

Discharge Medical Complexity, Change in Medical Complexity and Pediatric 30-day Readmission

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BACKGROUND: While medical complexity is associated with pediatric readmission risk, less is known about how increases in medical complexity during hospitalization affect readmission risk.

METHODS: We conducted a five-year retrospective, case-control study of pediatric hospitalizations at a tertiary care children's hospital. Cases with a 30-day unplanned readmission were matched to controls based on admission seasonality and distance from the hospital. Complexity variables included the number of medications prescribed at discharge, medical technology, and the need for home healthcare services. Change in medical complexity variables included new complex chronic conditions and new medical technology. We estimated odds of 30-day unplanned readmission using adjusted conditional logistic regression.

RESULTS: Of 41,422 eligible index hospitalizations, we included 595 case and 595 control hospitalizations. Complexity: Polypharmacy after discharge was common. In adjusted analyses, being discharged with

≥2 medications was associated with higher odds of readmission compared with being discharged without medication; children with ≥5 discharge medications had a greater than four-fold higher odds of readmission. Children assisted by technology had higher odds of readmission compared with children without technology assistance. Change in complexity: New diagnosis of a complex chronic condition (Adjusted Odds Ratio (AOR) = 1.75; 1.11-2.75) and new technology (AOR = 1.84; 1.09-3.10) were associated with higher risk of readmission when adjusting for patient characteristics. However, these associations were not statistically significant when adjusting for length of stay.

CONCLUSION: Polypharmacy and use of technology at discharge pose a substantial readmission risk for children. However, added technology and new complex chronic conditions do not increase risk when accounting for length of stay. *Journal of Hospital Medicine* 2019;14:474-481. © 2019 Society of Hospital Medicine

Hospitalizations are disruptive, stressful, and costly for patients and families.¹⁻⁵ Hospital readmissions subject families to the additional morbidity inherent to hospitalization and place patients at additional risk of hospital-acquired conditions or other harm.⁶⁻⁹ In pediatrics, hospital readmissions are common for specific conditions,¹⁰ with rates varying across institutions,^{10,11} and as many as one-third of unplanned pediatric readmissions are potentially preventable.¹²

Reducing pediatric readmissions requires a deeper understanding of the mechanisms through which readmissions occur. Medical complexity—specifically chronic conditions and use of medical technology—is associated with increased risk of readmission.^{13,14} Polypharmacy at discharge has also been associated with readmission.^{15,16} However, prior studies on polypharmacy and readmission risk examined the count of total medications and did not consider the nuances of scheduled versus as-needed medications, or the frequency of doses. These nuances may be critical to caregivers as discharge medical complexity can be overwhelming, even in diagnoses which are not traditionally considered complex.¹⁷ Finally, of potentially greater importance than medical complexity at discharge is a change in medical complexity during a hospitalization—for example, new diagnoses or new technologies that require additional education in hospital and management at home.

We sought to further understand the relationship between discharge medical complexity and readmission risk with re-

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gards to polypharmacy and home healthcare referrals at discharge. Specifically, we hypothesized that a change in medical complexity during an admission—ie, a new chronic diagnosis or new technology—would be a more prominent risk factor for readmission than discharge complexity alone. We examined these factors in the context of length of stay (LOS) since this is a marker of in-hospital severity of illness and a potentially modifiable function of time allowed for in-hospital teaching and discharge preparation.

METHODS

We conducted a retrospective, case-control study of pediatric hospitalizations at one tertiary care children's hospital. Children <18 years were eligible for inclusion. Normal birth hospitalizations were excluded. We randomly selected one hospitalization from each child as the index visit. We identified cases, hospitalizations at C.S. Mott Children's Hospital between 2008 and 2012 with a subsequent unplanned 30-day readmission,¹⁸ and matched them one to one with hospitalizations at the same hospital during the same period without subsequent readmission. We matched cases to controls based on the month of admission to account for seasonality of certain illnesses. We also matched on distance and direction from the hospital to the patient's home to account for the potential to have readmissions to other institutions. We utilized both distance and direction recognizing that a family living 30 miles in one direction would be closer to an urban area with access to more facilities, as opposed to 30 miles in another direction in a rural area without additional access. We subsequently performed medical record review to abstract relevant covariates.

Primary Predictors

Medical Complexity Models (Models 1 and 2):

We evaluated three attributes of discharge medical complexity abstracted by medical record review—discharge medications, technology assistance (ie, tracheostomy, cerebral spinal fluid ventricular shunt, enteral feeding tube, central line), and the need for home healthcare after discharge. We counted discharge medications based on the number of medications listed on the discharge summary separated into scheduled or as needed.¹⁹ We also considered the number of scheduled doses to be administered in a 24-hour period (see Appendix methods for more information on counting discharge medications). For assistance by technology, we considered the presence of tracheostomy, cerebral spinal fluid ventricular shunt, enteral feeding tube, and central lines. While we describe these technologies separately, for multivariable analyses we considered the presence of any of the four types of technology.

Change in Medical Complexity Models (Models 3 and 4)

We examined two aspects of change in medical complexity—the presence of a new complex chronic condition (CCC)²⁰ diagnosed during the hospitalization, and a new reliance on medical technology. The presence of new CCC was determined by comparing discharge diagnoses to past medical history abstracted by medical record review. A new CCC was

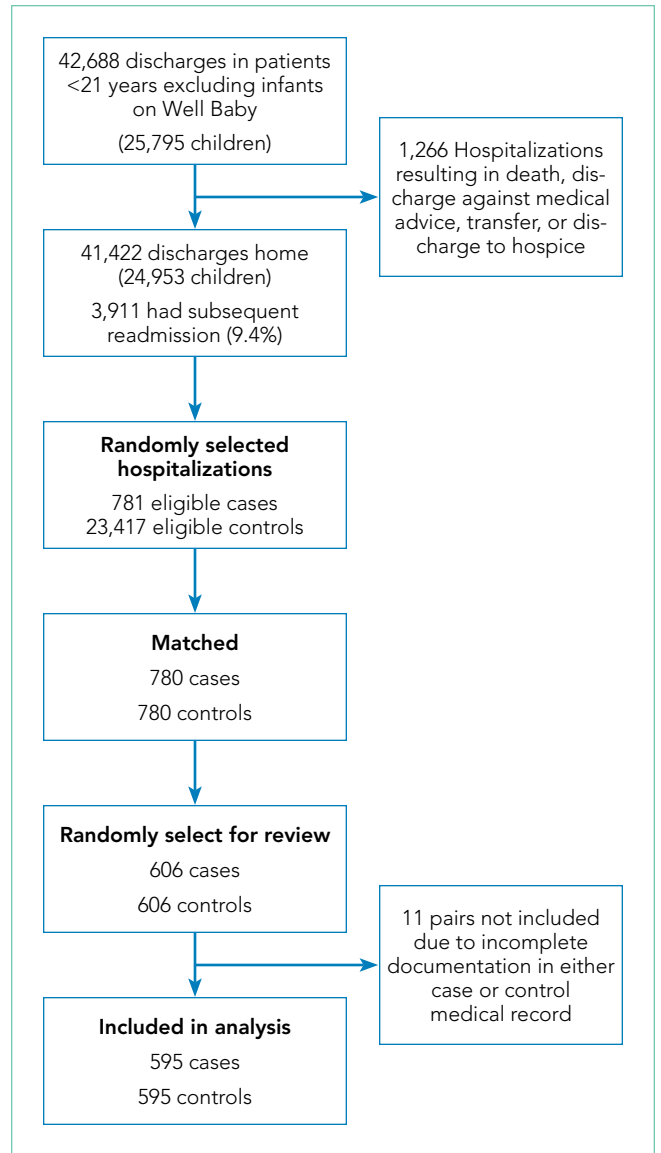


FIG. Cohort Derivation

defined as any complex chronic condition that was captured in the discharge diagnoses but was not evident in the past medical history. By definition, all CCCs coded during birth hospitalization (eg, at discharge from the neonatal intensive care unit) were assigned to "new" CCC. We calculated a kappa statistic to determine interrater reliability in determining the designation of new CCC. A sensitivity analysis examining these birth CCCs was also performed comparing no new CCC, new CCC, and new CCC after birth hospitalization. The methods appendix provides additional information on considering new CCCs. New technology, abstracted from chart review, was defined as technology placed during hospitalization that remained in place at discharge. If a child with existing technology had additional technology placed during the hospitalization (eg, a new tracheostomy in a child with a previously placed enteral feeding tube), the encounter was considered as having new technology placed.

TABLE 1. Patient Characteristics and Frequency of Medical Complexity

| Patient/ Hospitalization Characteristics | | Hospitalizations with Subsequent 30-day Readmission | | Hospitalizations without Subsequent 30-day Readmission | |
|--|--|---|---|--|--|
| | | Cases (n = 595) Number (column %) | Controls (n = 595) Number (column %) | | |
| Age | Newborn birth | 20 (3.4) | 34 (5.7) | | |
| | ≤1 year, non-newborn | 105 (17.7) | 121 (20.3) | | |
| | >1 to ≤5 | 105 (17.7) | 114 (19.2) | | |
| | >5 to ≤10 | 123 (20.7) | 90 (15.1) | | |
| | >10 ≤15 | 108 (18.2) | 118 (19.8) | | |
| | >15 years | 134 (22.5) | 118 (19.8) | | |
| | Race/ethnicity | Non-Hispanic White | 443 (74.5) | 458 (77.0) | |
| Non-Hispanic Black | | 73 (12.3) | 66 (11.1) | | |
| Hispanic or another race | | 62 (10.4) | 46 (7.7) | | |
| Unknown | | 17 (2.9) | 25 (4.2) | | |
| Gender | Female | 286 (48.1) | 276 (46.4) | | |
| Insurance | Private | 347 (58.3) | 386 (64.9) | | |
| | Medicaid | 160 (26.9) | 164 (27.6) | | |
| | Funds for children with medical complexity | 87 (14.6) | 43 (7.2) | | |
| | Self-pay/other including Medicare | 1 (0.2) | 2 (0.3) | | |
| Length of stay | 0-1 days | 96 (16.1) | 208 (35.0) | | |
| | 2-3 days | 192 (32.3) | 198 (33.3) | | |
| | 4-5 days | 86 (14.5) | 82 (13.8) | | |
| | 6-7 days | 58 (9.8) | 24 (4.0) | | |
| | 7-14 days | 82 (13.8) | 40 (6.7) | | |
| | >14 days | 81 (13.6) | 43 (7.2) | | |

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Covariates

We created different sets of multivariable models to account for patient/hospitalization characteristics. In Models 1 and 3, we examined the primary predictors adjusting for patient characteristics (age, race/ethnicity, sex, and insurance). In Models 2 and 4, we added the index hospitalization LOS into the multivariable models adjusting for patient characteristics. We chose to add LOS in a second set of models because it is a potentially important confounder in readmission risk: discharge timing is a modifiable factor dependent on both physiologic recovery and the medical team's perception of caregiver's readiness for discharge. We elected to present models with and without LOS since LOS is also a marker of illness severity while in the hospital and is linked to discharge complexity.

Statistical Analysis

A review of 600 cases and 600 controls yields 89% power to detect statistical significance for covariates with an odds ratio of 1.25 ($\beta = 0.22$) if the candidate covariate has low to moderate correlation with other covariates (<0.3). If a candidate covariate has a moderate correlation with other covariates (0.6), we have 89% power to detect an odds ratio of 1.35 ($\beta = 0.30$).²¹ We calculated odds of 30-days unplanned readmission using conditional logistic regression to account for matched case-control design. All the analyses were performed using STATA 13 (Stata Corp., College Station, Texas).

RESULTS

Of the 41,422 eligible index hospitalizations during the study period, 9.4% resulted in a 30-day unplanned readmission. Af-

TABLE 1. Patient Characteristics and Frequency of Medical Complexity (continued)

| | | | Hospitalizations with Subsequent 30-day Readmission | Hospitalizations without Subsequent 30-day Readmission |
|---------------------------------------|--|--------------|--|---|
| | | | Cases (n = 595) Number (column %) | Controls (n = 595) Number (column %) |
| Medical Complexity at Index Discharge | Number of scheduled medications ^a | 0 | 50 (8.4) | 143 (24.0) |
| | | 1 | 82 (13.8) | 142 (23.9) |
| | | 2 | 84 (14.1) | 99 (16.6) |
| | | 3 | 66 (11.1) | 79 (13.3) |
| | | 4 | 61 (10.3) | 33 (5.6) |
| | | 5+ | 252 (42.4) | 99 (16.6) |
| | Number of as-needed (prn) medications ^b | 0 | 204 (34.3) | 243 (40.8) |
| | | 1 | 177 (29.8) | 169 (28.4) |
| | | 2 | 94 (15.8) | 85 (14.3) |
| | | 3 | 75 (12.6) | 65 (10.9) |
| | | 4 | 29 (4.9) | 21 (3.5) |
| | | 5+ | 16 (2.7) | 12 (2.0) |
| | Number of scheduled doses per 24 hours | Median (IQR) | 6 (3,12) | 3 (0,7) |
| | Medical technology | Any | 249 (41.9) | 85 (14.3) |
| | Specific types of medical technology | Tracheostomy | 19 (3.2) | 8 (1.3) |
| Ventricular shunt | | 29 (4.9) | 11 (1.9) | |
| Surgically placed enteral tube | | 66 (11.1) | 19 (3.2) | |
| Nonsurgically placed enteral tube | | 44 (7.4) | 30 (5.0) | |
| Central line | | 158 (26.6) | 26 (4.4) | |
| Home healthcare after discharge | | 260 (43.7) | 138 (23.2) | |
| Change in Medical State Complexity | Any new complex chronic condition | 105 (17.7) | 60 (10.1) | |
| | Any new technology | 101 (17.0) | 43 (7.2) | |

^aMedian (IQR) of number of scheduled medications: Cases—4 (2,7) Controls—2 (1, 3)

^bMedian (IQR) of number of prn medications: Cases—1 (0, 2) Controls—1 (0, 2)

Abbreviations: IQR, interquartile range; PRN, pro re nata (as needed).

ter randomly selecting one hospitalization per child, there were 781 eligible cases. We subsequently matched all but one eligible case to a control. We randomly selected encounters for medical record review, reviewing a total of 1,212 encounters. After excluding pairs with incomplete records, we included 595 cases and 595 controls in this analysis (Figure). Patient/hospitalization characteristics are displayed in Table 1. The most frequent primary discharge diagnoses are displayed in Appendix Table 1.

Models of Medical Complexity at Discharge

Polypharmacy after discharge was common for both readmitted and nonreadmitted patients. Children who experienced unplanned readmission in 30 days were discharged with a median

of four different scheduled medications (interquartile range [IQR] 2,7) which translated into a median of six (IQR 3,12) scheduled doses in a 24-hour period. In comparison, children without an unplanned readmission had a median of two different scheduled medications (IQR 1,3) with a median of three (IQR 0,7) scheduled doses in a 24-hour period. Medical technology was more common in case children (42%) than in control children (14%). Central lines and enteral tubes were the most common forms of medical technology in both cases and controls. Home health referral was common in both cases (44%) and controls (23%; Table 1).

Many attributes of complexity were associated with an elevated readmission risk in bivariate analysis (Table 2). As the measures of scheduled polypharmacy (the number of sched-

TABLE 2. **Bivariate Logistic Regression Models**

| | | Odds Ratio (95% CI) | |
|---------------------------------------|--|-----------------------------------|-------------------|
| Medical Complexity at Index Discharge | Number of scheduled medications | 0 | REF |
| | | 1 | 1.70 (1.09-2.65) |
| | | 2 | 2.36 (1.50-3.72) |
| | | 3 | 2.62 (1.62-4.24) |
| | | 4 | 5.74 (3.25-10.14) |
| | | 5+ | 8.43 (5.36-13.25) |
| | Number of as needed (prn) medications | 0 | REF |
| | | 1 | 1.25 (0.94-1.64) |
| | | 2 | 1.34 (0.94-1.91) |
| | | 3 | 1.36 (0.93-2.01) |
| | | 4 | 1.67 (0.92-3.04) |
| | | 5+ | 1.62 (0.73-3.59) |
| | Number of scheduled doses per 24 hours | | 1.11 (1.08-1.13) |
| | Medical technology | Any | 4.49 (3.27-6.16) |
| | Specific types of medical technology | Tracheostomy | 2.57 (1.07-6.16) |
| | | Ventricular shunt | 2.80 (1.36-5.76) |
| | | Surgically placed enteral tube | 4.13 (2.35-7.26) |
| | | Nonsurgically placed enteral tube | 1.52 (0.93-2.47) |
| | | Central line | 7.60 (4.77-12.11) |
| | Home healthcare after discharge | | 2.77 (2.10-3.65) |
| Change in Medical Complexity | Any new complex chronic condition | 1.94 (1.37-2.74) | |
| | Any new technology | 2.81 (1.88-4.21) | |

uled medications and number of doses per 24 hours) increased, the odds of readmission also increased in a dose-response manner. Higher numbers of as-needed medications did not increase the odds of readmission. Being assisted with any medical technology was associated with higher odds of readmission. Specifically, the presence of a central line had the highest odds of readmission in unadjusted analysis (odds ratio [OR] 7.60 (95% confidence interval [CI]: 4.77-12.11). In contrast, the presence of a nonsurgically placed enteral feeding tube (eg, nasogastric tube) was not associated with readmission. Finally, in unadjusted analyses, home healthcare need was associated with elevated odds of readmission.

In Model 1 (adjusting only for patient characteristics; Table 3), being discharged on two or more scheduled medications was associated with higher odds of readmission compared to being discharged without medications, with additional medications associated with even higher odds of readmission. Children with any technology had higher odds of readmission than children without medical technology. Likewise, home health-

care visits after discharge were associated with elevated odds of readmission in multivariable analyses without LOS. However, after adding LOS to the model (Model 2), home healthcare visits were no longer significantly associated with readmission.

Change in Medical Complexity Models

The adjudication of new CCCs had good reliability ($K = 0.72$). New CCCs occurred in 18% and new technologies occurred in 17% of cases. Comparatively, new CCCs occurred in 10% and new technologies in 7% of hospitalizations in control children (Table 1). In bivariate analyses, both aspects of change in medical complexity were associated with higher odds of readmission (Table 2). In multivariate analysis with patient characteristics (Model 3; Table 3), all aspects of change in complexity were associated with elevated odds of readmission. A new CCC was associated with higher odds of readmission (adjusted OR (AOR) 1.75, 95% CI: 1.11-2.75) as was new technology during admission (AOR 1.84, 95%CI: 1.09-3.10). Furthermore, the odds of readmission for medical complexity variables

TABLE 3. **Multivariable Logistic Regression models^a**

| | | | Model 1: Complexity Model Adjusted for Patient Demographics Adjusted Odds Ratio (95%CI) | Model 2: Complexity Model Adjusted for Patient Demographics and LOS Adjusted Odds Ratio (95%CI) | Model 3: Change in Complexity Adjusted for Patient Demographics Adjusted Odds Ratio (95%CI) | Model 4: Change in Complexity Adjusted for Patient Demographics and LOS Adjusted Odds Ratio (95%CI) |
|--|---------------------------------------|---|--|--|--|--|
| Medical Complexity at Index Discharge | Number of scheduled medications | 0 | REF | REF | REF | REF |
| | | 1 | 1.39 (0.86-2.22) | 1.23 (0.75-2.01) | 1.34 (0.83-2.15) | 1.20 (0.73-1.98) |
| | | 2 | 2.13 (1.30-3.50) | 1.97 (1.17-3.32) | 2.10 (1.27-3.46) | 1.98 (1.17-3.36) |
| | | 3 | 1.86 (1.09-3.16) | 1.81 (1.03-3.18) | 1.86 (1.09-3.18) | 1.83 (1.04-3.22) |
| | | 4 | 3.78 (2.00-7.14) | 3.58 (1.81-7.09) | 3.61 (1.89-6.89) | 3.33 (1.67-6.66) |
| | | 5+ | 4.99 (2.99-8.35) | 4.63 (2.69-7.96) | 4.88 (2.91-8.18) | 4.54 (2.63-7.84) |
| | | Home healthcare after discharge | 1.48 (1.03-2.12) | 1.14 (0.77-1.70) | 1.53 (1.06-2.21) | 1.20 (0.80-1.80) |
| | Any medical technology | 2.60 (1.78-3.80) | 2.64 (1.78-3.92) | – | – | |
| Change in Medical Complexity | Any new complex chronic condition | | – | – | 1.75 (1.11-2.75) | 1.54 (0.95-2.52) |
| | Technology assistance | None | – | – | REF | REF |
| | | Preexisting technology | – | – | 3.00 (1.87-4.82) | 3.46 (2.11-5.68) |
| | | New technology during admission in children with or without preexisting technology | – | – | 1.84 (1.09-3.10) | 1.60 (0.92-2.80) |

^aAdjusted for patient demographics include age, race/ethnicity, sex, and insurance. Bolded values represent P < .05.

(polypharmacy and home healthcare need) remained largely unchanged when adding the change in medical complexity variables (ie, comparing Model 1 and Model 3). However, when accounting for LOS (Model 4), neither the acquisition of a new CCC nor the addition of new technology was associated with readmission. The most common form of new technology was central line followed by nonsurgically placed enteral tube (Appendix Table 2). Finally, in sensitivity analyses (results not detailed), separating new CCC acquired at birth and new CCCs in nonbirth hospitalizations, compared to hospitalizations with no new CCC, yielded similar results as the primary analyses.

DISCUSSION

Higher numbers of scheduled medications prescribed at discharge pose a progressively greater readmission risk for children. The presence of medical technology at admission is associated with subsequent readmission; however, added technology and home healthcare needs were not, when adjusting for patient characteristics and LOS. Additionally, the acquisition of a new CCC was not associated with readmission, when accounting for LOS.

We examined multiple attributes of polypharmacy—the number of scheduled medications, number of as-needed medications, and number of scheduled doses per 24 hours. Interestingly, only the scheduled medications (count of medication and number of doses) were associated with elevated readmission risk. As-needed medications have heterogeneity

in the level of importance from critical (eg, seizure rescue) to discretionary (eg, antipyretics, creams). The burden of managing these types of medications may still be high (ie, parents must decide when to administer a critical medication); however, this burden does not translate into increased readmission risk in this population.

Not surprisingly, greater medical complexity—as defined by higher numbers of scheduled discharge medications and technology assistance—is associated with 30-day readmission risk. Our analyses do not allow us to determine how much of the increased risk is due to additional care burden and risks of polypharmacy versus the inherent increase in complexity and severity of illness for which polypharmacy is a marker. Tailoring discharge regimens to the realities of daily life, with the goal of “minimally disruptive medicine”^{22,23} (eg, integrating manageable discharge medication routines into school and work schedules), is not a common feature of pediatric discharge planning. For adult patients with complex medical conditions, tailoring medication regimens in a minimally disruptive way is known to improve outcomes.²⁴ Similarly, adopting minimally disruptive techniques to integrate the polypharmacy inherent in discharge could potentially mitigate some of the readmission risks for children and adolescents.

Contrary to our hypothesis, new technologies and new diagnoses did not confer additional readmission risk when accounting for LOS and patient characteristics. One potential explanation is varying risks conveyed by different types of new

technologies placed during hospitalization. Central lines, the most common form of new technology, is associated with higher odds of reutilization in unadjusted analyses. However, the second most common form of new technology, nonsurgically placed enteral feeding tube, was not. Further analyses of the differential effects of new technology should be further examined in larger datasets. Additionally, the lack of additional readmission risk from new technology may relate to additional teaching and support provided to families of patients undergoing unfamiliar procedures offsets the risks inherent of greater complexity. If so, it may be that the more intensive teaching and postdischarge support provided to families with new technology or a new diagnosis could be replicated through refresher teaching during hospitalizations, when a patient's state of health is status quo for the family (ie, the child was admitted and discharged with the same technology and diagnoses). This notion is supported by prior work that demonstrated successful readmission reduction interventions for children with chronic conditions often rely on enhanced education or coaching.^{25,26}

We elected to present models both with and without LOS as a confounder because it is a potentially modifiable attribute of hospitalization. Change in medical complexity aspects were significantly associated with readmission in multivariable models without LOS. However, with the addition of LOS, they were no longer significant. Thus, the readmission risk of new complexity is accounted for by the readmission risk inherent in a longer LOS. This finding prompts additional questions that merit further study: is it that LOS is a general marker for heightened complexity, or is it that a longer LOS can modify readmission risk through additional in-hospital care and time for enhanced education?

Our study has several strengths. We were able to discern true complexity at the time of discharge through medical record review. For example, if a child had a peripherally inserted central catheter placed during hospitalization, it cannot be ascertained through administrative data without medical record review if the technology was removed or in place at discharge. Likewise, medical record review allows for identification of medical technology which is not surgically implanted (eg, nasogastric feeding tubes). Given the "fog" families report as part of their in-hospital experience and its threats to education and postdischarge contingency planning,¹⁷ we felt it important to evaluate medical technology regardless of whether or not it was surgically placed. Additionally, the more detailed and nuanced understanding gained of polypharmacy burden can better inform both risk prediction models and interventions to improve the transition from hospital to home.

This study should also be considered in the context of several limitations. First, the data was from a single children's hospital, so the generalizability of our findings is uncertain. Second, we utilized a novel method for counting new CCCs which compared information collected for clinical purposes (eg, obtaining a past medical history) with data collected for billing purposes (ie, discharge diagnoses). This comparison of information collected for different purposes potentially introduced uncertainty in the classification of diagnoses as new or not new; however,

the interrater reliability for adjudicating new diagnoses suggests that the process was reasonably reliable. Third, we did not have access to other hospitals where readmissions could have occurred. While this is a common limitation for readmission studies,^{10,12,14,15,18,27-29} we attempted to mitigate any differential risk of being readmitted to other institutions by matching on distance and direction from the hospital. Of note, it is possible that children with medical complexity may be more willing to travel further to the hospital of their choice; thus our matching may be imperfect. However, there is no established method available to identify preadmission medical complexity through administrative data. Finally, the case-control method of the study makes estimating the true incidence of a variety of elements of medical complexity challenging. For example, it is difficult to tell how often children are discharged on five or more medications from a population standpoint when this practice was quite common for cases. Likewise, the true incidence of new technologies and new CCCs is challenging to estimate.

CONCLUSION

Medical complexity at discharge is associated with pediatric readmission risk. Contrary to our hypothesis, the addition of new technologies and new CCC diagnoses are not associated with pediatric readmission, after accounting for patient and hospitalization factors including LOS. The dynamics of LOS as a risk factor for readmission for children with medical complexity are likely multifaceted and merit further investigation in a multi-institutional study.

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