### **Supplemental Online Content**

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#### eMethods.

**eFigure 1.** Distribution of Total Cost of Care Among Patients With 24-Month continuous Medicare Fee-for-Service (FFS) Coverage

eFigure 2. Mean Monthly Total Cost of Care Among Patients With 24-Month Continuous Medicare FFS Coverage

**eTable 1.** Proportion of Participants With Missing Medicare Reimbursement Data, Stratified by Completeness of Fee-for-Service (FFS) Coverage

**eTable 2.** Comparison of Baseline Characteristics Between Patients With Complete 24-Month Continuous Medicare FFS and Patients With Incomplete 24-Month Medicare FFS by Intervention Group

**eTable 3.** Total Cost of Care in the 12-Months Preceding and 12-Months During Enrollment Among Patients With 24-Month Continuous Medicare FFS Coverage

**eTable 4.** Heterogeneity of the Effect of the Care Ecosystem on Total Cost of Care From 1-6 Months and 7-12 Months After Trial Enrollment

**eTable 5.** Effect of the Care Ecosystem on Total Cost of Care From 7 to 12 Months After Enrollment With Outliers Trimmed at 97.5th to 99.5th Percentile

**eTable 6.** Effect of the Care Ecosystem on Total Cost of Care From 7 to 12 Months After Enrollment With Winsorization at 97.5th to 99.5th Percentile

**eTable 7.** Effect of the Care Ecosystem on Total Cost of Care From 7 to 12 Months After Enrollment Modeled Among Enrollees With 24-Month Continuous Medicare FFS Coverage Using a Hurdle Model

**eTable 8.** Community Resources and Other Topic Areas That Care Team Navigators Discussed With Caregivers

This supplemental material has been provided by the authors to give readers additional information about their work.

#### eMethods

#### **Intervention description**

The intervention entailed assignment to an unlicensed dementia care guide called a Care Team Navigator (CTN), who received training about dementia and who met with the dyad regularly (typically monthly) by phone or video. CTNs provided personalized information, support, and resources. During monthly calls, CTNs listened and asked questions as guided by standardized protocols to screen for safety hazards, medical issues, and psychosocial risks that required attention from a member of the Care Ecosystem dementia specialist team (nurse, social worker, or pharmacist) or another treating provider. These were addressed during weekly team meetings or individual meetings between the provider and caregiver. CTNs helped caregivers access community services, selected informational resources for them (eTable 8), and guided the caregivers to implement non-pharmacologic strategies to improve problematic behavioral and psychological symptoms of dementia.

#### **Claims files**

Claims files were obtained from the Medicare Inpatient, Outpatient, Carrier, Durable Medical Equipment, Skill Nursing Facility, Home Health Agency, and Hospice files. Part D payments were not included to limit increased variability in total cost of care related to high-priced brand-name specialty drugs, and because the Care Ecosystem intervention was hypothesized to have an effect on low-priced and over-the-counter as opposed to high-priced drugs.<sup>1,2</sup> Care Ecosystem participation was not a service reimbursable by Medicare thus total cost of care did not include the cost of participation among those randomized to receive the Care Ecosystem intervention.

#### Patient level variable details

Rural-Urban Community Area was coded as metropolitan: 1-3, or non-metropolitan, including small town or rural: 4+.<sup>3</sup> Charlson Comorbidity Index was derived from encounter-related ICD-10 codes associated with any encounter during the 12 month baseline period.<sup>4,5</sup> Medicare program enrollment included a monthly indicator of whether the patient had coverage under a Medicare fee-for-service (FFS) or Medicare Advantage (MA) plan.

#### References

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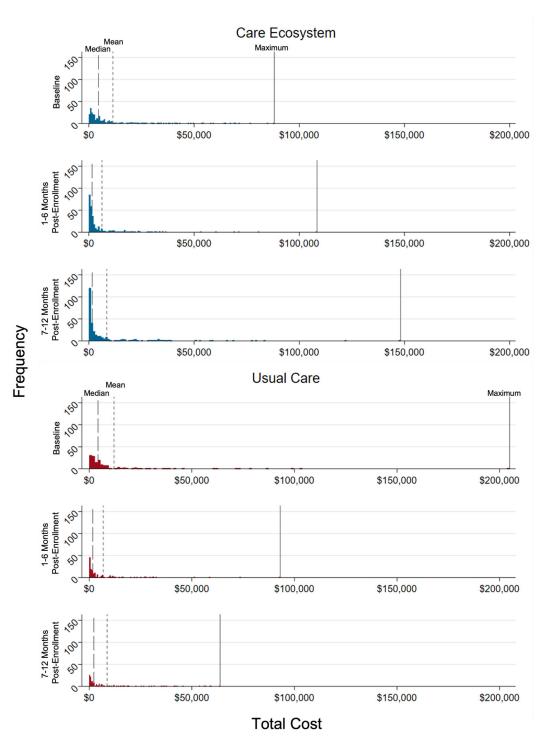
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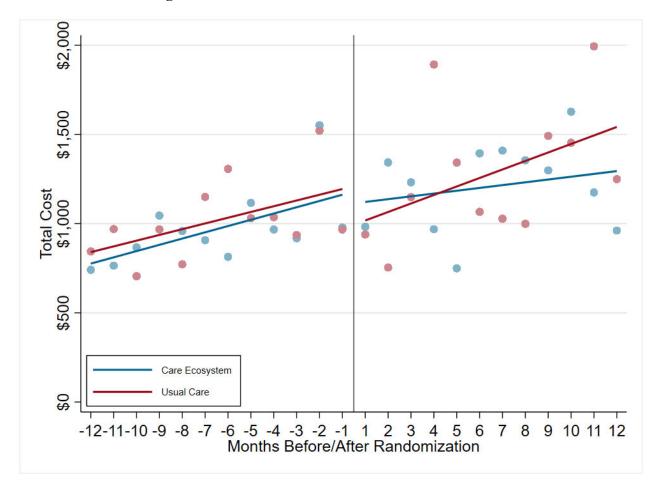
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# eFigure 1. Distribution of total cost of care among patients with 24-month continuous Medicare Fee-For-Service (FFS) coverage

eFigure 1 Legend: Total cost of care is presented separately for the intervention (blue) and usual care (red) arms by the three time periods: baseline (12 months preceding enrollment), 1-6 months post-enrollment, and 7-12 months post-enrollment. Dashed lines indicate median total cost of care, dotted lines indicate mean, and solid lines indicate maximum during the specified periods.



eFigure 2. Mean monthly total cost of care among patients with 24-month continuous Medicare FFS coverage

eFigure 2 Legend. Dots represent mean monthly total cost of care in the intervention (blue) and usual care (red) arm. Lines are fitted for each trial arm for the 12 months preceding and 12 months during trial enrollment.

eTable 1. Proportion of participants with missing Medicare reimbursement data, stratified by completeness of Fee-for-service (FFS) coverage

	Complete FFS	Incomplete FFS		
		Any MA	Other <sup>a</sup>	
	N = 460	N = 183	N = 140	
Missing cost data in 12 months preceding trial enrollment	2 (0.4%)	115 (62.8%)	48 (34.3%)	
Missing cost data in the 12 months during trial enrollment	1 (0.2%)	102 (55.7%)	47 (33.6%)	

<sup>a</sup> Other reasons for incomplete FFS coverage include death or lapse in FFS coverage during 24 month period or linkage failure due to non-Medicare insurance coverage (Medicaid, commercial, VA, etc.), early death, or failure of probabilistic linkage (N = 44).

<sup>b</sup> There were 3 participants with complete FFS coverage who did not have any claims during the year before enrollment (2) and the year after enrollment (1). These costs were set to 0 for the main analysis.

		Care Ecosystem			Usual Care			
		Complete FFS	Incomplete FFS	p-value <sup>a</sup>	Complete FFS	Incomplete FFS	p-value <sup>a</sup>	
		303	209		157	111		
Patient characteristics								
Any Medicare Advantage	;	0 (0)	120 (57.4)		0 (0)	63 (56.8)		
Age, median (IQR)		78.0 (73.0, 84.0)	78.0 (71.0, 85.0)	0.59	78.0 (72.0, 84.0)	78.0 (66.0, 85.0)	0.64	
Female sex, N (%)		171 (56.4%)	128 (61.2%)	0.28	85 (54.1%)	55 (49.5%)	0.46	
Race, N (%)				0.02			0.78	
	Asian	19 (6.6%)	15 (8.0%)		8 (5.3%)	7 (7.1%)		
	Black	8 (2.8%)	15 (8.0%)		6 (4.0%)	4 (4.0%)		
	White	259 (89.9%)	154 (82.4%)		135 (90.0%)	88 (88.9%)		
	Other	2 (0.7%)	3 (1.6%)		1 (0.7%)	0 (0.0%)		
Hispanic, N (%)		21 (6.9%)	33 (16.0%)	0.001	10 (6.4%)	19 (17.1%)	0.005	
Charlson comorbidity ind	ex, N							
(%)				0.97			0.50	
	0	47 (15.6%)	16 (16.5%)		22 (14.2%)	10 (21.3%)		
	1	111 (36.9%)	36 (37.1%)		55 (35.5%)	16 (34.0%)		
	<u>&gt;</u> 2	143 (47.5%)	45 (46.4%)		78 (50.3%)	21 (44.7%)		
Patient dementia severity								
<ul> <li>– QDRS baseline, median (IQR)</li> </ul>	n	10.5 (6.0, 15.5)	12.5 (8.0, 18.0)	< 0.001	10.5 (7.0, 15.6)	11.5 (8.0, 19.0)	0.16	
(IQIX)		10.5 (0.0, 15.5)	12.5 (8.0, 18.0)	<0.001	10.5 (7.0, 15.0)	11.5 (8.0, 19.0)	0.10	
ADI, median (IQR)		3.0 (1.0, 6.0)	3.0 (2.0, 5.0)	0.53	3.0 (1.0, 5.0)	3.0 (1.0, 5.0)	0.91	
Metropolitan, N (%)		257 (84.8%)	193 (92.3%)	0.01	129 (82.2%)	100 (90.1%)	0.07	
Caregiver characteristic	s	、 /	、 <i>/</i>		, <i>,</i> ,	、 / /		
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eTable 2. Comparison of baseline characteristics between patients with complete 24-month continuous Medicare FFS and patients with incomplete 24-month Medicare FFS by intervention group

Caregiver age, median (IQR)	68.0 (59.0, 75.0)	64.0 (55.0, 71.0)	< 0.001	65.0 (56.5, 73.0)	64.5 (56.5, 71.0)	0.87
Caregiver female sex, N (%) Caregiver relationship, N (%)	204 (67.3%)	153 (73.2%)	0.15 0.003	110 (70.1%)	86 (77.5%)	0.18 0.96
Spouse	183 (60.4%)	98 (46.9%)	0.000	88 (56.1%)	64 (57.7%)	0.50
Child	100 (33.0%)	83 (39.7%)		58 (36.9%)	39 (35.1%)	
Other	20 (6.6%)	28 (13.4%)		11 (7.0%)	8 (7.2%)	
Caregiver depression, N (%)			0.45			0.54
None	184 (60.7%)	120 (57.4%)		95 (60.5%)	63 (56.8%)	
Mild to severe	119 (39.3%)	89 (42.6%)		62 (39.5%)	48 (43.2%)	
High caregiver burden, N (%)	167 (55.1%)	120 (57.4%)	0.61	85 (54.1%)	71 (64.0%)	0.11
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<sup>a</sup>For categorical data, complete vs. incomplete FFS groups are compared using Pearson's chi-squared. Wilcoxon rank-sum is used where median and IQR is reported. Comparison of race is white vs. not white.

		ns preceding ollment	12-months during enrollment		
	<b>Care Eco</b>	<b>Usual Care</b>	<b>Care Eco</b>	<b>Usual Care</b>	
Ν	303	157	303	157	
Total cost of care, \$					
Median	4,579	4,284	4,507	5,764	
Mean	11,401	12,038	14,721	15,528	
25th %ile	1,660	1,847	1,622	1,696	
75th %ile	12,739	8,477	16,928	17,599	
Maximum	88,142	204,899	185,913	94,695	
Adjusted median <sup>a</sup>	4,540	3,542	4,362	4,223	
Adjusted mean <sup>a</sup>	10,785	6,390	12,254	9,781	
Adjusted maximum <sup>a</sup>	69,251	42,298	86,329	68,269	

eTable 3. Total cost of care in the 12-months preceding and 12-months during enrollment among patients with 24-month continuous Medicare FFS coverage

<sup>a</sup> Adjusted median, mean, and maximum of total cost of care reflects removal of individuals with DFBETA value above  $2/\sqrt{N}$ .

## eTable 4. Heterogeneity of the effect of the Care Ecosystem on total cost of care from 1-6 months and 7-12 months after trial enrollment

		Trial	Period 1-6 Mont	hs	Trial Period 7-12 Months		ths	Overall	
								Mean monthly cost reductio	
Subgroup	N	Effect <sup>a</sup>	95% CI	<b>p</b> <sup>b</sup>	Effect	95% CI	p°	n <sup>d</sup>	p <sup>e</sup>
Enrollees with 24- month FFS coverage	428	-3290.16	-6149.41, -430.91	0.02	-3026.71	-5899.07, -154.34	0.04	\$526	NA
Enrollees with 24- month FFS coverage or FFS coverage until time of death	484	-2806.92	-5613.35, -0.49	0.05	-3302.66	-6156.80, -448.51	0.02	\$509	NA
All enrolled Medicare beneficiaries <sup>f</sup>	589	-2536.71	-5013.91, -59.50	0.05	-3016.63	-5495.35, -537.90	0.02	\$463	NA
Patient dementia severity									
None to mild	257	-2673.35	-6440.52, 1093.82	0.64	-1708.02	-5484.33, 2068.30	0.20	\$365	0.50
Moderate to severe	171	-4063.24	-8496.85, 370.37	0.64	-4773.70	-9218.08, -329.33	0.30	\$736	0.59
ED Encounters during baseline <sup>g</sup>									
0	244	-1806.36	-5437.20, 1824.48	0.30	-527.86	-4246.86, 3191.14	0.07	\$195	0.19
1 or more	184	-4777.15	-9064.73, -489.57	0.30	-5944.31	-10335.99, -1552.63	0.07	\$893	0.18
Caregiver Burden									
Not high	193	-3275.21	-7539.31, 988.90	0.99	-2763.95	-7045.95, 1518.05	0.87	\$503	0.98
High	235	-3304.16	-7172.34, 564.02	0.99	-3246.11	-7130.52, 638.30	0.07	\$546	0.96
Caregiver Depression									
None	258	-2684.32	-6393.38, 1024.74	0.02	-658.32	-4386.23, 3069.58	0.05	\$279	0.12
Mild to severe	170	-4164.05	-8645.02, 316.92	0.62	-6555.72	-11059.46, -2051.98	0.05	\$893	0.12

 $^{\rm a}$  Interaction analyses were performed after excluding patients who exceeded the DFBETA cutoff value of 0.09

<sup>b</sup>p-value specifies the interaction effect of the characteristic (e.g. dementia severity) at 0-6 months after the patient was enrolled in the Care Ecosystem trial

<sup>c</sup> p-value specifies the interaction effect of the characteristic (e.g. dementia severity) at 7-12 months after the patient was enrolled in the Care Ecosystem trial

<sup>d</sup>Mean reduction in total costs on a monthly basis calculated by adding the effect of the intervention at 1-6 Months and 7-12 Months then dividing by 12

<sup>e</sup>p-value specifies the effect of the three-way interaction (e.g. dementia severity across all time periods)

<sup>f</sup>Participants in the Care Ecosystem trial with Medicare FFS or Medicare Advantage coverage who have non-missing costs in the 12 months preceding and 12 months during trial enrollment <sup>f</sup>Number of ED encounters in the 12 months preceding trial enrollment

eTable 5. Effect of the Care Ecosystem on total cost of care from 7 to 12 months after enrollment with outliers trimmed at 97.5th to 99.5th percentile

Trimming threshold	Ν	Coefficient	Lower CI	Upper CI	<b>P-value</b>
>97.5 percentile	428	-3620.63	-6349.24	-892.02	0.009
>98.0 percentile	434	-3907.13	-6761.55	-1052.71	0.007
>98.5 percentile	443	-3752.55	-6814.04	-691.05	0.016
>99.0 percentile	448	-3289.62	-6526.10	-53.14	0.046
>99.5 percentile	454	-2408.66	-5902.89	1085.57	0.177

eTable 6. Effect of the Care Ecosystem on total cost of care from 7 to 12 months after enrollment with winsorization at 97.5th to 99.5th percentile

Winsorization threshold	Ν	Coefficient	Lower CI	Upper CI	<b>P-value</b>
>97.5 percentile	460	-3422.15	-6153.57	-690.72	0.014
>98.0 percentile	460	-3724.22	-6608.79	-839.65	0.011
>98.5 percentile	460	-3617.34	-6716.80	-517.89	0.022
>99.0 percentile	460	-3222.27	-6477.24	32.71	0.052
>99.5 percentile	460	-2399.62	-5887.64	1088.40	0.178

eTable 7. Effect of the Care Ecosystem on total cost of care from 7 to 12 months after enrollment modeled among enrollees with 24-month continuous Medicare FFS coverage using a hurdle model<sup>a</sup>

	Estimate	Lower CI	Upper CI	P-value
Stage 1	Odds Ratio = 0.29	0.01	5.54	0.40
Stage 2	Coefficient = -0.25	-0.62	0.12	0.18
	Marginal predicted mean 12-month effect (\$) = - 1378.28	-4565.24	1808.67	0.40

<sup>a</sup> Performed as a mixed effects logistic regression model of any cost versus no cost followed by removing observations with zero costs (N = 27) and performing mixed effect linear regression model with log link function and gamma distribution to model total cost of care among enrollees with 24-month continuous Medicare FFS coverage, including outliers with a DFBETA value >  $2/\sqrt{N}$ 

# eTable 8. Community resources and other topic areas that Care Team Navigators discussed with caregivers

Community Resources
Support groups
Counseling
Educational workshops
Public benefits
Respite grants
Legal services
Home safety services
Exercise
Activity or day programs
In-home services
Residential care services
Other Topic Areas
Communication
Disease Information
Behavior Symptoms & Delirium
Caregiver Well-being
Safety (driving, falls, wandering, weight loss, choking)
Medications
Daily Care (activities, nutrition, bathing/grooming, incontinence)
Advance Care Planning
End of Life Care
Respite Care Options
Caregiving Supplies & Technology
Care Transitions (hospitalization, placement)