

Defining Ocular Surface Disease Activity and Damage Indices by an International Delphi Consultation

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Defining Ocular Surface Disease Activity and Damage Indices by an International Delphi Consultation

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Memberships of participating groups are listed at end of article.

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ABSTRACT Purpose: Unifying terminology for the description of ocular surface disease (OSD) is vital for determining treatment responses and ensuring robust clinical trial outcomes. To date, there are no agreed parameters describing ‘activity’ and ‘damage’ phases of disease. **Methods:** A working group of international experts in OSD, oculoplastics, and uveitis from a range of backgrounds (university, teaching, district general and private hospitals) participated in a modified Delphi consensus-building exercise (October 31, 2011 to March 20, 2015). Two steering group meetings took place in which factors based upon published literature were discussed and supplemented with anonymous web-based questionnaires to refine clinical indices according to ‘activity’ (reversible changes resulting directly from the inflammatory process) and/or ‘damage’ (persistent, >6 months duration) changes resulting from previously active disease that are cumulative and irreversible).

Results: The recommended set of clinical parameters for the assessment of OSD encompasses 68 clinical indices and 22 ancillary grading tools (in parenthesis) subdivided by anatomical domain as follows: 4(4) tear-film, eyelid 21(3), 17(3) conjunctiva, 15(10) cornea and 11(2) Anterior Chamber/Sclera. Of these; 17(2) were considered as measures of clinical activity, 27(3) as damage, 1(8) as measures of both activity and damage. Twenty-three clinical descriptors and 9 tools did not reach the threshold for inclusion into the main standard set. These were defined as ‘second tier’ parameters for use in special clinical settings. **Conclusion:** These core parameters provide the first description of ‘activity’ and ‘damage’ relevant to OSD and provide a platform for the future development of scoring scales for each parameter.

KEY WORDS cornea, conjunctiva, Delphi process, disease activity, disease damage, disease scoring, disease staging, ocular surface disease

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I. Introduction

The ocular surface is a specialized mucosa comprising an epithelium and stroma extending from the mucocutaneous junction at the eyelid margin and including the tarsal, fornical and bulbar conjunctiva, limbus, and cornea.¹ Obtaining nutrition largely from the tear film and vascular arcades, and for the cornea, from the aqueous humor, the system also includes associated adnexal structures, lacrimal glands, and eyelids. All components are linked by the continuity of the surface epithelia and through close interaction with innervation, endocrine, vascular, and immune systems. Disease processes affecting the ocular surface system, defined as *ocular surface disease (OSD)*, have a diverse range of underlying pathologies, encompassing a spectrum of clinical entities, often with overlapping pathogeneses. Documentation of OSD includes the recording of physical signs of all components of the ocular surface system but also the posterior layers of the cornea, sclera and anterior chamber.

The breadth of OSD sequelae is exemplified by conditions forming the progressive conjunctival scarring subgroup, such as mucous membrane pemphigoid (**MMP**), Stevens-Johnson syndrome/toxic epidermal necrolysis (**SJS-TEN**) and graft-versus-host disease (**GVHD**).^{2,3} These conditions are characterized by conjunctival inflammation often associated with destruction of the normal ocular surface architecture, fibrosis, dry eye disease and eyelid deformities leading to surface breakdown, vulnerability to infection, limbal epithelial stem cell destruction, corneal scarring, neovascularization and eventually ocular surface failure. Chronic SJS-TEN can additionally be accompanied by scleritis.⁴

While OcMMP, SJS-TEN and GVHD are relatively rare compared to many OSDs, they serve as model disease platforms for quantifying the wealth of clinical signs for stratification according to ‘activity’ and ‘damage’ that may be generalizable across the whole spectrum of OSD. A range of scoring systems have been proposed for OcMMP, including conjunctival inflammation,⁵ scarring,⁶⁻⁸ methods for quantifying scarring,⁹ and for SJS-TEN the use of temporal-spacial staging (acute or chronic)^{10, 11} accompanied by ordinal scales for tear film, eyelid, corneal and conjunctival involvement; these scoring systems cannot be applied across all OSDs.^{3,11} ‘Function’ can also be used for documenting OSD indices, as illustrated by the dry eye and meibomian gland workshops.¹¹⁻¹⁴ These existing scoring systems are not in routine clinical use.

Disease ‘activity’ or ‘damage’ can also be determined by classifying a range of clinical parameters into 1) ‘activity,’ in which clinical manifestations are reversible and result directly from the inflammatory process, in which disease remission occurs spontaneously or following treatment such as immunosuppression; or 2) ‘damage,’ in which clinical manifestations are persistent, i.e., are present for greater than 6 months duration, **and** result in permanent changes in anatomy, physiology, pathology or function.¹⁵ Damage results from previously active disease where changes are often cumulative and irreversible.¹⁵

This method of scoring specifically benefits patients who present with early disease when the diagnosis is uncertain, and detailed investigation and/or prolonged follow-up would be required before the phenotype of the disease would manifest a diagnosis. A similar model is obtained from rheumatologic conditions in which early diagnostic criteria and the distinctions between disease ‘activity’ and ‘damage’ have been established and extrapolated to disease entities such as systemic lupus erythematosus, rheumatoid arthritis, and Sjögren syndrome.¹⁶⁻²¹ In such conditions, these terms are critical for defining relapse, remission and progression of disease, together with documenting treatment response, developing novel treatment response guidance, or ensuring robust outcomes for both small- and large-scale clinical trials.^{12,20,22,23} OSD is not absolutely synonymous with rheumatological conditions, and patient perceptions of OSD can influence the ocular surface severity score. For instance, the OSD patient may describe neuropathic pain that outweighs observed clinical signs, but influences the patient-reported outcomes of a putative ocular surface severity score. Defining ‘activity’ and ‘damage’ parameters provides an excellent foundation to begin the process of developing an ocular surface disease scoring system (**OSDISS**).

To meet the same end point in OSD as has been achieved in rheumatology, unifying terminology to describe the stage of ‘activity’ and ‘damage’ is required in addition to agreement on the grading of each parameter to stage severity or progression of disease. One method for achieving this is a consensus statement derived through a Delphi process. Originally developed by the United States Air Force during the Cold War, the Delphi process is structured to obtain a consensus opinion from a group of experts.²⁴ The advantage of this approach is its ability to gather information from multiple experts without the risk of giving greater weight to input from senior or more vocal individuals.²⁵

The objective of our study is to integrate specialist OSD knowledge through use of a modified Delphi technique to obtain consensus on a set of core clinical domains for the assessment of OSD and categorize according to activity/damage domains. Ultimately, our aim is to develop measurable scales for each clinical parameter, evaluate patient-reported outcomes, validate, and internationally adopt an agreed OSDISS that could be employed generically across all OSDs, particularly in the early stages when the diagnosis is not established.

II. Methods

The modified Delphi process was designed to conform to best practice described by Sinha and colleagues.²⁵

A. Steering Group Formation

An expert steering group was established whose remit was to identify measurable disease parameters for consensus building. Experts were identified using inclusion criteria as described for the *Dysfunctional Tear Syndrome Delphi* process.¹² “Experts” met the following requirements:

1. Active clinicians (ophthalmologists).
2. Scientific contributions to clinical research on the ocular surface or expertise in ocular surface, as reflected by at least two of: peer-reviewed publications, other forms of written scientific communication, specialty meeting presentations, and membership in international ocular surface disease societies.
3. Comprised international representation.
4. Proficiency in English language to facilitate interaction.
5. Ability to respond to sets of questionnaires and available to steering group meetings at the University of Birmingham, United Kingdom.

Of the UK-based steering group members, panelists were selected to represent a cross-section of the geographical patient population centers identified through a surveillance study.²⁶ Additional advisors with expertise in oculoplastics and uveitis were included to increase breadth of knowledge required for describing the adnexae and inflammatory eye diseases as part of the spectrum ocular surface system disorders. All but one of the 22 invited panelists accepted the invitation.

B. Summary of the Modified Delphi Process

A modified version of the Delphi process was used to obtain group consensus. This included the essential features of a consensus method described by Hunter and Jones.²⁷

- Anonymity – to avoid dominance of one individual within the group.
- Multiple iterations – giving participants the chance to change their views.

- Controlled feedback – showing the distribution of the group response.
- Statistical group response – expressing judgment using summary measures of the whole group response.

The consensus building exercise was divided into 5 key work packages (WP).

Work Package 1 (WP1): Preliminary Ballot

The first web-based ballot (eFigure 1) consisted of a nonexhaustive itemization of possible clinical parameters for inclusion in a scoring tool, derived from modification of existing scoring systems for OSD.^{5-8,11-14,28-30} Steering group participants were encouraged to suggest additional variables for inclusion in the putative activity and damage OSD scoring system.

Work Package 2 (WP2): The First Steering Group Meeting

The Steering Group convened to discuss results of WP1 using the nominal group technique.^{27,31} During this meeting, each clinical parameter was discussed. The group was invited to vote again in light of the discussion as to whether the clinical parameter should be included. Each member of the group was given an opportunity to chair part of the session, ensuring even representation from all group members. In order to encourage the inclusion of as many of the parameters as possible, a majority vote was accepted for inclusion of the clinical variable to a putative list of activity and damage indicators.

Work Package 3 (WP3): Web-based International Consultation

The results of the first steering group meeting were used to create a web-based questionnaire (eFigure 2) and disseminated to an international group of OSD clinical specialists invited via an undisclosed recipient electronic-mail communication to members of the *International Ocular Surface Society* and *UK Bowman Club* via their administrative secretaries. Additional invitations were extended to advisory group specialists in uveitis (n,10) and oculo-plastics (n,10) extrapolated from the *International Uveitis Study Group (IUSG)* and *International Group of Oculoplastics Specialists*, using the criteria outlined for the members of the OSD steering group. This questionnaire formed an anonymous specialist consultation where a minimum

number of participants (n=30) commensurate with other published Delphi processes were defined to ensure validity of the consultation exercise.^{12,32,33}

OSD specialist recipients were asked to rank the 76 agreed indices and 30 ancillary grading tools over the 5 clinical domains (tear-film, eyelids, conjunctiva and fornices, cornea and sclera/anterior chamber [AC]) in the context of eight common or important clinical disease entities:

1. Blepharitis
2. Mucous membrane pemphigoid or other progressive cicatrizing conjunctivitis
3. Bacterial keratitis
4. Viral keratitis (e.g., herpetic keratitis)
5. Dry eye disease
6. Sjögren's syndrome (primary or secondary)
7. Corneal melt including peripheral ulcerative keratitis
8. Chemical injury

Participants were asked to consider the 'best' ancillary grading tool by "assuming you have all of these available" (see supplementary figure 2) and a gauge of the 'real world' scenario was examined by asking: "Which of these (these tools) do you have available?" Oculoplastics and uveitis specialist consultations were restricted to domains relevant to their subspecialties comprising tear film/conjunctiva-fornices/cornea and cornea/anterior chamber-sclera, respectively.

Based upon the variation among published 'cut-off' limits ranging from 60% to 80% to determine consensus,^{12,34-38} a 75% cut-off of the upper and lower quartiles in order to balance definite agreement ("agree" or "strongly agree"), disagreement ("disagree" or "strongly disagree") and areas of ambiguity was identified. A timeline of 10 weeks was set for completion. Information was anonymously populated into an Adobe Form Central data capture sheet (<https://new.acrobat.com>, 2013).

Work Package 4 (WP4): Second Steering Group Meeting

The results of the consultation questionnaire were presented at a second steering group meeting. The live anonymous web-based voting system using Adobe Form Central enabled unbiased arbitration during the following Expert workshops:

- *Workshop A (WSA): Unclassified Clinical Parameters.* Clinical parameters with $\geq 75\%$ consensus were included into the scoring system; otherwise they were classed as ‘second-tier.’ Second-tier parameters were defined as parameters for specialist situations (but not essential for a general ophthalmology setting).
- *Workshop B (WSB): ‘Activity’ and ‘Damage’.* The stratification of each parameter and grading tool into one of three activity-damage domain categories measuring 1) “activity”, 2) “damage”, and 3) both “activity and damage” was undertaken. A $\geq 75\%$ consensus was required for classification where *activity* was defined as reversible/medically modifiable manifestations or *damage* as manifestations that are persistent (>6 months) and result from previously active disease. Parameters and tools that could not be successfully classified into an activity-damage domain, formed a fourth domain termed ‘unclassified’ (either ‘activity’ or ‘damage’) and were subject to further consultation in WP5.

Work Package 5 (WP5): Final Consultation Defining ‘Activity’ and ‘Damage’.

Clinical parameters from WP5 Workshop B that remained unclassified were arbitrated through a final round of voting involving the OSD specialists where participants were asked to classify the ambiguous indices as 1) “activity,” 2) “damage,” and 3) both “activity and damage.” A $\geq 70\%$ consensus enabled direct definition of ‘activity’ and ‘damage’ domains to ensure maximal classification of remaining parameters. The outcome dataset from *Work Package 5* produced a consensus statement on a set of five core clinical domains for the assessment of OSD stratified across three activity-damage domains.

C. Approvals

Institutional review board approval was not required because the study did not involve patient or registry data and this was confirmed by the Research Support

Group Research Ethics Team (ref ERN_15-1195) at the University of Birmingham (UK).

ACCEPTED MANUSCRIPT

III. Results

A diagram of the work packages and summary of outcomes is outlined in Figure 1.

A. WP1 and WP2: Preliminary Ballot and The First Steering Group Meeting.

Seventy-six clinical indices were subdivided into five domains: Tear film; Eyelids, Lid Margins and Meibomian Glands; Conjunctiva and Fornices; Cornea; Anterior Chamber and Sclera were disseminated as part of *Work Package 1 (Preliminary Ballot)*. The Ballot ‘included’ the majority of proposed indices, with highest agreement in the fornix subset versus the tear film (lowest) (**Figure 2**).

Twelve clinical parameters were rejected from across all domains and two were reclassified as a new category, “ancillary grading tools,” i.e. methods of examining and quantifying disease (see below). Fifteen new clinical parameters were added, giving rise to a revised total of 76 clinical indices (Figure 2). It was agreed that a list of 30 additional “ancillary grading tools” should be constructed and considered as part of the WP3 international consultation exercise (Table 1). A scoping exercise to define the availability of these tools in the Delphi responders’ clinical practice was proposed to be integrated into WP3.

B. WP3: Web-based International Consultation

A total of 53 specialists responded (40 OSD, 5 oculoplastics, 8 uveitis) with most (47) practicing in a University Teaching Hospital. Specialists were from 8 countries (Australia 10, Belgium 1, Germany 3, Slovenia 1, New Zealand 1, Singapore 2, United Kingdom 32 and United States 2, one person did not specify their country of origin in the anonymous form)

Of the 76 clinical indices, 52 (68%) achieved $\geq 75\%$ ranking for inclusion and none reached the 75% threshold for exclusion.

Twenty-two of 30 ancillary grading tools achieved the 75% threshold (Table 1, Figure 3). Table 1 also highlights the availability 13 (43%) ancillary grading tools that were available in clinical practice to $\geq 75\%$ of participants.

C. WP4 and WP5: Second Steering Group Meeting and Final Consultation

Workshop A: Unclassified Clinical Parameters

Twenty-four of the 76 clinical parameters did not reach the $\geq 75\%$ threshold. After Steering Group discussion, anonymous web-based arbitration included a further 15 unclassified indices and 1 novel clinical entity (scleral thinning) was introduced giving a total of 68 indices (Figure 3). The ten remaining indices formed '2nd tier' clinical parameters.

The steering group arbitrated that ancillary grading tools that did not reach the $\geq 75\%$ threshold became '2nd tier' tools for use in refining scales in specialist situations (Figure 3)

Workshop B and WP5: Activity and Damage Stratification

During WP4 Workshop B, one parameter (palpebral aperture) was considered to be a physiological or anatomical descriptor that was not amenable to classification into activity and damage domains and was therefore not put forward to the final consultation forming WP5. This parameter formed an 'unclassified' category.

Following completion of the final round of questionnaires, 17 clinical parameters and two ancillary grading tools (in parenthesis) were classified as a measure of "activity": Tear film: 0(1), Lids: 4(0), Conjunctiva: 7(0), Cornea: 0(0), AC/Sclera: 6(1). Twenty-seven clinical parameters and 3 ancillary grading tools were classified as a measure of "damage": Tear film: 0(0), Lids: 10(2), Conjunctiva: 10(1), Cornea: 4(0), AC/Sclera: 3(0). One of the clinical parameters and 6 of the ancillary grading tools were classified as a measure of both "activity and damage" with photography appearing in 3 clinical domains (lids, conjunctiva, cornea). The remaining clinical parameters and ancillary grading tools formed 2nd tier for activity and damage: Tear film: 4(3), Lids: 7(0), Conjunctiva: 0(1), Cornea: 11(5), AC/Sclera: 1(0). (Figure 4, eTable 1).

IV. Discussion

This is the first international consensus statement on putative descriptors of OSD and discriminators of 'activity' and 'damage' providing a platform for standardizing terminology when describing disease staging and progression. This is of considerable importance for identifying response to treatment and to enable robust

outcome comparisons between clinical trials or gauging response to novel tissue specific therapeutic interventions.³⁹

Current OSD scoring systems are disparate, with multiple systems describing specific disease entities. While OSD-specific questionnaires relating to symptoms have been validated,³⁹ to date there is no generic consensus on accepted clinical indices and how these should be scored, independent of the underlying disease process. This is essential, as the overall end-stage for all OSD is ocular surface failure and, as such, it should be possible to assimilate a battery of clinical features and scales to apply broadly across numerous OSD processes, particularly at the early stages when the diseases may present with indistinguishable signs, e.g., red eye. To this end, we sought to establish an 'OSD Toolbox', where the clinician has the ability to select the most relevant 'tools' to accurately describe the clinical features of a patient who may not have a diagnosis or for whom the diagnosis is equivocal. This could provide the basis for stratifying the 'Tools' into descriptors of recognized disease entities. This process could allow retrospective analyses of clinical features at presentation, thereby identifying putative common features in stages of disease before the diagnosis is known. This methodology could enable clues to support earlier diagnosis and prediction of OSD clinical course thereby leading to earlier intervention and improved clinical outcomes.

The use of the Delphi approach in the healthcare setting is well established^{12,40-42} and has been used effectively across multiple specialties (rheumatology,^{16,32,43} ophthalmology,^{12,31} palliative care,⁴⁰ orthopedics,³⁵ and anesthesiology^{36, 37}). It excels when there is either a paucity of evidence, such as in the description of OSD, or when the available evidence is contradictory.^{27, 42} Limitations include reliability and reproducibility, possibly due to group selection.^{42,44,45} To overcome this, we identified participants based upon geographical distribution of the *British Ophthalmological Surveillance Unit* respondent data to ensure even expert representation of steering group attendees from across the United Kingdom.²⁶ As in other published literature, participant selection was non-random, so representativeness is not assured,⁴² but potential bias was minimized through the use of anonymous web-based questionnaires throughout the process. Where group discussions were held, opportunity was given for each participant to lead part of the group discussion to reduce the effect of a dominant individual.

Often the Delphi process is used when evidence is limited or absent in a given subject field, leading to the possibility of collective group error.⁴¹ Equally, the results may be in direct conflict with the available evidence. We included the majority of clinical parameters and ancillary grading tools, with the exclusion of very few indices. Since little has been excluded, collective group error has been attenuated, and the study has created an inclusive platform from which further refinements can be made. Work Package 2 saw a high level of agreement in Domain 5 (AC and Sclera), which has been the subject of a previous Delphi consensus endorsing reproducibility of the results of the Delphi process.²⁹

High levels of agreement were found for activity (conjunctival inflammation, foamy meibomian gland secretions, presence of anterior chamber cells, hypopyon) and damage (entropion, ectropion, horizontal forniceal involvement by fibrosis, iris atrophy). No clinical parameters or investigations were directly voted into the combined activity and damage domain. Some parameters failed reach sufficient consensus for classification as 'activity', 'damage' or 'activity and damage'. An example of this is tear film breakup time. This reached a 66.6% consensus for 'activity' in WP4, but only 50% consensus in WP5. A similar phenomenon occurred in the Cornea Domain, with none of the parameters being classified as measures of 'activity'. This is possibly because the votes were split across 'activity' and 'activity and damage'. Further iterations of the process are required for refinement. We acknowledge that setting a higher threshold resulted in exclusion of some parameters. The threshold was reduced to 70% for work package 5 to improve classification, but the Delphi process clearly highlighted significant disagreement among specialists, e.g., in corneal activity, and we believe this is likely to reflect the disparate nature of corneal disease. For this reason, no parameter was fully excluded, and those that did not meet a consensus were available in the reserve pool that was termed 'second tier'. We believe that corneal disease activity will be more easily defined in a disease specific context, beyond the remit of this exercise and will be focus for future validation work.

Following WP5, the distribution of votes among the remaining parameters highlighted sufficient uncertainty amongst participants such as the presence of filaments. This enabled these entities to default into a 2nd tier combined activity-damage domain rather than being excluded from the dataset.

The current list should be considered a platform for further development. Indeed, future work is required to refine the 'Tool Box' of clinical features describing OSD with scales for quantifying each parameter to enable its use in more specific disease processes.^{12, 35-37} This will necessitate further group discussions, literature review, and definition of severity scales. Prospective collection of patient data, including patient-reported outcomes encompassing vision related quality of life and neuropathic pain (a clinical feature that has recently gathered considerable interest), are essential composites to computing an activity and damage score and generating an OSDISS. We recognize that the outcome of this Delphi process provides a 'first step' to achieving this goal and should not be seen as a final arbiter. It is hoped that by defining these in a disease-specific context, it will be clearer how to relate scales to activity or damage or both. This specifically benefits patients who present with early disease when the diagnosis is uncertain and only after detailed investigation and/or prolonged follow-up, the phenotype of the disease manifests a diagnosis. Furthermore, with wider application of certain ancillary tools, e.g., AS-OCT, this may alter the profile of how parameters and diseases are quantified.

This study considered 'activity' and 'damage' in the context of inflammation. While inflammation is the major contributor in many ocular surface diseases, we acknowledge that dysfunctional innervation or the mechanical breakdown of the corneal surface, e.g., in recurrent corneal erosion syndrome, may themselves relate to 'activity' and 'damage' independent of overtly manifest inflammation per se. There is, however, increasing awareness that all processes at a molecular level have an underlying inflammatory component, as tissue injury of any nature, whether exogenous or endogenous, will release cytokines, inflammatory mediators and promote recruitment of inflammatory cells. This has, for example, been recently recognized in the context of progressive conjunctival scarring in clinically quiescent eyes where neutrophils have been identified as critical in mediating disease progression.^{46,47}

While the breadth of this toolbox may appear daunting, it is hoped that this Delphi process has taken the first step in pulling disparate indices into a single arena and for the first time gaining agreement. The most obvious utility for the OSDISS scales outside the research setting is the provision of a matrix from which electronic patient records (**EPR**) can be developed. The purpose of this study was not to create individual scales for each disease, enabling those in current use to be adopted for EPR

prior to disease specific validation exercises. Ultimately, the point of this exercise was to create an environment for further development. For instance, in forniceal scarring, we would propose the adoption of the fornix depth measurer to measure scarring, as this has been through an intra- and inter- observer validation exercise.^{9,48} In EPR development, this would be considered a continuous value that could be compared over time and for the purpose of recording damage. Conjunctival inflammation, however, may be considered through various ordinal scales, e.g., 1-4, 1-5 etc.^{5,49} Until such validation exercises have been completed, developers may elect to choose one or the other in order to facilitate EPR completion. In turn these may provide benchmark data for national data collection and audit exercise.

V. Conclusion

The validation of longitudinal collection of clinical ‘activity’ and ‘damage’ with grading scales correlated to measures of patient perception, experience, and reported outcomes of disease will provide a valuable objective resource for interrogating accurately described clinical features at presentation when diagnosis is equivocal. This will provide clues to earlier diagnosis, prediction of disease course, and improved clinical outcomes. It will also allow standardization of research data and a unified approach to objective assessment of treatment response, specifically to novel interventions in a clinical setting.

References

1. Gipson IK. The ocular surface: the challenge to enable and protect vision. *Invest Ophthalmol Vis Sci* 2007;48:4391-8
2. Chan LS, Ahmed AR, Anhalt GJ, et al. The first international consensus on mucous membrane pemphigoid: definition, diagnostic criteria, pathogenic factors, medical treatment, and prognostic indicators. *Arch Dermatol* 2002;138:370-9
3. Ogawa Y, Kim SK, Dana R, et al. International Chronic Ocular Graft-vs-Host-Disease (GVHD) Consensus Group: proposed diagnostic criteria for chronic GVHD (Part I). *Sci Rep* 2013;3:3419
4. De Rojas MV, Dart JK, Saw VP. The natural history of Stevens Johnson syndrome: patterns of chronic ocular disease and the role of systemic immunosuppressive therapy. *Br J Ophthalmol* 2007;91:1048-53

5. Elder MJ, Bernauer W. Monitoring of activity and progression in cicatrising conjunctivitis, in Bernauer W, Dart JKG, Elder MJ (eds). *Cicatrising Conjunctivitis*. Basel, Switzerland: Karger, 1997; v. 28
6. Tauber J, Jabbur N, Foster CS. Improved detection of disease progression in ocular cicatricial pemphigoid. *Cornea* 1992;11:446-51
7. Mondino BJ, Brown SI. Ocular cicatricial pemphigoid. *Ophthalmology* 1981;88:95-100
8. Foster CS. Cicatricial pemphigoid. *Trans Am Ophthalmol Soc* 1986;84:527-663
9. Williams GP, Saw VP, Saeed T, et al. Validation of a fornix depth measurer: a putative tool for the assessment of progressive cicatrising conjunctivitis. *Br J Ophthalmol* 2011;95:842-7
10. Power WJ, Ghorraishi M, Merayo-Llodes J, et al. Analysis of the acute ophthalmic manifestations of the erythema multiforme/Stevens-Johnson syndrome/toxic epidermal necrolysis disease spectrum. *Ophthalmology* 1995;102:1669-76
11. Sotozono C, Ang LP, Koizumi N, et al. New grading system for the evaluation of chronic ocular manifestations in patients with Stevens-Johnson syndrome. *Ophthalmology* 2007;114:1294-302
12. Behrens A, Doyle JJ, Stern L, et al. Dysfunctional tear syndrome: a Delphi approach to treatment recommendations. *Cornea* 2006;25:900-7
13. No authors listed. The definition and classification of dry eye disease: report of the Definition and Classification Subcommittee of the International Dry Eye WorkShop (2007). *Ocul Surf* 2007;5:75-92
14. Denniston AKO, Murray PI. *Oxford Handbook of Ophthalmology, 2nd ed.* Oxford: Oxford University Press, 2009; xl, 930 pp
15. Bowman SJ, Pillemer S, Jonsson R, et al. Revisiting Sjögren's syndrome in the new millennium: perspectives on assessment and outcome measures. Report of a workshop held on 23 March 2000 at Oxford, UK. *Rheumatology (Oxford)* 2001;40:1180-8
16. Seror R, Ravaud P, Bowman SJ, et al. EULAR Sjogren's syndrome disease activity index: development of a consensus systemic disease activity index for primary Sjogren's syndrome. *Ann Rheum Dis* 2010;69:1103-9

17. Bowman SJ, Sutcliffe N, Isenberg DA, et al. Sjogren's Systemic Clinical Activity Index (SCAI)--a systemic disease activity measure for use in clinical trials in primary Sjogren's syndrome. *Rheumatology (Oxford)* 2007;46:1845-51
18. Smolen JS, Strand V, Cardiel M, et al. Randomized clinical trials and longitudinal observational studies in systemic lupus erythematosus: consensus on a preliminary core set of outcome domains. *J Rheumatol* 1999;26:504-7
19. Felson DT, Anderson JJ, Boers M, et al. The American College of Rheumatology preliminary core set of disease activity measures for rheumatoid arthritis clinical trials. The Committee on Outcome Measures in Rheumatoid Arthritis Clinical Trials. *Arthritis Rheum* 1993;36:729-40
20. Tugwell P, Boers M, Brooks P, et al. OMERACT: an international initiative to improve outcome measurement in rheumatology. *Trials* 2007;8:38
21. Gordon C, Bertsias G, Ioannidis JP, et al. EULAR points to consider for conducting clinical trials in systemic lupus erythematosus. *Ann Rheum Dis* 2009;68:470-6
22. Douglas RS, Tsirbas A, Gordon M, et al. Development of criteria for evaluating clinical response in thyroid eye disease using a modified Delphi technique. *Arch Ophthalmol* 2009;127:1155-60
23. Naredo E, Wakefield RJ, Iagnocco A, et al. The OMERACT ultrasound task force--status and perspectives. *J Rheumatol* 2011;38:2063-7
24. Dalkey N, Helmar O. Experimental application of the Delphi method to the use of experts. *Management Sci* 1963;9:458-67
25. Sinha IP, Smyth RL, Williamson PR. Using the Delphi technique to determine which outcomes to measure in clinical trials: recommendations for the future based on a systematic review of existing studies. *PLoS Med* 2011;8:e1000393
26. Radford CF, Rauz S, Williams GP, et al. Incidence, presenting features, and diagnosis of cicatrising conjunctivitis in the United Kingdom. *Eye (Lond)* 2012;26:1199-208
27. Jones J, Hunter D. Consensus methods for medical and health service research. *BMJ* 1995;311(7001):376-80
28. Bron AJ. A simple scheme for documenting corneal disease. *Br J Ophthalmol* 1973;57:629-34

29. Jabs DA, Nussenblatt RB, Rosenbaum JT, et al. Standardization of uveitis nomenclature for reporting clinical data. results of the First International Workshop. *Am J Ophthalmol* 2005;140:509-16
30. Schwab IR, Linberg JV, Gioia VM, et al. Foreshortening of the inferior conjunctival fornix associated with chronic glaucoma medications. *Ophthalmology* 1992;99:197-202
31. Douglas RS, Tsirbas A, Gordon M, et al. for the International Thyroid Eye Disease Society. Development of criteria for evaluating clinical response in thyroid eye disease using a modified Delphi technique. *Arch Ophthalmol* 2009;127:1155-60
32. Bowman SJ, Jonsson R, Assmussen K, et al. . Revisiting Sjogren's syndrome in the new millennium: perspectives on assessment and outcome measures. Report of a workshop held on 23rd of March 2000 at Oxford, UK. *Rheumatology* 2001;40:1180-8
33. Arnaud L, Devilliers H, Peng SL, et al. The Relapsing Polychondritis Disease Activity Index: development of a disease activity score for relapsing polychondritis. *Autoimmun Rev* 2012;12:204-9
34. Olthof DC, van der Vlies CH, Joosse P, et al. Consensus strategies for the nonoperative management of patients with blunt splenic injury: a Delphi study. *J Trauma Acute Care Surg* 2013;74:1567-74
35. Diaz-Ledezma C, Higuera CA, Parvizi J. Success after treatment of periprosthetic joint infection: a Delphi-based international multidisciplinary consensus. *Clin Orthop Relat Res* 2013;471:2374-82
36. Boogaard S, Heymans MW, Patijn J, et al. Predictors for persistent neuropathic pain – a Delphi survey. *Pain Physician* 2011;14:559-68
37. Stanton TR, Latimer J, Maher CG, Hancock MJ. A modified Delphi approach to standardize low back pain recurrence terminology. *Eur Spine J* 2011;20:744-52
38. Morrison AP, Barratt S. What are the components of CBT for psychosis? A Delphi study. *Schizophr Bull* 2010;36:136-42
39. Schiffman RM, Christianson MD, Jacobsen G, et al. Reliability and validity of the Ocular Surface Disease Index. *Arch Ophthalmol* 2000;118:615-21
40. Woitha K, Van Beek K, Ahmed N, et al. Validation of quality indicators for the organization of palliative care: a modified RAND Delphi study in seven European countries (the Europall project). *Palliat Med* 2014;28:121-9

41. Armon K, Stephenson T, MacFaul R, et al. An evidence and consensus based guideline for acute diarrhoea management. *Arch Dis Child* 2001;85:132-42
42. Hasson F1, Keeney S, McKenna H. Research guidelines for the Delphi survey technique. *J Adv Nurs* 2000;32(4):1008-15
43. Shiboski SC, Shiboski CS, Criswell LA, et al. American College of Rheumatology classification criteria for Sjögren's syndrome: a data-driven, expert consensus approach in the SICCA cohort. *Arthritis Care Res* 2012;64:475-87
44. Burnand B, Vader JP, Froehlich F, et al.. Reliability of panel-based guidelines for colonoscopy: an international comparison. *Gastrointest Endosc* 1998;47:162-6
45. Campbell SM, Roland MO, Quayle JA, Shekelle PG. The effect of panel membership and feedback on ratings in a two-round Delphi survey: results of a randomized controlled trial. *Medical Care* 1999;37:964-8
46. Arafat SN, Suelves AM, Spurr-Michaud S, et al. Neutrophil collagenase, gelatinase, and myeloperoxidase in tears of patients with stevens-johnson syndrome and ocular cicatricial pemphigoid. *Ophthalmology* 2014;121:79-87
47. Williams GP, Southworth HS, Denniston AKO, et al. Conjunctival neutrophils predict progressive scarring in Ocular Mucous Membrane Pemphigoid. *Invest Ophthalmol Vis Sci* 2016, in press
48. Khan IJ, Ghauri AJ, Hodson J, et al. Defining the limits of normal conjunctival fornix anatomy in a healthy South Asian population. *Ophthalmology* 2014;121:492-7
49. Saw VP, Dart JK, Rauz S, et al. Immunosuppressive therapy for ocular mucous membrane pemphigoid strategies and outcomes. *Ophthalmology* 2008;115:253-61 e1

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Study group members contributed to steering group meetings and answered web-based questionnaires. They received reimbursement for travel expenses if they attended the steering group meetings. No other financial incentive was provided.

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(b) Data acquisition
(c) Data analysis and interpretation
- 2) (a) Drafting the article
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Figure legends

Figure 1: Diagrammatic representation summarizing the work packages forming the OSD Delphi process.

Figure 2: Summary of the results from WP1 and WP2. The results of the pre-meeting questionnaire (WP1) were discussed and categorized after anonymized voting at the first steering group meeting (WP2). Fifteen new parameters were added, 12 excluded and 2 were redeployed to a novel “ancillary grading tool” grouping (AGT). Some of the terms were altered at the first steering group meeting; ‘tear meniscus’ was changed to ‘tear meniscus height’, ‘anterior blepharitis’ was changed to ‘anterior lid margin disease’, ‘measurement of upper/lower fornix’ became ‘measurement of upper/lower fornix central depth’, ‘central corneal ulceration’ and ‘central corneal depth’ were changed to ‘localized corneal ulceration’ and ‘localized corneal depth’ respectively.

Figure 3: Results of the second steering group meeting (WP3 and WP4 Workshop A). Clinical parameters that met the $\geq 75\%$ consensus threshold in the Web-based International Consultation (WP3) together with equivocal parameters that were positively considered by the Steering Group (WP4 WSA) are shown in column 1. Equivocal clinical parameters that did not reach the $\geq 75\%$ consensus were placed in a “second tier”. Similarly, Ancillary grading tools that achieved $\geq 75\%$ consensus were included as tools essential for severity staging and the remainder were classified as ‘second tier’ to be used under specialist circumstances.

[[♦] added at the second meeting, ^{*} Term amended]

Figure 4: Work Package 4 Workshop B and Work Package 5: Defining

“activity” or “damage”. Attendees of the Second Steering Group Meeting (Work Package 4) were asked to participate in a web-based anonymous live voting exercise to determine clinical parameters and ancillary grading tools indicative of “activity” or “damage” or positively both “activity and damage”. A $\geq 75\%$ consensus was required to be classified as such for WP4 (percentage vote for classification is in parenthesis). The remaining clinical parameters and ancillary grading tools were arbitrated by a wider consultation of OSD specialists (WP5). A $\geq 70\%$ consensus was required for WP5 to ensure classification of a greater number of parameters. Parameters highlighted in red were successfully classified following WP5 and those that did not meet 70% consensus, were defaulted to ‘second tier’ “activity” and “damage” to be used under specialist circumstances.

eFigure 1: Preliminary ballot (Work Package 1). The first web-based questionnaire that was disseminated to participants prior to Work Package 2.

eFigure 2: Web-based International Consultation (WP3). Web-based International Questionnaire. Ocular Surface Disease Specialists were invited to vote on all 5 domains. Uveitis experts were asked to respond to Domains 4 (Cornea) and 5 (AC and Sclera) only. Oculoplastics experts were asked to respond to the Domains 1 (Tear film), 2 (Eyelids lid margins and Meibomian glands) and 4 (cornea).

Table 1: Summary of Ancillary Grading Tools

Considered	Included	Available in clinical practice
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<ul style="list-style-type: none"> • Tear function index • Tear film osmolarity • Schirmer's test • Fluorescein staining • Lissamine green staining • Meibography (meibomian gland drop out) • Fornix depth measurement • Confocal microscopy - tarsus • Confocal microscopy - conjunctiva • Confocal microscopy - cornea • Endothelial cell morphology and counting - specular microscopy 	<p>Domain 1: Tear film</p> <ul style="list-style-type: none"> • Tear film osmolarity • Schirmer's test • Fluorescein staining • Lissamine green staining <p>Domain 2: Lids, lid margins and meibomian glands</p> <ul style="list-style-type: none"> • Fornix depth measurement • Meibography • Photography <p>Domain 3: Conjunctiva and fornices</p> <ul style="list-style-type: none"> • Immunostaining • Fornix depth measurement • Photography 	<ul style="list-style-type: none"> • B Scan ultrasound • Corneal topography e.g. Orbscan, pentacam • Endothelial cell specular microscopy • Fluorescein staining • Fluorescein angiography (but only 20% had a specific protocol for the anterior segment) • Impression and brush cytology for retrieval of cells • Histological and cell staining • Immunostaining • OCT – spectral domain • Pachymetry • Photography
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<ul style="list-style-type: none"> • Tissue biomarkers (cells, proteins, lipids, gene expression) • Biofluid biomarkers (tears, aqueous humor, blood, serum) • Impression and brush cytology for retrieval of cells • Histological and cell staining • Immunostaining • Indocyanine green angiography • Fluorescein angiography 	Domain 4: Cornea <ul style="list-style-type: none"> • Aethesiometry Orbscan/Pentacam • OCT - spectral domain • Pachymetry • Fluorescein staining • Lissamine green staining • Impression and brush cytology • Histological and cell staining • Photography • Endothelial cell specular microscopy 	<ul style="list-style-type: none"> • Schirmer's test • Tonometry
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<ul style="list-style-type: none"> • Aesthesiometry (corneal sensation) • Corneal shape and thickness measurements; e.g. Orbscan and Pentacam • OCT - spectral domain • OCT - time domain • Wavefront aberrometry • Pachymetry • Corneal hysteresis • B-Scan Ultrasonography • High resolution anterior segment ultrasonography • Laser flare meter • Tonometry • Photography under standardized conditions 	Domain 5: AC and sclera <ul style="list-style-type: none"> • Laser flare meter • Anterior segment ultrasound
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Footnote: A list of all tools considered is shown together with a summary of the grading tools that met the 75% threshold after WP3 (International consultation) categorized according to clinical domain. Those that did not meet this threshold became second tier investigations. These were; biofluid biomarkers, B-scan ultrasonography, Confocal microscopy (conjunctival), confocal microscopy (corneal), confocal microscopy (tarsal), corneal hysteresis, fluorescein angiography, indocyanine green angiography, OCT (time domain), tear function index, tissue biomarkers, tonometry, wavefront aberrometry. Availability of listed tools in clinical practice for >75% respondents is also shown. (Abbreviations: OCT, optical coherence tomography).

Figure 1

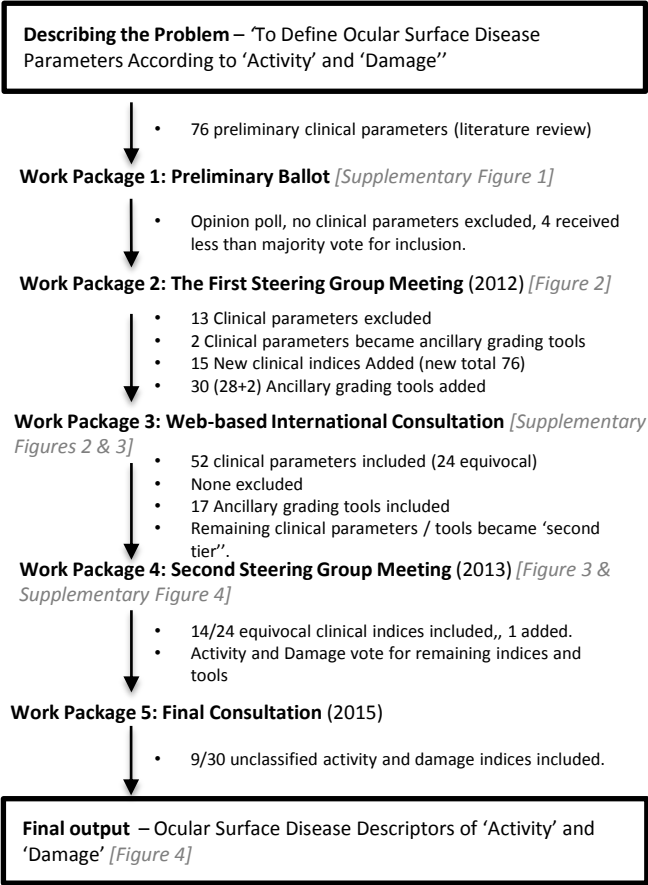


Figure 2

	Included	Added	Excluded	
Domain 1 Tear film	Tear film break up time Tear Meniscus Height Filaments	Tear clearance rate Tear film debris	InflammaDry Schirmer's test (to ancillary grading tools) Tear film osmolality (to ancillary grading tools)	
Domain 2 Lids and lid margins	Lid position and margins Lagophthalmos "close your eyes gently" Lid malposition Entropion Ectropion Lid margin irregularity Abnormal vascularity Anterior lid margin disease Medial Marx's line Central Marx's line Lateral Marx's line Lashes Lash loss Trichiasis	Dystichiasis Meibomian glands Pouting/plugging Opaque/scarred Meibomian gland orifice retroplacement Visibility of acini Concretions Chalazion formation Foam Quality of secretions	Palpebral aperture Lid margin reflex distances (one and two) Completeness of the blink cycle Standardized meibomian gland expressibility Lid margin thickening	Lid margins Upper puncta Lower puncta Mucocutaneous junction: Anteroplacement Retroplacement Mucosal absorption Ridging Meibomian glands: Cystoid dilation of the main duct
Domain 3 Conjunctiva & fornices	Conjunctiva Conjunctival inflammation (with each outer quadrant scored separately) Ulceration of bulbar conjunctiva Limbitis (with each quadrant scored separately) Conjunctival mucus Conjunctival keratinization Upper fornix Measurement of upper fornix central depth Upper fornix number of symblephara Upper fornix horizontal involvement by symblephara Upper fornix horizontal fibrosis	Lower fornix Measurement of lower fornix central depth Lower fornix number of symblephara Lower fornix horizontal involvement by symblephara Lower fornix horizontal involvement by fibrosis Ocular mobility Limitation in mobility	Punctate conjunctival staining Conjunctival papillae Conjunctival follicles	Peripheral ulcerative keratitis: degree of surface involvement Peripheral ulcerative keratitis: Depth.
Domain 4 Cornea	Sensation Punctate keratopathy Conjunctivalization Neovascularization (peripheral involvement) Neovascularization (encroaching on the visual axis) Corneal epithelial edema Corneal stromal edema Corneal opacification/scarring Corneal opacification/scarring encroaching on the visual axis Localized corneal thinning (no ulceration) Localized corneal ulceration	Localized corneal ulcer depth Descemet's folds Reduced endothelial cell count (clinically apparent) Graft-host interface changes	Corneal Ischemia Endothelial guttae	Endotheliitis Conjunctival keratinization (also on conjunctiva and kept there)
Domain 5 Anterior chamber & Sclera	Anterior Chamber Keratic precipitates Anterior chamber cells Flare Hypopyon Posterior synechiae Iris Iris nodules Sclera Anterior scleritis Posterior scleritis	Anterior synechiae (peripheral or central) Iris atrophy Intra-ocular pressure	Nil	

Figure 3

		> 75% Threshold for inclusion		Steering Group Vote for Inclusion		Second tier	
Domain 1	Tear film	Clinical	Tear film break up time		Tear Meniscus Height Tear Film Debris Filaments		Tear Clearance Rate
		Tool	Tear film osmolarity Fluorescein staining		Lissamine green staining Schirmer's test		Tear function index Biofluid biomarkers
Domain 2	Lids and lid margins	Clinical	Lid position Completeness of the blink cycle Lagophthalmos Lid malposition Entropion Ectropion Lid margins Lid margin irregularity Anterior lid margin disease	Lashes Trichiasis Dystrichiasis Meibomian glands Pouting/plugging Orifice retroplacement Quality of secretions	Lid position Palpebral aperture Lid margins Lid margin thickening Mucocutaneous junction (Marx's line)* Lashes Lash loss Meibomian glands Opaque/scarred Visibility of acini Chalazion formation Foam Expressibility		Lid margin reflex distances (one and two) Concretions Standardized meibomian gland expressibility Lid vascularity♦
		Tool	Fornix depth measurement Meibography Photography				
Domain 3	Conjunctiva & fornices	Clinical	Conjunctiva Inflammation Bulbar ulceration Limbitis Mucus Keratinization Punctate staining Papillae Follicles Ocular mobility Limitation in mobility	Upper fornix Number of symblephara Horizontal involvement by symblephara Horizontal involvement by fibrosis Lower fornix Central depth Number of symblephara Horizontal involvement by symblephara Horizontal involvement by fibrosis	Upper fornix central depth		Conjunctival chalasis♦
		Tool	Immunostaining Fornix depth measurement	Photography			Tarsal confocal microscopy Conjunctival confocal microscopy Tissue biomarkers
Domain 4	Cornea	Clinical	Sensation Punctate keratopathy Conjunctivalization Neovascularization (peripheral involvement) Neovascularization (encroaching on the central axis) Corneal epithelial edema	Corneal stromal edema Corneal opacification/scarring Corneal opacification/scarring encroaching on the visual axis Localized corneal thinning (no ulceration) Localized corneal ulceration Localized corneal ulcer depth Descemet's folds	Graft-host interface changes Endothelial guttata		Corneal ischemia Reduced endothelial cell count (clinically apparent)
		Tool	Endothelial cell specular microscopy Impression and brush cytology Histological and cell staining Aesthesiometry Corneal topography	Spectral domain OCT Pachymetry Fluorescein staining Lissamine green staining Photography			Corneal confocal microscopy Fluorescein angiography Indocyanine green angiography Time domain OCT Wavefront aberrometry Corneal hysteresis
Domain 5	Anterior chamber & Sclera	Clinical	Anterior Chamber Keratic precipitates Cells Flare Hypopyon Anterior synechiae (peripheral or central) Posterior synechiae Iris Iris atrophy	Sclera Anterior scleritis Posterior scleritis Other Intraocular pressure	Scleral thinning♦		Iris nodules
		Tool	High resolution anterior segment ultrasound Laser flare meter				B-scan ultrasonography Tonometry

Figure 4

	Activity	Damage	Activity and Damage	Activity and Damage: Second Tier
Domain 1	Tear film	Clinical		Tear film break up time Tear Meniscus Height Tear Film Debris Filaments
			<i>Tear film osmolarity (75%)</i>	<i>Fluorescein staining</i> <i>Lissamine green staining</i> <i>Schirmer's test</i>
Domain 2	Lids and lid margins	Clinical	Lid margins Anterior lid margin disease (83%) Meibomian glands Pouting/plugging (75%) Foam (100%) Quality of secretions (91.7%)	Lid position Entropion (91.7%) Ectropion (91.7%) Lid margin irregularity (83.3%) Lid Malapposition (80%) Lashes Lash loss (83.3%) Trichiasis (91.7%) Dystrichiasis (91.7%) Meibomian glands Opaque/scarred (75%) Orifice retroplacement (75%) Marx's line* (72.5%)
			<i>Formix depth measurement (83.3%)</i> <i>Meibography (83.3%)</i>	<i>Photography (91.6%)</i>
Domain 3	Conjunctiva & fornices	Clinical	Conjunctiva Inflammation (90.9%) Limbitis (90.9%) Mucus (81.8%) Follicles (81.8%) Bulbar ulceration (75%) Conjunctival papillae (82.5%) Punctate conjunctival staining (72.5%)	Upper fornix Central depth (81.8%) Number of symblephara (90.9%) Horizontal involvement by symblephara (90.9%) Horizontal involvement by fibrosis (100%) Lower fornix Central depth (90.9%) Number of symblephara (90.9%) Horizontal involvement by symblephara (90.9%) Horizontal involvement by fibrosis (100%) Bulbar keratinization (82.5%) Limitation of motility (72.5%)
			<i>Fornix depth measurement (82.5%)</i>	<i>Photography (82.8%)</i> <i>Immunostaining</i>
Domain 4	Cornea	Clinical	Conjunctivalization (90.9%) Corneal opacification/scarring (81.8%) Corneal opacification/scarring encroaching on the visual axis (81.8%) Endothelial guttata (90.9%)	Sensation Punctate keratopathy Neovascularisation (peripheral involvement) Neovascularization (visual axis) Corneal epithelial edema Corneal stromal edema Localized corneal thinning (no ulceration) Localized corneal ulceration (loss of epithelium) Localized corneal ulceration (thinned cornea) Descemet's folds Graft-host interface changes
				<i>Impression and brush cytology (81.8%)</i> <i>Histological and cell staining (81.8%)</i> <i>Photography (90.9%)</i> <i>Fluorescein staining (90.9%)</i> <i>Lissamine green staining (90.9%)</i> <i>Endothelial cell specular microscopy</i> <i>Aethesiometry</i> <i>Corneal topography</i> <i>Spectral domain OCT</i> <i>Pachymetry</i>
Domain 5	AC and Sclera	Clinical	Anterior Chamber Cells (100%) Flare (75%) Hypopyon (100%) Keratic precipitates (70%) Sclera Anterior scleritis (91.7%) Posterior scleritis (83.3%)	Anterior Chamber Anterior synechiae (peripheral or central) (75%) Iris Iris atrophy (100%) Sclera Scleral thinning (72.5%)
			<i>Laser flare meter (80%)</i>	<i>IOP (91.7%)</i> <i>High resolution anterior segment ultrasound (83.3%)</i>

	WP4 WSB	WP5	WP4 WSB	WP5	WP4 WSB	WP5	Final Category
	Activity (%)	Activity (%)	Damage(%)	Damage(%)	Activity and Damage (%)	Activity and Damage (%)	
Domain 1: Tear Film							
Tear film break up time	66.6	50	8.3	2.5	25	47.5	2nd Activity/Damage
Tear meniscus height	41.7	37.5	25	32.5	33.3	30	2nd Activity/Damage
Tear film debris	58.3	57.5	8.3	10	33.3	32.5	2nd Activity/Damage
Filaments	58.3	57.5	8.3	10	33.3	32.5	2nd Activity/Damage
<i>Tearfilm osmolarity</i>	75		8.3		16.7		Activity
<i>Schirmer's test</i>	16.7	32	41.7	40	41.7	35	2nd Activity/Damage
<i>Fluorescein staining</i>	25	42.5	25	10	50	47.5	2nd Activity/Damage
<i>Lissamine green staining</i>	25	42.5	25	10	50	47.5	2nd Activity/Damage

	WP4 WSB	WP5	WP4 WSB	WP5	WP4 WSB	WP5	Final Category
	Activity (%)	Activity (%)	Damage(%)	Damage(%)	Activity and Damage (%)	Activity and Damage (%)	
Domain 2: Eyelids, lid margins and meibomian glands							
Palpebral aperture	8.3		25		66.7		Unclassified
Completeness of the blink cycle	8.3	5	41.7	32.5	50	57.5	2nd Activity/Damage
Lagophthalmos	8.3	15	50	70	41.7	15	2nd Activity/Damage
Lid malposition	0	7.5	66.7	80	33.3	12.5	Damage
Entropion	8.3		91.7		0		Damage
Ectropion	8.3		91.7		0		Damage
Lid margin irregularity	8.3		83.3		8.3		Damage
Lash Loss	8.3		83.3		8.3		Damage
Trichiasis	8.3		91.7		0		Damage
Dystichiasis	8.3		91.7		0		Damage
Anterior lid margin disease	83		17		0		Activity
M. Gland pouting/plugging	75		8.5		16.7		Activity
M. Gland opaque/scarred	0		75		25		Damage
M. Gland orifice retroplacement	0		75		25		Damage
M. Gland visibility of acini	41.7	42.5	33.3	10	25	47.5	2nd Activity/Damage
Chalazion formation	41.7	42.5	25	47.5	33.3	10	2nd Activity/Damage
Foam	100		0		0		Activity

Quality of secretions	91.7		8.3		0		Activity
M.Gland expressibility	66.7	47.5	0	10	33.3	42.5	2nd Activity/Damage
Lid margin thickening	41.7	42.5	8.3	10	50	47.5	2nd Activity/Damage
Lid margin vascularity	41.7	42.5	8.3	10	50	47.5	2nd Activity/Damage
Position of Marx's line	8.3	5	58.3	72.5	33.3	12.5	Damage
<i>Fornix depth measurement</i>	0		83.3		16.7		Damage
<i>Meibography</i>	0		83.3		16.7		Damage
<i>Photography</i>	8.3		0		91.6		Activity/Damage

	WP4 WSB	WP5	WP4 WSB	WP5	WP4 WSB	WP5	Final Category
	Activity (%)	Activity (%)	Damage(%)	Damage(%)	Activity and Damage (%)	Activity and Damage (%)	
Domain 3: Conjunctiva and fornices							
Conjunctival inflammation	90.9		9.1		0		Activity
Bulbar ulceration	63.3	75	18.2	0	18.2	15	Activity
Limbitis	90.9		9.1		0		Activity
Mucus	81.8		18.2		0		Activity
Bulbar conjunctival keratinisation	9.1	10	72.7	82.5	18.2	7.5	Damage
Conjunctival punctate staining	63.3	72.5	18.2	5	18.2	22.5	Activity
Papillae	63.3	82.5	9.1	12.5	27.4	5	Activity
Follicles	81.8		9.1		9.1		Activity
Upper fornix central depth	9.1		81.8		9.1		Damage
Upper fornix number of symblephara	0		90.9		9.1		Damage
Upper fornix horizontal involvement by symblephara	0		90.9		9.1		Damage

Upper fornix horizontal involvement by fibrosis	0		100		0		Damage
Lower fornix central depth	0		90.9		9.1		Damage
Lower fornix number of symblephara	0		90.9		9.1		Damage
Lower fornix horizontal involvement by symblephara	0		90.9		9.1		Damage
Lower fornix horizontal involvement by fibrosis	0		100		0		Damage
Limitation of mobility	0	2.5	63.3	72.5	36.4	25	Damage
<i>Immunostaining</i>	54.5	27.5	9.1	52.5	36.4	20	2nd Activity/Damage
<i>Fornix depth measurement</i>	0	0	63.3	82.5	36.4	17.5	Damage
<i>Photography</i>	18.2		0		82.8		Activity/Damage

	WP4 WSB	WP5	WP4 WSB	WP5	WP4 WSB	WP5	Final Category
	Activity (%)	Activity (%)	Damage(%)	Damage(%)	Activity and Damage (%)	Activity and Damage (%)	
Domain 4: Cornea							
Corneal sensation	18.2	10	45.5	65	36.4	20	2nd Activity/Damage
Punctate keratopathy	54.5	65	36.4	5	9.1	30	2nd Activity/Damage
Conjunctivalisation	9.1		90.9		0		Damage
Neovascularisation (peripheral)	18.2	7.5	18.2	37.5	63.6	55	2nd Activity/Damage
Neovascularisation (visual axis)	18.2	7.5	18.2	37.5	63.6	55	2nd Activity/Damage
Epithelial oedema	18.2	50	18.2	15	63.6	35	2nd Activity/Damage
Stromal Oedema	9.1	32.5	18.2	17.5	72.7	50	2nd Activity/Damage
Corneal opacification/scarring (peripheral)	0		81.8		18.2		Damage
Opacification/scarring (visual axis)	0		81.8		18.2		Damage
Localised thinning (no ulceration)	9.1	7.5	72.7	55	18.2	37.5	2nd Activity/Damage

Localised ulceration	18.2	60	18.2	10	63.6	30	2nd Activity/Damage
Localised ulcer depth (thinned cornea)	18.2	17.5	27.2	30	54.5	52.5	2nd Activity/Damage
Descemet's folds	27.2	55	9.1	7.5	63.6	37.5	2nd Activity/Damage
Graft-host interface changes	9.1	7.5	27.2	27.5	63.6	65	2nd Activity/Damage
Endothelial guttae	0		90.9		9.1		Damage
<i>Specular microscopy</i>	9.1	0	72.7	55	18.2	45	2nd Activity/Damage
<i>Impression/brush cytology</i>	18.2		0		81.8		Activity/Damage
<i>Histological/cell staining</i>	18.2		0		81.8		Activity/Damage
<i>Anaesthesiometer</i>	0	5	54.5	67.5	45.5	27.5	2nd Activity/Damage
<i>Corneal shape/ thickness measurements</i>	0	5	54.5	47.5	45.5	47.5	2nd Activity/Damage
<i>Optical coherence tomography</i>	9.1	5	18.2	30	72.7	65	2nd Activity/Damage
<i>Pachymetry</i>	9.1	7.5	18.2	27.5	72.7	65	2nd Activity/Damage
<i>Fluorescein staining</i>	9.1		0		90.9		Activity/Damage
<i>Lissamine green staining</i>	9.1		0		90.9		Activity/Damage
<i>Photography</i>	0		9.1		90.9		Activity/Damage

	WP4 WSB	WP5	WP4 WSB	WP5	WP4 WSB	WP5	Final Category
	Activity (%)	Activity (%)	Damage(%)	Damage(%)	Activity and Damage (%)	Activity and Damage (%)	
Domain 5: AC and Sclera							
Keratic Precipitates	58.3	70	0	5	41.7	25	Activity
Anterior chamber cells	100		0		0		Activity
Flare	75		0		25		Activity
Hypopyon	100		0		0		Activity
Anterior synechiae (peripheral/central)	0		75		25		Damage
Posterior synechiae	0	12.5	50	50	50	27.5	2nd Activity/Damage
Iris atrophy	0		100		0		Damage
Intraocular pressure	8.3		0		91.7		Activity/Damage
Anterior Scleritis	91.7		0		8.3		Activity
Posterior scleritis	83.3		0		16.7		Activity
Scleral thinning	0	7.5	70	72.5	30	20	Damage
<i>Anterior segment ultrasonography</i>	16.7		0		83.3		Activity/Damage
<i>Laser flare meter</i>	50	80	0	20	50	0	Activity