Dynamic versus Adynamic Graciloplasty in Treatment of End-Stage Fecal Incontinence: Is the Implantation of the Pacemaker Really Necessary? 12-Month Follow-Up in a Clinical, Physiological, and Functional Study

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Purpose. The aim of the study is to compare functional results of end-stage fecal incontinence treatment with dynamic graciloplasty and adynamic graciloplasty augmented with transanal conditioning of the transposed muscle. Methods. A total of 20 patients were qualified for graciloplasty procedure due to end-stage fecal incontinence. 7 patients underwent dynamic graciloplasty (DGP), whereas 13 patients were treated with adynamic graciloplasty, with transanal stimulation in the postoperative period (AGP). Clinical, functional, and quality of life assessments were performed 3, 6, and 12 months after the procedures. Results. There were no intraoperative or early postoperative complications. The detachment of gracilis muscle tendon was observed in one patient in DGP group and two in AGP group. There was a significant improvement of Fecal Incontinence Quality of Life (FIQL) and Fecal Incontinence Severity Index (FISI) scores in both groups 12 months after procedure. Anorectal manometry showed improvement regarding basal and squeeze pressures in both groups, with significantly better squeeze pressures in AGP group. Conclusions. The functional effects in the DGP and AGP groups were similar. Significantly lower price of the procedure and avoidance of implant-related complication risk suggest the attractiveness of the AGP method augmented by transanal stimulation.

1. Introduction

Approximately 7–10% of the working population suffers from fecal incontinence, about 30% of which is affected by the end-stage of the disease. This includes patients with congenital absence of the anal sphincter, patients after surgical treatment of the anus and rectum, and also after major perineal trauma, the spinal injury or damage of peripheral innervation of sphincters [1], who in most cases do not respond to conservative measures, such as biofeedback training or electrical stimulation, and are not eligible for surgical repair, such as sphincteroplasty. To improve quality of life and avoid the end-stoma, this group requires the “last chance methods”: sphincter reconstruction by procedures involving transposition of autologous muscle or replacing anal sphincter with artificial prosthesis [2]. Regarding muscle transposition procedures, most commonly used procedures are the unstimulated graciloplasty (AGP) and dynamic (stimulated) graciloplasty (DGP), which were proven successful in highly selected patients [3].

Despite the fact that dynamic graciloplasty became more popular than its unstimulated variant, Rosen et al. showed that after initial conditioning of the transposed muscle even half of the patients no longer benefit from electrostimulation a year after the treatment [4]. Moreover, a large percentage of patients reported constipation as adverse effect of the procedure, even when the stimulation was turned off [5]. The data coming from literature reviews showed that the risk of complications was similar in both procedures; however the most severe complications were correlated with infection of the stimulator site or electrodes; thus revision and stimulator explantation rate in DGP is high [6]. The question emerged...
whether the pacemaker implantation is necessary for the success of the operation. There was no sufficient data comparing dynamic graciloplasty and adynamic (unstimulated) graciloplasty, partially because these procedures are rarely carried out, only in highly specialized units. Therefore, we designed a clinical study to compare the results of patients’ functionality and satisfaction after dynamic graciloplasty and adynamic graciloplasty.

2. Purpose

The aim of the study is to compare functional results of end-stage fecal incontinence treatment with dynamic graciloplasty and adynamic graciloplasty augmented with transanal conditioning of the transposed muscle.

3. Methods

This was a prospective observational study. In this study, based on the IFGD guidelines, the following stages of research were used

(i) identification and assessment of the epidemiological control of defecation based on subjective data, function tests, electrophysiological measurements, and imaging,
(ii) the patient division into two groups:
(a) patients treated by dynamic graciloplasty (DGP) and
(b) patients treated by adynamic graciloplasty (AGP),
(iii) analysis of complications during and after surgery and functional tests 3, 6, and 12 months after surgery,
(iv) assessment of the Fecal Incontinence Severity Index (FISI) [7] and Fecal Incontinence Quality of Life (FIQL) at 3, 6, and 12 months after surgery.

3.1. Inclusion Criteria

(i) End-stage fecal incontinence (>35 points in Fecal Incontinence Severity Index),
(ii) lack of sphincter or extensive sphincter defect (more than 1/3 of circumference of the muscle based on a transanal ultrasound examination),
(iii) unsuccessful previous treatments: electrostimulation (ES), biofeedback + ES, sphincteroplasty, and SNS,
(iv) sphincter innervation defect (verified by electrophysiological examination).

3.2. Exclusion Criteria

(i) Age less than 18 and more than 70 years,
(ii) contraindications to graciloplasty including patient’s refusal to undergo the procedure, peripheral neuropathy (e.g., diabetic and polyneuropathic), ischemic limb disorders, severe heart, kidney, liver conditions, and so forth,
(iii) contraindications to electrical stimulation usage.

A total of 20 patients were qualified for graciloplasty procedure during the period of 6 years, 14 women and 6 men; the mean age was 48 years (range 19–70 yrs). There were 8 Medtronic implantable stimulators on the Investigators’ disposal. Short expiry date of the devices and long qualification procedure forced the Investigators to withdraw initially planned randomisation, and to qualify first 8 consecutive patients to dynamic graciloplasty group instead. One of the stimulators was found defective prior to implantation, so there were a total of 7 patients in the DGP group, and 13 in AGP group. Both groups had similar gender and age characteristics (differences not significant).

3.3. Technique of Graciloplasty Procedure. In both groups of patients (DGP and AGP), the procedure was performed under general anesthesia. The gracilis muscle innervation was detected preoperatively by surface electromyography with a flexible array of 16 equally spaced silver bar electrodes and amplification system (ICU EMG16 Bioelettronica, Turin) as described by Bottin et al. [8]. Two 5 cm longitudinal incisions along the length of the thigh allowed the identification and mobilization of gracilis muscle. The classical method of operation described by Baeten et al. [9] or split-sling modification (Rosen et al.) [4] were performed. Gracilis muscle was dissected at the level of the neurovascular bundle (which was identified intraoperatively by electrostimulator needle: Neuro-pulse Aaron Medical Industries, Inc.). Gracilis muscle was formed into the alpha or gamma loop, depending on the anatomical conditions and fitted around the rectum creating thereby a new sphincter muscle (Figure 1).

In the DGP group the next step was the attachment of two monopolar electrodes to the muscle (model 4300 Medtronic) in the area of the main neurovascular bundle. Then the best parameters stimulation were defined. The most frequent parameters of the stimulation were: amplitude from 0.5 to 1.0 V, pulse width 0.210 ms, pulse frequency of 10 to 20 Hz (pacemaker IPG InterStim Medtronic 3023). Electrostimulation was not used for the first week after surgery. The elastic bandage was maintained on the leg for 2-3 weeks.
Drainage was removed from the wound 2–3 days after surgery. The patient was discharged home on average 5–7 days after surgery. In DGP group, two weeks after the procedure the prestimulation of muscle was started in order to achieve muscle fiber conversion (conditioning). Continuous stimulation was applied (voltage 1.5 V, pulse width 0.210 ms, frequency of 5.2 Hz). After 2 weeks, the stimulation was increased to 16 Hz (under stimulation parameters and anal canal pressures control) during the next 6 months. The output parameters of stimulation were maintained in case of 5 patients. In case of 1 patient the amplitude was increased to 2.0 V, and 1 patient required the electrodes polarity change, the increase of amplitude to 3.0 V, and frequency of stimulation. 3 patients with colostomy who had satisfactory functional results of treatment were qualified for stoma closure. It was performed in the period of 2–3 months after dynamic graciloplasty.

In AGP group the transanal stimulation by endoanal TNS stimulator SM1 (Schwa Medico) was implemented after healing of the wounds (about 2–3 weeks). The constant pulse frequency of 60 Hz were applied, 20–30 min three times a day.

3.4. Statistical Methods. Statistical analysis was performed by using the Mann-Whitney and Kruskal-Wallis test, which described the dependence of group-specific parameters, and which assumed the level of statistical significance $P < 0.05$. Friedmans’ test was used to determine the temporal relation to the individual parameters in the studied groups. Calculations were performed using STATISTICA software.

4. Results

There were no intraoperative or early postoperative complications in our survey. The detachment of gracilis muscle tendon was observed in one patient in DGP group. This patient was scheduled for reoperation—due to evident atrophy of the transposed muscle, observed intraoperatively, the patient underwent a repeated graciloplasty using the other muscle. The electrostimulation was performed in 6 weeks after operation and it gave good functional results. Six months after the second treatment a gradually rapid muscular function atrophy was observed and confirmed by manometric examination. The electrophysiological assessment revealed a total lack of response to stimulation and ultrasound tests showed fast atrophy of gracilis muscle. Therefore, the patient was qualified to artificial anal sphincter implantation and excluded from further follow-up in this study.

One patient in AGP group suffered from the limb wound suppuration. The wound was treated conservatively and the antibiotic was administered with a satisfying result. Additionally, the same patient had symptoms of deep vein thrombosis in operated limb several months after surgery. It was treated conservatively. In case of two patients from AGP group the detachment of the tendon was observed and required reoperation without further complications.

4.1. Assessment of Defecation Control Score (FISI), and Quality of Life Score (FIQL). Defecation control improvements were observed in both groups. In Jorge-Wexner questionnaire assessment, the patients scored from 11.5 points before surgery to 11.3 points in 6 months after surgery and 10.1 points in 12 months after surgery, the differences did not reach statistical significance (Friedman $P = 0.07$). However, the FISI score assessments ranged from 36.0 before to 29.8 points after treatment and were statistically significant ($P < 0.05$) (Figure 2). There was a significant ($P < 0.05$) improvement of Fecal Incontinence Quality of Life (FIQL) scores in both groups in 12 months after procedure. In DGP group patients felt embarrassed much more rarely (FIQL-E1.8 before versus 3.3 points after) than in AGP group (1.8 versus 2.2 points) and the difference was significant ($P < 0.05$).

4.2. Functional Examination and Electrophysiological Evaluation. There was no significant difference in the initial basal pressure (BAP) and squeeze pressure (SAP) values between the groups (Mann-Whitney, $P = 0.077$). After the procedure, mean BAP and SAP values increased gradually and significantly in both groups ($P < 0.05$). There were no significant differences in BAP values between DGP and AGP group during the follow-up study. In AGP group, beginning from assessment 3 months after surgery, SAP values were significantly higher than in DGP group. This difference increased during the observation 6 and 12 months after surgery (Table 1).

In DGP group the pressure in the anal canal increased on average by 37 cm H$_2$O above the basic pressure at the time of pacemaker activation. Interestingly, only 2 patients still used a stimulation permanently 6 months after operation. The remaining 4 patients activated the stimulator only occasionally. They claimed that it was not necessary for adequate control of defecation.

5. Discussion

During the last 50 years several European, American, and Australian centers used various modifications of muscle transposition or artificial implant in cases of patients with end-stage Fecal Incontinence. Belyaev and coworkers carried out a review of literature on transposition of the gracilis.
Table 1: Manometric examination results (BAP: basal anal pressure, SAP: squeeze anal pressure).

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<tr>
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<th>Before surgery</th>
<th>3-month follow-up</th>
<th>6-month follow-up</th>
<th>12-month follow-up</th>
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<tr>
<td></td>
<td>BAP (cmH₂O)</td>
<td>SAP (cmH₂O)</td>
<td>BAP (cmH₂O)</td>
<td>SAP (cmH₂O)</td>
</tr>
<tr>
<td>DGP</td>
<td>17 ± 11</td>
<td>35 ± 15</td>
<td>31 ± 12</td>
<td>55 ± 23</td>
</tr>
<tr>
<td>AGP</td>
<td>21 ± 15</td>
<td>39 ± 18</td>
<td>45 ± 21</td>
<td>82 ± 45</td>
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6. Conclusions

The presented results show that the rate of graciloplasty complications associated with pacemaker implantation can be potentially reduced by performing adynamic graciloplasty combined with simultaneous transanal stimulation. The functional effects of the DGP and AGP in presented survey were similar. Significantly lower price of the procedure and avoidance of implant-related complications risk, suggest the attractiveness of the AGP method augmented by transanal stimulation.

Conflict of Interests

None of the authors of the paper has any commercial associations that might pose a conflict of interests in connection with the submitted paper.
Authors’ Contribution

Piotr Walega designed the study, qualified the patients for the procedures, performed surgical operations and consulted the results. Michal Romaniszyn performed the manometric and electrophysiological examinations, analyzed the data, and coedited the paper. Benita Siarkiewicz gathered the data and wrote the paper. Dorota Zelazny prepared, performed, and analyzed Quality of Life and Fecal Incontinence assessments. This work was supported by Jagiellonian University statutory funding.

References
