Altered glycosylation of complexed native IgG molecules is associated with disease activity of systemic lupus erythematosus

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In addition to the redundancy of the receptors for the Fc portion of immunoglobulins, glycans result in potential ligands for a plethora of lectin receptors found in immune effector cells. Here we analysed the exposure of glycans containing fucosyl residues and the fucosylated tri-mannose N-type core by complexed native IgG in longitudinal serum samples of well-characterized patients with systemic lupus erythematosus. Consecutive serum samples of a cohort of 15 patients with systemic lupus erythematosus during periods of increased disease activity and remission were analysed. All patients fulfilled the 1982 American College of Rheumatology classification criteria. Sera of 15 sex- and age-matched normal healthy blood donors served as controls. The levels and type of glycosylation of complexed random IgG was measured with lectin enzyme-immunosorbent assays. After specifically gathering IgG complexes from sera, biotinylated lectins Aleuria aurantia lectin and Lens culinaris agglutinin were employed to detect IgG-associated fucosyl residues and the fucosylated tri-mannose N-glycan core, respectively. In sandwich-ELISAs, IgG-associated IgM, IgA, C1q, C3c and C-reactive protein (CRP) were detected as candidates for IgG immune complex constituents. We studied associations of the glycan of complexed IgG and disease activity according to the physician’s global assessment of disease activity and the systemic lupus erythematosus disease activity index 2000 documented at the moment of blood taking. Our results showed significantly higher levels of Aleuria aurantia lectin and Lens culinaris agglutinin binding sites exposed on IgG complexes of patients with systemic lupus erythematosus than on those of normal healthy blood donors. Disease activity in systemic lupus erythematosus correlated with higher exposure of Aleuria aurantia lectin-reactive fucosyl residues by immobilized IgG complexes. Top levels of Aleuria aurantia lectin-reactivity were found in samples taken during the highest activity of systemic lupus erythematosus. Our results showed that native circulating IgG complexes from active systemic lupus erythematosus patients expose fucosyl residues and their glycan core is accessible to soluble lectins. Two putative mechanisms may contribute to the increased exposure of these glycans: (1) the canonical N-glycosylation site of the IgG-CH2 domain; (2) an IgG binding non-IgG molecule, like complement or C-reactive protein. In both cases the complexed IgG may be alternatively targeted to lectin receptors of effector cells, e.g. dendritic cells.

Key words: IgG; immune complex; glycosylation; lectin-ELISA; Aleura aurantia; Lens culinaris; SLEDAI; physicians global assessment
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over time. The current view is that the SLE pathogenesis relates to abnormal apoptosis and deficient elimination of apoptotic material, such as chromatin. This eventually leads to autoantibody production and formation of circulating and/or tissue-bound IgG immune complexes (IC). IgG autoantibody-binding to tissue-exposed autoantigens and/or insufficient receptor-mediated clearance of circulating IC (CIC) via the reticuloendothelial system may impair hepatic IC clearance and cause extra-hepatic IC deposition. Altered clearance of apoptotic chromatin via CIC formation results in enhanced inflammation and tissue damage.

IgG is an example of a glycoprotein, characteristically carrying a complex N-linked biantennary glycan, consisting of a core heptasaccharide structure with variable additions of fucose and outer arm sugar residues like galactose and sialic acids, attached to asparagine-297 of the heavy chain CH2 domain. At least two dozen different Fc glycoforms, consisting of the heptasaccharide biantennary core with a combination of different numbers of core-fucose, galactose, bisecting N-acetyl glucosamine (GlcNAc) and terminal sialic acids, have been identified on polyclonal serum IgG and monoclonal IgG. Data from several studies indicate that these exposed glycans play important roles in structure and function of proteins; e.g. minor changes in a single glycan can affect protein folding and processing. In addition, the functionality of terminal sugars on IgG-N-glycans is of major biological importance. IgG without terminal galactose (G0) or core-fucose exhibit higher antibody dependent cell mediated cytotoxicity (ADCC) and monoclonal IgG. Data from several studies indicate that these exposed glycans play important roles in structure and function of proteins; e.g. minor changes in a single glycan can affect protein folding and processing. In addition, the functionality of terminal sugars on IgG-N-glycans is of major biological importance. IgG without terminal galactose (G0) or core-fucose exhibit higher antibody dependent cell mediated cytotoxicity (ADCC) and monoclonal IgG.

We employed the lectins *Aleuria aurantia* lectin (AAL) and *Lens culinaris* agglutinin (LCA) that bind to fucosyl residues and the fucosylated tri-mannose N-glycan core, respectively, to assess the glycosyl residues exposed and the fucosylated tri-mannose N-glycan core with a combination of different numbers of core-fucose, galactose, bisecting N-acetyl glucosamine (GlcNAc) and terminal sialic acids, have been identified on polyclonal serum IgG and monoclonal IgG.

The aims of the present study were to examine lectin targets associated with immobilized native complexed IgG in sera from patients with SLE. We analysed whether the exposure of fucosyl residues or the fucosylated tri-mannose N-glycan core of complexed IgG is related to SLE disease activity and disease phenotype. A regional Swedish register containing well-characterized SLE patients with longitudinal serum samples formed the basis of this investigation.

**Material and methods**

**Subjects**

15 SLE patients (13 female, two male; mean age 40.0 years; range 18–64) were recruited to the study on the basis of an episode of raised disease activity over a period of time. All patients took part in a prospective control programme (clinical lupus register in northeastern Gotha) at the Rheumatology department at Linköping University hospital and met the validated 1982 American College of Rheumatology (ACR) as well as the 2012 Systemic Lupus International Collaborating Clinics classification criteria. The mean number of fulfilled ACR criteria was 5.7 (range 4–7). Serial serum samples were drawn at each visit to the rheumatologist during the study period (2–10 visits per individual; mean 6) and the sera were kept frozen at −70°C until analysed. The physician’s global assessment of disease activity (PGA; 0–4) and the SLE disease activity 2000 (SLEDAI) was recorded at each visit. All patients showed signs of disease activity defined as a SLEDAI peak score of at least 6 (mean 11.9; range 6–30) for a median time of 16.9 months (range 6–26 months). The terms increased disease activity referred to the time-point of highest and decreased disease activity referred to the time-point of lowest SLEDAI score, respectively. Treatments with glucocorticoids and disease-modifying anti-rheumatic drugs were recorded. Clinical characteristics are summarized in Table 1. Fifteen sex- and age-matched normal healthy blood donors (NHD) (13 female, 2 male; mean age 40.3 years) served as controls.

**Routine laboratory analyses**

Laboratory analyses at all visits included erythrocyte, leukocyte and platelet counts, urine albumin/erythrocytes, creatinine, creatine kinase, anti-doublestranded (ds) DNA antibodies, erythrocyte sedimentation rate, C-reactive protein (CRP), complement protein C3 and C4, and classical haemolytic complement function. High sensitivity CRP (detection limit of 0.12 mg/L) was analysed by turbidimetry at the clinical chemistry of the Linköping
University Hospital. Complement analyses were performed at the University Hospitals in Uppsala or Linköping, Sweden. Microscope slides with fixed Crithidia luciliae (ImmunoConcepts) were used to analyse IgG class anti-dsDNA antibodies by indirect immunofluorescence microscopy (cut-off titre 1:10, corresponding to >99th percentile among healthy female blood donors). Anti-dsDNA antibody levels were end-point titration in two-fold dilution steps.

Analysis of CIC composition by IgG capture ELISA and determination of exposed IgG glycosyl residues by IgG capture lectin ELISA

To analyse the circulating nativecomplexed IgG, an IgG capture ELISA was performed. Ninety-six well microtitre plates (Nunc, F96 MaxiSorp) were coated with 2 μg/ml anti-human IgG F(ab')2-fragment (F(ab')2 goat anti-human serum IgG, H+L chain, Jackson Laboratories Immunoresearch) in coating buffer (0.1 M Na₂CO₃/NaHCO₃) pH 9.6 at 4°C overnight. Plates were blocked with 3% bovine serum albumin (Sigma-Aldrich) in phosphate buffered saline (PBS) containing 0.05% tween-20 (Roth) at 37°C for 2 hours. After blocking, patient sera or sera from normal healthy donors were diluted 1:1000 in PBS-tween and incubated at 37°C for 2 hours. These serum dilutions resulted in a saturating binding of IgG to the ELISA plate. After every incubation step the plates were washed three times with 200 μl PBS-tween. Horseradish-peroxidase (HRP)-labelled anti-human IgG-antibody (Southern Biotech), anti-human IgA-antibody (Southern Biotech), anti-human IgM antibody (Southern Biotech), anti-human C1q (Abcam), anti-human C3c (Abcam) or anti-human CRP (Abcam) in the concentration recommended by manufacturer were used for the detection of IgG and the other components of complexed native IgG, respectively.

In order to analyse the exposure of specific glycans of the glycoprotein IgG in the sera, an IgG capture lectin ELISA was performed (Figure 1(a)). After coating 96 well plates with the anti-human IgG F(ab')2-fragment described above, plates were blocked with deglycosylated blocking buffer (3% gelatine, 0.1% CaCl₂, 0.1% MgCl₂, 0.05% tween-20) at 37°C for 2 hours. This buffer was pre-treated for 24 hours with 1% periodic acid (Merck) to destroy glycosylation sites of the gelatine. The pH was restored by dialysis against tris buffered

### Table 1  Clinical characteristics of patients with SLE

<table>
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<tr>
<th>Patient #</th>
<th>Age/sex/ethnicity</th>
<th>Recent onset disease (yes/no)</th>
<th>Manifestation at flare</th>
<th>Fulfilled 1982 ACR criteria</th>
<th>Daily dosage of Prednisolone (at flare)</th>
<th>Other treatment at blood sampling</th>
<th>At flare</th>
<th>Anytime during the study</th>
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<td>Rtx</td>
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<td>Am</td>
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<td>Aza, Am</td>
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<td>Mmf</td>
<td>Rtx, Mmf</td>
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<td>15</td>
<td>Rtx</td>
<td>Aza, Am</td>
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<td>Mmf</td>
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<td>Mmf, Am</td>
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<td>Mmf, Am</td>
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<tr>
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<td>49/F/Caucasian</td>
<td>yes</td>
<td>Vasculitis, oral ulcers, headache</td>
<td>3, 4, 10, 11</td>
<td>30</td>
<td>Aza</td>
<td>Am</td>
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Am: antimalarials; Aza: azathioprine; Bmab: belimumab; Cyclo: cyclophosphamide; Mmf: mycophenolate mofetil; Mtx: methotrexate; Rtx: rituximab; Siro: sirolimus.
In order to detect carbohydrate variations of complexed IgG from serum of patients with SLE we performed Lectin-ELISAs. The plates were coated with goat anti-human IgG F(ab')2-fragment, treated with deglycosylated blocking buffer to avoid loading unspecific glycans onto the plates. After incubation with serum dilutions and several washing steps biotinylated lectins were employed to detect fucosyl residues and fucosylated tri-mannose N-glycan core sites of the complexed native IgG molecules (a). Ellipsometry was used to measure layer thickness of the proteins bound to the anti-human IgG F(ab')2-fragment. The hydrophobic surface constitutes the background. At least 1 nm of F(ab')2-fragments bound to the surface (mean value of 10 measurements shown). IgG from sera of five NHD and 11 SLE patients was immobilized to a similar extent of 5–7 nm; the data represent mean values of 10 separate measurements of each individual serum sample (b).
saline containing 0.1% CaCl$_2$, 0.1% MgCl$_2$ (TBS-Ca-Mg) until a pH-value of 7.4 was reached. After blocking, patient sera or sera from normal healthy donors were diluted 1:1000 in TBS-Ca-Mg and 0.05% tween-20 and incubated at 37°C for 2 hours. These serum dilutions resulted in a saturating binding of IgG to the ELISA plate. After every incubation step the plates were washed three times with 200 ml TBS-Ca-Mg-tween. Fifty ng/ml biotinlabelled AAL or LCA (Vector Laboratories) were used to detect glycosyl residues exposed by the captured IgG complexes and incubated at room temperature for 1 hour. After washing, plates were incubated with HRP-streptavidin (Jackson Laboratories Immunoresearch) at the recommended concentration for 1 hour at room temperature.

Detection was performed for both types of ELISA with the addition of substrate solution (0.1 M Na$_2$HPO$_4$, 0.05 M citrate acid monohydrate, 0.02% H$_2$O$_2$ and 100 µg/ml Tetra-methyl benzidine, pH=5, Merck) for 10 minutes and the colour reaction was stopped with 25% sulphuric acid (Merck). Optical density values were obtained with the ELISA-reader (Tecan infinite F200 Pro) employing a 450 nm/620 nm filter/reference pair.

**Ellipsometry**

Ellipsometry is a highly sensitive optical method that has been used since the 1960s to determine the thickness of an adsorbed bioorganic layer with a practical resolution of about 0.1 nm. Thus, we used ellipsometry to confirm the adsorption of F(ab’)$_2$-fragments and serum IgG onto surfaces of methylated (hydrophobic) silicon as previously described. Measurements were performed on: (1) bare methylated surface; (2) with F(ab’)$_2$-fragments adsorbed; (3) with sera adsorbed to F(ab’)$_2$-fragments.

**Ethics**

Oral and written informed consent was obtained from all subjects. The study protocol was approved by the Linköping University Ethical Review Board (M75-08/2008).

**Results**

Complexed IgG from patients with SLE expose fucosyl residues

Employing ellipsometry, we confirmed the adsorption of F(ab’)$_2$ of more than 1 nm onto the hydrophobic surface and this layer immobilized similar levels of serum proteins from patients and controls (Figure 1(b)). Once the immobilization of random serum IgG complexes was confirmed, their exposure of AAL and LCA binding glycoepitopes in consecutive serum samples of 15 patients with SLE and in sera of 15 NHD were measured employing a lectin-ELISA. We aimed to create a profile of the surface structure of IgG from patients with SLE using detection antibodies against IgM, IgA, C1q, C3c and CRP. The analysis confirmed equal and saturating IgG-coating by sera from patients with SLE and NHD. Table 2 shows that secondary antibodies against IgM, IgA, complement component C1q and CRP showed no significant difference between patients with SLE and the controls. However, we observed a significantly higher reactivity of the complexed IgG with the anti-C3c antibody ($p=0.000044$) from the sera of patients with SLE.

Figure 2 shows that the exposure of glycosyl residues was independent of the age of the patients (analysed at the first visit) or the controls. The exposure of glycan sites by complexed random IgG in the sera of patients with SLE varied during the course of the disease and is represented in Figure 3 as black circles. In general, the complexed IgG showed a higher binding of AAL (Figure 3(a), $p < 0.00001$) and LCA (Figure 3(b), $p < 0.05$) than NHD samples.

**Exposure of fucosyl residues on IgG is associated with high SLE disease activity and low serum C3 levels**

We also noticed that disease activity in SLE, demonstrated by PGA scores at each visit, was associated with higher levels of AAL reactivity (Figure 4(a)). It was observed that during times of increased disease activity (PGA 1–3) patients with SLE display significantly increased AAL reactivity than NHD ($p=0.00000844$). Interestingly, SLE patients clinically judged as non-active (PGA = 0,
i.e. remission) still showed higher AAL binding to complexed serum IgG than NHD ($p = 0.0038$). When comparing patients in remission (PGA = 0) with mild (PGA = 1), moderate (PGA = 2) and high (PGA = 3) disease activity, we found a significant increase of AAL-reactivity ($p = 0.0007$, 0.0011 and 0.016, respectively). The association of raised disease activity and higher degree of fucosyl accessibility of native complexed IgG was also demonstrated applying the SLEDAI (Figure 4(b)). Patients with inactive SLE (SLEDAI = 0) exhibit significantly higher AAL reactivity in comparison with NHD ($p = 0.0011$). Patients with SLE and SLEDAI scores higher than 0 show significantly higher levels of AAL binding than NHD ($p = 0.0000007$) and patients with SLE and a SLEDAI score of 0 ($p = 0.0002$).

We calculate the positive and negative predictive values (PPV and NPV respectively) of fucosyl residues exposed by random IgG and disease activity measured by SLEDAI and PGA, respectively. Samples with SLEDAI higher than 5 and PGA scores higher than 0 were considered as having increased disease activity. Positive values for AAL reactivity were considered as higher than 90% of the NHD. Considering SLEDAI as gold standard, the PPV was 70.3%, the NPV 69.8%, sensitivity 61.9% and specificity 77.1% for the detection of fucosyl residues in IgG IC by AAL. Similarly, for PGA-score we calculated PPV 89.2%, NPV 54.7%, sensitivity 57.9% and specificity 87.9%.

Correlation analysis revealed a strong negative association of the exposure of fucosyl residues with the levels of circulating complement component C3 ($p < 0.0001$; Figure 4(c)). Other laboratory parameters (C4, erythrocyte sedimentation rate, haemoglobin, leukocyte and thrombocyte counts) did not show relevant associations with the exposure of fucosyl residues on the immobilized IC.

**Increased disease activity associates with increased exposure of AAL-binding sites in individual patients**

We observed an association between the highest AAL reactivity of complexed random IgG of a patient and increased disease activity for the majority of the patients studied. In our follow-up cohort of 15 patients with SLE, 11 (73%) showed maximum levels of AAL-binding of complexed IgG during increased disease activity (Figure 5(a)). Using the PGA to quantify disease activity we noticed a similar outcome in seven out of 15 (47%) patients. Interestingly, it turned out that five patients with SLE showed no difference in PGA score while they reached the maximum and minimum levels of AAL-binding (Figure 5(b)). In order to closely observe the association between exposure of fucosyl residues by native IgG complexes and disease activity we illustrated the courses of AAL-reactivity and SLEDAI in the same plot (Figure 6). We included in this figure all SLE patients with at least five available longitudinal serum samples and a peak of SLEDAI between the first and the last determined SLEDAI scores (patient numbers 1–7, 9, 10, 12, 13). Eight from these 11 patients (numbers 2, 3, 4, 6, 9, 10, 12, 13) showed similarities between SLEDAI and

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**Figure 2** Binding of AAL and LCA to immobilized native IgG.

Lectin ELISA in sera from patients with SLE (close symbols) and normal healthy donors (open symbols). Binding of AAL and LCA to immobilized complexed IgG is shown in (a) and (b), respectively. Using all serum samples of the first visit of each patient no association among lectin binding, age and sex (female circles, male triangles) was to be observed.
AAL-reactivity, although the absolute levels were rather different between the individual patients.

Discussion

SLE is a disease of multifactorial genesis, in which immune complex formation by autoantibodies, complement activation and cytokine production lead to persistent inflammation and subsequent tissue damage in various organs. Enzymatic removal of the glycan structure of complexed Ig with endoglycosidase S has been shown to diminish various pro-inflammatory features of these CICs in autoimmune conditions. Activation of the complement system by classical pathway, induction of interferon-γ production and/or modulation of immune responses by binding to Fcγ receptors are the putative pathogenic mechanisms. This suggests the assumption that the molecular structures of CICs, which include their glycans strongly impact their functionality and thus their inflammation-driving potential. In rheumatoid arthritis (RA), a further chronic inflammatory disease, several studies have revealed that
Galactosylation of serum IgG correlates with disease activity and can be influenced by various types of therapy. The fucosylation of denatured IgG heavy chains was significantly increased in RA as measured by lectin binding. Furthermore, there is a positive correlation between galactosylation and sialylation of IgG with the improvement of RA during pregnancies and post-partum flares. Glycans are potential ligands for a plethora of lectin receptors found on immune effector cells, which are used as phagocytic receptors. Data about IgG glycosylation status and glycan exposure of native IgG in SLE during disease activity and remission are still scarce. In this study, we analysed serum samples of 15 well-characterized Swedish SLE patients. The samples were drawn at
consecutive visits during active and non-active periods of disease. The grade of disease activity was quantified for every sample with PGA and SLEDAI. The serum samples of 15 sex- and age-matched healthy blood donors served as controls.

In Lectin-ELISAs the binding of AAL and LCA to immobilized native serum IgG was quantified to detect exposure of fucosyl residues and of the fucosylated tri-mannose N-glycan core, respectively. For Lectin-ELISAs it was necessary to prepare a

Figure 5  Association between maximum and minimum AAL reactivity and clinical activity.
(a) Immobilized complexed IgG of patients with SLE display increased AAL binding during times of increased disease activity (SLEDAI). Four patients exhibit a higher SLEDAI score during lowest detectable AAL binding of serum IgG. Eleven patients (73%) present a higher SLEDAI score at the time point of the highest AAL binding. (b) PGA also demonstrates that IgG complexes of patients with SLE display higher levels of AAL binding during increased disease activity. For five patients with SLE there is no difference in PGA score for the time-points of highest and lowest detectable AAL reactivity. Three patients show a higher PGA score during time of lowest AAL binding of random serum IgG than during highest accessibility of fucosyl residues. Seven patients with SLE exhibit a higher PGA score at the time of the highest exposure of fucosylated sites of native IgG complexes.
special deglycosylated blocking buffer (treated with 1% periodic acid) to avoid loading unspecific glycan structures onto the polystyrene plates during the process of blocking.

Since several immune cells employ glycan-lectin recognition patterns, we aimed not to analyse the glycan of isolated and denatured IgG in the peripheral blood, but the glycans exposed by circulating native IgG complexes. We argue that a complex can be phagocytosed in a lectin dependent manner if any of the components exposes a corresponding glyco-epitope. This may even apply to the bound antigen. In contrast, internal glycans are going to be ignored when they are inaccessible due to steric hindrance. However, the latter are usually detected with techniques that require the denaturation of the target, like lectin blot or mass spectrometry.

Thus, we determined the level of immobilized native IgG, representing loading control, as well as the level of the IgG-associated candidate molecules IgA, IgM, C3c, C1q and CRP. This selection is by far not complete, but represents a couple of molecules that are reportedly found in circulating IgG complexes. The level of IgG-bound complement proteins correlates with its consumption. In these types of ELISA, we analysed native serum IgG complexes under preservation of their molecular integrity and thus tried to gain new information about the in vivo function of various glycosyl residues exposed by complexed, circulating immunoglobulins.

The analysis revealed that only the C3c epitope was significantly increased in patients with SLE. Since C3c is a stable cleavage product of C3 covalently bound to target structures, the detection of C3c gives a line on C3 metabolism. It is well-known that SLE is associated with an elevation of C3 turnover even during times of remission as a feature of chronic inflammation, whereby an upcoming exacerbation can be identified by a decline of C3, reflecting massive C3 consumption. The exposure of fucosyl residues on circulating IgG IC correlated highly significant with the consumption of serum complement C3. However, there was no association between C3c detected on IgG IC and AAL reactivity (data not shown). This suggests that the consumption of C3 may rather be associated to the deposition of IC in tissues and therefore the measured AAL reactivity may reflect indirectly an ongoing immune complex deposition.

Employing the lectins as detection tool, we observed a higher exposure of IgG fucosyl residues both in sera of patients with SLE in general, and

Figure 6: Exposure of fucosyl residues by random IgG during the course of disease. Association between Systemic Lupus Erythematosus Disease Activity Index-2K (SLEDAI-2K) (broken line) and AAL reactivity during the course of disease (solid line). In this figure the course of disease activity (raised SLEDAI-2K) and the course of AAL reactive serum IgG is shown in order to detect parallels. All patients with at least five consecutive serum samples and a SLEDAI-2K peak between the first and the last collected SLEDAI-2K score are included (patient numbers 1–7, 9–10, 12–13).
even more at time-points of increased disease activity. Figure 6 shows the marked parallels of increased accessibility of fucosylated sites of random IgG complexes and disease activity of some patients, giving a hint that there is an association between these two parameters. It has been shown in several studies that patients with RA display IgG with low galactosylation after flares, when compared to those in remission. The latter show levels of galactosylation similar to NHD. Fucosylation of IgG, conversely, turned out to be significantly increased throughout the whole course of RA. Under- and non-fucosylation of serum IgG is known to elevate the binding affinity to the FcγRIIIa and thus to enhance its proinflammatory activity. Former studies revealed that the binding affinity of defucosylated IgG to the FcγRIIIa might be 10-fold higher than that of immunoglobulins without decreased fucosylation. However, these results were obtained by employing genetic modification of host cells and denaturing conditions like treatment at 85°C to 100°C, mass spectrometry or electrophoretic separation of IgG heavy and light chains. Consequently, these techniques are prone to additionally detect glycan structures hidden in the native molecules. Applying these methods functional characteristics of the native IgG complexes are lost.

Our investigation revealed that patients with SLE exhibit a significantly higher exposure of fucosyl residues of complexed serum IgG both in times of decreased and even more during times of increased disease activity.

There are several possibilities that could explain the increased binding of AAL to circulating complexed IgG: (1) increased fucosylation; (2) increased exposure of fucosyl residues on random IgG; (3) a fucosylated glycan bound to the variable region of certain IgG molecules; (4) fucosyl residues exposed by the IgG associated molecules. Employing lectin-ELISA, we cannot distinguish which one of these mechanisms, if any, is responsible for our observations. Nevertheless, recognition of IC by immune cells may not distinguish the exact localization of the interacting fucosyl residues but just the presence of a glyco-epitope. Complexed IgG molecules exposing highly fucosylated sites may favour their uptake by binding to phagocyte surface lectin receptors and thus drive chronic inflammatory diseases like SLE and possibly others. After evaluating the performance of the test predicting periods of high disease activity we can consider the analysis of fucosylated sites exposed by circulating complexed IgG as an appropriate laboratory parameter to monitor patients with chronic inflammatory rheumatic diseases.

Besides the markedly elevated level of exposed fucosyl residues, we also detected a significant increase of LCA binding, suggesting more fucosylated tri-mannose N-glycan core sites in the complexed IgG of patients with SLE. It had been reported that highly mannosylated IgG strongly activate FcγRIIIa receptors and ADCC, fostering pro-inflammatory immune responses. But since those over-mannosylated IgGs also show reduced core fucose, it is not known which of these two features of glycosylation was responsible for the effect. Furthermore, highly mannosylated immunoglobulins exhibit increased affinity to the acute phase protein mannose-binding lectin (MBL), which usually binds to carbohydrate structures on the surface of pathogens and consecutively activates the complement system (lectin pathway). As a result, highly mannosylated immunoglobulins and microparticles display an increased rate of clearance. MBL comes along with higher susceptibility for SLE and it is presumed that MBL might foster the clearance of CIC, suspected to play an important role for the development of tissue damage in SLE. A possible explanation for the increased exposure of fucosylated tri-mannose N-glycan core sites by native IgG is that IgG with higher accessibility to mannosylated sites is synthesized in SLE to act protectively by raising the affinity between CICs and MBL and thus fostering accelerated clearance of potentially tissue damaging IC.

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Conflict of interest statement

The authors have no conflicts of interest to declare.
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