Alteration of Galectin-3 in Tears of Patients with Dry Eye Disease

Yuichi Uchino, Jerome Mauris, Ashley M. Woodward, Julia Dieckow, Francisco Amparo, Reza Dana, Flavio Mantelli, and Pablo Argüeso

Schepens Eye Research Institute and Massachusetts Eye and Ear, Department of Ophthalmology, Harvard Medical School, Boston, Massachusetts (Y.U., J.M., A.M.W., J.D., F.A., R.D., P.A.); and the Campus Bio Medico University of Rome, Rome, Italy (F.M.)

Abstract

Purpose—To investigate the expression, release, and proteolytic degradation of galectin-3 in patients with dry eye disease.

Design—Observational case series with a comparison group.

Methods—Tear washes and conjunctival impression cytology specimens were collected through standard procedures from 16 patients with dry eye and 11 age-matched healthy subjects. Galectin-3 content in tears was analyzed by quantitative Western blot, using recombinant galectin-3 protein to generate a calibration curve. The relative expression of galectin-3 and matrix metalloproteinase 9 (MMP9) was evaluated by quantitative polymerase chain reaction. The cleavage of galectin-3 was studied in vitro using activated recombinant MMP9 and protease inhibitors.

Results—The concentration of galectin-3 protein in tears, but not galectin-3 expression in conjunctival epithelium, was significantly higher in tears of patients with dry eye (0.38 ng/μg total protein, range 0.04-1.36) compared to healthy subjects (0.12 ng/μg total protein, range 0.00-0.41) (P < .01). By Western blot, an intact (∼28.0 kDa) galectin-3 band was identified in tear samples.
from healthy subjects, whereas 50% of the dry eye samples were characterized by the additional presence of a partially degraded form (∼25.4 kDa). In our experiments, elevated expression of MMP9 in dry eye subjects correlated with the ability of active MMP9 to cleave galectin-3 from recombinant origin. Interestingly, cleavage of endogenous galectin-3 in tear samples was impaired using a broad-spectrum proteinase inhibitor cocktail, but not the pan-specific MMP inhibitor GM6001, suggesting the presence of proteases other than MMPs in promoting galectin-3 degradation in dry eye.

Conclusions—Our results indicate that release of cellular galectin-3 into tears is associated with epithelial dysfunction in dry eye, and that galectin-3 proteolytic cleavage may contribute to impaired ocular surface barrier function.

Keywords
galectin-3; tears; dry eye disease; ocular surface barrier function; proteolytic activity

Introduction

Disruption of epithelial barrier function is associated with a wide range of disorders that includes dry eye, a disease affecting millions of people worldwide and whose pharmacological treatment remains unsatisfactory. While the intercellular junctions that connect individual cell membranes on mucosal surfaces have been clearly defined as components of the paracellular barrier, research performed during the last decade has redefined and highlighted the contribution of apical cell membranes as an additional layer of protection—the transcellular or glycocalyx barrier. Indeed, both molecular and structural changes of the glycocalyx in apical cells within the stratified squamous epithelium have been observed in drying diseases, and appear to correlate with loss of barrier function at the ocular surface.

Transmembrane mucins (MUCs), such as MUCs 1, 4 and 16, are an essential component of the glycocalyx barrier in mucosal epithelia. They have single membrane-spanning regions with large extracellular glycosylated domains that form rod-like structures extending 200–500 nm above the plasma membrane, far above other glycoconjugates in the glycocalyx. Extensive research has shown that O-glycans, a form of glycosylation resulting from the addition of N-acetylgalactosamine to serine or threonine residues in the mucin ectodomain, are critical to preventing epithelial damage and infection of mucosal surfaces. At the ocular surface, transmembrane mucins contribute to provide boundary lubrication and to prevent the penetrance of extracellular molecules and pathogenic microorganisms onto the epithelia.

It is now clear that transmembrane mucins and their associated glycans promote the integrity of the epithelial barrier at the ocular surface by interacting on the apical glycocalyx with galectin-3. Galectin-3 is a multivalent β-galactoside-binding lectin known to promote formation of plasma membrane lattices with unique functions, concentrating specific glycoproteins within the lattice while excluding other molecules. Similarly to other members of the galectin family, galectin-3 is defined by the presence of an evolutionarily conserved carbohydrate-binding domain that mediates receptor binding. Critical to the
multimerizing function of galectin-3 is the presence of an N-terminal domain that allows self-association through intermolecular interactions when bound to a multivalent ligand.\textsuperscript{18} Using site-directed mutagenesis, we have recently shown that maintenance of ocular surface glycocalyx integrity \textit{in vitro} requires the N-terminal multimerization domain of galectin-3.\textsuperscript{14} To date, however, much remains unknown about the synthesis and structural integrity of galectin-3 in patients with altered ocular surface barrier function. In the present study, we investigated whether alteration of galectin-3 is associated with dry eye, by evaluating the expression and release of galectin-3 onto the ocular surface, and examining whether galectin-3 undergoes proteolytic degradation in tears.

Methods

Study population

This observational case series study was conducted in compliance with good clinical practice, Institutional Review Board approval (Campus Bio Medico University of Rome, IRB # 07/06.PARComEtCBM; Massachusetts Eye and Ear Infirmary, IRB #12-1591-1), informed-consent regulations and the tenets of the Declaration of Helsinki. A total of sixteen patients (10 women, 6 men) with dry eye disease were recruited from Fondazione GB Bietti and Massachusetts Eye and Ear Infirmary, with a mean age of 56.9±11.0 years (range, 35–84 years). Within this population, twelve patients were aqueous-deficient without meibomian gland disease (MGD), three were aqueous-deficient with MGD, and one was aqueous-sufficient with MGD (Supplementary Table). Within the MGD patients, one had Sjögren’s syndrome and two had graft versus host disease (GVHD). None of the patients had chalasis. The subjects completed an IRB approved questionnaire regarding history of ocular allergies, disease, surgery, contact lens wear, current medications, type and frequency of symptoms of dry eye and dry mouth, and the use of dry eye therapy. Inclusion criteria were symptoms of dry eye and at least one of the following objective signs: Schirmer I <10 mm at 5 minutes, decreased tear breakup time (TBUT) of ≤5 seconds, and positive vital dye staining of the conjunctiva or cornea. The degree of positive staining used for the diagnosis of dry eye corresponds to scores higher than 1 in the Oxford Scheme, or higher than 3 in the Bijsterveld score. Patients with a history of ocular surgery, topical anti-glaucoma or anti-inflammatory therapy, or eye allergies were excluded. Eleven age- and sex-matched healthy subjects (7 women, 4 men) without dry eye symptoms, eye diseases, or history of eye surgery or contact lens wear were used as a control group. The mean age of the control group was 55.6±11.5 years (range, 34–72 years).

Sample collection

Tear fluid and conjunctival epithelium from healthy subjects and patients with dry eye disease were collected to determine galectin-3 protein levels, and galectin-3 and matrix metalloproteinase 9 (MMP9) messenger ribonucleic acid (mRNA) expression, respectively. Tear fluid was collected from unanesthetized eyes by pipetting 60 μL sterile saline (AddiPak\textsuperscript{®} 0.90% sodium chloride sterile solution) into the inferior fornix using a sterile transfer pipet, followed by movement of the eyes to mix the tear fluid content as described.\textsuperscript{10} The volumes acquired ranged between 14.0 and 72.6 μL (average 42.9 μL). Then, a drop of topical anesthetic (proparacaine hydrochloride, 0.5%, Alcaine\textsuperscript{®}; Alcon Inc.;
Ft. Worth, TX) was applied to the eye. A sterile disc of nitrocellulose filter paper, 10 mm in diameter, was placed on the temporal bulbar conjunctiva, gentle pressure was applied to the disc for 15 seconds, and the disc was then carefully transferred into an Eppendorf tube containing TRIzol reagent (Gibco; Grand Island, NY). All samples were promptly frozen at -80°C until the time of analyses.

Individual tear samples were centrifuged for 30 minutes at 18,000 × g at 4°C using a microcentrifuge (centrifuge 5424R, Eppendorf, Hamburg, Germany). Protein concentration was determined using the Micro bicinchoninic acid (BCA) Protein Assay Kit (Thermo Scientific, Waltham, MA), according to the manufacturer’s protocol.

Production of Recombinant Human Galectin-3

Recombinant human galectin-3 (rhGal-3) was expressed as previously reported. Briefly, heterologous protein expression was induced by addition of 0.3 mM of isopropyl β-D-1-thiogalactopyranoside (IPTG) to Rosetta E. coli clones carrying a rGal-3 vector, followed by incubation overnight at 15°C with shaking. The rhGal-3 was purified from lysates by affinity chromatography using lactosyl sepharose. To eliminate contaminating bacterial endotoxins, rGal-3 was further purified by polymyxinB affinity chromatography. The absence of lipopolysaccharide was confirmed using ToxinSensor™ Chromogenic LAL Endotoxin Assay Kit (GenSript, Piscataway, NJ, USA). Protein solutions were concentrated by filtration (VIVASPIN, Littleton, MA, USA), dialyzed against phosphate-buffered saline (PBS) buffer containing 10% glycerol and stored at -20°C. Protein concentration was determined using the BCA Protein Assay Kit (Thermo Scientific).

Immunoblotting

Tear samples containing 1-10 μg of total protein were separated by electrophoresis on 10% sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) gels and transferred onto nitrocellulose membranes (BioRad Laboratories). Membranes were then blocked with 5% nonfat dry milk in Tris buffered saline with Tween (TBST) for 2 hours at room temperature, followed by incubation overnight at 4°C with the following primary antibodies diluted in 5% nonfat dry milk in TBST: anti-galectin-3 (H160; 1:2-3,000; Santa Cruz Biotechnology; Dallas, TX), anti-MMP9 (ab38898; 1:2,000; Abeam; Cambridge, MA). Following incubation with the corresponding peroxidase-conjugated secondary antibody (1:2,000; Santa Cruz), positive binding was visualized using the SuperSignal West Pico Chemiluminescent Substrate (Thermo Scientific) on HyBlot CL autoradiography film (Denville Scientific; Inc., Metuchen, NJ). When no band intensity was detected using the West Pico substrate, membranes were subsequently incubated with an enhanced and more sensitive substrate (SuperSignal West Femto Maximum Sensitivity Substrate; Thermo Scientific), which enables detection of low femtogram amounts of protein. Band intensities were quantified by densitometry (ImageJ software; National Institutes of Health; Bethesda, MD; in the public domain available at http://rsb.info.nih.gov/nihimage).

Quantification of protein concentrations in tear fluid by immunoblotting was performed as previously described. Galectin-3 content (ng/μg total protein) in tears was determined
using a standard curve based on signal intensity generated with different concentrations of rhGal-3, and two internal controls of rhGal-3 per gel when analyzing experimental samples.

**RNA Isolation, cDNA Synthesis and Quantitative Polymerase Chain Reaction**

Total RNA was extracted from conjunctival impression cytology specimens using an extraction reagent (TRIzol; Life Technologies) according to the manufacturer's protocol. The integrity and concentration of the total RNA were determined using a NanoDrop 2000 Spectrophotometer (Thermo Scientific). One microgram of total RNA was treated with amplification-grade DNase I (Life Technologies) to eliminate residual genomic DNA and reverse transcribed into cDNA using random hexamer primers and reverse transcriptase (iScript; Bio-Rad; Hercules, CA) according to the manufacturer's protocol.

Detection of gene expression in conjunctival impression cytology specimens was performed by quantitative polymerase chain reaction (qPCR) using PrimePCR™ primers for galectin-3 (Unique Assay ID: qHsaCID0008552) and MMP9 (Unique Assay ID: qHsaCID0011597). The qPCR reactions were carried out in triplicates in a 20 μL reaction volume using 1 μL of cDNA, 1 μL primers and the SYBR® Fast master mix (KARA Biosystems; Wilmington, MA) in a Mastercycler® ep realplex² thermal cycler (Eppendorf; Hauppauge, NY). The following parameters were used: 2 minutes at 95°C, followed by 40 cycles of 5 seconds at 95°C and 30 seconds at 60°C. All samples were normalized using glyceraldehyde-3-phosphate dehydrogenase (GAPDH) housekeeping gene expression (PrimePCR™ GAPDH primers; Bio-Rad). The comparative C_T method was used for relative quantitation of the number of galectin-3 and MMP9 transcripts, selecting the relative mRNA levels in samples from healthy subjects as the calibrator.

**Protease Cleavage Assays**

For recombinant galectin-3 protease cleavage assay, 10 ng of rhGal-3 were incubated with increasing concentrations of activated recombinant human MMP9 (rhMMP9, Calbiochem, San Diego, CA) in Dulbecco's phosphate-buffered saline supplemented with magnesium and calcium (Thermo Scientific, Waltham, MA) at 37 °C for various time periods as indicated. Reactions were stopped by the addition of Laemmli buffer and then subjected to boiling for 5 minutes. For endogenous galectin-3 protease cleavage assay, tear samples (3-5 μg total protein) from three patients with dry eye disease were incubated in PBS buffer at 37 °C for 6 hours. In additional experiments, samples were supplemented with the pan-specific MMP inhibitor GM6001 (2 μM for rhGal-3; 40 μM for human tears) or a broad-spectrum proteinase inhibitor cocktail (complete™ EDTA free; Roche Applied Science; Mannheim, Germany). The reaction products were analyzed by electrophoresis on 10% SDS-PAGE gels and visualized by immunoblotting.

**Statistical Analyses**

Statistical significance between normal and dry eye groups was evaluated with the Mann-Whitney U test for nonparametric data, using a p value of 0.05 as the threshold for significance. Correlations between TBUT and galectin-3 protein concentration or gene expression were determined using the Spearman rank correlation test. All statistical analyses were performed using InStat3 software (GraphPad Software, La Jolla, CA).
Results

Galectin-3 Levels Increase in Tears of Dry Eye Patients

Altered levels of galectin-3 in biological fluids and sera have been associated with the pathophysiology of a number of diseases, such as asthma, heart failure, and cancer. To study a potential role of galectin-3 in dry eye disease, we developed a defined immunoblot method to quantify the levels of endogenous galectin-3 in human tear samples. For this purpose, a standard curve from a two-fold serial dilution series of rhGal-3 was produced to determine the linear range of detection for the galectin-3 antibody. Densitometric analysis of the protein bands revealed linear responses between 0.6 and 18 ng of rhGal-3 (Figure 1, upper left). Next, we obtained the relative intensities of endogenous galectin-3 in human tears, normalized to the amount of total protein as previously described. We found a significant increase in galectin-3 levels in tears of dry eye patients. As shown in Figure 1 (upper right) and Supplementary Table, the mean concentration of galectin-3 in control eyes was 0.12±0.04 ng/μg total protein (range 0.00-0.41), whereas in dry eye it was 0.38±0.08 ng/μg total protein (range 0.04-1.36) (P < .01).

To determine whether enhanced expression of galectin-3 mRNA at the ocular surface was responsible for the increase of galectin-3 protein in tears, we performed qPCR on samples of human conjunctival epithelium obtained by impression cytology from the same population groups. Interestingly, no significant differences in expression were observed between the healthy and dry eye groups (Figure 1, lower left). These results were consistent with data from a public database of microarray gene expression (http://www.functionalglycomics.org), showing no statistical differences in signal intensity for galectin-3 in conjunctiva of normal subjects and patients with dry eye disease (Figure 1, lower right).

Galectin-3 Release into Tears Correlates with Tear Film Instability

A key step in dry eye disease is the occurrence of an unstable tear film that eventually breaks up when in contact with an altered ocular surface epithelium. To determine a potential association between the amount of galectin-3 in tears and the occurrence of an unstable tear film, we performed a Spearman rank correlation test. As shown in Figure 2 (upper panel), we found a significant correlation between galectin-3 protein concentration in tears and tear instability as measured by tear film breakup time (R=-0.4988; P < .01). In contrast, there was no correlation between galectin-3 gene expression in human conjunctival epithelium and tear film breakup time (Figure 2, lower panel).

Galectin-3 is Partially Degraded in Tears of Dry Eye Patients

Proteolytic degradation of galectin-3 is thought to impair the ability of the molecule to multimerize in the presence of binding partners, and to abrogate biological functions that depend on the ability of the lectin to form lattices. The human gelatinases MMP2 and MMP9 have been shown to induce cleavage of galectin-3 at the Ala62-Tyr63 bond in the N-terminus, resulting in a 22 kDa cleavage product lacking the multimerization domain known as galectin-3C (Figure 3, upper panel). In our experiments, analysis of galectin-3 in human tears samples by immunoblot revealed the presence of a major galectin-3 band with a molecular weight of 28.0 kDa (Figure 3, lower left). Interestingly, we also detected the
presence of a partially degraded form with a molecular weight of 25.4 kDa. Quantification of the ratio between galectin-3C and galectin-3 revealed a significant increase in tears of dry eye patients compared to tears of healthy subjects (Figure 3, lower right; Supplementary Table). Further, whereas higher values of galectin-3 were not associated with a particular etiology, the ratio between galectin-3C and galectin-3 in tears of patients with Sjögren's syndrome and GVHD was significantly higher than in other patients (data not shown).

Degradation of Galectin-3 in Human Tears Involves Protease Activity

Previous studies have shown that MMP9 activity is significantly higher in patients with dry eye disease.\textsuperscript{28, 29} Therefore, to further explore the mechanism by which galectin-3 is degraded in human tears, we analyzed the levels of MMP9 expression in conjunctival impression cytology samples collected from our study population. Consistent with previous data, we found a significantly higher expression of MMP9 mRNA in samples from dry eye patients compared to normal subjects (Figure 4, upper panel). In subsequent experiments, we tested whether active human rhMMP9 would cleave rhGal-3. Our results showed that active MMP9 promotes galectin-3 cleavage in a concentration- and time-dependent manner, producing a 25-kDa cleavage product (Figure 4, middle and lower panels). As expected, the pan-specific MMP inhibitor GM6001 attenuated the degradation of full-length galectin-3 in control experiments.

Based on data using recombinant protein, we next hypothesized that cleavage of endogenous galectin-3 at the ocular surface would depend on the activity of MMP9 in human tears. To test this possibility, we developed an \textit{in vitro} galectin-3 degradation assay with human tears (Figure 5). In this assay, tear samples were incubated at 37 °C for 6 hours to promote degradation of endogenous galectin-3. Analyses of tear samples from three dry eye patients revealed that, under these conditions, endogenous full-length galectin-3 was partially or completely degraded. Moreover, we also observed the appearance of galectin-3C in two of the three samples following incubation at 37 °C for 6 hours. Unexpectedly, we found that GM6001 could not prevent the proteolytic degradation of endogenous galectin-3 in human tears, in contrast to \textit{in vitro} data using rhMMP9 and rhGal-3 (Figure 4, middle and lower panels). However, cleavage of endogenous galectin-3 could be partially prevented in samples from two patients by using the broad-spectrum proteinase inhibitor cocktail cOmplete\textsuperscript{TM}, suggesting the presence of proteases other than MMPs in promoting galectin-3 degradation at the ocular surface.

Discussion

The traditional model of mucosal protection portrays transmembrane mucins as passive molecules on the surface of epithelial cells. They emanate from apical plasma membranes, well above other glycoproteins, acting primarily as a physical barrier that prevents the influx of toxic agents and pathogens. It is now clear that transmembrane mucins contribute via glycosidic residues in an active manner to promote barrier function through interactions with galectin-3 on the ocular surface epithelial glycocalyx.\textsuperscript{13, 14} Here, we report that release of cellular galectin-3 into tears is associated with epithelial dysfunction in patients with dry eye.
Galectin-3 is a carbohydrate-binding protein expressed in multiple epithelial tissues where it controls biological processes such as cell–cell and cell–extracellular matrix adhesion, cell growth and differentiation, cell-cycle, signaling, and apoptosis. Alteration in galectin-3 expression has been predominantly studied in cancer and inflammatory disease, but is commonly observed in many other pathological states. At the ocular surface, Hrdlickova-Cela et al. used non-quantitative immunoblotting to report that galectin-3 was absent in tear fluid from four normal subjects, but present in tears of a limited cohort (mostly one subject per condition) of patients with diverse ocular surface disorders including bullous keratopathy, alkali burn, chronic blepharitis and adenoviral conjunctivitis. Our data showing increased levels of galectin-3 in tears of patients with dry eye disease are consistent with these findings reporting galectin-3 immunoreactivity in tears of patients with ocular surface disease. However, contrary to the results of Hrdlickova-Cela et al., we could also detect the presence of galectin-3 in normal tears, which could be explained by the analysis of a larger pool of subjects in our study population, and the use of a more sensitive detection assay for galectin-3. In our analyses, the severity of the disease did not correlate with the amount of galectin-3 or the cleavage of galectin-3 in tears (data not shown). It is possible to speculate that activation of compensatory mechanisms, such as increased glycosylation, in severe dry eye could lead to partial retention of galectin-3 on cell surfaces. Increased expression of glycosyltransferases has been observed before ocular surface keratinization in patients with ocular cicatricial pemphigoid. Further, the protease responsible for galectin-3 degradation in tears has yet to be identified, and its activity correlated with the severity of disease.

Interestingly, we found that the increase in galectin-3 levels in tear samples of dry eye patients did not correlate with an altered expression of galectin-3 mRNA in conjunctival impression cytology samples. We hypothesize that the discrepancy between galectin-3 protein and mRNA levels at the ocular surface could be explained by the disruption of the epithelial glycocalyx in dry eye disease. Mucins, like many other glycoproteins on cell surfaces, are subject to changes in glycosylation during pathological conditions that include inflammation. Further, it has been established that mucin glycosylation is altered in dry eye disease. Therefore, alterations on transmembrane mucin glycosylation in corneal and conjunctival epithelia could lead to loss of galectin-3 binding affinity and release of cellular galectin-3 into the tear film. It is also possible to speculate that galectin-3 in serum could diffuse into tears as a consequence of increased vascular and conjunctival epithelial permeability in dry eye, and that other cell types, such as those of the lacrimal gland or inflammatory cells present within the epithelium or tear fluid, could likewise contribute to secrete galectin-3 in these patients. In our assays, the tear fluid was subjected to low rate centrifugation to reduce the possibility of cellular contamination. However, it will be exciting to determine in future studies the individual contribution of increased permeability and the role of different cell types to the content of galectin-3 at the ocular surface in dry eye.

Multimerization is a common feature of extracellular galectin-3 that leads to ligand cross-linking and the formation of lattice-like structures on cell surfaces. Evidence suggests that degradation of galectin-3 results in the formation of a dominant-negative inhibitor, galectin-3C, that lacks the ability to multimerize and competes with extracellular full-length...
Truncated forms of galectin-3 have been detected in pathological conditions and have been shown to inhibit a variety of processes such as tumor growth and wound healing. Importantly, galectin-3C has a much higher binding affinity for the receptors compared to the full-length protein. In our experiments, we found that the ratio of galectin-3C to galectin-3 increased in tears of patients with dry eye. Although we do not know the biological consequences of an increased level of galectin-3C in dry eye disease, we speculate that it might play an important role in the impairment of ocular surface glycocalyx barrier function by interfering with the formation of transmembrane mucin lattices on apical cells within the stratified squamous epithelium. This hypothesis is supported by in vitro data indicating that treatment with a dominant-negative inhibitor of galectin-3 polymerization lacking the N-terminal domain, but not full-length galectin-3, prevented the recovery of barrier function in stratified human corneal epithelial cells.

Previous reports have shown that the collagen-like domain of galectin-3 is susceptible to rapid and efficient cleavage by MMP2 and MMP9. Consistent with previous data, we found a significantly higher expression of MMP9 mRNA in samples from dry eye patients that correlated with the ability of active MMP9 to cleave galectin-3 from recombinant origin in a dose- and time-dependent manner. Interestingly, we also found that cleavage of endogenous galectin-3 in tear samples could be impaired by using a broad-spectrum proteinase inhibitor cocktail, but not the pan-specific MMP inhibitor GM6001, suggesting the presence of proteases other than MMPs in promoting galectin-3 degradation in tears of patients with dry eye. As with human tears, gelatinolytic activity and MMP9 immunoreactivity have been identified in human seminal plasma, however, endogenous metalloproteases purified from seminal plasma have failed to cleave galectin-3. It is now clear that other proteases, such as prostate specific antigen, MMP13, neutrophil elastase, and leishmanolysin, can also degrade galectin-3. Exploring whether these or other candidates are responsible for the proteolytic degradation of galectin-3 at the ocular surface could bring novel insights into the mechanisms of epithelial dysfunction in dry eye disease. Further, data from these studies indicate that galectin-3 could potentially be used as a biomarker for the evaluation and management of patients with ocular surface pathology. Assays to measure galectin-3 are available and have been found to aid in the prognosis of human disease.

**Supplementary Material**

Refer to Web version on PubMed Central for supplementary material.

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References


Biographies

Dr. Yuichi Uchino graduated from Keio University School of Medicine, Tokyo, Japan in 2001. Afterward, he worked as a resident in Department of Ophthalmology, Keio University School of Medicine, Tokyo, Japan. From 2012, he has been a postdoctoral research fellow at the Schepens Eye Research Institute, Harvard Medical School, Boston, Massachusetts. Dr. Uchino’s primary research interests are the etiology of dry eye disease and the mechanism of glycocalyx barrier function.

Dr. Pablo Argüeso received his Ph.D. (Biochemistry) in Spain in 1997. He is currently an Associate Professor of Ophthalmology at Harvard Medical School, and Associate Scientist at The Schepens Eye Research Institute/Massachusetts Eye and Ear. His research focuses on one of the last frontiers of molecular biology—glycobiology—analyzing the sugars that coat the surface of the eye and deciphering their role in protecting the eye against dehydration and pathogen invasion.
Galectin-3 levels increase in tears of dry eye patients. (Upper left) Standard curve from a two-fold serial dilution series of recombinant human galectin-3 (rhGal-3). (Upper right) Quantification of galectin-3 concentrations in tear fluid from 11 normal subjects (11 eyes) and 16 dry eye patients (20 eyes) by immunoblotting. The relative intensities of endogenous galectin-3 in human tears were normalized to the amount of total protein. Data for individual samples are presented, with mean value indicated by horizontal bar for each group. (Lower left) Quantification of galectin-3 mRNA in conjunctival impression cytology samples from 10 normal subjects (10 eyes) and 15 dry eye patients (17 eyes) by qPCR. Data for individual samples are presented, with mean value indicated by horizontal bar for each group. (Lower right) Microarray analysis of impression cytology samples showing galectin-3 mRNA levels in human conjunctiva of normal and dry eye patients. Detailed data on glycogene expression can be found at http://www.functionalglycomics.org/glycomics/publicdata/microarray.jsp; Accession # MAEXP_272_042605. **P < .01; n.s., not significant. Error bars indicate the standard deviation.
Figure 2.
Galectin-3 release into tears correlates with tear film instability. Scattergrams illustrating the correlations between the levels of galectin-3 in tears (upper panel) or galectin-3 mRNA (lower panel) from the same individuals and TBUT using the Spearman rank correlation test. Significance was achieved when correlating galectin-3 in tears (R = -0.4988; P < .01),
but not galectin-3 mRNA (R = -0.09697, P = 0.6304), with TBUT. Black plots, dry eye patients; white plots, normal subjects.
Galectin-3 is partially degraded in tears of dry eye patients. (Upper panel) Schematic diagram illustrating the MMP-mediated cleavage of galectin-3 at the Ala_62-Tyr_63 bond in the N-terminus to produce a 22 kDa cleavage product lacking the multimerization domain (galectin-3C). (Lower left) By immunoblot, a partially degraded form of galectin-3 with a molecular weight of 25.4 kDa is detected in tears of patients with dry eye. (Lower right) Densitometric analyses revealed a significant increase in the ratio between galectin-3C and galectin-3 in tears of dry eye patients compared to tears of healthy subjects. Data for individual samples are presented. PGR, proline, glycine, and tyrosine-rich domain; CRD, carbohydrate recognition domain; N-term, N-terminus; aa, amino acid. *P < .05.
Figure 4.
Matrix metalloproteinase 9 promotes galectin-3 cleavage *in vitro*. (Upper panel) Quantification of MMP9 mRNA in conjunctival impression cytology samples from 10 normal subjects (10 eyes) and 15 dry eye patients (17 eyes) by qPCR. Data for individual samples are presented, with mean value indicated by horizontal bar for each group. rhMMP9 promoted galectin-3 cleavage in a concentration- and time-dependent manner (middle and
lower panels, respectively), producing a 25-kDa cleavage product. Samples in the middle panel were incubated at 37 °C for 81 min. Samples in the lower panel were incubated at 37 °C with 1.0 ng/μl rhMMP9. *P < .05.
Figure 5.
Degradation of galectin-3 in human tears involves protease activity. Immunoblots demonstrating partial or complete degradation of endogenous galectin-3 in an \textit{in vitro} assay using tears from three patients with dry eye. The broad-spectrum proteinase inhibitor cocktail \textsc{complete}\textsuperscript{TM}, but not the pan-specific MMP inhibitor GM6001, impaired cleavage of galectin-3 in samples from two patients. The presence of the latent (pro-MMP9) and active forms of MMP9 in these tear samples was confirmed by immunoblot. MMP9 blots were from the same tear samples used in the galectin-3 blots above.