

William T. Grant Foundation

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Consultation and Software Tools

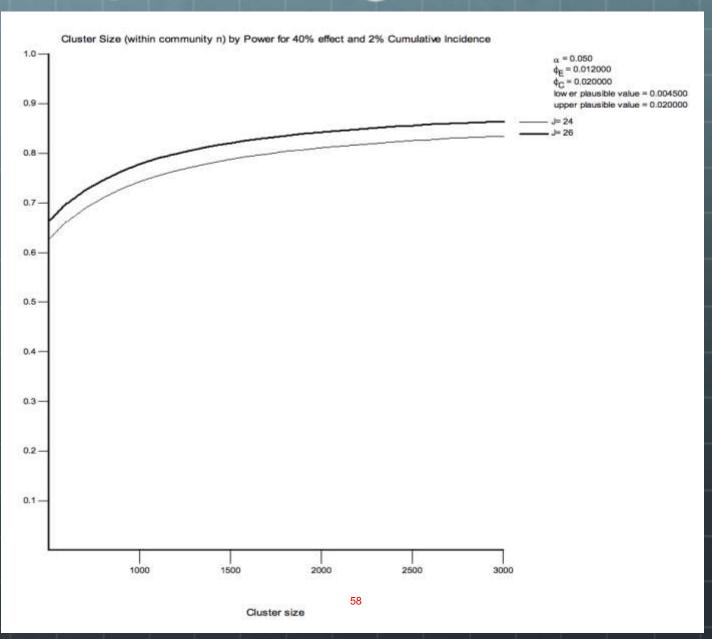
We support a consultation service and software to aid in the design of group-randomized studies. The consultation service can also be used for help with assessing the reliability and validity of measures of social settings.

- Consultation Service
- Optimal Design Software

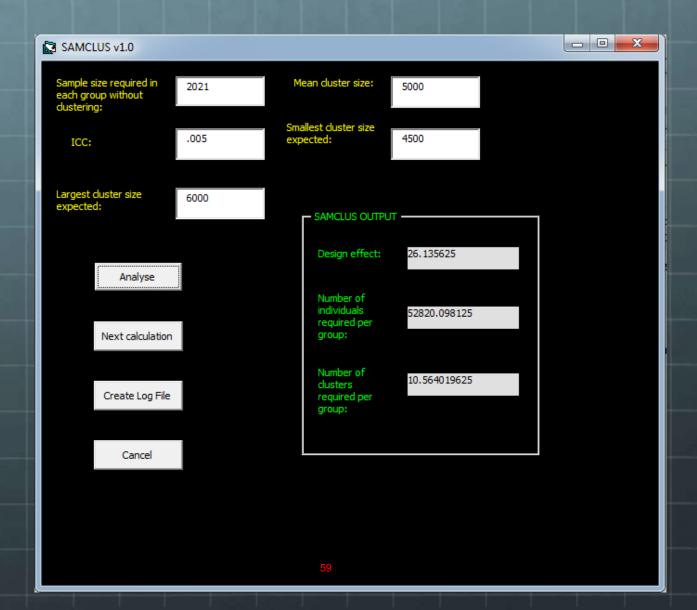
Consultation on Design of Group-Randomized Studies

The William T. Grant Foundation, in collaboration with Stephen Raudenbush and his colleagues, supports a free consultation service. The team encourages researchers, funders, and policymakers to send questions about the design of group-randomized studies of interventions designed to promote the development of youth between ages 8-25, and/or the assessment of reliability and validity of measures of social settings. Designs not involving randomization of groups or measures of social settings are beyond the scope of this service.

Optimal Design Software



SamClus Software



SampClus in Stata

```
0 0
                         ampclus.txt
. sampsi .02 .01, power(.8) onesided
Estimated sample size for two-sample comparison of proportions
Test Ho: p1 = p2, where p1 is the proportion in population 1
                   and p2 is the proportion in population 2
Assumptions:
        alpha = 0.0500  (one-sided)
         power = 0.8000
           p1 = 0.0200
           p2 = 0.0100
        n2/n1 = 1.00
Estimated required sample sizes:
           n1 =
                    2021
                    2021
           n2 =
sampclus, obsclus(5000) rho(.005)
Sample Size Adjusted for Cluster Design
  n1 (uncorrected) = 2021
  n2 (uncorrected) = 2021
   Intraclass correlation
                             = .005
  Average obs. per cluster = 5000
   Minimum number of clusters = 22
  Estimated sample size per group:
     n1 (corrected) = 52536
```

60

n2 (corrected) = 52536

Hayes Equations in R

```
[Workspace restored from /Users/echarlebois/.RData]
> get_clusters_matched_proportions <- function(p0, p1, k, m, za=1.96, zb=0.84) { 2 + (za + zb)^2 * (((p0*(1-p0)/m) + (p1*(1-p1)/m) + k^2 * (p0^2 + p1^2))/ (p0 - p1)^2) }
> get_clusters_matched_rates <- function(l0, l1, k, y, za=1.96, zb=0.84) { 2 + (za + zb)^2 * (((l0 + l1)/y) + k^2 * (l0^2 + l1^2) / (l0 -l1)^2) }
> get_clusters_unmatched_proportions <- function(p0, p1, k0, k1, m, za=1.96, zb=0.84) { 1 + (za + zb)^2 * (((p0*(1-p0)/m) + (p1*(1-p1)/m) + (k0^2 * p0^2 + k1^2 * p1^2))/ (p0 -
> get_clusters_unmatched_rates <- function(l0, l1, k0, k1, y, za=1.96, zb=0.84) { 1 + (za + zb)^2 * (((l0 + l1)/y) + (k0^2 * l0^2 + k1^2 * l1^2) / (l0 -l1)^2) }
> get_clusters_matched_proportions(.02,.01,.5,1000)
[1] 14.1128
kseq<- seq(0,.55,by=.05)</p>
> i<-0
> cluster<-1:11
> for (thisk in kseq) {
   cluster[i]<-get_clusters_matched_proportions(.02,.01,thisk,1000)</pre>
   i<-i+1
kseq
[1] 0.00 0.05 0.10 0.15 0.20 0.25 0.30 0.35 0.40 0.45 0.50 0.55
[1] 4.4108 4.7048 5.1948 5.8808 6.7628 7.8408 9.1148 10.5848 12.2508 14.1128 16.1708
```

[R.app GUI 1.31 (5537) i386-apple-darwin9.8.0]

Matched Pair Cluster Randomized Trial Cluster Sample Sizes				
Total Sites	Cluster Size @ 80% Power	Coefficient of Variation Km	HIV Incidence in Intervention Arm	HIV Incidence in Control Arm
10	4.4108	0	0.01	0.02
10	4.7048	0.05	0.01	0.02
12	5.1948	0.1	0.01	0.02
12	5.8808	0.15	0.01	0.02
14	6.7628	0.2	0.01	0.02
16	7.8408	0.25	0.01	0.02
20	9.1148	0.3	0.01	0.02
22	10.5848	0.35	0.01	0.02
26	12.2508	0.4	0.01	0.02
30	14.1128	0.45	0.01	0.02
34	16.1708	0.5	0.01	0.02
8	3.6268	0	0.015	0.03
8	3.9208	0.05	0.015	0.03
10	4.4108	0.1	0.015	0.03
12	5.0968	0.15	0.015	0.03
12	5.9788	0.2	0.015	0.03
16	7.0568	0.25	0.015	0.03
18	8.3308	0.3	0.015	0.03
20	9.8008	0.35	0.015	0.03
24	11.4668	0.4	0.015	0.03
28	13.3288	0.45	0.015	0.03
32	15.3868	0.5	0.015	0.03
14	6.7628	0	0.005	0.01
16	7.0568	0.05	0.005	0.01
16	7.5468	0.1	0.005	0.01
18	8.2328	0.15	0.005	0.01
20	9.1148	0.2	0.005	0.01
22	10.1928	0.25	0.005	0.01
24	11.4668	0.3	0.005	0.01
26	12.9368	0.35	0.005	0.01
30	14.6028	0.4	0.005	0.01
34	16.4648	6 245	0.005	0.01
38	18.5228	0.5	0.005	0.01

Simulation

- Using simulation to predict likely ranges of clustering effects
- Varying cluster effects, contamination, unmeasured external influences

Learning Objectives

Hopefully you now have knowledge of:

- a basic understanding of cluster randomized trials,
- what are the big issues in designing and analyzing CRTs, and
- where do you go to learn more and get the software tools you need to do sample size planning for CRTs.

Discussions

- Your experiences with Cluster Randomized Trials
- Other resources and experts
- Clarifications
- Getting copies of the resource items and locations

