# Treatment with Implant-Supported Fixed Dental Prostheses in Patients with Congenital and Acquired Neurologic Disabilities: A Prospective Study

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Purpose: To study the medium- to long-term outcome of implant treatment in patients with neurologic disabilities. *Materials and Methods:* Twenty-seven patients with different disabilities and in need of prosthodontic treatment were treated with various implant-supported prostheses. Altogether, 88 threaded titanium implants were placed. General anesthesia was used in 21 patients and local anesthesia in 6 patients. Patients were recalled every 3 months by a dental hygienist and annually by a prosthodontist. **Results:** Five of the original 27 patients died during the 5- to 10-year follow-up period (mean, 7.2 years), but the remaining 22 patients with 70 implants could be clinically examined at the final follow-up. Twelve implants (14%) were lost, 3 before loading and 9 after insertion of the implant-supported fixed prostheses. The cumulative survival rate for placed implants was 85.8% after 10 years. Perimucositis was diagnosed in 10 patients and for 14 of the 70 implants. Three of the 15 patients with measurable radiographs and 4 implants were diagnosed with peri-implantitis. Several prosthodontic complications occurred, from minor and easily correctable to severe and requiring retreatment. **Conclusions:** Patients with different neurologic disabilities present more problems during implant treatment and maintenance compared with healthy patients. Nevertheless, it was possible to carry out treatment, and outcomes were relatively favorable. The results indicate that implant treatment can be a valid option in oral rehabilitation of patients with neurologic disabilities. Int J Prosthodont 2013;33:517-524. doi: 10.11607/ijp.3511

The successful use of dental implants for the treatment of completely and partially edentulous patients is well documented in numerous studies and books.<sup>1-6</sup> Survival rates of  $\leq$  95% have been found in systematic reviews of studies with follow-ups of at least 5 years.<sup>7,8</sup> Reports of implant treatment in individuals with rare diseases are limited and mainly consist of anecdotal reports and case presentations.<sup>9,10</sup>

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A search in PubMed for articles on oral rehabilitation using dental implants in patients with rare disorders up to July 2012, supplemented with a manual search of references in the retrieved papers, revealed the continuing publication of case reports but few controlled studies.<sup>11</sup> One study described the relatively favorable outcome 3 to 113 months after implant loading in 18 patients with various degrees of physical and mental impediments.<sup>12</sup> A case series study of 24 special care patients demonstrated a cumulative survival rate after 5 years of 93.4%; the survival rate of the prostheses was 100%.13 A retrospective study of 1 to 16 years (mean, 4.6  $\pm$  3.1 years) after implant treatment in 61 patients with severe epilepsy and additional motor and/or intellectual impairments reported good results.<sup>14</sup> All three retrospective studies concluded that dental implants offer a viable option for special care patients with various disabilities.

A prospective study of dental implant treatment for individuals with neurologic disabilities described the early experience as containing both possibilities and difficulties.<sup>15</sup> This study concluded that it was possible to carry out treatment with relatively good

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| Patient         | Sex | Age at<br>delivery (v) | Diagnosis                                      | Prosthodontic treatment   | Implant location*                            | Function (v) |
|-----------------|-----|------------------------|--|---|--|--------------|
| 1†              | Μ   | 48                     | Down syndrome, epilepsy                        | Tooth-supported fixed prosthesis in maxilla,<br>ISFP in partially edentulous mandible | 46, 44, 43                                   | 6            |
| 2               | F   | 41                     | Mental retardation                             | Single-tooth implant  | 11   | 10           |
| 3               | F   | 51                     | Mental retardation, epilepsy                   | Single-tooth implant  | 11, 21                                       | 9.5          |
| 4               | F   | 55                     | Mental retardation                             | ISFP in completely edentulous mandible  | 44, 43, 31, 33, 34                           | 8            |
| 5               | Μ   | 53                     | Fragile X                                      | ISFP in partially edentulous maxilla  | 16, 14, 13                                   | 9.5          |
| 6               | М   | 24                     | Autistic syndrome, epilepsy                    | Single-tooth implant  | 21   | 9.5          |
| 7               | F   | 46                     | Down syndrome                                  | Single-tooth implant  | 21   | 9            |
| 8               | F   | 55                     | Mental retardation, epilepsy, schizophrenia    | ISFP in completely edentulous maxilla   | 15, 13, 11,<br>21, 23, 25                    | 8            |
| 9†              | F   | 36                     | Retts syndrome, epilepsy                       | Single-tooth implant  | 21   | 2            |
| 10              | F   | 53                     | Mental retardation, epilepsy                   | ISFP in partially edentulous maxilla  | 13, 11, 23                                   | 8            |
| 11              | F   | 19                     | Mental retardation,<br>multiple disabilities   | ISFP in partially edentulous maxilla  | 12, 22                                       | 8            |
| 12              | М   | 50                     | Mental retardation                             | Single-tooth implant  | 24   | 8            |
| 13              | F   | 53                     | Mental retardation, epilepsy                   | Single-tooth implant  | 24   | 8            |
| 14              | М   | 33                     | Mental retardation                             | Single-tooth implant  | 21   | 8.5          |
| 15              | F   | 35                     | Severe mental retardation                      | Single-tooth implant  | 21   | 7.5          |
| 16              | Μ   | 58                     | Dystrofia myotonica                            | ISFP in completely edentulous mandible  | 44, 43, 41, 33, 34                           | 7.5          |
| 17              | Μ   | 41                     | Aspberger syndrome                             | ISFP in completely edentulous mandible  | 45, 43, 31, 33, 34                           | 7.5          |
| 18              | Μ   | 56                     | Mental retardation                             | ISFP in partially edentulous maxilla  | 14, 15, 16                                   | 7            |
| 19              | Μ   | 47                     | Mental retardation, epilepsy, short in stature | ISFP in completely edentulous maxilla and mandible                                    | 15, 13, 11, 21, 23, 25<br>45, 43, 31, 33, 34 | 7            |
| 20              | М   | 58                     | Mental retardation                             | ISFP in completely edentulous mandible  | 45, 43, 41, 33, 34                           | 6.5          |
| 21†             | М   | 54                     | Down syndrome                                  | Single-tooth implant  | 11   | Not loaded   |
| 22              | М   | 19                     | Down syndrome                                  | Single-tooth implant  | 13, 23                                       | 6.5          |
| 23†             | Μ   | 80                     | Mental retardation                             | Implant-supported overdenture in com-<br>pletely edentulous mandible                  | 43, 31, 33                                   | 3            |
| 24              | Μ   | 22                     | Cerebral palsy,<br>mental retardation          | ISFP in partially edentulous maxilla  | 13, 11, 21, 22                               | 5.5          |
| 25              | F   | 52                     | Prader Willi                                   | ISFP in partially edentulous maxilla<br>ISFP in completely edentulous mandible        | 12, 21, 23, 25<br>45, 43, 31, 33, 34         | 5.5          |
| 26 <sup>†</sup> | F   | 55                     | Dystrofia myotonica                            | ISFP in partially edentulous maxilla  | 24, 25                                       | 2            |
| 27              | F   | 49                     | Mental retardation                             | 2 ISFPs in partially edentulous maxilla   | 14, 15<br>12, 11, 21, 23                     | 5            |

#### Table 1 Details of the Study Population and Treatments

ISFP = implant-supported fixed prosthesis.

\*FDI system. \*Deceased.

<sup>T</sup>Deceased.

results despite the fact that all patients had severe disabilities including mental retardation and different degrees of autistic behavior. This short-term report dealt with a group of 14 patients with a follow-up of 1 year or more for only 6 of them. The present paper will cover the entire group of patients followed for > 5 years. The purpose of this prospective study was to provide a medium- to long-term report of implant treatment in patients with neurologic disabilities.

# **Materials and Methods**

The 27 patients had been referred to the National Orofacial Resource Centre with different disabilities, mainly neurologic disorders, causing various orofacial dysfunction problems (Table 1). They were completely or partially edentulous and considered suitable for prosthodontic treatment with implants. The background of the study and details of the early

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| Length (mm) | Ma<br>RP + NP | TiUnite<br>Mk III + IV<br>RP | Replace<br>Select<br>RP + NP | Speedy Groovy<br>tapered<br>RP | Lost before<br>loading | Lost after<br>loading |
|-------------|---------------|------------------------------|------------------------------|--------------------------------|------------------------|-----------------------|
| 7           |               | 2                            |                              |                                | 1 (TiU)                | 1 (TiU)               |
| 8.5         |               | 2                            |                              |                                |                        |                       |
| 10          | 1             | 3                            |                              |                                |                        | 1 (TiU)               |
| 11.5        |               | 3                            |                              |                                |                        | 1 (TiU)               |
| 13          | 4             | 12                           | 2                            |                                |                        |                       |
| 15          | 6             | 41                           |                              | 2                              | 2 (Ma)                 | 1 (Ma) 5 (TiU)        |
| 16          |               |                              | 4                            |                                |                        |                       |
| 18          | 2             | 4                            |                              |                                |                        |                       |
| Total       | 13            | 67                           | 6                            | 2                              | 3                      | 9                     |

 Table 2
 Distribution of Implants Placed and Lost

Ma = machined; RP = regular platform; NP = narrow platform; TiU = TiUnite.

experiences of the first 14 treated patients have been previously published.<sup>15</sup> The Swedish Parliament decided on "necessary dental care" in 1998 to increase financial support and service given to persons with disabilities who were dependent on nursing staff.

The inclusion criterion for the study population was patients with different neurologic disorders, either congenital or acquired, including mental retardation and with different degrees of autistic behavior. They were referred to the clinic because they were judged to need treatment with implant-supported prostheses. Patients were excluded if they themselves or their legal guardian or personal caregiver had negative views of prosthodontic treatment with implants. Despite sometimes serious disabilities, all patients in this study lived in their own homes or in residential housing with the help of personal caregivers.

The study was approved by the Regional Ethics Committee in Region Västra Götaland, Sweden, Dnr 151–11, on April 20, 2011.

The prospective study was planned for 30 patients and completed for 27 patients referred to the clinic between 2000 and 2006.

The prosthodontist first examined all patients and planned their treatments. Before implant placement, the prosthodontist once again examined all patients together with the oral surgeon and discussed the treatment concept. Because of behavior problems among some of these patients, panoramic and other relevant radiographs were not always available at the time of examination and had to be taken under general anesthesia at the time of implant placement.

Oral surgeons made all implant placements except in one patient in whom the first author placed the implants. The first author did most of the abutment operations and all prosthodontic treatment. The implants were placed under general anesthesia in 21 patients and with local anesthesia in 6 patients.

In the first 17 patients, a very strict surgical protocol for Brånemark implant placement (Nobel Biocare) was used. It included a two-stage procedure using antibiotics pre- and postoperatively, chlorhexidine for mouth rinse, analgesic when needed, and frequent postoperative controls. A two-stage procedure was also used in 4 more patients with limited bone quantity and quality. In 3 patients with an edentulous mandible, a one-stage procedure was used with loading of the implants within 2 months. In 2 more patients, a one-stage procedure was used but with a healing period of 6 months. One patient was treated with immediate loading (Nobel Guide).

The types and lengths of implants placed varied, but the great majority of them were 15- or 13-mm long TiUnite Mark III and IV with a regular platform (Rp, Nobel Biocare) (Table 2).

Radiographic examination was planned after placement of the implant and at control visits up to the final follow-up. Because of severe problems due to the physical and mental condition of some patients, radiographs of acceptable quality could be obtained from only 15 of the 27 patients, including 52 implants. A trained radiologist made all estimations of periimplant bone loss by measuring changes in exposed threads from the radiographs taken at placement of the implant-supported restoration up to the last examination.

The following variables were recorded: presence or absence of parafunction, tooth wear, implant failure, implant bone level, surgical and/or prosthetic complications, as well as the design of the prostheses and number of visits. The patients and/or their caregivers were asked questions on patient experience of the implant treatment and satisfaction with the restorations. The anatomical form, surface, color, and fit of the fixed prostheses were recorded according to the California Dental Association (CDA) system for quality evaluation for dental care.<sup>16</sup> Oral hygiene in general and visible plaque on abutments were also recorded by a dental hygienist. Soft tissue pathology, such as perimucositis or peri-implantitis, and fistulas was recorded according to recent definitions.<sup>17,18</sup>

Implant success was determined using the criteria established by Albrektsson and Zarb.<sup>19</sup> Implant stability was checked relative to the actual implant and superstructure, in combination with inspection of the peri-implant mucosa and radiographs of the implants. However, the prosthesis was not removed at the evaluation of implant stability. The bone quality and shape of the alveolar crest were classified according to Lekholm and Zarb.<sup>20</sup>

Parafunctions such as daytime bruxism were recorded according to a scale with five degrees: no, little, sometimes strong, strong, and continuous bruxism. The data were obtained through questioning the patients' personal caregivers. Tooth wear was assessed at the clinical examinations using a five-point scale.<sup>21</sup>

Each implant placed was given a prognostic score from 1 to 4 (1 = uncertain and 4 = very good). The criteria evaluated for each implant were: (a) implant placed in bone of good quality, (b) implant shows good initial stability, (c) no exposed threads, and (d) placement was done according to the standard protocol. The implant was given a score of 4 if all four criteria were fulfilled.

All patients and their caregivers were given an individual prophylactic program by a dental hygienist, including a chart of photographs presenting suitable toothbrushes for optimal cleaning of the prostheses. Patients were recalled every 3 months for an oral hygiene check-up by a dental hygienist (MZ) and annually by a prosthodontist (AE) up to the year 2007. Thereafter, an experienced hospital dentist made the examinations and performed any necessary treatment. Final clinical and radiographic examinations of all available patients were performed in 2012.

Cumulative survival rates for the implants were calculated through life table analysis.<sup>22</sup>

#### **Results**

All patients participating in this study had some degree of mental retardation from neurologic impairment, congenital defect, trauma, or genetic syndromes such as Down syndrome. Often, the patients had other medical disorders such as thyroid dysfunction or epilepsy, sometimes in combination with different degrees of autistic behavior (Table 1). Many of the patients had a list of medications that included substances such as thyroid hormone, antiepileptics, antidepressives, megaloblastics, neuroleptics, and tranquillizers.

Some persons had developed finger and/or oral habits, eg, tongue movements that increased the risk of postsurgical complications. Frequent check-ups and the use of a soft splint to cover the surgical area postoperatively were found to be valuable.

Five of the original 27 patients died during the observation period, but all remaining 22 patients with 70 implants were clinically examined at the final examination. The implant-supported prostheses had then been in function 5 to 10 years after implant treatment.

The distribution of the scores of bone quality and shape of the alveolar crest, according to Lekholm and Zarb,<sup>20</sup> for the 29 treated arches at implant placement were: A1 (n = 2), A3 (n = 2), A4 (n = 1), B2 (n = 4), B3 (n = 6), C2 (n = 3), C3 (n = 10), and D4 (n = 1).

In all, 12 implant-supported single crowns were placed and none lost; 17 implant-supported fixed dental prostheses were placed and three had to be remade due to implant loss. One overdenture was in use until the death of the patient. The survival rate of prosthetic constructions was 27/30 (90%).

The implant-supported fixed prostheses (n = 8) in edentulous arches were fabricated using Nobel Procera (Nobel Biocare, titanium framework and acrylic resin teeth). All fixed partial prostheses (n = 10) were made of metal-ceramic with a framework of gold. The single-tooth restorations (n = 12) were either made as all-ceramic crowns using the CeraOne System (Nobel Biocare) (n = 10) or with individual abutments in titanium (n = 2).

### **Biologic Complications and Implant Failure**

Twelve implants (14%) were lost, three before loading and nine after insertion of the implant-supported fixed prostheses (ISFPs) (Table 2). Both of the 7-mm-long implants failed, whereas none of the 10 longest (16 and 18 mm) failed. The prognostic score did not prove fully reliable even if the most optimistic score, 4, had the lowest failure rate (5%), whereas the rate for the other scores (3, 2, and 1) varied (25%, 11%, and 14%, respectively) (Table 3).

#### **Biologic Complications and Bone Loss**

Among the 22 patients examined at the final followup, 10 (45%) exhibited perimucositis according to the definition of bleeding on probing (BoP) and pocket depth  $\ge$  4 mm; 14 of the 70 implants (20%) had this

#### **Table 3** Abnormal Incidents Noted in Surgical Records of 88 Originally Placed Implants in Relation to the Prognostic Score\* and

| and Implant Loss |                    |                    |                                |           |                           |                          |  |
|------------------|--------------------|--------------------|--------------------------------|-----------|---------------------------|--------------------------|--|
| Score            | No. of<br>implants | Exposed<br>threads | Rupture of mucoperiosteal flap | Infection | Lost<br>before<br>loading | Lost<br>after<br>loading |  |
| 4                | 40                 |                    | 2                              | 4         | 1                         | 1                        |  |
| 3                | 32                 | 11                 | 2                              | 2         | 2                         | 6                        |  |
| 2                | 9                  | 2                  | 1                              |           |                           | 1                        |  |
| 1                | 7                  |                    |                                |           |                           | 1                        |  |

\*Range, 1 to 4 where 4 is very good and 1 is uncertain.

| Table 4    | Peri-implant Bone Loss and Number |
|------------|-----------------------------------|
| of Implant | as According to Time in Function  |

| Exposed | Exposed Time |         | Time (y) | 1е (у)  |        |
|---------|--------------|---------|----------|---------|--------|
| threads | 5.0-5.9      | 6.0-6.9 | 7.0-7.9  | 8.0-8.9 | 9.0-10 |
| ≤ 0     | 8            | 3       | 4        | 5       | 7      |
| 0.1-1.0 |              |         | 6        | 4       |        |
| 1.1-2.0 |              | 1       | 4        |         |        |
| 2.1-3.0 |              | 1       | 2        | 2       |        |
| 3.1-4.0 |              |         | 2        |         |        |
| 4.1-5.0 |              |         | 1        | 1       |        |
| 5.1-6.0 |              |         |          |         |        |
| 6.1-7.0 |              |         |          |         |        |
| 7.1-8.0 |              |         |          | 1       |        |
| Total   | 8            | 5       | 19       | 13      | 7      |

| Table 5 | Prosthodontic | Complications | and Their | Management |
|---------|---------------|---------------|-----------|------------|
|---------|---------------|---------------|-----------|------------|

| Dationt | Complication   | Management   |
|---------|--|--|
| Falleni | Complication   | Ivianagement   |
| 1       | ISFP loose, 1 implant failed before loading  | Failed implant removed, ISFP shortened and later reinserted<br>on new implant and new ISFP made  |
| 4       | Loss of 1 implant before loading, another one when the ISFP was to be inserted   | ISFP inserted on 3 implants awaiting new implants to be placed and new ISFP constructed  |
| 5       | Fracture of porcelain crown  | Polishing of fractured area  |
| 12      | Extremely hard biting individual: porcelain fracture, crown loose (thrown away by patient), fracture of neighbor tooth (intact canine) | New implant-supported crown, which fractured after 6 months, crown on fractured tooth  |
| 17      | Self-destructive behavior (banged head against walls),<br>ISFP loosened several times, ISFP fractured, implants lost,<br>severe wear   | Several reinsertions of ISFP, removal of failed implants,<br>back to implant overdenture, and, finally, conventional<br>complete denture |
| 19      | Fractures of abutment and fixed partial denture screws, fracture of acrylic parts of superstructure at epileptic seizures              | ISFP rebuilt with occlusal contact on titanium splint to protect the acrylics  |
| 23      | Implant overdenture poor retention (patient removed it repeatedly with his tongue), overdenture fractured                              | Overdenture repaired but only used at rare social events   |
| 25      | Implant lost after 4 years, ISFP loose   | ISFP reinserted awaiting new implant placement   |

ISFP = implant-supported fixed prosthesis.

diagnosis. Three of the 15 patients with measurable radiographs (20%) and 4 implants (8%) were diagnosed with peri-implantitis using the definition bone loss  $\geq$  3 threads and BoP.<sup>17</sup>

The peri-implant bone loss was small to moderate in most patients (Table 4). It should be noted that the seven implants in the four patients followed for 9 to 10 years showed no measurable bone loss. Altogether, 27 (52%) of the 52 implants with acceptable radiographs had no measurable bone loss after 5 to 10 years, and for a further 10 (19%), the bone loss corresponded to, at most, one thread during this period. The greatest peri-implant bone loss observed, corresponding to eight threads, occurred in one patient before loading, early after placement, probably as a consequence of a postoperative dehiscence of the

mucoperiosteal flap. This implant remained stable and no more bone loss was seen up to the last examination shortly before the death of the patient after 6 years. In another patient with a maxillary full-arch ISFP, bone loss was above average around three implants (eight, five, and three threads, respectively), whereas the bone loss measured one thread around the remaining 3 implants. The ISFP was stable and well functioning all through the 9-year follow-up period.

#### **Prosthodontic Complications**

Several prosthodontic complications occurred, from minor and easily correctable to severe requiring retreatment (Table 5). The most severe complication occurred in a patient with Aspberger syndrome and



**Fig 1** Extensive wear related to hard biting and epileptic seizures associated with patient no. 19's disorders. ISFP rebuilt with occlusal contact on titanium splint to protect the acrylics (mandible).

**Table 6**Quality Assessment According toCalifornia Dental Association Criteria

| Assessment  | Surface<br>and color | Anatomical<br>form  | Margin<br>integrity |
|---|----------------------|---------------------|---------------------|
| Satisfactory<br>Excellent<br>Acceptable                     | 12<br>13             | 10<br>12            | 13<br>12            |
| Not acceptable<br>Repair or correct<br>Replace <sup>b</sup> | 2                    | 3 <sup>a</sup><br>2 | 2                   |

<sup>a</sup>Porcelain fractures (all three became acceptable after polishing). <sup>b</sup>ISFP failed after multiple implant failures (provided with an implant overdenture and eventually a complete denture; single-tooth implant crown: all porcelain fractured (provided with a new crown).

| Table 7   | Life Table / | Analysis  | of Placed | Implants |
|-----------|--------------|-----------|-----------|----------|
| (Re-opera | ated Implan  | ts Not In | cluded)*  |          |

| Time period (y)   | No.<br>followed | No.<br>failed | Success rate<br>within group<br>(%) | CSR<br>(%) |
|-------------------|-----------------|---------------|-------------------------------------|------------|
| Placement-loading | 88              | 3             | 96.6                                | 96.6       |
| Load-1            | 84              | 1             | 98.8                                | 95.4       |
| 1–2               | 80              | 2             | 97.5                                | 93.0       |
| 2-3               | 77              | 0             | 100                                 | 93.0       |
| 3-4               | 77              | 1             | 98.7                                | 91.8       |
| 4–5               | 76              | 3             | 96.1                                | 88.2       |
| 5-6               | 73              | 2             | 97.3                                | 85.8       |
| 6-7               | 70              | 0             | 100                                 | 85.8       |
| 7–8               | 49              | 0             | 100                                 | 85.8       |
| 8-9               | 24              | 0             | 100                                 | 85.8       |
| 9–10              | 7               | 0             | 100                                 | 85.8       |

CSR = cumulative survival rate.

\*Five patients died and their nine implants were not included in further calculations.

self-destructive behavior. He often banged his head against walls, which repeatedly led to the loosening of the ISFP and, eventually, fracture and loss of both



Fig 2 Maxilla in patient no. 19.



**Fig 3** The oral hygiene was not optimal in patient no. 19 despite the assistance of personal caregivers.

implants and the prosthesis. An overdenture was constructed on one remaining implant, but this implant was also lost, resulting in a return to a complete denture, which the patient seldom uses.

The great majority of the ISFPs still in use were satisfactory according to the CDA quality assessment. Because of various complications, five prostheses (19%) were considered not acceptable, although three porcelain fractures became acceptable after polishing (Table 6).

All patients and/or caregivers were satisfied with the appearance and function of the ISFPs and agreed that they would consider implant treatment again, if necessary.

It was difficult to obtain reliable answers to questions on parafunctional habits. However, some of the prosthodontic complications were probably related to various parafunctions and/or extremely hard biting (Table 5). The same cause was probable for a few patients with severe tooth wear (Figs 1 to 3).

#### Survival Rate

The cumulative survival rate (CSR) for placed implants was 85.8% after 5 to 10 years. No implant was lost after 6 years (Table 7).

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# Discussion

There is no doubt that the group of patients with different neurologic and other disabilities, often including mental retardation, presented/offered greater problems during implant treatment and maintenance compared with healthy patients. Nevertheless, it was possible to carry out the treatment, and the outcome was relatively favorable and led to improved dental appearance and oral function.

The implant failure rate (12 of 88 placed implants lost = 14%) was much greater than in a series of healthy patients.<sup>7,23</sup> However, the CSR (Table 7) changed little from 5 years (88.2%) to 10 years (85.8%). The 10-year results are somewhat lower than the outcomes presented in systematic reviews of implant therapy. For example, according to meta-analyses, the 10-year estimated survival of ISFPs was 93.1%.<sup>8</sup> This may indicate that the problems in disabled patients are greatest during the first period after implant placement, after which the situation can stabilize.

However, the relationship between implant failure and the patients' diagnoses can only be speculated. As discussed earlier,<sup>15</sup> the implant loss in patient no. 1 might be associated with the reduced resistance to infections in subjects with Down syndrome in combination with macroglossia and oral habits. Two of the new implants placed in this patient after the early loss of one implant also failed.

An increased risk of implant failures in patients with Down syndrome has been reported in a study of 18 patients with various handicaps, 4 of whom had Down syndrome. Of four implants placed in 2 patients with Down syndrome, three failed.<sup>12</sup> The suitability of patients with Down syndrome as candidates for implant placement has been questioned because of macroglossia, osteoporotic-like alveolar bone, and a tendency toward poor cooperation.<sup>24</sup>

In contrast, in a study of special care patients, including three subjects with Down syndrome, no implants failed.<sup>13</sup> In this study, there were three more patients with Down syndrome who did not exhibit any implant failure, although one of them died before the implant was loaded. Several of the prosthodontic complications including different types of fractures were probably caused by oral habits, trauma, hard biting, and epileptic seizures associated with the patients' disorders (Table 5). Surprisingly, a retrospective study covering 1 to 16 years of patients with severe epilepsy reported excellent results with few implant failures and other complications and no progression of the frequent perimucositis to peri-implant bone loss.<sup>14</sup> It should be mentioned that only a small percentage of the patients were followed for more than 6 years.

It is well known today that good oral hygiene is important to maintain healthy conditions around teeth and implants. Most of the patients in this study could not maintain good oral care on their own but required help from caregivers and regular control and service by a dental hygienist. Still, many of the patients had a large amount of plaque and almost half of them exhibited perimucositis (Fig 3). Other series of disabled patients have reported similar problems with poor oral hygiene and a high prevalence of peri-mucosal inflammation but good functional results from implant treatment.<sup>12-15</sup>

The more severe condition, peri-implantitis, including bone loss, was diagnosed in 20% of patients and for 8% of implants, according to the definition used.<sup>17</sup> The prevalence, definition/diagnosis, clinical importance, as well as the etiology of peri-implantitis are controversial.<sup>25-29</sup>

Even if much is not yet known regarding these conditions, it is agreed that bacterial inflammation is an essential part and the disease may develop from perimucositis to peri-implantitis.<sup>30</sup> Therefore, maintaining good oral hygiene is important but difficult in this group of patients. It is necessary that these patients are helped by informed caregivers in daily oral care and also given regular professional support.

The results of the present study and of earlier reports<sup>11-15</sup> indicate that implant treatment can be a valid option in the oral rehabilitation of patients with neurologic disabilities, although maintenance often requires the management of more complications compared with healthy implant patients.

#### Conclusion

Patients with different neurologic and other disabilities, often including mental retardation, present greater problems during implant treatment and maintenance phases than healthy patients generally do. Nevertheless, it was possible to carry out the treatment, and the outcomes were relatively favorable. The results indicate that implant treatment can be a valid option in oral rehabilitation of patients with neurologic disabilities.

#### **Acknowledgments**

The authors wish to thank Lisbeth Heijel-Berndtsson, Clinical Coordinator, National Orofacial Resource Centre for Rare Disorders, Gothenburg, for organizing the clinical examination of these patients. The authors also wish to thank the team of oral surgeons: Drs Göran Widmark, Carl Johan Ivanoff, Cecilia Larsson, and Lucy Kartous, and oral radiologist Dr Eva Borg, Specialist Clinics of Oral and Maxillofacial Surgery and Oral and Maxillofacial Radiology, Mölndal Hospital, Mölndal. We also wish to thank

the staff at the Department of Oral and Maxillofacial Radiology, Institute of Odontology, University of Gothenburg, Sweden. This study has been supported by grants from Praktikertjänst, Stockholm, Sweden.

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