Eosinophilic esophagitis: Comparison of clinical, endoscopic and histological scoring systems

Eosinophile Ösophagitis: Vergleich klinischer, endoskopischer und histologischer Scoringsysteme

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ABSTRACT

Background Eosinophilic Esophagitis (EoE) has received increasing attention as a disease entity, and it is now recognized as an important disorder of the Upper Gastrointestinal Tract. Topical corticosteroids (tCS) are effective in clinical-pathological remission induction (RI) and remission maintenance (RM) of active EoE. With scoring systems, such as clinical (SDI), endoscopic (EREFS), and histological (EoEHSS) systems, EoE can be graded, and its disease activity can be assessed.

Objective To discover how closely results within each of the three scoring systems SDI, EREFS, and EoEHSS are correlated between initial diagnosis (ID), RI, and RM, and to determine how well scores from the three systems are intercorrelated at each time point.

Methods Retrospective cohort analysis of patients with active EoE was performed between 2006 and 2020, with follow-up for up to 6 years. SDI, EREFS and EoEHSS scores were recorded at ID, at RI, and in RM. Evaluation employed descriptive statistics, the Friedman test, and Bonferroni-corrected post hoc pairwise comparisons.

Results At RI 29 and at RM 19 EoE patients provided data. Significant correlations were found between EREFS and EoEHSS at RI and in RM. Pairwise comparisons showed significant differences between ID and RI for SDI, for EREFS, and for EoEHSS.

Conclusion The scoring systems tested did not show intercorrelation at ID. Comparison revealed significant differences for SDI, EREFS, and EoEHSS between the systems at ID and RI, but not in RM, during tCS treatment. These results underline the efficacy of tCS (at RI and RM) in the treatment of active EoE.

ZUSAMMENFASSUNG

Hintergrund Die EoE als eigenständige Krankheitsentität gewann in den letzten 2 Jahrzehnten zunehmend an Aufmerksamkeit und etablierte sich seither zu einem wichtigen Krankheitsbild des oberen Gastrointestinaltraktes. Topische Corticosteroide (tCS) sind effektiv in der klinisch-pathologischen Remissionsinduktion (RI) und -erhaltung (RE) bei aktiver EoE. Unter Zuhilfenahme klinischer (SDI), endoskopischer (EREFS) sowie histologischer (EoEHSS) Bewertungssysteme kann die EoE graduiert und ihrer Krankheitsaktivität zugeordnet werden. Wie korrelieren SDI, EREFS und EoEHSS zum Zeitpunkt der Erstdiagnose (ED), nach RI und in RE untereinander sowie im Therapieverlauf?

Methodik Retrospektive Kohortenanalyse von 2006–2020 bei Patienten mit einer aktiven EoE mit Follow-Up-Intervallen bis zu 6 Jahren. Die Erhebung des SDI, EREFS und EoEHSS erfolgte zum Zeitpunkt der ED, nach der RI und in der RE. Die
Auswertung erfolgte mittels deskriptiver Statistik, Friedman-Test und Bonferroni-korrigierter Post-hoc-Vergleiche.

**Ergebnisse** Zum Zeitpunkt der RI konnten bei 29 sowie zur RE 19 EoE-Patienten analysiert werden. Alle EoE-Patienten wurden im Mittel über 13 Wochen bis zur RI bzw. im Mittel über 21 Monate mit tCS bis zur RE behandelt. Signifikante Korrelationen zeigte der EREFS zum EoEHSS bei RI sowie bei RE. Die Paarvergleiche mittels Bonferroni-korrigierter Post-hoc-Tests ergaben signifikante Unterschiede zwischen ED und RI für SDI, EREFS und EoEHSS.

**Schlussfolgerungen** Die Ergebnisse zeigen, dass bei ED die untersuchten Bewertungssysteme zur Bestimmung der Krankheitsaktivität nicht untereinander korrelieren. Der Vergleich der Bewertungssysteme erbringt signifikante Unterschiede zu den Zeitpunkten ED und RI für SDI, EREFS und EoEHSS, jedoch nicht in der RE unter tCS-Therapie. Diese Ergebnisse unterstreichen die Effektivität der tCS in der RI- und RE-Therapie der aktiven EoE.

**Introduction**

Eosinophilic esophagitis (EoE) is a chronic inflammatory disorder of the esophagus, mediated by type 2 T-helper cells. It is manifested clinically by symptoms of esophageal dysfunction and histologically by eosinophil-dominant infiltration of the esophageal mucosa. Reliable diagnosis requires the exclusion of other possible causes of esophageal eosinophilic infiltration[1, 2]. To objectify the clinical symptoms, various scoring systems are available, some validated [3, 4, 5] and some not; the latter include the Straumann Dysphagia Instrument (SDI) [6]. To take account of the highly variable picture obtained by endoscopy, endoscopic diagnosis can be standardized by use of the Endoscopic Reference Score (EREFS), which shows moderate, but adequate, inter- and intra-observer consistency [7, 8]. From the histopathological side, the Eosinophilic Esophagitis Histology Scoring System (EoEHSS) serves as a further instrument of assessment, allowing classification of the extent and severity of histologically determined EoE activity. Furthermore, the EoEHSS allows distinction between untreated and treated EoE patients [9], and it shows high inter-rater and intra-rater consistency [10]. Topically administered corticosteroids (tCSs) are efficacious in both the induction [11, 12, 13] and the maintenance [14, 15, 16, 17] of clinical-pathological remission of EoE. Several studies have shown that the clinical symptoms and the histological activity of EoE are frequently only poorly correlated with one another [18, 19, 20]. In this study we first investigated, separately for each of the three assessment systems SDI, EREFS, and EoEHSS, the extent to which results at the times of initial diagnosis (ID), induction of remission (RI), and maintenance of remission (RM) were correlated (see ▶ Fig. 1A). We then investigated, for each of these time points, how the results from ID, RI, and RM were correlated with one another (▶ Fig. 1B).

**Materials and methods**

**Study design**

This monocentric, retrospective observational study was conducted at the Magdeburg University Hospital over a period from 2006 to 2020. We investigated a cohort of EoE patients, who after ID were treated with tCSs and subsequently achieved RI and then RM, by conducting assessments at the times mentioned. The cohort analysis was approved by the Ethics Committee of Magdeburg University Hospital (AZ R18–18).

EoE was defined according to the current consensus recommendations [1, 2]. Symptoms of esophageal dysfunction and of eosinophilic esophageal infiltration (≥ 15 eosinophil granulocytes per high-power field, EoS/HPF) in at least one out of a maximum of six esophagus biopsies after exclusion of other possible causes of esophageal eosinophilia. Non-adherence to the therapy, changes in diet or lifestyle, inadequately assessable follow-up examinations, or fewer than two evaluable Oesophageo-Gastro-Duodenoscopies (OGDs) likewise resulted in exclusion. Earlier endoscopic interventions such as dilations, or incidental findings of axial hernias, were accepted. Patients of either sex and any age and ethnicity could be included. Clinical-histological remission was defined as a decrease in SDI by ≥ 3 points accompanied by eosinophilic infiltration of the esophageal mucosa of ≤ 15 EoS/HPF.

**Data acquisition**

Demographic data, symptom frequency and severity, any allergic comorbidities, conspicuous endoscopic findings, and relevant disease characteristics for the scoring systems under study were recorded.
Clinical score (SDI)
The SDI [6] was used for assessment of dysphagia. It quantifies and sums the severity and the frequency of dysphagic events, as follows. Frequency: “none” = 0, “once per week” = 1, “several times per week” = 2, “once per day” = 3, “several times per day” = 4. Severity: “swallowing unhindered” = 0, “slight sensation of resistance” = 1, “slight retching with delayed passage” = 2, “short period of obstruction necessitating intervention (e.g., drinking, breathing)” = 3, “longer-lasting period obstruction only removable by vomiting” = 4, “long-lasting complete obstruction requiring endoscopic intervention” = 5. The total possible score is thus 0–9.

Endoscopic score (EREFS)
Data were recorded retrospectively by reference to 60 complete digital images (56 statical, 4 videos) of the esophageal sections (proximal, middle, and distal), 17 uncomplete digital images and to the acquired verbal descriptions of the OGDs. Results were classified according to the validated) modified EREFS criteria [7]: “fixed rings” = 0–3, “strictures” = 0–1, “reddish furrows” = 0–1, “off-white exudates” = 0–2, “mucous membrane oedema” = 0–1, “crepe paper esophagus” = 0–1. The total possible score is thus 0–9.

Histological score (EoEHSS)
Histological assessment represents a central feature in the diagnosis of EoE [19]. The uniformly standardized evaluation of histological sections was performed according to the validated EoEHSS [9]. The graduation was performed in a standardized manner by staining the microscopic preparations with haematoxylin–eosin, and additionally with Giemsa stain, from distal through middle to proximal. For each esophageal section one pair was used for each location: two for proximal, two for central, and two for distal (in total six biopsies).

Severity (grade) and extent (stage) are each assessed by the use of eight histological features, each on a scale from 0 to 3, for total six biopsies). If a feature cannot be assessed in staging biopsy, then the maximum total score is automatically reduced, independently of the other features, by 3 points. To take account of missing features, the score is expressed as a ratio: the score actually obtained is divided by the greatest possible score for the biopsy in question; for example, if all features are assessable and each has the highest possible value, then the result for severity and extent is 24/24 = 1.00, while if only seven of the eight features can be assessed and the total score is 12, then the result is 12/21 = 0.57.

The mean result of a biopsy, for each section of the esophagus, is then calculated as an average of extent score [E] and severity score [S], i.e., (E + S)/2. The mean total score for all three locations is similarly: \( E_{\text{proximal}} + E_{\text{central}} + E_{\text{distal}} + S_{\text{proximal}} + S_{\text{central}} + S_{\text{distal}} \)/6.

Thus, the total overall score lies in the range 0–1, with 1 as the worst possible value.

Statistical analysis
All calculations were performed with GraphPad PRISM (version 7.05). Descriptive statistics were used to characterize the cohorts. For each patient and each point in time (ID, RI and RM) the scores for SDI, EREFS and EoEHSS were determined. The statistical significance of differences for the ordinarily scaled scoring systems were determined for pairs of samples by using the Wilcoxon pair-difference test. For groups of more than two samples the Friedman test was used and a two-factor repeated-measures analysis of variance (ANOVA) was conducted. Post hoc testing with the Dunn–Bonferroni correction and subsequent calculation of the size of the effect was performed. The correlations were assessed by means of Spearman’s coefficient \( r_s = 0.1 \) poor correlation, \( r_s = 0.3 \) middle correlation, \( r_s = 0.5 \) strong correlation. The significance level was set to \( p = 0.05 \).

Results
Patient characteristics
Sixty EoE patients were included according to the in- and exclusion criteria (see \( \text{Fig. 2} \)). Of these, 29 (48 %) were assessable for two time points (ID and RI) and 19 (32 %) for all three time points (ID, RI and RM). Demographic and disease data for the 29 assessable patients are shown in \( \text{Table 1} \). Previously, none of them had responded to a 6–8 week high-dose proton pump inhibitors (PPI) treatment to rule out PPI-responsive esophagitis. On average, these patients were treated for 13 weeks with tCSs in their remission-inducing therapy; 19 (65 %) received budesonide-containing orodispersible tablets (BUD-SKT), 8 (28 %) a budesonide suspension (BUD-S), and 2 (7 %) oral fluticasone (sFLU). As 10 of the patients did not provide follow-up data (i.e., for remission), there remained 19 complete patient-data sets for assessment in RM. The mean duration of RM with tCS was 21 months; 15 (79 %) of the 19 patients were treated with BUD-SKT, 1 (5 %) with BUD-S, and 3 (16 %) with sFLU.

| Assessed for eligibility \((N = 60)\) | Inclusion criteria not met \((N = 6)\) |
|-------------------------------------|--------------------------------------|
| Eligible \((N = 44)\)                | No follow-up \((N = 10)\)            |
| Two complete clinical, endoscopic and histological follow-ups \((N = 29)\) | Elimination diet \((N = 4)\) |
| Lost to follow-up \((N = 11)\)      | No tCS treatment \((N = 11)\)        |

\( \text{Fig. 2} \) Flow diagram for in-/exclusion of patients.
Of the 29 study patients, 22 (76 %) reported atopical comorbidities; allergic rhinitis (15 patients, 68 %) and allergic bronchial asthma (6 patients, 27 %) were the most common of these.

At ID all the 29 study patients reported symptoms of esophageal dysfunction. The most common of these were dysphagic complaints (28 patients, 97 %) and retrosternal pain independent of meal-times (19 patients, 68 %). For 11 patients (38 %), 8 (28 %) and 10 (34 %), ID took place respectively 0–5, 5–10 and 10–15 years after first symptoms. The most common additional endoscopic findings in anamnesis at ID were stenoses (14 patients, 48 %) and axial hernia (10, 34 %). Two patients had earlier undergone balloon dilation of the esophagus. For further details see ▶ Table 1.

### EoEHSS at time points ID, RI, and RM

At ID, a total of 174 biopsies from the proximal, central, and distal location were taken, at RI 174 and at RM 14. Of these, 168 (97 %, from 29 patients) at ID, 172 (99 %, from 29 patients) at RI, and 112 (98 %, from 19 patients) at RM were evaluable. The EoEHSS was 0.41 ± 0.21 at ID and at RI it had decreased with statistical significance to 0.16 ± 0.16 (N = 29, p < 0.001; shown in ▶ Fig. 3A). Dunn–Bonferroni-corrected pairwise comparisons, performed subsequently, showed a consistently low EoEHSS score of 1.8 ± 1.8 (N = 19, p = 0.186; shown in ▶ Fig. 3B).

### Clinical-histological remission

Clinical remission was attained (time point RI) on average after 13 weeks for 24 (83 %) of the 29 EoE patients; 23 (79 %) of these were in histological remission. With a combined criterion for clinical-histological remission, 18 (62 %) patients were identified as being in remission at RI. At time point RM, all the patients were symptom-free and had on average been in clinical remission for 21 months after their treatment. Two patients showed histological recurrence during treatment. Five patients were late responders who only attained histological remission at time point RM, so that – after discounting the patients with missing follow-up examinations after remission-maintenance tCS treatment – histological remission was observed in 17 (90 %) of the 19 patients assessed at RM. Clinical-histological remission at RM was found for 17 (90 %; not the same 17 as above) of these patients.

### Mutual correlations between the ID, RI and RM total scores

When the scores at ID and RI were compared for the 29 EoE patients, no correlation was found. When the time points RI and RM were compared, strong correlation was found between EREFS and EoEHSS: at RI, r = 0.56, p < 0.05 (▶ Fig. 4A) and at RM r = 0.61, p < 0.05 (▶ Fig. 4B).

### Discussion

In this study we investigated the relationships within and between clinical, endoscopic, and histological assessment systems in the...
short and long term. Clinically significant results on the efficacy of tCSs, using the SDI after RI, have been demonstrated in placebo-controlled studies, which showed reductions from 2.7 [12] to 3.39 [6] score points. Our results support these, and in our study the SDI score was even more strongly reduced (by 4.2 points). Regarding the development of the SDI score in maintenance therapy, available data are sparse [21]. Greuter et al. have shown that among 23 EoE patients the mean SDI score decreased by 5.8 points from ID to RM [15]. Our own results confirmed this, with a mean decrease in SDI score of 5.4 points. The EREFS – as an important, established and validated endoscopic scoring system – has often been used as a surrogate marker for determining the usefulness of endoscopy in EoE patients [22, 23, 24], and it has been employed as a secondary endpoint for assessing the efficacy of modern biologicals in treatment of active EoE [25].

Significant differences between endoscopic assessment systems have been observed. The patient cohorts examined have included: children in short- and long-term therapy [14], and adults after RI following treatment with tCSs [13] and/or dietetic therapy [26] and after RM [17]. Because of separate recording of distal and proximal EREFS results [27, 28], and because of omission [29, 30, 31] or alteration [26, 32] of the assessment criteria, study results are difficult to compare. Our investigation showed a significant improvement in EREFS after RI for adults, with scores remaining constant during the maintenance phase. The results of a randomized, placebo-controlled study at RI [13] and of another prospective observational study at RM [15] gave comparable results in respect of improvement in EREFS after medication-based induction or maintenance of remission. In contrast to this situation, for EoEHSS only a few studies have been reported [28, 32, 33]. Frequently, the peak eosinophil count (PEC) has been at the centre of histological investigations in adult [29, 31] and child [34] EoE patients. This results from the current consensus guideline recommendation, according to which eosinophil count is a

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**Fig. 3** Differences in the scoring systems between pairs of time points: **A** ID and RI, **B** RI and RM.

**Fig. 4** Correlation between EREFS and EoEHSS: **A** at time point RI, **B** at time point RM.
order to obtain clinical, endoscopic, and histological findings from EoE patients; this may have led to some distortion of results through selection bias. Some of the study data were acquired before the publication of the clinical [6], endoscopic [7], and histopathological [9] scoring systems, so that distortions could have arisen through incorrect classification (information bias). Our most recent data set from 2006 was acquired before the introduction of SDI (in 2010) and also before DSQ [3] and EEsAI PRO [5] were established (in 2013 and 2014 respectively). Our clinical analysis was made in detail and was documented; it was aimed specifically at recording the intensity and frequency of dysphagia, at discerning possibilities in the daily course of the patients’ disorder, and at determining the types of meal that gave rise to dysphagic events. This enabled us to determine these retrospectively from our data sets before the introduction of SDI. Both DSQ and EEsAI are less value in everyday practice, i.e., outside controlled clinical studies, on account of their complexity. Both of these assessment instruments need to be applied almost daily over a period of 14 [37] or respectively 7 days [5]. Moreover, EEsAI is recommended mainly for prospective studies [5].

Nonetheless, the strengths of our study lie in the long observation and treatment periods covered during RM, which alongside the clinical assessment was assessed by EREFS and EoEHSS, both of which were developed and validated specially for EoE. The clinical and endoscopic scoring was performed by an EoE expert (U.v.A.) and the data for them were acquired under her supervision. EoEHSS was assessed by an experienced pathologist (P.C.). According to current consensus conditions, a histological remission is defined as one showing <15 Eos/HPF. In the literature, the use of stricter definitions has been reported; nonetheless, the limit values reported in this paper have generally been found.

Regarding clinical-histological remission, reports are not in uniform agreement. To the best of our knowledge, no long-term results, based on a combination of clinical and histological assessment tools, have so far been published. We therefore use in this work the definition of clinical-histological remission as used in a published study and provide one of the first reports of results of clinical-histological remission in the short and especially in the long term [13].

Our results show that a tCS-based induction therapy is – both clinically and histologically – highly effective and that this efficacy can be maintained during a long-term tCS therapy. This result demonstrates that adult EoE patients profit from tCS in the long term, as also found by Greuter et al. [30]. A further important finding in this study is that symptoms and histological activity in the maintenance phase are not intercorrelated. This has consequences for the surveillance of EoE patients; a regular observation of histological activity during maintenance therapy is indicated, in the interest of avoiding long-term complications.

In summary, our study has shown that results from the clinical, endoscopic and histological scoring systems were not intercorrelated at ID. They differed significantly after induction of remission, but not in therapy maintenance. During RM all the systems showed stable, low scores. These results underline the efficacy of tCS in both the short-term and the long-term treatment of active EoE.

Conflict of Interest

The authors declare that they have no conflict of interest.
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