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CHOICE

# Glucose tolerance, insulin sensitivity and $\beta$ -cell function in patients with rheumatoid arthritis treated with or without low-to-medium dose glucocorticoids

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## ABSTRACT

**Objectives** To compare glucose tolerance and parameters of insulin sensitivity and  $\beta$ -cell function between chronic glucocorticoid (GC)-using and GC-naïve patients with rheumatoid arthritis (RA).

**Methods** Frequently sampled 75 g oral glucose tolerance tests were performed in 58 chronic GC-using and 82 GC-naïve patients with RA with established disease, with no known type 2 diabetes mellitus (T2DM), and 50 control subjects of comparable age with normal glucose tolerance. The associations between cumulative GC dose and disease characteristics and glucose tolerance state, insulin sensitivity and  $\beta$ -cell function were tested using multivariate linear and logistic regression models, correcting for patient characteristics.

**Results** Glucose tolerance state, insulin sensitivity and  $\beta$ -cell function did not differ between the two RA populations; de novo T2DM was detected in 11% and impaired glucose metabolism in 35% of patients with RA. In patients with RA, cumulative GC dose was associated with T2DM, which seemed mostly driven by the effects of cumulative GC dose on insulin resistance; however, the association decreased when corrected for current disease activity. Patients with RA had decreased insulin sensitivity and impaired  $\beta$ -cell function compared with controls, and multivariate regression analyses showed a negative association between the presence of RA and insulin sensitivity.

**Conclusions** GC-using and GC-naïve patients with RA had comparable metabolic parameters, and had decreased insulin sensitivity and  $\beta$ -cell function as compared with healthy controls. Although cumulative GC dose was shown to have a negative impact on glucose tolerance state and insulin sensitivity, confounding by indication remains the main challenge in this cross-sectional analysis.

## INTRODUCTION

Patients with rheumatoid arthritis (RA) are at increased risk of developing cardiovascular disease, comparable to the risks seen in subjects with type 2 diabetes mellitus (T2DM).<sup>1</sup> Additional impairment of glucose metabolism may contribute significantly to the accelerated atherogenesis in patients with RA.<sup>2</sup> Fasting and postprandial glucose metabolism are determined by  $\beta$ -cell function (insulin production) and by the peripheral effects of insulin (insulin sensitivity) which increases glucose uptake in skeletal muscle and decreases glucose production by the liver. Glucocorticoids (GCs) impair hepatic and peripheral insulin sensitivity and induce

$\beta$ -cell dysfunction; however, the precise underlying mechanisms are still being investigated.<sup>3–4</sup> Previously, patients with RA were shown to have impaired fasting insulin sensitivity (homeostatic model assessment of insulin resistance (HOMA-IR) and fasting  $\beta$ -cell function (HOMA-B)), which correlated with disease activity and markers of inflammation.<sup>5–7</sup> Consequently, prevalent diabetes was estimated to be up to 15–19% in patients with RA,<sup>8–9</sup> an increased number as compared with the estimated T2DM prevalence of 4–8% in middle-aged men and women in the general population.<sup>10</sup>

The role of GCs in glucose intolerance in patients with RA has been one of paradox. Other than the electrolyte balance-regulating activity of their family member mineralocorticoids, GCs derived their name through their carbohydrate-regulating abilities.<sup>11</sup> GCs play essential roles in glucose, lipid and protein metabolism in the fasted state, providing substrate for oxidative metabolism by increasing adipose tissue lipolysis (glycerol and non-esterified free fatty acids), skeletal muscle proteolysis and hepatic glucose production.<sup>12</sup> The effects of GCs on glycogenolysis and glycogen synthesis are at present less clear. It is evident that GCs increase endogenous glucose production by the liver by enhancing gluconeogenesis. The contribution of glycogenolysis to endogenous glucose production is less well established.<sup>13</sup> On the one hand, in animal models and in short-term clinical trials in healthy subjects, GCs were shown to impair glucose metabolism by weakening hepatic and peripheral insulin sensitivity and by inducing  $\beta$ -cell dysfunction.<sup>13</sup> In retrospective, population-based studies, GC therapy was associated with incident diabetes,<sup>14</sup> and the need for blood-glucose-lowering treatment in a cumulative dose-dependent way.<sup>15</sup> In retrospective studies in patients with RA, GC exposure was shown to correlate with insulin resistance,<sup>16</sup> and to predict diabetes.<sup>17</sup> On the other hand, the use of GCs in chronic inflammatory states may improve glucose tolerance by their anti-inflammatory effects, as was demonstrated in a number of short-term studies using GC treatment;<sup>18–19</sup> this was also shown in a study using methotrexate.<sup>20</sup> In addition, confounding by indication should be kept in mind when evaluating the relation between GC use and glucose tolerance in patients with RA in observational studies. This is the possibility that patients with higher cumulative inflammation (ie, higher disease activity), resulting in a priori increased insulin resistance, were more likely to be given high-dose GCs

than those with less inflammation (disease) activity. Thus, the impact of GC treatment on glucose metabolism in patients with RA requires further clarification.

Previous studies that have examined the effects of GCs on glucose tolerance, insulin sensitivity and  $\beta$ -cell function in patients with RA included a small number of patients,<sup>18</sup> or relied solely on fasting parameters, that is, HOMA-IR and HOMA-B.<sup>16</sup> As mentioned above, as insulin sensitivity and insulin secretion are inter-related, the use of the HOMA formulas, both of which use the same fasting variables—that is, fasting plasma insulin and glucose, may not be appropriate to discern changes in insulin sensitivity from changes in insulin secretion.<sup>21</sup> For instance, if two subjects have the same fasting glucose level, but, one person achieves this with higher fasting insulin levels, this person is more insulin resistant (thus has lower insulin sensitivity), which is expressed by a higher HOMA-IR score. Similarly, from fasting glucose and insulin levels, a HOMA-B score is calculated which gives an impression of  $\beta$ -cell function. Although these model-derived indices are well validated, they provide no information about the stimulated, postload state.<sup>21</sup> From dynamic tests, such as the frequently sampled oral glucose tolerance test (OGTT), indices of postload insulin sensitivity and glucose-stimulated  $\beta$ -cell function may be calculated, in order to provide more detailed information on glucose metabolism.<sup>22</sup>

Therefore, in this study, we compared glucose tolerance and (fasting and dynamic) parameters of insulin sensitivity and  $\beta$ -cell function from frequently sampled OGTTs in a large group of chronic GC-using patients with RA versus GC-naïve patients with RA. Furthermore, we included a control group of comparable age to create a perspective of our OGTT findings in patients with RA, and to assess the association of RA itself with measures of insulin sensitivity and  $\beta$ -cell function in subjects with normal glucose tolerance. Finally, we assessed the association between cumulative GC dose and disease characteristics with these metabolic parameters.

## METHODS

### Population

Patients with RA with established disease—that is, defined as having a disease duration of >2 years, were recruited in four rheumatology clinics in the region of Utrecht, The Netherlands. Patients were either current and chronic GC users (RA+GC), which indicated GC treatment for at least 3 months, or they were GC usage naïve (RA–GC). Known T2DM (defined as receiving treatment) was an exclusion criterion. We included a control group (controls) with normal glucose tolerance and without first-degree relatives with T2DM, consisting of individuals who had undergone an OGTT for screening purposes for other studies at the Diabetes Centre of the VU University Medical Centre in Amsterdam. Accordingly, this group consisted of relatively overweight predominantly male individuals. We did not match the control population to the RA population, since our main focus was on studying the effects of GCs in patients with RA (particularly compared with GC-naïve patients with RA). The healthy control group in our study served to show the perspective of the values in the patients with RA. An independent ethics committee approved the study and all subjects provided written informed consent before participation; the protocol was according to the 'Declaration of Helsinki'.

### Protocol

Participants visited the clinic after an overnight fast of a minimum of 10 h. A physical examination, including recording

of height, weight and waist circumference was performed and fasting blood tests were carried out in all patients. In the patients with RA, in addition, the Disease Activity Score (DAS28 and DAS28-C-reactive protein)<sup>23,24</sup> was calculated; history of disease-modifying antirheumatic drug (DMARD) taking, laboratory measurement of anti-citrullinated protein antibodies (ACPA) and x-ray examinations of hands and feet (to detect erosive damage) were also performed. Finally, all participants underwent a 2 h 75 g OGTT. Blood samples for determination of glucose, insulin and C-peptide were collected at times 0, 10, 20, 30, 60, 90 and 120 min, starting immediately after the ingestion of the 75 g glucose solution. Since insulin clearance may vary considerably between subjects,<sup>25</sup> plasma C-peptide levels may provide additional information on  $\beta$ -cell function.

### Analytical determinations

Plasma glucose was measured using a chemical technique on a DXC-800 analyser (Beckman Coulter, Los Angeles, California, USA). Plasma insulin was measured using an immunometric technique on an IMMULITE 1000 Analyzer (Siemens Medical Solutions Diagnostics, Los Angeles, California, USA). Plasma C-peptide was measured using an electrochemiluminescence immunoassay on the Modular E170 (Roche Diagnostics GmbH, Mannheim, Germany).

### Data analysis

Normal glucose tolerance was defined as fasting plasma glucose (FPG) <6.1 mmol/l and a 2 h glucose value <7.8 mmol/l; impaired glucose metabolism (IGM) as FPG between 6.1 and 7.1 mmol/l or a 2 h glucose value between 7.8 and 11.1 mmol/l; T2DM as FPG >7.1 mmol/l or a 2 h glucose value >11.1 mmol/l. Areas under the 2 h glucose ( $AUC_{gluc}$ ), insulin ( $AUC_{ins}$ ) and C-peptide ( $AUC_{c-pep}$ ) curves were determined using the trapezoidal rule. Insulin sensitivity in the fasted state was computed by HOMA-IR.<sup>26</sup> Estimated metabolic clearance rate ( $MCR_{est}$ /Stumvoll Index) and oral glucose insulin sensitivity (OGIS) were used to estimate postload insulin sensitivity.<sup>27</sup> Various measures of  $\beta$ -cell function were calculated: HOMA-B was derived from fasting measures.<sup>26</sup> Dynamic measures of  $\beta$ -cell function were derived from OGTT data and included:  $AUC_{c-pep}/AUC_{gluc}$  ratio over the 2 h period and the Insulinogenic Index (IGI):  $(insulint=30 - insulint=0)/(gluct=30 - gluct=0)$ , as measures of early insulin secretion.<sup>28</sup> The oral Disposition Index (DI) was calculated by multiplying IGI and OGIS, to adjust insulin secretion for insulin sensitivity. Insulin clearance was calculated by dividing  $AUC_{c-pep}$  and  $AUC_{ins}$ .<sup>25</sup>

### Statistical analysis

#### Comparison of parameters of glucose tolerance state, insulin sensitivity and $\beta$ -cell function of RA $\pm$ GC groups and controls

Data were presented as mean values  $\pm$  SD and as median (IQR) for non-normal distribution. Intergroup differences in continuous outcomes were tested by analysis of variance, and with the Kruskal–Wallis tests in cases of non-normal distribution. Differences between groups in dichotomous outcomes were tested with the  $\chi^2$  test. Post hoc Bonferroni correction was applied to multiple testing by multiplying the p value by 2 (3 groups minus 1).

#### Associations between patient and disease characteristics and parameters of glucose tolerance state, insulin sensitivity and $\beta$ -cell function

Associations between known determinants of insulin sensitivity and  $\beta$ -cell function (age, waist circumference, body mass

**Table 1** Baseline characteristics

	Controls	RA–GCs	RA + GCs	p Value controls*	
				Vs RA–GCs	Vs RA + GCs
N	50	82	58	–	–
Age (years)	56±8	57±12	59±12	1.0	0.4
Female (%)	38	71	71	<0.001	<0.001
BMI (kg/m <sup>2</sup> )	29±4	25±4	26±6	<0.001	0.008
Waist circumference male (cm)	104±10	95±8	94±10	0.002	0.005
Waist circumference female (cm)	100±12	82±11	91±16	<0.001	0.03
Increased waist† (%)	62	24	35	<0.001	0.003
SBP (mm Hg)	125±10	124±18	125±17	1.0	1.0
DBP (mm Hg)	80±7	73±10	73±10	0.001	0.002
Hypertension‡ (%)	10	23	26	0.6	0.3
Antihypertensive drugs (%)	–	24	29	–	–
FPG (mmol/l)	5.4±0.5	5.5±0.7	5.3±0.7	0.6	1.0
Triglycerides (mmol/l)	1.2±0.4	1.0±0.5	1.2±0.7	0.2	1.0
LDL (mmol/l)	3.3±0.9	3.4±0.9	3.4±1	1.0	1.0
HDL male (mmol/l)	1.4±0.3	1.1±0.3	1.4±0.4	0.02	1.0
HDL female (mmol/l)	1.6±0.5	1.5±0.4	1.6±0.4	0.4	1.0
Total cholesterol (mmol/l)	5.3±0.9	5.2±1.0	5.4±1.2	1.0	1.0
Dyslipidaemia‡ (%)	50	82	62	<0.001	0.3
Hypercholesterolaemia‡ (%)	78	76	76	0.6	0.7
Statin use (%)	–	12	7	–	–
<b>RA-characteristics</b>				<b>p Value</b>	
Duration of RA (years)		13±8	13±8	0.6	
Diabetes§ (%)		9	14	0.3	
IGM§ (%)		37	33	0.6	
Current DMARD use					
Synthetic (%/n)		89/1.2	78/1.2	0.07	
Biological (%)		21	55	<0.001	
Historic DMARD use					
Synthetic (%/n)		71/1.9	71/2.8	1.0	
Biological (%/n)		7/1.3	24/1.9	0.005	
DAS28 (no dimension)		2.8±1.3	3.5±1.2	0.002	
Tender joint count		0 (0–3)	2 (0–5)	0.004	
Swollen joint count		0 (0–1)	1 (0–2)	0.09	
General well-being (VAS 0 (good) to 100)		26±21	38±25	0.002	
ESR (mm/h)		11 (8–21)	14 (9–28)	0.1	
Anti-CCP positive (%)		71	71	0.9	
Any erosive damage at x-ray of hands or feet (%)		72	81	0.2 (only hand erosions p=0.06)	
Cumulative dose GCs (g. prednisone equivalent)		0	13 (7–27)	–	
Daily dose (mg)		0	6.3 (5–10)	–	
Dexamethasone pulse (% ever used pulse/mean n pulses)		0	19/2	–	

Data represent means±SD or median (IQR) when data were not normally distributed. Intergroup differences in continuous outcomes were tested by ANOVA, and with both the Kruskal–Wallis and Mann–Whitney tests in cases of non-normal distribution. Differences between groups in dichotomous outcomes were tested with the  $\chi^2$  test. Post hoc Bonferroni correction was applied in cases of multiple testing (p value  $\times 2$ ).

\*p Value controls is the intergroup difference tested by ANOVA, Mann–Whitney or  $\chi^2$  test with the Bonferroni post hoc test. p Values of the RA–GC RA+GC difference were not depicted; the only significant/trend differences were male HDL p=0.07; dyslipidaemia p=0.009.

†Increased waist circumference was defined as >102 cm in men and >88 cm in women. Hypertension was defined as  $\geq 140$  mm Hg systolic or 90 mm Hg diastolic pressure.

‡Dyslipidaemia was defined as triglycerides >1.7 mmol/l and/or HDL-cholesterol < 0.9 mmol/l (male) and <1 mmol/l (female).

Hypercholesterolaemia was defined as total cholesterol >5 mmol/l and/or LDL-cholesterol >3 mmol/l.

§Diabetes was defined as either FPG $\geq 7.1$ , or  $\geq 11$  at 120 min of OGTT; IGM was defined as either FPG (<7.1 and >6.1), or impaired glucose tolerance (<11 and >7.8 at 120 min of OGTT).

ANOVA, analysis of variance; BMI, body mass index; CCP, cyclic citrullinated peptide; DAS28, Disease Activation Score using 28 joints; DBP, diastolic blood pressure; DMARD, disease-modifying antirheumatic drug; ESR, erythrocyte sedimentation rate; FPG, fasting plasma glucose; GC, glucocorticoid; HDL, high-density lipoprotein; IGM, impaired glucose metabolism; LDL, low-density lipoprotein; OGTT, oral glucose tolerance test; RA, rheumatoid arthritis; RA + GCs, patients with rheumatoid arthritis, currently using glucocorticoids; RA–GCs, patients with rheumatoid arthritis, glucocorticoid-naïve; SBP, systolic blood pressure; VAS, Visual Analogue Scale.

index (BMI) and insulin clearance) and gender, and, within the RA populations, cumulative and daily GC dose and disease activity (DAS28 and its components, DMARD use, hand or feet erosions on x-ray examination, ACPA) and parameters of insulin sensitivity and  $\beta$ -cell function were investigated with linear regression analysis; associations of the above

factors with IGM and T2DM were investigated with logistic regression analyses. Non-normally distributed variables were log transformed when used in multivariate linear regression analysis. In the multivariate analyses with OGTT outcomes as dependent variable, where the RA populations were compared with the controls, patients with RA with previously unknown

T2DM, IGM or with first-degree relatives with T2DM were excluded (ie, fitting the exclusion criteria of controls). SPSS for Mac version 16.0 (SPSS, Chicago, Illinois, USA) was used for all statistical analyses. A p value <0.05 was considered statistically significant.

**RESULTS**

**Baseline characteristics**

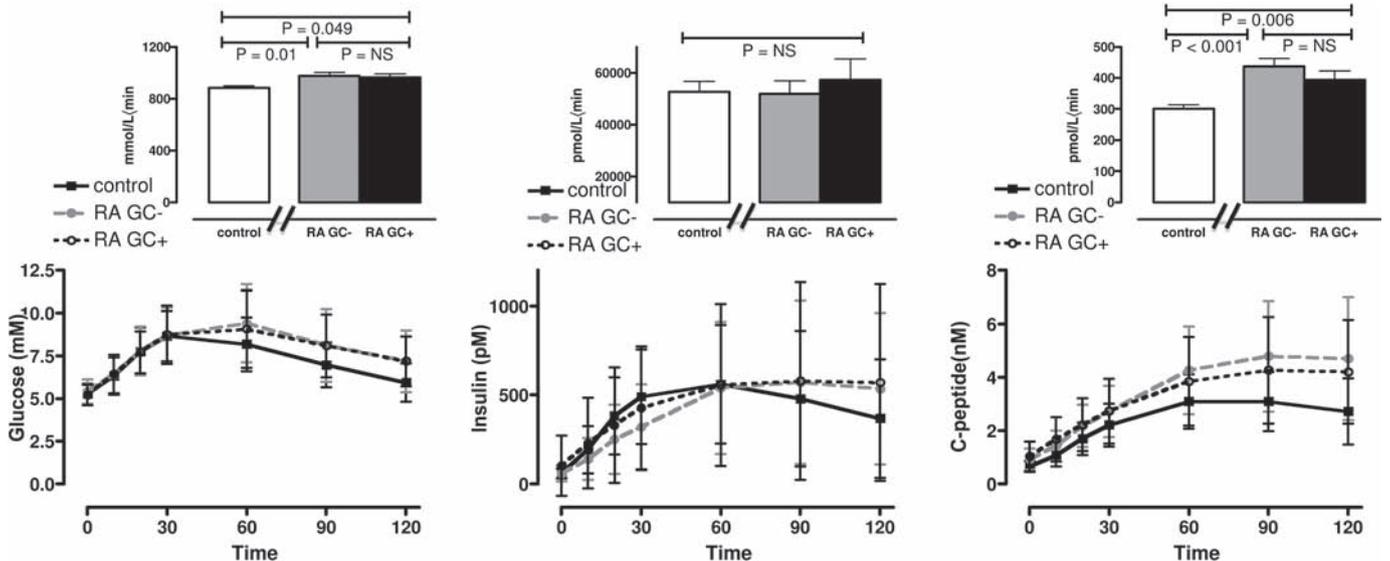
After screening 167 patients with RA, a total of 140 middle-aged patients with established RA were included; 82 were GC naïve (RA-GC) and 58 were current GC users (RA+GC). Of the 27 excluded patients with RA, 16 were patients with known T2DM (11 RA-GC and 5 RA+GC). In addition, 50 controls with comparable age were recruited. Subject characteristics are provided in table 1. As compared with patients with RA, controls had a higher percentage of male gender, and had a higher BMI and waist circumference; these factors were corrected for in the multivariate models. The RA groups had similar anthropometrics, but RA+GC had higher disease activity than RA-GC (table 1).

**Glucose tolerance state**

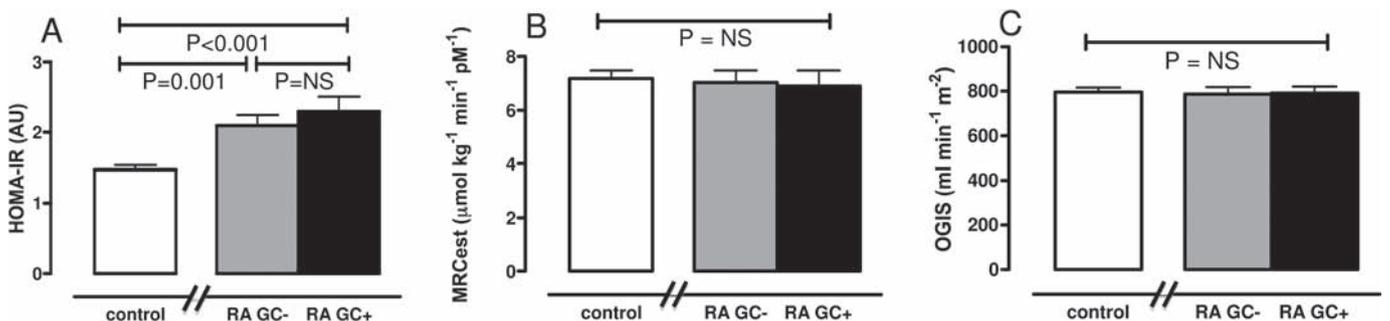
The prevalence of previously unknown T2DM was comparable between the two RA groups (table 1). If those patients with RA who were excluded because of known T2DM (n=27) had been included, then the prevalence of T2DM would have been 19% (RA-GC 18% vs RA+GC 21%, p=0.9). Within the RA groups, both cumulative and daily prednisolone dose was associated with incident T2DM in univariate analyses (OR cumulative dose (g): 1.04; p=0.002; daily dose (mg): 1.13; p=0.048). This association was sustained after adjusting for disease activity and patient characteristics (DAS28, ACPA, erosions, DMARD history, disease duration, age, BMI, waist circumference and gender; OR cumulative dose (g): 1.04; p=0.03; daily dose (mg): 1.15; p=0.13), whereas it was decreased and less significant (a trend) after adjustment for current disease activity alone and patient characteristics (DAS28, age, BMI, waist circumference and gender; OR cumulative GC dose (g): 1.02; p=0.08; daily dose (mg): 1.11; p=0.3).

**Metabolic responses during OGTT**

Glucose levels during the OGTT did not differ between the RA groups, whereas AUC<sub>gluc</sub> was higher in patients with RA than in



**Figure 1** Glucose and insulin levels during oral glucose tolerance test (OGTT). Mean (±SD) of glucose and C-peptide levels, respectively, are shown during an OGTT for control subjects (control), GC-naïve patients with RA (RAGC-) and GC-using patients with RA (RAGC+). Intergroup differences were tested by analysis of variance, and with the Kruskal–Wallis test in cases of non-normal distribution. Post hoc Bonferroni correction was applied in cases of multiple testing (p value × 2). GC, glucocorticoid; RA, rheumatoid arthritis.



**Figure 2** Insulin sensitivity indices. Mean (±SD) of insulin sensitivity indices (A) HOMA-IR (Fasting Index), and of the dynamic parameters (B) MCR<sub>est</sub> Index and (C) oral glucose insulin sensitivity (representing glucose clearance during a 2 h oral glucose tolerance test) are shown. Intergroup differences were tested by analysis of variance, and with the Kruskal–Wallis test in case of non-normal distribution. Post hoc Bonferroni correction was applied in cases of multiple testing (p value × 2). GC, glucocorticoid; HOMA-IR, homoeostatic model assessment of insulin resistance; MCR<sub>est</sub>, estimated metabolic clearance rate; RAGC-, GC-naïve patients with RA; RAGC+, GC-using patients with RA.

**Table 2** Association of risk factors and disease characteristics with HOMA-IR and MCR<sub>est</sub>

Patient characteristics	$\beta$ (95% CI) (RA-patients (n=56), controls (n=50))	$\beta$ (95% CI) (RA-patient only (n=140))	$\beta$ (95% CI) (RA-patient only (n=140))
Three multivariate regression analysis models with HOMA-IR as dependent variable			
<b>RA–GC*</b>	<b>0.6 (0.08, 1.0)</b>		
<b>RA+GC*</b>	<b>1.0 (0.6, 1.5)</b>		
<b>Waist circumference</b>	0.02 (–0.01 to 0.04)	0.02 (–0.003 to 0.05)	<b>0.03 (0.0002 to 0.06)</b>
Age	0.002 (–0.02 to 0.02)	0.008 (–0.007 to 0.02)	0.006 (–0.01, 0.02)
BMI	0.03 (–0.05 to 0.1)	0.05 (–0.2 to 0.1)	0.03 (–0.04 to 0.1)
Female gender	0.2 (–0.2 to 0.6)	–0.1 (–0.5 to 0.3)	–0.06 (–0.5 to 0.4)
<b>Cumulative GC dose (mg)†</b>		<b>0.01 (0.003 to 0.02)</b>	<b>0.01 (0.003 to 0.02)</b>
Daily GC dose† (mg)		0.01 (–0.04 to 0.05)	0.005 (–0.04 to 0.05)
DAS28‡			0.1 (–0.04 to 0.3)
Any erosions of the hands or feet			–0.3 (–0.7 to 0.2)
Past DMARDs (n)§			–0.06 (–0.2 to 0.05)
ACPA			0.2 (–0.7 to 0.2)
Disease duration (years)			0.02 (–0.008 to 0.04)
Four multivariate regression analysis models with MCR <sub>est</sub> as dependent variable			
<b>RA–GC*</b>	0.4 (–0.5 to 1.3)		
<b>RA+GC*</b>	0.4 (–0.5 to 1.2)		
<b>Waist</b>	–0.02 (–0.07 to 0.03)	–0.06 (–0.1 to 0.01)	<b>–0.09 (–0.2 to –0.02)</b>
<b>Age</b>	–0.03 (–0.06 to 0.005)	<b>–0.07 (–0.1 to –0.03)</b>	<b>–0.07 (–0.1 to –0.03)</b>
<b>BMI</b>	<b>–0.3 (–0.4 to –0.1)</b>	<b>–0.2 (–0.4 to –0.06)</b>	<b>–0.1 (–0.3 to –0.06)</b>
Female gender	–0.8 (–1.6 to 0.08)	–0.08 (–1.3 to 1.1)	–0.5 (–1.7 to 0.7)
Cumulative GC dose (g)†		0.002 (–0.01 to 0.04)	0.01 (–0.01 to 0.04)
Daily GC dose† (mg)		–0.07 (–0.19 to 0.05)	–0.04 (–0.2 to 0.08)
<b>DAS28‡</b>			<b>–0.5 (–0.9 to –0.1)</b>
Any erosions of the hands or feet			0.8 (–0.3 to 2.0)
Past DMARDs (n)§			0.2 (–0.06 to 0.5)
ACPA			–0.7 (–1.8 to 0.4)
Disease duration (years)			0.03 (–0.04 to 0.1)

Three multivariate models, correcting for age, BMI and female gender were used to test associations with insulin sensitivity parameters HOMA-IR and MCR<sub>est</sub>: (1) tested RA populations (as compared with healthy controls), (2) tested cumulative and daily GC dose and (3) tested disease activity (DAS28 and its components, DMARD use, hand or feet erosions on x-ray examination, ACPA). Significant associations ( $p < 0.05$ ) are depicted in bold

\*RA–GC and RA+GC populations are tested with the healthy control group used as the reference population; only patients with RA with normal glucose tolerance during OGTT and without first-degree relatives with type 2 diabetes mellitus were included in the model for the comparison with healthy controls.

†Cumulative GC dose, daily GC dose were separately tested in models of the RA population only.

‡Disease parameters (DAS28, DAS28-CRP, CRP, ESR) were separately tested in the model of the RA population only; only DAS28 is depicted, whereas ESR was also significantly associated with MCR<sub>est</sub> in this model.

§Number of DMARDs (both synthetic and biological; not GCs) that were used by a patient in the past.

ACPA, anti-citrullinated protein antibodies; BMI, body mass index; CRP, C-reactive protein; DAS28, Disease Activation Score measured by questioning general health, physical examination of 28 joints and ESR or CRP; DMARD, disease-modifying antirheumatic drug; ESR, erythrocyte sedimentation rate; GC, glucocorticoid; HOMA-IR, homeostatic model assessment of insulin resistance; MCR<sub>est</sub>, Insulin Sensitivity Index by Stumvoll; OGTT, oral glucose tolerance test; RA, rheumatoid arthritis.

controls, which was mostly driven by higher glucose levels during the final 90 min of the test (figure 1A). Insulin levels were comparable between all groups (figure 1B); however, C-peptide secretion was higher in the RA groups than in controls (figure 1C), with no difference between the RA groups, indicating increased insulin clearance in patients with RA as compared with controls (data not shown). Insulin clearance was decreased in RA+GC as compared with RA–GC (data not shown).

### Parameters of insulin sensitivity

Parameters of both fasting (figure 2A) and postload insulin sensitivity (figure 2B,C) were comparable between the RA groups. In multivariate linear regression analyses (correcting for age, gender, BMI, waist circumference and disease activity) the presence of RA, waist circumference and cumulative GC dose were independent predictors of HOMA-IR (table 2). MCR<sub>est</sub> was negatively associated with DAS28, erythrocyte sedimentation rate, age, BMI and waist circumference (table 2); a similar pattern was observed for OGIS (data not shown). The healthy control group was more insulin sensitive in the fasted

state, but had similar postload insulin sensitivity to that of the RA groups.

### Parameters of $\beta$ -cell function

HOMA-B was higher in RA+GC than in RA–GC (figure 3A), while all dynamic measures of  $\beta$ -cell function were comparable between the RA groups (figure 3B,D). Positive associations between the presence of RA and cumulative GC use with HOMA-B were found (corrected for age, gender, BMI, waist circumference and disease activity; data not shown). Age and both RA–GC and RA+GC were negatively associated with IGI (corrected for age, gender, BMI, waist circumference; data not shown), whereas no patient characteristics correlated with the DI (table 3). As compared with healthy controls, patients with RA had higher basal C-peptide secretion (higher HOMA-B), but impaired early insulin secretion, also when corrected for insulin sensitivity (figure 3B,C). The total amount of C-peptide secreted during the entire OGTT relative to glucose levels, was higher in patients with RA than healthy controls (figure 3D).

**Table 3** Association of risk factors and disease characteristics with the DI

Independent variable	$\beta$ (95% CI) (RA-patients (n=56), controls (n=50))	$\beta$ (95% CI) (RA-patient only (n=140))	$\beta$ (95% CI) (RA-patient only (n=140))
Three multivariate regression analysis models with DI as dependent variable			
RA – GC*	–42489 (–96745 to 11767)		
RA + GC*	–23040 (–75891 to 29810)		
Waist circumference	–1778 (–4895 to 1338)	315 (–1831 to 2462)	274 (–2009 to 2557)
<b>Age</b>	<b>–3295 (–5219 to –1370)</b>	<b>–1280 (–2502 to –57)</b>	–1100 (–2385 to 186)
BMI	6205 (–1930 to 14340)	–2771 (–8193 to 2650)	–3070 (–8893 to 2752)
Female gender	–8287 (–52574 to 36000)	25773 (–10358 to 61904)	31014 (–6984 to 69013)
Cumulative GC dose (mg)†		–482 (–1307 to 343)	–284 (–1150 to 582)
Daily GC dose†		–885 (–4669 to 2899)	140 (–3731 to 4010)
DAS28‡			–4759 (–16525 to 7007)
Any erosions of the hands or feet			–14480 (–50730 to 21770)
Past DMARDs (n)§			662 (–7572 to 8896)
ACPA			24336 (–9300 to 57973)
Disease duration (years)			–1392 (–3447 to 664)

Three multivariate models, correcting for age, BMI and female gender were used to test associations with the  $\beta$ -cell parameter DI: (1) tested RA populations (as compared with healthy controls), (2) tested cumulative and daily GC dose and (3) tested disease activity (DAS28 and its components, DMARD use, hand or feet erosions on x-ray, ACPA).

Significant associations ( $p < 0.05$ ) are depicted in bold

\*RA – GC and RA + GC populations are tested against the control subject population; only patients with RA with normal glucose tolerance during OGTT and without first-degree relatives with type 2 diabetes mellitus were included for the comparison with control subjects. RA + GC was significantly associated with DI when these analyses were performed with log-transformed DI.

†Cumulative GC dose, daily GC dose were separately tested in models of the RA population only.

‡Disease parameters (DAS28, DAS28-CRP, CRP, ESR) were separately tested in the model of the RA population only; only DAS28 and significant correlations are depicted.

§Number of DMARDs (both synthetic and biological; not GCs) that were used by a patient in the past.

ACPA, anti-citrullinated protein antibodies; BMI, body mass index; CRP, C-reactive protein; DAS28, Disease Activation Score measured by questioning general health, physical examination of 28 joints and ESR or CRP; DI, Disposition Index; DMARD, disease-modifying antirheumatic drug; ESR, erythrocyte sedimentation rate; GC, glucocorticoid; RA, rheumatoid arthritis; RA + GCs, patients with rheumatoid arthritis currently using glucocorticoids; RA – GCs, glucocorticoid-naïve patients with rheumatoid arthritis.

### Effect modification

As compared with patients with RA, controls had a higher percentage of male gender, and had a higher BMI and waist circumference; in addition, RA+GC patients had higher disease activity than RA-GC patients (table 1). These factors were studied for effect modification using interaction terms in the regression models mentioned below, and were shown not to modify the effects of patient and disease characteristics on glucose metabolism outcomes in the multivariate models (data not shown).

### DISCUSSION

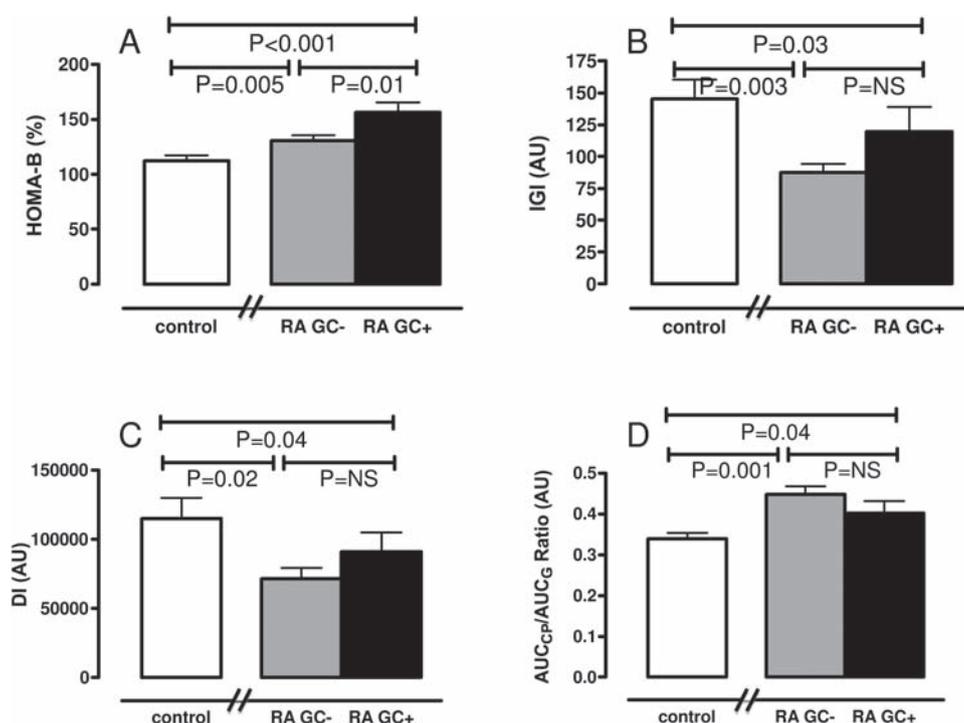
In this study of patients with RA with established disease, chronic GC users and GC-naïve patients had similar insulin sensitivity and  $\beta$ -cell function parameters; however, high cumulative or daily GC dose was associated with T2DM. In addition, in all patients with RA IGM and T2DM were frequently diagnosed, suggesting that glucose intolerance remains an underestimated problem in RA. As compared with a healthy control group, patients with RA had impaired insulin sensitivity and  $\beta$ -cell dysfunction, explaining their impaired metabolic state.

To our knowledge, this is the first study that has examined glucose metabolism in a relatively large sample of patients with RA with established disease in such detail. A few studies investigated glucose tolerance state in RA. One study showed an increased prevalence of T2DM in comparison with age-matched controls<sup>29</sup>; another study reported a 19% diabetes prevalence as part of a longitudinal medical record cohort on cardiovascular risk.<sup>8</sup> Both studies relied on self-reported T2DM and did not perform glucose measurements. Because of the OGTT measurements of glucose at 0 and 120 min we were now able to register 11% T2DM prevalence in patients with RA with established disease without known T2DM, and in addition, identify high-risk patients, by detecting 35% prevalence of IGM. This shows that glucose intolerance is a considerable and underestimated problem in patients with RA with established disease, and might (partially) explain their increased cardiovascular risk.<sup>1</sup>

The subject of insulin resistance in RA has been examined in recent years, but was only evaluated by the fasting measure HOMA-IR.<sup>2 5 6</sup> Unlike these studies, we were able to show a negative association of DAS28 with insulin sensitivity after correcting for potential confounders and other risk factors, which might have been because we also used stimulated measures of insulin sensitivity and because our sample size was larger.

Another important finding of our study was impaired  $\beta$ -cell function in patients with RA as compared with controls, as was shown by the decreased dynamic parameters IGI and DI, also when correcting for age, BMI and waist circumference (in the case of IGI). This indicates impaired insulin secretion during the early phase after glucose stimulation. So far, a limited number of other studies have reflected upon  $\beta$ -cell function in RA, and only used the fasting state measure HOMA-B. In one retrospective study HOMA-B was reduced in patients with RA with a higher level of inflammation in comparison with patients with RA with a lower level of inflammation,<sup>6</sup> which seems in line with our findings of impaired  $\beta$ -cell function in patients with RA compared with the control population. In our analysis, HOMA-B was higher in patients with RA, which indicates increased basal C-peptide secretion. This seems contradictory in view of the decrease in  $\beta$ -cell function parameters obtained in the stimulated state. However, HOMA-B should always be interpreted in the context of prevailing insulin resistance.<sup>21</sup> In our study, RA+GC participants had higher HOMA-IR values—that is, they were more insulin resistant. In order to maintain fasting glucose levels within the normal range, more insulin was secreted in the fasted state, which resulted in a higher HOMA-B score. However, this higher HOMA-B score does not imply improved  $\beta$ -cell function, but merely indicates the potential to compensate for reduced insulin sensitivity.

In addition, we examined the specific role of (cumulative or daily) GC dose and disease characteristics within patients with



**Figure 3** Parameters of  $\beta$  cell function. Mean ( $\pm$ SD) of  $\beta$  cell indices (A) HOMA-B (Fasting Index), and of the dynamic parameters (B) Insulinogenic Index and (C) Disposition Index, and of the (D)  $AUC_{c-pep}/AUC_{gluc}$  ratio are shown. Intergroup differences were tested by analysis of variance, and with the Kruskal–Wallis and Mann–Whitney tests in cases of non-normal distribution. Post hoc Bonferroni correction was applied in cases of multiple testing ( $p$  value  $\times 2$ ). AUC, area under the curve; DI, Disposition Index; GC, glucocorticoid; HOMA-B, homeostatic model assessment of  $\beta$  cell function; IGI, Insulinogenic Index; RAGC–, GC-naïve patients with RA; RAGC+, GC-using patients with RA.

RA and found strong indications that RA+GC patients were less glucose tolerant in a dose-dependent manner: Although no relation was shown between (cumulative or daily) GC dose and dynamic tests of insulin sensitivity and  $\beta$ -cell function, cumulative and daily GC dose were associated with previously unknown T2DM, and negatively affected fasting insulin sensitivity (HOMA-IR), independently of age, gender, BMI, waist circumference and disease activity. In addition, RA+GC patients had a decreased insulin clearance, indicating hepatic insulin resistance, which is explained by the steeper insulin curve in the first part of the OGTT. Our results are in line with one other retrospective study of non-diabetic patients with RA<sup>16</sup>; that study analysed a successive group of patients with RA and showed that ever having taken oral prednisone and/or high doses of pulsed GCs was independently associated with decreased insulin sensitivity independently of BMI.

We acknowledge some limitations in our study design; first, the difference in anthropometrics between controls (with normal glucose tolerance) and patients with RA—that is, controls were primarily recruited for other studies at the VUMC Diabetes Centre and therefore consisted of more male subjects and had a relatively high BMI, and lower insulin clearance. Compared with these control subjects, patients with RA were more insulin resistant and had more  $\beta$ -cell dysfunction. Although the use of this control population, as compared with more lean, insulin-sensitive controls, may be suboptimal, it is likely that the impact of RA and the associated proinflammatory state on glucose metabolism as described here, may even be underestimated. Besides, in multivariate analyses we corrected for these anthropometrics and, furthermore, the control subjects served mainly to create a perspective for the insulin resistance and  $\beta$ -cell parameters of patients with RA. A second point is confounding by indication that might have caused the effects of GCs on

glucose metabolism, since cumulative GC use might be a proxy for long-term disease activity, which itself influences glucose metabolism. This was shown also in our study by a decrease of the regression coefficient ( $\beta$ ) for the association between cumulative or daily GC dose and T2DM when disease activity (DAS28) was added to the multivariate regression model.

In conclusion, because of (1) the stimulated-state measurement of glucose metabolism parameters, (2) the size of our population and (3) the contrast with control subjects, we were able to draw firm conclusions about the prevalence of glucose tolerance abnormalities in patients with RA with established disease and to confirm the relation between RA (activity) and insulin resistance and  $\beta$ -cell dysfunction. Chronic GC use was associated with metabolic toxicity in a dose-dependent way, but this association was difficult to assess owing to confounding by indication.

Until there is more clarity about the problem of glucose intolerance in GC-using patients with RA, it remains important to keep the duration of GC use short and to use the lowest possible dose, as is advised by the European League Against Rheumatism recommendations on RA treatment,<sup>30</sup> and on systemic GC use.<sup>31</sup> The question of how harmful the diabetogenic effects of long-term GCs are in patients with established RA needs further assessment in longitudinal (randomised) trials. These trials should measure glucose metabolism with stimulated-state measures, and examine whether GCs exert direct metabolic toxicity or secondary toxicity owing to other GC-related phenomena, such as abdominal fat and adipocytokines, which are known mediators of metabolic toxicity in RA.<sup>32</sup>

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## REFERENCES

- Peters MJ, van Halm VP, Voskuyl AE, *et al*. Does rheumatoid arthritis equal diabetes mellitus as an independent risk factor for cardiovascular disease? A prospective study. *Arthritis Rheum* 2009;**61**:1571–9.
- Dessein PH, Joffe BI, Stanwix A, *et al*. The acute phase response does not fully predict the presence of insulin resistance and dyslipidemia in inflammatory arthritis. *J Rheumatol* 2002;**29**:462–6.
- Giorgino F, Almahfouz A, Goodyear LJ, *et al*. Glucocorticoid regulation of insulin receptor and substrate IRS-1 tyrosine phosphorylation in rat skeletal muscle in vivo. *J Clin Invest* 1993;**91**:2020–30.
- Lambillotte C, Gilon P, Henquin JC. Direct glucocorticoid inhibition of insulin secretion. An *in vitro* study of dexamethasone effects in mouse islets. *J Clin Invest* 1997;**99**:414–23.
- Chung CP, Oeser A, Solus JF, *et al*. Inflammation-associated insulin resistance: differential effects in rheumatoid arthritis and systemic lupus erythematosus define potential mechanisms. *Arthritis Rheum* 2008;**58**:2105–12.
- Dessein PH, Joffe BI. Insulin resistance and impaired beta cell function in rheumatoid arthritis. *Arthritis Rheum* 2006;**54**:2765–75.
- Shahin D, Eitoraby E, Mesbah A, *et al*. Insulin resistance in early untreated rheumatoid arthritis patients. *Clin Biochem* 2010;**43**:661–5.
- Maradit-Kremers H, Nicola PJ, Crowson CS, *et al*. Cardiovascular death in rheumatoid arthritis: a population-based study. *Arthritis Rheum* 2005;**52**:722–32.
- Simard JF, Mittleman MA. Prevalent rheumatoid arthritis and diabetes among NHANES III participants aged 60 and older. *J Rheumatol* 2007;**34**:469–73.
- Wild S, Roglic G, Green A, *et al*. Global prevalence of diabetes: estimates for the year 2000 and projections for 2030. *Diabetes Care* 2004;**27**:1047–53.
- Buttgereit F, da Silva JA, Boers M, *et al*. Standardised nomenclature for glucocorticoid dosages and glucocorticoid treatment regimens: current questions and tentative answers in rheumatology. *Ann Rheum Dis* 2002;**61**:718–22.
- McMahon M, Gerich J, Rizza R. Effects of glucocorticoids on carbohydrate metabolism. *Diabetes Metab Rev* 1988;**4**:17–30.
- van Raalte DH, Ouwens DM, Diamant M. Novel insights into glucocorticoid-mediated diabetogenic effects: towards expansion of therapeutic options? *Eur J Clin Invest* 2009;**39**:81–93.
- Gulliford MC, Charlton J, Latinovic R. Risk of diabetes associated with prescribed glucocorticoids in a large population. *Diabetes Care* 2006;**29**:2728–9.
- Gurwitz JH, Bohn RL, Glynn RJ, *et al*. Glucocorticoids and the risk for initiation of hypoglycemic therapy. *Arch Intern Med* 1994;**154**:97–101.
- Dessein PH, Joffe BI, Stanwix AE, *et al*. Glucocorticoids and insulin sensitivity in rheumatoid arthritis. *J Rheumatol* 2004;**31**:867–74.
- Wolfe F, Michaud K. Corticosteroids increase the risk of diabetes mellitus in rheumatoid arthritis and contribute to the risk of myocardial infarction and heart failure (abstract). *Ann Rheum Dis* 2004;**63**(Suppl 1):495.
- Svenson KL, Lundqvist G, Wide L, *et al*. Impaired glucose handling in active rheumatoid arthritis: effects of corticosteroids and antirheumatic treatment. *Metab Clin Exp* 1987;**36**:944–8.
- Hällgren R, Berne C. Glucose intolerance in patients with chronic inflammatory diseases is normalized by glucocorticoids. *Acta Med Scand* 1983;**213**:351–5.
- Dessein PH, Joffe BI, Stanwix AE. Effects of disease modifying agents and dietary intervention on insulin resistance and dyslipidemia in inflammatory arthritis: a pilot study. *Arthritis Res* 2002;**4**:R12.
- Wallace TM, Levy JC, Matthews DR. Use and abuse of HOMA modeling. *Diabetes Care* 2004;**27**:1487–95.
- Pacini G, Mari A. Methods for clinical assessment of insulin sensitivity and beta-cell function. *Best Pract Res Clin Endocrinol Metab* 2003;**17**:305–22.
- Prevoot ML, van 't Hof MA, Kuper HH, *et al*. Modified disease activity scores that include twenty-eight-joint counts. Development and validation in a prospective longitudinal study of patients with rheumatoid arthritis. *Arthritis Rheum* 1995;**38**:44–8.
- Wells G, Becker JC, Teng J, *et al*. Validation of the 28-joint Disease Activity Score (DAS28) and European League Against Rheumatism response criteria based on C-reactive protein against disease progression in patients with rheumatoid arthritis, and comparison with the DAS28 based on erythrocyte sedimentation rate. *Ann Rheum Dis* 2009;**68**:954–60.
- Kotronen A, Vehkavaara S, Seppälä-Lindroos A, *et al*. Effect of liver fat on insulin clearance. *Am J Physiol Endocrinol Metab* 2007;**293**:E1709–15.
- Levy JC, Matthews DR, Hermans MP. Correct homeostasis model assessment (HOMA) evaluation uses the computer program. *Diabetes Care* 1998;**21**:2191–2.
- Mari A, Pacini G, Murphy E, *et al*. A model-based method for assessing insulin sensitivity from the oral glucose tolerance test. *Diabetes Care* 2001;**24**:539–48.
- Kosaka K, Hagura R, Kuzuya T, *et al*. Insulin secretory response of diabetics during the period of improvement of glucose tolerance to normal range. *Diabetologia* 1974;**10**:775–82.
- Han C, Robinson DW, Jr, Hackett MV, *et al*. Cardiovascular disease and risk factors in patients with rheumatoid arthritis, psoriatic arthritis, and ankylosing spondylitis. *J Rheumatol* 2006;**33**:2167–72.
- Smolen JS, Landewé R, Breedveld FC, *et al*. EULAR recommendations for the management of rheumatoid arthritis with synthetic and biological disease-modifying antirheumatic drugs. *Ann Rheum Dis* 2010;**69**:964–75.
- Hoes JN, Jacobs JW, Boers M, *et al*. EULAR evidence-based recommendations on the management of systemic glucocorticoid therapy in rheumatic diseases. *Ann Rheum Dis* 2007;**66**:1560–7.
- Rho YH, Chung CP, Solus JF, *et al*. Adipocytokines, insulin resistance, and coronary atherosclerosis in rheumatoid arthritis. *Arthritis Rheum* 2010;**62**:1259–64.