

Clinical presentation and outcomes of Haitian youth with diabetes receiving continuity of care in a specialized clinic in Montrouis, Haiti



Dumas M-P, MD¹; Sainvil M, MD²; Altenor K, MD³; von Oettingen JE, MD PhD MMSc¹, for the DESIDE study group

¹McGill University Health Center – Research Institute, ²University of Massachusetts, ³Kay Mackenson Clinic, Haiti

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Retinopathy

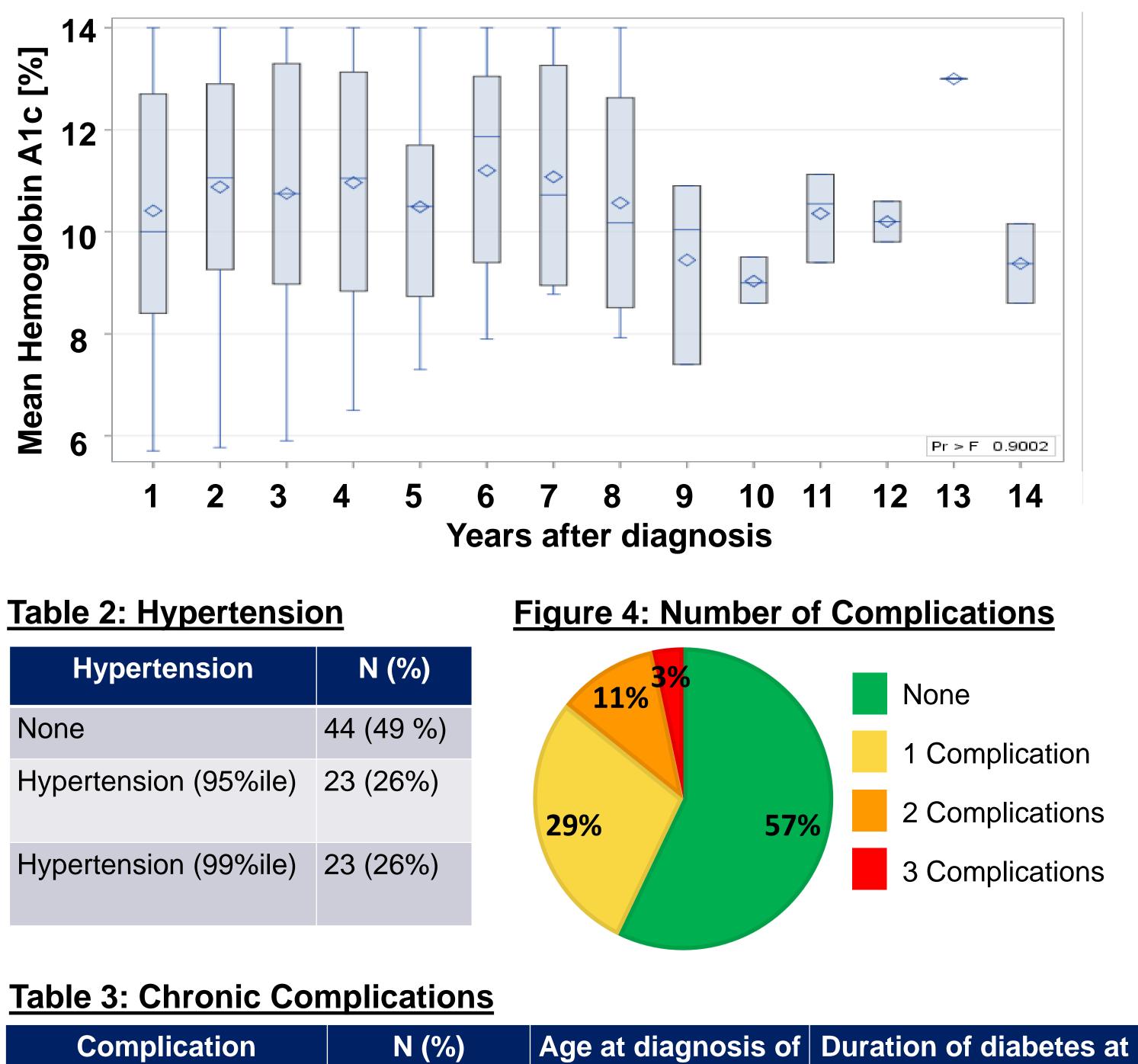
Introduction

Type 1 Diabetes – Haiti and low-income countries:

- Insufficient data on prevalence in these settings. (1)
- High risk of mortality, frequent diabetic ketoacidosis (DKA), high rates of early diabetes complications. (2-3)
- Clinical presentation and genetic markers for type 1 diabetes in non-Caucasian populations are not well described. (4)
- Clinical observation suggests that **phenotypes**, **auto-immune** and genetic markers may be distinct. (5)

Results (continued)

Figure 3: Mean A1c distribution over time



Objective

To describe the clinical presentation, glycemic control, and **chronic complications** of diabetes and their predictors in Haitian youth residing in Haiti.

Methods

Study group

DEterminants **S**ociaux et **I**ndividuels de santé en Diabète pEdiatrique (DESIDE)



Setting

- Kay Mackenson clinic in Montrouis, Haiti (second largest clinic for pediatric patients with non-communicable diseases in Haiti).
- Access to specialized care, insulin & medical supplies (Life for a) child supported) and education free of charge.

Study Design and Population

- Retrospective chart review between Jan 2013 and May 2018.
- Insulin-dependent diabetes in youth <25 years old at diagnosis.

Outcome measures:

- <u>Clinical phenotypes</u>: anthropometrics, symptoms, suspected diabetic ketoacidosis (DKA) / coma at diagnosis, yearly total daily insulin dose yearly after diagnosis
- Quarterly Hemoglobin A1c: proxy of glycemic control
- <u>Complications</u>: hypertension, chronic complications (cataract, retinopathy, nephropathy, neuropathy) and their predictors

Statistics:

• Predictors: descriptive statistics; linear and logistic regression to determine predictors of mean A1c and of complications

Results

Table 1: Baseline Characteristics

Variable	N (%) or Mean±SD	
Ν	91	
Female	55 (60%)	
Mean age [y]	18 ± 5	
Diabetes Duration [y]	4 ± 3.5	
Suspected ketoacidosis / coma at presentation	55 (60%) / 18 (20%)	
BMI z-score, at presentation	-1.7 ± 1.4	
BMI z-score, most recent	-0.9 ± 1.0	
Total daily insulin dose, most recent [IU/kg/day]	0.48 ± 0.28	

Cataract	13 (15.3 %)	16 (13.6, 18.9)	3.13 (1.6, 4.4)
Nephropathy	18 (24.3 %)	18.2 (16.7, 22.6)	3.3 (2.0, 7.7)
Microalbuminuria Macroalbuminuria	5 (20.3 %) 3 (4.1 %)		
Neuropathy	9 (13.6 %)	16.3 (15.8, 21.9)	2.8 (0.9, 5.8)
By Microfilament	9 (13.6 %)		
By Vibration Testing	1 (1.4 %)		

complication [y]

Mean (SD)

17.6 (16.1, 19.7)

complication [y]

Median (IQR)

3.06 (1.7, 4.4)

Table 4: Predictors of the presence of any chronic complication

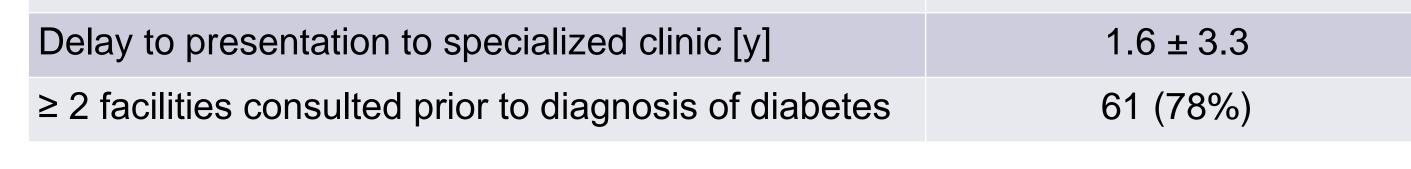
15 (17.4 %)

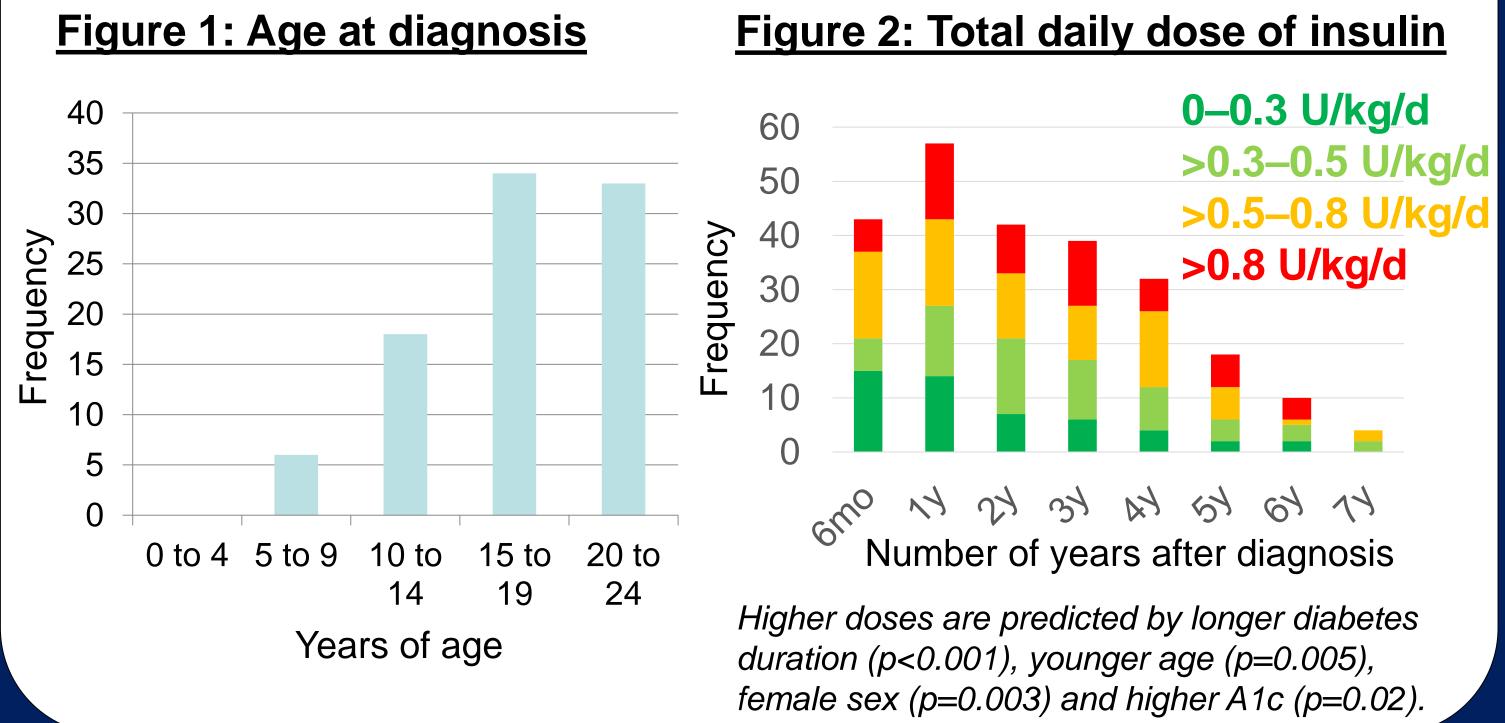
Predictor	Odds Ratio	95% Confidence Limits	P-value
Diabetes Duration	1.81	1.26 – 2.58	0.001
Initial BMI Z-score	0.64	0.41 - 0.99	< 0.05
DKA at diagnosis	3.71	1.13 – 12.24	0.03
Most recent Total Daily Insulin Dose (TDD)	1.02	0.81 – 1.28	0.88

Confounders: TDD is a confounder for diabetes duration and BMI z-score **Non-significant predictors** : Mean A1c, hypertension, age at diagnosis, sex

Conclusions

• Haitian youth with diabetes are **older at diagnosis**, present to care late, and often experience **DKA** and coma at diagnosis.





- Glycemic control is suboptimal, despite access to care.
- Diabetes complications occur frequently and precociously, are not predicted by mean A1c, and only partly by diabetes duration.
- Prolonged symptoms and extreme cachexia prior to diagnosis, and long periods of low insulin requirements post diagnosis may suggest prolonged exposure to hyperglycemia prior to diagnosis, predisposing to early complications.

Future directions

• Evaluation of the nutritional, psychosocial, and socioeconomic determinants of glycemic control, complications and quality of life. Validation of findings in prospective cohort study.

References

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