



USE OF THE **LIPOD OPTIMIZATION TOOL (LOT)** TO ACHIEVE VARIOUS LDL TARGETS IN COMMUNITY PRACTICE

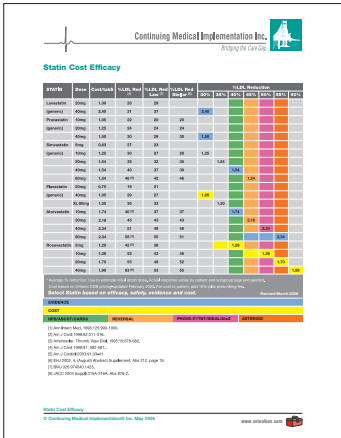
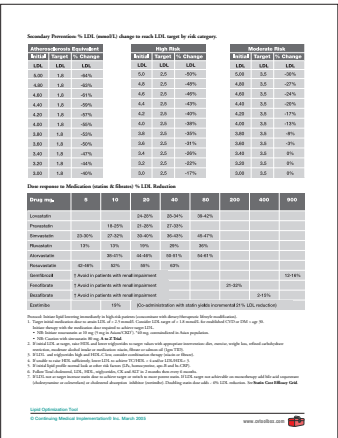
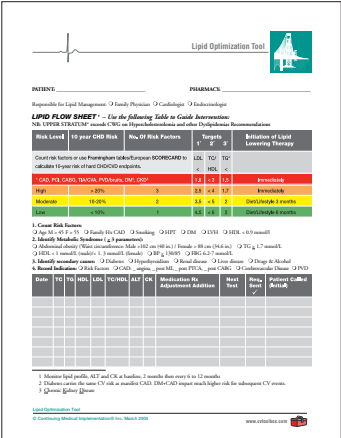
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Background (Objectives)

Lipid optimization has the potential to significantly reduce cardiovascular events and CHD death in various at risk populations. Failure to reach LDL target is commonplace and compromises cardiovascular outcomes. Our objective was to audit the effectiveness of the **Lipid Optimization Tool** as available on www.cvtoolbox.com, in achieving dyslipidemia control measured as LDL target achievement, in the high volume community practice at OCC.

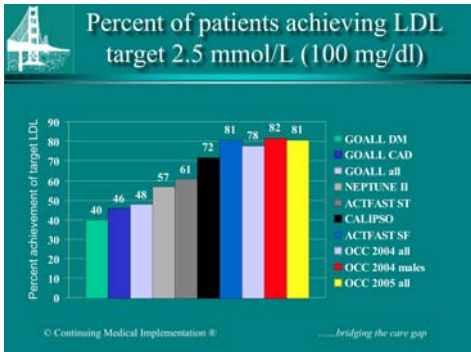
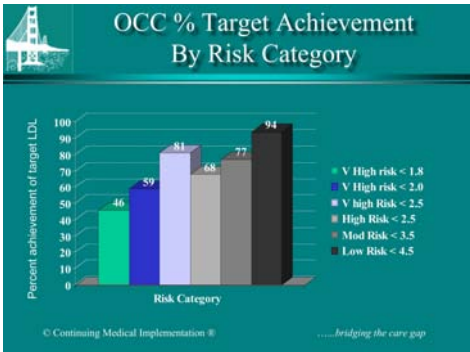
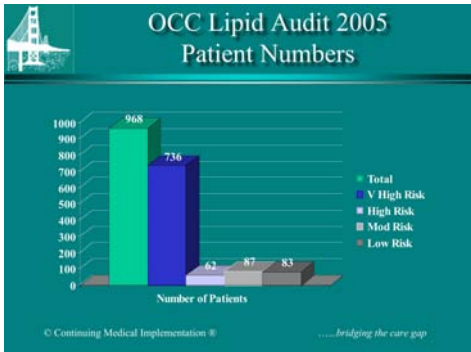
Methods

OCC provides comprehensive follow-up to approximately 27,000 cardiovascular patients and structured lipid management and Telemonitoring for approximately 7000 dyslipidemia patients. The LOT has been used to guide lipid management at OCC since 1999. LOT flow sheets for patients of the 9 OCC specialists (6 cardiologists and 3 internists) were sequentially analyzed from July to December 2005. LDL control rates were calculated for the current Canadian Working Group (CWG) targets of 4.5 mmol/L for low risk (LR), 3.5 mmol/L for moderate risk (MR) and 2.5 mmol/L for high risk (HR) patients as well as the anticipated revised CWG LDL HR target of 2.0 mmol/L and the optional very high risk (VHR) NCEP ATP-III LDL target of 1.8 mmol/L.



Results

Data was collected on 963 patients; 78 LR, 87 MR, 62 HR and 736 VHR. LDL control rates were LR 94% (target < 4.5 mmol/L), MR 77% (target < 3.5 mmol/L), HR 68% (target < 2.5 mmol/L) and VHR 81% (target < 2.5 mmol/L), 59% (target < 2.0 mmol/L) and 46% (target 1.8 mmol/L).



1) RT Yan et al. Guideline Oriented Approach in Lipid Lowering (GOALL) Registry data presented at CCC Symposium Oct 2005.
2) MH Davidson et al. Results of the National Cholesterol Education (NCEP) Program Evaluation Project Utilizing Novel e-Technology (NEPTUNE) II Survey and Implications for Treatment Under the Recent NCEP Writing Group Recommendations. Am J Cardiol 2005 Aug 15;96(4):556-63.
3) A Langer et al. Targeted Dosing of Atorvastatin Achieves Cholesterol Targets Quickly in Subjects with Diabetes or the Metabolic Syndrome (The ACTFAST Studies). Can J Cardiol 2005; Vol 21 Suppl C: Abstract 826, 253C.
4) C Bourgault et al. Statin Therapy in Canadian Patients with Hypercholesterolemia: The Canadian Lipid Study – Observational (CALIPSO). Can J Cardiol 2005; 21(13):1187-1193.

Conclusions

Use of the LOT management protocol in contemporary community practice at OCC achieves amongst the best lipid control rates in the world.

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PHARMACY:

ogist ○ Endocrinologist

LIPID FLOW SHEET¹ – Use the following Table to Guide Intervention:

NB: UPPER STRATUM* exceeds CWG on Hypercholesterolemia and other Dyslipidemias Recommendations

Risk Level	10 year CHD Risk	No. Of Risk Factors	Targets ^a	Initiation of Lipid Lowering Therapy
			1 ^b 2 ^c 3 ^d	
		Count risk factors or use Framingham tables/European SCORECARD to calculate 10-year risk of hard CHD/CVD endpoints.	LDL TG*	
			< HDL	<
			1.8 < 3	1.5
High	> 20%	3	2.5 < 4	1.7
Moderate	10-20%	2	3.5 < 5	2
Low	< 10%	1	4.5 < 6	2
				Immediately
				Immediately
				Diet/Lifestyle 3 months
				Diet/Lifestyle 6 months

- 1. Count Risk Factors:**
 ○ Age: $M \geq 45$ F > 55 ○ Family Hx CAD ○ Smoking ○ HPT ○ DM ○ LVH ○ HDL < 0.9 mmol/l
- 2. Identify Metabolic Syndrome (≥ 3 parameters):**
 ○ Abdominal obesity (Waist circumference: Male ≥ 102 cm (40 in.) / Female ≥ 88 cm (34.6 in.) ○ TG ≥ 1.7 mmol/L ○ HDL < 1 mmol/L (male) < 1.3 mmol/L (female) ○ BP $\geq 130/85$ ○ FPG ≥ 6.27 mmol/L
- 3. Identify secondary causes:**
 ○ Diabetes ○ Hypothyroidism ○ Renal disease ○ Liver disease ○ Drugs & Alcohol
- 4. Record Indications ○ Risk Factors**
 ○ CAD: angina, post MI, post PTCA, post CABG ○ Cerebrovascular Disease ○ PVD

[illegible]

- 1 Monitor lipid profile, ALT and CK at baseline, 2 months then every 6 to 12 months
- 2 Diabetes carries the same CV risk as manifest CAD. DM+CAD impart much higher risk for subsequent CV events.
- 3 Chronic Kidney Disease

Lipid Optimization Tool

Secondary Prevention: % LDL (mmol/L) change to reach LDL target by risk category.

Atherosclerosis Equivalent				High Risk			Moderate Risk		
Initial	Target	% Change		Initial	Target	% Change	Initial	Target	% Change
LDL	LDL	LDL		LDL	LDL	LDL	LDL	LDL	LDL
5.00	1.8	-64%		5.0	2.5	-50%	5.00	3.5	-30%
4.80	1.8	-63%		4.8	2.5	-48%	4.80	3.5	-27%
4.60	1.8	-61%		4.6	2.5	-46%	4.60	3.5	-24%
4.40	1.8	-59%		4.4	2.5	-43%	4.40	3.5	-20%
4.20	1.8	-57%		4.2	2.5	-40%	4.20	3.5	-17%
4.00	1.8	-55%		4.0	2.5	-38%	4.00	3.5	-13%
3.80	1.8	-53%		3.8	2.5	-35%	3.80	3.5	-8%
3.60	1.8	-50%		3.6	2.5	-31%	3.60	3.5	-3%
3.40	1.8	-47%		3.4	2.5	-28%	3.40	3.5	0%
3.20	1.8	-44%		3.2	2.5	-22%	3.20	3.5	0%
3.00	1.8	-40%		3.0	2.5	-17%	3.00	3.5	0%

Dose response to Medication (statins & fibrates) % LDL Reduction

Drug mg.	5	10	20	40	80	200	400	900
Lovastatin			24-28%	28-34%	39-42%			
Praavastatin		18-25%	21-28%	27-33%				
Sinvastatin	23-30%	27-32%	30-40%	36-43%	45-47%			
Fluvastatin	13%	13%	19%	29%	38%			
Atorvastatin		38-41%	44-46%	50-51%	54-61%			
Rosuvastatin	42-46%	52%	55%	63%				
Gemfibrozil	$\dagger$ Avoid in patients with renal impairment							
Fenofibrate	$\dagger$ Avoid in patients with renal impairment							
Bazalifbrate	$\dagger$ Avoid in patients with renal impairment							
Ezetimibe		19%	(Co-administration with statins yields incremental 21% LDL reduction)			21-32%	2-15%	12-16%

Protocol: Initiate lipid lowering immediately in high-risk patients (concomitant with dietary/therapeutic lifestyle modification).

1. Target initial medication dose to attain LDL of ≤ 2.5 mmol/L. Consider LDL target of ≤ 1.8 mmol/L for established CVD or DM + age ≥ 30 . Initiate therapy with the medication dose required to achieve target LDL.
2. If LDL is not attained at 10 mg (5 mg in Asian CCKD):
 - a. Niaspan: initiate treatment at 10 mg (5 mg in Asian CCKD), 40 mg contraindicated in Asian population.
 - b. Niaspan: target initial LDL and lower triglycerides in target values with appropriate intervention: diet, exercise, weight loss, refined carbohydrate restriction, moderate alcohol intake or medication: niacin, fibrate or statoin if (upon TTD).
 - c. TLDL and triglycerides high and HDL-C low: consider combination therapy (statoin or fibrate).
3. If LDL and triglycerides high and HDL-C low: target to achieve TG/HDL < 4 and/or LDL/HDL < 3 .
4. If initial lipid profile normal/low at other risk factors (LDL, homocysteine, apo-B, and/or CVD):
 - a. Consider statoin monotherapy, CKD and AHA ≥ 2 months from entry.
 - b. If LDL target not achievable on monotherapy, add lipid acid equivalent (obestatin or odestatin) or cholesterol absorption inhibitor (ezetimibe). Doubling statoin dose adds $\sim 6\%$ LDL reduction. See Statoin Cost Efficacy Grid.

ipid Optimization Tool

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Statin Cost Efficacy

STATIN	Dose	Cost/tab\$ ⁽¹⁾	%LDL Red ⁽¹⁾	%LDL Red Law ⁽²⁾	%LDL Red Stellar ⁽³⁾	%LDL Reduction						
						30%	35%	40%	45%	50%	55%	60%
Lovastatin	20mg	1.30	26	29								
	(generic)	40mg	2.40	31	37	2.40						
	Pravastatin	10mg	1.05	22	20	20						
	(generic)	20mg	1.25	24	24	24						
Simvastatin	40mg	1.50	30	29	30	1.50						
	5mg	0.63	27	23								
	10mg	1.25	30	27	28	1.25						
	(generic)	20mg	1.54	35	32	35	1.54					
Fluvastatin	40mg	1.54	40	37	39			1.54				
	80mg	1.54	48 ⁽²⁾	42	46				1.54			
	20mg	0.75	19	21								
	(generic)	40mg	1.05	29	27		1.05					
Atorvastatin	XL 80mg	1.30	36	33			1.30					
	10mg	1.74	40 ⁽³⁾	37	37			1.74				
	20mg	2.18	45	43	43				2.18			
	40mg	2.34	51	49	48					2.34		
Rosuvastatin	80mg	2.34	58 ⁽⁴⁾	55	51						2.34	
	5mg	1.29	42 ⁽³⁾	38					1.29			
	10mg	1.38	52	43	46						1.38	
	20mg	1.70	55	48	52							1.70
	40mg	1.99	63 ⁽³⁾	53	55							1.99

Select Statin based on efficacy, safety, evidence and cost.

EVIDENCE	COST		PROVE-IT/TNT/IDEAL/a10z	ASTEROID
HPS/JASCOT/CARDS	REVERSAL			

- [1] *Ann Intern Med.* 1996;125:990-1000.
- [2] *Am J Cardiol*:1998;82:311-316.
- [3] *Arterioscler Thromb Vasc Biol.* 1995;15:678-682.
- [4] *Am J Cardiol* 1998;81: 582-587.
- [5] *Am J Cardiol* 2003;91:33-41
- [6] *EHJ* 2002, 4 (August) Abstract Supplement: Abs 212, page 19.
- [7] *BMJ* 326 974040:1423.
- [8] *JACC* 2003 (suppl):315A-316A. Abs 875-2.

Statin Cost Efficacy

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