

Implementation of MIND at Home Program for People Living with Dementia in Primary Care

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Study Objective: Evaluate the feasibility of implementing MIND at Home dementia care coordination into primary care in two large clinics in Iowa and North Carolina.

Background

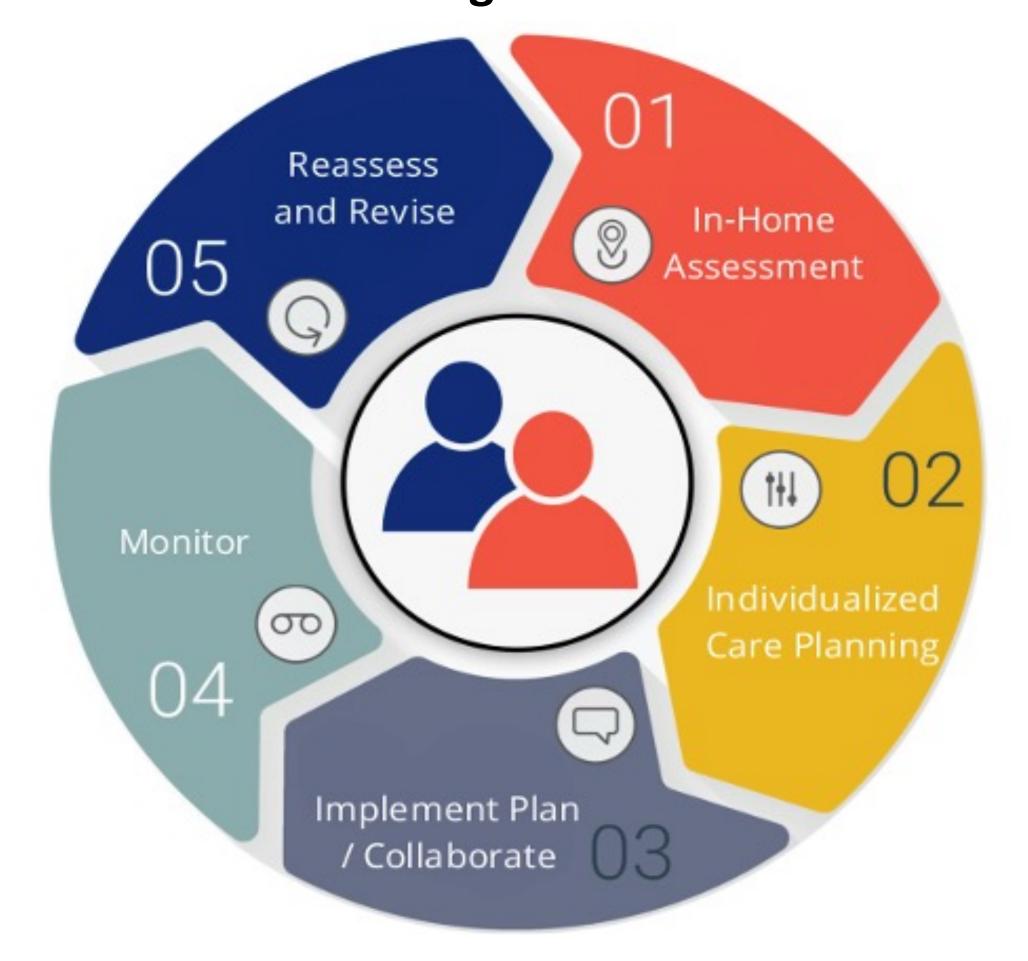
- The prevalence of people living with dementia (PLWD) is growing and dementia is one of the highest burden, and highest cost chronic conditions in the US.¹⁻⁵
- Care management support within primary care practices is one solution to provide patient- and care partner (CP)-centered care, improve patient care outcomes, and reduce high healthcare costs.⁶
- The Maximizing Independence at Home (MIND at Home) dementia care coordination program is a comprehensive evidence-based approach that combines the benefits of clinic-based health care with home-based supportive services for PLWD, families, and care partners.

Methods

- 100 primary care patients and their CPs recruited for a 3-month intervention period
- Program delivery included:
- Specialized initial and ongoing training in dementia care for primary care team
- Comprehensive in-home needs assessment for PLWD and CP assessing 13 domains of needs
- Development and implementation of individualized care plan for PLWD and CP
- One home visit per month (goal)
- Program-specific educational resources and program tools/assessments
- Interdisciplinary dementia specialist expert case consultation to primary care providers
- Process outcomes included: (1) patients enrolled/declined; (2) PLWD and CP needs identified, met.
- Clinical outcomes: (1) hospital transfers; (2) ED visits; and (3) polypharmacy and appropriate medication use.

Intervention

Figure 1. MIND at Home Program Process



Case Examples

- <u>Fall-risk management/home safety</u>: Rugs were removed, wandering was addressed, safety pendants were recommended.
- Aging-in-preferred-setting: PLWD was able to remain in her home longer because of CP support provided.
- Challenging behaviors: CP was coached on behavior management problem solving techniques to reduce agitation
- Referrals to social and community services: CP was connected to several social services because of MIND.
- Caregiver Education and coaching: Provided CP with local dementia support group that meets monthly.
- <u>Transition care:</u> Provided education, coaching, and referrals for long term care services and supports following a hospitalization

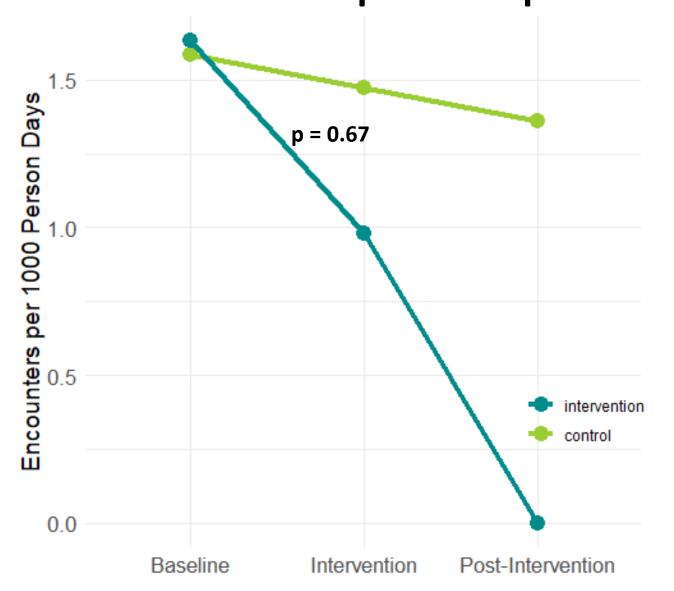
Preliminary Results

Table 1. Characteristics of study and control PLWD

	Intervention (n=34)	Control (n=98)	p-value
Age, mean (range)	81 (64–92)	80 (60-95)	0.68
Male	18 (53%)	50 (51%)	1
Non-Hispanic White	33 (97%)	96 (98%)	1
RUCA, mean (range)	3.56 (9)	3.51 (9)	0.94
Enrolled in Medicare	19 (56%)	56 (57%)	1
CCI, [†] mean score (range)	1.71 (0-9)	1.56 (0-9)	0.73
Dementia-related Rx [‡]	26 (77%)	70 (71%)	0.73
Dementia diagnosis	31 (91%)	64 (65%)	0.01*
Had MMSE [§] (median score)	61% (24)		
Had SLUMS [¶] (median score)	35% (12)		

*Denotes statistical significance; †Charlson Comorbidity Index; †Dementia-related Rx includes memantine and acetylcholinesterase (AChE) inhibitors; §Mini Mental State Examination; ¶Saint Louis University Mental Status Examination for Detecting Mild Cognitive Impairment

Figure 2. Number of hospital transfers/ED visits among intervention and control PLWD per 1000 person days



Although the intervention group saw a steeper decrease in non-ambulatory encounters compared to the control group, the difference over time between the two groups was not significantly different (p = 0.67)

Table 2. PLWD and Care Partners' Identified Needs and Needs Met

Need Type	Need Identified n	Need Met n (% needs met)
PLWD Needs	330	317 (96%)
Home and Personal Safety	106	105 (99%)
General Health Care	70	69 (99%)
Daily and Meaningful Activities	55	48 (87%)
Behavioral Symptoms	40	40 (100%)
Legal and Advanced Care Planning	36	35 (97%)
Cognitive Symptoms	13	13 (100%)
Care Financing	10	7 (70%)
Care Partner Needs	135	114 (84%)
Education	96	80 (83%)
Informal and Emotional Support	21	19 (90%)
Daily Living	7	7 (100%)
Decision Making and Legal Documents	6	5 (83%)
Mental Health	3	1 (33%)
Health	2	2 (100%)

Patient/Caregiver Satisfaction

- Patients/care partners liked education and access to a compassionate care coordinator.
- Patients/care partners wished they could receive continued support after the program or in some other way.
- 75% of patients/caregivers rated that they trusted their care coordinator completely.

Lessons Learned

- Underdiagnosis of dementia, especially among rural and lower SES/educ.
- Recruitment was challenging. Trust was a key factor; easier to recruit when patient was part of another established program or when providers referred directly. Caregivers were "in survival mode" post COVID.
- Expansion of enrollment criteria to include dementia-related Rx was necessary due to hesitancy of primary care providers to dx dementia.
- Home visits provided extensive information on daily living (compared to clinic visits alone) and were valuable to comprehensive care planning.
- Interdisciplinary team-based case discussions were viewed as important, and attendance was high throughout pilot.
- Capturing time at home using EMR records is difficult; ascertainment of other outcomes was feasible.

Discussion and Conclusions

- MIND at Home was successfully implemented in the primary care setting and key outcomes could be ascertained through EHRs.
- Pragmatic modifications including optimization of eligibility indicators; streamlining of program assessment and documentation within electronic systems and waiving in-home visit requirement when needed were necessary to meet the local needs and context of each health system.
- An embedded pragmatic clinic trial (ePCT) is needed to fully evaluate the impact of MIND at Home in the clinical setting on targeted outcomes.

Acknowledgements

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