Bariatric Surgery versus Intensive Medical Therapy for Diabetes — 3-Year Outcomes

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Abstract

BACKGROUND—In short-term randomized trials (duration, 1 to 2 years), bariatric surgery has been associated with improvement in type 2 diabetes mellitus.

METHODS—We assessed outcomes 3 years after the randomization of 150 obese patients with uncontrolled type 2 diabetes to receive either intensive medical therapy alone or intensive medical therapy plus Roux-en-Y gastric bypass or sleeve gastrectomy. The primary end point was a glycated hemoglobin level of 6.0% or less.

RESULTS—The mean (±SD) age of the patients at baseline was 48±8 years, 68% were women, the mean baseline glycated hemoglobin level was 9.3±1.5%, and the mean baseline body-mass index (the weight in kilograms divided by the square of the height in meters) was 36.0±3.5. A total of 91% of the patients completed 36 months of follow-up. At 3 years, the criterion for the primary end point was met by 5% of the patients in the medical-therapy group, as compared with 38% of those in the gastric-bypass group (P<0.001) and 24% of those in the sleeve-gastrectomy group (P = 0.01). The use of glucose-lowering medications, including insulin, was lower in the surgical groups than in the medical-therapy group. Patients in the surgical groups had greater mean percentage reductions in weight from baseline, with reductions of 24.5±9.1% in the gastric-bypass group and 21.1±8.9% in the sleeve-gastrectomy group, as compared with a reduction of 4.2±8.3% in the medical-therapy group (P<0.001 for both comparisons). Quality-of-life measures were significantly better in the two surgical groups than in the medical-therapy group. There were no major late surgical complications.

CONCLUSIONS—Among obese patients with uncontrolled type 2 diabetes, 3 years of intensive medical therapy plus bariatric surgery resulted in glycemic control in significantly more patients

than did medical therapy alone. Analyses of secondary end points, including body weight, use of glucose-lowering medications, and quality of life, also showed favorable results at 3 years in the surgical groups, as compared with the group receiving medical therapy alone. (Funded by Ethicon and others; STAMPEDE ClinicalTrials.gov number, NCT00432809.)

Bariatric surgery has recently emerged as a potentially useful treatment for type 2 diabetes mellitus.[1] Observational studies[2–5] and randomized, controlled trials[6–10] have shown that procedures including Roux-en-Y gastric bypass, sleeve gastrectomy, gastric banding, and biliopancreatic diversion significantly improve glycemic control and favorably affect cardiovascular risk factors.

In the Surgical Treatment and Medications Potentially Eradicate Diabetes Efficiently (STAMPEDE) trial, we found that 1 year after randomization, gastric bypass and sleeve gastrectomy were superior to intensive medical therapy alone in achieving glycemic control and reducing cardiovascular risk factors while decreasing dependency on pharmacotherapy for diabetes management.[7] Although bariatric surgery yields short-term improvements in glycemic control, questions remain regarding the durability of the metabolic benefits of surgery, long-term safety, quality of life, and effects on diabetes-related end-organ disease. The current report provides results of the 3-year follow-up analyses from the STAMPEDE trial and addresses other unanswered questions about the durability of the benefits of bariatric surgery as compared with intensive medical therapy for treating diabetes mellitus.

METHODS

STUDY DESIGN

The rationale, design, and methods of the study have been reported previously.[7,11] The complete protocol was approved by the institutional review board at the Cleveland Clinic and is available with the full text of this article at NEJM.org. Briefly, the trial was a three-group, randomized, controlled, single-center study involving 150 obese patients, in which the effects of intensive medical therapy were compared with those of gastric bypass or sleeve gastrectomy. With the use of block randomization, patients were assigned, in a 1:1:1 ratio, to one of the three study groups, with stratification according to the baseline use of insulin. Eligibility criteria included an age of 20 to 60 years, a glycated hemoglobin level of more than 7.0%, and a body-mass index (BMI, the weight in kilograms divided by the square of the height in meters) of 27 to 43. All patients provided written informed consent.

STUDY OUTCOMES

The primary outcome was a glycated hemoglobin level of 6.0% or less, with or without the use of diabetes medications.[7,11] This report provides 3-year outcomes in the study patients, including measures of glycemic control, weight loss, blood pressure, lipid levels, renal function, carotid intima–media thickness,[12] medication use, adverse events, disease-related complications, and quality of life (as evaluated with the use of the RAND 36-Item Health Survey).[13] The strategy for all three groups was the adjustment of medical therapy (every 3 months for 2 years and every 6 months thereafter) with the goal of achieving the therapeutic target of a glycated hemoglobin level of 6.0% or less, without unacceptable side effects associated with medical treatment.

STUDY OVERSIGHT

This investigator-initiated trial was financially supported by Ethicon, with additional support from LifeScan, the Cleveland Clinic, and the National Institutes of Health. The sponsors participated in discussions regarding study design but had no role in data accrual, data analysis, or manuscript preparation. The first author wrote the first draft of the manuscript. All the authors had full and independent access to all the data and vouch for the integrity and the accuracy of the analysis and its fidelity to the protocol. Complete study governance is outlined in the Supplementary Appendix, available at NEJM.org.

STATISTICAL ANALYSIS

We report all continuous variables with a normal distribution as means and standard deviations. Variables with a non-normal distribution are reported as medians and interquartile ranges. Categorical variables are summarized with the use of frequencies. We used the chi-square test to evaluate the primary end point of a glycated hemoglobin level of 6.0% or less at 3 years. We used an analysis of variance to analyze continuous laboratory measurements and to perform comparisons among the three study groups. For glycemic measures and body weight, a mixed model for repeated measures was used to analyze the change from baseline, and least-square means with corresponding standard errors were plotted graphically.

A stepwise multivariable logistic model was used to determine factors associated with achieving the primary end point. Factors that were considered in the model included age, sex, insulin use, duration of diabetes, baseline glycated hemoglobin and glucose levels, C-peptide levels, baseline BMI, change in BMI, blood pressure, and lipid measures. No adjustments were made for multiple comparisons, since these were exploratory analyses. Analyses were performed with the use of SAS software, version 9.2 (SAS Institute).

RESULTS

STUDY PATIENTS

Of the 150 patients who underwent randomization from March 2007 through January 2011, a total of 9 patients withdrew from the trial immediately after randomization or during the initial 6 months (8 patients in the medical-therapy group and 1 patient who did not undergo sleeve gastrectomy because of severe anemia); 4 patients were lost to follow-up. The remaining 137 patients (91.3%) were evaluated in the 3-year assessment of safety and efficacy.

The baseline characteristics of the 150 patients were reported previously.[7] In the current analysis, 68% of the patients were women, and 74% were white. The mean (±SD) age was 48±8 years, and the mean BMI was 36±3.5; 49 patients (36%) had a BMI of less than 35. The mean glycated hemoglobin level was 9.3±1.5%, and the average duration of diabetes was 8.3±5.1 years, with 43% of patients requiring insulin at baseline. There were no significant differences between the study groups at baseline (Table S1 in the Supplementary Appendix).

PRIMARY END POINT

At 3 years, the target glycated hemoglobin level of 6.0% or less was achieved in 5% of the patients in the medical-therapy group, as compared with 38% of those in the gastric-bypass group (P<0.001) and 24% of those in the sleeve-gastrectomy group (P = 0.01) (Table 1). The percentage of patients who had a glycated hemoglobin level of 6.0% or less at 1 year but did not maintain this level of glycemic control at 3 years (which was defined as a glycemic relapse) was 80% in the medical-therapy group, as compared with 24% in the gastric-bypass group (P = 0.03) and 50% in the sleeve-gastrectomy group (P = 0.34). In the entire cohort, a reduction in the BMI was the only significant predictor of achieving the primary end point (odds ratio for each 1-unit decrease in BMI, 1.41; 95% confidence interval [CI], 1.22 to 1.64; P<0.001) (Table S2 in the Supplementary Appendix). In the two surgical groups, meeting the criterion for the primary end point was predicted both by a reduction in the BMI (odds ratio, 1.33; 95% CI, 1.15 to 1.56; P<0.001) and by a duration of diabetes of less than 8 years (odds ratio, 3.3; 95% CI, 1.2 to 9.1; P = 0.02).

GLYCEMIC CONTROL

After 3 years, each of the two surgical procedures was superior to intensive medical therapy alone in achieving exploratory targets for glycated hemoglobin of 6.5% and 7.0%, with or without the use of diabetes medications (P<0.05 for all comparisons) (Table 1). Median levels of fasting plasma glucose were significantly lower in the two surgical groups than in the medical-therapy group (P<0.01 for both comparisons) (Table 1). There were more rapid, larger, and more sustained reductions in levels of glycated hemoglobin and fasting plasma glucose and in the use of glucose-lowering medications in the two surgical groups than in the medical-therapy group (Table 2, Fig. 1A and 1C, and Fig. S1 in the Supplementary Appendix). The reductions in glycated hemoglobin levels, medication use, and BMI in the surgical groups were similar in patients with a BMI of less than 35 and those with a BMI of 35 or more, and the reductions in both BMI subgroups of the surgical group were greater than the reductions in either BMI subgroup of the medical-therapy group (Fig. 1B, and Fig. S2 and S3 in the Supplementary Appendix).

DIABETES MEDICATIONS

At 3 years, the use of glucose-lowering medications including insulin was reduced from baseline in the two surgical groups (Table 2). Patients in the gastric-bypass group required fewer glucose-lowering medications per day than did those in the sleeve-gastrectomy group (0.48±0.80 vs. 1.02±1.01). The proportion of patients who were not taking any glucose-lowering medications was significantly higher in the gastric-bypass group than in the sleeve-gastrectomy group (Table 2).

WEIGHT LOSS

At 3 years, reductions in body weight, BMI, waist circumference, and waist-to-hip ratio were greater after gastric bypass and sleeve gastrectomy than after intensive medical therapy (Table 1, Fig. 1D, and Table S3 in the Supplementary Appendix). The reduction in body weight was greater after gastric bypass than after sleeve gastrectomy (P = 0.02).

CARDIOVASCULAR BIOMARKERS AND MEDICATIONS

The decrease in triglyceride levels and increase in high-density lipoprotein (HDL) cholesterol levels that had been observed after the two surgical procedures, as compared with intensive medical therapy, were sustained at 3 years (Table 1). There were no significant differences in blood pressure and low-density lipoprotein (LDL) cholesterol levels among the three study groups, although there was a significant reduction in the number of medications needed to treat hyperlipidemia and hypertension in the surgical groups (Table S4 in the Supplementary Appendix). There were no significant differences among the three groups in maximal carotid intima–media thickness at baseline or at 24 months (Table S5 in the Supplementary Appendix).

RENAL OUTCOMES

At 3 years, the urinary albumin-to-creatinine ratio (as measured in milligrams of albumin to grams of creatinine) decreased from a median of 9 to 6 in the gastric-bypass group (P = 0.08) and from 12 to 7 in the sleeve-gastrectomy group (P<0.001), as compared with 6.5 to 5.5 in the medical-therapy group (P = 0.77). The reduction in the albumin-to-creatinine ratio from baseline in the two surgery groups was significant, as compared with the medical-therapy group (P<0.04 for both comparisons) (Table S5 in the Supplementary Appendix). Of the patients with baseline albuminuria, a return to normal values at 3 years occurred in 8 of 13 patients (62%) in the gastric-bypass group (P = 0.04 for the within-group comparison), 8 of 10 patients (80%) in the sleeve-gastrectomy group (P = 0.11), and 1 of 4 patients (25%) in the medical-therapy group (P = 1.00). There was no significant difference in the serum creatinine level and the estimated glomerular filtration rate among the three groups during follow-up (Table S5 in the Supplementary Appendix).

QUALITY OF LIFE

There were significant improvements in five of eight mental and physical domains among patients in the gastric-bypass group and in two of eight domains among patients in the sleeve-gastrectomy group, as compared with the medical-therapy group (Fig. 2, and Table S6 in the Supplementary Appendix).

ADVERSE EVENTS

Additional surgical interventions were required in 4 patients within the first 12 months after randomization,[7] but none were performed thereafter. There was no excessive weight loss or hypoalbuminemia and no life-threatening complications or deaths in any of the groups (Table 3, and Tables S7 and S8 in the Supplementary Appendix). Excessive weight gain (>5% from baseline) was observed in 7 of 43 patients (16%) in the medical-therapy group and in no patients in the two surgical groups (P<0.05 for both comparisons).

DISCUSSION

The results of this follow-up analysis show that 3 years after randomization, bariatric surgery, as compared with intensive medical therapy alone, was associated with superior and sustained glycemic control and weight reduction. Patients who underwent gastric bypass or sleeve gastrectomy were significantly more likely to achieve and maintain a glycated

hemoglobin level of 6.0% or less than were those who received intensive medical therapy alone. More than one third of the patients in the gastric-bypass group and a fifth of those in the sleeve-gastrectomy group, as compared with no patients in the medical-therapy group, had a glycated hemoglobin level of 6.0% or less without the use of diabetes medications. Patients in the two surgical groups had a significant absolute decrease of 2.5 percentage points in glycated hemoglobin levels, a reduction that was sustained for 3 years, as compared with a reduction of 0.6 percentage points in the medical-therapy group. The results of surgery are particularly striking in this population with relatively long-standing uncontrolled disease. The surgically treated patients had superior glycemic control for 3 years while also reducing dependency on oral diabetes medications and insulin. More than 90% of surgical patients had glycemic control without the use of insulin. Weight loss and a shorter duration of diabetes were the main predictors of having a glycated hemoglobin level of 6.0% or less after surgery.

Analysis of secondary end points, including BMI, body weight, waist circumference, and levels of triglycerides and HDL cholesterol, also showed favorable results at 3 years in the surgical groups, as compared with the group receiving intensive medical therapy alone. Patients in the two surgical groups had a significant reduction in the use of antihypertensive and lipid-lowering agents, even though there were no significant changes from baseline measurements in blood pressure or LDL cholesterol among the three study groups. Some adverse effects of surgical treatment were observed in this study but were modest in severity and relatively uncommon after the first year.

Although observational studies have shown impressive improvements in glycemic control after bariatric surgery, with rates of improvement in diabetes ranging from 55 to 95%,[1–5] direct comparisons with intensive medical therapy have been limited.[6–10] The Swedish Obese Subjects (SOS) study, a nonrandomized, prospective trial comparing bariatric surgery with conventional medical treatment, showed higher rates of diabetes remission after surgery than after conventional medical treatment at 2, 10, and 20 years. The SOS study also showed significant reductions in long-term complications, including rates of death from any cause and major cardiovascular events, with surgical treatment.[14–16]

Five short-term, randomized, controlled trials compared bariatric surgery with medical treatment with respect to type 2 diabetes, with 1 to 2 years of follow-up.[6–10] The initial STAMPEDE report showed that at 1 year, patients had better glycemic control (defined as a glycated hemoglobin level of 6.0% or less, with or without the use of medications) after gastric bypass (42% of patients) or sleeve gastrectomy (37%) than after intensive medical therapy (12%) (P<0.001 for both comparisons).[7] All five trials showed that standard bariatric procedures, as compared with medical treatment alone, were associated with few major complications and resulted in superior glycemic control, weight reduction, and reductions in cardiovascular risk factors. Our findings show continued durability of glycemic control and persistent reductions in cardiovascular risk factors at 3 years after surgery. The diabetes remission rates after surgery in our study are similar to those reported by Ikramuddin et al.[9] but are lower than those reported by Mingrone et al.[8] Such discrepancies could be explained by the greater severity and longer duration of diabetes in our population, as well as a stricter definition of remission.

Observational studies have suggested that bariatric surgery may reduce long-term renal impairment associated with diabetes.[4,17] Diabetes and obesity are independent risk factors for the development of albuminuria, which is associated with an increased risk of death from cardiovascular causes and end-stage renal disease.[18] We noted improvement in albuminuria in the surgical groups despite a reduction in the use of renin–angiotensin system blockers, which suggests that bariatric surgery may have a role in the prevention of further renal parenchymal damage. Although there was no significant change in serum creatinine levels or the glomerular filtration rate, these findings do not imply a lack of benefit of weight loss on renal function, since creatinine and the glomerular filtration rate are influenced by loss of muscle mass associated with weight loss.[19] Cumulatively, our data should be considered to be hypothesis-generating and suggest the need for further long-term studies examining the effects of bariatric surgery on renal function in diabetes.

Obese patients with diabetes have a reduced quality of life and ability to cope with associated chronic diseases. Using a validated quality-of-life instrument, we found significant improvements in five of eight mental and physical domains among patients in the gastric-bypass group and in two of eight domains among patients in the sleeve-gastrectomy group. Intensive medical therapy resulted in no significant improvements from baseline in quality of life.

Metabolic and weight-loss outcomes were generally similar in the two surgical groups at 1 year, although some advantages of gastric bypass over sleeve gastrectomy have emerged during longer follow-up, including a greater likelihood of reaching a glycated hemoglobin level of 7.0% or less (a therapeutic goal of the American Diabetes Association) with no use of diabetes medications, a reduced requirement for diabetes and cardiovascular medications, greater reductions in weight and BMI, and a greater improvement in quality of life. Some differences between the gastric-bypass group and the sleeve-gastrectomy group did not reach statistical significance, although the study was not adequately powered to detect modest differences between these procedures. In a prespecified substudy analysis of beta-cell function, insulin sensitivity, and body composition in a subgroup of patients, we found that at 2 years, gastric bypass was superior to sleeve gastrectomy with respect to insulin secretion, insulin sensitivity, and relative reduction in truncal fat as compared with subcutaneous fat.[20]

Most clinical guidelines and insurance coverage for bariatric surgery limit access to the surgery to patients with a BMI of 35 or more, presumably because of insufficient studies evaluating outcomes in patients with a BMI of less than 35. In our study, 49 of 137 patients (36%) had a BMI of 27 to 34, and these patients had an improvement in glycemic control and durability that was similar to that in patients with a BMI of 35 or more. Other trials and observational studies involving patients with only mild obesity showed similar improvements in glycemic control.[21,22] Accordingly, some guidelines for diabetes management are beginning to endorse the use of bariatric surgery in patients with diabetes and a BMI of 30 to 34, especially those who have poor glycemic control despite receiving the best available medical therapy.[23,24]

Important limitations of our study include an inadequate sample size and duration to detect differences in the incidence of diabetes complications, such as myocardial infarction, stroke, or death. The protocol specifies further follow-up at 5 years for all patients, which should allow additional assessment of even longer-term efficacy and safety.

Despite these limitations, we conclude that bariatric surgery represents a potentially useful strategy for the management of type 2 diabetes, allowing many patients to reach and maintain therapeutic targets of glycemic control that otherwise would not be achievable with intensive medical therapy alone. Some patients in our study had complete diabetes remission, whereas others had a marked reduction in the need for pharmacologic treatment. The reduction in cardiovascular risk factors was sustained, allowing for reductions in lipid-lowering and antihypertensive therapies. Other benefits of surgery included a significant improvement in the quality of life. The question as to whether the documented benefits will reduce microvascular and macrovascular morbidity and mortality, as shown in nonrandomized studies, can be adequately answered only through larg0065r, multicenter clinical-outcome trials.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgments

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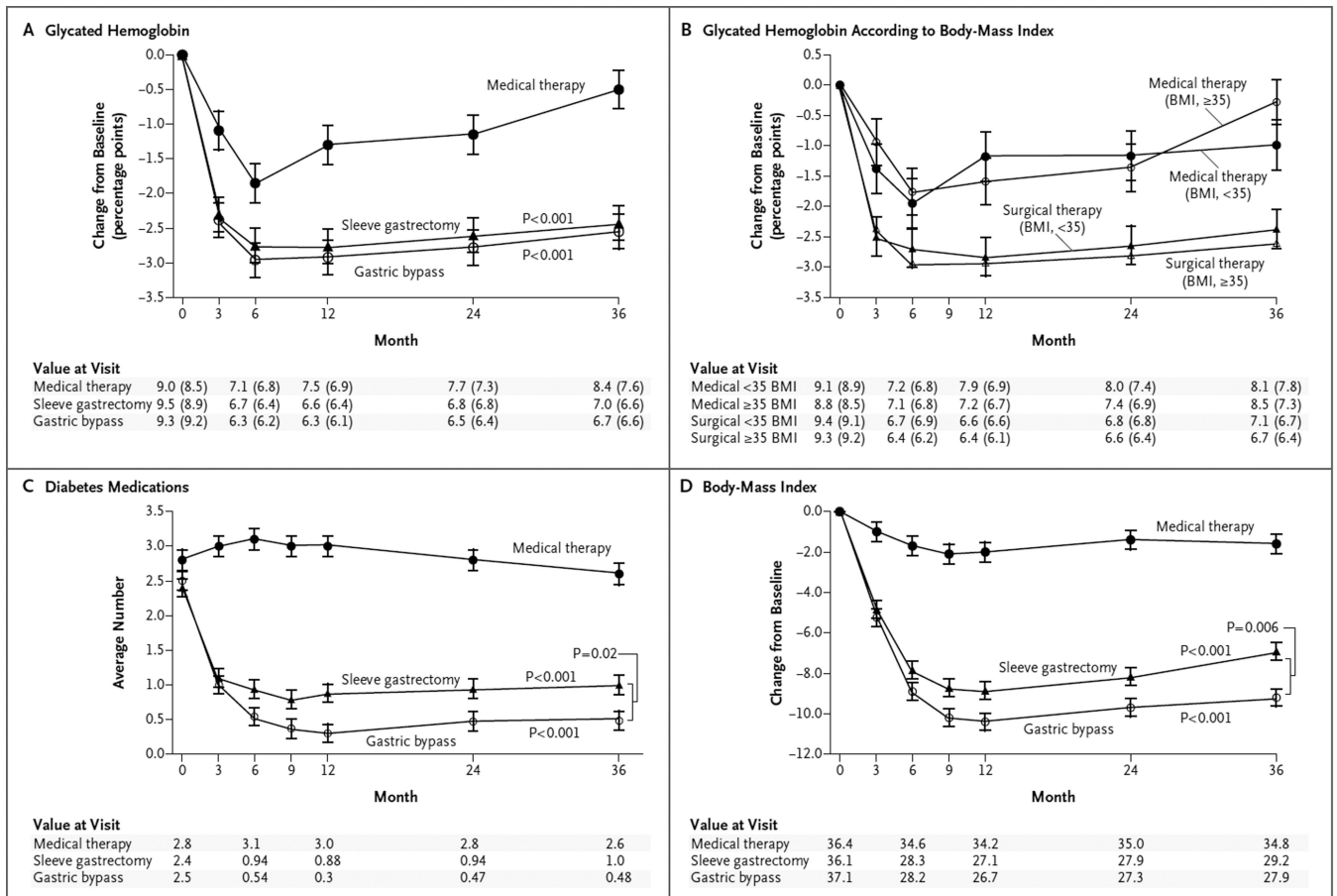
Figure 1. Mean Changes in Measures of Diabetes Control from Baseline to 3 Years
Shown are the percentage change in glycated hemoglobin levels (Panel A), the percentage change in glycated hemoglobin levels according to body-mass index (BMI) (Panel B), the average number of diabetes medications during the study period (Panel C), and the changes in BMI (Panel D) over a 3-year period among patients receiving intensive medical therapy only, sleeve gastrectomy, or gastric bypass. I bars indicate standard errors. Mean values in each group are provided below the graphs; in Panels A and B, median values are also provided in parentheses. P values are for the comparison between each surgical group and the medical-therapy group in Panels A, C, and D. In Panel B, P = 0.008 for the comparison between the surgical groups and the medical-therapy group for the subgroup of patients with a BMI of less than 35; P<0.001 for the comparison for the subgroup with a BMI of 35 or more.
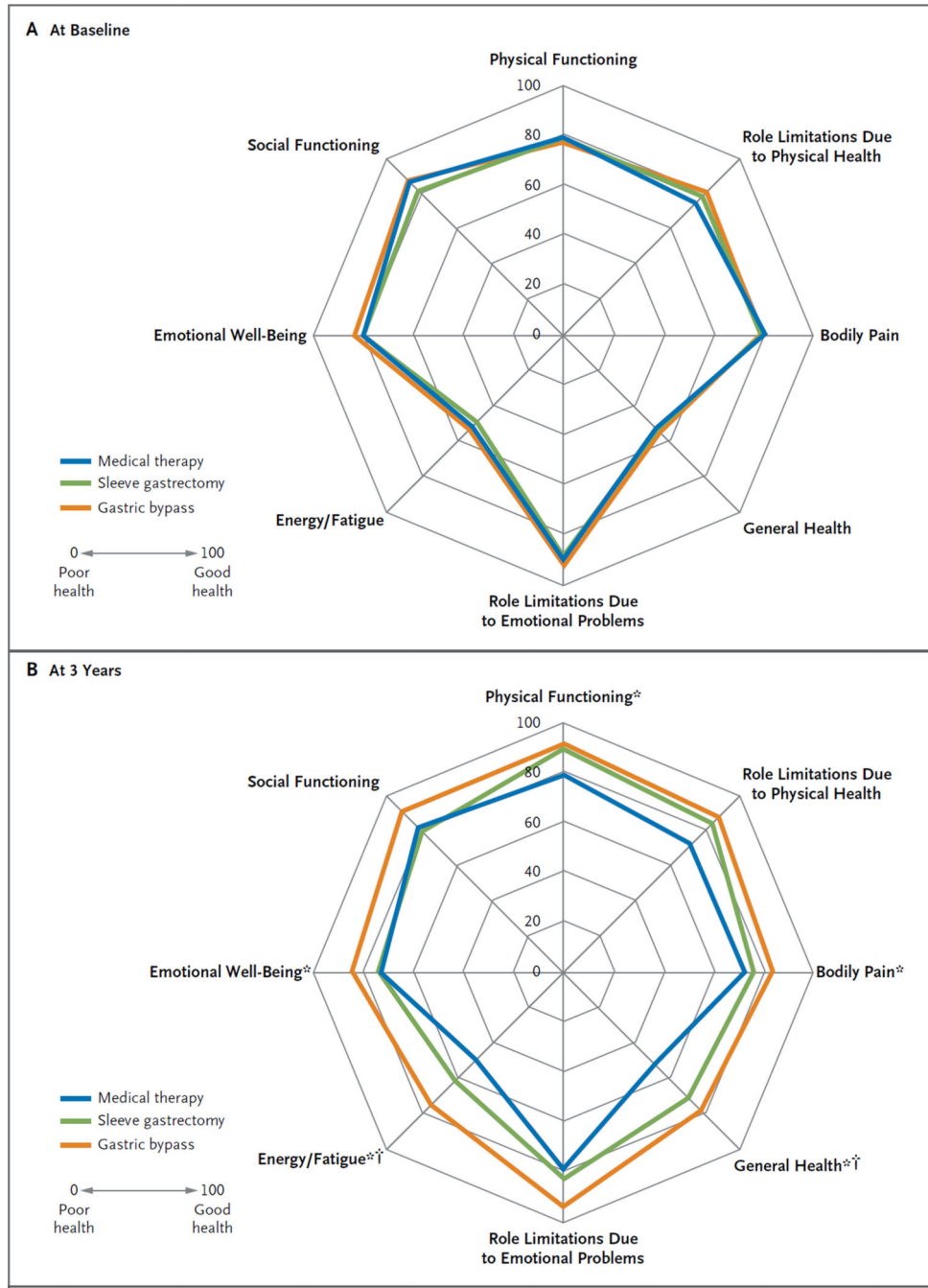
Figure 2. Polar Chart of Scores for Quality of Life at Baseline and 3 Years after Randomization
The scores on the RAND 36-Item Health Survey range from the worst score of 0 (poor health) to the best score of 100 (good health). Asterisks indicate P<0.05 for the comparison between the gastric-bypass group and the medical-therapy group; daggers indicate P<0.05 for the comparison between the sleeve-gastrectomy group and the medical-therapy group. The minimally important difference (MID) in scoring for this survey is unknown.

Table 1

Primary and Secondary End Points at 3 Years.[*]

End Point	Medical Therapy (N = 40)	Gastric Bypass (N = 48)	Sleeve Gastrectomy (N = 49)	P Value		
				Gastric Bypass vs. Medical Therapy	Sleeve Gastrectomy vs. Medical Therapy	Gastric Bypass vs. Sleeve Gastrectomy
Glycated hemoglobin						
Level — no. of patients (%)						
6%	2 (5)	18 (38)	12 (24)	<0.001	0.01	0.17
6% without diabetes medications	0	17 (35)	10 (20)	<0.001	0.002	0.10
6.5%	7 (18)	23 (48)	23 (47)	0.003	0.003	0.92
6.5% without diabetes medications	0	22 (46)	14 (29)	<0.001	<0.001	0.08
7%	16 (40)	31 (65)	32 (65)	0.02	0.02	0.94
7% without diabetes medications	0	28 (58)	16 (33)	<0.001	<0.001	0.01
At baseline — %						
Mean	9.0±1.4	9.3±1.4	9.5±1.7			
At 3 yr — %						
Mean	8.4±2.2	6.7±1.3	7.0±1.3	<0.001	<0.001	0.42
Median (IQR)	7.7 (6.7 to 9.6)	6.6 (5.8 to 7.4)	6.6 (6.1 to 7.5)			
Change from baseline — percentage points	−0.6±2.5	−2.5±1.9	−2.5±2.1	<0.001	<0.001	0.99
Relapse — no./total no. (%)[†]						
Glycemic control	4/5 (80)	5/21 (24)	9/18 (50)	0.03	0.34	0.09
Diabetes	NA	8/21 (38)	6/13 (46)	NA	NA	0.64
Median (IQR) fasting plasma glucose — mg/dl						

End Point	Medical Therapy (N = 40)	Gastric Bypass (N = 48)	Sleeve Gastrectomy (N = 49)	P Value		
				Gastric Bypass vs. Medical Therapy	Sleeve Gastrectomy vs. Medical Therapy	Gastric Bypass vs. Sleeve Gastrectomy
At baseline	157 (115 to 199)	193 (142 to 236)	164 (132 to 224)			
At 3 yr	132 (104 to 179)	100 (87 to 141)	106 (86 to 136)	0.001	0.007	0.92
Change from baseline†	−6.0 (−68.5 to 56.0)	−85.5 (−122.0 to −21.5)	−46.0 (−113.0 to −21.0)	0.001	0.006	0.24
Body weight						
At baseline — kg	104.5±14.2	106.8±14.9	100.6±16.5			
At 3 yr — kg	100.2±16.6	80.6±15.5	79.3±15.1	<0.001	<0.001	0.69
Change from baseline — kg	−4.3±8.8	−26.2±10.6	−21.3±9.7	<0.001	<0.001	0.02
% Change from baseline	−4.2±8.3	−24.5±9.1	−21.1±8.9	<0.001	<0.001	0.06
Other risk factors						
% Change from baseline in low-density lipoprotein cholesterol	2.5±29.9	16.9±54.5	14.5±52.2	0.14	0.20	0.82
% Change from baseline in high-density lipoprotein cholesterol	4.6±20.7	34.7±27.3	35.0±31.0	<0.001	<0.001	0.96
Median % change from baseline in triglycerides (IQR)	−21.5 (−45.4 to 16.4)	−45.9 (−61.0 to −7.5)	−31.5 (−52.1 to −6.9)	0.01	0.01	0.18
Change from baseline in blood pressure — mm Hg						
Systolic	0.63±22.63	1.29±20.38	−4.43±20.69	0.88	0.27	0.17
Diastolic	−6.48±12.33	−4.25±10.57	−6.27±13.30	0.36	0.94	0.41

*
Plus–minus values are means ±SD. To convert the values for glucose to millimoles per liter, multiply by 0.05551. To convert the values for cholesterol to millimoles per liter, multiply by 0.02586. To convert the values for triglycerides to millimoles per liter, multiply by 0.01129. IQR denotes interquartile range, and NA not applicable.

†
Relapse of glycemic control was defined as having met the primary end point for glycated hemoglobin of 6% or less at 1 year but not at 3 years. Relapse of diabetes was defined as having met the primary end point for glycated hemoglobin of 6% or less with the use of no antidiabetic medications at 1 year but not at 3 years. None of the patients in the medical-therapy group had a complete remission of diabetes, and thus, these patients were not evaluated for relapse.

†For skewed data, the median of the change is not the numerical difference between the group-level median at baseline and the median value at 3 years.

Table 2

Medication Use at Baseline and at 3 Years. *

Medication	At Baseline			At 3 Years		
	Medical Therapy (N = 40)	Gastric Bypass (N = 48)	Sleeve Gastrectomy (N = 49)	Medical Therapy (N = 40)	Gastric Bypass (N = 48)	Sleeve Gastrectomy (N = 49)
Diabetes medications						
No. of medications	2.80±1.11	2.50±1.15	2.45±1.19	2.60±1.10	0.48±0.80 [†]	1.02±1.01 [†‡]
Insulin — no. of patients (%)	21 (52)	22 (46)	22 (45)	22 (55)	3 (6) [†]	4 (8) [†]
Not taking this class of medication — no. of patients (%)	1 (2)	1 (2)	1 (2)	1 (2)	33 (69) [†]	21 (43) [†‡]
Cardiovascular medications						
No. of medications	2.70±1.22	2.73±1.32	2.18±1.09	2.63±1.31	0.96±1.15 [†]	1.35±1.40 [†]
ACE inhibitor or ARB — no. of patients (%)	25 (62)	36 (75)	30 (61)	22 (55)	11 (23) [§]	13 (27) [§]
Not taking this class of medication — no. of patients (%)	0	3 (6)	2 (4)	1 (2)	20 (42) [†]	19 (39) [†]
Any medication						
No. of medications	5.50±1.71	5.23±1.76	4.63±1.67	5.23±1.86	1.44±1.49 [†]	2.37±1.82 [†‡]
Difference from baseline to 3 yr — no.				−0.28±2.03	−3.79±1.81 [†]	−2.27±1.99 [†‡]

*
Plus–minus values are means ±SD. All P values are for the comparison with the medical-therapy group unless otherwise indicated. ACE denotes angiotensin-converting enzyme, and ARB angiotensin-receptor blocker.

[†]
P<0.001.

[‡]
P<0.05 for comparison between gastric bypass and sleeve gastrectomy.

[§]
P<0.01.

Table 3

Complications at 3 Years.[*]

Complication	Medical Therapy (N = 43)	Gastric Bypass (N = 50) *number of patients (percent)*	Sleeve Gastrectomy (N = 49)
Gastrointestinal			
Bowel obstruction	1 (2)	1 (2)	1 (2)
Stricture	0	1 (2)	1 (2)
Ulcer	1 (2)	4 (8)	0
Leak	0	0	1 (2)
Intraabdominal bleeding	0	2 (4)	0
Dumping syndrome	0	4 (8)	1 (2)
Gallstone diseases	0	1 (2)	1 (2)
Microvascular or macrovascular			
Stroke	0	0	1 (2)
Retinopathy	0	1 (2)	2 (4)
Nephropathy [†]	4 (9)	7 (14)	5 (10)
Foot ulcer	0	2 (4)	1 (2)
Nutritional and metabolic			
Anemia	6 (14)	8 (16)	15 (31)
Intravenous treatment for dehydration	3 (7)	7 (14)	4 (8)
Hypoglycemic episode	39 (91)	32 (64)	40 (82)
Severe hypoglycemia requiring intervention	0	1 (2)	0
Excessive weight gain[‡]	7 (16)	0	0
Other			
Wound infection	0	1 (2)	0
Hernia	1 (2)	3 (6)	1 (2)
Pneumonia	0	2 (4)	1 (2)
Renal calculus	6 (14)	5 (10)	4 (8)
Cancer	2 (5)	2 (4)	2 (4)

[*] Not included in the safety analysis were seven patients in the medical-therapy group who withdrew immediately after randomization and one patient in the sleeve-gastrectomy group who had anemia before withdrawing from the study before surgery. Five patients who started the study but later withdrew or were lost to follow-up are included in this analysis until their discontinuation.

[†] Nephropathy was defined as any one of the following criteria: doubling of the serum creatinine level or a decrease in the glomerular filtration rate of more than 20%; development of macroalbuminuria (urine albumin-to-creatinine ratio, >300 [as measured in milligrams of albumin to grams of creatinine]); or renal transplantation, initiation of dialysis, or an increase in the serum creatinine level of more than 3.3 mg per deciliter (290 μmol per liter) in the absence of an acute reversible cause.

[‡] Excessive weight gain was defined as a 5% increase in body weight over baseline. $P < 0.05$ for the comparison between the medical-therapy group and each of the surgical groups.