Primary care physician involvement during hospitalisation: a qualitative analysis of perspectives from frequently hospitalised patients

Erin Yildirim Rieger 1, Josef N S Kushner,2 Veena Sriram 3, Abbie Klein 4, Lauren O Wiklund 5, David O Meltzer 4, Joyce W Tang 4

ABSTRACT

Objective To explore frequently hospitalised patients’ experiences and preferences related to primary care physician (PCP) involvement during hospitalisation across two care models.

Design Qualitative study embedded within a randomised controlled trial. Semistructured interviews were conducted with patients. Transcripts were analysed using qualitative template analysis.

Setting In the Comprehensive Care Programme (CCP) Study, in Illinois, USA, Medicare patients at increased risk of hospitalisation are randomly assigned to: (1) care by a CCP physician who serves as a PCP across both inpatient and outpatient settings or (2) care by a PCP as outpatient and by hospitalists as inpatients (standard care).

Participants Twelve standard care and 12 CCP patients were interviewed.

Results Themes included: (1) Positive attitude towards PCP; (2) Longitudinal continuity with PCP valued; (3) Patient preference for PCP involvement in hospital care; (4) Potential for in-depth involvement of PCP during hospitalisation often unrealised (involvement rare in standard care); in CCP, frequent interaction with PCP fostered patient involvement in decision making; and (5) PCP collaboration with hospital-based providers frequently absent (no interaction for standard care patients; CCP patients emphasising PCP’s role in interdisciplinary coordination).

Conclusion Frequently hospitalised patients value PCP involvement in the hospital setting. CCP patients highlighted how an established relationship with their PCP improved interdisciplinary coordination and engagement with decision making. Inpatient–outpatient relational continuity may be an important component of programmes for frequently hospitalised patients. Opportunities for enhancing PCP involvement during hospitalisation should be considered.

INTRODUCTION

Before the mid-1990s, primary care physicians (PCPs) in the USA typically oversaw care for their own patients when they were hospitalised. Since then, the number of hospitalists has significantly increased within the USA. This shift in care delivery model was motivated by perceptions about increased hospitalist efficiency, availability, specialised expertise, and possible cost and mortality reductions. Despite such advantages, the hospitalist model may increase fragmentation between inpatient and outpatient care, particularly for patients who are frequently hospitalised. Previous studies found that hospitalised patients frequently had limited knowledge about their diagnosis, care plan or postdischarge instructions. Other studies identified discrepancies between hospitalised patients and their inpatient physicians in perceived goals of care, and limited opportunities for shared decision making. A possible contributing factor to these communication barriers is lack of an established relationship between hospitalised patients and their inpatient physicians. In comparison, PCPs with whom patients have ongoing relationships often have intimate knowledge of patients’ preferred communication style, values, family context and care preferences, but...
independently communicate directly with patients during hospitalisation. Communication between PCPs and inpatient providers during hospitalisation may also be limited in frequency and scope. 

Particularly for patients with complex needs, PCP involvement during hospitalisation may greatly impact patient experience due to their familiarity with their patients’ complex health history and established relationship with patients. However, few studies directly compare the hospitalisation experiences of patients cared for by their PCP versus by hospitalists. These are limited to quantitative comparisons related to satisfaction. There is a need for qualitative patient perspectives on the role of a PCP in the hospital setting.

We conducted a qualitative study of frequently hospitalised patients’ experiences and preferences related to PCP involvement during hospitalisation. This qualitative study was embedded within a larger randomised controlled trial, the Comprehensive Care Programme (CCP) Study. Patients at increased risk of hospitalisation are randomly assigned to one of two care models: (1) a CCP physician who serves as both the outpatient PCP and hospital attending (intervention arm) or (2) outpatient care from a PCP and hospital care from hospitalists (standard care). This study context provides a unique opportunity to explore and compare the experiences and preferences surrounding PCP involvement during hospitalisation between patients cared for by hospitalists as compared with a patient’s own PCP.

**METHODS**

**Setting**

The CCP study at the University of Chicago Medicine (UCM) is a randomised controlled trial assessing the effect of an interdisciplinary care team for patients at high risk of hospitalisation. The overall CCP study recruited Medicare part A and B enrollees with at least one hospitalisation at UCM within the previous twelve months. Patients randomised to the intervention group were cared for by PCPs with limited panels of approximately 200 patients to enable them to care for their patients as the primary attending in both inpatient and outpatient settings. As outpatients, the patients also receive care from a social worker, two nurses and a clinic coordinator. Patients randomised to the control group received ‘standard of care,’ which included following with their prior PCP (or were offered assistance in obtaining one if they did not have one) and being treated by hospitalist physicians if admitted to UCM.

The broader CCP study compares clinical outcomes, healthcare costs and experiences of patients in CCP versus standard care. This embedded study used qualitative interviews with a subset of both CCP and standard care patients to better understand and compare patients’ experiences and preferences surrounding the role of their PCP during hospitalisation.

**Participant selection**

The participants for the embedded qualitative study were drawn from the broader CCP study. Additional inclusion criteria included, participation in the CCP study for at least 1 year, and having at least three hospitalisations within the previous twelve months (based on self-report during quarterly phone surveys), with the most recent hospitalisation occurring at UCM. For intervention group patients, medical records were screened to confirm that their assigned CCP physician served as their primary attending during the most recent hospitalisation. Patients were recruited by a research assistant or medical student in-person or by phone between July 2017 and August 2018. Recruitment continued until data saturation was reached.

**Development of interview guide**

The semistructured interview guide was developed by an interdisciplinary team including a CCP physician (JWT), a medical student without ties to CCP (JNSK), a CCP social worker (NG) and a research assistant without ties to CCP (JH). Two members of the team (JWT, JH) had prior experience in qualitative research methods. The interview guide was further modified after review by three patients in the intervention arm of the CCP study. The final interview guide (online supplemental appendix A) focused on patients’ care experiences during and after their most recent hospitalisation at UCM, with an emphasis on: (1) communication with physicians and nurses in the hospital setting, particularly surrounding goals of the hospitalisation and decision making; (2) postdischarge care and (3) relationship with their PCP and their PCP’s role during hospitalisation.

**Data collection**

Semistructured interviews, approximately 30 min in length, were conducted in-person at UCM by a medical student and a research assistant, neither with ties to CCP (JNSK or AK). Patients provided verbal consent for the interview and received a US$30 gift card for participation. All patients had previously provided written consent for the broader CCP study. Patient characteristics including sex, age and healthcare utilisation were collected from the medical record.

**Data analysis**

The interviews were digitally recorded and transcribed by the research team; identifiable personal data were redacted. The interview transcripts were analysed using template analysis, a methodology developed by Crabtree and Miller. Template analysis was selected as a systematic yet flexible methodology that lends itself to analysis across the two groups of subjects. The qualitative analysis team (JWT, EYR and JNSK) was composed of one CCP physician and two medical students. The three team members separately reviewed five interview transcripts (three control group; two CCP group) and engaged in discussions to develop a preliminary ‘template’ (coding
guide), a hierarchical organisation of the identified themes. Some codes were identified inductively and others were rooted in the interview guide questions. Through an iterative process, additional codes were added to the template as they arose from the five sample transcripts. The three team members then applied the initial template to code the transcripts using NVivo V.11 (QRS International) software. Two coders reviewed each of the 24 transcripts. During analysis, the team met weekly to resolve discrepancies in coding through discussion and to revise the template. After the template was finalised, themes were developed through repeated review of codes and discussion. These themes were described, and representative quotes were selected and agreed on by the entire research team. The qualitative analysis team practised reflexivity through open communication about their preconceptions and how their roles in patient care relate to their perspective.

Patient and public involvement
Three CCP patients provided feedback on the interview guide during development. As a check on the validity of the analysis, results were reviewed and discussed with the CCP study patient and community advisory board and with CCP physicians, team members and administrators.

RESULTS

Patient sample
Twenty-four interviews were conducted, 12 with CCP patients and 12 with standard care patients. Patient characteristics are shown in table 1.

For CCP and standard care groups combined, 50% were female and the average age was 53 years. Patients had been enrolled in the CCP study for an average of 2.7 years at the time of interview. All patients had two or more chronic medical conditions. Of the standard care patients, 67% received primary care from internal medicine resident physicians; all CCP patients were cared for by attending physicians.

Theme 1: positive attitude towards PCP
Themes and additional quotes from interviews are summarised in table 2.

Patient is comfortable talking with their PCP
A majority of CCP patients and a few standard care patients described feeling comfortable conversing with their PCP. Patients in both groups valued that their PCP listened to what they said. Several CCP patients, but only a few of the standard care patients, thought discussions were better with their PCP than with hospital providers. A common perception was that the patient could speak more openly with their PCP.

It’s good because I can openly talk to him and not be afraid to tell him if something is not going right. (Female, standard care)

Several CCP patients also described engaging with their PCP about topics outside of medicine, including challenging social issues.

You want to be straight up with your primary about things. You want to tell him everything, what's giving you problems. Well my wounds are giving me problems, do you have any other issues? And you tell him: well depression issues, housing issues. (Male, CCP)

<table>
<thead>
<tr>
<th>Characteristics</th>
<th>All patients (n=24)</th>
<th>Standard care patients (n=12)</th>
<th>CCP patients (n=12)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age in years* (mean (SD))</td>
<td>53 (14)</td>
<td>57 (15)</td>
<td>49 (11)</td>
</tr>
<tr>
<td>Female (n (%))</td>
<td>12 (50)</td>
<td>6 (50)</td>
<td>6 (50)</td>
</tr>
<tr>
<td>Years since CCP study enrolment* (mean (SD))</td>
<td>2.7 (1.2)</td>
<td>2.7 (1.3)</td>
<td>2.7 (1.1)</td>
</tr>
<tr>
<td>Chronic medical conditions (n (%))</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Heart failure</td>
<td>11 (46)</td>
<td>7 (58)</td>
<td>4 (33)</td>
</tr>
<tr>
<td>Coronary artery disease</td>
<td>6 (25)</td>
<td>4 (33)</td>
<td>2 (17)</td>
</tr>
<tr>
<td>Diabetes</td>
<td>11 (46)</td>
<td>7 (58)</td>
<td>4 (33)</td>
</tr>
<tr>
<td>End-stage renal disease</td>
<td>10 (42)</td>
<td>5 (42)</td>
<td>5 (42)</td>
</tr>
<tr>
<td>Chronic lung disease†</td>
<td>8 (33)</td>
<td>7 (58)</td>
<td>1 (8)</td>
</tr>
<tr>
<td>Cancer</td>
<td>2 (8)</td>
<td>1 (8)</td>
<td>1 (8)</td>
</tr>
<tr>
<td>Rheumatologic disease‡</td>
<td>2 (8)</td>
<td>0</td>
<td>2 (17)</td>
</tr>
<tr>
<td>Other‡</td>
<td>2 (8)</td>
<td>1 (8)</td>
<td>1 (8)</td>
</tr>
</tbody>
</table>

*At time of interview.
†Chronic obstructive pulmonary disease, Asthma, Interstitial lung disease, Pulmonary arterial hypertension, Cystic fibrosis.
‡Scleroderma, Crohn’s disease.
§Sickle cell disease, Spina bifida.
CCP, Comprehensive Care Programme.
PCP is caring towards patient
Across CCP and standard care, several patients expressed that they felt cared for by their PCP. Patients appreciated that the PCP was concerned about their health and that the physician offered their time and attention.

She sits and talks to me, she […] connects with me, you know. Not just like business type thing, but a family type thing. (Male, CCP)

Shared trust between patient and PCP
Half of CCP patients, but none of the standard care patients, mentioned shared trust with their PCP. In several instances, CCP patients stated that their PCP trusted them to make their own decisions about care. Participants also described an increased level of trust in their CCP’s judgement over time.

See we have this really good relationship, and she knows that I am an informed patient. I know my body and I can pretty much tell her more than she can tell me about my body, and so she trusts me the same way I trust her. (Female, CCP)

**Theme 2: longitudinal continuity with PCP valued**
Many standard care and CCP patients emphasised that it takes time to build a relationship with a PCP. A majority of standard care participants described discontinuity with their PCPs. A few patients attributed the frequent changes to receiving primary care from residents who graduated and transitioned their patients to new trainees.

[...] this’ll be my 4th primary care doctor in 10 years. But, you know, you guys do your 2 or 3 year residency, then you’re off […] But at some point you’ve got to learn, like what you’re doing with me. (Male, standard care)

In comparison, CCP patients experienced greater relational continuity with their physician. A majority of CCP
patients mentioned that the relationship with their CCP physician improved over time as they got to know each other. Two participants described a difficult start that morphed into a positive relationship.

Once you get to know a person it gets better because you know the first couple of times it’s not going to be smooth sailing. (Male, CCP)

**Theme 3: patient preference for PCP involvement in hospital care**

A majority of standard care patients and nearly all CCP patients stated that their PCP should be involved during hospitalisation. However, there was variation in the preferred form of involvement. A majority of standard care patients thought that ideally a PCP would remain in communication with their patient during hospitalisation.

It shows they care, they’re concerned about my being, my health. And that’s a good thing. (Female, standard care)

Patients commonly preferred that this interaction be in person, with a few specifically stating that they would prefer to be treated by their PCP while in the hospital. A few standard care patients noted that their PCP could have been able to offer emotional support during hospitalisation.

[…] he probably could have informed me and let me know more about what was going on, and I would have had less anxiety. I would have felt more relieved. (Male, standard care)

Further, a few standard care patients thought that a PCP should remain in communication with the patient’s care team during hospitalisation. PCP involvement was considered beneficial due to their prior knowledge of the patient’s health conditions.

I don’t think he should just be in the office all the time. I think he should know about me being in the hospital and why and help me to maintain my health. I mean, these other doctors, they don’t really see me that much. They don’t know much about me. They just know what I came in and I complain about, and then they fix that and then they send me on my way? What if I have another issue that they’re not aware of that my primary care doctor is aware of? (Female, standard care)

On the other hand, some standard care patients stated that they chose not to reach out to their PCP during hospitalisation due to lack of a relationship with their PCP. Others shared that they had no expectation of PCP involvement during hospitalisation. These patients thought it was sufficient for the PCP to view records following discharge or answer questions if contacted by the hospital team.

I just concentrate on the doctors at hand, and I know they’re making notes so he see it on the chart. I don’t have it where I can text him and let him know each time I’m in the hospital […] So, by the time I see him, I’m quite sure there’s a flag somewhere to let him know I’ve been in the hospital, and he can read the chart and see that I’ve been in the hospital so. (Female, standard care)

Among CCP patients, all described a preference to be treated by their PCP in the hospital setting. For several, a PCP’s knowledge about their health and personal preferences was thought to expedite care and improve adherence to previously developed plans.

Besides my opinion, she should be able to make the decision. She should be the one running stuff. Should no other doctor be running nothing or make no decisions because you don’t know me. (Female, CCP)

A few CCP patients also pointed to the shared trust with their PCP. Due to previous hospitalisations or office visits patients perceived that the PCP had greater understanding of the patient’s preferences, and the patient felt comfortable with the PCP’s plan.

[…] I feel like with any other doctor, it would be like: ‘You were just ready to go home, now all of a sudden I say this and you’re not feeling well.’ I think she knows that it’s not just necessary that I’m not saying I’m not feeling well, I think she knows what I’ve told her already, why I like the blood transfusions, so she don’t look at it like a ploy. (Female, CCP)

**Theme 4: potential for in-depth involvement of PCP during hospitalisation often unrealised**

There was considerable variation between the two groups in interaction between the patient and PCP during hospitalisation. Most standard care patients did not interact with their PCP during hospitalisation.

He usually gets the report after I’m out […] I just go through whatever doctor sees me in the emergency room, and then they send to the floor. (Female, standard care)

For the few patients who described interaction, some initiated, and some were contacted by their PCP. The form of PCP–patient interaction also varied. A few patients received in-person visits from their PCP and one talked with their PCP over the phone. Patients expressed positive feelings towards their PCP visiting them in the hospital. However, there was little elaboration about how and to what extent the PCP actively participated in their care during hospitalisation.

For CCP patients, nearly all described frequent in-person interaction with their PCP during hospitalisation. Most patients discussed their plan of care with the PCP. Half described making decisions with their PCP about treatment options or the timing of discharge.

Mainly I talked to my Comprehensive doctor. She’s like the main authority over all of that. They have
to talk to her first, you know to see if I’m okay with leaving. She’ll come ask me: ‘How do you feel about leaving today?’ If I say ‘I don’t feel like leaving,’ she’ll be like: ‘You can stay an extra day.’ […] (Male, CCP)

**Theme 5: PCP collaboration with hospital-based providers frequently absent**

There was also significant variation between the two groups in experience of interaction between the PCP and hospital providers. Only two standard care patients described being aware of interaction between their PCP and hospital providers, such as providing guidance to the inpatient care team. A few standard care patients openly expressed uncertainty and concern about whether their PCP was contacted during their hospitalisation.

I don’t know for sure that they’re calling him and letting him know or if he’s getting the reports or any of that. I need to know that he’s getting this information to know I’m there. (Female, standard care)

Among CCP patients, a majority described their PCP being in communication with other hospital providers. Several of these patients referred to the PCP as leader of their healthcare team in the hospital. Patients described their PCPs keeping specialists informed and interfacing with the other providers when the patient had a concern or conflict.

She’s my main doctor, so she makes sure everybody gets the email when I’m in the hospital. They’ll know that: ‘OK I gotta go see how he’s doing, and see if I can give him any help for his pain or anything.’ So that’s the best thing I can ask for. That’s probably why I switched from another hospital. Since I can just ask my Comprehensive Care anything wrong with me, she’ll make sure that all my other doctors know too, so I ain’t gotta be worrying about it, like my pain, or if I miss an appointment they’ll all be informed that I’m in the hospital […] (Male, CCP)

**DISCUSSION**

The main aim of this study was to explore frequently hospitalised patients’ experiences and preferences related to PCP involvement during hospitalisation. A unique contribution of this study was the qualitative comparison of perspectives of standard care patients who were cared for by hospitalists or housestaff teams to those of CCP patients being treated by their own PCP during hospitalisation.

Both standard care and CCP patients expressed a preference for repeated interactions with their PCP over time to build a relationship and shared knowledge. While CCP patients described consistent relationships with their PCPs that benefited from shared experiences across inpatient and outpatient settings, many standard care patients described relational discontinuity with PCPs, which sometimes weakened these relationships. These results were consistent with prior research that patients prefer, and may benefit from, relational continuity of care with physicians, and that patients’ trust in their PCP was associated with the duration of their relationship.

It is concerning that in this study of patients with frequent hospitalisation and multiple chronic conditions, many in standard care may not experience the benefits of long-term relational continuity. Most of the patients experiencing discontinuity received care in a resident clinic characterised by frequent turnover. It is possible that the purposive sampling of this embedded qualitative study disproportionately selected for standard care patients with resident PCPs. Unfortunately, these patients’ experiences with PCP discontinuity are not unique. Previous studies found that, as compared with patients with attending PCPs, patients with resident PCPs were more likely to have multiple health conditions, and be non-white, of low socioeconomic status, and on Medicare or Medicaid insurance. Patients who transition care to a new resident reported challenges including missed tests and difficulty building a relationship with a new provider. Patients may also experience PCP discontinuity due to the resident clinic schedule.

A vast majority of patients in this study wanted their PCP to be involved during hospitalisation, a preference consistent with previous findings. Despite the overall preference for PCP involvement during hospitalisation, few standard care patients described actual involvement of their PCP during hospitalisation; when involved, the PCP role was usually limited to single visits or brief conversations with the patient or hospital providers. The finding that a majority of the standard care patients did not have interaction with their PCPs during hospitalisation echoes previous research.

In contrast, consistent with the structure of the programme, CCP patients described substantial involvement of their PCPs during hospitalisation. A major contribution of this study was in highlighting the value of PCP involvement in the hospital setting through the lens of patients in CCP. Specifically, patients in CCP emphasised the PCP’s role as a leader of their care team. Patients found it reassuring to have their PCP working to align the knowledge and goals of the various hospital providers. CCP patients expressed that shared trust with their PCP allowed for more patient involvement in care decisions due to greater patient comfort to voice disagreement, and PCP respect for the patient’s input. As the CCP model is further developed and disseminated to other care settings, longitudinal relationships and direct patient engagement in the inpatient setting will be important components to uphold.

Inpatient–outpatient relational continuity is a component of other interdisciplinary programmes for frequently hospitalised patients. The nature of team involvement in the inpatient setting varies. For instance, the University of Colorado intensive outpatient clinic team collaborates with hospital providers to develop care plans. In
the CareMore Health System, hospitalists treat high-risk patients for a limited duration across the transition from inpatient to rehabilitation or community settings. Social workers in the Northwestern University Complex High Admission Management Programme provide continuity by rounding on their admitted patients. It is unknown if and to what extent findings from this CCP study may translate to programmes with inpatient–outpatient continuity involving a non-PCP provider, or inpatient involvement that is not direct care. Incorporating patient perceptions into evaluation plans for these interdisciplinary programmes could refine our understanding of the nature of involvement needed.

In practice, it is uncommon for patients in the USA to be treated by their PCP while hospitalised. In a sample of 2013 Medicare data, PCPs cared for their own patient in only 14.2% of hospital admissions. However, for frequently hospitalised patients, increasing PCP engagement in the inpatient setting may improve patient experiences, even if the PCP is not providing direct care. PCPs can use their relationship with the patient to help assess preferences and identify needs. This may benefit the patient by encouraging patient engagement in decision-making, strengthening the patient–PCP relationship, and improving interdisciplinary coordination across settings. To achieve this, a first challenge is ensuring that PCPs receive information when their patient is hospitalised.

Healthcare systems may also consider how to provide PCPs with time and compensation for communicating with their hospitalised patients and their inpatient care teams by phone or in-person visit. There are several limitations of the patient sample and analysis. First, patients recruited for the embedded qualitative study may have been a healthier and more engaged group than the overall study population. In the case of CCP patients, those with positive feelings towards the programme may have been most likely to participate. Second, while all CCP patients had attending physicians as PCPs, 67% of the standard care patients had resident physicians as PCPs. Although a limitation, it also reflects the reality that complex, vulnerable, patients who experience frequent hospitalisations often receive primary care from residents. Third, the exclusive focus on PCPs in the analysis is a limitation. CCP PCPs may share similar roles or characteristics with specialists or other providers who see patients across care settings.

**CONCLUSION**

In summary, this study was a valuable contribution to the existing literature on PCP involvement during hospitalisation due to the qualitative comparison of perspectives of standard care and CCP patients. Specifically, the results suggested that for frequently hospitalised patients, active inpatient involvement by a consistent PCP with knowledge of the patient’s health and personal preferences could improve patient experience with interdisciplinary coordination and engagement in care during hospitalisation. For frequently hospitalised patients not being treated in the hospital by their PCP, future research is needed to clarify which forms of PCP engagement may be most likely to confer these benefits.

**Twitter** Erin Yildirim Rieger @ereriyrieger, Veena Sriram @veena_sriram and Joyce W Tang @joycewtang

**Acknowledgements** Thank you to the patients interviewed in the study for their time and willingness to discuss their medical care. Thank you to Nicole Gier and Jannelle Highland for their assistance with developing the interview guide. Thank you also to Andrea Flores, MA for her role in developing the sampling approach used in this study.

**Contributors** EYR: formal analysis, writing—original Draft. JN:K: methodology, formal analysis, investigation, writing—original Draft. VS: methodology, writing—review & editing. DOM: conceptualisation, writing—review and editing, funding acquisition. JWT: conceptualisation, methodology, formal analysis, writing—original draft, supervision, guarantor.

**Funding** This work was supported by: Centers for Medicare and Medicaid Services, Center for Medicare and Medicaid Innovation grant CMS-101-12-0001; Agency for Healthcare Research and Quality, Patient Centered Outcomes Research Faculty Development Program grant K12; and National Institutes of Health, National Institute on Aging grant 5T35AG029795-12.

**Competing interests** None declared.

**Patient consent for publication** Not applicable.

**Ethics approval** This study received approval from the University of Chicago Institutional Review Board (IRB12-1440).

**Provenance and peer review** Not commissioned; externally peer reviewed.

**Data availability statement** Data are available on reasonable request. Full transcripts have not been shared to protect anonymity of patients.

**Supplemental material** This content has been supplied by the author(s). It has not been vetted by BMJ Publishing Group Limited (BMJ) and may not have been peer-reviewed. Any opinions or recommendations discussed are solely those of the author(s) and are not endorsed by BMJ. BMJ disclaims all liability and responsibility arising from any reliance placed on the content. Where the content includes any translated material, BMJ does not warrant the accuracy and reliability of the translations (including but not limited to local regulations, clinical guidelines, terminology, drug names and drug dosages), and is not responsible for any error and/or omissions arising from translation and adaptation or otherwise.

**Open access** This is an open access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited, appropriate credit is given, any changes made indicated, and the use is non-commercial. See: http://creativecommons.org/licenses/by-nc/4.0/.

**ORCID iDs** Erin Yildirim Rieger http://orcid.org/0000-0002-0888-5056
Veena Sriram http://orcid.org/0000-0001-7235-253X
Abbie Klein http://orcid.org/0000-0002-1077-1269
Lauren O Wiklund http://orcid.org/0000-0001-9987-0813
David O Meltzer http://orcid.org/0000-0002-6762-8729
Joyce W Tang http://orcid.org/0000-0003-4792-924X

**REFERENCES**