FPIN's Clinical Inquiries

Fish Oil for Treatment of Dyslipidemia

RAMA NARLA, MD, and SHANNON B. PECK, PhD, Baylor Family Medicine Residency, Garland, Texas KEFENG (MAYLENE) QIU, MLIS, University of Pennsylvania Biomedical Library, Philadelphia, Pennsylvania

Clinical Inquiries provides answers to questions submitted by practicing family physicians to the **Family Physicians Inquiries** Network (FPIN). Members of the network select questions based on their relevance to family medicine. Answers are drawn from an approved set of evidence-based resources and undergo peer review. The strength of recommendations and the level of evidence for individual studies are rated using criteria developed by the **Evidence-Based Medicine** Working Group (http:// www.cebm.net/?o=1025).

The complete database of evidence-based questions and answers is copyrighted by FPIN. If interested in submitting questions or writing answers for this series, go to http://www.fpin. org or email: questions@fpin.org.

A collection of FPIN's Clinical Inquiries published in *AFP* is available at http://www.aafp.org/afp/ fpin.

Clinical Question

How effective is fish oil in lowering lipid levels in adults with dyslipidemia?

Evidence-Based Answer

Supplementation with omega-3 fatty acids decreases triglyceride and very low-density lipoprotein cholesterol levels. However, it can also increase low-density lipoprotein (LDL) cholesterol levels. Treatment with omega-3 fatty acids does not decrease total mortality, cardiovascular events, or cancer incidence, and therefore should not be recommended to patients to decrease their risk of dyslipidemia. (Strength of Recommendation: A, based on meta-analyses of randomized controlled trials [RCTs].)

Evidence Summary IMPROVEMENTS IN LABORATORY VALUES

A Cochrane review of 23 RCTs involving 1,075 participants with type 2 diabetes mellitus showed that supplementation with fish oil (average dosage of 3.5 g per day for an average of 8.9 weeks) lowered triglyceride levels by 8.1 mg per dL (0.1 mmol per L; P < .00001). In treated patients, very lowdensity lipoprotein cholesterol levels decreased by 1.26 mg per dL (0.03 mmol per L; P = .04), and LDL cholesterol levels increased by 1.98 mg per dL (0.05 mmol per L; P = .05). There was no change in high-density lipoprotein cholesterol levels.

An earlier Cochrane review of 17 RCTs (N = 3,918) sought to determine the effect of omega-3 fatty acid supplementation on rates of mortality and cardiovascular events.² The meta-analysis of studies that included adults at risk of cardiovascular disease found that omega-3 fatty acid supplementation significantly reduced serum triglyceride levels (weighted mean

difference = -7.2 mg per dL [-0.1 mmol per L];95% confidence interval [CI], -10.08 to -4.14). Total and high-density lipoprotein cholesterol levels did not change significantly. LDL cholesterol significantly increased (weighted mean difference = 2.34 mg per dL [0.06 mmol per L; 95% CI, 0.54 to 3.96]). It should be noted that this study examined omega-3 intake from fish and plant sources.

A systematic review of four RCTs concluded that supplementation with two omega-3 fatty acids (docosahexaenoic acid [DHA] and eicosapentaenoic acid [EPA]) in doses of 1.7 g or greater reduced triglyceride levels by at least 10% compared with baseline.³ Patients with higher baseline triglyceride levels had the greatest benefit.

NO IMPROVEMENTS IN PATIENT OUTCOMES

Two meta-analyses investigated the effects of fish oil supplementation on mortality rates. A meta-analysis of 17 RCTs (N = 63,279) identified no relationship between omega-3 fatty acid intake and mortality (relative risk = 0.96; 95% CI, 0.91 to 1.02).⁴ Follow-up for participants in the included studies ranged from one to 6.2 years. A Cochrane review of 48 RCTs (N = 36,913) and 41 cohort studies evaluating participants over six months showed no reduction in rates of overall mortality, cardiovascular events, or cancer.² However, the length of follow-up may have been inadequate to establish effectiveness.

Recommendations from Others

The American Heart Association recommends daily supplementation with 2 to 4 g of EPA/DHA for patients with elevated triglyceride levels.³ The amount of EPA/DHA ranges from 300 to approximately 850 mg, ▶

Clinical Inquiries

depending on the brand.3 The National Cholesterol Education Program expert panel has not recommended a specific dosage of omega-3 fatty acids for supplementation.⁵

Copyright Family Physicians Inquiries Network. Used with permission.

Address correspondence to Rama Narla, MD, at rama. narla@baylorhealth.edu. Reprints are not available from the authors.

Author disclosure: No relevant financial affiliations.

REFERENCES

1. Hartweg J, Perera R, Montori V, Dinneen S, Neil HA, Farmer A. Omega-3 polyunsaturated fatty acids (PUFA) for type 2 diabetes mellitus. Cochrane Database Syst Rev. 2008;(1):CD003205.

- 2. Hooper L, Thompson RL, Harrison RA, et al. Omega 3 fatty acids for prevention and treatment of cardiovascular disease. Cochrane Database Syst Rev. 2004;(4):CD003177.
- 3. Balk E, Chung M, Lichtenstein A, et al. Effects of omega-3 fatty acids on cardiovascular risk factors and intermediate markers of cardiovascular disease. Evidence report/technology assessment no. 93. Rockville, Md.: Agency for Healthcare Research and Quality;
- 4. Rizos EC, Ntzani EE, Bika E, Kostapanos MS, Elisaf MS. Association between omega-3 fatty acid supplementation and risk of major cardiovascular disease events: a systematic review and meta-analysis. JAMA. 2012:308(10):1024-1033.
- 5. Third report of the National Cholesterol Education Program (NCEP) Expert Panel on Detection, Evaluation, and Treatment of High Blood Cholesterol in Adults (Adult Treatment Panel III) final report. Circulation. 2002;106(25):3143-3421.

Straightforward. Accessible. Manageable. TAKE THE GUESSWORK OUT OF SAMS.

Register early to save \$40

Course Date / Subject / Location

April 12 / Cerebrovascular / Las Vegas, NV April 13 / Preventive Care / Las Vegas, NV July 18 / Asthma / Myrtle Beach, SC July 19 / Diabetes/ Myrtle Beach, SC Sept. 20 / Coronary Artery Disease / Chicago, IL (Rosemont)

Sept. 21 / Childhood Illness / Chicago, IL (Rosemont)

Sept. 20 / Hypertension / Albuquerque, NM (Santa Ana Pueblo)

Sept. 21 / Health Behavior / Albuquerque, NM (Santa Ana Pueblo)

Oct. 11 / Hospital Medicine / Hershey, PA

Oct. 12 / Heart Failure / Hershey, PA

Oct. 22 / Diabetes / Washington, D.C.

Oct. 23 / Asthma / Washington, D.C.

Oct. 24 / Preventive Care / Washington, D.C.

Oct. 25 / Health Behavior / Washington, D.C.

Nov. 15 / Heart Failure / Las Vegas, NV

Nov. 16 / Well Child Care / Las Vegas, NV

Attend an AAFP SELF-ASSESSMENT MODULE

WORKING GROUP and complete the knowledge assessment portion of your MC-FP Part II with ease.

In one day, assess and review the 60 core competency questions, led by expert family physicians currently working in the field, and decide the best answer as a group. The AAFP will report your successful knowledge assessment score to the ABFM on your behalf.

Participate with your colleagues in a distraction-free, focused work environment that fosters interaction and promotes collaboration, while you earn CME*.

Prices start at \$225 for AAFP members aafp.org/completesams (800) 274-2237



ONE DAY.

^{*}To complete the SAM process and earn CME, you must also complete the clinical simulation on the ABFM website within one year (separate fee to the ABFM required).