

ORIGINAL RESEARCH

Having a Primary Care Provider is the Strongest Predictor of Successful Follow-up of Participants in a Clinical Trial

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Purpose: Ethnic minorities, women, and those of low socioeconomic status are widely underrepresented in clinical trials. Few studies have explored factors associated with successful follow-up in these historically difficult-to-reach patients. This study's objective was to identify patient characteristics and methods of contact that predict successful contact for follow-up in an urban, predominantly ethnic minority, majority-women, poor population to help devise strategies to improve retention.

Methods: We retrospectively reviewed records from a prospective randomized control trial of 400 hospitalized chest pain patients to determine which characteristics were associated with successful telephone follow-up at 1 year after enrollment. We assessed demographic variables, medical history, and social factors by using bivariate analyses. A multivariate analysis was performed using variables from the bivariate analysis with $P \leq .2$.

Results: The overall successful 1-year follow-up rate was 95% (381/400). Study participants who completed follow-up were significantly more likely to have a primary care physician (PCP) (88% [337/381] versus 68% [13/19]), speak English natively (52% [199/381] versus 26% [5/19]), have a higher Charlson comorbidity index score, and identify as women (64.0% [244/381] versus 42.1% [8/19]). Having a PCP and native English language remained significant at multivariate analysis. Socioeconomic status score, quantity of contact information recorded at recruitment, and insurance status were not significantly associated with successful follow-up.

Conclusions: Patients engaged with the health care system by having a PCP are significantly more likely to achieve follow-up. Successful follow-up is also associated with native English speaking. The potential of improving follow-up by facilitating connections with health care providers requires further study. (J Am Board Fam Med 2020;33:431–439.)

Keywords: Ethnic Groups, Minority Groups, Retrospective Studies

Introduction

Clinical research informs decision making and provides the foundation for evidence-based medicine and best practices.¹ Comparative effectiveness

research with randomized clinical trials is a preferred method for testing an intervention for a particular population or setting.² This helps clinicians to maximize outcomes, knowing that the evidence they are using is specific for the task at hand and tested under comparable conditions.

The length of follow-up and retention are particularly important, as truncated follow-up can erroneously indicate that an intervention is more or

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less efficacious than it would be with long-term, more realistic use.^{3,4} A significant challenge in clinical research is keeping participants within a study for an extended period. Loss of research subjects in follow-up occurs for numerous reasons.^{3,5} It is useful to understand why this happens so that efforts can be made to mitigate that potential loss.⁶

It is well documented that racial/ethnic minorities are underrepresented in clinical trials^{7,8} and, in particular, cardiovascular studies. Some of the often-cited reasons include a distrust of medical research, economic burdens, and poor access to care.^{8,9} With regard to cardiovascular health in general, racial/ethnic minorities have both a higher prevalence of disease and worse outcomes.¹⁰ These inequalities are particularly problematic because the existing literature provides suboptimal guidance for the treatment of some patients. Women are widely underrepresented within clinical trials.¹¹⁻¹³ The dominance of white males in cardiovascular clinical trials has resulted in data that, in many cases, apply to less than half of the population. As a result, funding organizations have placed renewed emphasis on the explicit study of minority populations and women.¹⁴

Some of the challenges involved in performing the multicultural and gender-balanced clinical research needed to inform practice are already apparent. Patient characteristics, including male gender,¹⁵ black race,¹⁶ and young children in the household¹⁷ are associated with reduced retention rates in clinical trials. Patient involvement in the medical system and language preferences are potential determinants of success in follow-up. Different methods of patient recruitment, community involvement, and cultural tailoring have been studied to improve trial retention.¹⁸ Identifying correlates of poor follow-up allows a specific direction of efforts to retain at-risk patients and appropriate allocation of study resources.

The purpose of the present study is to assess the factors associated with successful 1-year follow-up from a multiethnic, majority-women, urban, cardiovascular randomized controlled trial to inform strategies to improve patient retention.

Methods

The Clinical Trial is registered at <https://clinicaltrials.gov/ct2/show/NCT00705458> under ClinicalTrials.gov Identifier NCT00705458.

Study Design

This study is a secondary analysis of the PROSPECT trial, which was conducted between July 2008 and December 2013.^{19,20} PROSPECT was a randomized controlled trial of 400 intermediate-risk chest pain patients admitted to telemetry at Montefiore Medical Center in the Norwood section of Bronx, NY. There were no recruitment restrictions regarding gender, race/ethnicity, native language, or health insurance. Participants were randomized to receive coronary computed tomography or radionuclide stress myocardial perfusion imaging, with the primary outcome variable being cardiac catheterization not leading to revascularization within 1 year. There was no statistically significant difference between the 2 imaging modalities with regard to the primary outcome. The trial design and primary outcome variable required 1 year of follow-up after recruitment.

Enrollment and Follow-up

Enrollment occurred between July 2008 and March 2012. At the time of enrollment, patients were asked for multiple phone numbers and the name and phone number of a relative or other close contact. Patients were called for telephone questionnaires at 6 and 12 months after enrollment to discover subsequent cardiac events and hospitalizations. If a patient could not be reached by telephone after at least 4 attempts either directly or via a close contact, further attempts were made through a primary care physician (PCP) or cardiologist identified from electronic medical records. Providers and facilities outside of Montefiore were contacted by telephone for records when patients provided this information and their permission. Additionally, continuous telephone and electronic (ie, e-mail and texting) follow-up after 1 year was conducted in a chronological fashion based on enrollment date. At the end of the study period, patients were contacted again to determine whether they experienced major adverse cardiovascular events.

Follow-up Outcome and Variables of Interest

The current study's primary outcome variable was defined as successful contact with the patient at or beyond 12 months after enrollment. This exploratory analysis assessed potential predictor variables, including sociodemographic data (age; race/ethnicity; gender; native or nonnative English speaker; socioeconomic status (SES) score based on

Table 1. Patient Baseline Characteristics (n = 400)

Characteristic	Values*
Age (y)	57 ± 11
BMI (kg/m ²) [†]	31 ± 6.4
SES [‡]	-3.95 (-6.58, -1.55)
Gender	
Men	148 (37.0%)
Women	252 (63.0%)
Ever smoked	
Yes	169 (42.3%)
No	231 (57.8%)
Current smoker	
Yes	59 (14.8%)
No	341 (85.3%)
Diabetes	
Yes	127 (31.8%)
No	273 (68.3%)
Hypertension	
Yes	288 (72.0%)
No	112 (28.0%)
Dyslipidemia	
Yes	206 (51.5%)
No	194 (48.5%)
PCP [§]	
Yes	350 (87.5%)
No	50 (12.5%)
Close Contact	
Yes	359 (89.8%)
No	41 (10.3%)
Number of phone numbers	2 (2, 3)
Charlson score	0 (0, 1)
Interpreter used	
Yes	103 (25.85)
No	297 (74.3%)
Native Language	
English	204 (51.0%)
Non-English	196 (49.0%)
Race	
Asian	18 (4.5%)
Black	143 (35.8%)
Hispanic	215 (53.8%)
Multiracial	6 (1.5%)
White	18 (4.5%)

Continued

employment, housing, and education in their ZIP code, normalized to the state average), medical comorbidities (diabetes; obesity; hypertension; dyslipidemia; current or prior smoking; Charlson comorbidity score), health insurance status, having a PCP (defined by current provider listed in the electronic medical record or stated by the participant),

Table 1. Continued

Characteristic	Values*
Insurance	
Commercial	118 (29.5%)
Medicaid	177 (44.3%)
Medicare	98 (24.5%)
Self pay	7 (1.8%)

*Categorical variables are presented as frequencies and percentages. Continuous variables are presented as either means and standard deviations or medians and interquartile ranges, depending on distribution.

[†]BMI, body mass index.

[‡]SES, socioeconomic score. Displayed as a Z score calculated based on the person's address and 6 socioeconomic variables for the neighborhood by zip code (log of median household income; log of median value of housing units; the percentage of households receiving interest, dividend, or net rental income; education; the percentage of adults who completed college; and the percentage of employed individuals in executive, managerial or professional positions) and normalized to the New York State average.

[§]PCP, primary care physician.

number of phone numbers provided for follow-up, having a close contact, and whether an interpreter was used during enrollment. All modifiable variables (eg, insurance status) were captured at time of enrollment and were not updated over the course of follow-up.

The secondary outcome was defined as the percentage of potential follow-up that was actually achieved. The potential follow-up period was calculated by determining the number of days from each patient's enrollment date to the end date of the study (December 26, 2013). The actual follow-up period was calculated by determining the number of days from the enrollment date until the date the patient was most recently followed-up by electronic medical records or telephone.

Analysis

To assess association with follow-up, bivariate analyses using χ^2 or Fisher's exact test for categorical variables, and 2-tailed t test or Wilcoxon-Mann-Whitney U test for continuous variables was performed.

Logistic regression analysis using a stepwise model selection strategy was performed, with a significance of 0.2 required for variable inclusion into the model and a significance of 0.05 to stay in the model. The odds ratio of having successful follow-

Table 2. Bivariate Analysis of Independent Risk Factors for Loss of Follow-up*

Factor	No One-Year Follow-up (n = 19)	Successful One-Year Follow-up (n = 381)	P Value
Age (y)	56 ± 10	57 ± 11	.7512
BMI (kg/m ²) [†]	29 ± 6.7	31 ± 6.4	.1862
SES [‡]	-3.1371 (-6.56945, -1.675089)	-3.9734 (-6.6542, -1.647682)	.5627
Gender			.0532
Men	11 (57.9%)	137 (36.0%)	
Women	8 (42.1%)	244 (64.0%)	
Ever smoked			.6435
Yes	9 (47.4%)	160 (42.0%)	
No	10 (52.6%)	221 (58.0%)	
Current smoker			.5025
Yes	4 (21.1%)	55 (14.4%)	
No	15 (78.9%)	326 (85.6%)	
Diabetes			.4493
Yes	4 (21.1%)	123 (32.3%)	
No	15 (78.9%)	258 (67.7%)	
Hypertension			.7218
Yes	13 (68.4%)	275 (72.2%)	
No	6 (31.6%)	106 (27.8%)	
Dyslipidemia			.1902
Yes	7 (36.8%)	199 (52.2%)	
No	12 (63.2%)	182 (47.8%)	
PCP [§]			.0010
Yes	13 (68.4%)	337 (88.5%)	
No	6 (31.6%)	44 (11.5%)	
Close contact			.7071
Yes	18 (94.7%)	341 (89.5%)	
No	1 (5.3%)	40 (10.5%)	
Number of phone numbers	2 (2, 2)	2 (2, 3)	.4856
Charlson score	0 (0, 1)	1 (0, 1)	.0480
Interpreter used			.2572
Yes	7 (36.8%)	96 (25.2%)	
No	12 (63.2%)	285 (74.8%)	
Native Language			.0274
English	5 (26.3%)	199 (52.2%)	
Non-English	14 (73.7%)	182 (47.8%)	
Race			.4468
Asian	0 (0.0%)	18 (4.7%)	
Black	4 (21.1%)	139 (36.5%)	
Hispanic	14 (73.7%)	201 (52.8%)	
Multiracial	0 (0.0%)	6 (1.6%)	
White	1 (5.3%)	17 (4.5%)	
Insurance			.9068
Commercial	5 (26.3%)	113 (29.7%)	
Medicaid	10 (52.6%)	167 (43.8%)	
Medicare	4 (21.1%)	94 (24.7%)	
Self pay	0 (0.0%)	7 (1.8%)	

*Categorical variables are presented as frequencies and percentages. Continuous variables are presented as either means and standard deviations or medians and interquartile ranges, depending on distribution.

[†]BMI, body mass index.

[‡]SES, socioeconomic score. Displayed as a Z score as in the previous table.

[§]PCP, primary care physician.

up, given 1 unit increase/category change of the patient factors, was estimated.

With regard to the secondary outcome, the percentage of follow-up completed, separated into tertiles, was the dependent variable. Bivariate analysis for 2 category variables used a 2-tailed *t* test, and analysis of variance was used for variables with more than 2 categories. Linear regression was used for continuous variables. The degree of follow-up was separated into tertiles for the purpose of an ordered logistic regression analysis, which was applied to variables that were significant or near-significant on bivariate analysis. A *P* value of $<.05$ was considered significant.

Results

Follow-up

At the midpoint of recruitment ($n = 200$), 64% (63/99) of patients eligible for 6-month follow-up were reached. By the completion of the study, 95.3% (381/400) had successful 1-year follow-up. No patients died within 12 months of recruitment.

Group Characteristics

The average age for cohort was 57 years; 63% were women, 53.8% were Hispanic, 35.8% were black, and 49.0% had a non-English native language (see Table 1).

Participants who had successful follow-up were significantly more likely to have a PCP (88.5% vs 68.4%, $P < .009$) and report a native language of English (52.2% vs 26.3%, $P < .027$) than those lost to follow-up (Table 2). Although specific medical diagnoses were not associated with follow-up, those who had successful follow-up had a higher Charlson comorbidity score (median, 1 vs 0; $P < .0480$). Gender was close to being a statistically significant variable, as women were more likely to achieve follow up ($P = .532$).

Both PCP and native English language remained significant at logistic regression analysis (Table 3). With native language held constant, having a PCP

resulted in 3.5 times the odds of completing the year of follow-up (odds ratio = 3.5; 95% CI, 1.25–9.75). Alternatively, with PCP held constant, having a native language of English was associated with 3.0 times the odds of completing the year of follow-up (odds ratio = 3.0; 95% CI, 1.06–8.62).

Patients who had a PCP were more likely to be in the higher follow-up tertiles ($P = .0046$) (Table 4). Women had an increased representation in the higher follow-up tertiles than men ($P = .019$). Although the secondary analysis produced multiple significant variables at the bivariate stage, no variables remained significant for the secondary outcome after logistic regression analysis.

Discussion

Successful follow-up and clinical trial retention of patients from an urban, predominantly minority, majority-women, poor population is challenging. In the present study, we found that having a PCP made a patient more than 3 times as likely to complete a year of follow-up, with almost nine-tenths of patients with PCPs having successful follow-up compared with just more than two-thirds of patients without a PCP. Having a PCP was also associated with achieving follow-up over a greater proportion of the potential follow-up period (86% versus 76%). Having a PCP may keep a patient more connected with the health care system, thus serving as an important liaison between clinical trial staff and research participants. An implication for research practice is that a patient's PCP status could allow trial designers to focus more resources on maintaining follow-up with participants who have fewer connections to their health care system. An upstream effect of this could be an increased emphasis on connecting patients to PCPs. With regard to current clinical practice, increasing the role of primary care in medicine is an important topic that has ramifications for cost and health outcomes more generally.²¹

It is likely that having a PCP is a proxy for a multitude of factors, including SES, social demands, and personal interest in physical health. Having a PCP may signify higher levels of commitment and flexibility, as well as satisfaction with the health care system. Some studies suggest that health care satisfaction and commitment play a positive role in trial enrollment and retention of minorities.^{22,23} External factors, such as transit options, the local health care

Table 3. Logistic Regression Analysis of Factors Associated with Successful Follow-up

Factor	Odds Ratio (95% CI)	<i>P</i> Value
PCP,* Yes	3.487 (1.247–9.753)	.0173
Native Language, English	3.030 (1.063–8.621)	.0381

*PCP, primary care physician; CI, confidence interval.

Table 4. Bivariate Analysis of Independent Variables for Achieving Complete Follow-up*

Variable	% Follow up 1 st Tertile	% Follow up 2 nd Tertile	% Follow up 3 rd Tertile	P Value
Possible Follow-up (days)	1,359 ± 378	1,391 ± 377	1,446 ± 376	.1289
Age (y)	55 ± 11.2	60 ± 10.9	55 ± 10.9	.0016
BMI (kg/m ²) [†]	31 ± 7.2	31 ± 6.3	30 ± 5.8	.9513
SES [‡]	-4.211	-4.0430	-4.0625	.8651
Gender				.0019
Women	71 (53.4%)	100 (74.1%)	81 (61.4%)	
Men	62 (46.6%)	35 (25.9%)	51 (38.6%)	
Ever smoked				.2245
No	83 (62.4%)	73 (54.1%)	75 (56.8%)	
Yes	50 (37.6%)	62 (45.9%)	57 (43.2%)	
Current smoker				.6544
No	115 (86.5%)	112 (83.0%)	114 (86.4%)	
Yes	18 (13.5%)	23 (17.0%)	18 (13.6%)	
Diabetes				.0191
No	99 (74.4%)	80 (59.3%)	94 (71.2%)	
Yes	34 (25.6%)	55 (40.7%)	38 (28.8%)	
Hypertension				.2592
No	42 (31.6%)	31 (23.0%)	39 (29.5%)	
Yes	91 (68.4%)	104 (77.0%)	93 (70.5%)	
Dyslipidemia				.0027
No	69 (51.9%)	32 (30.5%)	63 (47.7%)	
Yes	64 (48.1%)	73 (69.5%)	69 (52.3%)	
PCP [§]				.0046
No	24 (18.0%)	7 (5.2%)	19 (14.4%)	
Yes	109 (82.0%)	128 (94.8%)	113 (85.6%)	
Close contact				.2400
No	9 (6.8%)	15 (11.1%)	17 (12.9%)	
Yes	124 (93.2%)	120 (88.9%)	115 (87.1%)	
Number of phone numbers	2 (2, 3)	2 (2, 3)	2 (2, 3)	.3205
Charlson score	0 (0, 1)	1 (0, 2)	1 (0, 1)	.5851
Interpreter used				.9130
No	97 (72.9%)	101 (74.8%)	99 (75.0%)	
Yes	36 (27.1%)	34 (25.2%)	33 (25.0%)	
Native language				.2019
English	67 (50.4%)	62 (45.9%)	75 (56.8%)	
Non-English	66 (49.6%)	73 (54.1%)	57 (43.2%)	
Race				.4560
Asian	4 (3.0%)	6 (4.4%)	8 (6.1%)	
Black	49 (36.8%)	44 (32.6%)	50 (37.9%)	
Hispanic	72 (54.1%)	77 (57.0%)	66 (50.0%)	
Multiracial	2 (1.5%)	4 (29.6%)	0 (0.0%)	
White	6 (4.5%)	4 (29.6%)	8 (6.1%)	
Insurance				.0619
Commercial	49 (36.8%)	29 (21.5%)	40 (30.3%)	
Medicaid	51 (38.3%)	58 (43.0%)	68 (51.5%)	
Medicare	29 (21.8%)	48 (35.6%)	21 (15.9%)	
Self pay	4 (3.0%)	0 (0.0%)	3 (2.3%)	

*For each bivariate analysis, % follow-up complete was divided into tertiles. Categorical variables are presented as frequencies and percentages. Continuous variables are presented as either means and standard deviations or medians and interquartile ranges, depending on distribution.

[†]BMI, body mass index.

[‡]SES, socioeconomic score. Displayed as a Z score as in the previous table.

[§]PCP, primary care physician.

infrastructure, and social policies, also play a role. At the individual level, establishing care takes time and, often, money; yet, it is notable that SES score was not a significant predictor of follow-up. It is also notable that having a PCP remained a strong factor when included in a multivariate logistic regression with native language, another significant predictor of follow-up. This suggests that it is not just the ease of communication but rather the process of establishing and maintaining care that is associated with follow-up. When a patient is enrolled in a study, the lack of a PCP indicates that they may be less able to meet the trial's requirements and could require additional support.

Native English speaking was also a significant predictor of successful follow-up. Numerous languages, including Arabic, Bengali, Creole (Haiti), Greek, Khmer (Cambodia), Mende (Sierra Leone), Serbian, Spanish, Tagalog (Philippines), Twi (Ghana), Vietnamese, and Wolof (The Gambia), were preferred by our study patients. Native language may be a proxy for culture, and cultural nuances also play roles in the emphasis on social support, deference to physicians/researchers, and preferential modes of communication. In addition, some of these patients may have been visitors and noncitizens who returned to their country of origin and are, hence, difficult to locate. Some were without regular access to a telephone. In a notable example, 1 trial participant was deaf and preferred follow-up by physical mail correspondence.

Although being a woman did not reach significance as a predictor of follow-up in the primary analysis, women completed a significantly larger percentage of their potential follow-up period in the secondary analysis. Previous studies of retention in communities of racial/ethnic minorities show that men are more likely to be lost.¹⁵ Gender stereotypes suggest that women are more often caretakers in the home and men may work outside of the home more frequently or for longer hours. In addition, women more commonly present to their PCP for routine care and during illness.²⁴ In fact, women represent the majority of people receiving care at our medical center. The relative ease of contacting women research participants is a somewhat unexpected boon for efforts favoring improved trial design to adequately represent women.

We anticipated that 3 variables in particular would be significantly associated with 1-year

follow-up: SES score, contact information (both number of phone numbers and having a close contact), and insurance status. Surprisingly, none of these variables were significant. The SES score represents how a patient's ZIP code's income, education, and employment relate to the New York State average. The trial population as a whole was 3 standard deviations below the NY average,²⁰ and it may be difficult to detect differences in our uniformly low SES score population. Our observation is consistent with prior research in low-income minority populations.^{17,25}

We surmised that contact information would play a role—if a participant provides 1 phone number versus another who gives out 4, there ought to be a difference in ease of follow-up 12 months later. However, the electronic medical record served as a fallback in our study, providing physician contact information and care updates. Hence, patients who provided fewer phone numbers could achieve follow-up via electronic records. We do not have data to show which particular resources were used to contact each participant for their 12-month follow-up. The widespread use of cell phones likely mitigates the historic need for multiple phone numbers, as cell phone numbers typically remain constant. Similar to our need for various methods of contact, having multiple methods to contact participants and collect data has been discussed as a potential avenue of increasing retention in trials.²⁵ A recent assessment of retention strategies in the United Kingdom revealed a similar line of thinking, with various methods used, including contacting PCPs and SMS text messaging.²⁶ The effectiveness of phone calls to participants is uncertain, as 1 recent study showed that regular phone calls from a coordinating center did not improve retention.²⁷ Our study did not use phone calls as a variable; yet, this offers an interesting contrast.

We anticipated that patients who did not have insurance would be less likely to receive follow-up. These participants have fewer resources, so a lack of time and money could result in difficulty communicating. However, we did not find insurance status to be significant. Patients with Medicaid comprised a nonsignificantly greater percentage of the group that did not achieve follow-up.

We expected that patients with a 1-year potential follow-up duration would more readily achieve complete follow-up than those with a potential follow-up duration of 5 years. This was not the case,

suggesting that the follow-up period is not as important as the characteristics of the population. This supports the feasibility of trials with longer follow-up periods.

Limitations

The moderate-sized cohort (n = 400) is a limitation of this study. In particular, the group that did not receive follow-up totaled only 19 patients. Several variables had small subgroups. Despite the limitation in size, the relationship between having a PCP and being a native English speaker were statistically significantly related to successful follow-up and are likely to be clinically meaningful. A larger study would be beneficial in terms of adding multiple centers with different encatchments, as this trial was performed at a single site with a single, albeit diverse, patient mix. The population that was studied (minority, low SES, majority-women) varies considerably from the demographics of many areas. Potential interventions to improve follow up were not employed in this study, nor did we look at whether patients with PCPs might respond differently to said interventions than patients without PCPs. We were reliant on participants informing us of their use of outside doctors and hospitals; not having consistent access to participants' entire medical networks was another constraint. In addition, there was no third-party verification of whether participants had a PCP; the reliance on electronic medical records and self-reporting of PCP status is an important limitation. Although the variable of having a PCP was statistically significant, PCPs were also themselves contacted to assist with follow-up, resulting in possible confounding variables. However, the various phone numbers and close contacts were used for follow-up and neither was statistically significant. Therefore, we suggest that methods of contacting patients did not impact follow-up as much as inherent patient characteristics.

Future Directions

In conclusion, we found that successful follow-up was associated with patients having a PCP and being native English speakers. When designing clinical trials, especially in an urban, culturally diverse, underserved environment, it is vital to assess whether patients are connected to the health care system and that their barriers to care, both internal and external, are addressed. Most importantly, exploring whether participants have a PCP

should allow researchers to customize follow-up protocols. Other work has recently asked whether routine clinical care could improve retention,²⁷ and indeed, this question warrants additional study. Successful follow-up depends on an awareness of a patient's individual needs and communication preferences. It is also important that language and cultural differences are considered. This can be difficult in the fast-paced environment of medical data gathering, but the rigor and generalizability of outcomes research is at stake.

To see this article online, please go to: <http://jabfm.org/content/33/3/431.full>.

References

1. Lord SJ, Irwig L, Simes RJ. When is measuring sensitivity and specificity sufficient to evaluate a diagnostic test, and when do we need randomized trials? *Ann Intern Med* 2006;144:850–5.
2. Lauer MS. Comparative effectiveness research: the view from the NHLBI. *J Am Coll Cardiol* 2009;53:1084–6.
3. Akl EA, Briel M, You JJ, et al. Potential impact on estimated treatment effects of information lost to follow-up in randomised controlled trials (LOST-IT): systematic review. *BMJ* 2012;344:e2809–e2809.
4. Altman DG. Missing outcomes in randomized trials: addressing the dilemma. *Open Med* 2009;3:e51–3.
5. Fleming TR. Addressing missing data in clinical trials. *Ann Intern Med* 2011;154:113–7.
6. Bower P, Brueton V, Gamble C, et al. Interventions to improve recruitment and retention in clinical trials: a survey and workshop to assess current practice and future priorities. *Trials* 2014;15:399.
7. Coakley M, Fadiran EO, Parrish LJ, Griffith RA, Weiss E, Carter C. Dialogues on diversifying clinical trials: successful strategies for engaging women and minorities in clinical trials. *J Womens Health (Larchmt)* 2012;21:713–6.
8. Zhang T, Tsang W, Wijeyesundera HC, Ko DT. Reporting and representation of ethnic minorities in cardiovascular trials: a systematic review. *Am Heart J* 2013;166:52–7.
9. Paskett ED, Reeves KW, McLaughlin JM, et al. Recruitment of minority and underserved populations in the United States: the Centers for Population Health and Health Disparities experience. *Contemp Clin Trials* 2008;29:847–61.
10. Davis AM, Vinci LM, Okwuosa TM, Chase AR, Huang ES. Cardiovascular health disparities: a systematic review of health care interventions. *Med Care Res Rev* 2007;64:29S–100S.

11. Pilote L, Dasgupta K, Guru V, et al. A comprehensive view of sex-specific issues related to cardiovascular disease. *CMAJ* 2007;176:S1–44.
12. Harris DJ, Douglas PS. Enrollment of women in cardiovascular clinical trials funded by the National Heart, Lung, and Blood Institute. *N Engl J Med* 2000;343:475–80.
13. Ghare M, Chandrasekhar J, Mehran R, et al. Sex disparities in cardiovascular device evaluations: strategies for recruitment and retention of female patients in clinical device trials. *JACC Cardiovasc Interv* 2019;12:301–8.
14. National Institutes of Health. Inclusion of women and minorities as participants in research involving human subjects. Available from: https://grants.nih.gov/grants/funding/women_min/women_min.htm.
15. Warner ET, Glasgow RE, Emmons KM, et al. Recruitment and retention of participants in a pragmatic randomized intervention trial at three community health clinics: results and lessons learned. *BMC Public Health* 2013;13:192.
16. Senturia YD, McNiff Mortimer K, Baker D, et al. Successful techniques for retention of study participants in an inner-city population. *Control Clin Trials* 1998;19:544–54.
17. Nicholson LM, Schwirian PM, Klein EG, et al. Recruitment and retention strategies in longitudinal clinical studies with low-income populations. *Contemp Clin Trials* 2011;32:353–62.
18. Yancey AK, Ortega AN, Kumanyika SK. Effective recruitment and retention of minority research participants. *Annu Rev Public Health* 2006;27:1–28.
19. Levsky JM, Travin MI, Spevack DM, et al. Rationale and design of a randomized controlled trial comparing stress myocardial perfusion imaging with coronary CT angiography as the initial imaging study for intermediate-risk patients admitted with chest pain. *J Cardiovasc Comput Tomogr* 2009;3:264–71.
20. Levsky JM, Spevack DM, Travin MI, et al. Coronary computed tomography versus radionuclide myocardial perfusion imaging in chest pain patients admitted to telemetry: a randomized, controlled trial. *Ann Intern Med* 2015;163:174–83.
21. Bazemore A, Petterson S, Peterson L, et al. Higher primary care physician continuity is associated with lower costs and hospitalizations. *Ann Fam Med* 2018;16:492–7.
22. Gadegbeku CA, Stillman PK, Huffman MD, Jackson JS, Kusek JW, Jamerson KA. Factors associated with enrollment of african americans into a clinical trial: results from the african american study of kidney disease and hypertension. *Contemp Clin Trials* 2008;29:837–42.
23. Janson SL, Alloto ME, Boushey HA; Asthma Clinical Trials Network. Attrition and retention of ethnically diverse subjects in a multicenter randomized controlled research trial. *Control Clin Trials* 2001;22:236S–43S.
24. Bertakis KD, Azari R, Helms L, Callahan E, Robbins J. Gender differences in the utilization of health care services. *J Fam Pract* 2000;49:147–52.
25. Nabil-Khorazaty MN, Johnson AA, Kiely M, et al. Recruitment and retention of low-income minority women in a behavioral intervention to reduce smoking, depression, and intimate partner violence during pregnancy. *BMC Public Health* 2007;7:233.
26. Kearney A, Daykin A, Shaw ARG, et al. Identifying research priorities for effective retention strategies in clinical trials. *Trials* 2017;18:406.
27. Glassman AB, Stockdale CR, et al. Effect of telephone calls from a centralized coordinating center on participant retention in a randomized clinical trial. *Clin Trials*. In press.
28. Brunson D, Biesty L, Brocklehurst P, et al. What are the most important unanswered research questions in trial retention? A James Lind Alliance Priority Setting Partnership: the PRioRiTty II (Prioritising Retention in Randomised Trials) study. *Trials* 2019;20:593.