Hypothalamic-pituitary-adrenal axis suppression in asthma: A glucocorticoid receptor polymorphism may protect

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Abstract

Background: Asthmatic children on corticosteroids can develop hypothalamic-pituitary-adrenal axis suppression (HPAS). Single nucleotide polymorphisms (SNPs) rs242941 and rs1876828 of the corticotrophin-releasing hormone receptor 1 (CRHR1) gene were associated with lower stimulated cortisol (F) levels, whereas rs41423247 of the glucocorticoid receptor (NR3C1) gene was associated with higher basal F levels. The objective of the current study was to confirm whether these three SNPs are associated with HPAS in asthmatic children. Methods: DNA was extracted from saliva obtained from 95 asthmatic children, who had previously undergone basal F and metyrapone testing. Thirty-six children were classified as suppressed. Non-suppressed children were sub-classified according to their post-metyrapone ACTH (PMTP ACTH) level into a middle (106-319 pg/ml) and a high (>319 pg/ml) ACTH response group. TaqMan® polymerase chain reaction assays were utilized. Results: Only rs41423247 was inversely associated with HPAS (OR = 0.27 [95% CI 0.06-0.90]). Its GC genotype was inversely associated with HPAS (log odds = -1.28, p = 0.021). PMTP ACTH was associated with CC (effect size = 10.85, p = 0.005) and GC genotypes (effect size = 4.06, p = 0.023). The C allele is inherited as a dominant trait (effect size = -1.31 (95% CI -2.39 – -0.33; p = 0.012). In the high ACTH response group, both genotypes affected the PMTP ACTH (effect sizes 1.41 and 15.46; p-values 0.023 and < 2x10-26 for GC and CC respectively). Conclusions: The C allele of rs41423247 was found to be protective against HPAS. CC genotype is associated with the highest PMTP ACTH response.
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Abstract and keywords

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Conclusions: The C allele of rs41423247 was found to be protective against HPAS. CC genotype is associated with the highest PMTPACTH response.
Key words: Asthma, hypothalamus, pituitary, adrenal, polymorphism, receptors, glucocorticoids, steroids/adverse events, adrenal insufficiency, pituitary-adrenal-function tests

Main text

Key Messages

Rs41423247 (G/C) of the NR3C1 gene was found to be inversely associated with hypothalamic-pituitary-adrenal axis suppression (HPAS) in asthmatic children on steroids. The CC genotype is associated with the highest ACTH levels. If confirmed, the CC genotype could well be a marker of HPAS protection, thereby allowing for personalized treatment decisions to be made in asthma care.

1 BACKGROUND

It has been shown that 16.3 % of asthmatic children treated with inhaled corticosteroids (ICS) and nasal steroids (NS) in Cape Town, South Africa, can develop hypothalamic-pituitary-adrenal axis suppression (HPAS) when assessed with the metyrapone (MTP) test. If partial forms of suppression are included, even two-thirds of children may be affected. A higher body mass index (BMI) was found to be protective against HPAS. Unfortunately suitable screening tests for HPAS are not available. The only viable alternative thus would be to identify gene variants, which would help to predict which children are either prone or resistant to developing HPAS. In Greek asthmatic children on inhaled corticosteroids, homozgyotes for the single nucleotide polymorphisms (SNPs) rs242941 (TT) and rs1876828 (AA) of the corticotrophin-releasing hormone receptor 1 (CRHR1) gene were associated with lower stimulated cortisol (F) levels when tested with the low dose adrenocorticotropic (ACTH) stimulation test (0.5 μg/1.73m²). Heterozygotes (GC) for rs41423247 (Bcl 1 restriction length polymorphism) of the glucocorticoid receptor (NR3C1) gene, on the other hand, were found to have higher basal F levels. In the quest for universally applicable predictors of HPAS in asthmatic children on corticosteroids, the findings of the Greek study needed to be confirmed by a study, utilizing the gold standard metyrapone test, to determine whether the SNPs rs242941 and rs1876828 of the CRHR1, and rs41423247 of the NR3C1 gene are associated with HPAS in asthmatic school children on corticosteroids.

2 METHODS

2.1 Participants

Asthmatic children, 5-18 years old, treated with ICS and NS at the Lung Institute, Tygerberg and Red Cross Children’s Hospitals in Cape Town, South Africa, whose hypothalamic-pituitary-adrenal axis (HPA) was previously assessed with a morning basal serum F and MTP testing, were re-recruited. HPAS was diagnosed, if F < 83 nmol/l or the post-MTP (PMTP) ACTH < 106 pg/ml (23.5 pmol/l), 11-deoxycortisol (11DOC) < 208 nmol/l and 11DOC+C < 400 nmol/l.

2.2 Study design and methodology

A cross-sectional study was performed. Height, weight, sex and age were recorded as measured previously. The BMI z-score (Centre of Disease Control) was computed. Salivary samples were collected with an Oragene DNA collection kit (OG-500). Samples were stored at -20°C. Once all the samples were collected, DNA was extracted. Genotyping for rs242941, rs1876828 and rs41423247 was performed by using TaqMan® polymerase chain reaction (PCR) assays. Ethical approval was granted by the ethics committees of both Stellenbosch University and the University of Cape Town. The study conformed to the standards of the Declaration of Helsinki. All participants and their parents gave their informed consent prior to inclusion of the study.

2.3 Statistical analysis and sample size considerations

2.3.1 Sample size

In the previous study1 30 patients had hypocortisolaemia or HPAS. The cross-sectional design allowed for an equal number of patients to be analysed across the full spectrum of post-MTP responses i.e. low, midrange and high (30 subjects each). The ACTH mean and standard deviation (SD) were calculated (ACTH
mean = 330 pg/ml [73.3 pmol/l], SD = 280 pg/ml [62.2 pmol/l]) from the previous data. The proportional distribution of the genetic subgroups was derived from the paper of Tsartsali et al, table 4.7. Power analysis with a sample size of 90 was done across the three ACTH groups. Significance level was taken as 0.05. A one-way analysis of variance test (ANOVA) was considered for the power analysis. The following mean values were specified under the alternative hypothesis: group 1 (low range) 230 pg/ml [51.1 pmol/l], group 2 (mid-range) 330 pg/ml [73.3 pmol/l], group 3 (high range) 430 pg/ml [95.5 pmol/l]. This gives a 100 ACTH units difference between two adjacent groups. This is a relative difference of 36% of the SD. [(100/280) = 0.36] which is a conservative expected effect size. Even for unbalanced groups such as n1=24, n2=35 and n3=40, the power will be 81%. Thus the power is good if the sample size distribution across the genetic subgroups is moderately unbalanced. Power is improved by square root transformations of ACTH to stabilise the within group variances.

2.3.2 Statistical analysis

An online statistical tool (https://ihg.helmholtz-muenchen.de/cgi-bin/hw/hwa1.pl) and the R statistical package were used. Fisher’s Exact and Pearson’s Chi-square tests were conducted to test for statistical difference of minor allele counts. The Hardy-Weinberg package version 1.6.3 installed in RStudio version 3.5.1 was used to assess whether the SNPs were in Hardy-Weinberg equilibrium. To test for significance at SNP and genotype level, the Wald’s test in multinom() function from the nnet package in R was utilized. PMTP ACTH data were square root transformed and a one-way ANOVA performed. The Wilcoxon test was used for pairwise multiple comparisons of differences in mean [?]?PMTP ACTH per SNP. Linear analysis for BMI z-score, [?]?PMTP ACTH and binomial analysis for HPAS were conducted with the glm() function in R. The best genetic model for rs41423247 was selected on the basis of the lowest significant Akaike information criterion. Significance level for all tests was taken as 0.05.

3 RESULTS

A total of 96 patients (94 from previous studies1,2 and two new patients) were recruited. One child had to be excluded, because of a discrepancy between the basal F level and the MTP test. The demographics, therapy and HPAS status of the 95 enrolled patients are listed (table 1). Only one child was on oral prednisone at the time of testing. According to the basal F and the PMTP ACTH response, 35 children were classified in a suppressed (F < 83 nmol/l or ACTH < 106 pg/ml [23.5 pmol/l]), 29 in a middle (ACTH 106-319 pg/ml [23.5-70.8 pmol/l]) and 31 in a high (ACTH > 319 pg/ml [70.8 pmol/l]) ACTH response group.

All three SNPs were in Hardy-Weinberg equilibrium (rs242941 A/C [suppressed ACTH response group p=0.830; middle/high ACTH response group p=0.580], rs1876828 A/T [suppressed ACTH response group p=0.791; middle/high ACTH response group p=0.481] and rs41423247 G/C [suppressed ACTH response group p=0.445; middle/high ACTH response group p=0.948]). The rs41423247 (G/C) SNP was inversely associated with HPAS (OR = 0.27 [95% CI 0.06-0.90]). Both rs242941 (A/C) and rs1876828 (C/T) were not associated with HPAS, their OR being 0.99 (95% CI 0.43-2.27) and 0.54 (95% CI 0.05-3.48) respectively. Similarly, only the rs41423247 GC and GC+CC genotypes were inversely associated with the suppressed ACTH response group(OR=0.310 [95% CI: 0.137-0.920] and OR = 0.296 [95% CI: 0.1116 - 0.756]). The homozygous CC genotype just missed statistical significance (OR = 0.101 [95% CI: 0.005-1.908], p = 0.041). The mean [?]?PMTP ACTH of the CC genotype was significantly higher (p = 0.002) than for the GC and GG genotypes (fig 1). The respective means and their 95% CIs were 22.59 (16.18-29.00), 16.00 (13.00-19.00) and 11.82 (10.11-13.63). These correspond to real ACTH mean levels of 553.0 pg/ml (122.9 pmol/l) for the CC, 331.0 pg/ml (73.6 pmol/l) for the GC and 187.5 pg/ml (41.7 pmol/l) for the GG genotype.

The frequency of the guanosine (G) allele (the major allele) of rs41423247 was 77%, while the frequency of the cytidine (C) allele (the minor allele) was 23%. The genotype frequencies for GG, GC and CC were 60, 34 and 6% respectively.

BMI was associated with the CC genotype (table 2) while the heterozygous GC was associated with HPAS,
independently of BMI (table 3). Both the homozygous CC and the heterozygous GC genotypes were associated with \[?\]PMTP ACTH with CC having a significant larger impact than GC (table 4). The observed ACTH effect was independent of age, sex, height and weight (and hence of BMI). The higher the ACTH response, the greater and more significant the genotype (CC) effect (table 5).

After correcting for age, sex, height and weight, both the additive and the dominant genetic models were the best fit to describe the protective effect of the C allele on HPAS. The effect size for the additive model was -1.30 (-2.32 – -0.42; p = 0.007) and for the dominant model, the effect size was -1.31 (95% CI -2.39 – -0.33; p = 0.012). Similarly, the effect of the C allele on \[?\]PMTP ACTH is best described by an additive (effect size = 4.42 [95% CI 1.77 – 7.07; p = 0.002]) and a dominant model (effect size = 4.79 [95% CI 1.54-8.04; p = 0.005]).

4 DISCUSSION

The SNP rs41423247 (GC) of theNR3C1 gene was found to be protective against HPAS, while no association with HPAS was found with rs242941 and rs1876828 of the CRHR1 gene. In contrast, in Greek asthmatic children basal or stimulated F levels were different for all three SNPs.\(^5\) The discrepancy in the findings between results of the two studies can be explained by the use of different adrenal function tests. In the Greek study the low dose ACTH stimulation test (not the gold standard) was used without defining definitive cut-offs\(^3,6\) for basal and stimulated F levels. Relying on statistical differences of these levels vis-a-vis their respective genotypes could have led to erroneous conclusions. Furthermore, serum F is a very crude test, which is influenced by many variables\(^3\) e.g. by Circadian rhythm. There is no record in Tsatsali et al’s paper whether the basal F levels were taken in the early morning (which is essential). The MTP test, utilized in this study, is a gold standard adrenal function test with clearly defined criteria for HPAS. Analyses based on this test can be expected to produce valid results.\(^3\)

In a recently published genome-wide association study (GWAS) investigating susceptibility to HPAS in asthmatic children, screening was not performed for rs41423247 (GC).\(^7\) Instead rs591118, an intronic variant of the PDGFD gene, was significantly associated with HPAS. This particular SNP was not included in the current study, because the study was already completed when the paper reporting rs591118 was published.

In the current study, CC of rs41423247 is associated with higher ACTH levels when compared to the GC and GG genotypes. In adults with depression on the other hand, a dexamethasone/corticotropin-releasing hormone (CRH) test revealed that a poor response to anti-depressive therapy was associated with marginally lower median peak ACTH levels in patients with the CC genotype.\(^8\) The results of this study could however have been confounded by the use of a variable dose of triiodothyronine given to some patients, which could have induced unrecognised adrenal insufficiency in individuals at risk.\(^9\) Tsartsali et al\(^5\) and others\(^10\) reported that the GC genotype was associated with a significantly higher basal F level than the CC genotype. The GG genotype conversely was associated with lower F levels after a dexamethasone suppression test in Dutch elderly individuals.\(^11\) The authors interpreted this as “hypersensitivity” to glucocorticoids. However, on re-analysis of their paper, CC is associated with a significantly higher F level post-dexamethasone, suggesting glucocorticoid resistance of the CC genotype.

Both the HPAS-protective effect and the ACTH response seem to be inherited in an additive and dominant fashion. A similar phenotype is therefore expected with the heterozygous GC and homozygous CC genotype, except that the phenotypic effect of the latter should be more pronounced. Although the overall sample size was probably adequate, the low minor allele frequency of 23% (compared to 38% in Europe\(^12\)) and the low CC genotype frequency of 6% (compared to 40% in Greek children), limit the interpretation of the study. This resulted in overlapping CIs of the GC and CC genotypes and an incaulculable lower bound of the CC CI. However, the additive genetic effect accounted for a huge effect size and a highly significant p-value (<2 x 10\(^{-16}\)), which even far exceeds the required significance threshold for a GWAS (5 x 10\(^{-8}\)).\(^13\) Hence, pending confirmation in a larger study, the most suitable marker for HPAS protection would still be rs41423247(CC) of theNR3C1 gene.

The biochemical phenotype attributed to both CC and GC genotypes of rs41423247 is akin to the one de-
scribed in the syndrome of primary generalized glucocorticoid resistance (PGGR).\textsuperscript{14-16} In this syndrome there is dysregulation of F feedback at hypothalamic and pituitary level, resulting in elevated ACTH and possibly F levels. In the current study, early morning basal F levels of the children with rs41423247 were not elevated (results not shown). Clinical features of this syndrome e.g. hypertension, hirsutism or premature pubarche were not detected. Hence, the biochemical phenotype of rs41423247 (GC and CC) could represent tissue-specific glucocorticoid resistance.\textsuperscript{16} More likely, it is just a common variant in the population (40\% in our sample) presenting with reduced glucocorticoid sensitivity.\textsuperscript{16}

PGGR is associated with 26 mutations\textsuperscript{16} and 8 pathogenic SNPS\textsuperscript{17} in the exome of NR3C1 at 5q31.3, while rs41423247 falls within the intronic regulatory region of the gene,\textsuperscript{12} confirmed independently by variant browser analysis (result not shown). Mechanistically, impaired promoter function would lead to impaired DNA transcription, RNA translation and protein production.

In a recently published study on asthma children in Turkey, the FEV\textsubscript{1} of children with the GG genotype improved more after a high dose of ICS than children with the CG + CC genotype.\textsuperscript{18} In the current study, three of five children with the CC genotype had a suboptimal FEV\textsubscript{1} (result not shown). Unfortunately the number is too small to draw any valid conclusions from it.

In the same study population, it was previously shown that, as the BMI z-score rises, the ACTH level rises.\textsuperscript{1} In the current study, an association between CC genotype and BMI z-score could be demonstrated. No association of genotype was found with overweight or obesity (results not shown). In Dutch elderly the GG genotype was associated with a lower BMI\textsuperscript{11}, in line with the current study. However, others have found the opposite.\textsuperscript{19} The higher BMI z-score associated with rs41423247 is probably mediated through periodically elevated F levels. However, as shown, the BMI z-score does not confound the SNP effect on the ACTH level.

As findings in Greek and South African children (and possibly Turkish children and Dutch adults) are similar, it is likely that the results of this study are generalizable across all populations and ages. Pathophysiological and molecular explanations of the observed phenotype should also be equally applicable to all humans. However, inadvertently the study is limited by the small sample of children with the CC genotype. Hence, a larger study should be performed to confirm the finding. In addition, a “proof of concept” study (screening asthmatic children for rs41423247 (CC), and when positive, testing them with the MTP test to confirm the phenotype) should be done. If confirmed, rs41423247 (CC), as a marker of HPAS protection, should be screened for in asthmatic children on glucocorticoids. This would allow clinicians to choose which children can be safely treated with higher doses of glucocorticoids – a first step towards personalized medicine in asthma therapy.

In conclusion, rs41423247 (GC) of the NR3C1 gene was found to be protective against HPAS. The CC genotype was associated with the highest ACTH levels. The C allele is inherited as an additive and dominant trait. The GC genotype was independently associated with BMI z-score. If confirmed by a larger and a “proof of concept” study, rs41423247 (CC) could well be a marker of HPAS protection, thereby allowing for personalized treatment decisions to be made in asthma care.

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