3237 Relation of Salivary Elastase with Periodontal Conditions and Treatment Response

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Saliva contains egress from gingival crevice, and thus saliva may be used as a medium for overall assessment of periodontal conditions. Objective: This study was to determine whether salivary granulocyte elastase activity could be used in the assessment of periodontal conditions and treatment response. Methods: The participants were 32 nonsmoking patients with untreated chronic periodontitis in a clinical trial and 12 periodontally healthy subjects. Patients were randomly assigned to test (n=16) and control (n=16) groups, both groups receiving scaling and root debridement, while the test group also used Chlorhexidine. Full-mouth bleeding on probing and probing depth (PD) were recorded by the Floride Probe® at baseline, 1, 3 and 6 months post-treatment. Stimulated whole saliva was collected by a standard spitting method immediately prior to clinical examination. Salivary granulocyte elastase was analyzed with a granulocyte-specific substrate (pGluProVal-pNA), and the maximal rate of elastase activity (MR-EA, mAbs/min) was calculated. The statistical methods used included ANOVA, repeated measures ANOVA and correlation analysis with two-tailed significance testing. Results: Baseline salivary MR-EA in patients (19.136±2.704) was significantly higher than in healthy subjects (0.006±0.001, p<0.001). No significant difference was found in baseline clinical parameters and MR-EA between test and control groups. Clinical parameters significantly improved with a concomitant reduction in salivary MR-EA (p<0.001) post-treatment, and no difference was found between groups. At baseline, MR-EA was positively correlated with mean PD (r=0.43, p=0.014) and with % sites PD \geq 5.0mm (r=0.42, p=0.018). The change of MR-EA was positively correlated with the concomitant change in PD at 3 months (r=0.39, p=0.028) and 6 months (r=0.37, p=0.044). Conclusion: The study suggests that salivary granulocyte elastase activity seems to reflect overall periodontal conditions, and might serve as a biochemical test in the assessment of periodontal conditions and treatment responses. Supported by the Hong Kong Research Grant Council (RGC, HKU 7287/97M and 7310/00M).

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