Summary. Aims: The purpose of this study was to draw up a TMJ disc histopathological score that is a semi-quantitative transcription of the entire spectrum of TMJ degenerative diseases related to changes in disc tissue, and then validate the proposed grading, in order to contribute to a standardized histopathological diagnosis. Methods: Sections from sixty two temporomandibular joint disc specimens affected by tissue degenerative changes and stained with Hematoxylin & Eosin (H&E) were collected from among those found in the archives of the Department of Dentistry at Catania University. Specimens, included anterior, intermediate and posterior disc bands. Based on a literature search regarding the most frequent histopathological changes detected in TMJ disc tissue a grading score was designed. This score takes into account pathological disc tissue transformation i.e. collagen bundles, non-specific degenerative changes and the presence of blood vessels. This grading system results in a score ranging from 0 up to 8 for heavily degenerated disc tissue. Two observers performed the assessment of the TMJ disc H&E stained sections. Each specimen was scored twice by each observer after a minimum interval of 1 week. Results: The average TMJ disc degeneration score of the TMJ disc sample was 3.89±1.37. There was an almost perfect agreement, between the two observers, and degeneration scores from the two observers were highly correlated. Conclusions: The introduction of this validated degeneration score system may be of major importance for future research and collaboration between different centers in order to improve knowledge in TMJ disc histopathology.

Key words: TMJ disc, Degeneration, Histopathology, Grading, Score

Introduction

The human temporomandibular joint (TMJ) articular disc consists of a compact fibrous tissue, in which rare fibroblast-like cells can be found among the compact and regularly arranged collagen fibres (Caltabiano et al., 1991; Marchetti et al., 1999; Leonardi et al., 2001, 2002b). This disc separates the articulating surface of the glenoid fossa and eminence from that of the mandibular condyle, dividing the joint cavity into two compartments (Kapila et al., 1995). The functional importance of the disc is highlighted in various TMJ arthropathies especially, when aberrations to its position, morphology, and composition are accompanied by symptoms, which range from joint crepitus and clicking to a limited range of mandibular movements and pain (Haskin et al., 1995; Sato et al., 1997). Several studies have also demonstrated that, subsequent to TMJ arthropathies, the discal tissue presents degenerative changes, which are the result of maladaptation to increased joint loading (Kirk, 1990; Haskin et al., 1995). These degenerative changes are influenced by the type and degree of disc displacement, in that the more it is displaced the more the severe histopathological changes can be observed (Carlsson et al., 1967; Bean et al., 1977; Leonardi et al., 2007).

Several investigations have been published describing the histopathological findings of the degenerated TMJ disc and their relationship with cytokine, growth factors and biomolecular mediators (Paegle et al., 2003; Hamada et al., 2008; Leonardi et al., 2000, 2008; Matsumoto et al., 2008), however, so far, no molecular basis has been identified as a cause for this
A TMJ disc degeneration grading score

degenerative disorder (Luder, 2002). Therefore, the pathology and/or mechanism regarding the development and progression of disc degeneratve change, is still not fully understood (Kang et al., 2006). Besides the difficulties encountered by previous investigation in the study of TMJ disease already described (Haskin et al., 1995), there is also the lack of a histopathological score system. Consequently, data analysis has been limited to descriptive or subjective estimates (Paegle et al., 2002). The lack of this score system has represented one of the major obstacles when trying to understand tissue changes at a molecular level and inter-relate the findings from different research teams. To the knowledge of the authors, no study has been devoted to developing a histopathological score to evaluate TMJ disc degeneration and until now no relative literature has been found.

The purpose of this study was to draw up a TMJ disc histopathological score that is a semi-quantitative transcription of the entire spectrum of TMJ disc degenerative diseases related to changes in disc tissue, and then validate the proposed grading, in order to contribute to a standardized histopathological diagnosis.

**Materials and methods**

**Specimens**

Sections from sixty-two temporomandibular joint disc specimens from among those stained with Hematoxylin & Eosin (H&E) and affected by tissue degenerative changes were collected from previously studied materials used in our earlier investigations (Leonardi et al., 2000a,b, 2001, 2002a,b, 2003, 2007, 2008). Eight virtually unaffected human TMJ discs (control sample) were studied along with the pathological material. Approval by the Ethics committee and informed consent were obtained for each patient before tissue collection.

The discs affected by degenerative changes (32 from the right TMJ and 30 from the left) were taken from 51 female and 11 male patients, the mean age value of all patients was 37.2±9.4 years. All the selected discs, extirpated during surgery as previously described (Widmark et al., 1997) were not mutilated when removed or incompletely excised or removed in fragments. Every disc was macroscopically deformed. None of them had a normal biconcave shape.

The autopsy specimens (control sample) which were collected and their findings thoroughly examined so as to decide whether or not to include them in the TMJ degeneration score (Pritzker, 1977; Pritzker et al., 2006; Roberts et al., 1989). According to these aforementioned premises, a grading score was designed.

This score takes into account three different pathological disc tissue transformation parameters (collagen bundles, non-specific degenerative changes and the presence of blood vessels).

As regards collagen bundles, in normal human TMJ the disc is composed of superimposed layers of compact bundles of collagen (mainly type I) fibres disposed sagitally and obliquely and alternated with each other (Minarelli and Liberti, 1997). Moreover the organization and function of the collagen fiber system in the human temporomandibular joint disc and its attachments are different according to the disc regions, as demonstrated under the polarizing microscope (Scapino et al., 2006).

In the degenerated disc sample, collagen bundle fragmentation is seen first, while, in the even more degenerated discs tears appear. These latter ones are characterized by defects following the orientation of the collagen fibers bundles. In heavily degenerated discs, splitting can be appreciated. This is defined as a disc fissure of the disc, lined with a layer of cells (Fig. 1). The lining of cells on the inner surface of the split differentiates it from an artifact produced during specimen sectioning (Kurita et al., 1989).

As far as the presence of blood vessels in youngsters and adults is concerned, all the central, posterior and anterior portions of the disc, excluding the posterior (bilaminar zone) and anterior disc attachments are avascular (Yoshida et al., 1998; Paegle et al., 2002). In contrast, a high density of blood vessels is described in disc specimens taken at any stage of internal derangement of the joint (Yoshida et al., 1999; Paegle et al., 2002; Leonardi et al., 2003). Newly formed patients and autopsy specimen included the anterior and intermediate bands, as well as the posterior disc band.

**Histopathological characteristics of the TMJ disc degeneration score**

In order to establish principles and proposed standards for the scoring system three items were considered: 1) it has to be simple and reproducible (simplicity); 2) The system should be equally useful for the assessment of both clinical and experimental TMJ disc histopathology models (utility); 3) The histopathology grading system should be capable of being harmonized, eventually, with histological assessment systems from other joints (comparability) (Pritzker et al., 2006). To create the score, a literature search on human TMJ discs with normal appearance and degenerative changes, was carried out and pertinent articles were retrieved. The more frequent histopathological findings were collected, as well those for normal disc appearance. Other grading score systems for degenerative joint changes other than TMJ were also collected and their findings thoroughly examined so as to decide whether or not to include them in the TMJ degeneration score (Pritzker, 1977; Pritzker et al., 2006; Roberts et al., 1989). According to these aforementioned premises, a grading score was designed.

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Fig. 1. Hematoxylin & Eosin stained sections of TMJ discs affected by different degrees of degeneration. **a)** Fragmentation of collagen bundles (fc); **b)** tear (t) and **c)** splitting (s). Scale bars: a, b, 0.23 mm; c, 0.19 mm.

Fig. 2. Hematoxylin & Eosin stained sections of TMJ discs affected by different degrees of degeneration. **a)** an ingrowth of capillaries can be appreciated in a mild degenerated disc tissue; **b)** capillaries, arterioles and venules are detected in severe damaged disc. Scale bar: a, 0.27 mm; b, 0.11 mm.
capillaries are observed at first, and arteries and venules become evident as disc degeneration progresses (fig.2).

Non specific degenerative changes (fatty degeneration, calcified areas, hyalinization and chondroid metaplasia) were also included in the degeneration score, as they are very frequently reported in the literature (Kurita et al., 1989; McKay et al., 1992; Luder, 2002; Leonardi et al., 2007).

This grading system results in a score ranging from 0 up to 8 for badly degenerated disc tissue. The parameter values are summarized and interpreted according to Table 1. A score from 0 to 3 was allocated in two of the three categories (Table 1) while the third category contains the 0 to 2 score. These were added together to give a final score out of 8. The three groups were: no/minimal degeneration (degeneration score of 0 or 1), moderate degeneration (score 2-3), and severe degeneration (score 4-8).

### Sample scoring

All samples were presented blinded with respect to tissue origin and examined in random order under direct light microscopy by each of the observers. First, each H&E stained section was examined at a 40x magnification to obtain information about the entire section and to determine the extent of lesions present. Secondly, the entire section was examined at a 100x magnification to give more detail of the disc’s appearance. Two observers A and B, with at least 10 years experience in TMJ disc tissue histopathological evaluation, performed the assessment of the TMJ disc H&E stained sections. Each specimen was scored twice by each observer after a minimum interval of 1 week. The mean average of the two scores from each observer was used in whatever analysis was performed.

### Statistical analysis

The score values of this score are on an ordinal scale of measurement. Median values and standard deviations were chosen for descriptive statistics. Cohen’s kappa was calculated to determine an agreement between the two expert diagnosis. Pearson’s correlation coefficient was used to investigate the correlation between the grading results of the two observers. All data were analyzed with the SPSS program (SPSS® release 16.0, Chicago, IL, USA).

### Results

The average TMJ disc degeneration score (sum of the scores obtained by the two observers) of the TMJ disc sample was 3.89±1.37. The mean scores for each observer were very close respectively, 3.90±0.18 and 3.88±0.16. The Cohen’s kappa (percentage of agreement) value obtained was 0.95, which is commonly interpreted as an almost perfect agreement. Also, the Pearson correlation coefficient (r=0.971) showed that scores obtained by independent observers were statistically significant.

### Discussion

Several articles have described the histopathological anatomy of human TMJ discs (Axelsson et al., 1987; Haskin et al., 1995; Holmlund, 2007; Leonardi et al., 2007, 2008; Matsumoto et al., 2008), but a histopathological degeneration grading score system has not yet been published, even though it is needed. A universally accepted system for the histopathologic grading of degenerative TMJ disc pathological findings is an important prerequisite for the generation of histological, histochemical, and immunohistochemical data in TMJ disc research, data that can be compared by different laboratories. So far, it has been claimed that maintaining a consistent observer performance in the evaluation of TMJ disc histopathology is difficult (Kurita et al., 1989) due to the lack of a validated score system (Hall et al., 1984). As matter of fact, some grading systems have been proposed to quantify the degree of degeneration of intervertebral disc (IVD) (Pritzker, 1977; Roberts et al., 1989), but these cannot be used for the TMJ disc as its anatomy clearly differs from IVD.

Therefore, the present study aimed at introducing, for the first time, a scoring system for TMJ disc...
degenerative pathology, in order to correlate disc histopathology clinical, medical and research findings, and so gain a better understanding of this pathology. The TMJ disc degeneration score proposed here focuses on the above histopathological changes because these can be easily identified in routine H&E-stained slides. This proposed system uses an 8 point score system based on a combination of tissue architectural changes.

Cell types and their distribution were not accounted for in our scoring system, as TMJ disc cells are poorly characterized and frequently referred to by various names, (Kapila et al., 1995). Moreover, it seems that there is a heterogeneously distributed subpopulation (Leonardi et al., 2002b; Detamore et al., 2006), whose percentage has not yet been well established in normal or degenerated tissue. In fact, due to the lack of investigation concerning disc cells subpopulation, no universally accepted and objective findings on a morphological basis have been reported in literature. The only morphometric study on degenerated TMJ discs describes an increase of chondrocyte-like cells on ID discs compared to the control, but this was not statistically significant. Another issue with disc cell subpopulation is that their exact quantification requires time and a grid mounted onto the eye piece of an optical microscope (Kurita et al., 1989; Paegle et al., 2002) which is rather time-consuming (Gynther et al., 1998). As far as inflammatory cells are concerned, no significant inflammation has ever been reported in human TMJ disc specimens (Hall et al., 1984; Kurita et al., 1989) contrary to the synovial living (Gynther et al., 1998), thus no score was provided in our grading system, as cellular signs inflammation are not a characteristic of TMJ disc degeneration.

On the other hand, three items were taken into account during the design of our disc degeneration score system, i.e. collagen bundle integrity or damage, degree of neo-vascularization and non specific degenerative changes: i.e. fatty degeneration, calcified areas, hyalinization (fibrosis), and chondroid metaplasia, on the basis that they have been the most frequent findings reported by studies on TMJ disc histopathology and can be easily detected on H & E stained section (Kurita et al., 1989; Leonardi et al., 2001, 2003). Immunohistochemistry which may improve the accuracy of the histologic diagnosis, was not taken into consideration in our score as it cannot be used regularly due to its relatively high cost.

The score presented here shows a very good interobserver reliability, as the percentage of agreement between the two observers was almost perfect and data were highly consistent. The introduction of the present validated degeneration score system may be of major importance for future research collaboration between different centers so that consensus exists regarding definition and classifications. Moreover, findings from several investigations can be easily compared using this score, with the hope of improving our knowledge of TMJ disc degeneration etiology.

References


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