

## Reviewing Grants

### Rationale

Hypothesis clearly stated with relevant comparison condition and testable  
Likelihood hypothesis is correct  
Both positive and negative outcomes of interest  
Congruence with prior studies  
Should more basic questions be answered first  
Lit review – focused, comprehensive citation (including in press, presentations),  
balanced review vs argue one's own point, adequate interpretation of results  
Alternate explanations consider  
Is the hole you are filling important. If so, why?

### Significance

Magnitude of problem  
Novelty  
Increase understanding- effect on theories about behavior  
Improve public health/clinical outcomes  
Magnitude of impact on above

### Experience

Prior productivity  
Content expertise  
Pilot work  
Used methods previously

### Participants

How selected/recruited  
Generalizability: Are groups over/underrepresented?  
Inclusion criteria too strict/lenient  
What is group of interest?  
S expectancies  
Flowchart of S attrition (see Consort)  
Will Ss comply  
Ability to recruit  
Compensation is adequate  
Sample size adequacy – use estimated effect size, base rates usually smaller  
than you think

### Design

What is control group or base rates  
Use multiple control/comparison groups?  
Think factorially  
Prefer to do within-Ss  
What is the adequacy of test – esp if obtain negative results

How does test map onto hypothesis – are constructs well-operationalized  
Control/comparison group will do better than you think  
Possible confounds/moderators, interactions needed to show effect  
Dismantle in later studies  
Time, order, testing effects  
Biases if not random assignment  
Groups differ on variables other than variable of interest  
Regression to the mean  
Control groups: placebo, no drug, historical, multiple baseline, usual care, optimal care, standard care  
Use positive control group; e.g. another tx  
What are historical, epidemiological base rates  
Pure vs stratified randomization  
Will diffusion, demoralization, occur – See Campbell book  
Randomize 2:1 if one group of more interest?  
Anticipated dropout rate- reasons for dropouts  
Baseline stability and noise, will practice or order or carryover effects occur  
Usually there is initial noise in behavior change- focus on later data?  
Is intervention well-operationalized  
How handle missing contacts  
Debriefing of participants. Ask questions to help interpret results

#### Interventions

Adequate dose, duration and timing of intervention  
Adequate training of therapists  
Practicality  
Both conditions same emphasis/quality  
Monitor compliance  
Tx those in control group after study  
Blindness maintained

#### Measures

Designated major outcome  
If multiple outcomes are they expected to be convergent and, if so, how handle inconsistent results across outcomes  
How well operationalized  
Actual behavior>self-report of behavior>subjective reports>intentions, attributions  
What is not being measured  
Proxy vs real measures  
Process/mechanism measures  
Blinding of assessments  
Add pre and post qualitative interviews  
In what ways do measures map onto concepts  
Test-retest and interrater reliability, predictive validity, sensitivity  
Stability of outcome  
Use measures hypothesized not to change as specificity test

What is likely effect of demand bias and political correctness  
Use challenge or eliciting tests  
Measure functional status  
Are questions ambiguous

### Results

A priori criterion for success  
Stats tied to hypotheses  
Magnitude of effect- clinical significance  
Outliers  
Type of scale – continuous, ordinal, categorical, nominal  
Distribution for stats  
Covariates included  
Examine raw data  
Avoid transformations when possible  
Do not chop up continuous to ordinal/categorical unless clinically meaningful  
Is nominal made to seem ordinal  
Sufficient variability for correlational analyses  
Linear vs curvilinear vs threshold effects  
Dose-responsivity  
Adverse events – clinically significant, cause dropouts  
Amount of overlap of groups  
Intent-to-tx, when designated as S

### Interpretation

Causality – within subjects crossover > parallel groups RCT >prospective  
prediction > cross sectional association  
Relate to others work  
Is it actually a conceptual replication  
Limitations of generalizability, qualifiers needed  
Was it conservative or liberal test  
Impact on field  
Alternate explanations  
Most conservative interpretation  
Significance for explaining vs intervening

### Human Ss

Safety, especially for subpopulations  
Confidentiality  
Ethics of control group  
Subject concerns  
Legal issues  
Steps to minimize risks  
Alternate treatments  
How handle emergencies, break code  
Consistent with usual care

