

WEEKLY EPIDEMIOLOGICAL REPORT

A publication of the Epidemiology Unit Ministry of Health

231, de Saram Place, Colombo 01000, Sri Lanka Tele: + 94 11 2695112, Fax: +94 11 2696583, E mail: epidunit@sltnet.lk Epidemiologist: +94 11 2681548, E mail: chepid@sltnet.lk Web: http://www.epid.gov.lk

Vol. 42 No. 08

14th – 20th February 2015

Hyperlipidaemia (Part I) This is the first of a series of two articles on Hy• A total chol perlipidemia. dL (5.17 mn

Background

Hyperlipidemia refers to increased levels of lipids (fats) in the blood, including cholesterol and triglycerides.

Although Hyperlipidemia does not cause symptoms, it can significantly increase the risk of developing cardiovascular disease; including disease of blood vessels supplying the heart (coronary artery disease), brain (cerebrovascular disease), and limbs (peripheral vascular disease). These conditions can in turn lead to chest pain, heart attacks, strokes and other problems. Because of these risks, treatment is often recommended for people with Hyperlipidemia.

Lipid Types

The term lipids includes cholesterol and triglycerides. There are many different types of lipids (also called lipoproteins). Blood tests can measure the level of lipoproteins. The standard lipid blood tests include a measurement of total cholesterol, LDL (low density lipoproteins), HDL (high density lipoproteins), and triglycerides.

Total cholesterol — A high total cholesterol level can increase the risk of cardiovascular disease. However, decisions about when to treat high cholesterol are usually based upon the level of LDL or HDL cholesterol, rather than the level of total cholesterol.

- A total cholesterol level of less than 200 mg/ dL (5.17 mmol/L) is normal.
- A total cholesterol level of 200 to 239 mg/dL (5.17 to 6.18 mmol/L) is borderline high.
- A total cholesterol level greater than or equal to 240 mg/dL (6.21 mmol/L) is high.

The total cholesterol level can be measured any time of day. It is not necessary to fast (avoid eating for 12 hours) before testing.

LDL cholesterol — The low density lipoprotein (LDL) cholesterol (sometimes called "bad cholesterol") is a more accurate predictor of cardiovascular disease than total cholesterol. Higher LDL cholesterol levels increase the risk of cardiovascular disease.

Most healthcare providers prefer to measure LDL cholesterol after you have not eaten (fasted) for 12 to 14 hours. A test to measure LDL in people who have not fasted is also available, although the results may differ slightly.

Some healthcare providers make decisions about how to treat lipids based on a goal LDL cholesterol. If the healthcare provider uses this strategy, the goal for LDL cholesterol will depend on several factors, including any history of cardiovascular disease and the risk of developing cardiovascular disease in the future. People at higher risk are often assigned a lower LDL cholesterol goal.

10-year risk of developing coronary artery disease-The 10-year risk score is based on in-

Contents	Page
1. Leading Article – Hyperlipidemia - (Part I)	1
2. Summary of selected notifiable diseases reported - $(07^{th} - 13^{th} February 2015)$	3
3. Surveillance of vaccine preventable diseases & AFP - (07 th – 13 th February 2015)	4

WER Sri Lanka - Vol. 42 No. 08

14th February 20th 2015

formation from the Framingham Heart Study, a large study that has followed participants, as well as their children and grandchildren, for greater than 50 years. The 10-year risk can be calculated for women and for men.

Triglycerides — High triglyceride levels are also associated with an increased risk of cardiovascular disease, although this association is not typically important once other risk factors are taken into account. Triglyceride levels are divided as follows:

- Normal less than 150 mg/dL (1.69 mmol/L)
- Borderline high 150 to 199 mg/dL (1.69 to 2.25 mmol/L)
- High 200 to 499 mg/dL (2.25 to 5.63 mmol/L)
- Very high greater than 500 mg/dL (5.65 mmol/L)

Triglycerides should be measured after fasting for 12 to 14 hours.

HDL cholesterol — Not all cholesterol is bad. HDL cholesterol is considered "good cholesterol" because it helps remove LDL cholesterol from the arteries Elevated levels of HDL cholesterol actually lower the risk of cardiovascular disease. A level greater than or equal to 60 mg/dL or 1.55 mmol/L is excellent, while levels of HDL cholesterol less than 40 mg/dL or 1.03 mmol/L are lower than desired. There are no treatments for raising HDL cholesterol that has been proven to reduce the risk of heart attacks and strokes.

Similar to total cholesterol, the HDL-cholesterol can be measured on any blood specimen. It is not necessary to be fasting.

Non-HDL cholesterol — Non-HDL cholesterol is calculated by subtracting HDL cholesterol from total cholesterol. Since total cholesterol and HDL cholesterol can be measured without fasting, so can non-HDL cholesterol. Non-HDL cholesterol is a good predictor of cardiovascular risk and is a better predictor of risk than LDL cholesterol in people with type 2 diabetes and in women.

An appropriate non-HDL cholesterol goal can be calculated by adding 30 mg/dL (0.78 mmol/L) to the LDL cholesterol goal. As discussed, the LDL cholesterol goal depends on a number of factors.

Lp(a) Cholesterol

Lp(a) is a genetic variation of LDL (bad) cholesterol. A high level of Lp(a) is a significant risk factor for the premature development of fatty deposits in arteries. Lp(a) isn't fully understood, but it may interact with substances found in walls of arteries and contribute to the buildup of fatty deposits.

Sources

High cholesterol and lipids, available at <u>http://</u> www.uptodate.com/contents/high-cholesterol-and-lipidshyperlipidemia-beyond-the-basics

Hyperlipidemia, available at <u>http://www.heart.org/HEARTORG/</u> <u>Conditions/Cholesterol/aboutCholesterol/</u> <u>Hyperlipidemia_UCM_434965_Article.jsp#</u>

Compiled by Dr. C U D Gunasekara of the Epidemiology Unit

District	MOH areas	No: Expected *	No: Received
Colombo	12	72	33
Gampaha	15	90	NR
Kalutara	12	72	NR
Kalutara NIHS	2	12	NR
Kandy	23	138	NR
Matale	12	72	NR
Nuwara Eliya	13	78	22
Galle	19	114	32
Matara	17	102	5
Hambantota	12	72	NR
laffna	11	66	0
Kilinochchi	4	24	0
Manner	5	30	30
/avuniya	4	24	0
Mullatvu	4	24	6
Batticaloa	14	84	0
Ampara	7	42	34
Trincomalee	11	66	NR
Kurunegala	23	138	18
Puttalam	9	54	60
Anuradhapura	19	114	NR
Polonnaruwa	7	42	0
Badulla	15	90	77
Moneragala	11	66	63
Rathnapura	18	108	42
Kegalle	11	66	19
Kalmunai	13	78	NR

To be Continued...

WER Sri Lanka - Vol. 42 No. 08

14th February 20th 2015

Table 1: Selected notifiable diseases reported by Medical Officers of Health

07th - 13th Feb 2015 (07th Week)

	1: 3		ecte	u III	oum		u	004								•					-	• •	0/m - 1		эт ге		eb 2015 (
WRCD	* č	31	23	38	4	15	23	25	œ	0	0	75	20	0	40	14	43	33	26	38	37	14	35	6	33	27	54	26
WF	*	69	47	62	96	85	77	75	92	100	100	25	80	100	60	86	57	67	74	62	63	86	65	91	67	73	46	74
nani-	B	0	0	0	H	2	0	0	40	15	0	0	0	0	1	0	0	0	15	0	29	6	0	9	ω	0	0	121
Leishmani- asis	A	0	0	0	0		0	0	4	ъ	0	0	0	0	0	0	0	0	2	0	2	7	0		0	0	0	17
litis	в	З	m	5	m	1	6	12	2	7	1	0	0	0	1	2	ε	1	4	2	9	8	8	2	4	6	0	96
Meningitis	۲	0	0	0	H	0	4	1	0	0	0	0	0	0	0	0	0	1	1	0		0	0		0	7	0	12
	в	52	13	27	58		11	42	11	40	18	0	0	2	0	5	34	5	65	9	19	20	19	19	4	27	24	522
Chickenpox	A	6	2	2	29	0	4	3	4	9	2	0	0	0	0	1	11	1	16	0	4	m		2	0	4		105
	B	1	0	-	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	2
Human Rabies	A	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0
Viral Hepatitis	B	11	21	5	48	m	21	1	ø	7	5	0	0	1	0	0	0	0	6	1	m	2	21	7	55	21	0	250
Hep <	۲	с	2	0	2	7		0	2		2	0	0	1	0	0	0	0	2	0	0		ω	2	0	2	0	31
s Fever	в	1	2	0	17	1	11	6	7	6	307	3	9	8	2	0	0	1	8	m	4	0	13	13	13	2	0	440
Typhus Fever	A	0	0	0	m	0	∞	1	m	m	24	0	0	0	1	0	0	0	0	0		0	4	7		0	0	51
	B	30	39	47	13	12	ы	35	17	27	7	0	9	8	2	1		4	56	6	63	28	ø	54	43	36		552
Leptospirosi s	A	ъ	~	2	0	0	0	ъ		m	1	0		0	0	0	0	0	e	0	4		7	2	2	2	0	41
bi Ding	B	11	2	9	0	0	0	9	0	19	4	25	H	2		0	0	22	0	0	ы	0	m	2		0	∞	118
Food Poisoning	∢	1	0	2	0	0	0	0	0	0	1	0	0	0	0	0	0	0	0	0	0	0	m		0	0	m	듺
Fever	в	8	4	9	8	2	m	1	4	2	73	2	4	9	1	1	0	8	с	0	0	m		ъ	7	16	0	168
Enteric Fever	◄	1		0	-		0	0		0	13	0	1	1	0	0	0	2		0	0	0	0	0			0	25
halit	в	1	2	-	0	0	0	0	0	0	4	0	0	2	ц.	1	0	0	1	0	0		0	0	m	2	0	19
Encephalit is	۲	0	0	0	0	0	0	0	0	0	1	0	0	0		0	0	0	0	0	0		0	0	0	0	0	m
	B	33	б	14	29	12	38	15	ъ	11	117	13	2	5	ъ	26	13	ε	34	6	15	7	30	28	69	12	24	578
Dysentery	◄	e		2	ω	9	ы	2	0	2	17	2	1	1	0	6	0	0	4	0	4	0	m		0		0	64
Fever	в	2387	982	344	349	216	51	190	65	104	727	19	59	41	45	477	11	156	414	293	162	64	223	59	210	132	267	8047
Dengue Fever	A	298	75	28	52	38	10	26	15	14	60	0	m	З	7	85	1	20	63	19	18	11	13	8	23	13	18	921 8
RDHS Division		Colombo	Gampaha	Kalutara	Kandy	Matale	NuwaraEliya	Galle	Hambantota	Matara	Jaffna	Kilinochchi	Mannar	Vavuniya	Mullaitivu	Batticaloa	Ampara	Trincomalee	Kurunegala	Puttalam	Anuradhapura	Polonnaruwa	Badulla	Monaragala	Ratnapura	Kegalle	Kalmune	SRILANKA

•T=Timeliness refers to returns received on or before 13th February, 2015 Total number of reporting units 337 Number of reporting units data provided for the current week: 253 C**-Completeness

Terriliane and an and a second a se

14th February 20th 2015

Table 2: Vaccine-Preventable Diseases & AFP

07th - 13th Feb 2015 (07th Week)

Disease			N	lo. of Cas	es by P	rovince			Number of cases during current	Number of cases during same	Total number of cases to date in	Total num- ber of cases to date in	Difference between the number of cases to date	
	W C S N F NW NC U Sab weeki							week in 2015	week in 2014	2015	2014	in 2014& 2015		
AFP*	00	00	00	00	00	00	01	01	00	02	01	09	09	%
Diphtheria	00	00	00	00	00	00	00	00	00	00	00	00	00	%
Mumps	01	00	02	00	01	01	00	02	01	08	10	51	128	-60.1%
Measles	16	01	05	00	01	02	01	02	01	29	40	189	365	-48.3%
Rubella	00	00	00	00	00	00	00	00	00	00	00	02	01	+100%
CRS**	00	00	00	00	00	00	00	00	00	00	00	00	00	%
Tetanus	00	00	00	00	00	00	00	00	00	00	00	02	02	%
Neonatal Teta- nus	00	00	00	00	00	00	00	00	00	00	00	00	00	%
Japanese En- cephalitis	00	00	00	01	00	00	00	00	00	01	02	03	13	-77%
Whooping Cough	00	00	00	01	00	01	01	00	00	03	01	13	08	62.5%
Tuberculosis	76	39	23	02	00	00	08	10	01	159	148	1241	1528	-18.8%

Key to Table 1 & 2

Provinces: W: Western, C: Central, S: Southern, N: North, E: East, NC: North Central, NW: North Western, U: Uva, Sab: Sabaragamuwa.

RDHS Divisions: CB: Colombo, GM: Gampaha, KL: Kalutara, KD: Kandy, ML: Matale, NE: Nuwara Eliya, GL: Galle, HB: Hambantota, MT: Matara, JF: Jaffna,

KN: Killinochchi, MN: Mannar, VA: Vavuniya, MU: Mulláitivu, BT: Batticaloa, AM: Ampara, TR: Trincomalee, KM: Kalmunai, KR: Kurunegala, PU: Puttalam, AP: Anuradhapura, PO: Polonnaruwa, BD: Badulla, MO: Moneragala, RP: Ratnapura, KG: Kegalle.

Data Sources:

Weekly Return of Communicable Diseases: Diphtheria, Measles, Tetanus, Neonatal Tetanus, Whooping Cough, Chickenpox, Meningitis, Mumps., Rubella, CRS, Special Surveillance: AFP* (Acute Flaccid Paralysis), Japanese Encephalitis

CRS** =Congenital Rubella Syndrome

AFP and all clinically confirmed Vaccine Preventable Diseases except Tuberculosis and Mumps should be investigated by the MOH

In	Influenza Surveillance in Sentinel Hospitals - ILI & SARI														
	1	Human			Animal										
IV	lonth	No Received	ILI	SARI	Infl A	Infl B	Pooled samples	Serum Samples	Positives						
Ja	nuary	4453	61	18	08	13	1130	873	0						

Source: Medical Research Institute & Veterinary Research Institute

PRINTING OF THIS PUBLICATION IS FUNDED BY THE WORLD HEALTH ORGANIZATION (WHO).

Comments and contributions for publication in the WER Sri Lanka are welcome. However, the editor reserves the right to accept or reject items for publication. All correspondence should be mailed to The Editor, WER Sri Lanka, Epidemiological Unit, P.O. Box 1567, Colombo or sent by E-mail to chepid@sltnet.lk. Prior approval should be obtained from the Epidemiology Unit before publishing data in this publication

ON STATE SERVICE

Dr. P. PALIHAWADANA CHIEF EPIDEMIOLOGIST EPIDEMIOLOGY UNIT 231, DE SARAM PLACE COLOMBO 10