

Update on the Management of Diabetes

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FOR THE 7TH ANNUAL OPHALMOLOGY SOCIETY OF JAMAICA'S CONFERENCE MARCH 19 2017

Prevalence of select NCDs in Jamaican, adults 15-74 years old

Disease Condition	JHLS-2000 (%)	JHLSII-2008 (%)
Diabetes Mellitus	7.2	1 7.9
Hypertension	20.9	^ 25.2
Pre-hypertension	29.9	† 35.3
High Cholesterol	14.6	↓ 11.7
Depression	_	20.3
Asthma (self-reported)	-	7.0

Mortality Figures for Jamaica

Table 55 Ten Leading Causes of Death among Males 5 Years Old and Over: 2011 and Comparative Figures for 2009 and 2010

ICD 10 Code	Cause of Death	2011	2010	2009
V01-Y89	External Causes	2,092	1,994	1,916
160-169	Cerebrovascular Diseases	989	396	920
E10-E14	Diabetes Mellitus	885	823	676
C61	Malignant Neoplasm of Prostate	616	590	587
20-125	Ischaemic Heart Diseases	557	563	537
10-114	Hypertensive Diseases	553	457	496
C33-C34	Malignant Neoplasm of the Larynx, Trachea, Bronchus and Lung	319	296	369
J40-J47	Chronic Lower Respiratory Diseases	301	317	320
B20-B24	Human Immunodeficiency Virus (HIV) Disease	285	293	256
126-151	Other Heart Diseases	283	343	314
	TOTAL	6,880	6,642	6,391

Source: Registrar General's Department

Notes: (i) External causes include sudden and violent cases reported by the police but not yet registered by the Registrar General's Department

Table 56	Ten Leading Causes of Death among Females 5 Years Old and Over: 201
	and comparative Figures for 2009 and 2010

ICD 10 Code	Cause of Death	2011	2010	2009
E10-E14	Diabetes Mellitus	1,381	1,224	1,103
160-169	Cerebrovascular Diseases	1,127	1,087	1,098
110-114	Hypertensive Diseases	588	622	602
20-125	Ischaemic Heart Diseases	529	481	541
126-151	Other Heart Diseases	380	287	336
C44-49, C51-52, C57-60, C62-66, C68-69, C73-81, C88, C96-97	Remainder of Malignant Neoplasm	356	334	352
C50	Malignant Neoplasm of the Breast	294	300	291
V01-Y89	External Causes	290	265	351
C53	Malignant Neoplasm of Cervix uteri	191	158	162
B20-B24	Human Immunodeficiency Virus (HIV) Diseases	178	201	156
	TOTAL	5,314	4,959	4,992

Source: Registrar General's Department

Notes: (i) External causes include sudden and violent cases reported by the police but not yet registered by the Registrar General's Department

Statistical Institute of Jamaica

DIABETIC COMPLICATIONS

COMPLICATIONS

STROKE / TIA

AMPUTATION

BLINDNESS ISCHAEMIC HEART DISEASE

RENAL INSUFFICIENCY









Prevalence of Micro/ Macrovascular complications



1. Mowatt L. Middle East Afr J Ophthalmol. 2013 Oct-Dec;20(4):321-6. doi: 10.4103/0974-9233.120017.

DCCT: Relative Risk of Developing Chronic Complications of Diabetes in Relation to HbA1c Relative Risk



Adapted by Skyler J, Endocr & Metab. N. Amer, 1996: 25:250

Mean Glucose Levels for Specified A1C Levels

		Mean Plasma	Glucose*	Mean Fasting Glucose	Mean Premeal Glucose	Mean Postmeal Glucose	Mean Bedtime Glucose
	A1C%	mg/dL	mmol/L	mmol/L	mmol/L	mmol/L	mmol/L
	6	126	7.0				
	<6.5			6.8	6.6	8.0	7.6
_							
	6.5-6.99			7.9	7.7	9.1	8.5
	7	154	8.6				
/	7.0-7.49			8.4	8.4	9.8	9.8
_	7.5-7.99			9.3	8.6	10.5	9.7
	8	183	10.2				
	8-8.5			9.9	9.9	11.4	12.3
	9	212	11.8				
	10	240	13.4				
	11	269	14.9				
	12	298	16.5				

These estimates are based on ADAG data of ~2,700 glucose measurements over 3 months per A1C measurement in 507 adults with type 1, type 2, and no diabetes. The correlation between A1C and average glucose was 0.92. A calculator for converting A1C results into estimated average glucose (eAG), in either mg/dL or mmol/L, is available at http://professional.diabetes.org/eAG.

ADA. 6. Glycemic Targets. Diabetes Care. 2015;38(suppl 1):S35; Table 6.1

TO PREVENT OR DELAY DIABETES COMPLICATIONS

Haemoglobin A1c

KEEP THE HBA1C TO LESS THAN 7 %

REDUCE THE CHRONIC COMPLICATIONS BY 25-30%

UKPDS, Lancet, 1998; DCCT, NCJM, 1993

For every percentage point decrease in HbA1c there is a :

25% reduction in diabetes-related deaths

7% reduction in all-cause mortality

18% reduction in combined fatal and nonfatal myocardial infarction

American Association of Clinical Endocrinologists and American College of Endocrinology Clinical Practice Guidelines for Developing a Diabetes Mellitus Comprehensive Care Plan

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ENDOCRINE PRACTICE Vol 21 No. 4 April 2015

Outpatient Glucose Targets for Nonpregnant Adults

Parameter	Treatment Goal		
	Individualize on the basis of age, comorbidities, duration of disease, and hypoalycemia risk:		
AIC, %	 In general, ≤6.5 for most* 		
	 Closer to normal for healthy Less stringent for "less healthy" 		
FPG, mmol/l	<6.1		
2-Hour PPG, mmol/l	<7.8		
FPG = fasting plasma glucose; PPG = postprandial glucose. *Provided taraet can be safely achieved.			



PREDIABETES ALGORITHM

IFG (100–125) | IGT (140–199) | METABOLIC SYNDROME (NCEP 2005)





Diet Nutrition Therapy: AIMS To achieve and maintain near normal blood glucose To achieve and maintain a healthy body weight To achieve and maintain a favourable blood lipid profile To minimise complications such as hypoglycaemia To provide appropriate nutrition / meeting

metabolic and growth needs

Caribbean Food and Nutrition Institute, PAHO/WHO 2004





Caribbean Food and Nutrition Institute, PAHO/WHO 2004

Diet

Parameter

• Eat regular meals and snacks; avoid fasting to lose weight

Treatment Goal

- Consume plant-based diet
- High in fiber
- Low calories
- Low glycemic index
- High in phytochemicals and or antioxidants
- Understand Nutrition Facts
- Understand Label information

Therapeutic Lifestyle Changes

Parameter	Treatment Goal
Weight loss For the overweight and obese patients	Reduce weight by 5% to 10%

Therapeutic Lifestyle Changes



Pharmacological Agents

Class	Primary Mechanism of Action	Agent(s)	Available as
α- Glucosidase inhibitors	 Delay carbohydrate absorption from intestine 	Acarbose Miglitol	GLUCOBAY
DOSAGE	• 50-100MG TDS		
SIDE EFFECT	• GI		

Garber AJ, et al. Endocr Pract. 2013;19(suppl 2):1-48. Inzucchi SE, et al. Diabetes Care. 2012;35:1364-1379.

Glucobay

- Affects postprandial hyperglycaemia
- Has some effect on Cardiovascular disease
- Lowers HBA1c by 0.8%
- Avoids hypoglycaemia
- Avoids weight gain
- Main side effect is flatulence (78% of patients) and diarrhoea (14% of patients)
- Not to be used in inflammatory bowel disease or bowel obstruction
- Not to be used in renal failure



http://www.glucobay.com/scripts/pages/en/professionals_home/key_facts/what_is_glucobay/index.php

Class	Primary Mechanism of Action	Agent(s)	Available as
Amylin analogue	 Decrease glucagon secretion Slow gastric emptying Increase satiety 	Pramlintide	Symlin SymlinP en 120, SymlinP en 60
SUBCUTANEOUS			
Garber AJ, et al. <i>Endocr Pr</i>	act. 2013;19(suppl 2):1-48. Inzucchi SE, et al. <i>Diabetes Care</i> . 2012;35:13	364-1379.	

Class	Primary Mechanism of Action	Agent(s)	Available as
Biguanide	 Decrease Hepatic Glucose Production (HGP) Increase glucose uptake in muscle 	Metformin	Glucophage or generic
SIDE EFFECTS	GILACTIC ACIDOSIS	NOT TO BE USED IN CKD CCF, AMI , LIVER FAILURE	

Garber AJ, et al. Endocr Pract. 2013;19(suppl 2):1-48. Inzucchi SE, et al. Diabetes Care. 2012;35:1364-1379.

Metformin

- May prevent <u>cardiovascular disease</u> in diabetes
- May prevent <u>cancer</u> complications in diabetes
- Low risk of hypoglycaemia
- Weight neutral
- Common side effects include <u>diarrhea</u>, <u>nausea</u> and abdominal pain
- 500mg to 2000mg OD to TDS
- DRUG OF CHOICE IN OVERWEIGHT / OBESE DIABETICS



Class	Primary Mechanism of Action	Agent(s)	Available as
Bile acid sequestrant	 Decrease HGP? Increase incretin levels? 	Colesevelam	WelChol
Garber AJ, et	al. Endocr Pract. 2013;19(suppl 2):1-48. Inzucchi SE, et al. Diabetes Care. 2012;	35:1364-1379.	

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Class	Primary Mechanism of Action	Agent(s)	Available as
DPP-4 inhibitors	 Increase glucose- dependent insulin secretion Decrease glucagon secretion 	Alogliptin Linagliptin Saxagliptin Sitagliptin Valdagliptin	Nesina Trajenta Onglyza Januvia Galvus

DPP-4 = dipeptidyl peptidase; HGP = hepatic glucose production.

Garber AJ, et al. Endocr Pract. 2013;19(suppl 2):1-48. Inzucchi SE, et al. Diabetes Care. 2012;35:1364-1379.

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I. Januvia 25-100 mg OD II. Galvus 50 mg BID III.Trajenta 5 mg OD IV.Risk of pancreatitis V.Risk of pancreatic cancer VI.Weight neutral VII.Low risk of hypoglycaemia VIII.Can be used in renal failure X.No risk of cardiovascular disease



Antidiabetic

Boehringer &

Class	Primary Mechanism of Action	Agent(s)	Available as
Dopamine- 2 agonist	 Activates dopaminergic receptors 	Bromocriptine	Cycloset

Garber AJ, et al. Endocr Pract. 2013;19(suppl 2):1-48. Inzucchi SE, et al. Diabetes Care. 2012;35:1364-1379.

Class	Primary Mechanism of Action	Agent(s)	Available as
Glinides	 Increase insulin secretion 	Nateglinide Repaglinide	Starlix Novonorm

Garber AJ, et al. Endocr Pract. 2013;19(suppl 2):1-48. Inzucchi SE, et al. Diabetes Care. 2012;35:1364-1379.

REPAGLINIDE 0.5-2.0mg TDS



NATEGLINIDE 60- 120 mg TDS



Class	Primary Mechanism of Action	Agent(s)	Available as
GLP-1 receptor agonists	 Increase glucose- dependent insulin secretion Decrease glucagon secretion Slow gastric emptying Increase satiety 	Albiglutide Dulaglutide Exenatide Exenatide XR Liraglutide	Tanzeum Trulicity Byetta Bydureon Victoza

Garber AJ, et al. Endocr Pract. 2013;19(suppl 2):1-48. Inzucchi SE, et al. Diabetes Care. 2012;35:1364-1379.

 Subcutaneous (SC) injection in the thigh, abdomen, or upper arm.

- 5 mcg administered twice daily (BID) at any time within the 60-minute period before the morning and evening meals
- Increased to 10 mcg twice daily after 1 month of therapy
- Associated with acute pancreatitis
- Not to be used in creatinine clearance <30 mL/min) or end-stage renal disease



Nausea (44%) , vomiting (13%), diarrhoea (13%) Decreased appetite Decrease HBA1c by 0.9% Decrease weight No hypoglycaemia







Dose 5-10 mg OD





100 - 300 mg OD Moderate renal impairment with an eGFR of 45 to less than 60 mL/min/1.73 m2.

https://www.drugs.com/dosage/invokana.html

Sulfonylureas Increase insulin secretion Glipizide Diamic MR Sulfonylureas Glipizide Amary Glucos Gliphenclamide Glypuride Glypuride Sulfonylureas Glypuride Glypuride Glipizide Glypuride Glypuride Glypuride Glypuride Glypuride Glypuride Glypuride Glypuride Glypuride Glypuride Glypuride Glipizide Glypuride Glypuride Glypuride Glypuride Glypuride <td>cron /l trol se b</td>	cron /l trol se b



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Glimepiride	
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Voie orale / Oral use	
100 comprimés sécables / scored tablets	capoli aventis

Class	Primary Mechanism of Action	Agent(s)	Available as
Thiazolidinediones	 Increase glucose uptake in muscle and fat Decrease HGP 	Pioglitazone Rosiglitazone	Actos Avandia
	 Decrease HGP 		

Garber AJ, et al. Endocr Pract. 2013;19(suppl 2):1-48. Inzucchi SE, et al. Diabetes Care. 2012;35:1364-1379.





Efficacy of Agents Available for T2D



AGI = α -glucosidase inhibitors; BCR-QR = bromocriptine quick release; Coles = colesevelam; DPP4I = dipeptidyl peptidase 4 inhibitors; FPG = fasting plasma glucose; GLP1RA = glucagon-like peptide 1 receptor agonists; Met = metformin; Mod = moderate; PPG = postprandial glucose; SGLT2I = sodium-glucose cotransporter 2 inhibitors; SU = sulfonylureas; TZD = thiazolidinediones.

*Mild: albiglutide and exenatide; moderate: dulaglutide, exenatide extended release, and liraglutide.

Continued on next slide

Side effects of Agents Available for T2D



Side - Effects of Agents Available for T2D

	Met	GLP1RA	SGLT2I	DPP4I	TZD	AGI	Coles	BCR- QR	SU/ Glinide	Insulin	Pram
Renal impair- ment/ GU	Contra- indicated in stage 3B, 4, 5 CKD	Exenatide contra- indicated CrCl <30 mg/mL	GU infection risk	Dose adjust- ment (except lina- gliptin)	May worsen fluid retention	Neutral	Neutral	Neutr al	Increased hypo- glycemia risk	Increased risks of hypo-glycemia and fluid retention	Neutral
GI adverse effects	Mod	Mod*	Neutral	Neutral*	Neutral	Mod	Mild	Mod	Neutral	Neutral	Mod
СНҒ	Neutral	Neutral	Neutral	Neutral†	Mod	Neutral	Neutral	Neutr al	Neutral	Neutral	Neutral
CVD	Possible benefit	Neutral	Neutral	Neutral	Neutral	Neutral	Neutral	Safe	Ś	Neutral	Neutral
Bone	Neutral	Neutral	Bone loss	Neutral	Mod bone loss	Neutral	Neutral	Neutr al	Neutral	Neutral	Neutral

Monotherapy, Dual The Therapy for T2D	rapy, and Triple
Monotherapy*	
Metformin	
GLP1RA	
SGLT2I	
DPP4I	
AGI	
TZD [†]	
SU/glinide [†]	

Monotherapy, Dual Therapy, and Triple Therapy for T2D
Dual therapy*
Metformin (or other first-line agent) plus
GLP1RA
SGLT2I
DPP4I
TZD [†]
Basal insulin [†]
Colesevelam
BCR-QR
AGI
SU/glinide [†]

COMBINATION THERAPY Metformin and













Triple Therapy for T2D

Triple therapy*
First- and second-line agent plus
GLP1RA
SGLT2I
TZD [†]
Basal insulin [†]
DPP4I
Colesevelam
BCR-QR
AGI
SU/glinide [†]



GLYCEMIC CONTROL ALGORITHM





PROGRESSION OF DISEASE

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Insulin Regimens

Insulin is required for survival in T1D

Physiologic regimens using insulin analogs should be used for most patients

Multiple daily injections (MDI)

1-2 injections basal insulin per day
Prandial insulin injections before each meal

Continuous subcutaneous insulin infusion (CSII)

 Insulin pump using rapid acting insulin analog Management of Diabetic Retinopathy

- Slow retinopathy progression by maintaining optimal control of
 - Blood glucose
 - Blood pressure
 - Lipids
- For active retinopathy, refer to ophthalmologist as needed
 - For laser therapy
 - For vascular endothelial growth factor therapy

SUMMARY



Thank you