

Prevalence of Healthcare Conditions and Services Used by Patients with Myotonic Dystrophy (DM) Pre- and Post-Diagnosis: a Real-World Data Analysis



John Day, MD, PhD¹; Kathryn A. Munoz, PhD, MPH²; Richard A. Brook, MS, MBA³; Bradley McEvoy, DrPH²; Kelly DiTrapani, BA, BSN²; Nathan L. Kleinman, PhD³; Chao-Yin Chen, PhD²; Mark C. Stahl, MD, PhD²; Li-Jung Tai, MD, PhD²

¹Stanford University Medical Center; ²Avidity Biosciences, Inc.; ³Better Health Worldwide. All authors have met authorship criteria.

Objectives

Describe the changes in outcomes (healthcare conditions, services, costs, and care days) for patients with DM compared with matched controls (MCs) 2 years post-diagnosis versus 2 years pre-diagnosis

Background

- DM is a rare, dominantly inherited, monogenic, multisystem disease that causes myotonia, progressive muscle weakness, and atrophy, along with respiratory, gastrointestinal, cardiac, and central nervous systems dysfunction, which significantly impacts quality of life¹⁻³
- There are two major types of DM (type 1 and type 2)¹
- Patients experience significant physical limitations, pain, fatigue, and a negative impact on wellbeing^{4,5}
- Currently there is no cure or targeted treatment for DM³
- Real-world data characterizing the patients' pre- and post-diagnosis changes are limited

Methods

- Retrospective database analysis to compare outcomes for patients with DM versus MCs
 - Database: IQVIA US PharMetrics[®] Plus
 - Timeframe: January 2010 through March 2021
- The DM cohort is identified as having ≥2 DM claims ≥30 days apart
 - Claims identified by International Classification of Disease Ninth Revision (ICD-9) code 359.21 or Tenth Revision (ICD-10) code G71.11, which does not differentiate between DM subtypes
- The first diagnosis date was used for the index date
- DM patients were matched to a 5% random sample of eligible non-DM controls
 - Matching was done using R's Matchit procedure, with nearest neighbor matching (exact matching on month of index date)
- Cohorts were matched (5-MC:1-DM) on index month and baseline age, region, gender, plan, and payer types
- All subjects (patients and MCs) were required to have a minimum of 48 months of continuous data:
 - 24 months prior to their index date
 - 24 months following ("post") their index date (includes the index date)
 - The index date was the beginning of the post-index evaluation period
- Changes in outcomes were measured as 2 years post-diagnosis minus 2 years pre-diagnosis using:
 - Location of care data for overall care
 - 283 US Agency for Healthcare Research and Quality (AHRQ) condition categories
- Post-pre changes were compared within cohorts (using McNemar tests) and between cohorts (using t-tests)
- All presented comparisons were significant ($p < 0.05$) unless noted
- Based on the 283 comparisons, those p -values < 0.000177 (0.05/283) are considered highly significant

Results

We identified 519 DM patients and 2,595 MCs

- Descriptive characteristics were similar between cohorts (Table 1)
- The cohorts had significant ($p < 0.0001$) differences for the Charlson Comorbidity Index (Table 2)
- The cohorts had changes (Post-Pre) in costs and days of service by location of care (Table 3)
- DM patients':
 - AHRQ prevalence changed significantly in 58 categories (↑57 ↓1, Figure 1)
 - Costs changed significantly in 7 AHRQ categories (↑6 ↓1, Figure 2)
 - Number of services per person per year changed significantly in 21 AHRQ categories (↑20 ↓1, Figure 3)

Table 1: Age, US Region, Insurance, and Payer Types Were Similar Between DM Patients and MCs

Descriptive Characteristics	DM Patients (N=519)
Gender, % Female	47.6%
Age, mean years (SD)	43.8 (18.0)
Age, years	
<18	11.0%
≥18 to <35	17.3%
≥35 to <45	18.9%
≥45 to <55	21.8%
≥55 to <65	22.2%
≥65	8.9%
US Region	
South	30.1%
Midwest	26.8%
Northeast	22.7%
West	18.1%
Unknown	2.3%
Insurance Type	
Preferred Provider Organization	69.9%
Health Maintenance Organization	20.8%
Point-of-Service Plan	4.0%
Consumer-Directed Health Care	2.1%
Indemnity/Traditional Plan	1.3%
Unknown Plan	1.7%
Payer Type	
Commercial	60.9%
Self-Insured	22.7%
Medicaid	6.2%
Medicare Advantage	4.2%
Medicare Supplemental	4.6%
Unknown Line of Business	1.3%

There were no significant differences between DM patients and MCs $p > 0.05$

Table 2: Before and After Diagnosis, Charlson Comorbidity Index Scores Were Higher for DM Patients Versus MCs

Descriptive Characteristics	DM Patients (N=519)	Matched Non-DM Patients (N=2,595)
Pre-index (before diagnosis)		
Mean (SD) score	1.08 (1.83)	0.57 (1.30)
Percent with values >1, %*	24.3%	13.0%
Post-index (after diagnosis)		
Mean (SD) score	1.58 (2.27)	0.71 (1.63)
Percent with values >1, %*	33.5%	14.8%

*All comparisons highly significant $p < 0.0001$

Abbreviations:

AHRQ, US Agency for Healthcare Research and Quality; DM, myotonic dystrophy; MC, matched control; PMPY, per member per year; SD, standard deviation.

References:

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Conclusions

- Healthcare utilization increased significantly in DM patients following diagnosis and was higher both overall and in different AHRQ categories than in MCs
 - This likely reflects the need to investigate and manage previously unsuspected manifestations of DM following formal diagnosis
- Future research should confirm if these findings hold true in longer-term follow-up
- These data highlight the many unmet needs for DM patients, including higher costs, more days of care, more prevalent and costly comorbidity management and the need for therapeutic interventions
- Based on the high unmet need, Avidity Biosciences is investigating AOC 1001 for the potential treatment of myotonic dystrophy type 1

Table 3: Costs and Number of Days of Service (Before, After, and Changes [Post minus Pre]) Were Higher for DM Patients for "Emergency Department," "Inpatient" and "All Locations of Care"

Location of Care	Cost (PMPY)			Days of Service (PMPY)		
	DM Patients (SD)	MCs (SD)	Difference	DM Patients (SD)	MCs (SD)	Difference
Post-diagnosis						
Emergency Department	\$558 (\$2,017)	\$208 (\$1,057)	\$350*	0.59 (1.66)	0.22 (0.61)	0.37*
Inpatient	\$8,242 (\$36,117)	\$1,482 (\$8,460)	\$6,760*	2.65 (10.56)	0.41 (2.64)	2.24*
All Medical Locations of Care [^]	\$21,130 (\$56,447)	\$5,182 (\$14,272)	\$15,949*	26.03 (27.06)	9.80 (13.59)	16.24*
All Medical and Drug	\$25,594 (\$59,365)	\$6,684 (\$16,709)	\$18,910*	39.80 (35.41)	17.48 (20.30)	22.32*
Pre-diagnosis						
Emergency Department	\$449 (\$1,623)	\$211 (\$1,242)	\$238*	0.42 (1.25)	0.22 (0.76)	0.20*
Inpatient	\$6,965 (\$39,174)	\$1,023 (\$6,192)	\$5,942*	1.60 (8.65)	0.37 (3.27)	1.23*
All Medical Locations of Care [^]	\$15,534 (\$47,773)	\$4,276 (\$11,264)	\$11,257*	18.40 (23.73)	8.88 (12.21)	9.52*
All Medical and Drug	\$18,705 (\$51,537)	\$5,640 (\$13,277)	\$13,065*	29.87 (30.87)	16.03 (18.85)	13.84*
Change (Post-Pre)						
Emergency Department	\$108 (\$1,856)	-\$4 (\$1,075)	\$112	0.17 (1.39)*	0.00 (0.67)	0.17*
Inpatient	\$1,277 (\$50,185)	\$459 (\$9,331) [§]	\$818	1.05 (8.71)*	0.04 (3.12)	1.01*
All Medical Locations of Care [^]	\$5,597 (\$67,294)	\$905 (\$13,991) [‡]	\$4,691	7.63 (20.88)*	0.92 (11.15)*	6.71*
All Medical and Pharmacy	\$6,889 (\$69,332) [§]	\$1,044 (\$14,926) [‡]	\$5,845	9.93 (22.31)*	1.45 (12.58)*	8.48*

Change=Post- minus Pre-Diagnosis. [^]Emergency Department, [‡]Inpatient, [§]Laboratory, Office, [¶]Other, ^{||}Outpatient, and ^{||}Pharmacy
Between Cohorts: * $p < 0.0001$; [†] $p < 0.01$ Within Cohorts: [‡] $p < 0.0001$; [§] $p < 0.01$; ^{||} $p < 0.05$

Figure 1: DM Patients Had Higher Increases in Prevalence For AHRQ Categories of "Other Nervous System Disorders," "Cardiac Dysrhythmias," and "Other Lower Respiratory Disease" Versus MCs

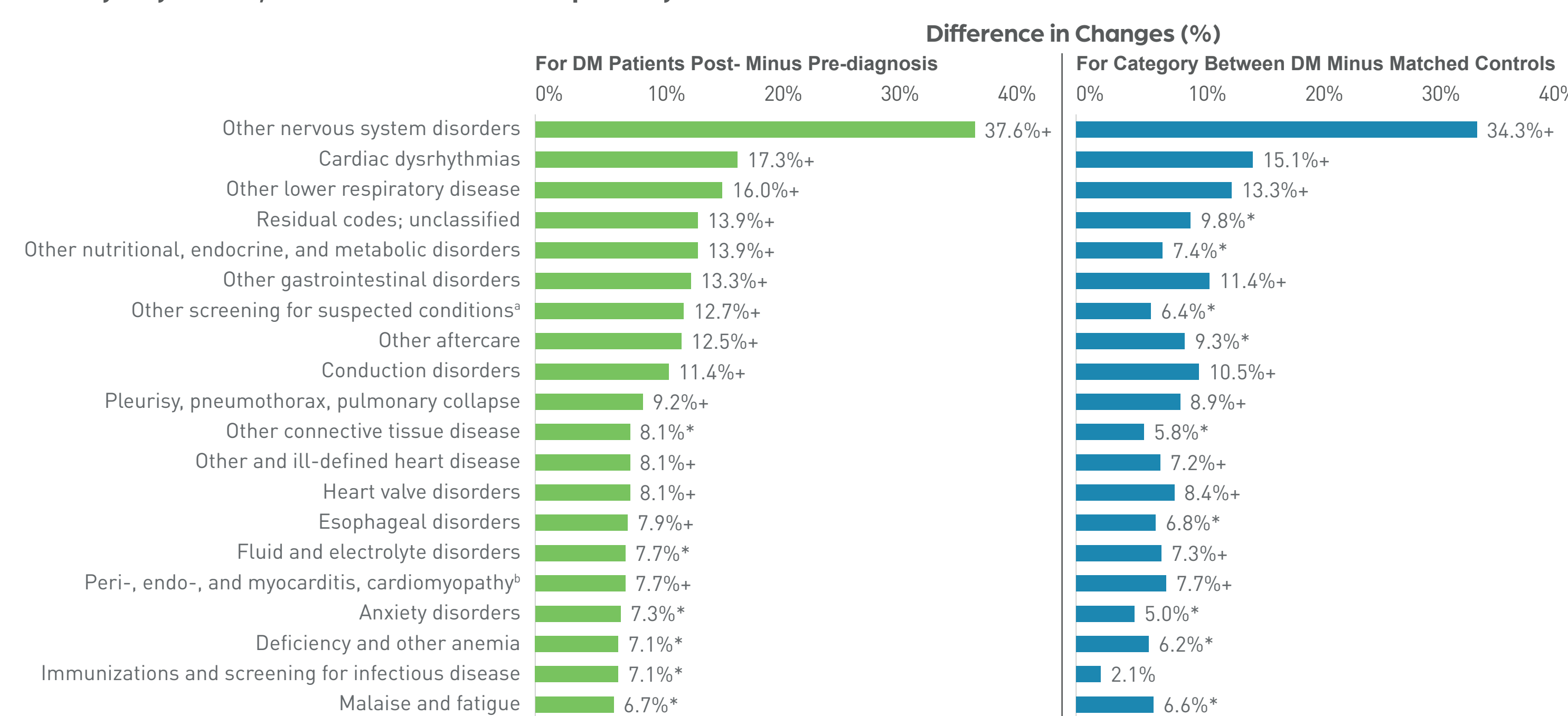


Figure 2: DM Patients Had Higher Increases in Costs in AHRQ Categories of "Other Nervous System Disorders," "Unclassified Codes," "Other Fractures," and "Developmental Disorders" Versus MCs

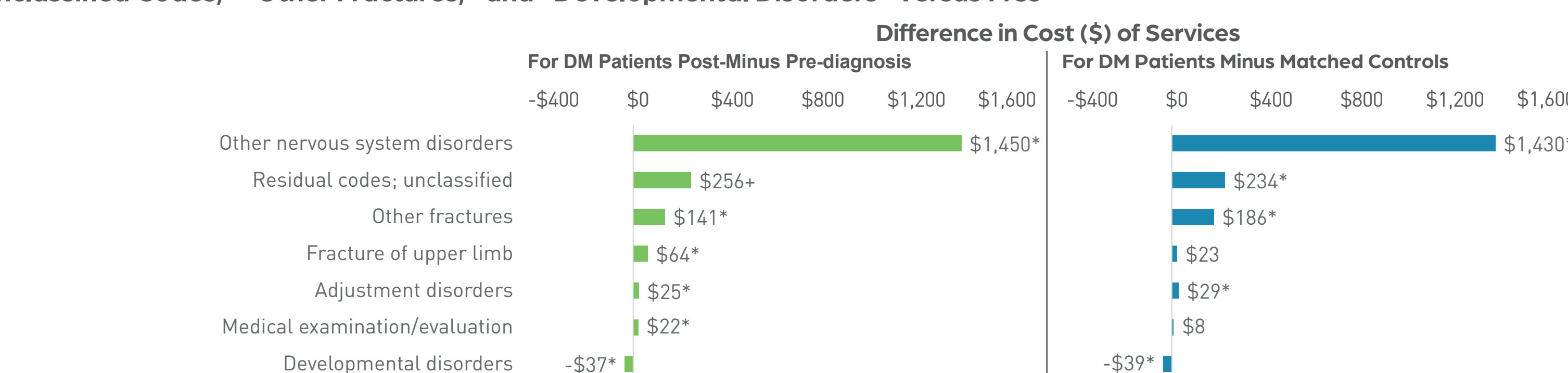
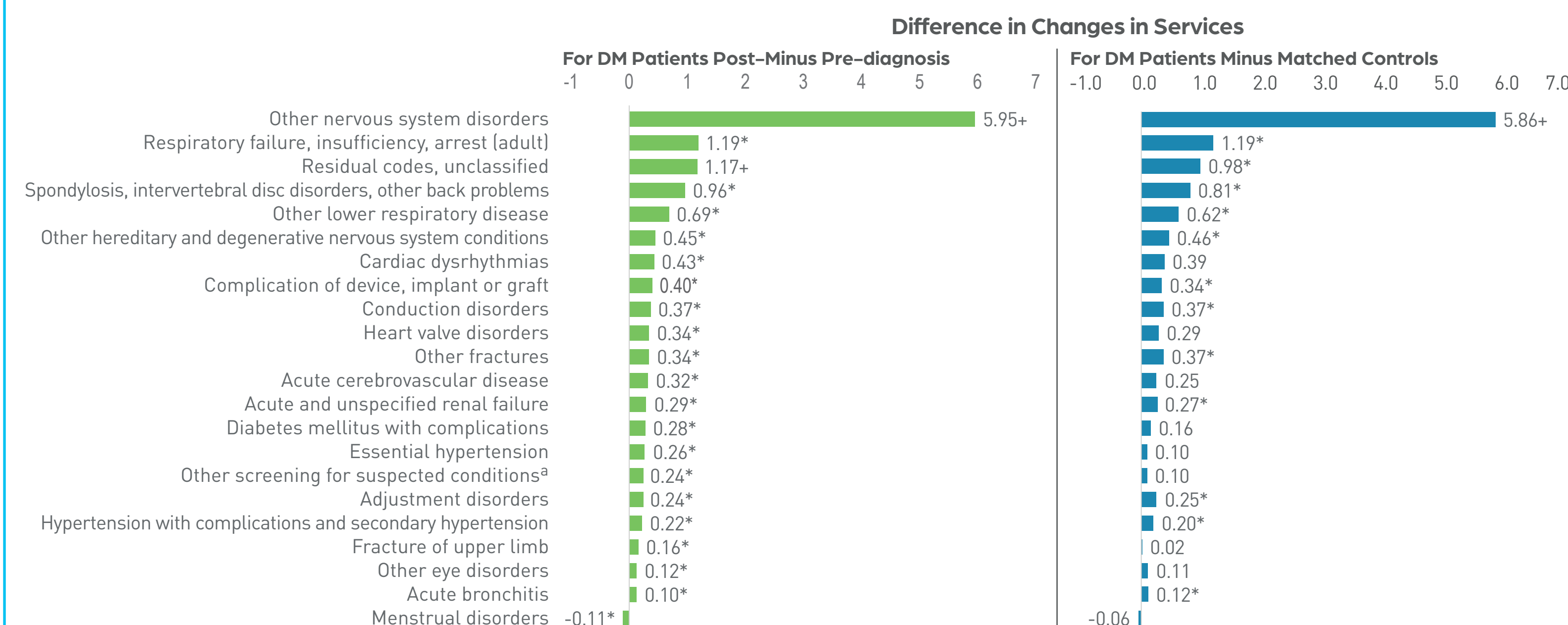


Figure 3: DM Patients Had Higher Increases in "Other Nervous System Disorders" and "Respiratory-Related" Services Versus MCs



* $p < 0.05$; [†] $p < 0.000177$ [threshold for multiple comparisons (0.05/283 comparisons)]

[†]Not mental disorders or infectious disease; [‡]Except that caused by tuberculosis or sexually transmitted disease