease. *World J Gastroenterol* **18**, 5839–5847.
- Dietary recommendations for omega-3 fatty acids (2014) www.omega3dressings.com/Dietary_Recommendation.html
- Scientific Advisory Committee on Nutrition and Committee on Toxicity (2004) Advice on fish consumption: benefits and risks. Norwich, UK, The Stationery Office. http://www.sacn.gov.uk/pdfs/fics_sacn_advice_fish.pdf
- Alaska seafood by-products: potential products, markets and competing products. Report prepared for Alaska Fisheries Development Foundation Anchorage, Alaska January 8, 2009, by Anthony P. Bimbo. <http://seafod.oregonstate.edu/pdf/%20Links/Alaska-Seafood-By-Products-Potential-Products-Markets-and-Competing-Products.pdf>
- Myers RA & Worm B (2003) Rapid worldwide depletion of predatory fish communities. *Nature* **423**, 280–283.
- Petrie JR, Shrestha P, Zhou XR *et al.* (2012) Metabolic engineering plant seeds with fish oil-like levels of DHA. *PLoS One* **7**, e49165.
- Strobel C, Jahreis G & Kuhnt K (2012) Survey of *n*-3 and *n*-6 polyunsaturated fatty acids in fish and fish products. *Lipids Health Dis* **11**, 144.
- Nichols PD, Petrie J & Singh S (2010) Long-chain omega-3 oils—an update on sustainable sources. *Nutrients* **2**, 572–585.
- Walker CG, Jebb SA & Calder PC (2013) Stearidonic acid as a supplemental source of ω -3 polyunsaturated fatty acids to enhance status for improved human health. *Nutrition* **29**, 363–369.
- Surette ME (2013) Dietary omega-3 PUFA and health: stearidonic acid-containing seed oils as effective and sustainable alternatives to traditional marine oils. *Mol Nutr Food Res* **57**, 748–759.
- Lemke SL, Maki KC, Hughes G *et al.* (2013) Consumption of stearidonic acid-rich oil in foods increases red blood cell eicosapentaenoic acid. *J Acad Nutr Diet* **113**, 1044–1056.
- Miller MR, Nichols PD & Carter CG (2007) Replacement of dietary fish oil for Atlantic salmon parr (*Salmo salar* L.) with a stearidonic acid containing oil has no effect on omega-3 long-chain polyunsaturated fatty acid concentrations. *Comp Biochem Physiol B Biochem Mol Biol* **146**, 197–206.
- Miller MR, Bridle AR & Nichols PD (2008) Increased elongase and desaturase gene expression with stearidonic acid enriched diet does not enhance long-chain (*n*-3) content of seawater Atlantic salmon (*Salmo salar* L.). *J Nutr* **138**, 2179–2185.
- Torstensen BE, Bell JG, Rosenlund G *et al.* (2005) Tailoring of Atlantic salmon (*Salmo salar* L.) flesh lipid composition and sensory quality by replacing fish oil with a vegetable oil blend. *J Agric Food Chem* **53**, 10166–10178.
- Venegas-Calcrón M, Sayanova O & Napier JA (2010) An alternative to fish oils: metabolic engineering of oil-seed crops to produce omega-3 long chain polyunsaturated fatty acids. *Prog Lipid Res* **49**, 108–119.
- Ruiz-Lopez N, Haslam RP, Napier JA, *et al.* (2013) Successful high-level accumulation of fish oil omega-3 long-chain polyunsaturated fatty acids in a transgenic oil-seed crop. *Plant J* **77**, 198–208.
- Hong WK, Rairakhwada D, Seo PS *et al.* (2011) Production of lipids containing high levels of docosahexaenoic acid by a newly isolated microalga, *Aurantiochytrium* sp. KRS101. *Appl Biochem Biotechnol* **164**, 1468–1480.
- Martins DA, Custódio L, Barreira L *et al.* (2013) Alternative sources of *n*-3 long-chain polyunsaturated fatty acids in marine microalgae. *Mar Drugs* **11**, 2259–2281.
- Ryckebosch E, Bruneel C, Muylaert K *et al.* (2012) Microalgae as an alternative source of omega-3 long chain polyunsaturated fatty acids. *Lipid Technol* **24**, 128–130.
- Zhou PP, Lu MB, Li W *et al.* (2010) Microbial production of docosahexaenoic acid by a low temperature-adaptive strain *Thraustochytridae* sp. Z105: screening and optimization. *J Basic Microbiol* **50**, 380–387.
- Qu L, Ji XJ, Ren LJ *et al.* (2011) Enhancement of docosahexaenoic acid production by *Schizochytrium* sp. using a two-stage oxygen supply control strategy based on oxygen transfer coefficient. *Lett Appl Microbiol* **52**, 22–27.
- Amiri-Jami M & Griffiths MW (2010) Recombinant production of omega-3 fatty acids in *Escherichia coli* using a gene cluster isolated from *Shewanella baltica* MAC1. *J Appl Microbiol* **109**, 1897–1905.
- Araujo P, Zhu H, Breivik JF *et al.* (2013) Determination and structural elucidation of triacylglycerols in krill oil by chromatographic techniques. *Lipids* **49**, 163–172.



32. Ward OP & Singh A (2005) Omega-3/6 fatty acids: alternative sources of production. *Process Biochem* **40**, 3627–3652.
33. Lee JH, O’Keefe JH, Lavie CJ *et al.* (2008) Omega-3 fatty acids for cardioprotection. *Mayo Clin Proc* **83**, 324–332.
34. Gog A, Senila L, Roman M *et al.* (2012) Oil extraction and fatty acid characterization of *Nannochloropsis oculata* microalgae for biodiesel applications. *Stud Univ Babeş-Bol* **57**, 111–118.
35. Lum KK, Kim J & Lei XG (2013) Dual potential of microalgae as a sustainable biofuel feedstock and animal feed. *J Anim Sci Biotechnol* **4**, 53 [Epub ahead of print].
36. Hamilton ML, Haslam RP, Napier JA *et al.* Metabolic engineering of *Phaeodactylum tricorutum* for the enhanced accumulation of omega-3 long chain polyunsaturated fatty acids. *Metab Eng* **22**, 3–9.
37. Mühlroth A, Li K, Røkke G *et al.* (2013) Pathways of lipid metabolism in marine algae, co-expression network, bottlenecks and candidate genes for enhanced production of EPA and DHA in species of Chromista. *Mar Drugs* **11**, 4662–4697.
38. Winther B, Hoem N, Berge K *et al.* (2011) Elucidation of phosphatidylcholine composition in krill oil extracted from *Euphausia superba*. *Lipids* **46**, 25–36.
39. Phleger CF, Nelson MM, Mooney BD *et al.* (2002) Interannual and between species comparison of the lipids, fatty acids and sterols of Antarctic krill from the US AMLR elephant Island survey area. *Comp Biochem Physiol B Biochem Mol Biol* **131**, 733–747.
40. Ulven SM, Kirkhus B, Lamglait A *et al.* (2011) Metabolic effects of krill oil are essentially similar to those of fish oil but at lower dose of EPA and DHA, in healthy volunteers. *Lipids* **46**, 37–46.
41. Miller MR, Nichols PD & Carter CG (2008) *n*-3 Oil sources for use in aquaculture – alternatives to the unsustainable harvest of wild fish. *Nutr Res Rev* **21**, 85–96.
42. Naylor RL, Hardy RW, Bureau DP *et al.* I (2009) Feeding aquaculture in an era of finite resources. *Proc Natl Acad Sci USA* **106**, 15103–15110.