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A table in the article “Lipid-regulating and anti-atherosclerotic therapy: current options and future approaches” (*Cleveland Clinic Journal of Medicine* 1996; 63:31–41) contained an error. In Table 6 on page 37, the values for the effects of the various drugs on HDL-C and LDL-C were reversed through an editing error. The corrected table appears below.

<b>Bile-acid sequestrants</b>		
Lipid effects:	LDL-C:	↓ 15%–30%
	HDL-C:	↑ 3%–5%
	TG:	↑ or no effect
Drugs and daily dose:	Cholestyramine	4–24 g
	Colestipol	5–30 g
<b>HMG-CoA reductase inhibitors</b>		
Lipid effects:	LDL-C:	↓ 20%–40%
	HDL-C:	↑ 5%–15%
	TG:	↓ 10%–20%
Drugs and daily dose:	Fluvastatin	20–40 mg
	Lovastatin	10–80 mg
	Pravastatin	10–40 mg
	Simvastatin	5–40 mg
<b>Nicotinic acid (NA)</b>		
Lipid effects:	LDL-C:	↓ 10%–25%
	HDL-C:	↑ 15%–35%
	TG:	↓ 20%–50%
Drugs and daily dose:	Crystalline NA	1.5–6 g
<b>Fibric-acid derivatives<sup>†</sup></b>		
Lipid effects:	LDL-C:	↓ 10%–15% (may↑)
	HDL-C:	↑ 10%–15%
	TG:	↓ 20%–50%
Drugs and daily dose:	Gemfibrozil	1200 mg
	Clofibrate	2000 mg
	Fenofibrate	300 mg

<sup>‡</sup>LDL-C, low-density lipoprotein cholesterol; HDL-C, high-density lipoprotein cholesterol; TG, triglyceride  
<sup>‡</sup>Clofibrate is not considered a first-line agent because of associated toxicity; fenofibrate is approved but not currently available in the United States