# **BOHR**

# Differences Between Hyperglycemic Hyperosmolar Non-Ketotic Syndrome and Diabetic Ketoacidosis

## Gudisa Bereda

*Department of Pharmacy, Negelle Health Science College, Negelle, Oromia regional state, Ethiopia E-mail: lionheart.bereda@yahoo.com* 

Abstract. Since high blood glucose levels might result from inadequate insulin levels, diabetic ketoacidosis is more common in those with insulin-dependent diabetes mellitus. Diabetic ketoacidosis develops only when there is insufficient insulin in the body to convert blood sugar into energy. The liver then uses the acids created by this process, known as ketones, to breakdown fat for energy. Osmotic diuresis, which results in significant amounts of urine production and volume depletion, as well as dehydration, occurs when an excessive amount of glucose enters the renal tubules of an individual with diabetes. Non-insulin-dependent diabetes mellitus can be more likely to cause hyperglycemic hyperosmolar nonketotic syndrome. More often than not, people with non-insulin-dependent diabetes mellitus who do not have their diabetes under control will experience hyperglycemic and hyperosmolar non-ketotic syndrome. The majority of cases of hyperglycemic hyperosmolar non-ketotic syndrome occur in people with non-insulin-dependent diabetes mellitus who also have another condition that reduces fluid intake.

**Keywords:** Diabetic ketoacidosisis, hyperglycemic hyperosmolar non-ketotic syndrome, insulin-dependent diabetes mellitus, non-insulin-dependent diabetes mellitus.

# INTRODUCTION

The primary characteristics of diabetic ketoacidosis are ketoacidosis and hyperglycemia, whereas hyperglycemia in hyperglycemic hyperosmolar non-ketotic syndrome is typically more severe than diabetic ketoacidosis but lacks ketoacidosis [1]. The range of 285–295 mOsm/L is considered typical for plasma osmolality [2]. The differences between hyperglycemic hyperosmolar non-ketotic syndrome and diabetic ketoacidosis are explored in turn below.

# DIABETIC KETOACIDOSIS

As a result of excessively high blood glucose levels, which might result from inadequate insulin, diabetic ketoacidosis is more common in people with insulin-dependent diabetes mellitus [3, 4]. When the body produces high quantities of blood acids called ketones, extremely high blood sugar, and low insulin levels, it can lead to a dangerous complication of diabetes called diabetic ketoacidosis [5]. Due to its severe, quick, and 24-h rapid onset, diabetes-related ketoacidosis is regarded as an acute complication [6]. Blood sugar levels are greater than or equal to 300 mg/dl in diabetic ketoacidosis [7]. Diabetic ketoacidosis only develops when there is insufficient insulin in the body to convert blood sugar into energy. The acids produced by this process-known as ketonesare subsequently used by the liver to digest fat for energy. Diabetic ketoacidosis has a plasma osmolality of less than 320 mOsm/L. In diabetic ketoacidosis, plasma osmolality is frequently increased to more than 290 mOsm/L) [8]. The arterial blood's pH is less than or equal to 7.3 when a person has diabetic ketoacidosis. Although a milder form of diabetic ketoacidosis may present with a bicarbonate level between 15 and 18 mmol/L, a lower PH is frequently associated with a fall in bicarbonate to 15 mmol/L or less [9]. The relationship between blood pH and the partial pressure of carbon dioxide in diabetic ketoacidosis is shown in Figure 1 [10]. In diabetic ketoacidosis, the blood ketone level was extremely high [11]. High blood glucose levels are caused by glucose that cannot enter the cells, which builds up in diabetic ketoacidosis [12]. Diabetic ketoacidosis also causes abnormally high urine ketone levels. Osmotic diuresis, which causes a high amount of urine to be produced and causes volume depletion and





Figure 1. Pathophysiology of diabetic ketoacidosis.

dehydration, occurs when an excessive amount of glucose reaches the renal tubules in diabetic ketoacidosis [13]. Lack of insulin causes too many ketones to accumulate in the blood and eventually "spill over" into the urine [14]. The production of ketone bodies in the liver increases when people are fasting or have medical disorders like diabetes mellitus that cause the body to produce more ketone bodies than it can use. When people urinate, the body tries to get rid of them, which causes ketonuria, or excessive ketone levels in urine [15]. Diabetic ketoacidosis causes the breath to smell sweet or like acetone (nail polish), and the urine to smell like rotting apples [17]. Urine that smells sweet or fruity is an indication of diabetes mellitus or high blood sugar. The sweetness in the urine is caused by sugar and indicates the body is attempting to eliminate surplus sugar in the blood [18]. Normally nonexistent sugar and ketones can build up in the urine in advanced insulin-dependent diabetes mellitus and produce a pungent odor [19]. When the body breaks down fatty acids for energy, ketosis is formed. The liver then excretes ketones as waste products, compromising acetone [20]. Because the body is releasing acetone as fat is broken down, the breath may smell nicer [21]. Low potassium levels, also known as hypokalemia; cerebral edema, which is an enlargement of the brain; and pulmonary edema, which is an enlargement of the lungs, are complications of diabetic ketoacidosis [22] (Figure 1).

### HYPERGLYCEMIC HYPEROSMOLAR NON-KETOTIC SYNDROME

Non-insulin-dependent diabetes mellitus may be more likely to cause hyperglycemic hyperosmolar nonketotic syndrome [23]. People with diabetes mellitus have excessive blood levels of glucose (sugar). Their bodies either do not produce enough insulin or have difficulties using the insulin they do produce, which causes the blood glucose to rise. More frequently, those with non-insulin-dependent diabetes mellitus who do not have their diabetes under control will experience hyperglycemic, hyperosmolar, nonketotic syndrome [24]. The majority of cases of hyperglycemic hyperosmolar non-ketotic syndrome are seen in people with non-insulin-dependent diabetes mellitus who also have another concurrent condition that reduces fluid intake [25]. Hyperglycemic hyperosmolar non-ketotic syndrome develops slowly rather than suddenly [26]. The blood sugar level in the hyperglycemic hyperosmolar non-ketotic syndrome is greater than or equivalent to 600 mg/dl. When blood glucose levels are extremely high, excess glucose is excreted in the urine [27]. In the



Figure 2. Pathophysiology of hyperglycemic hyperosmolar non-ketotic syndrome.

hyperglycemic hyperosmolar non-ketotic condition, the plasma osmolality exceeds 320 mOsm/L. Higher blood osmolality in hyperglycemic hyperosmolar non-ketotic syndrome is associated with significant impairment of the state of consciousness [28]. The PH of arterial blood is greater than or equal to 7.3 in hyperglycemic hyperosmolar non-ketotic syndrome. Acidosis is mostly brought on by dehydration and impaired end-organ perfusion in hyperglycemic hyperosmolar non-ketotic syndrome [29]. Hypernatremia is haphazardly associated with a water deficit in people with hyperglycemic hyperosmolar non-ketotic syndrome because of osmotic diuresis-induced hypotonic losses, which cause a loss of water greater than a loss of sodium. Because blood ketones increased when blood glucose levels were high due to a lack of insulin, which is necessary to allow glucose to enter the cells for energy, the hyperglycemic hyperosmolar non-ketotic syndrome showed little or no change in blood ketones [30]. Ketosis may not occur in non-insulin-dependent diabetes mellitus patients; ketones may not be produced, or the absence of ketosis may be caused by a relative rather than an absolute lack of insulin, which inhibits the production of ketones. The body attempts to eliminate extra glucose in the urine more often as blood glucose levels rise, which exacerbates dehydration [31]. Ketone levels are frequently normal or only slightly elevated because the pancreas produces just enough insulin to maintain fat in fat cells and prevent ketone production [32]. In the hyperglycemic, hyperosmolar, non-ketotic condition, the urine ketone level was normal or slightly elevated [33]. In the hyperglycemic, hyperosmolar, non-ketotic condition, breath and urine odors are normal [34]. The hyperglycemic, hyperosmolar, non-ketotic condition can be complicated by shock, blood clot development, cerebral edema (swelling of the brain), an elevated blood acid level, or lactic acidosis [35] (Figure 2).

#### CONCLUSION

When the body produces excessive amounts of blood acids called ketones, extremely high blood sugar, and low insulin levels, it can result in diabetic ketoacidosis, a serious complication of the disease. High blood glucose levels occur as a result of the inability of glucose to enter the cells in diabetic ketoacidosis. In non-insulin-dependent diabetes mellitus, hyperglycemic hyperosmolar non-ketotic syndrome is more common. In contrast to a sudden onset, the hyperglycemic hyperosmolar non-ketotic syndrome develops gradually.

#### **COMPETING INTERESTS**

The author has no financial or proprietary interest in any of material discussed in this article.

#### REFERENCES

- Bereda G. Case Report: Diabetic Ketoacidosis during Pregnancy Due to Insulin Omission. Open Access Emergency Medicine 2022:14; 615– 618.
- [2] Rasouli M. Basic concepts and practical equations on osmolality: biochemical approach. Clinical biochemistry. 2016 Aug 1; 49(12):936– 41.
- [3] Muñoz C, Floreen A, Garey C, Karlya T, Jelley D, Alonso GT, McAuliffe-Fogarty A. Misdiagnosis and diabetic ketoacidosis at diagnosis of type 1 diabetes: patient and caregiver perspectives. Clinical Diabetes. 2019 Jul 1; 37(3):276–81.
- [4] Balaji R, Duraisamy R, Kumar MP. Complications of diabetes mellitus: A review. Drug Invention Today. 2019 Jan 15; 12(1).
- [5] Dhatariya KK, Glaser NS, Codner E, Umpierrez GE. Diabetic ketoacidosis. Nature Reviews Disease Primers. 2020 May 14; 6(1):1–20.
- [6] Holt RI, DeVries JH, Hess-Fischl A, Hirsch IB, Kirkman MS, Klupa T, Ludwig B, Nørgaard K, Pettus J, Renard E, Skyler JS. The management of type 1 diabetes in adults. A consensus report by the American Diabetes Association (ADA) and the European Association for the Study of Diabetes (EASD). Diabetes Care. 2021 Nov; 44(11):2589–625.
- [7] Ayub A, Ijaz S, Qudrat SM, Rani T, Raziq A, Ali M, Butt TA. The Impact of COVID 19 Pandemic on Type 1 Diabetes Mellitus: An experience of a Tertiary Care Hospital in a resource limited country. ESPE Abstracts. 2021 Sep 6; 94.
- [8] Bereda G. Diabetic Ketoacidosis: Precipitating Factors, Pathophysiology, and Management. Biomed J Sci & Tech Res, 2022; 44(5): 35843–35848. BJSTR. MS.ID.007105.
- [9] Mahler SA, Conrad SA, Wang H, Arnold TC. Resuscitation with balanced electrolyte solution prevents hyperchloremic metabolic acidosis in patients with diabetic ketoacidosis. The American journal of emergency medicine. 2011 Jul 1; 29(6):670–4.
- [10] Bereda G. Clinical Management of Gestational Diabetes Mellitus. Journal of Diabetic Nephropathy and Diabetes Management. 2022;1(1):1–10.
- [11] Karrar HR. Diabetic Ketoacidosis: A Review Article. World Family Medicine. 2022; 20(6):66–71.
- [12] Dwivedi M, Pandey AR. Diabetes mellitus and its treatment: An overview. J. Adv. Pharmacol. 2020; 1:48–58.
- [13] Ravindran S, Munusamy S. Renoprotective mechanisms of sodiumglucose co-transporter 2 (SGLT2) inhibitors against the progression of diabetic kidney disease. Journal of Cellular Physiology. 2022 Feb; 237(2):1182–205.
- [14] MacFarlane B. Need to know vs good to know: The glucuretics. AJP: The Australian Journal of Pharmacy. 2016 Mar; 97(1148):92–5.
- [15] De Cabo R, Mattson MP. Effects of intermittent fasting on health, aging, and disease. New England Journal of Medicine. 2019 Dec 26; 381(26):2541–51.
- [16] Panova TI, Bortnikova AK. Ketosis level as a factor determining addictive behavior of Alcoholized rats. Neurophysiology. 2016 Aug; 48(4):252–8.

- [17] Warren C. What the dog knows: scent, science, and the amazing ways dogs perceive the world. Simon and Schuster; 2015 Mar 10.
- [18] Karamanou M, Protogerou A, Tsoucalas G, Androutsos G, Poulakou-Rebelakou E. Milestones in the history of diabetes mellitus: The main contributors. World journal of diabetes. 2016 Jan 1; 7(1):1.
- [19] Vistoli G, De Maddis D, Cipak A, Zarkovic N, Carini M, Aldini G. Advanced glycoxidation and lipoxidation end products (AGEs and ALEs): an overview of their mechanisms of formation. Free radical research. 2013 Aug 1; 47(sup1):3–27.
- [20] van der Kolk JH, Gross JJ, Gerber V, Bruckmaier RM. Disturbed bovine mitochondrial lipid metabolism: A review. Veterinary quarterly. 2017 Jan 1; 37(1):262–73.
- [21] Mathew TL, Pownraj P, Abdulla S, Pullithadathil B. Technologies for clinical diagnosis using expired human breath analysis. Diagnostics. 2015 Feb 2; 5(1):27–60.
- [22] Bereda. Risk Factors, Complications and Management of Diabetes Mellitus. Am J Biomed Sci & Res. 2022; 16(4): 409–412. AJBSR.MS.ID.002245.
- [23] Tittel SR, Sondern KM, Weyer M, Poeplau T, Sauer BM, Schebek M, Ludwig KH, Hammer F, Fröhlich-Reiterer E, Holl RW. Multicentre analysis of hyperglycaemic hyperosmolar state and diabetic ketoacidosis in type 1 and type 2 diabetes. Actadiabetologica. 2020 Oct; 57(10):1245–53.
- [24] Fayfman M, Pasquel FJ, Umpierrez GE. Management of hyperglycemic crises: diabetic ketoacidosis and hyperglycemic hyperosmolar state. Medical Clinics. 2017 May 1; 101(3):587–606.
- [25] Garg SK, Peters AL, Buse JB, Danne T. Strategy for mitigating DKA risk in patients with type 1 diabetes on adjunctive treatment with SGLT inhibitors: a STICH protocol. Diabetes technology & therapeutics. 2018 Sep 1; 20(9):571–5.
- [26] Bereda G, Bereda G. The Incidence and Predictors of Poor Glycemic Control among Adults with Type 2 Diabetes Mellitus in Ambulatory Clinic of Mettu Karl Referral Hospital, South Western, Ethiopia: A Prospective Cross Sectional Study. Int Arch Endocrinol Clin Res. 2021; 7:024.
- [27] Hensen J. Diabetic Coma: Current Therapy of Diabetic Ketoacidosis and Non-Ketoacidotic Hyperosmolar Coma. InType 2 Diabetes 2016 Apr 19 (pp. 251–264). CRC Press.
- [28] Rosenbloom AL. Hyperglycemic comas in children. InPediatric Endocrinology 2006 Dec 26 (pp. 181–194). CRC Press.
- [29] Whitmore SP, Gunnerson KJ. Acid-Base and Electrolyte Disorders in Emergency Critical Care. InEmergency Department Critical Care 2020 (pp. 301–329). Springer, Cham.
- [30] Dhatariya KK, Vellanki P. Treatment of diabetic ketoacidosis (DKA)/hyperglycemic hyperosmolar state (HHS): novel advances in the management of hyperglycemic crises (UK versus USA). Current diabetes reports. 2017 May; 17(5):1–7.
- [31] Bereda G. Hyperosmolar Hyperglycemic State: Background, Precipitating Factors, Pathophysiology and Management. In J Dia It Compl. 2022; 1 (1): 1–6.
- [32] Kolb H, Kempf K, Röhling M, Lenzen-Schulte M, Schloot NC, Martin S. Ketone bodies: From enemy to friend and guardian angel. BMC medicine. 2021 Dec; 19(1):1–5.
- [33] Pasquel FJ, Umpierrez GE. Hyperosmolar hyperglycemic state: a historic review of the clinical presentation, diagnosis, and treatment. Diabetes care. 2014 Nov 1; 37(11):3124–31.
- [34] Bereda G. Complication of Diabetes Mellitus: Microvascular and Macrovascular Complications. Int J Diabetes, 2022; 3(1): 123–128
- [35] Wolfsdorf JI, Glaser N, Agus M, Fritsch M, Hanas R, Rewers A, Sperling MA, Codner E. ISPAD Clinical Practice Consensus Guidelines 2018: Diabetic ketoacidosis and the hyperglycemic hyperosmolar state. Pediatric diabetes. 2018 Oct; 19:155–77.