Anti-tissue transglutaminase antibodies from coeliac patients inhibit transglutaminase activity both in vitro and in situ

C Esposito, F Paparo, I Caputo, M Rossi, M Maglio, D Sblattero, T Not, R Porta, S Auricchio, R Marzari, R Troncone

**Background and aims:** Coeliac disease (CD) is a multifactorial disorder which has an autoimmune component characterised by the occurrence of disease specific autoreactive antibodies against the enzyme tissue transglutaminase (tTG). The aim of this study was to investigate whether binding of antibodies to the enzyme influences tTG activity.

**Methods:** tTG activity was assayed in the presence of immunoglobulin A (IgA) and immunoglobulin G (IgG) purified from the serum of coeliac patients, CUB 7402 (an anti-tTG mouse monoclonal antibody), and human anti-tTG monoclonal antibodies derived from both intestinal lymphocytes from three patients with CD and from peripheral blood lymphocytes from healthy subjects. For our studies we used calcium treated and untreated recombinant human tTG. Furthermore, the effects of antibodies were determined by immunohistochemical detection of tTG activity in sections of human umbilical cord.

**Results:** IgG and IgA from CD patients inhibited tTG activity in vitro in a dose dependent manner, with a different rate of inhibition among patients. The monoclonal antibody CUB 7402 and human monoclonal antibodies displayed a dose dependent inhibitory effect towards the catalytic activity of the enzyme, both in vitro and in situ. Preincubation of tTG with CaCl2 caused loss of the inhibitory effect due to CUB 7402 but not that caused by human monoclonal antibodies.

**Conclusions:** Purified CD IgA, IgG, as well as human anti-tTG monoclonal antibodies inhibited the enzymatic activity of human tTG both in vitro and in situ.

**T**issue transglutaminase (tTG, EC 2.3.2.13) belongs to a family of Ca2+ dependent enzymes that catalyse post-translational modification of peptides and proteins with the formation of isopeptide bonds between the γ-carboxamide group of protein bound glutaminyl residues and ε amino group of lysyl residues, present in the same or different polypeptide chain, or primary amines. tTG is widely distributed in vertebrate tissue and is a strongly regulated enzyme both transcriptionally and functionally. It appears to play a critical role in controlling cell and tissue homeostasis, regulating the cell cycle through its involvement in proliferation, terminal differentiation, and apoptosis. It is also present in extracellular compartments where it has a role in the stabilisation of the extracellular matrix and tissue repair by forming cross links between various substrate proteins. Finally, tTG is involved in the transduction of extracellular signals. In fact, tTG is a bifunctional enzyme with a transamidating catalytic activity on peptidyl glutamine residues as well as a guanosine triphosphate (GTP) hydrolysing activity, analogous to that of G proteins, in adrenergic receptor transduction pathways.

Other than its physiological roles, tTG mediated post-translational modifications of proteins may represent a pathogenic mechanism in human diseases characterised by autoimmune phenomena, including coeliac disease (CD). It is considered to be the result of a dysregulated T cell mucosal immune response to wheat gliadin and related prolamines of other toxic cereals (barley, rye, and possibly oats). It affects genetically susceptible individuals, with human leucocyte antigen genes playing a major role. Autoimmunity is an increasingly recognised feature of the disease; in fact, CD not only appears to be strongly associated with a series of autoimmune conditions, such as insulin dependent diabetes mellitus, thyroiditis, and Addison’s disease, but is itself characterised by autoimmune phenomena. It has long been known that disease specific antibodies recognise proteins of the extracellular matrix and the target was recently identified by Dieterich et al as being tTG. This observation has had a large impact on diagnostic strategies for CD as well as providing new perspectives in the understanding of the disease mechanisms at both the local and systemic levels, reflecting the role of tTG in many crucial biological processes.

Recently, Marzari and colleagues isolated a series of antibodies to tTG by selecting phage display antibody libraries derived from either intestinal lymphocytes or peripheral blood lymphocytes from three patients with CD. They showed that whereas anti-gliadin responses could be selected from all libraries, the anti-tTG response was restricted to intestinal lymphocytes, involving the recognition of two main tTG epitopes. Here, we have investigated the effect of these antibodies, as well as that of immunoglobulin (Ig) purified from coeliac serum, on tTG catalytic activity. We report data showing that the interaction between anti-tTG antibodies and tTG inhibits the transamidating activity of the enzyme both in vitro and in situ. These results are discussed in relation to the still obscure role played by these autoantibodies in the pathogenesis of CD.

**Abbreviations:** CD, coeliac disease; tTG, tissue transglutaminase; Ig, immunoglobulin; GTP, guanosine triphosphate; MDCK, Madin-Darby canine kidney; EDTA, ethylenediamine-tetraacetic acid; PBS, phosphate buffered saline; BSA, bovine serum albumin; RT, room temperature; EGTA, ethylene glycol bis(β-aminoethyl ether) N,N,N′,N′-tetraacetic acid; DMC, N,N-dimethylated casein; bio-MDC, biotinylated monodansyl-cadaverine; TGF-β, transforming growth factor β.
MATERIALS AND METHODS
Cell culture and DNA transfection
Madin-Darby canine kidney (MDCK) cells (European Cell Collection, 85011435) were grown in minimum essential medium (Life Technologies, Milan, Italy) supplemented with 10% fetal bovine serum, 10%Earl’s balanced salt solution, 50 U/ml penicillin, 50 µg/ml streptomycin, and 1% non-essential amino acids. Cells were maintained in a 5% CO₂, humidified atmosphere. Human tTG CDNA,22 cloned in the eukaryotic expression vector pSG5 (Stratagene, La Jolla, California, USA), was used to transfect MDCK cells by calcium phosphate precipitation. A stable cell clone, MDCK-tTG, expressing recombinant tTG was obtained by cotransfecting MDCK cells with pSV2-Neo (Clontech, Palo Alto, California, USA) in a 1:10 ratio. Cotransfected cells were cultured in a selective medium containing 400 µg/ml G418 (Life Technologies).

MDCK-tTG cells were harvested in 10 mM Tris/HCl pH 7.5, 1 mM ethylenediamine-tetraacetic acid (EDTA), and sonicated for 10 seconds. Protein content was estimated by the method of Bradford with bovine serum albumin (BSA) (Sigma, St Louis, Missouri, USA) as the standard.22

IgG and IgA purification
IgG and IgA antibodies from normal and CD patients were purified using Sepharose beads conjugated with protein A or rabbit antibodies to human IgA (Sigma). Serum was buffered with 0.1 M Tris/HCl pH 8.0 and applied to the anti-human IgA Sepharose equilibrated in the same buffer. The flow through fraction, containing IgG, was collected and subsequently applied to protein A Sepharose. Bound antibodies were eluted with 0.1 M glycine buffer pH 3.0; fractions were collected into tubes containing 1 M Tris/HCl pH 8.0 to prevent antibody denaturation. Protein content was determined using the method of Bradford.22

Monoclonal antibodies
Monoclonal antibodies (single chain antibody fragments) to tTG were prepared as described previously.22 Briefly, total RNA was prepared from intestinal biopsies from three previously untreated CD adult patients, cDNA was synthesised using random hexamers, and Ig V regions were amplified using specific V region primers and assembled before cloning into the phagemid vector pDAN5.22 Antibodies to tTG were isolated by recursive cycles of binding to recombinant human tTG, washing, and elution of phage particles expressing antibody fragments on their surface. In the present work, clones 2.18, 3.7, and 4.2 were isolated from CD libraries whereas clone D51 was selected from a naïve library22 made from peripheral lymphocytes from healthy donors. The CD antibodies are reported according to the donor patient, and given reference numbers 2, 3, and 4 followed by the individual clone reference number.

Soluble antibodies were obtained by expression in HB2151 (K12, ara Δ(lac-pro), thi/F’ proA’ B’, lacPZΔM15, a non-suppressor Escherichia coli strain). Phages from individual colonies were infected into HB2151, grown to OD₆₀₀ 0.2, induced with 1 mM isopropyl-β-D-thiogalactopyranoside, and further grown overnight at 28°C. Antibodies were purified from supernatants of induced bacterial cultures by Ni-NTA chromatography (Qiagen, Germany) and dialysed against phosphate buffered saline (PBS) to obtain a concentration of approximately 0.2 mg/ml.

Enzyme linked immunosorbent assay (ELISA) for anti-tTG antibodies
To measure serum IgA and IgG anti-tTG titres, we used an ELISA method. U bottomed microtitre plates (Falcon, Franklin Lakes, New Jersey, USA) were coated with 1 µg of human recombinant tTG per well in 100 µl of coating buffer (50 mM Tris/HCl, 150 mM NaCl, 5 mM CaCl₂, pH 7.5) for two hours at 37°C. Wells were washed three times with washing buffer (50 mM Tris/HCl, 150 mM NaCl, 10 mM EDTA, 0.1% Tween-20, pH 7.4). The plates were incubated in washing buffer overnight at 4°C. Sera, diluted 1:100 in the same buffer, were added to the wells and incubated for one hour at room temperature (RT). After three washes, the wells were incubated with 100 µl of peroxidase conjugated rabbit anti-human IgA or peroxidase conjugated rabbit anti-human IgG (Dako, Denmark) diluted respectively 1:1000 and 1:3000 in the same buffer for one hour at RT. Unbound antibodies were removed with three washes and colour was developed by addition of 100 µl of 1 M sodium citrate, 1 M citric acid, 1 mg/ml α-phenylenediamine-hydrochloride (Sigma), and 0.06% H₂O₂ (Carlo Erba, Italy), pH 5, at RT for 30 minutes in the dark. Absorbances were read on an ELISA reader (SpectraCount, Packard, Milan, Italy) at 450 nm. All samples were tested in duplicate. Values were expressed as percentage of a reference pool of sera obtained from untreated coeliac patients.

In vitro tTG assay
Enzyme activity was assayed by a radiometric method based on Ca²⁺ dependent incorporation of [³H]spermidine (Nen Life Science Products, Boston, Massachusetts, USA) into N,N,N’-dimethylated casein (DMC). Briefly, the assay mixtures (100 µl) containing 125 mM Tris/HCl buffer, pH 8.0, 10 mM dithiothreitol, 5 mM CaCl₂, 0.16 µM [³H]spermidine (15 Ci/mmol), 0.2 mg DMC and, as enzymatic source MDCK-tTG cell homogenate (7 µg of proteins), were incubated at 37°C for one hour. The inhibition studies were performed by preincubation of the enzyme with different amounts of IgG and IgA in 25 mM Tris/HCl pH 8.0 (final volume 20 µl) at 20°C for 10 minutes. Preincubation of tTG with different amounts of monoclonal antibodies was performed in the absence or presence of 5 mM CaCl₂. Blanks were simultaneously run with 5 mM ethylene glycol-bis(β-aminoethyl ether) N,N,N’,N” -tetraacetic acid (EGTA). The reactions were stopped with 10% trichloroacetic acid containing 2 mM unlabelled spermidine. After centrifugation, the precipitates were washed twice with the same solution, dissolved in 0.5 ml of 0.2 N NaOH, and finally counted in 5 ml of Pico-Fluor 40 scintillation mixture (Packard).

In situ tTG activity assay
Umbilical cord cryostat sections (7 µm thick) were mounted on slides with 2% triethoxysilyl-propylamin (Merck, Germany) in acetone, air dried for 60 minutes at RT, and fixed in acetone for 10 minutes. Sections were preincubated with 1% BSA in 100 mM Tris/HCl, pH 7.4, for 30 minutes at RT, and then for two hours at RT with biotinylated monodansylcadaverine (bio-MDC) (Molecular Probes, Leiden, the Netherlands). This substrate solution was prepared by adding 10 µl of 10 mM bio-MDC and 25 µl of 200 mM CaCl₂ solution to 965 µl of 100 mM Tris/HCl, pH 7.4. The tTG reaction was stopped with 25 mM EDTA for five minutes. Incorporation of labelled substrate was visualised by incubation in a humid chamber for 30 minutes at RT with streptavidin fluorescein isothiocyanate conjugate (Dako) diluted 1:50 in PBS. Finally, sections were washed twice in PBS and mounted in 10% glycerol in PBS. In control sections, 200 mM CaCl₂ was replaced by 200 mM EDTA. Photographs were taken using an Axioskop (Zeiss, Germany) and colour reversal film (Kodak, UK).

RESULTS
Effects of CD patient IgG and IgA on tTG activity in vitro
Table 1 shows that addition of 10 µg of IgG and IgA from six and five CD patients, respectively, into the assay mixture inhibited the incorporation of [³H]spermidine into DMC catalysed by recombinant tTG expressed in the MDCK-tTG cell line. A different rate of inhibition among patients was also observed. In fact, the extent of inhibition due to IgG and IgA...
The monoclonal antibody CUB 7402 displayed a dose dependent inhibitory effect towards the catalytic activity of the enzyme expressed in MDCK-ITG cells with respect to basal values, as shown in fig 2. Because calcium ions influence iTG activity by inducing structural changes in the protein, we preincubated the enzyme, before the enzymatic assay, with different amounts of antibodies in 25 mM Tris/HC1 pH 8.0 in the presence of 5 mM CaCl2. Preincubation caused loss of the inhibitory effect due to CUB 7402.

The same experiments were performed using monoclonal antibodies to iTG derived from both intestinal lymphocytes from three patients with CD and from peripheral blood lymphocytes from healthy subjects. The results reported in fig 3 show that the dose related inhibition caused by 4.2, 2.18, and 3.7 clones was almost the same (ranging from 20% to 40%). However, when preincubation was performed in the presence of CaCl2 inhibition was still present; in particular, when 2.18 clone was used, enhanced inhibition was observed.

Finally, only a weak inhibitory effect was evoked by the anti-ITG monoclonal antibody derived from peripheral blood lymphocytes from healthy subjects, suggesting that the anti-ITG antibodies found in CD have special properties which may play a role in the disease.

### Table 1: Effects of IgG and IgA purified from the serum of patients with coeliac disease and controls on tissue transglutaminase (ITG) activity

<table>
<thead>
<tr>
<th>Antibody</th>
<th>Patients</th>
<th>% inhibition*</th>
<th>Titres†</th>
</tr>
</thead>
<tbody>
<tr>
<td>IgG</td>
<td>Control</td>
<td>0 (0.5)</td>
<td>2</td>
</tr>
<tr>
<td>IgG</td>
<td>2</td>
<td>13.3 (3.7)</td>
<td>39</td>
</tr>
<tr>
<td>IgG</td>
<td>3</td>
<td>16.0 (2.3)</td>
<td>37</td>
</tr>
<tr>
<td>IgG</td>
<td>4</td>
<td>8.3 (4.3)</td>
<td>89</td>
</tr>
<tr>
<td>IgG</td>
<td>5</td>
<td>24.0 (1.0)</td>
<td>37</td>
</tr>
<tr>
<td>IgG</td>
<td>6</td>
<td>14.0 (3.1)</td>
<td>54</td>
</tr>
<tr>
<td>IgG</td>
<td>7</td>
<td>11.7 (2.0)</td>
<td>90</td>
</tr>
<tr>
<td>IgA</td>
<td>Control</td>
<td>0 (0.5)</td>
<td>2</td>
</tr>
<tr>
<td>IgA</td>
<td>2</td>
<td>39.0 (1.0)</td>
<td>92</td>
</tr>
<tr>
<td>IgA</td>
<td>3</td>
<td>22.0 (2.0)</td>
<td>46</td>
</tr>
<tr>
<td>IgA</td>
<td>4</td>
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<td>53</td>
</tr>
<tr>
<td>IgA</td>
<td>5</td>
<td>19.5 (5.0)</td>
<td>30</td>
</tr>
<tr>
<td>IgA</td>
<td>6</td>
<td>11.1 (1.5)</td>
<td>29</td>
</tr>
</tbody>
</table>

*Values represent the mean of two replicates in two independent sets of experiments; 1 µg of purified IgG or IgA was used in each in vitro assay tube.

†Titres were measured by ELISA and expressed as percentage of a reference pool of sera.

### Figure 1: Effect of purified serum IgG from coeliac patients (Nos 3 and 4) and from healthy controls on tissue transglutaminase (ITG) activity

- Patient No 4
- Patient No 3
- Controls

### Figure 2: Effect of the monoclonal antibody anti-tissue transglutaminase (ITG) (CUB 7402) on ITG activity in response to Ca2+. Increasing concentrations of CUB 7402 were preincubated at 20°C for 10 minutes in 20 µl of 25 mM Tris/Cl, pH 8.0, in the presence or absence of 5 mM CaCl2, with 7 µg of proteins from Madin-Darby canine kidney (MDCK)-ITG cells. The samples were further incubated for one hour at 37°C in a 100 µl final volume. The activity of ITG was calculated as a percentage of basal activity (no CUB 7402 added) within MDCK-ITG cells. Data are presented as means (SD) from three separate experiments performed in duplicate.

### Figure 3: Effect of single chain monoclonal antibodies on tissue transglutaminase (ITG) activity in response to Ca2+. Human monoclonal antibodies (0.5, 1.0, and 1.5 µg) were preincubated at 20°C for 10 minutes in 20 µl of 25 mM Tris/Cl, pH 8.0, in the absence or presence of 5 mM CaCl2, with 7 µg of proteins from Madin-Darby canine kidney (MDCK)-ITG cells. The samples were further incubated for one hour at 37°C in a 100 µl final volume. The activity of ITG was calculated as a percentage of basal activity (no antibodies added) within MDCK-ITG cells. Data are presented as means (SD) from three separate experiments performed in duplicate.


**Figure 4**  Effect of specific antibodies on tissue transglutaminase (tTG) activity in human umbilical cord. The signal corresponds to incorporation of substrate biotinylated monodansyl-cadaverine (A). Sections treated with the anti-tTG CUB 7402, diluted 1:25 (B), and with monoclonal antibody 4.2, undiluted (C), showed a significantly lower signal (original magnification x400).

**Effect of antibodies on tTG activity in situ**

This effect was detected in umbilical cord sections by incorporation of the labelled substrate bio-MDC, resulting in the presence of fluorescent signals (fig 4A). Such activity was clearly affected by addition of antibodies. Anti-tTG CUB 7402 (fig 4B) and single chain monoclonal antibodies from the intestinal mucosa of coeliac patients (fig 4C) significantly decreased signalling. Both IgG and IgA from coeliac serum, but not from controls, significantly decreased the signalling (data not shown).

**DISCUSSION**

The recent identification of tTG as the main endomysial autoantigen recognised by IgA from patients with CD allowed the implementation of ELISA based tests, initially based on guinea pig tTG and subsequently on human tTG, in the clinical diagnosis of CD. Interestingly, IgA from coeliac sera preferentially bind Ca\(^{2+}\) activated tTG, suggesting the recognition of conformational epitopes in the enzyme. The mechanism of formation of these antibodies is not presently understood. It has been proposed that gliadin specific T cells provide cognate help for their production by tTG specific B cells to explain the strict gluten dependence. Similarly, the possible role of these autoantibodies in the pathogenesis of mucosal damage is uncertain. Although most evidence suggests a gliadin triggered Th1 mechanism of damage, it is very important to address the question of their possible contribution to the intestinal lesion, given the strong disease specific presence of tTG antibodies and the role played by this enzyme in several crucial biological processes. So far only indirect evidence has been provided: it has recently been shown that transforming growth factor β (TGF-β) dependent differentiation of T84 crypt epithelial cells in a fibroblast coculture model can be prevented by the addition of anti-tTG IgA. These authors suggested that blocking tTG prevented the generation of the active form of TGF-β which relies on tTG for its maturation, and they further speculated that anti-tTG antibodies may also affect the differentiation of the coeliac epithelium. In this paper, we have directly addressed the issue of whether anti-tTG antibodies can modulate the activity of the enzyme in any way.

Our results demonstrate for the first time that IgA and IgG from serum of CD patients as well as monoclonal anti-tTG antibodies, especially those obtained from CD patients, display inhibitory effects towards the catalytic activity of human tTG both in vitro and in situ. Inhibition of recombinant human tTG activity was shown in all sera tested although the degree of inhibition was different from case to case. Moreover, the extent of such inhibition did not correlate with the titre of antibodies measured by ELISA. Take together, the data indicate that more than one epitope is recognised by sera from different patients. Also interesting were the data obtained with monoclonal antibodies produced by B cells present in the coeliac lesion. Antibodies obtained from a phage antibody library produced both from peripheral blood and from CD patients recognised different tTG epitopes. These antibodies were isolated from three CD patients and shown to recognise two main conformational tTG epitopes. In the present study, four monoclonal antibodies were chosen for the inhibition experiments: clones 2.18 and 4.2 belonging to the two categories as above, 3.7 with uncertain attribution, and D51 derived from healthy donor peripheral lymphocytes. Interestingly, all of the monoclonal antibodies from CD patients showed inhibitory activity whereas D51, obtained from non-coeliac subjects, did not show any inhibitory activity, implying that tTG antibodies from healthy subjects may not bind to epitopes related to substrate combining sites of tTG.

TGF-β is subject to fine regulation, involving calcium and GTP which, respectively, activate and inhibit transamidating activity by inducing conformational changes. Recently, using experimental approaches and computer modelling, a three dimensional model was proposed in which the protein core domain, containing the catalytic site, becomes accessible to macromolecular substrates as a consequence of calcium binding. The catalytic domain extends from amino acid 140 to 434, also containing the three putative calcium binding regions comprising residues 146–162, 229–234, and 434–452, while the GTP binding site was located in the region 1–180. We have tested the consequences of the presence of calcium ions during preincubation of recombinant human tTG with monoclonal antibodies from patients. Calcium did not interfere with the inhibitory activity implying that the change in conformation induced by calcium ions did not affect the antibody binding site: indeed, in one case stronger inhibition was observed. In contrast, preincubation with calcium ions before assay blocked the inhibition induced by CUB 7402. It is known that CUB 7402 maps the linear epitope 447–478 of tTG which overlaps the calcium binding region 343–452: calcium ions
could thus mask the binding site of the antibody interfering with its inhibitory activity.

Although inhibition was shown in in vitro experiments, the potential importance of this finding in the in vivo situation is suggested by in situ inhibition of tTG activity in umbilical cord tissue. Even so, it is difficult to predict the relevance in the pathogenesis of the coeliac lesion. Firstly, the inhibition we have observed is far from complete and the residual enzymatic activity could well be sufficient to fulfill all tTG biological functions; and secondly, it has recently been reported that tTG knockout mice have a normal small intestinal architecture, suggesting that the biological function of tTG may be replaced by other enzymes, although it cannot be excluded that anti-tTG antibodies may also inhibit these putative enzymes.

In conclusion, we have shown that CD related anti-tTG autoantibodies can inhibit enzyme activity. Further research is required to clarify to what extent this could play a role in the modulation of T cell activity and/or, independent of T cells, contribute to the mucosal lesion in CD.

ACKNOWLEDGEMENTS

This study was carried out with the financial support of Telethon–Italy (grant No E.11411); the commission of the European Communities, specific RTD programme “Quality of Life and Management of Living Resources” (QLK–1999–00387). Evaluation of the prevalence of coeliac disease and its genetic components in the European population. The work was partly supported also by MURST (PRIN), and by Regione Campania (Ricerca Sanitaria Finalizzata).

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Gut 2002 51: 177-181
doi: 10.1136/gut.51.2.177

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