Characterization of Heparin-binding Site of Tissue Transglutaminase

ITS IMPORTANCE IN CELL SURFACE TARGETING, MATRIX DEPOSITION, AND CELL SIGNALING

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Background: TG2 is a multifunctional matrix protein and cross-linking enzyme.

Results: Identification and mutation of its heparan sulfate (HS)-binding site blocks matrix deposition of TG2 as do inhibitors of syndecan shedding.

Conclusion: Coordinated binding to cell surface heparan sulfates facilitates TG2 cell surface trafficking and deposition into the ECM.

Significance: Blocking heparan sulfate binding provides an avenue for regulating the pathological roles of the enzyme.

Tissue transglutaminase (TG2) is a multifunctional Ca\(^{2+}\)-activated protein cross-linking enzyme secreted into the extracellular matrix (ECM), where it is involved in wound healing and scarring, tissue fibrosis, celiac disease, and metastatic cancer. Extracellular TG2 can also facilitate cell adhesion important in wound healing through a nontransamidating mechanism via its association with fibronectin, heparan sulfates (HS), and integrins. Regulating the mechanism how TG2 is translocated into the ECM therefore provides a strategy for modulating these physiological and pathological functions of the enzyme. Here, through molecular modeling and mutagenesis, we have identified the HS-binding site of TG2 and demonstrated how the HS-binding site for translocation of TG2 into the ECM through a mechanism involving cell surface shedding of HS. By synthesizing a peptide NPFLKNAGRDCSRRSS corresponding to the HS-binding site within TG2, we also demonstrate how this mimicking peptide can in isolation compensate for the RGD-induced loss of cell adhesion on fibronectin via binding to syndecan-4, leading to activation of PKCo, pFAK-397, and ERK1/2 and the subsequent formation of focal adhesions and actin cytoskeleton organization. A novel regulatory mechanism for TG2 translocation into the extracellular compartment that depends upon TG2 conformation and the binding of HS is proposed.

Overexpression of the Ca\(^{2+}\)-activated tissue transglutaminase (TG2) is closely related to a wide range of pathological processes, such as wound healing and scarring, fibrosis, celiac disease, multiple sclerosis, and tumor metastasis. Under stress, cells overexpress TG2, leading to enhanced externalization of the enzyme and the increased deposition of TG2 into the extracellular matrix (ECM). A huge amount of effort has been devoted to investigating the mechanism of TG2 secretion, but how it reaches the cell surface and is then translocated into the ECM is still unknown. Evidence suggests that TG2 is externalized via an unconventional secretion pathway, which may involve endosome binding and association with the cell surface receptors and/or the ECM protein fibronectin (FN). Once externalized, the enzyme can be deposited into the ECM where it forms a hetero-complex with FN and facilitates cell adhesion via an RGD-independent pathway, which during wound healing and matrix turnover can rescue RGD peptide-induced anoikis and subsequent loss of FN deposition through a syndecan-4/2 and \(\alpha_5\beta_1\) integrin signalling pathway.

However, in many pathological conditions, the continuous deposition of TG2 into the matrix results in increased protein cross-linking leading to the onset of and progression of tissue fibrosis and scarring, celiac disease, tumor metastases, and multiple sclerosis. Hence, modulating the mechanism of TG2 translocation into the ECM provides an important basis for developing therapeutic strategies that are able to regulate its physiological and pathological functions.

Interestingly, cell surface heparan sulfate proteoglycans (HSPG), for which TG2 has a high affinity binding, are also shed from the cell surface and can be deposited into the matrix where, like TG2, they are associated with a number of similar physiological and pathological conditions such as bone growth and development, wound healing and scarring, tumor progression, and angiogenesis. The HSPG family is composed of four major syndecans, syndecans 1–4, of which syndecan-1 and syndecan-4 are major resources for shedding into the ECM. By activation of cell surface matrix metallopro-
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Table 1

| Antigen                  | Host species | Clone   | Company          |
|--------------------------|--------------|---------|------------------|
| TG2                      | Mouse        | Monoclonal | Thermo Fisher   |
| p-FAK397                 | Mouse        | Monoclonal | Millipore       |
| Total FAK                | Rabbit       | Polyclonal | Santa Cruz Biotechnology |
| p-ERK1/2                 | Rabbit       | Polyclonal | Santa Cruz Biotechnology |
| Total ERK1/2             | Rabbit       | Polyclonal | Santa Cruz Biotechnology |
| Syndecan-4              | Rabbit       | Polyclonal | Invitrogen       |
| Syndecan-4 ectodomain   | Rabbit       | Polyclonal | Santa Cruz Biotechnology |
| Syndecan-2              | Rabbit       | Polyclonal | Invitrogen       |
| β1 Integrin             | Rabbit       | Polyclonal | Santa Cruz Biotechnology |
| α-Tubulin               | Mouse        | Monoclonal | Sigma            |
| Vinculin                 | Mouse        | Monoclonal | Sigma            |
| His6 tag                | Rabbit       | Monoclonal | Invitrogen       |
| α5β1 integrin blocking antibody and Isotype control antibody | Mouse | Monoclonal | Biologend      |
| HRP-conjugated anti-mouse secondary | Rabbit | Sigma    |                  |
| HRP-conjugated anti-rabbit secondary | Swine | Dako    |                  |
| TRITC-conjugated anti-mouse secondary | Rabbit | Dako    |                  |

and their universal negative control siRNA were obtained from Qiagen (Crawley, UK). KOD HOT Start DNA polymerase was obtained from Merck, and the QuikChange II site-directed mutagenesis kit was from Stratagene (Cheshire, UK). Cell transfection kits and reagents, including Nucleofector kit R used for transfection of NIH 3T3 and OK cells, were from Lonza Ltd. (Wokingham, Berkshire, UK). Guinea pig liver transglutaminase (gplTG2) was purified as described previously (14). The synthetic peptides GRGDTP, GRADSP, and the PKCa specific inhibitor Go6976 were obtained from Calbiochem. The P1 peptide (NPKFLKNAGRDCSRRSS) and scrambled control peptide P1s (FNRADLKPRCGSSNKRSS) the peptide corresponding to the N-terminal end of TG2, AEELVLERCDLELE (P2) and the scrambled peptide EECRELAEELLEDVL (P2s), the G2K1 peptide (GENPIYKSAVTTP-VNPKYEGRKRQIKWFQNNRMMWK) and its scrambled control peptide (GTAKINEPYSVTVPYGKNNVQRQIKWQNNRMMWK) fused to the antennapedia third helix sequence (15) were synthesized by Peptide Protein Research, UK. Fibronectin and its purified 45- and 70-kDa fragments, heparin, heparinase, and chondroitinase were purchased from Sigma. MMPs inhibitors were from Merck. Antibodies used in this work are listed in Table 1.

Experimental Procedures

Mammalian Cell Culture—Cell lines used in this study include human kidney epithelial cells HEK293T/17 (ATCC CRL-11268) and mouse embryo fibroblasts NIH 3T3 (ATCC CRL-11268), Chinese hamster ovary cells CHO K1 (ATCC CCL-61), and the heparan sulfate-deficient CHO-K1 derivative pgsD-677 (ATCC CRL-2244) (HS-mutant CHO), which were obtained from the American Type Culture Collection (ATCC, Manassas, VA). Additionally, human osteoblasts (HOB) were kindly provided by Prof. S. Downes (University of Nottingham, Nottingham, UK). HEK293T/17 cells, NIH 3T3 cells, HOB, and opossum kidney (OK) cells were cultured in Dulbecco’s modified Eagle’s medium (DMEM) containing 10% (v/v) heat-inactivated fetal bovine serum (FBS), 2 mM l-glutamine, nonessential amino acids, 100 units/ml penicillin, and 100 μg/ml streptomycin. CHO-K1 and HS mutant CHO cells were cultured in a mixture of F-12 (Ham’s) medium (Sigma) supplemented with 10% (v/v) FBS. Tetracycline (tet)-inducible Swiss 3T3 cells transfected with wild type human TG2 were cultured, and TG2 expression was induced as documented previously (13). Cells were all maintained at 37 °C in a 5% CO2-humidified atmosphere.

Vector, Antibodies, Kits, and Reagents—The pcDNA3.1 vector and the DH5α strain of Escherichia coli were purchased from Invitrogen. All restriction enzymes were obtained from New England Biolabs (Knowl Piece, UK). Wizard® Plus SV minipreps DNA purification system and Wizard® SV gel and PCR clean-up system were obtained from Promega (Southampton, UK). The endotoxin-free plasmid DNA maxi purification kit, and the human and mouse syndecan-4 targeting siRNAs and their universal negative control siRNA were obtained from Qiagen (Crawley, UK). KOD HOT Start DNA polymerase was obtained from Merck, and the QuikChange II site-directed mutagenesis kit was from Stratagene (Cheshire, UK). Cell transfection kits and reagents, including Nucleofector kit R used for transfection of NIH 3T3 and OK cells, were from Lonza Ltd. (Wokingham, Berkshire, UK). Guinea pig liver transglutaminase (gplTG2) was purified as described previously (14). The synthetic peptides GRGDTP, GRADSP, and the PKCa specific inhibitor Go6976 were obtained from Calbiochem. The P1 peptide (NPKFLKNAGRDCSRRSS) and scrambled control peptide P1s (FNRADLKPRCGSSNKRSS) the peptide corresponding to the N-terminal end of TG2, AEELVLERCDLELE (P2) and the scrambled peptide EECRELAEELLEDVL (P2s), the G2K1 peptide (GENPIYKSAVTTP-VNPKYEGRKRQIKWFQNNRMMWK) and its scrambled control peptide (GTAKINEPYSVTVPYGKNNVQRQIKWQNNRMMWK) fused to the antennapedia third helix sequence (15) were synthesized by Peptide Protein Research, UK. Fibronectin and its purified 45- and 70-kDa fragments, heparin, heparinase, and chondroitinase were purchased from Sigma. MMPs inhibitors were from Merck. Antibodies used in this work are listed in Table 1.

Generation of Wild Type TG2 and TG2 Mutants—Wild type (WT) human (16) and C277S mutant (17) TG2 were amplified by PCR using primers TG2-F/TG2-R and cloned into the KpnI/Ncol sites of pcDNA3.1. The WT TG2 plasmid was then used to generate a set of TG2 mutants. These were constructed by either PCR in the case of Δ1–15 TG2 or by using the QuikChange II site-directed mutagenesis kit (Stratagene, UK) for point mutations. Primers are shown in Table 2. The identity and proper arrangement of the TG2 mutants were verified by restriction analysis and nucleotide sequencing.

Transient Transfection and Expression of Wild Type TG2 and TG2 Mutants in Human HEK293T/17 Cells, Mouse NIH 3T3 Cells, and OK Cells—HEK293T/17, NIH 3T3, and OK cells, which express very low levels of endogenous TG2, were transiently transfected with wild type TG2 and the TG2 mutants. HEK293T/17 cells were transfected by the calcium phosphate procedure, whereas NIH 3T3 and OK cells were transfected by electroporation (Lonza Nucleofector kit R). Transfected cells
were grown for 48 h at 37 °C in a 5% CO₂-humidified atmosphere to allow expression prior to analysis.

**Syndecan-4 Silencing by siRNA Transfection**—The HP genome-wide siRNA sequence targeting human syndecan-4 (SI00046816) (5) and siRNAs targeting mouse syndecan-4 (SI02691710, SI02714285, SI00201341, and SI02671543) and the nonsilencing (NS) control siRNA were obtained from Qiagen (UK). The target sequences are non-homologous for any other syndecan types or cell surface receptors. The transfection was performed according to the manufacturer’s protocol. Briefly, 3 × 10⁵ cells/well HOB or tet-inducible Swiss 3T3 cells were seeded into 6-well plates for 24 h to reach 50–80% confluency. 150 ng of siRNAs were used for each transfection by using HiPerFect transfection reagents. Following 30–48 h siRNA transfection, cells were used in cell adhesion assays (5).

**Western Blotting**—Cells were lysed with cell lysis buffer (Santa Cruz Biotechnology). Lysates containing 50 µg of protein were dissolved in 2× Laemmli buffer (Sigma) and separated by SDS-PAGE. Western blotting was performed using specific primary antibodies as described above. Primary antibodies were detected using the appropriate secondary antibody conjugated to horseradish peroxidase. Detection was performed by using ECL chemiluminescence (ECL™ Western blotting system, GE Healthcare), where applicable α-tubulin was used for normalization of protein loadings using densitometry.

**Biotinylation of Cell Surface Proteins**—Cell surface proteins were labeled by biotinylation as described previously (6). Briefly, cell monolayers were rinsed three times with ice-cold PBS, pH 8.0, and labeled with 0.8 mM sulfo-NHS-LC-biotin dissolved in PBS, pH 8.0, at 4°C for 20 min. Cells were then washed with 50 mM Tris-HCl, pH 8.0, and lysed with 1% SDS at 4 °C in PBS, pH 8.0. Following denaturation at 95 °C, cell lysates were clarified by centrifugation at 14,000 × g at 20 °C, and 200 µg of protein was incubated overnight at 4 °C with NeutrAvidin-agarose resin. After washing three times with PBS, pH 8.0, the biotin-labeled proteins were dissolved in 2× Laemmli buffer separated by SDS-PAGE and subjected to Western blotting. Cell surface levels of TG2 shown were normalized, using densitometry, to the total cell lysate values using densitometry after the latter was corrected for protein loading using α-tubulin. The WT cells used as the 100% control. Value shown represents the mean values from two separate experiments unless stated otherwise.

**Detection of TG2 in the ECM**—Following transfection, cells were incubated for 24 h with 10% (v/v) serum, which was replaced with 1% (v/v) serum, and TG2 expression and matrix deposition allowed to proceed for a further 24 h were then detached with 2 mM EDTA in PBS, pH 7.4, and the ECM was extracted with 0.1% (w/v) deoxycholate in PBS, pH 7.4. The residual deoxycholate-insoluble ECM proteins were dissolved in 2× Laemmli buffer for further analysis by SDS-PAGE and Western blotting using Cub7402.

To detect the presence of TG2 in the ECM of TG2 transfected tet-inducible Swiss 3T3 cells, the cells were seeded into 6-well plates at the density of 3 × 10⁵ cells/well. TG2 expression was induced by withdrawing tetracycline from the culture system (tet− cells), whereas cells cultured in the presence of tetracycline (tet+ cells) were used as the control. After induction of TG2 expression in the presence of certain treatments for 48 h, the cells were lifted with 2 mM EDTA in PBS, pH 7.4, lysed in cell lysis buffer, and used as the cell lysate fractions. The ECM fractions were washed once and collected into 2× Laemmli buffer. Western blotting was performed to detect the target proteins by using specific antibodies. The treatments included the following: general MMP inhibitor GM6001, MMP-2/9 inhibitor, MMP-3 inhibitor, and MMP-8 inhibitor at the concentration of 10 µM, 10 µg/ml heparin, 15 milliunits/ml heparinase or chondroitinase, 0.5 µM PMA or 5 µM PKCα inhibitor Go6976. Matrix levels of TG2 shown were normalized to the total cell lysate values using densitometry after the latter was corrected for protein loading using α-tubulin. The WT cells or for the Swiss 3T3 cells the untreated cells were used as the 100% control with values shown represented as a percentage of this. Values shown represent the mean values from two separate experiments unless stated otherwise.

**Dot Blotting for ECM Syndecan-4**—To detect the presence of the shed syndecan-4 ectodomain in the ECM, modified dot blotting was applied. Briefly, tet-inducible 3T3 cells were seeded into 12-well plates at the density of 2 × 10⁵ cells/well and induced for TG2 expression for 48 h with the treatments (including PMA, Go6976, and the MMP-2/9 inhibitor) as described for TG2. The matrix fractions were then collected as described for the matrix TG2 and then dissolved into 2× Laemmli buffer. The ECM fractions containing equal amounts of protein were then loaded onto nitrocellulose membranes under vacuum using a dot blotter, washed, and blocked with blocking buffer (5% (w/v) dried-fat-free milk dissolved in TBS/Tween, pH 7.4), and the presence of syndecan-4 was detected by immunoprobeing with a specific antibody recognizing the ectodomain of syndecan-4.

**Measurement of TG2 Binding to FN by ELISA**—Microtiter 96-well plates were coated with 50 µl of 5 µg/ml FN in 50 mM Tris-HCl, pH 7.4, at 4 °C overnight. The wells were blocked for
30 min with 3% BSA in PBS, pH 7.4, and washed twice with PBS/Tween, pH 7.4, and once with PBS, pH 7.4. Aliquots (100 μl) of cell lysate containing 60 μg of protein were added to the FN-coated wells and incubated for 1 h at 37 °C. After washing three times with PBS, pH 7.4, wells were blocked with 100 μl of 3% BSA in PBS, pH 7.4, for 30 min at room temperature and then incubated with 100 μl of Cub7402 (1:1000 dilution in blocking buffer) for 2 h at 37 °C. After washing three times with PBS, pH 7.4, the wells were incubated with 100 μl of rabbit anti-mouse IgG-HRP-conjugated antibody (1:1000 dilution in blocking buffer) for 2 h at 37 °C. HRP was detected by the addition of 100 μl of o-phenylenediamine substrate solution (Fast OPD, Sigma). Color development was terminated by the addition of 100 μl of 2.5 m H₂SO₄, and absorbance at 490 nm was measured. The amount of TG2/mutant TG2 binding to the FN was normalized between samples by using the densitometry values from Western blotting for the TG protein found in each of the cell lysates used. α-Tubulin was used to normalize for any differences in protein loading.

**Inhibition of Purified TG2 Binding to FN by TG2-derived Peptides**—96-Well Microtiter plates were coated overnight with either full-length FN (5 μg/ml) or its N-terminal fragments of 45-kDa (54 μg/ml) or 70-kDa (45 μg/ml) that contain the TG2-binding site, in 50 mM Tris-HCl, pH 7.4, as described previously (6, 19). Wells were blocked with 3% BSA (w/v) in TBS, pH 7.6. After three washes with PBS, pH 7.4, competitive peptides diluted in PBS, 2 mM EDTA, pH 7.4, were added to the wells at concentrations from 10 μM to 1 mM and incubated for 1 h at room temperature. Wells were washed three times with PBS, pH 7.4, and purified human recombinant TG2 was added at a final concentration of 2 μg/ml in the presence of the different concentrations of P2 or P2s peptides and incubated for 1 h at room temperature. After three washes with PBS, pH 7.4, bound TG2 was measured via ELISA as introduced above.

For the binding ability of TG2 when in either the fully closed or open conformation, gpITG (20 mg/ml) was reduced by pre-incubation with 1 mM DTT and then diluted (at least 20-fold) in PBS, pH 7.4. The reduced gpITG 20 μg/ml was preincubated with 1 mM GTP or GTPγS (closed conformation) or TG2 site-directed irreversible inhibitors R281 or R283 (extended conformation) at the concentration of 500 μM (in the presence of 10 mM Ca²⁺) at room temperature for 30 min. Treated enzymes were then incubated with the FN-coated wells at 37 °C for 1 h as described under “Experimental Procedures.” 20 μg/ml gpITG in 2 mM EDTA in PBS, pH 7.4, was used as the positive control, and 2 mM EDTA in PBS, pH 7.4, was used as the negative control. TG2 bound to FN was detected via ELISA as described. Values represent the mean ± S.D. absorbance at 450 nm from three experiments.

**Binding of TG2 to Heparin-Sepharose**—HEK cells, transiently transfected with wild type and mutant TG2, were washed twice with ice-cold PBS, pH 7.4, and lysed by the addition of 150 μl of 20 mM Tris-HCl, pH 7.4, 10 mM EGTA, 2 mM EDTA, 1 mM NaF, and 1 mM Na₂VO₃. After clarification by centrifugation (300 × g for 10 min), cell lysates were mixed with 450 μl of 50 mM Tris-HCl, 1 mM EDTA, 1 mM DTT, pH 7.5, and applied to a 5-ml heparin-Sepharose column (GE Healthcare) equilibrated in 50 mM Tris-HCl, 1 mM EDTA, 1 mM DTT, pH 7.5 (buffer), at a flow rate of 1 ml/min. For the analysis of GTP-bound TG2 (closed conformation), lysates were preincubated for 1 h at room temperature with 0.5 mM GTP in 50 mM Tris-HCl, 2 mM MgCl₂, 1 mM EDTA, 1 mM DTT, pH 7.5. For the analysis of inhibitor-reacted TG2 (open conformation), lysates were incubated with 0.5 mM irreversible inhibitor R281 (20) and 10 mM CaCl₂. The column was washed with 25 ml of buffer, and protein was eluted with a linear gradient of increasing NaCl concentrations (0–1 M) in buffer. Fractions were assayed for TG2 activity and analyzed for the presence of TG2 antigen by SDS-PAGE and Western blotting using Cub7402.

**Measurement of TG Activity in Cell Lysates and Heparin-Sepharose Fractions**—TG activity in column fractions was measured by biotin-X-cadaverine incorporation into N,N’-dimethylcasein as described previously (21). For the analysis of TG activity in cell lysates, 100 μl of 10 mg/ml N,N’-dimethylcasein in 50 mM Tris-HCl, pH 8.0, plates were washed with TBS/Tween, pH 7.6, and TBS, pH 7.6, and 50 μl of column flow-through as well as each eluted fraction were added into the coated wells. Additionally, 50 μl of 100 mM Tris-HCl, pH 8.0, 0.25 mM biotin X-cadaverine, 10 mM DTT, 20 mM CaCl₂ (or 5 mM EDTA as control) was added into each well. The reaction was allowed to proceed for 1 h at 37 °C. The plate was then washed once with TBS/Tween, pH 7.6, and TBS, pH 7.6, before being blocked with 100 μl of 3% (w/v) BSA in TBS, pH 7.6, for 30 min at 37 °C. After another wash, biotin X-cadaverine incorporation into N,N’-dimethylcasein was detected by incubation for 1 h at 37 °C with 100 μl of Extravidin-peroxidase (Sigma) diluted 1:2000 in 3% (w/v) BSA in TBS, pH 7.6. After another set of washes, TG2 activity was measured using Sigma Fast OPD, and plates were dissolved in 20 ml of distilled H₂O. The color was developed by adding 2.5 m H₂SO₄, and the absorbance at 490 nm was measured using a microplate reader.

For cell lysates the incorporation of biotin-X-cadaverine into FN as described previously was used (21). Briefly, 96-well plates were coated with 5 μg/ml FN in 50 mM Tris-HCl, pH 7.4, and incubated overnight at 4 °C. After washing with 50 mM Tris-HCl, pH 7.4, wells were blocked with 3% (w/v) BSA in 50 mM Tris-HCl, pH 7.4, for 30 min at 37 °C. Enzyme reactions contained 50 μg of cell lysate protein in 50 mM Tris-HCl, pH 7.4, buffer containing 5 mM CaCl₂ (or 5 mM EDTA as control), 10 mM DTT, 0.132 mM biotin X-cadaverine each done in triplicate. Reactions were allowed to proceed for 2 h at 37 °C. Plates were then processed as described above.

**Cell Adhesion Assays**—The cell adhesion assay was carried out as described previously (6). Briefly, 96-well plastic tissue culture plates were coated with 5 μg/ml FN in 50 mM Tris-HCl, pH 7.4, at 4 °C overnight, and after washing three times with 100 μl of 50 mM Tris-HCl, pH 7.4, gpITG (20 μg/ml) in 2 mM EDTA in PBS, pH 7.4, was added to the wells for 1 h at 37 °C, and the wells were washed three times with 50 mM Tris-HCl, pH 7.4. Serum-starved (for 16 h) HOB cells with different treatments as introduced below in serum-free medium were detached by trypsinization and then treated with trypsin inhibitor. Cells were washed three times with serum-free medium and seeded onto either FN or TG-FN matrix for 20–40 min. Peptides P1 or scrambled P1 (P1s) were added at concentrations between 100 and 300 μg/ml. To determine the RGD-in-
dependent cell adhesion, assays were performed by incubating cells with either RGD or RAD peptide (100 μg/ml) unless stated otherwise in the presence of the P1 or P1s peptides for 20 min prior to seeding of the cells. Attached cells were washed once with PBS, pH 7.4, and then fixed with 3.7% paraformaldehyde, further permeabilized with 0.1% (v/v) Triton X-100 in PBS, and co-stained with May-Grunwald and Giemsa stains as described previously (5). Images of stained cells from nonoverlapping fields of view were photographed at ×20 magnification and analyzed using the imaging analysis program Scion Image (National Institutes of Health). Cell attachment and spreading were quantified, and the number of cells per image was assessed as described previously (19). Cell attachment on FN without peptide was considered as the control value for all the experiments unless stated otherwise. The mean number of attached cells from at least three wells was calculated and that of the control was considered as 100%. The mean number of attached cells (cell attachment) for each sample was then expressed as the percentage of cell attachment on FN. The mean percentage of attached cells that are spread (cell spreading) for each sample was determined separately, and the mean percentage of spread cells on FN control was expressed as 100%. The mean percentage of spread cells for each sample was then normalized against that of FN control. Cell attachment on FN without the RGD peptide was considered as the control value for most of the experiments unless stated otherwise. The mean number of attached cells from three wells was calculated and that of the control was considered as 100%. The mean number of attached cells (cell attachment) for each sample was then expressed as the percentage of cell attachment on FN. The mean percentage of attached cells that are spread (cell spreading) for each sample was determined separately, and the mean percentage of spread cells on FN control was expressed as 100%. The mean percentage of spread cells for each sample was then normalized against that of FN control. Cell pretreatments included the following: PKCα inhibitor Go6976 (5 μM) or GSK21 peptide (8 μM, which blocks the interaction between PKCα and the intracellular domain of β1 integrins) for 1 h in serum-free medium prior to cell detachment; heparinase (15 milliunits/ml) to cells in suspension in serum-free medium for 1 h or a5β1 integrin blocking antibody N1K1-SAM-1 and its isotype control antibody (20 μg/ml) in serum-free medium for 1 h. For the detection of the signaling molecules via Western blotting, the cell adhesion assay was performed in 60-mm Petri dishes, and cells were collected into cell lysis buffer (Santa Cruz Biotechnology) as described previously and precleared by centrifugation at 300 × g for 10 min. Western blotting was performed with specific anti-p-397 FAK or p-ERK1/2 antibodies. Membranes were stripped (19), and total FAK and ERK1/2 were detected using anti-FAK and anti-ERK1/2 antibodies as listed in Table 2, whereas α-tubulin was used as an equal loading standard (5).

Fluorescence Staining—Cell adhesion on FN matrix in the presence of either the P1 or the P1s peptide (100 μg/ml) was performed as described above. Cells were seeded in 8-well glass chamber slides (8 × 104 cells/well) previously coated with FN and allowed to attach and spread for 20–40 min. Cells were fixed and permeabilized as described above and then blocked with 3% BSA in PBS, pH 7.4 (the blocking buffer), for 30 min. For the actin stress fiber staining, the cells were incubated with FITC-labeled phalloidin (20 μg/ml) in blocking buffer. For the focal contact staining in HOB cells, after treatment with blocking buffer, the cells were incubated with anti-vinculin antibody (1:100 dilution) followed by anti-mouse IgG-TRITC at 37 °C for 2 h/incubation. Slides were mounted with Vectashield mountant (Vector Laboratories) and examined via confocal microscopy (5).

For the FITC-cadaverine incorporation in situ assay, tet-inducible Swiss 3T3 cells were seeded into 8-well chambers at the density of 7 × 104 cells/well. Together with the withdrawal of the tetracycline, the cells were treated with PKCα inhibitor Go6976, MMP-2/9 inhibitor, MMP-8 inhibitor, heparin, and heparinase as described above. After a 48-h induction of TG2 expression, fresh medium with 0.5 mM FITC-cadaverine (with or without the treatment) was incubated with the cells at 37 °C for 1 h. The wound was introduced into the mono-cell layer using a sterile plastic pipette tip, and the cells were incubated for another 1 h as described previously (22). After washing three times with PBS, pH 7.4, the cells were fixed with methanol at −20 °C for 10 min and then mounted in Vectorshiel mountant medium, and the fluorescence signal was detected by fluorescence microscopy.

Co-immunoprecipitation Assay for the Interaction between TG2 and Syndecans—Tet-inducible Swiss 3T3 cells (5 × 104 cells/well) were seeded into a 6-well plate, and TG2 expression was induced by withdrawing tetracycline from the cell culture system. 200 μg/ml P1 or P1s peptide was used together with the TG2 induction. The cell lysates were collected into the co-immunoprecipitation buffer, and co-immunoprecipitation was performed to detect the interaction between TG2 and syndecan-4 as introduced previously (14). Briefly, 0.5 μg of anti-syndecan-4 antibody was used to pull down the syndecan-4-immunocomplex from the precleared cell lysates, and Western blotting was carried out to detect the presence of TG2 in the immunocomplex by using specific anti-TG2 antibody Cub7402. The noninduced Swiss 3T3 cells with the undetectable TG2 expression were used as the negative control.

Solid Binding Assay for P1 Peptide Binding to Syndecans—To detect the binding specificity of P1 peptide toward syndecan-4 and syndecan-2, ELISA plates were coated with P1 peptide at the concentration of 1 μg/ml in Na2CO3 solution, pH 9.6, at 4 °C overnight. The wells were blocked with 3% BSA in PBS, pH 7.4, after washing with 50 mM Tris-HCl, pH 7.4. 2 μg/ml recombinant human syndecan-4 or syndecan-2 (with the His tag) in 50 mM Tris-HCl, pH 7.4, was incubated with the wells at 37 °C for 1 h. The presence of syndecan-4 or syndecan-2 bound to the peptide-coated wells was detected by using anti-His tag antibody and anti-mouse secondary antibody. The signals were detected by using OPD substrate, and the absorbance was measured at 490 nm.

Docking Studies—The crystal structures 1KV3 and 2Q3Z were downloaded as PDB files from the Protein Data Bank and opened in the software CAChe WorkSystem Pro version 7.50.0.85 (Fujitsu Ltd.). Hydrogen was added, and water and ions were deleted. The hydrogen atom positions were relaxed by conducting an MM2 geometry optimization for each structure.
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where all the non-hydrogen atom positions were locked. For the HS1 site, docking sites were defined by selecting all the amino acid residues within either 5 or 8 Å of residues Lys-600, Arg-601, and Lys-602. For the HS2 site, docking sites were defined by selecting all the amino acid residues within either 5 or 8 Å of residues Lys-202, Lys-205, Arg-209, Arg-213, and Arg-222. Both crystal structures contained missing residues, but these were far enough away from the defined docking sites such that their absence would not interfere with the docking studies. Using the same software, three ligand structures (a dimer, a pentamer, and a hexamer) were defined by taking residues 2–3, 2–6, and 2–7, respectively, from the glycosaminoglycan structure PDB code 1HPN. Using the Project Leader module and the Active Site docking component of the same software, the three ligands were each docked three times into the defined docking sites of both proteins using the flexible ligand and flexible active site side chain options. Other parameters and options included the following: Use Amber van der Waals; population size 50; maximum generations 3000; crossover Rate 0.8; mutation Rate 0.2; elitism number 5; local search rate 0.06; maximum iterations local search 300.

Data Analysis—Results shown are the mean ± S.D. unless otherwise stated. The differences between data calculated were undertaken by Student’s t test and indicated as significant when p < 0.05.

RESULTS

Identification of Putative Heparin-binding Motifs in TG2—HS-binding motifs may be composed of basic amino acid-containing sequences, such as XBBXBX or XBBXXBXY, where B is a basic amino acid whose side chain is exposed on the protein surface and X is a neutral or hydrophobic amino acid, whose side chain is directed toward the protein interior (23). Examination of the primary amino acid sequence of TG2 for these linear consensus HS-binding motifs revealed one such sequence 261LRRWK266 close to the active site of TG2 (Fig. 1A) (24). Examination of the crystal structure of TG2 (PDB 1KV3) shows that 261LRRWK266 is part of an α-helix, whereas the XBBXBX consensus must be in a β-sheet in order for the basic residues to face the same direction. A common structural theme of linear HS-binding motifs is that there are two basic residues ~20 Å apart to accommodate a pentasaccharide, facing in opposite directions on an α-helix (25). Because 261LRRWK266 is too short to satisfy this requirement, it is unlikely to be able to bind HS and so was ruled out. The three-dimensional arrangement of basic amino acid residues is likely to be more important than linear clustering, such that many HS-binding motifs can also be composed of sequence-distant basic amino acid residues (26). Examination of the crystal structure of TG2 revealed two likely candidate motifs, KIRILGEPKQKRK602 (HS1), which is located at the tip of C-terminal β-barrel 2, and another composed of 202KFLKNA GRDCSRRSSPPYYGR222 with Lys-387 (HS2), forming a shallow pocket lined with basic residues (Fig. 1, A–C). Importantly, both of these sites are likely to allow simultaneous binding of fibronectin (Fig. 1A) as described previously (6, 19, 27).

Heparin-derived Oligosaccharide Docks into a Characteristic Heparin Binding Pocket in TG2—The HS1 and HS2 regions were docked with heparin-derived oligosaccharides. An iduronic acid-2-sulfate-glucosamine-2,6-disulfate disaccharide is the most common repeating unit of heparin. This docked better with the HS2 region than the HS1 region of the closed form of TG2 (PDB 1KV3), although both sites docked well. A pentasaccharide and a hexasaccharide with the same repeating units bound with a similar enthalpy to both the HS2 and HS1 site (see Fig. 1, D and E). The same oligosaccharides failed to dock as successfully with the open conformation of TG2 (PDB 2Q3Z) at both the HS1 and HS2 sites, although a slight preference was shown for the HS1 site (supplemental Table 1).

Expression of the FN and HSPG TG2 Mutants in Mammalian Cell Systems—Mutant TG2 enzymes for the binding of FN (D94A,D97A), the N-terminal deletion (Δ1–15), and HS mutants HS1 (K600A,R601A,K602A) and HS2 (K205A,R209A) were generated by mutagenesis of plasmid pcDNA3.1-TG2. Mutants were confirmed by nucleotide sequencing and transferred into HEK293T/17 and NIH 3T3 cells to assess expression. All mutants were expressed in both cell types (Fig. 2, A and D), and Western blotting showed bands of the expected molecular weight (Fig. 2, A and D), although at slightly different levels, with the Δ1–15 TG2 mutant showing significantly less expression in both cell types when compared with the WT cells (Fig. 2, B, C, E, and F). Comparable transamidating activities were found with all mutants when normalized to TG2 protein (Fig. 2, C and F) apart from the Δ1–15 TG2 mutant that was significantly lower (~50%) than the WT enzyme in HEK293T/17 cells but still present, suggesting no evidence for gross misfolding.

High Affinity Heparin-binding Site of TG2 Is Located in the Catalytic Core Domain—TG2 binds to heparin-Sepharose with high affinity and has been used for its purification. The binding strength of TG2 and mutants HS1 and HS2, expressed in HEK293T/17 cells, to heparin-Sepharose was determined by elution with an increasing salt gradient (Fig. 3A). TG2 was eluted over a very broad peak ranging from 100 to 500 mM NaCl with optimum elution at 330 mM NaCl. To rule out that the observed low affinity binding was due to denatured TG2, the transglutaminase activity of eluted fractions was determined, and this showed that specific activity did not differ significantly between fractions (supplemental Fig. 1).

Mutant HS1 bound heparin identically to TG2, with low and high affinity, whereas HS2 lost its high affinity binding to heparin, with a large percentage of the mutant showing no binding to the heparin column and the remainder showing low affinity binding, eluting at a NaCl concentration of 60 mM (Fig. 3A), suggesting that residues Lys-205 and Arg-209 contribute to the high affinity binding to heparin. Because mutant HS1 had a higher net charge reduction than HS2, this alteration in HS2 binding was not simply due to electrostatic interaction. Comparable studies with the D94A,D97A FN mutant showed this mutant had a comparable elution profile to the WT enzyme. However, the Δ1–15 TG2 FN-binding mutant showed only a low affinity binding to the heparin column with the major peak eluting at around 250 mM NaCl.
High Affinity Heparin Binding Is Dependent on TG2 Conformation—Because TG2 can adopt two extremes of conformation in the presence or absence of GTP, the effect of GTP binding on the association of TG2 to heparin was also investigated (Fig. 3A). In the presence of GTP, which results in a compact globular conformation, all of the TG2 bound with high affinity to heparin with the major peak eluting at about 330 mM NaCl. In contrast, after reaction with the irreversible peptidic inhibitor R281 (20) or mutation of the active site Cys-277 to Ser, which restricts the conformation to an extended form, the TG2 bound to heparin with lower affinity with the major peak eluting at 250 mM NaCl. Because the GTP-bound globular form of TG2 bound to heparin with high affinity, although the extended R281-bound form still retained an affinity greater than that of the HS2 mutant, this suggests that the loss of high affinity binding of HS2 is not solely due to an altered conformation. Hence, TG2 residues Lys-205 and Arg-209 are very likely to be directly involved in high affinity heparin binding.

Studies with the HS2 Peptide NPKFLKNAGRDCSRRSS (P1)—Our previous studies (6, 19, 27) indicated that TG2, when regulating cell adhesion via syndecans, interacted directly with syndecan-4, but not syndecan-2, which was activated indirectly via PKCa. We therefore tested the binding specificity of the TG2 HS2 region toward these two syndecans. A peptide 200NPKFLKNAGRDCSRRSS216 (P1), which mimics the heparin binding domain in TG2, was synthesized. This peptide was chosen for its potential to fold correctly, and the hydrophobic C-terminal sequence PVYVGR, which may affect solubility, was excluded. A solid binding assay was undertaken to study the binding ability of this peptide toward syndecan-4 and syndecan-2.
decan-2, in which His-tagged recombinant human syndecan-4 or syndecan-2 was assessed for their binding to the immobilized P1 peptide and then detected by using anti-His tag antibody. As shown in Fig. 3B, strong preference was shown for syndecan-4 binding to the P1 peptide when compared with the binding of syndecan-2, suggesting that the binding specificity of the P1 peptide toward syndecan-4 confirms our previous findings that TG2 binds preferentially to the heparan sulfate chains on the cell surface syndecan-4 and not syndecan-2 (6). Using immunoprecipitation, we further tested the competitive effect of the P1 peptide for the interaction between syndecan-4 and TG2. To confirm this specificity and to rule out any potential toxicity of the P1 peptide to the cells, a scrambled peptide (P1s, FNRADLKPRCGSSNKSR) was also used. Fig. 3C shows that the immunoprecipitation of TG2 with anti-syndecan-4 anti-body was reduced around 50% in the presence of the P1 peptide, but no effect was found with the scrambled analogue, thus confirming the specificity of the P1 peptide for syndecan-4.

Using cell studies, we then investigated if the peptide P1 can mimic TG2 in cell adhesion studies and as such can either substitute for or abolish the TG2-mediated compensation of the RGD-mediated loss of cell adhesion. In our earlier studies using both HOB and mouse embryonic fibroblast (MEF) cells, it was shown that extracellular TG2 bound to matrix FN could compensate for the loss of integrin-mediated cell adhesion in the presence of RGD peptides in a process requiring cell surface syndecan-4 but not its transamidase activity (27). Because direct interaction of TG2 with syndecan-4 is essential for this process, the HSPG binding properties of TG2 are therefore critical. The P1 peptide was tested for its ability to compensate for
the RGD-induced loss of cell adhesion on FN in HOB cells. At a peptide concentration between 0.01 and 200 μg/ml, significant compensation started as low as 5–10 μg/ml with maximum compensation achieved between 50 and 100 μg/ml (Fig. 4A).

We then investigated whether the P1 peptide, when used at a concentration of 100 μg/ml, could compete for the binding of syndecan-4 and abrogate the compensatory effects of TG-FN when the cells were plated onto the TG-FN matrix in the presence of the RGD peptide. Fig. 4B shows that P1 peptide only has a small but not significant dose-dependent negative effect on the attachment and spreading of HOB cells on TG-FN in the presence of the RGD peptide, whereas the scrambled control peptide P1s had almost no effect. This suggests that even though the peptide may compete with TG2 for the syndecan-4-binding site, its ability to mimic TG2 in compensating for RGD-induced loss of cell adhesion means no large changes in cell adhesion were likely to be observed. Neither peptide significantly affected the binding of cells to FN alone when used at similar concentrations of 100–300 μg/ml, although a small (~10%) but significant enhancement of adhesion was found for P1 peptide when compared with the P1s peptide (Fig. 4B). Importantly, the TG2 mimicking P1 peptide was able to restore actin cytoskeleton formation and focal adhesion assembly, which was disrupted with the RGD peptide (Fig. 5, A and B).

FIGURE 3. A, differences in the binding strength of TG2 mutants to heparin-Sepharose. Cell lysates from HEK293/T17 cells transfected with TG2 and mutants were applied to a heparin-Sepharose column (5 ml) and eluted with a NaCl gradient as shown. Both flow-through and resulting fractions (1–25 as shown) were assayed for TG2 by Western blotting with Cub7402. B and C, binding specificity of P1 peptide toward syndecan-4. B, solid binding assay was carried on to detect the binding between P1 peptide and recombinant human syndecan-4 or syndecan-2. The presence of the recombinant protein was detected by using anti-His tag antibody as described under “Experimental Procedures.” C, representative Western blot performed with TG2 tetracycline-inducible Swiss 3T3 cells (tet−) to detect the presence of TG2 in the syndecan-4 immunocomplex in the presence of P1 or P1s peptide as described under “Experimental Procedures.” CNTL, control.
although the scrambled analogue FNRADLKPRCGSSNKSR (P1s) showed no effect on either of these.

**P1 Peptide Acts via Binding and Activation of Syndecan-4-mediated Cell Signaling**—Further confirmation that the P1 peptide can act like extracellular TG2 in compensating for $\alpha_5\beta_1$ integrin, loss (6) of cell adhesion was shown by preincubated cells with the P1 peptide prior to incubation with the $\alpha_5\beta_1$ integrin blocking antibody (NKI-SAM-1). The P1 peptide led to a significant increase in cell adhesion in the presence of the inactivating antibody when compared with the control cells incubated with the antibody in the presence and absence of the P1 scrambled peptide (Fig. 6A).

To test the importance of cell surface HS in the binding of P1 peptide, HOB cells were pretreated with heparinase or chondroitinase prior to the cell adhesion assay. Treatment with heparinase, but not chondroitinase, abolished the compensatory effect of the P1 peptide on the RGD-induced loss of cell adhesion (Fig. 6B). To confirm that the P1 peptide is binding to cell surface syndecan-4 molecules, HOB cells were treated with syndecan-4 siRNA and scrambled control siRNA as documented previously (5). As found previously (5), treatment of HOB cells with syndecan-4 siRNA led to a 50% reduction in protein expression (supplemental Fig. 2A) without affecting either syndecan-2 or $\alpha_5\beta_1$ integrin expression, another two major players in the TG-FN complex-mediated signaling pathway (6).

This loss in expression of syndecan-4 led to a comparable but significant reduction in the compensatory effect for the P1 peptide on the RGD-induced loss of cell adhesion (Fig. 7A). The scrambled siRNA had no significant effect on either expression or cell adhesion, further confirming the essential role of syndecan-4 in the P1 peptide-related cell adhesion process.

Our next step was to demonstrate the activation of protein kinase Cα (PKCa) in the signaling effects mediated by the P1 peptide. The PKCa inhibitor Go6976 and the GK21 peptide (which is reported to compete with the PKCa-binding site on $\beta_1$ integrin (15)) both inhibit PKCa activation events leading to a significant loss of the compensatory effect of the P1 peptide on the RGD-induced loss of cell adhesion (Fig. 7B and C), strongly suggesting that the P1 peptide is acting in a comparable signaling manner to that of TG2. To confirm our observations that

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**FIGURE 4. Effect of P1 peptide on interaction between TG2 and syndecan-4 and the RGD-induced loss of cell adhesion on FN.**

A. compensatory effect of P1 peptide on loss of cell adhesion with the RGD peptide. HOB cells were treated with P1 peptide (0.01–200 $\mu$g/ml) with RAD or RGD peptides on FN. RAD-treated HOB cells were used as the control (the 0 group). Mean percentage value of the control attached and spread cells on FN ± S.D. (control) was set at 100%. B. effect of P1 peptide on the cell adhesion on TG-FN matrix. FN or TG-FN matrix was prepared as introduced before, and HOB cell adhesion assay was performed in the presence of P1 or P1s peptide (100–300 $\mu$g/ml) on FN or TG-FN matrix as described under “Experimental Procedures.”
the P1 peptide is inducing intracellular signaling pathways comparable with the TG-FN matrix, we looked at the activation of FAK by phosphorylation at tyrosine 397 (Fig. 8, A and B) and phosphorylation of extracellular signal-regulated kinase1/2 (ERK1/2) (Fig. 8, C and D). In each case, in the presence of RGD and P1 peptide, the compensation of the RGD-induced loss of adhesion was paralleled by a significantly greater phosphorylation of p-FAK397 than that found with the P1s peptide, thus confirming the ability of the P1 peptide to restore focal adhesion assembly and activation of p-ERK1/2 (Fig. 8, A–D).

FIGURE 5. A, co-staining of the actin cytoskeleton and vinculin of P1- or P1s-treated HOB cells on FN was undertaken as described under “Experimental Procedures.” B shows higher magnification of actin and vinculin staining in the presence of the P1 peptide and RGD with single cell image representing the co-localization of vinculin and actin. Arrows indicate presence of focal adhesion points.
N Terminus of TG2 Is Involved in FN Binding—The H9004 1–15 and D94A,D97A TG2 mutants were assessed for their ability to bind to FN in a solid phase binding assay utilizing purified FN and clarified lysates from HEK293T/17 cells transfected with the different mutants. The D94A,D97A mutant described previously (28) demonstrated significantly less binding (~50%) compared with wild type TG2, whereas the H9004 1–15 deletion mutant previously described (28) had 60% less (~p < 0.05) binding capacity of the wild type (Fig. 8E). A competitive peptide corresponding to the N-terminal deletion site of TG2 2AEELVLERCDLELE15 (P2) (TG2 undergoes N-terminal post-translational modification of the N-terminal methionine) was tested for its ability to inhibit wild type TG2 binding to FN and the smaller TG2 binding N-terminal FN fragments of 45 and 70 kDa in a solid binding assay. The P2 peptide resulted in a moderate but significant inhibition of binding to FN and its smaller fragments at higher concentrations (500 μM to 1 mM) of the peptide not shown with the scrambled P2s peptide (Fig. 8F and G), suggesting that the N-terminal residues 1–15 of TG2 may be important for FN binding, in addition to the published D94A,D97A site. However, the limited inhibition seen with this peptide suggests that the FN-binding site around the N-terminal site of TG2 may be more complex and that a larger structural unit than that described previously (28) may be involved in FN binding.

HS Binding Is Required for ECM Localization of TG2—Because TG2 has been shown to have a high affinity binding for heparan sulfates (29), we investigated the ability of the FN- and HS-binding mutants to be secreted to the cell surface and/or deposited into the ECM. Using transfected NIH 3T3 fibroblasts, the cellular localization of the D94A,D97A FN mutant, the H9004 1–15 mutant, and the HS1 and HS2 TG2 mutants were tested for their presence at the cell surface using the cell surface biotinylation assay. For detecting the presence of these mutant TG2s in the ECM, transfected NIH 3T3 cells were used, which actively deposit ECM. The TG2 H9004 1–15 mutant was not detected on the cell surface of the NIH3T3 cells (Fig. 9A), whereas the HS2 mutant was detected but in very reduced amounts compared with the wild type TG2. The levels of the FN mutant D94A,D97A and the HS1 mutant were found at comparable levels on the cell surface with that shown for the wild type TG2 (Fig. 9A). A comparable picture was observed for the presence of these mutant TG2s in the ECM, transfected NIH 3T3 cells were used, which actively deposit ECM. The TG2 ∆1–15 mutant was not detected on the cell surface of the NIH3T3 cells (Fig. 9A), whereas the HS2 mutant was detected but in very reduced amounts compared with the wild type TG2. The levels of the FN mutant D94A,D97A and the HS1 mutant were found at comparable levels on the cell surface with that shown for the wild type TG2 (Fig. 9A). A comparable picture was observed for the presence of the TG2 WT and mutants in the ECM of NIH 3T3 cells apart for the HS2 mutant and the H9004 1–15 mutant that were barely or not detectable, respectively, within the matrix (Fig. 9B).

Cellular Localization of TG2 in CHO WT and CHO-HS-M Cells—To confirm the importance of cell surface HS in the cell surface distribution of TG2, the HS-deficient Chinese hamster ovary-K1 (CHO-K1) cell derivative pgsD-677 was compared.
with WT CHO-K1 cells with respect to their distribution of TG2 at the cell surface (Fig. 9, C and D), which clearly demonstrates that the amount of TG2 found at the cell surface in the CHO mutant cells is considerably reduced when compared with the WT cells.

Involvement of Syndecan Shedding in Deposition of Matrix TG2—We next determined whether cell surface syndecan shedding may be a possible mechanism for translocating HS-bound TG2 into the extracellular matrix. Because MMPs are reported to be involved in the shedding of cell surface HS into the ECM (12), a general MMP inhibitor GM6001 was first used to treat TG2-transfected Swiss 3T3 cells. TG2 expression can be induced in these cells via the tet-inducible promoter and is externalized and deposited into the matrix in large amounts (27). Following induction of TG2 by withdrawal of tetracycline (tet−) from the cell culture system, the matrix-deposited TG2 is detectable after a 48-h induction (Fig. 10A). In the presence of 100 μg/ml P1 or P1s peptide as described under “Experimental Procedures.” Mean percentage value of the DMSO-treated attached and spread control cells on FN ± S.D. (control (CNTL)) was set at 100%.

FIGURE 7. Involvement of syndecan-4 and its downstream signaling molecule PKCα in the P1 peptide-mediated RGD-independent cell adhesion. A, requirement of syndecan-4 by the P1 peptide. Cell adhesion assays using HOB cells treated with syndecan-4 targeting or NS siRNA on FN in the presence of P1 or P1s control peptide (100 μg/ml) were performed as introduced under “Experimental Procedures.” Mean percentage value of the control attached and spreading cells on FN ± S.D. was set at 100%. B and C, importance of PKCα in P1 peptide-mediated cell adhesion. Cell adhesion and spreading of HOB cells on FN with RGD or RAD were performed using PKCα inhibitor Go6976 or the GK21 peptide, in the presence of 100 μg/ml P1 or P1s peptide as described under “Experimental Procedures.” Mean percentage value of the control attached and spread control cells on FN ± S.D. (control (CNTL)) was set at 100%.
Heparan Sulfate-related TG2 Localization

effect on TG2 expression and deposition (Fig. 10A). To clarify which member(s) of the MMP family members are involved in the TG2 deposition, specific MMP inhibitors against MMP-2/9, MMP-3, MMP-8 were used. As shown in Fig. 10B, the MMP-2/9 inhibitor reduced the matrix TG2 deposition by around 50%, which was not apparent in the other MMP inhib-
itor-treated cells. This suggests that the MMP-2/9 inhibitors, which are the major MMP members participating in wound healing and HS shedding (12, 30), are also involved in ECM TG2 deposition. Different treatments (12, 31) known to affect HS shedding were then used in the tet-inducible 3T3 cells. Fig. 10C shows that PMA, an enhancer for HS shedding (31), slightly increased the amount of matrix TG2, whereas the PKCa inhibitor Go6976 (an inhibitor for HS shedding (12)) blocked deposition of TG2 into the ECM. Heparin (at the concentration of 10 μg/ml), a known protector of MMP-dependent HS shedding (32), reduced the presence of TG2 in the matrix, whereas heparinase (which digests the cell surface HS chains) completely inhibited matrix TG2 deposition. In contrast, chondroitinase did not show any effect on matrix TG2 deposition (Fig. 10C). By using syndecan-4 siRNA targeting mouse syndecan-4, the expression of syndecan-4 was reduced to around 50% compared with the NS control (supplemental Fig. 2B), which led to a comparable reduction of ECM-deposited TG2 deposition (Fig. 11A). To confirm that, levels of syndecan-4 comparable with that found for TG2 are found in the ECM following treatment of cells with the same key modulators of syndecan shedding. The matrix of the treated cells was collected and immunopробed for the presence of syndecan-4. As shown for TG2, PMA, an enhancer of syndecan shedding, increased the amount of syndecan-4 present in the ECM, whereas the inhibitors of shedding PKCa inhibitor Go6976 and the inhibitor of MMP-2/9 reduced the amount of syndecan-4 present in the ECM (Fig. 11B).

To assess whether HS shedding may contribute to the increased TG2 activity found in the matrix following wounding (22, 33), fluorescence staining using FITC-cadaverine as a measure of TG2 in situ activity was used. After wounding the tet-inducible 3T3 cells in a scratch assay, cells were incubated for a 1-h period with FITC-cadaverine. As shown in Fig. 12, the PKCa inhibitor Go6976, MMP-2/9 inhibitor, heparin, and heparinase treatments and the peptidic TG2 irreversible inhibitor R281 all reduced the presence of in situ TG2 activity found at the edge of the wound when compared with the nontreated control. This reduction was not seen with either MMP-8 or chondroitinase treatment, thus confirming the importance of syndecan shedding in delivering increased TG2 activity to the matrix after wounding.

DISCUSSION

Identifying and subsequently being able to regulate the mechanism of TG2 translocation into the ECM provides a novel strategy for modulating the pathological functions of the enzyme. Earlier studies suggest that cell surface HSPGs might be important in the trafficking of TG2 onto the cell surface (29). However, evidence for an HS-binding site on TG2 has not been proven. We therefore identified the HS-binding site(s) within TG2 by using a combination of amino acid sequence analysis for known HS-binding motifs and by analysis of the available crystal structures of TG2. Two potential binding sites were identified (HS1 and HS2), which were docked with heparin-derived oligosaccharides consisting of an iduronic acid-2-sulfate-glucosamine-2,6-disulfate disaccharide, which is the most common repeating unit of heparin. Both the HS1 and HS2 sites docked well with all the oligosaccharides when TG2 was in its closed conformation, although the HS2 site was the preferred site. In comparison, neither site docked well when the open conformation of TG2 was used, although slightly better docking was observed with the HS1 site. As a consequence of these results, it was decided to subject both sites to the site-directed mutagenesis of the key surface-exposed basic residues. By using a heparin affinity chromatography column, we showed that the HS2 site, consisting of residues 2022FLKKNAGRDCC

FIGURE 8. Identification of the intracellular signaling molecules in P1 peptide-mediated signaling transduction. A–D, phosphorylation of FAK and ERK1/2 in P1 peptide-mediated signaling transduction. HOB cell adhesion was undertaken with the P1 or P1s peptides (100 μg/ml) as described under “Experimental Procedures.” P-FAK (p) and p-ERK1/2 (D) were detected by Western blotting. Membranes were reprobed to detect the total FAK or ERK1/2, whereas α-tubulin was used as the standard for equal loading. Relative amounts of p-397 FAK (B) and p-ERK1/2 (D) compared with total FAK or ERK1/2 were measured by densitometry and normalized to tubulin and then represented as a percentage of that calculated for FN in the presence of RAD. Values are shown as the mean ± S.D. from three separate experiments. E–G, affinity of TG2 and the binding site mutant (D94A,D97A), the N-terminal deletion product of TG2 (Δ1–15), and the HS2 mutant (K205A,R209A) for binding to FN. E, clarified cell lysates (60 μg protein) from transiently transfected cells were added to FN-coated plates and the TG2 proteins detected as described under “Experimental Procedures.” Data show the mean values ± S.D. from three experiments. F and G, human recombinant TG2 (2 μg/ml) was added to microtiter plates previously coated with FN or the N-terminal 70- or 45-kDa FN fragments. Prior to addition of the TG2, wells were blocked, washed, and then incubated with either the P2 peptide (F) or the scrambled peptide (G, P2s) as described under “Experimental Procedures.” Data show the mean values ± S.D. from three experiments.
eluted enzyme showed full TG activity that could be inhibited by GTP indicating that the HS2 mutant is still able to bind to this nucleotide (supplemental Fig. 1). Hence, the affinity of the Lys-205 and Arg-209 mutant for heparin is significantly lower than that of either GTP-bound or R281-reacted wild type TG2, and it is likely that little binding would occur at physiological ionic strength. In support of this, NIH 3T3 cells expressing the HS2 mutant showed much reduced amounts of the HS2 mutant on the cell surface, and it was found to be barely present in the ECM of the NIH 3T3 cells. In view of a very recent paper (35) indicating that the N-terminal \( \beta \)-sandwich domain of TG2 is essential for TG2 localization.

**FIGURE 9.** A and B, detection of extracellular TG2 in NIH 3T3 cells transfected with wild type TG2 and TG2 mutants. C and D, presence of extracellular TG2 in CHO-K1 cells and the HS-deficient CHO-K1 derivative pgsD-677. A, cell surface TG2 and mutants were detected by Western blotting after treatment of cells with sulfo-NHS-LC-biotin. Densitometry values (mean value from two experiments) for the different TGs are from Western blots normalized for loading and TG2 expression levels as described in Fig. 2. B, for TG2 and TG2 mutants in the ECM, transfected NIH 3T3 cells were grown for 24 h post-transfection in full DMEM and for a further 24 h in 1% serum. TG2 antigen in the ECM was detected by Western blotting with Cub7402. The bars in the histogram represents the mean values for matrix TG2 and its mutants as a percentage of the WT TG2 after normalizing for the relative expression of TG2 in the cell lysates. Data are from two independent experiments as described under “Experimental Procedures.” C, cell surface TG2 in CHO cells and its HS-deficient CHO-K1 derivative were detected after treatment of cells with sulfo-NHS-LC-biotin as described above. D, relative amounts of cell surface TG2 expressed as a percentage of the WT CHO cells measured by densitometry of the Western blots (mean values from two experiments) after normalizing to the expression of TG2 as described in A above.
secretion to the cell surface in kidney OK epithelial cells, we also measured the amount of the cell surface HS2 mutant and matrix-associated HS2 in these same cells (supplemental Fig. 3). Again, despite a good expression level of this mutant, there was little HS2 mutant present at the cell surface, although the enzyme was absent in the matrix, thus agreeing with our data for the HS2 mutant-transfected NIH 3T3 cells. Support for the importance of HS in the trafficking of TG2 to the cell surface was obtained using TG2-expressing HS mutant CHO cells (which are unable to synthesize HS but are still able to maintain the presence of the core proteins of the proteoglycans in the cell membrane (36)). A multiple alignment of TG peptide sequences, including TG2 from different species and human TG isoforms (supplemental Fig. 4), shows that the high affinity heparin binding domain is conserved among TG2 enzymes but is absent from other iso-

FIGURE 10. HS shedding and its effects on the trafficking of TG2 into the ECM. General MMP inhibitor GM6001 (GM) (A) or specific MMP inhibitors for MMP-2/9, MMP-3, and MMP-8 (10 μM) (B) were used to treat the TG2 induced (tet +) and noninduced (tet −) 3T3 cells. Cell lysates and matrix fractions after 48 h were Western-blotted to detect TG2 antigen in the fractions. The bars in the histogram represent the mean values for matrix TG2 represented as a percentage of the nontreated control cells after normalizing for the relative expression of TG2 in the cell lysates. Data are from two independent experiments as described under “Experimental Procedures.” C, effect of HS on TG2 deposition. Cell lysates and matrix fractions from tet-inducible 3T3 cells induced (tet −) or not induced (tet +) for TG2 were incubated with PMA (50 nM), PKCα inhibitor Go6976 (Go) (5 μM), heparin (HN) (10 μg/ml), heparinase (HNase), or chondroitinase (CD) (15 milliunits/ml) and were used in Western blotting to detect TG2 and levels (mean of two experiments) expressed as a percentage of the nontreated control, calculated as described in B above. CNTL, control.
forms with residues corresponding to human TG2 positions 202, 205, 213, and 222 conserved as basic residues among all the TG2 sequences analyzed.

Syndecan shedding is found in many physiological and pathological situations, where up-regulation of TG2 expression also occurs, e.g. during wound healing, in cancer migration, and in bone differentiation and mineralization (9, 37). To study the involvement of syndecan shedding in the trafficking of TG2 into the matrix, we used the well characterized stably transfected Swiss 3T3 cells, in which TG2 expression, which results in easily detectable deposition of TG2 into the matrix, is under the control of the tet-inducible promoter (13). Using both general and specific inhibitors of MMPs in these cells, we first demonstrated that MMP-2 and MMP-9, the two major MMPs involved in HS shedding (9, 38, 39), are required for the deposition of TG2 into the ECM. By protecting or inhibiting the cell surface HS from shedding by treating cells with heparin or the PKCα inhibitor Go6976, TG2 deposition into the matrix was reduced, whereas PMA, an accelerator for HS shedding, increased the amount of TG2 deposited. Moreover, removal of the HS chains in the TG2-transfected tet-inducible 3T3 cells completely blocked the deposition of TG2 into the matrix. Parallel studies undertaken to measure the amount of syndecan-4 present in the cell matrix after treatment of cells with PMA, PKCα inhibitor Go6976, and the inhibitor of MMP2/9 showed comparable levels of syndecan-4 in the matrix to that found for TG2 when treated with the same key modulators of syndecan shedding. Importantly, knocking down the expression of syndecan-4 in these 3T3 cells by siRNA significantly reduced the trafficking of TG2 into the ECM, further confirming the importance of syndecan-4 shedding in matrix TG2 deposition. It is reported that wounding of cells or tissues results in a rapid but transient increase in the TG2 activity found in the matrix surrounding the wound area (22, 33). We therefore looked to see if syndecan shedding may also account for this rapid increase in TG2 activity found around the wound area. Our data showed that the increased TG2 activity found around the wound area of TG2-inducible Swiss 3T3 cells could be either partially or totally blocked by agents known to affect syndecan shedding, thus supporting our claims for the importance of syndecan shedding in the trafficking of TG2 into the matrix following its increased expression in cells. Importantly, this finding also fits with the suggested role of TG2 as a stress/wound-response enzyme (40).

It has been suggested that the mechanism of TG2 secretion from cells may be dependent on conformation and its transition between the closed and open forms (3, 41). Using this informa-
tion and the data obtained for binding of the TG2 mutants to heparin and the importance of syndecan shedding in the trafficking of TG2 into the matrix, one might envisage a mechanism whereby TG2 is first externalized in its closed GTP-bound conformation. It is then retained at the cell surface by its high affinity binding to HS at physiological ionic strength. Immediate binding to the negatively charged cell surface HS may also be responsible for maintaining the closed conformation of TG2 by sequestration of surrounding Ca$^{2+}$ until the enzyme is translocated into the ECM via syndecan shedding. Once shed and exposed to Ca$^{2+}$, TG2 will adopt an open conformation, and the affinity for HS is sig-

FIGURE 12. Effect of HS in wound-induced TG2 deposition. In situ activity of TG2 in TG2-transfected tet-inducible (tet$^-$) 3T3 cells seeded into 8-well chambers detected by incorporation of FITC-cadaverine after different treatments was visualized using fluorescence microscopy as described under “Experimental Procedures.” The dotted line marks the edge between the cell area (marked C in the image) and the wound areas (marked W). CNTL, control.
nificantly reduced, whereas its affinity for FN is increased (supplemental Fig. 5) enabling it to bind to and cross-link its matrix substrate proteins such as FN (Fig. 13) (42). Subsequent oxidation or nitrosylation of the matrix-bound enzyme then further modulates its transamidating activity (43, 44), such that the enzyme may then act as a novel FN-bound cell adhesion protein, either through its interaction with cell surface syndecan-4 or β integrins (18, 19).

Strong support for the role of TG2 as a novel cell adhesion protein working via interaction with syndecan-4 comes from the finding that the peptide representing the proposed binding pocket on TG2 for HS (P1 peptide) acted in a comparable manner to TG2 in compensating RGD-induced loss of cell adhesion when cells were plated onto FN, using a previously described osteoblast model (5). At concentrations as low as 5–10 μg/ml, compensation of cell adhesion was noted. Moreover, when an α5β1 integrin blocking antibody was used to induce loss of cell adhesion, the P1 peptide, like the full TG2 protein, was able to compensate for the loss of adhesion, indicating the potency of this interaction and its ability to mimic TG2 (27). Fluorescence co-staining of F-actin and vinculin further demonstrated the ability of the P1 peptide to restore loss of actin cytoskeleton organization and focal adhesion formation induced by the RGD peptide treatment. Importantly, this compensatory effect of the peptide could be abrogated by pretreatment of cells with heparinase or treatment of cells with syndecan-4 siRNA, indicating that P1 peptide like TG2 binds to cell surface syndecan-4. These data and the finding that the binding of the P1 peptide to syndecan-4 stimulates activation of PKCα, FAK at Tyr-397, and subsequent activation of ERK1/2 confirm the interaction of TG2 with cell surface syndecan-4 (57). One explanation for this is that the enzyme may then act as a novel FN-bound cell adhesion protein, either through its interaction with cell surface syndecan-4 or β integrins (18, 19).

In summary, our findings with the HS-binding peptide and the discovery of a novel TG2 HS-binding site confirm the importance of matrix-bound TG2-syndecan-4 interactions in cell adhesion. They also provide a new mechanism for the rapid translocation of TG2 into the ECM involving syndecan shedding. Moreover, the demonstration of the potency of the HS-binding peptide in modulating cell adhesion may also have future potential applications in regulating cell behavior in TG2-mediated pathologies.

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