Oral Rehabilitation with Osseointegrated Implants in a Patient with Oromandibular Dystonia with Blepharospasm (Brueghel’s Syndrome): A Patient History

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Oromandibular dystonia with blepharospasm (also known as Brueghel’s syndrome, Meige’s syndrome, or idiopathic orofacial dystonia) is characterized by intense and involuntary spasms of the orofacial muscles, with a frequent loss of teeth and occlusal alterations that worsen the dystonic manifestations and cause mucosal lesions that can lead to complete edentulism. The history of a patient with oromandibular dystonia who was rehabilitated with mandibular overdentures supported by endosteal implants is presented. Oral rehabilitation with implant-supported overdentures improved the situation, despite serious problems with instability. Mandibular overdentures supported by endosteal implants were satisfactorily used to re-establish occlusion, ensuring prosthetic stability and improving the dynamics of the masticatory muscles. (INT J ORAL MAXILLOFAC IMPLANTS 2001;16:115–117)

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Oromandibular dystonia with blepharospasm (also known as Brueghel’s syndrome, Meige’s syndrome, or idiopathic orofacial dystonia) is an adult disorder (the usual age of onset is over 50 years) with an unknown etiology characterized mainly by prolonged symmetric dystonic contractions of the orofacial muscles. It occurs more frequently in women. Patients with this disorder exhibit bizarre facial contortions and are frequently disabled. In Brueghel’s syndrome, the orofacial dystonic movements are characteristically slow and intense and last for 20 to 30 seconds. The movements can be triggered by emotional stress, reading, watching television, fatigue, or speech.¹ Both blepharospasm and orofacial dystonia may occur in patients with known diseases of the basal ganglia, such as those that cause symptomatic torsion dystonia, or are subjected to treatment with antipsychotic agents. The condition may also present as isolated dystonia in middle or late adult life in otherwise normal individuals.² The patient presented here demonstrated spasms and contractions of the facial, lingual, and masticatory muscles typical of idiopathic oromandibular dystonia.²

Blepharospasm consists of involuntary shutting of the eyelids and is characterized by the presence of tonic bilateral and more or less symmetric and arrhythmic contractions of the orbital muscles.³ Oromandibular dystonia in turn consists of prolonged spasmodic contractions of the muscles of the mouth and jaws; these dystonic movements appear to be distinct from the chewing, lip-smacking, and tongue-rolling choreiform movements seen in more familiar orofacial dyskinesia.²

The present article describes a patient with oromandibular dystonia who was rehabilitated with mandibular overdentures supported by endosteal implants.

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A 67-year-old woman who had suffered oromandibular dystonia for the previous 2 years presented with repeated traumatic ulcerations of the lower lip. There were no other personal or family antecedents of interest. The dystonia was being treated with clonazepam (6 mg/day) and amitriptyline (75 mg/day). She presented small, involuntary, continuous positional dystonic movements of the mandible, lips, and tongue (Fig 1). Oral examination revealed numerous edentulous spaces in both jaws, which prevented stable occlusion and caused multiple ulcerations of the mucosa of the alveolar processes (Fig 2).

The remaining teeth were removed, with alveoloplasty of the maxillary and mandibular processes. After a 4-week healing period, 2 cylindric 3i implants (Implant Innovations, Inc, West Palm Beach, FL) were surgically positioned within the chin region. The postoperative course was uneventful. Four months later, a mandibular overdenture supported by 2 endosteal implants, as well as a conventional complete maxillary denture, were positioned (Fig 3).

The patient reported that the oral rehabilitation improved mastication and speech, although the oromandibular spasms progressed. Associated blepharospasm developed 2 years later, and although botulin toxin infiltration afforded initial improvement, the movements subsequently became refractory to treatment. Five years after placement of the implants, the latter show osseointegration with prosthesis stability, and the patient maintains adequate occlusion.

The significance of dental alterations in dystonia is not entirely clear, although tooth loss is thought to cause a decrease in afferent proprioceptive impulses and an alteration of the modulation mechanisms of the central nervous system, which may trigger or worsen the dystonia.4–6 According to Fletcher et al,4 dystonia may be precipitated by peripheral injuries. In some patients, oromandibular dystonia has been seen to develop after dental treatment, and in blepharospasm, a history of preceding local ocular disease has been documented in about 12% of cases. Similar observations apply to tardive dyskinesia, which is facilitated or exacerbated by edentulism7; occlusal rehabilitation may thus afford improvement of anomalous movements in these patients.8
Management of the dental aspects of the condition should seek to preserve as much tissue as possible. Prosthetic rehabilitation is important in helping to ensure a normal occlusion that is capable of securing a stable mandibular position and adequate muscle rest without worsening the dystonic movements. In this sense, complete denture rehabilitation poses serious difficulties, especially in the mandible, because of the marked instability involved. This problem can be resolved by placing mandibular implants that facilitate adequate prosthetic stability and the restoration of occlusion.

In the present situation, as in the patient with tardive dyskinesia reported by Kelleher et al., partial tooth loss and occlusal alterations preceded the appearance of involuntary movements, and therefore it was decided to utilize a mandibular overdenture supported by endosteal implants. Both the surgical phase and the prosthetic rehabilitation period were uneventful and, in the 5 years of follow-up, the patient slowly stabilized her oromandibular dystonic movements, with improved function and esthetic results. In contrast, the blepharospasm worsened. A review of the literature yielded no descriptions of patients with idiopathic orofacial dystonia rehabilitated with implants.

REFERENCES