



ELSEVIER

Contents lists available at ScienceDirect

Data in Brief

journal homepage: www.elsevier.com/locate/dib

Data Article

Data for serum 1,5 anhydroglucitol concentration in different populations



Marciane Welter ^a, Kátia C. Boritza ^a,
 Mauren I. Anghebem-Oliveira ^b, Railson Henneberg ^b,
 Aline B. Hauser ^b, Fabiane G.M. Rego ^{a,b,*}, Geraldo Picheth ^{a,b}

^a Post Graduate Program in Pharmaceutical Sciences, Federal University of Parana, Curitiba, Parana, Brazil

^b Department of Clinical Analysis, Federal University of Parana, Rua Prefeito Lothário Meissner, 632, 80210-170 Curitiba, Parana, Brazil

ARTICLE INFO

Article history:

Received 18 July 2018

Received in revised form

10 August 2018

Accepted 27 August 2018

Available online 1 September 2018

ABSTRACT

1,5 anhydroglucitol (1,5-AG), is a nonmetabolized 1-deoxy form of glucose, originate mainly from the diet. 1,5-AG is a biomarker to detect and magnify hyperglycemic excursions (postprandial hyperglycemia) in diabetic patients. Concentrations of 1,5-AG has been applied as supporting biomarker to diagnosis of the major forms of diabetes (type 1, type 2, and gestational). The serum 1,5-AG reference interval is relevant to the appropriate clinical application of this biomarker. This article contains data regards to serum concentration of the biomarker primarily for healthy subjects, capture from the literature, in different populations. Correlation analysis between 1,5-AG and markers associated with diabetes and its complication were presented. The data was complementary to the study “Reference intervals for serum 1,5-anhydroglucitol in children, adolescents, adults, and pregnant women” (Welter et al., 2018). The data present in this article improve the comparisons for 1,5-AG in different conditions and methodologies.

© 2018 The Authors. Published by Elsevier Inc. This is an open access article under the CC BY license (<http://creativecommons.org/licenses/by/4.0/>).

DOI of original article: <https://doi.org/10.1016/j.cca.2018.07.018>

* Corresponding author at: Department of Clinical Analysis, Federal University of Parana, Rua Prefeito Lothário Meissner, 632, 80210-170 Curitiba, Parana, Brazil.

E-mail address: rego@ufpr.br (F.G.M. Rego).

<https://doi.org/10.1016/j.dib.2018.08.165>

2352-3409/© 2018 The Authors. Published by Elsevier Inc. This is an open access article under the CC BY license (<http://creativecommons.org/licenses/by/4.0/>).

Specifications Table

Subject area	Clinical laboratory
More specific subject area	<i>Biomarkers for clinical chemistry associated with diabetes</i>
Type of data	<i>Tables</i>
How data was acquired	<i>Capture from literature data for 1,5-AG concentration and methodology. Comparison reference interval data was obtained with enzymatic colorimetric assay (Glycomark) measured in automate system (LabMax 400, Labtest, Brazil)</i>
Data format	<i>Analyzed</i>
Experimental factors	<i>Blood samples (serum or plasma EDTA) measured with different methodologies</i>
Experimental features	<i>Compilation, calculations, analysis (descriptive statistics) and comparison of literature data</i>
Data source location	<i>Federal University of Parana, Curitiba, Brazil.</i>
Data accessibility	<i>Data is in published papers and in this article.</i>
Related research article	<i>M. Welter, K.C. Boritza, M.I. Anghebem-Oliveira, R. Henneberg, A.B. Hauser, F.G.M. Rego, G. Picheth, Reference intervals for serum 1,5-anhydroglucitol in children, adolescents, adults, and pregnant women, CCA (2018) [1].</i>

Value of the data

- Data will facilitate the comparison of 1,5-AG in different studies.
- The data showed the correlation among 1,5-AG and relevant parameters associated with diabetes.
- The data provide comparison between 1,5-AG reference interval in different ethnicities, ages, gender and methodologies.
- These data provide information to researchers and clinical laboratory professionals to improve the 1,5-AG diagnostic use.

1. Data

In this article, we provide complementary data (urea, total protein and lipid profile) of the studied groups (Table 1), correlation analysis (Table 2) and comparisons from the literature for serum 1,5-anhydroglucitol (1,5-AG) concentration, to our study. We proposed a reference interval (Tables 3 and 4) for this biomarker in children, adolescent, adults and pregnant women [2].

Table 1

Complementary laboratory characteristic of studied groups.

Parameters	Children n = 580	Adolescents n = 496	Adults n = 922	Pregnant women n = 305
Sex (M/F)	242/338	192/304	460/462	–
Urea, mmol/L	3.8 (3.1–4.5)	3.8 (3.2–4.3)	4.3 (3.5–5.3)	3.4 (2.8–4.3)
Total Protein, g/L	81 (74–89)	78 (71–85)	71 (67–74)	69 (64–74)
Cholesterol, mmol/L	4.1 (3.6–4.6)	3.9 (3.4–4.4)	4.4 (4.0–5.4)	5.2 (4.1–6.2)
HDL-c, mmol/L	1.4 (1.2–1.6)	1.3 (1.0–1.5)	1.4 (1.1–1.5)	1.2 (1.0–1.6)
LDL-c, mmol/L	2.1 (1.7–2.5)	2.1 (1.7–2.6)	2.5 (2.0–3.1)	3.0 (2.5–4.1)
Triglycerides, mmol/L	1.1 (0.8–1.5)	0.8 (0.6–1.1)	1.4 (1.0–2.0)	1.3 (1.0–2.0)

Values are median (25–75%; interquartile range); M, male and F, female

HDL-c, high density lipoprotein-cholesterol; LDL-c, low density lipoprotein-cholesterol

Abbreviations: BMI, Body mass index; n, sample size.

Table 2

Significative ($P < 0.05$) Spearman rank order correlation of 1,5 anhydroglucitol with glycemia, HbA1c, age, body mass index (BMI) and creatinine.

Groups	Sex	1,5-AG correlation (R)				
		Glycemia	HbA1c	Age	BMI	Creatinine
Children (0–14 y)	Male	NS	NS	0.133	NS	NS
	Female	NS	0.128	NS	NS	0.163
Adolescents (14–18 y)	Male	NS	0.221	0.153	0.151	NS
	Female	NS	NS	0.221	NS	NS
Adults (≥ 18 y)	Male	NS	NS	–0.144	NS	NS
	Female	NS	NS	–0.102	NS	NS
Combining all		NS	NS	–0.310	–0.187	0.112
Pregnant women	Gestation weeks					
n = 110	< 23 weeks	NS	–	NS	NS	NS
n = 106	24–28 weeks	NS	–	NS	NS	NS
n = 52	29–32 weeks	NS	–	NS	NS	0.402
n = 37	> 32	NS	–	NS	NS	NS
Combining pregnant		0.095	–	–0.155	NS	NS

NS, non-significant; –, data no available.

Y, years old.

We studied healthy Euro-Brazilian subjects, classified as children (0–14 years old), adolescent (> 14 and < 18 years old) and adults (≥ 18 years old). Additionally, we analyzed pregnant women in four gestational periods, < 23 weeks; 24–28 weeks, 29–32 weeks and > 32 weeks of gestation. 1,5-AG was measure by enzymatic colorimetric method (Glycomark™; Tomen America, New York, NY, USA) in automated system Labmax 400 analyzer (Labtest Diagnostic).

The laboratory parameters, markers for kidney function (urea), nourishment (total protein) and lipid profile were compatible with healthy subjects (Table 1).

The correlation in healthy subjects between 1,5-AG and glycemia, HbA1c, age, BMI and creatinine were weak or none (Table 2).

2. Experimental design, materials and methods

2.1. Study population

The population comprises 2303 unrelated Euro-Brazilian healthy subjects from Curitiba, State of Parana, South of Brazil [1]. All samples were obtained with the approval of the Ethics Committee of the Federal University of Parana.

Adult samples ($n = 922$) were collected from blood bank donors. Children ($n = 580$) and adolescent samples ($n = 496$) were obtained from Public Schools. Healthy pregnant women ($n = 305$) samples were obtained from the Curitiba Government Laboratory.

The normoglycemic criteria applied for selected subjects in the study were fasting glycemia < 5.5 mmol/L with an HbA1c range of 20.2–36.6 mmol/mol (4.0–5.7%) for children, adolescents, and adults. For pregnant women a fasting blood glucose < 5.1 mmol/L was applied to exclude gestational diabetes.

All subjects declared that were not using any medications or drugs.

2.2. Samples

Samples were serum obtained in non-fasting state for adults, children and adolescents, and fasting for those who were pregnant. Blood were collected in BD vacutainers SST II advance vacutainer with

Table 3
Serum 1,5-anhydroglucitol reference intervals and concentrations in different healthy populations.

Sex Subjects	1,5-AG, $\mu\text{mol/L}$				Studies/Methodology
	Male		Female		
	R.I.	n	R.I.	n	
Children and adolescents (5–18 years)	(92–298)	432	(84–278)	642	Our study [2] Enzymatic colorimetric GlycoMark™ [3] Enzymatic colorimetric GlycoMark™
US adolescents (12–18 years)	(95–178) [88–212] 150 \pm 31 (95–178) [102–198] 150 \pm 24 158 \pm 40	6	(140–172) [120–180] 150 \pm 15	5	
US young (10–29 years)	(63–271) [78–238] Total (n = 136) 151 \pm 39 (54–271); [73–229] (< 18 years) 158 \pm 35 [88–228] (> 18 years) 137 \pm 42 [53–221]	82	143 \pm 37 (54–227) [69–217]	54	[4] Enzymatic colorimetric GlycoMark™
Adults (19–79 years)	(80–260)	460	(62–241)	462	Our study [2] Enzymatic colorimetric GlycoMark™ [5] Gas chromatography (GC)
Finland adults (25–50 years)	93 mean 81 mean; 10–146 range (n = 139)	29	77 mean	110	
US adults (18–39 years)	(61–207)	224	(37–195)	224	[6] Enzymatic colorimetric GlycoMark™
US adults (18–39 years)	(52–178)	875	(50–166)	924	[7]
Australian adults (40 \pm 13 years)	125 \pm 41 (n = 95) [43–207]				[8] Enzymatic colorimetric GlycoMark™
German adults	157 \pm 44 (n = 116) [69–245]				[9] Liquid chromatography–mass Spectrometric (LC–MS)
Chinese Mauritians	144 \pm 51 (n = 82) [42–246] (98–195)				[10] Enzymatic pyranose oxidase
Chinese adults (22–80 years)	182 \pm 39 [104–260] 92–294 (n = 57)		159 \pm 52 [55–263]		[11] Enzymatic pyranose oxidase
Chinese adults > 20 and < 40 years > 50 years	176 \pm 46 [84–268] 166 \pm 67 [32–300]	82 9	116 \pm 35 [46–186] 122 \pm 41 [40–204]	185 14	[12] Liquid chromatography negative ion electrospray tandem mass spectrometry (LC–MS/MS)
Chinese adults	(83.1–240.7) 161.9 \pm 40.2 [81.5–242.3] (n = 120)				[13] Enzymatic

Chinese adults (22–78 years)	190 ± 54 [82–298] (69–278) [67–279] 173 ± 53 (107–367)	254	160 ± 49 [62–258]	322	[14] Enzymatic colorimetric GlycoMark™
Chinese adults (20–79 years)	226.3 ± 60.7 [104.9–347.7]	226	(79–306) 175.2 ± 55.8 [63.6–286.8]	232	[15] Enzymatic Medical system, Ningbo, China
Japanese (18–81 years)	132 ± 36 (n = 45) [60–204]				[16] Gas–liquid chromatography (GLC)
Japanese (mean 47 years)	145 ± 44 (n = 229) [57–233]				[17] Gas–liquid chromatography (GLC)
Japanese (27–68 years)	(114–215) 158 ± 38 [82–234]				[18] Gas–liquid chromatography (GLC)
Japanese adults (23–76 years)	159.8 ± 9.8 (n = 20) [140–179]				[19] Enzymatic Nippon–Kayaku
Japanese adults (30–79 years)	140 ± 56 [28–252]	991	122 ± 43 [36–208]	1104	[20] Enzymatic Kyowa Medex Co.
Japanese adults	137.1 ± 8.2 [120.7–153.5] (50.5 ± 9.7 y) 124.3 ± 45.1 [34.1–214.5] (74.5 ± 5.8 y)	181 231	120.6 ± 39 [42.6–198.6] (54.1 ± 10.3) 115.1 ± 40.2 [34.7–195.5] (75.3 ± 6.7 y)	203 519	[21] Enzymatic Lana Nippon–Kayaku

US, Americans from United States; UK, English; n, sample size.

Values were reference interval (2.5th–97.5th); [95% calculated as mean ± 2–SD]; mean ± SD or median (IQR, interquartile range, 25–75%).

Table 4
Serum 1,5-anhydroglucitol reference intervals and concentrations in pregnancy in different populations.

Pregnant	1,5-AG, $\mu\text{mol/L}$ R.I.	n	Studies/Methodology
Pregnant Women < 23 weeks of gestation	(56–298)	110	Our study [2] Enzymatic colorimetric GlycoMark™
Pregnant Women > 24 weeks of gestation	(33–181)	195	Our study [2] Enzymatic colorimetric GlycoMark™
Japanese healthy non-pregnant women	113 \pm 32 [49–197]	25	[22] Gas-liquid chromatography (GLC)
Japan pregnant women at 36 weeks gestation	62 \pm 28 [6–118]	543	
Japan women on 5th day of puerperium	66 \pm 23 [20–112]	543	
Japan women on 30th day of puerperium	87 \pm 21 [45–129]	543	
Japan pregnant Women at > 24 weeks of gestation	128 (IQR 102–160)		
UK Normoglycemic women with glycosuric pregnancy (~31 weeks)	46 (IQR 30–56)	16	[23] High-performance liquid chromatography (HPLC)
UK Normoglycemic women without glucosuric pregnancy (~31 weeks)	72 (IQR 55–79)	16	
UK Normoglycemic women (~31 weeks)	55 (IQR 31–72)	32	
Chinese pregnant women (16–45 years) at 26–28 weeks of gestation	133.0 \pm 52.9 [27.2–238.8]	44	[24] Enzymatic pyruvate oxidase

US, Americans from United States; UK, English; n, sample size.

Values were reference interval (2.5th–97.5th); [95% calculated as mean \pm 2 – SD]; mean \pm SD or median (IQR, interquartile range, 25–75%).

silica clot activator/gel (Becton Dickinson Co.). Bloods were separated in less than two hours from venipuncture and the serum stored in an ultrafreezer (-80°C).

2.3. Analytical methods

Concentrations of 1,5-AG were measured enzymatically with the Glycomark reagent (GlycoMark, Tomen America, New York, NY Inc.) in an automated system (Labmax 400 analyzer; Labtest. Diagnostics). The reaction details and methodology performance were described in Nowatzke et al. [6].

2.4. Clinical and laboratory parameters

Clinical data acquisition and analytical procedures, for laboratory data, have been reported previously [2].

2.5. Data analysis

Descriptive statistics, correlation analysis and reference intervals were calculated with MedCalc MedCalc version 17.6 (MedCalc Statistical Software bvba, Ostend, Belgium). Probability values (p -values) less than 5% ($p < 0.05$) were considered significant for all tests.

Acknowledgements

This project was supported by CNPq and the Araucaria Foundation.

Transparency document. Supporting information

Transparency data associated with this article can be found in the online version at <http://dx.doi.org/10.1016/j.dib.2018.08.165>.

References

- [1] M. Welter, K.C. Boritzka, M.I. Anghebem-Oliveira, R. Henneberg, A.B. Hauser, F.G.M. Rego, G. Picheth, Reference intervals for serum 1,5-anhydroglucitol in children, adolescents, adults, and pregnant women, *Clin. Chim. Acta; Int. J. Clin. Chem.* 486 (2018) 54–58.
- [2] M. Welter, K.C. Boritzka, M.I. Anghebem-Oliveira, R. Henneberg, A.B. Hauser, F.G.M. Rego, G. Picheth, Reference intervals for serum 1,5-anhydroglucitol in children, adolescents, adults, and pregnant women, *Clin. Chim. Acta* 486 (2018) 54–58.
- [3] T.M. Nguyen, L.M. Rodriguez, K.J. Mason, R.A. Heptulla, Serum 1,5-anhydroglucitol (Glycomark) levels in children with and without type 1 diabetes mellitus, *Pediatr. Diabetes* 8 (4) (2007) 214–219.
- [4] S.N. Mehta, N. Schwartz, J.R. Wood, B.M. Svoren, L.M. Laffel, Evaluation of 1,5-anhydroglucitol, hemoglobin A1c, and glucose levels in youth and young adults with type 1 diabetes and healthy controls, *Pediatr. Diabetes* 13 (3) (2012) 278–284.
- [5] E. Pitkanen, Serum 1,5-anhydroglucitol in normal subjects and in patients with insulin-dependent diabetes mellitus, *Scand. J. Clin. Lab. Investig.* 42 (5) (1982) 445–448.
- [6] W. Nowatzke, M.J. Sarno, N.C. Birch, D.F. Stickle, T. Eden, T.G. Cole, Evaluation of an assay for serum 1,5-anhydroglucitol (GlycoMark) and determination of reference intervals on the Hitachi 917 analyzer, *Clin. Chim. Acta; Int. J. Clin. Chem.* 350 (1–2) (2004) 201–209.
- [7] E. Selvin, B. Warren, X. He, D.B. Sacks, A.K. Saenger, Establishment of community-based reference intervals for fructosamine, glycated albumin, and 1,5-anhydroglucitol, *Clin. Chem.* 64 (5) (2018) 843–850.
- [8] A.S. Januszewski, C. Karschimkus, K.E. Davis, D. O'Neal, G. Ward, A.J. Jenkins, Plasma 1,5 anhydroglucitol levels, a measure of short-term glycaemia: assay assessment and lower levels in diabetic vs. non-diabetic subjects, *Diabetes Res. Clin. Pract.* 95 (1) (2012) e17–e19.
- [9] C. Hess, B. Stratmann, W. Quester, B. Madea, F. Musshoff, D. Tschoepe, Clinical and forensic examinations of glycemic marker 1,5-anhydroglucitol by means of high performance liquid chromatography tandem mass spectrometry, *Forensic Sci. Int.* 222 (1–3) (2012) 132–136.

- [10] D.A. Robertson, K.G. Alberti, G.K. Dowse, P. Zimmet, J. Tuomilehto, H. Gareeboo, Is serum anhydroglucitol an alternative to the oral glucose tolerance test for diabetes screening? The Mauritius noncommunicable diseases study group, *Diabet. Med. J. Br. Diabet. Assoc. MC: Diabet. Med.* 10 (1993) 56–60.
- [11] H. Shi, J. Fang, X. Yang, Z. Shen, X. Zhu, Serum 1,5-anhydro-D-glucitol as a new clinical marker for glucose metabolism in type 2 diabetics, *Chin. Med. J.* 112 (6) (1999) 571–573.
- [12] S. Li, X. Heng, H. Sheng, Y. Wang, C. Yu, Determination of glycemc monitoring marker 1,5-anhydroglucitol in plasma by liquid chromatography-electrospray tandem mass spectrometry, *J. Chromatogr. B Anal. Technol. Biomed. Life Sci.* 875 (2) (2008) 459–464.
- [13] F. Jing, L. Jun, Y. Wang, M. Zhu, Z. Yong, X. Fei, J. Zhang, A novel fully enzymatic method for determining glucose and 1,5-anhydro-D-glucitol in serum of one cuvette, *Appl. Biochem. Biotechnol.* 150 (3) (2008) 327–335.
- [14] Y. Wang, Y.L. Zhang, Y.P. Wang, C.H. Lei, Z.L. Sun, A study on the association of serum 1,5-anhydroglucitol levels and the hyperglycaemic excursions as measured by continuous glucose monitoring system among people with type 2 diabetes in China, *Diabetes/Metab. Res. Rev.* 28 (4) (2012) 357–362.
- [15] Q. Zhou, D.B. Shi, L.Y. Lv, The establishment of biological reference intervals of nontraditional glycemc markers in a Chinese population, *J. Clin. Lab. Anal.* (2016) 1–7. <https://doi.org/10.1002/jcla.22097>.
- [16] T. Yamanouchi, H. Akanuma, T. Asano, C. Konishi, I. Akaoka, Y. Akanuma, Reduction and recovery of plasma 1,5-anhydro-D-glucitol level in diabetes mellitus, *Diabetes* 36 (6) (1987) 709–715.
- [17] T. Yamanouchi, H. Akanuma, T. Nakamura, I. Akaoka, Y. Akanuma, Reduction of plasma 1,5-anhydroglucitol (1-deoxyglucose) concentration in diabetic patients, *Diabetologia* 31 (1) (1988) 41–45.
- [18] T. Yamanouchi, Y. Tachibana, H. Akanuma, S. Minoda, T. Shinohara, H. Moromizato, H. Miyashita, I. Akaoka, Origin and disposal of 1,5-anhydroglucitol, a major polyol in the human body, *Am. J. Physiol.* 263 (2 Pt 1) (1992) E268–E273.
- [19] T. Niwa, L. Dewald, J. Sone, T. Miyazaki, M. Kajita, Quantification of serum 1,5-anhydroglucitol in uremic and diabetic patients by liquid chromatography/mass spectrometry, *Clin. Chem.* 40 (2) (1994) 260–264.
- [20] M. Watanabe, Y. Kokubo, A. Higashiyama, Y. Ono, Y. Miyamoto, T. Okamura, Serum 1,5-anhydro-D-glucitol levels predict first-ever cardiovascular disease: an 11-year population-based cohort study in Japan, the Suita study, *Atherosclerosis* 216 (2) (2011) 477–483.
- [21] M. Ouchi, K. Oba, H. Yamashita, M. Okazaki, M. Tsunoda, M. Ohara, K. Sekimizu, K. Watanabe, T. Suzuki, H. Nakano, Effects of sex and age on serum 1,5-anhydroglucitol in nondiabetic subjects, *Experimental and clinical endocrinology & diabetes: official journal, Ger. Soc. Endocrinol. Ger. Diabetes Assoc.* 120 (5) (2012) 288–295.
- [22] M. Tetsuo, T. Hamada, K. Yoshimatsu, J. Ishimatsu, T. Matsunaga, Serum levels of 1,5-anhydro-D-glucitol during the normal and diabetic pregnancy and puerperium, *Acta Obstet. Et. Gynecol. Scand.* 69 (6) (1990) 479–485.
- [23] E.S. Kilpatrick, B.G. Keevilt, K.L. Richmond, P. Newland, G.M. Addison, Plasma 1,5-anhydroglucitol concentrations are influenced by variations in the renal threshold for glucose, *Diabet. Med. J. Br. Diabet. Assoc. MC: Diabet. Med.* 16 (1999) 496–499.
- [24] W.H. Tam, M.S. Rogers, T.K. Lau, M. Arumanayagam, The predictive value of serum 1,5-anhydro-D-glucitol in pregnancies at increased risk of gestational diabetes mellitus and gestational impaired glucose tolerance, *BJOG: Int. J. Obstet. Gynaecol.* 108 (7) (2001) 754–756.