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ABSTRACT

BACKGROUND AND PURPOSE: Stent-assisted coiling may result in less aneurysm recanalization but more complications than coiling alone. We evaluated outcomes of coiling with and without stents in the multicenter Matrix and Platinum Science Trial.

MATERIALS AND METHODS: All patients in the Matrix and Platinum Science Trial with unruptured intracranial aneurysms treated per protocol were included. Baseline patient and aneurysm characteristics, procedural details, neurologic outcomes, angiographic outcomes, and safety data were analyzed.

RESULTS: Overall, 137 of 361 (38%) patients were treated with a stent. Stent-coiled aneurysms had wider necks (≥4 mm in 62% with stents versus 33% without, $P < .0001$) and lower dome-to-neck ratios (1.3 versus 1.8, $P < .0001$). Periprocedural serious adverse events occurred infrequently in those treated with and without stents (6.6% versus 4.5%, $P = .39$). At 1 year, total significant adverse events, mortality, and worsening of mRS were similar in treatment groups, but ischemic strokes were more common in stent-coiled patients than in coiled patients (8.8% versus 2.2%, $P = .005$). However, multivariate analysis confirmed that at 2 years after treatment, prior cerebrovascular accident (OR, 4.7; $P = .0089$) and aneurysm neck width ≥4 mm (OR, 4.5; $P = .02$) were the only independent predictors of ischemic stroke. Stent use was not an independent predictor of ischemic stroke at 2 years (OR, 1.1; $P = .94$). Stent use did not predict target aneurysm recurrence at 2 years, but aneurysm dome size ≥10 mm (OR, 9.94; $P < .0001$) did predict target aneurysm recurrence.

CONCLUSIONS: Stent-coiling had similar outcomes as coiling despite stented aneurysms having more difficult morphology than coiled aneurysms. Increased ischemic events in stent-coiled aneurysms were attributable to baseline risk factors and aneurysm morphology.

ABBREVIATIONS: MAPS = Matrix and Platinum Science Trial; TAR = target aneurysm recurrence

As intracranial aneurysm treatment has shifted in the past 30 years from exclusively surgical to predominantly endovascular, aneurysm morphologies once considered untreatable endovascularly are now treatable with coils, stents, and flow diverters.[1-3] Particularly for saccular aneurysms with broad necks and short domes, stent-assisted coiling has become a common technique.[4-8] Prior studies have reported that stent-coiling may result in less aneurysm recanalization over time but more complications—both intraprocedurally and in a delayed fashion—than coiling alone.[9-13] A recent large, single-institution, retrospective series described higher morbidity and mortality rates associated with the stent-coiling technique as compared with coiling either with or without balloon assistance.[14]

Given that prospective data on stent-coiling are limited, we analyzed data from the prospective, randomized, multicenter Matrix and Platinum Science (MAPS) Trial (NCT00396981, www.clinicaltrials.gov). The MAPS Trial was primarily designed to determine whether polymer-modified coils or platinum bare metal coils result in lower aneurysm recanalization, lower aneurysm rupture or rerupture, or lower aneurysm retreatment. Although patients were

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From the Departments of Radiology and Biomedical Imaging (S.W.H., C.F.D.) and Neurology (S.C.J.), University of California, San Francisco, San Francisco, California; Department of Radiology (A.T.), Medical University of South Carolina, Charleston, South Carolina; Departments of Neurology and Radiology (J.D.E.), California Pacific Medical Center, San Francisco, California; Department of Neurosurgery (J.M.), Vanderbilt University Medical Center, Nashville, Tennessee; Department of Neurosurgery (C.P.), University of Medicine and Dentistry of New Jersey, Newark, New Jersey; Department of Radiology (G.N., S.B.), Oregon Health and Science University, Portland, Oregon; Department of Biostatistics (S.G.G., J.N.J.), Stryker Neurovascular (K.C.), Fremont, California; Department of Neurosurgery (Y.M.), Jikei University, Tokyo, Japan; Department of Neuroradiology (A.G.), Newcastle Upon Tyne Hospital, Newcastle Upon Tyne, United Kingdom; Department of Neurosurgery (D.L.), Rush University Medical Center, Chicago, Illinois; and Department of Neurosurgery (C.M.), Barrow Neurological Institute, Phoenix, Arizona.

Please address correspondence to Steven W. Hetts, MD, Interventional Neuroradiology, 505 Parnassus Ave, L-352, San Francisco, CA 94143–0628; e-mail: steven.hetts@radiology.ucsf.edu

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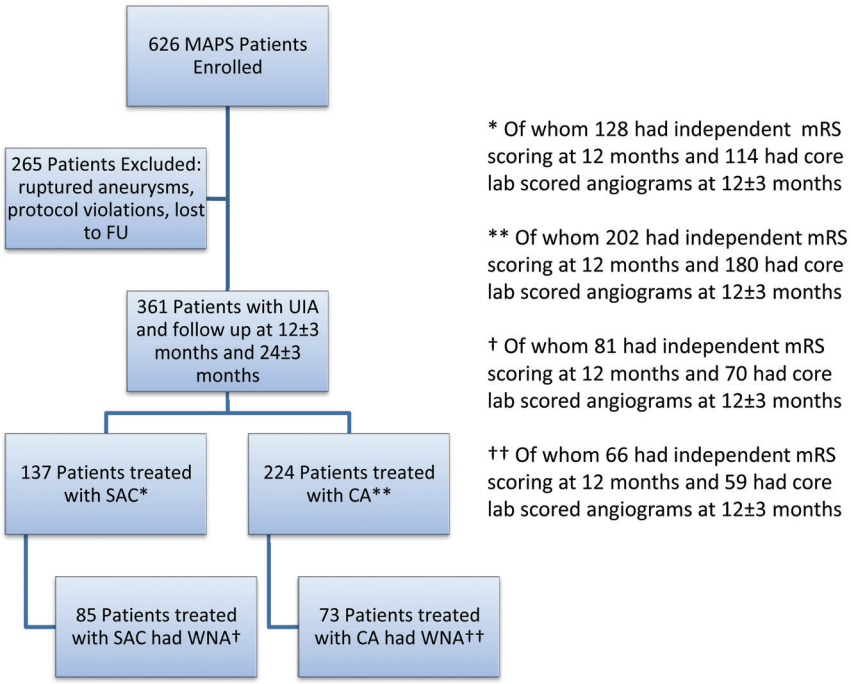

FIG 1. Patient flow for MAPS stent substudy. FU indicates follow-up; UIA, unruptured intracranial aneurysm; SAC, stent-assisted coiling; CA, coiling alone; WNA, wide-neck aneurysm.

randomly assigned to platinum bare metal coil or polymer-modified coil implantation, adjunctive devices (including balloons and stents) could be used in any case at the discretion of the operating physician.

We compared baseline patient and aneurysm characteristics, procedural details, safety data, neurologic outcomes, and angiographic outcomes in MAPS patients with unruptured intracranial aneurysms who were treated with stent-coiling or coiling without stent placement. Additionally, we analyzed results for the subset of patients with wide-neck aneurysms.

MATERIALS AND METHODS

The MAPS Trial was conceived and designed by the investigators, with advice provided by the sponsor, and was approved by all local institutional review boards. The study was conducted in accordance with the *International Conference on Harmonization (ICH) Good Clinical Practice (GCP): Consolidated Guideline*, the Declaration of Helsinki, EN ISO14155 Clinical Investigations of Medical Devices for Human Subjects, and the applicable regulations from the US Food and Drug Administration. The primary end point of the MAPS Trial was target aneurysm recurrence (TAR), designed to be a clinically relevant composite end point that comprised aneurysm rupture after treatment, retreatment, or death from an unknown cause. The trial was designed to study the TAR rate for the study device and to investigate how TAR correlated with the angiographic surrogates that are widely used in endovascular aneurysm treatment studies to evaluate outcomes. Clinical and angiographic evaluations were completed at the time of initial aneurysm treatment and within 12 ± 3 months after initial aneurysm treatment. Further clinical follow-up of subjects by telephone interview will continue annually until 5 years after initial aneurysm treatment.

Study Subjects

The study population for the current analysis included subjects 18–80 years of age with a baseline mRS score of 0–3 who had a single documented, untreated, unruptured intracranial aneurysm (4–20 mm in maximum dimension on DSA) for which both polymer modified coils (Matrix2, Stryker Neurovascular, Fremont, California) and platinum bare metal coils (GDC, Stryker Neurovascular) were treatment options and for which primary coiling treatment was planned to be completed during a single procedure. Stent placement (Neuroform stent, Stryker Neurovascular), as a separate preliminary procedure antecedent to the primary coiling, was allowed, as was stent placement in conjunction with the primary coiling procedure. Although the overall MAPS Trial prospectively enrolled patients with ruptured and unruptured aneurysms, only patients with unruptured aneurysms were included in the present post hoc data analysis.

We chose to analyze the unruptured aneurysm cohort because the stent-coiling technique is primarily applied in clinical practice to patients with unruptured aneurysms. The administration of dual antiplatelet medications typically indicated in stent-coiling is relatively contraindicated in patients with ruptured aneurysm who might need additional interventions such as ventricular drain placement. On this basis, all patients with ruptured aneurysms in the MAPS trial were excluded from our current analysis, consisting of 6 patients treated with stent-coiling and 201 patients treated with coiling (Fig 1).

A total of 361 patients in the MAPS Trial with unruptured intracranial aneurysms were treated per protocol. Data were analyzed post hoc for all unruptured intracranial aneurysms and for the wide-neck (≥4 mm) aneurysm subgroup of unruptured intracranial aneurysms. Note that because this is a post hoc analysis of the MAPS Trial, patients were not randomly assigned to stent-coiling or coiling and may be dissimilar, especially within the cohort including all unruptured intracranial aneurysms.

As defined above, the primary outcome measure was TAR. Secondary outcome measures, all defined a priori, included angiographic assessment as assessed by enrolling sites and core imaging laboratory; neurologic assessments (mRS at 12 ± 3 months and as change from baseline performed in-person by an independent certified practitioner at a scheduled clinic visit); and technical procedural success, defined as the successful placement of coils in the target aneurysm. Target aneurysm reintervention was defined as any further treatment of the aneurysm, with the retreatment decision being at the discretion of the operator.

All sites graded their own angiographic outcomes on the basis of the modified 3-point Raymond Scale after the procedure and at follow-up.[15-17] All sites also recorded an assessment of perceived change from baseline (same, better, worse) at follow-up. Digitized

copies of the angiograms were created for all cases and stored at an independent angiographic core laboratory located at the University of California, San Francisco. The core laboratory assessed all treatment and 1-year follow-up angiograms blinded to the treatment technique. Core laboratory evaluations used the same angiographic scales as did the enrolling sites. All angiographic data presented in this analysis are from core laboratory evaluations performed by 2 neurointerventionalists, with adjudication of differences in angiographic scoring by a third neurointerventionalist.[17] The core laboratory also evaluated angiograms of patients with wide-neck aneurysms receiving stent-coiling specifically for any stent migration between immediate postprocedure DSA and follow-up DSA.

An independent steering committee was responsible for overall oversight of the science and execution of the trial. Patient safety data were reviewed at regular intervals by an independent Data Monitoring Committee. An independent Clinical Events Committee was responsible for reviewing and adjudicating all deaths and neurologic events. On-site monitoring and source document verification of case report forms against original patient records were completed for more than 40% of patients at the completion of the 1-year follow-up, including all patients who had been treated with stent-coiling.

Statistical Methods

The primary end point (TAR) rate was calculated by use of Kaplan-Meier estimates in each group at the end of a 12 ± 3–month window (455 days) and a 24 ± 3–month window (820 days). Time to event was based on the real time to rupture/rerupture, retreatment, or unknown cause of death, whichever happened first for each subject. Subjects who had not had an event were censored at their last clinical visit or at 820 days, whichever came earlier.

The protocol prespecified additional univariate and multivariate regression models to analyze the time to TAR, changes in mRS from baseline to the 12-month assessment, and subgroup analyses. Additional post hoc multivariate regression analysis was performed to evaluate the contribution of baseline risk factors, aneurysm characteristics, and use of stents to stroke rates and target aneurysm recurrence. Complete data on ischemic stroke and TAR rates were included in predictor analyses, including data from all enrolling centers.

A Student *t* test was used to test distributions of continuous variables between the groups. Either χ^2 or the Fisher exact test was used to analyze binary variables according to standard statistical practice. For ordinal variables, such as the mRS, recanalization, and mRS scores, the Wilcoxon rank sum test was used to test the distribution between the groups. The differences between the groups were presented with the 95% CI estimated by the normal approximation. For the binary outcomes, the relative risks as well as its 95% CIs were also presented. All statistical analyses were performed with the use of SAS, version 9.2 (SAS Institute, Cary, North Carolina).

RESULTS

Of 626 patients in the MAPS Trial, overall, 361 with unruptured intracranial aneurysms were treated per protocol (Fig 1). Of these 361 patients with unruptured intracranial aneurysms, 137 were treated with a Neuroform stent and either platinum bare metal coils or polymer modified coils (stent-coiling group) and 224 were treated with platinum bare metal coils or polymer modified coils without a stent (coiling group). Within the unruptured intracranial aneurysm cohort, 158 patients had wide-neck aneurysms with necks ≥4 mm in diameter (wide-neck aneurysm subset). Within this wide-neck aneurysm subset, 85 patients received stent-coiling and 73 received coiling. Because stent use was at the operating physician's discretion on a case-by-case basis, some enrolling sites used many stents and some used none (Fig 2). Stent use was particularly inhomogeneous geographically: 88.3% of stent-coiling cases were performed in North American centers versus 65.6% of coiling cases without stent use ($P = .0001$, Table 1).

Baseline Demographics

Patients with stent-coiling trended toward being sicker at baseline than did coiling patients in the entire unruptured intracranial aneurysm cohort (Table 1), with coronary artery disease in 19% of stent-coiling versus 13.1% of coiling ($P = .14$). Coronary artery disease was significantly more frequent among the patients with stent-coiling with wide-neck aneurysms (22.4%) as compared with patients with wide-neck aneurysms treated with coiling (5.6%, $P = .003$). Prior stroke was also marginally more frequent in the patients with stents and coils (17.9%) versus the patients with coils alone in the wide-neck aneurysms (8.3%, $P = .08$). Otherwise, both groups were similar at baseline.

Aneurysm Characteristics

Aneurysms in the stent-coiling group had more technically challenging morphologies than those in the coiling group (Table 2). Among unruptured intracranial aneurysms, 62% of stent-coiling versus 33% of coiling were wide-neck aneurysms ($P < .0001$), and the dome-to-neck ratio was lower for stent-coiling than for coiling (1.3 versus 1.8, $P < .0001$). Among wide-neck aneurysms, patients receiving stent-coiling had smaller aneurysms (maximum dimension >10 mm in 30.6% versus 47.9%, $P = .03$), with larger necks (mean, 5.6 versus 5.0 mm; $P = .004$), and lower dome-to-neck ratios (1.2 versus 1.6, $P < .0001$). In both the overall unruptured intracranial aneurysm cohort as well as the wide-neck aneurysm subset, aneurysms treated with stent-coiling were less likely to be located on the circle of Willis than those treated with coiling.

Procedural Characteristics

Among patients with unruptured intracranial aneurysms, the stent-coiling procedure took slightly longer than did the coiling procedure (mean, 134.2 versus 117.8 minutes; $P = .02$). This time difference was not significant in the more technically challenging wide-neck aneurysm subset (mean, 147.5 versus 135.2 minutes; $P = .30$) (Table 3).

For unruptured intracranial aneurysms, coil packing attenuation trended higher with stent-coiling (26.2% versus 24.2%, $P = .07$). Among wide-neck aneurysms, packing attenuation for stent-coiling was higher than packing attenuation for coiling without stent placement (26.4% versus 21.1%, $P = .002$) despite a trend

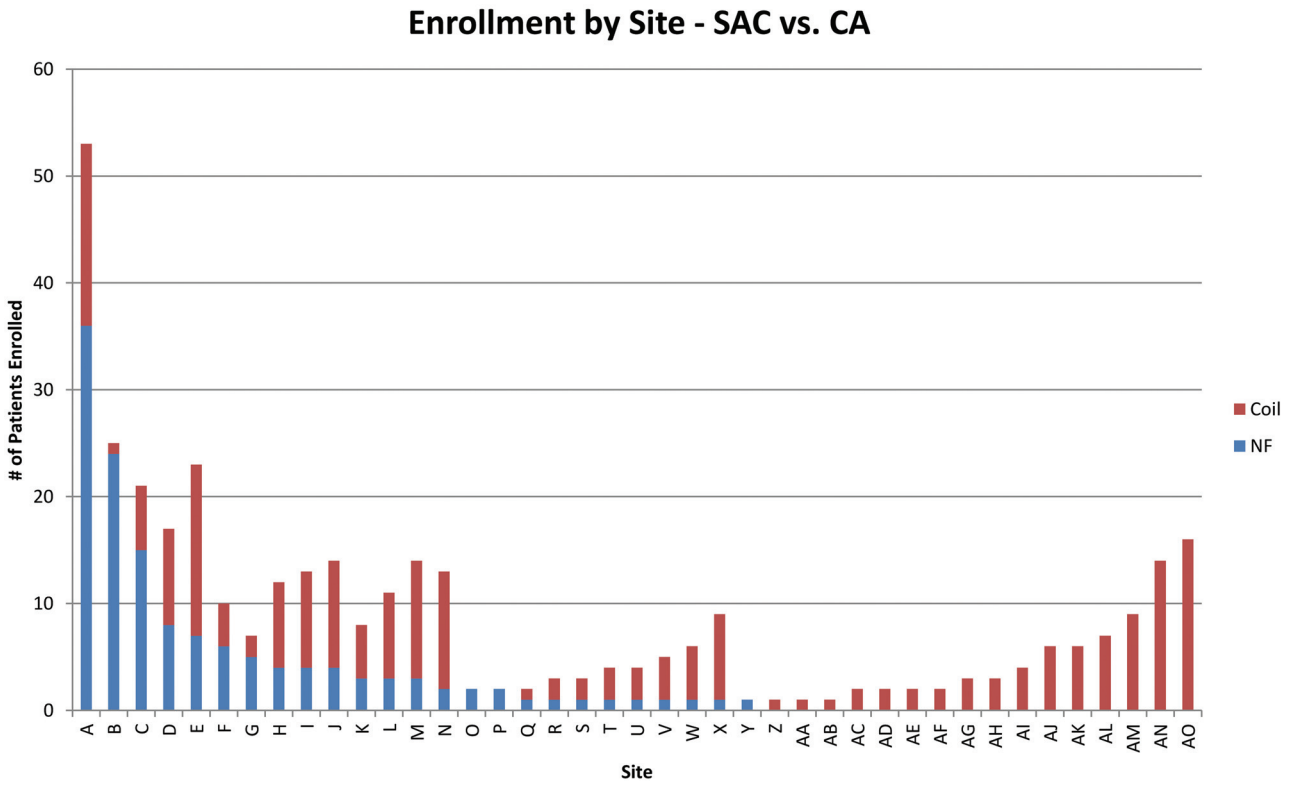

FIG 2. Stent use by enrolling site. SAC indicates stent-assisted coiling, CA, coiling alone, NF, Neuroform stent.

Table 1: Baseline patient demographics

	All Unruptured Aneurysms			Wide-Neck Aneurysms		
	Stent-Coil (*n* = 137)	Coil Alone (*n* = 224)	*P* Value	Stent-Coil (*n* = 85)	Coil Alone (*n* = 73)	*P* Value
North American	88.3%	65.6%	.0001	89.4%	61.6%	<.0001
Age, y	56.5	56.7	.90	58.4	57.7	.68
Female	76.6	76.8	.98	72.9	79.5	.34
Coronary artery disease	19.0%	13.1%	.14	22.4%	5.6%	.003
≥2 Cerebrovascular risk factors[a]	32.1%	25.9%	.20	34.1%	26.0%	.27
Prior CVA	16.9%	14.5%	.54	17.9%	8.3%	.08
Preprocedure mRS						
0	80.3%	84.4%	>.99	80.0%	90.4%	NA
1	15.3%	11.6%		14.1%	9.6%	
2	4.4%	3.6%		5.9%	0.0%	
3	0.0%	0.4%	>.99	0.0%	0.0%	NA

Note:—CVA indicates cerebrovascular accident; NA, not applicable.

[a] Cerebrovascular risk factors include hypertension, hyperlipidemia, coronary artery disease, and diabetes mellitus.

Table 2: Aneurysm characteristics

	All Unruptured Aneurysms			Wide-Neck Aneurysms		
	Stent-Coil (*n* = 137)	Coil Alone (*n* = 224)	*P* Value	Stent-Coil (*n* = 85)	Coil Alone (*n* = 73)	*P* Value
Maximum dimension	7.6	7.8	.46	8.9	10.1	.007
Height is max	43.8%	54.9%	.04	42.4%	50.7%	.30
Width is max	29.9%	21.4%	.07	30.6%	23.3%	.30
Depth is max	26.3%	23.7%	.58	27.1%	26.0%	.88
Max dimension >10 mm	19.7%	21.0%	.77	30.6%	47.9%	.03
Neck ≥4 mm	62.0%	32.6%	<.0001	100%	100%	NA
Average neck	4.7	3.5	<.0001	5.6	5.0	.004
Dome/neck radio	1.3	1.8	<.0001	1.2	1.6	<.0001
Dome/parent artery ratio	2.0	2.1	.33	2.2	2.6	.01
Circle of Willis location	44.5%	64.7%	.0002	48.2%	72.6%	.002

Note:—Max indicates maximum.

toward use of fewer coils (8.5 mean versus 10.1 mean, $P = .07$). Overall technical success was high for all groups but marginally lower for stent-coiling in the unruptured intracranial aneurysm cohort (97.8%) as compared with coiling in the unruptured intracranial aneurysm cohort (100%, $P = .054$). Among unruptured intracranial aneurysms, complete angiographic obliteration

Table 3: Procedural Characteristics

	All Unruptured Aneurysms			Wide-Neck Aneurysms		
	Stent-Coil (*n* = 137)	Coil Alone (*n* = 224)	*P* Value	Stent-Coil (*n* = 85)	Coil Alone (*n* = 73)	*P* Value
Aspirin alone	8.0%	23.2%	.0002	4.7%	28.8%	<.0001
Clopidogrel alone	5.1%	4.5%	.78	3.5%	1.4%	0.62
Aspirin + clopidogrel	78.1%	37.1%	<.0001	82.4%	37.0%	<.0001
Antiplatelet use not recorded[a]	8.8%	35.3%	<.0001	9.4%	32.9%	.0003
Procedure time, min	134.2	117.8	.02	147.5	135.2	.30
No. of coils	6.9	7.1	.70	8.5	10.1	.07
Bare metal coils	49.6%	50.9%	.82	48.2%	53.4%	.52
Matrix coils	50.4%	49.1%	.82	51.8%	46.6%	.52
Packing density	26.2%	24.0%	.07	26.4%	21.1%	.002
Technical success	97.8%	100%	.054	97.6%	100%	.50
Occlusion assessment						
Raymond 1	21.1%	33.9%	.02[b]	18.6%	27.1%	.25[c]
Raymond 2	25.4%	25.6%	.98[b]	27.1%	30.5%	.67[c]
Raymond 3	53.5%	40.6%	.03[b]	54.3%	42.4%	.18[c]

[a] No antiplatelet use recorded in these subjects: for an individual subject, this could mean no aspirin or clopidogrel was used or that data are missing.
[b] Core lab occlusion assessment for the 81% of patients who had assessable angiograms after the procedure.
[c] Core lab occlusion assessment for the 82% of patients who had assessable angiograms after the procedure.

Table 4: Safety: Stroke and Other Significant Adverse Events

	All Unruptured Aneurysms			Wide-Neck Aneurysms		
	Stent-Coil (*n* = 137)	Coil Alone (*n* = 224)	*P* Value	Stent-Coil (*n* = 85)	Coil Alone (*n* = 73)	*P* Value
Periprocedural significant adverse event rate	6.6%	4.5%	.39	3.7%	0.89%	.11
1-Year hemorrhagic stroke rate	2.9%	0.4%	.07	2.4%	0.0%	.50
1-Year ischemic stroke rate	8.8%	2.2%	.005	11.8%	4.1%	.08
if problem site is excluded[a]						
1-Year ischemic stroke rate	6.2%	2.2%	.11	8.8%	4.1%	.31

[a] One enrolling site accounted for 5 of 12 subjects with ischemic stroke in the unruptured aneurysm group. All ischemic strokes at that site occurred ≥7 days after the procedure.

Table 5: Safety: mRS at baseline and 1 year after the procedure

	All Unruptured Aneurysms					Wide-Neck Aneurysms				
	Stent-Coil		Coil Alone			Stent-Coil		Coil Alone		
12-Month mRS[a]	Base (*n* = 137)	1 Year (*n* = 128)	Base (*n* = 224)	1 Year (*n* = 202)	*P* Value	Base (*n* = 85)	1 Year (*n* = 81)	Base (*n* = 73)	1 Year (*n* = 66)	*P* Value
0	80.3%	80.5%	84.4%	87.6%		80.0%	80.2%	90.4%	92.4%	
1	15.3%	14.1%	11.6%	6.9%		14.1%	12.3%	9.6%	6.1%	
2	4.4%	1.6%	3.6%	1.0%		5.9%	2.5%	0.0%	0.0%	
3	0.0%	1.6%	0.4%	2.5%		0.0%	2.5%	0.0%	1.5%	
6	0.0%	2.3%	0.0%	2.0%		0.0%	2.5%	0.0%	0.0%	
mRS worse than base		12.5%		8.4%	.23		13.6%		4.5%	.06

[a] Twelve-month mRS scores were available for 128, 202, 81, and 66 patients, respectively, across all subsets.

of aneurysm filling at the conclusion of treatment was lower for stent-coiling than for coiling (21.1% versus 33.9%, $P = .02$). Although a similar trend was present in the wide-neck aneurysm subset, it did not reach statistical significance. Dual-antiplatelet use at the time of treatment was higher for stent-coiling than for coiling because it is the practice of most centers to use both aspirin and clopidogrel at the time of stent-coiling and for a variable period thereafter.

Safety: Stroke, Other Significant Adverse Events, and Neurologic Disability

Although total periprocedural significant adverse events did not differ between stent-coiling and coiling (6.6% versus 4.5%, $P = .39$), the rate of stroke within 1 year of treatment did differ between these groups (8.8% versus 2.2%, respectively; $P = .005$, Table 4). Of note, 42% of stent-coiling ischemic strokes occurred at one enrolling site, the exclusion of which brings the comparative ischemic stroke rates to 6.2% versus 2.2%, respectively ($P = .11$). Within the wide-neck aneurysm subgroup, there was no significant difference in ischemic stroke rates between stent-coiling and coiling, though there was a trend toward more ischemia in the stented patients ($P = .08$). Among all patients with unruptured intracranial aneurysms, 1-year hemorrhagic strokes also trended toward a higher rate in the stent-coiling group (2.9%) compared with the coiling group (0.4%, $P = .07$). At 1 year, total significant adverse events, mortality, and worsening of mRS scores were not different between stent-coiling and coiling (Table 5) in the entire unruptured intracranial aneurysm cohort, though there was a trend toward worsening mRS scores in the wide-neck aneurysm subset ($P = .06$).

Five stent-coiled patients had periprocedural strokes (<7 days after the procedure) and 7 stent-coiled patients had delayed strokes (On-line Table). Of the periprocedural strokes, one was thought related to preprocedure angioplasty or intraprocedure parent artery coil prolapse. A patient with extensive cardiovascular risk factors had both a periprocedural

Table 6: Clinical outcomes 1 year after the procedure

	All Unruptured Aneurysms			Wide-Neck Aneurysms		
	Stent-Coil (*n* = 137)	Coil Alone (*n* = 224)	*P* Value	Stent-Coil (*n* = 85)	Coil Alone (*n* = 73)	*P* Value
TAR	8.8%	8.5%	.93	14.1%	13.7%	.94
Delayed bleed	0.0%	0.4%	<.99	0.0%	1.4%	.46
Retreatment	8.8%	8.5%	.93	14.1%	13.7%	.94

Table 7: Angiographic outcomes 1 year after the procedure

	All Unruptured Aneurysms			Wide-Neck Aneurysms		
	Stent-Coil (*n* = 114)	Coil Alone (*n* = 180)	*P* Value	Stent-Coil (*n* = 70)	Coil Alone (*n* = 59)	*P* Value
Occlusion assessment						
Raymond 1	51.8%	44.4%	.22	45.7%	27.1%	.03
Raymond 2	21.1%	23.9%	.57	17.1%	30.5%	.07
Raymond 3	27.2%	31.7%	.41	37.1%	42.4%	.55
Change assessment						
Better	51.8%	31.1%	.0004	45.7%	20.3%	.003
Same	31.6%	35.6%	.48	32.9%	28.8%	.62
Worse	16.7%	33.3%	.002	21.4%	50.8%	.0005

Note:—81% of subjects in the "all unruptured aneurysms" group and 82% of subjects in the "wide-neck aneurysms" subgroup had angiograms assessable by the core lab at 1-year follow-up.

Table 8: Multivariate predictors of ischemic stroke at 1 year and 2 years

Parameter	1-Year OR (95% CI)	1-Year *P* Value	2-Year OR (95% CI)	2-Year *P* Value
Prior cerebrovascular accident	3.84 (1.29–11.4)	.0159	4.71 (1.47–15.0)	.0089
Aneurysm neck size ≥4 mm	3.70 (1.09–12.5)	.0359	4.51 (1.27–16.0)	.0196
Stent used	1.85 (0.61–5.59)	.2732	1.05 (0.34–3.27)	.9351

Note:—Complete data on ischemic stroke rates were included in predictor analysis, including data from all enrolling centers.

Table 9: Multivariate predictors of target aneurysm recurrence at 1 year and 2 years

Parameter	1-Year OR (95% CI)	1-Year *P* Value	2-Year OR (95% CI)	2-Year *P* Value
Aneurysm dome size ≥10 mm	10.1 (4.06–24.9)	<.0001	9.94 (4.12–24.0)	<.0001
Aneurysm neck size ≥4 mm	2.34 (0.94–5.81)	.0664	2.17 (0.93–5.06)	.0729
Stent used	0.89 (0.38–2.10)	.7855	0.83 (0.36–1.88)	.6505

Note:—Complete data on target aneurysm recurrence rates were included in predictor analysis, including data from all enrolling centers.

stroke and a myocardial infarction and died. Three patients had delayed strokes after documented or suspected antiplatelet medication noncompliance. One patient each had a delayed stroke after hernia surgery, after aneurysm retreatment, and after surveillance angiography. One patient with a basilar tip aneurysm had a pontine infarct thought to be related to small-vessel ischemic disease.

Outcomes at 1 Year and 2 Years

Clinical outcomes at 1 year and 2 years, on the basis of the primary MAPS composite end point of TAR, were similarly excellent in stent-coiling and coiling patients in the unruptured intracranial aneurysm cohort as well as in the wide-neck aneurysm subset (Table 6). Only a single patient from the coiling group had aneurysm rupture; all remaining TAR events were aneurysm retreatments at the discretion of the operating physician.

Complete angiographic obliteration rates for stent-coiling were significantly higher than for coiling in the wide-neck aneurysm subset at 1 year (Raymond 1 occlusion, 45.7% versus 27.1%, $P = .03$) (Table 7). Angiographic worsening (on the better-same-worse scale comparing 12 ± 3–month follow-up angiograms with immediate posttreatment angiograms) was lower for stent-coiling than for coiling in both the unruptured intracranial aneurysm (16.7% versus 33.3%, $P = .002$) and wide-neck aneurysm groups (21.4% versus 50.8%, $P = .0005$). Concomitantly, angiographic improvement at 1 year in treated aneurysms was more common for stent-coiling versus coiling in the unruptured intracranial aneurysm cohort (51.8% versus 31.1%, $P = .0004$) as well as in the wide-neck aneurysm subset (45.7% versus 20.3%, $P = .003$). Core lab analysis revealed no significant stent migration at 1 year. Because angiographic follow-up was not mandated as part of the trial beyond 12 ± 3 months, 2-year angiographic data were not collected.

Multivariate Analysis of Stroke and TAR

The higher ischemic stroke rate in patients receiving stents was attributable to a higher proportion of stent-coiling patients having a baseline history of cerebrovascular accident and a higher proportion of stent-coiling patients having aneurysms with wide necks (Table 8) both at 1 year and 2 years after treatment. TAR, which consisted almost entirely of aneurysm retreatments, was predicted by baseline aneurysm morphologic characteristics, including dome ≥10 mm and neck ≥4 mm, at both 1 year and 2 years of follow-up (Table 9). Stent use was not an independent predictor of TAR.

DISCUSSION

Self-expanding stents have greatly broadened the range of aneurysm morphologies amenable to endovascular treatment. Although the MAPS Trial was designed to evaluate polymer modified coils versus platinum bare metal coils, it allows us to analyze high-quality prospective data on patient outcomes after stent-coiling.

There was a high technical success rate for both stent-coiling and coiling. Stent-coiling was used more frequently in aneurysms with morphologies that typically limit the use of coils because of the risk of parent artery coil prolapse, including low dome-to-neck ratios and wide aneurysm necks. Aneurysm neck ≥4 mm has been associated with a higher risk of aneurysm recanalization[18,19]; in the MAPS Trial, all 12 stent-coiled aneurysms and 10 of 19 coiled aneurysms retreated within 1 year had necks ≥4 mm (Table 6). Conversely, no stent-coiled aneurysms with a <4-mm neck (and only 4% of coiled aneurysms with <4-mm neck) were retreated within 1 year. Wide-neck stent-coiled aneurysms also had higher packing densities, perhaps because the stent lessens concern for parent artery coil prolapse.[20] This higher packing attenuation may explain why wide-neck stent-coiled aneurysms had superior aneurysm occlusion (Raymond Scale scores) at 1 year as compared with wide-neck coiled aneurysms.

Although stent-coiled aneurysms had worse Raymond occlusion scores immediately after treatment than did coiled aneurysms in both the total unruptured intracranial aneurysm cohort and the wide-neck aneurysm subset, they also had more improvement in angiographic appearance at follow-up. There are 4 possible explanations. First, starting with worse initial angiograms will bias follow-up readings on the better-same-worse scale toward more improvement. Second, procedural dual-antiplatelet use is significantly more frequent in stent-coiled patients; therefore it is possible that more interstitial filling is present immediately after treatment in stent-coiled patients as compared with coiled patients. Third, stents were used in more morphologically challenging aneurysms than in the coiled group, and it may not have been possible to treat these aneurysms without a stent. For example, in a very broad-neck, low-domed, shallow aneurysm, it might be difficult to herniate coils into the proximal and distal corners of the aneurysm neck even with a stent in place; such aneurysms could not have been treated with coils alone. Fourth, stents may help to prevent coil compaction within an adjacent aneurysm, perhaps directly by acting as partial parent artery flow diverters, or indirectly by allowing practitioners to confidently pack more coils into stented aneurysms (especially at the neck), or by providing a scaffold for endothelialization across the neck. Because angiographic appearance on follow-up significantly influences aneurysm retreatment decisions and initial core laboratory angiographic score predicts retreatment at 1 year (McDougall et al, *AJNR* in press), it will be important to determine how well immediate posttreatment aneurysm occlusion ultimately predicts aneurysm retreatment over the course of the entire MAPS Trial.

Although periprocedural total significant adverse events were low and similar in stent-coiled and coiled patients, the higher delayed ischemic stroke rate observed in stent-coiled patients is concerning. It is reassuring, however, that multivariate analysis including patients from all enrolling centers confirms that this increased stroke risk is attributable to the presence of more patients in the stent-coiled group having a history of cerebrovascular accident and the presence of more wide-neck aneurysms in the stent-coiled group, as opposed to being caused by stent use per se. Other investigators have reported increased thromboembolic events with the use of intracranial stents in the treatment of aneurysms.[14] Whenever metal is placed in the parent artery, use of antiplatelet agents such as aspirin and/or clopidogrel is prudent to reduce platelet aggregation before the stent becomes endothelialized. Many centers use dual-antiplatelet medications (eg, aspirin and clopidogrel) for either a specified period of time after intracranial stent deployment (eg, 6 weeks or 6 months), until a follow-up angiogram, or indefinitely. Practitioner variability in postprocedure antiplatelet medication may significantly influence delayed ischemic risk to stent patients. This is suggested by poor patient antiplatelet compliance accounting for at least 3 of 12 ischemic strokes in the stent-coiled group and 3 additional ischemic strokes taking place immediately around the time of surgery, aneurysm retreatment, and follow-up angiography, during which antiplatelet medication regimens were uncertain (On-line Table). In addition to medication compliance and lack of consensus on antiplatelet regimen after intracranial stent placement, studies also suggest that inherent biologic resistance to the effects of aspirin and/or clopidogrel may also play a role in delayed ischemic events.[21]

The primary goal of aneurysm treatment is to prevent subarachnoid hemorrhage. Given the very low hemorrhage rate (1 of 361 patients within 2 years), it is too soon to speculate on the overall utility of stent-coiling compared with coiling in protecting patients with unruptured intracranial aneurysms from aneurysm rupture. Not surprisingly, aneurysm dome size ≥10 mm predicted TAR, possibly as the result of the greater opportunity for coil compaction in large aneurysms.

There are several limitations to our study. First, this is a post hoc data analysis from a prospective trial designed to evaluate polymer modified coils versus platinum bare metal coils, not stent-coiling versus coiling. Therefore, stent use was at the operating physician's discretion. Some centers used many stents and some used none. Stent use was significantly higher in North America as opposed to outside North America, suggesting that other geography-specific confounders may be present. Second, some aneurysm morphologies probably could not be treated with coiling alone and could only be treated with stent-coiling, thus biasing stent-coiling cases toward aneurysms with particularly wide necks and low dome-to-neck ratios, known predictors of aneurysm recanalization and procedural complications. Third, postprocedure antiplatelet medication management was not uniform. Because the delayed ischemic stroke rate for stent-coiling may be associated with antiplatelet management, this is a significant limitation. Fourth, the core angiographic laboratory could not score almost 20% of angiograms, most frequently because of image quality, nonmatched comparison views between immediate posttreatment and follow-up, and lack of digital subtraction. Fifth, because the core angiographic laboratory evaluated primarily DSA images, although stent proximal and distal markers could usually be visualized, stent struts could not be directly visualized. Given recent reports of increased delayed thromboembolic complications arising in cases in which stents do not fully appose the wall of the parent artery,[22,23] this also limits our ability to evaluate delayed ischemic risk in stent-coiled patients.

CONCLUSIONS

Stent-coiling had outcomes similar to coiling, despite stented aneurysms having more difficult morphology than did coiled aneurysms. Increased ischemic events in stent-coiled aneurysms were attributable to baseline risk factors and aneurysm morphology, underscoring the overall safety of the stent-coiling technique.

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