

Higher hydrocortisone dose increases bilirubin in hypopituitary patients- results from an RCT

Jorien Werumeus Buning*, Jenny E. Kootstra-Ros[†], Pauline Brummelman*, Gerrit van den Berg*, Melanie van der Klauw*, Bruce H. R. Wolffenbuttel*, André P. van Beek* and Robin P. F. Dullaart*

*Department of Endocrinology, [†]Laboratory Center, University of Groningen, University Medical Center Groningen, Groningen, The Netherlands

ABSTRACT

Background Bilirubin has anti-oxidative and anti-inflammatory properties, which may explain its proposed protective effects on the development of cardiometabolic disorders. Glucocorticoids affect heme oxygenase regulation *in vitro*, which plays a key role in bilirubin production. Effects of variations in glucocorticoid exposure on circulating bilirubin levels in humans are unknown. Here we tested whether a higher hydrocortisone replacement dose affects circulating bilirubin in hypopituitary patients.

Materials and methods A randomized double-blind cross-over study (ClinicalTrials.gov, number NCT01546992) was performed in 47 patients with secondary adrenal failure [10-week exposure to a higher hydrocortisone dose (0.4–0.6 mg/kg body weight) vs. 10 weeks of a lower hydrocortisone dose (0.2–0.3 mg/kg body weight)].

Results Plasma total bilirubin was increased by 10% from 7 to 8 μ M in response to the higher hydrocortisone dose ($P = 0.033$). This effect was inversely related to age ($P = 0.042$), but was unaffected by sex, obesity and (replacement for) other hormonal insufficiencies. The higher hydrocortisone dose also resulted in lower alkaline phosphatase ($P = 0.006$) and aspartate aminotransferase activities ($P = 0.001$).

Conclusion Bilirubin is modestly increased in response to higher glucocorticoid exposure in humans, in conjunction with lower alkaline phosphatase and aspartate aminotransferase activities, which are supposed to represent biomarkers of a pro-inflammatory state and enhanced liver fat accumulation.

Keywords Alkaline phosphatase, bilirubin, hydrocortisone, secondary adrenal insufficiency, transaminases.

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Introduction

Bilirubin is able to scavenge peroxy radicals, to diminish low-density lipoprotein (LDL) oxidation, to decrease the expression of cellular adhesion molecules and to inhibit the production of pro-inflammatory cytokines [1–3]. Furthermore, circulating bilirubin levels are inversely related to the inflammation markers, high-sensitivity C-reactive protein and serum amyloid A, as well as to a composite measure of circulating glycosylated acute-phase proteins [4–6]. Among other mechanisms, anti-oxidative and anti-inflammatory properties are believed to explain the proposed protective effects of bilirubin on the development of cardiometabolic disorders [3]. In line, bilirubin is inversely associated with intima media thickness, a marker of subclinical atherosclerosis [7,8], as well as with incident cardiovascular disease [9,10].

The heme oxygenase (HO) system is crucial for the generation of bilirubin from heme [3,11,12]. Expression of the HO-1

isoenzyme is inducible by a variety of factors, whereas the HO-2 isoenzyme is considered to be expressed in a predominantly constitutive fashion [11,12]. Interestingly, the HO-2 gene contains a functional glucocorticoid response element [11]. As a result, HO-2 transcripts and protein levels are upregulated by glucocorticoids *in vitro* at least in neonatal rat brain and testis [13–15]. On the other hand, upregulation of HO-1 by interleukin-6 may be counteracted by glucocorticoids in endothelial cells [16].

Given the alleged influence of bilirubin on cardiometabolic disorders [3], it is pathophysiologically relevant to determine the contribution of glucocorticoids to bilirubin regulation, but no data are available with respect to effects of variation in glucocorticoid exposure on circulating bilirubin levels in humans. We have recently conducted a randomized cross-over study in hypopituitary patients with secondary adrenal insufficiency (ClinicalTrials.gov, number NCT01546992) which is aimed at assessing effects of a higher compared to a lower

hydrocortisone replacement dose on cognitive function, quality of life and metabolic parameters [17]. The present ancillary analyses were performed to test whether the degree of hydrocortisone exposure affects circulating bilirubin levels in this patient category.

Patients and methods

Participants and study design

Reporting of the study conforms to CONSORT and the broader EQUATOR guidelines [18]. This randomized double-blind cross-over study was performed in a university hospital setting in Groningen, the Netherlands. The study has been registered with ClinicalTrials.gov, number NCT01546992, and has been approved by the medical ethics committee of the University Medical Center Groningen, the Netherlands. All participants provided written informed consent. The rationale and design of the study have been provided in detail elsewhere [17]. Figure S1 shows eligibility, inclusion and follow-up of the participants. In brief, hypopituitary patients with established secondary adrenal insufficiency (based on internationally accepted criteria) were eligible for this randomized double-blind cross-over study. They were recruited from the outpatient endocrinology clinic of the University Medical Center, Groningen, the Netherlands. The following inclusion criteria were applied: treatment for pituitary disorder (surgery and/or radiotherapy) at least 1 year before study entry, stable replacement therapy for other pituitary hormone deficiencies for at least 6 months (thyroid hormone deficiency, growth hormone deficiency, testosterone/estradiol deficiency, diabetes insipidus), age between 18 and 75 years and body weight between 50 and 100 kg [17]. Main exclusion criteria were major cognitive impairment, drug abuse, current psychiatric disorders, shift work, malignancy, previous Cushing's disease, a history of frequent episodes of hypocortisolism, medically treated diabetes possibly leading to hypoglycaemia (insulin, sulfonylurea) and anti-epileptic drugs [17]. Subjects experiencing a hospital admission during the study were also excluded.

Before entry in the study, all patients who were treated with cortisone acetate were switched to hydrocortisone in a bioequivalent dose during a 4-week run-in period [17]. The hydrocortisone dose administered during the run-in period averaged 25 mg (0.31 mg/kg body weight) per day. The participants were then randomized to a lower hydrocortisone dose during 10 weeks followed by a higher hydrocortisone dose during another 10-week period or vice versa. The lower hydrocortisone dose was 0.20–0.30 mg hydrocortisone per kg body weight per day divided into three doses, taken before breakfast, before lunch and before dinner. A double dose, also divided into three doses and taken at the same time points, was administered during the higher hydrocortisone dose period.

Hydrocortisone administration was performed in a double-blind fashion using 5-mg tablets in the lower and 10-mg tablets in the higher hydrocortisone replacement period. In case of intercurrent illnesses, doubling or tripling the hydrocortisone dose was allowed for 7 days maximally (+10% of the cumulative study dose). Compliance to hydrocortisone tablet intake was verified as described [17]. Sixty-three patients were initially randomized of whom 47 completed the study. Reasons for withdrawal were described elsewhere [17]. All these 47 subjects were included in the present analysis. Twenty-two patients received the lower hydrocortisone dose first, whereas 25 participants were first given the higher hydrocortisone dose. Detailed clinical characteristics of the participants, including medical history, cause of secondary adrenal insufficiency and other hormone replacement therapies, have been provided elsewhere [17].

Venous blood was obtained after an overnight fast at 8 a.m.; that is, 1 h after the morning hydrocortisone dose was taken. Twenty-four-hour urine collections were obtained 1 day before the scheduled visits at the end of the lower and higher hydrocortisone periods. Body mass index was calculated as body weight (in kg) divided by length (in m) squared.

Laboratory analyses

Plasma and serum samples were prepared by centrifugation at 4 °C. The samples were then stored at –80 °C until analysis.

Total bilirubin was measured in heparinized plasma by a colorimetric assay (Roche, Mannheim, Germany). The detection limit was 1.7 µM. The intra-assay coefficient of variation (CV) was < 2% at the lower normal range. In healthy subjects, bilirubin is most abundantly present in plasma in its unconjugated form [19]. In a validation experiment ($n = 80$), a strong correlation between total bilirubin and unconjugated bilirubin (Spearman's $r = 0.92$, $P < 0.001$), as well as between total bilirubin and conjugated direct bilirubin (Spearman's $r = 0.82$, $P < 0.001$) was observed. For this study, we only used total bilirubin in agreement with other reports [5,6,8–10].

Plasma alkaline phosphatase (ALP), alanine transferase (ALT), aspartate aminotransferase (AST) and gamma-glutamyltransferase (GGT) were routinely measured on a Roche Modular platform. All tests were performed according to the International Federation of Clinical Chemistry recommendations. ALT and AST were measured with pyridoxal phosphate activation. The intra-assay coefficients of variation (CVs) were < 2.5%.

Serum cortisol was measured by electrochemiluminescence immunoassay (Roche Modular Systems, Mannheim, Germany; intra-assay CV < 2%). Urinary free cortisol was analysed by automated online solid phase extraction in combination with liquid chromatography tandem-mass spectrometry (XLC-MS/MS) (intra-assay CV < 2.5%).

Statistical analysis

Sample size calculation and information regarding the random allocation sequence and its implementation have been provided elsewhere [17]. Data are presented as mean ± SD or as median (interquartile range). Because of skewed distribution total bilirubin, ALP, ALT, AST, GGT and urinary free cortisol excretion were natural log-transformed to achieve approximately normal distributions. Changes are presented as mean (95% confidence intervals) for parametrically distributed variables and as median (interquartile range) for nonparametrically distributed variables. The effect of higher vs. lower hydrocortisone dose on the various variables was compared by Student's t-tests. The univariate relationships of changes in bilirubin with continuous and dichotomous baseline variables were tested using Pearson's correlation coefficients and unpaired Student's t-tests, respectively. To check for period and treatment by period interaction effects (carry-over effect), the procedure developed by Altman was used [20].

Results

Twenty-nine men and 18 women completed the study. At entry, mean age at entry was 52 ± 14 years, and mean body mass index was 27.0 ± 4.0 kg/m². Thyroid hormone was diagnosed in 92% of patients and replaced in all; growth hormone deficiency was diagnosed in 66% and replaced in 68%; testosterone deficiency was diagnosed 79% of men and

replaced in all; eight women were premenopausal in whom estradiol deficiency was diagnosed in 50% and treated in all (postmenopausal estradiol deficiency not replaced); and diabetes insipidus was diagnosed in 19% and replaced in all. Baseline clinical characteristics were provided elsewhere [17]. The various hormonal substitution regimens were left unchanged during the 4-week run-in period, as well as during the 10-week lower and higher hydrocortisone replacement periods. A doubling of the hydrocortisone dose as stress-related dose adjustment was reported 159 times on the low dose (1.6% of the total dose administrations) and 146 times on the high dose (1.5% of the total dose administrations).

Doubling of the hydrocortisone dose elicited an increase in serum cortisol, as well as in urinary free cortisol excretion (Table 1). Body mass index was slightly increased with the higher hydrocortisone dose, but plasma glucose was unchanged (Table 1). On average, total bilirubin was increased by 10% during the higher dose of hydrocortisone, whereas serum ALP and AST activity were slightly decreased (Table 1; Fig. 1). ALT and GGT activity did not significantly change. In univariate analysis, the increase in total bilirubin was inversely related to age ($r = -0.298, P = 0.042$), but was related neither to sex, BMI, the use of thyroid hormone, the use of sex steroids, growth hormone deficiency, diabetes insipidus and plasma glucose ($P > 0.29$ for all), nor to the sequence of the lower and higher hydrocortisone doses ($P = 0.35$). No period or treatment by period interaction effects was found for all the outcome measures (all $P > 0.05$).

Table 1 Body mass index, estimates of cortisol metabolism, plasma glucose, plasma total bilirubin, alkaline phosphatase activity, transaminase activities and γ -glutamyltransferase (GGT) activity during randomized periods of lower and higher hydrocortisone replacement doses in 47 hypopituitary patients with secondary adrenal insufficiency.

	Lower dose	Higher dose	Change	P-value
Hydrocortisone dose (mg/day)	17.93 ± 2.23	35.85 ± 4.46	17.93 (17.29–18.56)	< 0.001
Body mass index (kg/m ²)	27.0 ± 4.0	27.2 ± 4.1	0.2 (0.0–0.3)	0.026
Serum cortisol (1 h after intake; nM)	643 ± 241	918 ± 245	275 (206–345)	< 0.001
Urinary free cortisol (nmol/24 h)	78 (46–111)	274 (199–408)	207 (131–314)	< 0.001
Fasting plasma glucose (mM)	5.24 ± 0.9	5.21 ± 0.8	-0.03 (-0.18–0.12)	0.70
Plasma total bilirubin (μM)	7 (5–12)	8 (6–11)	1 (-1–2)	0.033
Plasma ALP activity (U/L)	60 (48–73)	55 (47–73)	-2 (-9–2)	0.006
Plasma AST activity (U/L)	26 (22–31)	22 (20–28)	-3 (-7–0)	0.001
Plasma ALT activity (U/L)	21 (16–33)	19 (15–29)	-1 (-7–3)	0.085
Plasma GGT activity (U/L)	23 (16–33)	23 (18–34)	1 (-2–4)	0.30

Data in mean ± SD or median (interquartile range). Changes are given in mean with 95% confidence intervals (CI) for parametrically distributed data or as median (interquartile range) for nonparametrically distributed data. ALP, alkaline phosphatase activity; ALT, alanine transferase activity; AST, aspartate aminotransferase activity; GGT, γ -glutamyltransferase activity.

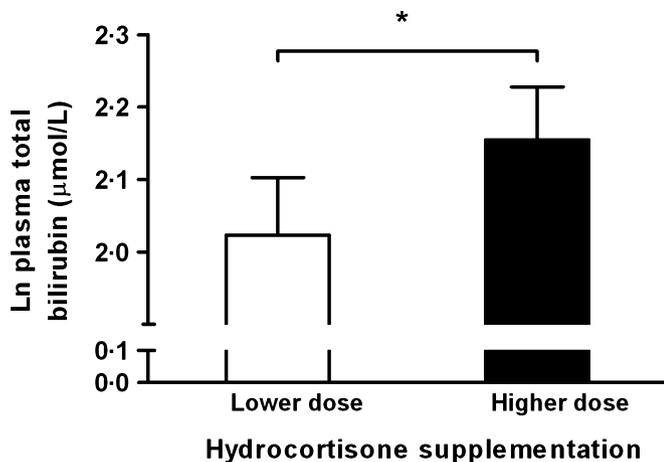


Figure 1 Plasma total bilirubin in response to higher vs. lower hydrocortisone replacement doses. Natural log-transformed (Ln) values of plasma bilirubin are shown. Bars indicate mean \pm SE values. * $P = 0.033$.

Discussion

This randomized double-blind cross-over study demonstrates to our knowledge for the first time that a higher hydrocortisone replacement regimen increases circulating bilirubin levels in hypopituitary subjects with secondary adrenal insufficiency. This effect appeared to be less outspoken in older individuals, but was unrelated to sex, obesity, fasting glucose and (replacement for) other hormonal deficiencies. Moreover, we could rule out confounding due to carry-over effects of the higher and the lower hydrocortisone replacement doses. The present findings are, therefore, consistent with the notion that variations in glucocorticoid exposure affect bilirubin metabolism in humans.

In view of the supposition that bilirubin levels reflect overall status of HO activity [21], our results consent with the hypothesis that glucocorticoids stimulate HO expression in humans. While this possibility agrees with previous results in rodent models [13–15], it is obvious that the current findings do not allow to discern between differential effects of hydrocortisone on specific HO isoforms nor in specific tissues. Furthermore, it should be noted that bilirubin levels are also determined by biliary elimination via uridine diphosphate glucuronosyltransferase 1A1, which conjugates bilirubin and enables bilirubin elimination [3]. This enzyme is induced by glucocorticoids in HepG2 cells [22,23]. Although an opposing contribution to the glucocorticoid increasing effects on circulating bilirubin via uridine diphosphate glucuronosyltransferase 1A1 stimulation cannot be excluded, short-time glucocorticoid administration does not affect biliary bilirubin output in humans [24].

Doubling the hydrocortisone replacement dose expectedly increased serum cortisol and urinary free cortisol excretion, and may have resulted in mild hypercortisolism in some individuals [25]. The increase in serum bilirubin in response to the higher hydrocortisone replacement regimen averaged 10%. While this previously unreported bilirubin response may have pathophysiological relevance, its clinical significance is uncertain. Based on large prospective cohort studies and meta-analysis, the risk of incident cardiovascular disease has been estimated to be about 18% lower per doubling of the bilirubin level [9,10]. In this context, the impact of the observed bilirubin increase on cardiovascular risk is likely to be small. Of note, we also documented a drop in ALP activity in response to a higher hydrocortisone replacement dose. As ALP is regarded as an inflammatory biomarker [26,27], this finding may point to less chronic low-grade inflammation consequent to modestly higher glucocorticoid exposure [28]. However, a limitation of our study is that we did not measure other inflammation markers such as high-sensitivity C-reactive protein.

It is pertinent that overt hypercortisolism coincides with hepatic fat accumulation [29,30]. However, plasma AST activity was also modestly decreased after the higher hydrocortisone replacement dose. Considering that this enzyme may provide a biomarker of liver fat accumulation [31,32], this finding would be consistent with some beneficial effect on nonalcoholic fatty liver disease (NAFLD) development, putatively consequent to higher bilirubin levels [33]. In addition, it seems possible that the increasing effect of a higher hydrocortisone dose on plasma bilirubin could provide a potentially relevant additional mechanism which contributes to the anti-inflammatory role commonly ascribed to glucocorticoids [28].

Several other methodological issues and limitations of our study need to be discussed. First, our trial was performed in hypopituitary patients with established secondary adrenal insufficiency to rule out confounding due to coexisting mineralocorticoid deficiency – in particular on cognitive parameters – as much as possible. Consequently, extrapolation to other patient categories is limited. Second, our study was not designed to establish glucocorticoid dose- and time-dependent effects on circulating bilirubin. However, the present comparison of a lower and higher hydrocortisone replacement regimen puts the changes in bilirubin in a physiological perspective as much as possible. Additionally, it seems plausible that 10-week administration periods are sufficient to establish a steady state of altered glucocorticoid exposure to various metabolic processes. We believe the same holds true for a 4-week run-in period in particular given the fact that the total daily hydrocortisone dose during the run-in period was lower than that during the higher hydrocortisone dose replacement period. Third, all participants received adequate replacement treatment for other pituitary hormone deficiencies if deemed clinically

necessary. Replacement doses with thyroid hormone, growth hormone, sex steroids and desmopressin were left unchanged during the study. This approach was chosen to limit coexisting effects of other hormonal factors on bilirubin metabolism [34,35].

In conclusion, circulating bilirubin is modestly increased in response to higher glucocorticoid exposure. Our findings support the hypothesis that glucocorticoids contribute to bilirubin metabolism in humans.

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Conflict of interest statement

The study is investigator-driven, and the authors do not have any financial or other conflict of interests to declare.

Contributors

Werumeus Buning, van Beek and Dullaart had full access to all of the data in the study and take responsibility for the integrity of the data and the accuracy of the data analysis. The manuscript was completely drafted by Werumeus Buning, van Beek and Dullaart. Brummelman, Wolffenbittel, van Beek and Dullaart conceptualized and designed the study. Werumeus Buning, Kootstra-Ros, Brummelman, van den Berg, van der Klauw, Wolffenbittel, van Beek and Dullaart collected, analysed and interpreted the data. Werumeus Buning, van Beek and Dullaart drafted the manuscript. Werumeus Buning, Koostra-Ros, Brummelman, van den Berg, van der Klauw, Wolffenbittel, van Beek and Dullaart critical revised the manuscript for important intellectual content. Werumeus Buning and Dullaart statistically analysed the data. Van Beek supervised the study.

Address

Department of Endocrinology, University of Groningen, University Medical Center Groningen, P.O. Box 30.001, Groningen, 9700 RB, The Netherlands (J. Werumeus Buning, P. Brummelman, P. Brummelman, G. van den Berg, M. van der Klauw, B. H. R. Wolffenbittel, A. P. van Beek, R. P.F. Dullaart); Laboratory Center, University of Groningen, University Medical Center Groningen, P.O. Box 30.001, Groningen, 9700 RB, The Netherlands (J. E. Kootstra-Ros).

Correspondence to: Robin P. F. Dullaart, MD, PhD, Department of Endocrinology, University Medical Center Groningen,

P.O. Box 30-001, Groningen, 9700 RB, The Netherlands. Tel.: +31 503613731; fax: +31 503619392; e-mail: r.p.f.dullaart@int.umcg.nl

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Supporting Information

Additional Supporting Information may be found online in the supporting information tab for this article:

Figure S1. Eligibility, inclusion and follow-up of the participants.