# Anaesthetic Management of Diabetic Child

# Essay

Submitted for Partial Fulfillment of the Master Degree in Anaesthesia

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#### **List of Abbreviations**

Abbr. Full-term

**ACR** : Albumin/creatinine ratio

**ADA** : American Diabetes Association

**BG** : Blood glucose

**BOHB** : Blood glucose and measure blood β-hydroxybutyrate

**cIMT** : Carotid intima-media thickness

**CKD** : Chronic kidney disease

**CSII** : Continuous subcutaneous insulin infusion

**CVD** : Cardiovascular disease

**DHC** : Diabetes healthcare

**DKA** : Diabetic ketoacidosis

**DN** : Diabetic neuropathy

**ECG** : Electrocardiogram

**eGFR** : Estimated GFR

**GA** : General anesthetic

**GAD** : Glutamic acid decarboxylase

**I-A2** : Insulinoma-associated antigen 2

**IAA** : Insulin auto antibodies

**ICA** : Islet cell antibodies

**ICU** : Intensive care unit

**IV** : Intravenous

# List of Abbreviations (Cont.)

Abbr. Full-term

**IFG** : Impaired fasting glycaemia

**IGT** : Impaired glucose tolerance

**IMCL** : Intramyocellular lipid content

**IV** : Intravenous

**LDL-C**: Low-density lipoprotein cholesterol

**MODY** : Maturity onset diabetes of the young

**NPH** : Neutral protamine Hagedorn

**NPO**: Nil per os

**OGTT** : Oral glucose tolerance test

**OHAs** : Oral hypoglycaemic agents

**PG**: Plasma glucose; SC, subcutaneous.

**PPI** : Preproinsulin

**T1D** : Type 1 diabetes

TCRs : T cell antigen receptors

**TDD** : Total daily dose

**TPN**: Total parenteral nutrition

**Treg** : T regulatory

**VDD** : Vitamin D deficiency

**ZnT8** : Zinc transporter

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#### Introduction

Diabetes mellitus is a group of metabolic diseases characterised by chronic hyperglycemia resulting from defects in insulin secretion, insulin action, or both. If ketones are present in blood or urine, treatment is urgent, because ketoacidosis can evolve rapidly (*Craig et al.*, 2009).

Type 1 diabetes results from the autoimmune destruction of insulin-producing  $\beta$  cells in the pancreas. Genetic and, as yet undefined, environmental factors act together to precipitate the disease. The excess mortality associated with the complications of type 1 diabetes and the increasing incidence of childhood type 1 diabetes emphasize the importance of therapeutic strategies to prevent this chronic disorder (*Kathleen, 2006*). Type 2 diabetes in youth was almost unheard of only two decades ago. However, tracking the recent dramatic rise in childhood obesity, type 2 diabetes has become increasingly prevalent (*Gemmill et al., 2011*).

The incidence of both type 1 and type 2 diabetes mellitus in children is increasing worldwide. The use of insulin pumps and various multicomponent insulin regimens has increased the complexity of perioperative management of children with diabetes. Anesthesiologists must carefully consider the pathophysiology of the disease, as well as each child's specific diabetes treatment regimen, glycemic control, intended surgery, and anticipated postoperative course, when devising an

appropriate perioperative management plan. Standardized algorithms for perioperative diabetes management improve care without significantly increasing costs (*Krane et al., 2013*).

When children with diabetes require surgery or other requiring procedures sedation or anaesthesia. optimal management should maintain adequate hydration and near to normal glycemia, while minimising the risk of hypoglycemia. The stress of surgery may cause acute hyperglycemia, which increases the risk of postoperative infection (Betts et al., 2009). The management of the pediatric patient before, during and after surgery or diagnosis is aimed at maintaining a normal glucose level; i.e. plasma sugar between 100 y 200 mg/dL. The goal is to keep the patient within this range to reduce the risk of osmotic diuresis, dehydration, hydroelectrolytic imbalance, metabolic acidosis, infection and hypoglycemia. The diabetic child scheduled for minor surgical interventions can be admitted on the same day of surgery, while those scheduled for major surgical procedures should be admitted one day before surgery (Ana et al., 2011).

Once the child is able to resume oral nutrition, resume the child's usual diabetes treatment regimen. Give short- or rapid-acting insulin (based on the child's usual insulin: carbohydrate ratio and correction factor), if needed, to reduce hyperglycemia or to match food intake (*Erinn et al.*, 2014).

# **Aim of the Study**

The aim of this study is to provide clear and standardised guidelines for all staff caring for paediatric patients with type 1 and 2 diabetes requiring a general anaesthetic or sedation for surgery.

# Chapter (1): Pathophysiology of Diabetes Mellitus

#### **Definition:**

Diabetes mellitus is a general term for heterogeneous disturbances of metabolism for which the main finding is chronic hyperglycaemia. The cause is either impaired insulin secretion or impaired insulin action or both. The definition includes the following criteria:

- HbA1c  $\geq$ 6.5% ( $\geq$ 48mmol/mol)
- Random plasma glucose ≥200 mg/dl (≥11.1 mmol/l)
- Fasting plasma glucose ≥126mg/dl (≥7.0mmol/dl)
- OGTT 2-hour glucose in venous plasma ≥200mg/dl (≥11.1mmol/l)

These guidelines have been recommending the use of HbA1c for diagnosing diabetes since 2010 (*Kerner et al.*, 2014).

# Classification

**Table (1):** Etiologic classification of diabetes mellitus (*Drouin et al., 2013*)

| ☐ Type 1 di   | abetes (β-cell destruction, usually leading to absolute insulin |
|---------------|---|
| deficienc     | y)  |
| A. Imn        | une mediated  |
| B. Idio       | pathic  |
|               | liabetes (may range from predominantly insulin resistance with  |
| relative insu | lin deficiency to a predominantly secretory defect with insuli- |
| resistance)   |   |
| □ O41         | if a town as  |
| □ Other spe   | * *   |
| A. A. C       | Senetic defects of β-cell function                              |
|               | 1. Chromosome 12, HNF-1α (MODY3)                                |
|               | 2. Chromosome 7, glucokinase (MODY2)                            |
|               | 3. Chromosome 20, HNF-4α (MODY1)                                |
|               | 4. Chromosome 13, insulin promoter factor-1 (IPF-1; MODY4)      |
|               | 5. Chromosome 17, HNF-1β (MODY5)                                |
|               | 6. Chromosome 2, NeuroD1 (MODY6)                                |
|               | 7. Mitochondrial DNA  |
|               | 8. Others   |
|               | etic defects in insulin action                                  |
|               | 1. Type A insulin resistance                                    |
|               | 2. Leprechaunism  |
|               | 3. Rabson-Mendenhall syndrome                                   |
|               | 4. Lipoatrophic diabetes  |
|               | 5. Others   |
| C. Dise       | ases of the exocrine pancreas                                   |
|               | 1. Pancreatitis   |
|               | 2. Trauma/pancreatectomy  |
|               | 3. Neoplasia  |
|               | 4. Cystic fibrosis  |
|               | 5. Hemochromatosis  |
|               | 6. Fibrocalculous pancreatopathy                                |
|               | 7. Others   |
|               | ocrinopathies   |
|               | 1. Acromegaly   |
|               | 2. Cushing's syndrome   |
|               | 3. Glucagonoma  |

#### Delitus (1): Pathophysiology of Diabetes Mellitus

4. Pheochromocytoma 5. Hyperthyroidism 6. Somatostatinoma 7. Aldosteronoma 8. Others E. Drug or chemical induced 1. Vacor 2. Pentamidine 3. Nicotinic acid 4. Glucocorticoids 5. Thyroid hormone 6. Diazoxide 7. β-adrenergic agonists 8. Thiazides 9. Dilantin 10. γ-Interferon 11. Others F. Infections 1. Congenital rubella 2. Cytomegalovirus 3. Others G. Uncommon forms of immune-mediated diabetes 1. "Stiff-man" syndrome 2. Anti-insulin receptor antibodies 3. Others H. Other genetic syndromes sometimes associated with diabetes 1. Down syndrome 2. Klinefelter syndrome 3. Turner syndrome 4. Wolfram syndrome 5. Friedreich ataxia 6. Huntington chorea 7. Laurence-Moon-Biedl syndrome 8. Myotonic dystrophy 9. Porphyria 10. Prader-Willi syndrome 11. Others Gestational diabetes mellitus