

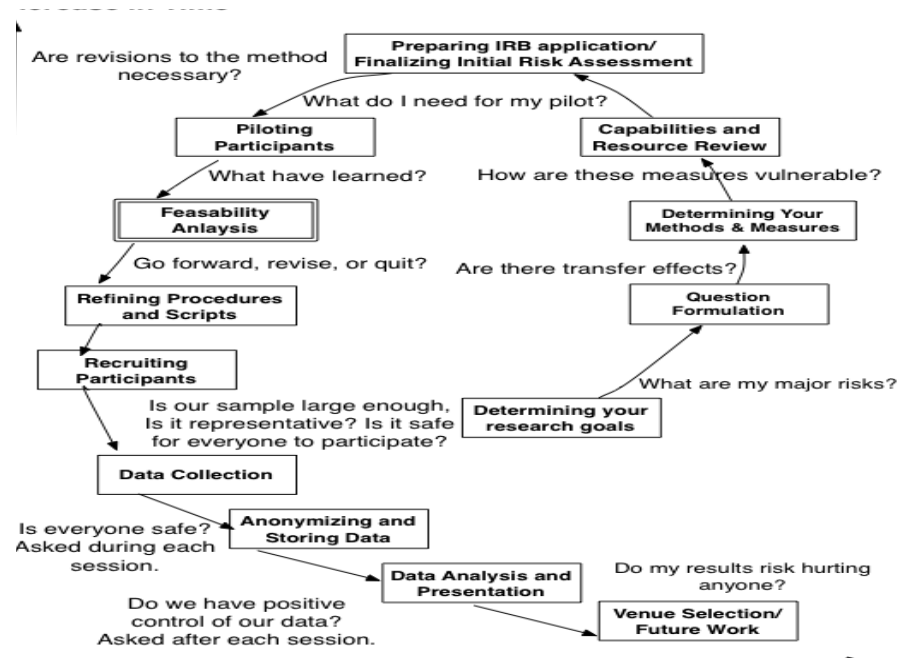
A Risk Driven Approach to Experimental Design and Practice

Frank E. Ritter and Jonathan H. Morgan (slides)

The College of IST
Penn State
and

Jong W. Kim and Richard Carlson (book)

Psychology, U. of Central Florida, and Psychology, Penn State



Overview (TB, p. 3)



acs.ist.psu.edu/papers

1400-1415 (0) Orientation

1415-1445 (1) An overview of risk-driven experimental design

1445-1515 (2) Preparation for running an experiment

1515-1540 break

1540-1615 (3) Ethical challenges in the experimental process

1615-1645 (4) Risks to validity, with class participation

1645-1700 Slack

1700-1715 (5) Conducting an experiment

1715-1730 (6) Concluding a study and reporting results

What to get out of this Tutorial

1) Some feeling for how to run a study

- Cognitive science may be modeling + data
So, to use data you have to know how it was gathered
- Modeling is slow, so data publication helps modelers
- If you are a computer scientist, you won't have taste in this area
=> Help you develop a green thumb
- Not how to *design* a study, but related

2) Some tools to help you set up a study

3) Materials

Draft book on this topic (please let me know if you use it for a class)

Handout

Example problems

4) A break at ~1515 pm (local)

5) A greater appreciation for mistakes to avoid and a theory of how to avoid them

Who are you?

- 1) Name, organization, background, number of studies, what you want to get from this
- 2) Please form into pairs for later exercise

Ch 1. Overview

Some Terms used

A study, varying an Independent variable (IV, e.g., amount of practice), to see the effect on a dependent variable (DV)

Worth reading a methods book(s)

Subjects (Ss) or Participants (Ps), Users, learners, students, Experimenters (Es)

See APA manual and also Roediger (2006) for arguments for S and P and U/L/S

Example studies

Multi-lingual fonts

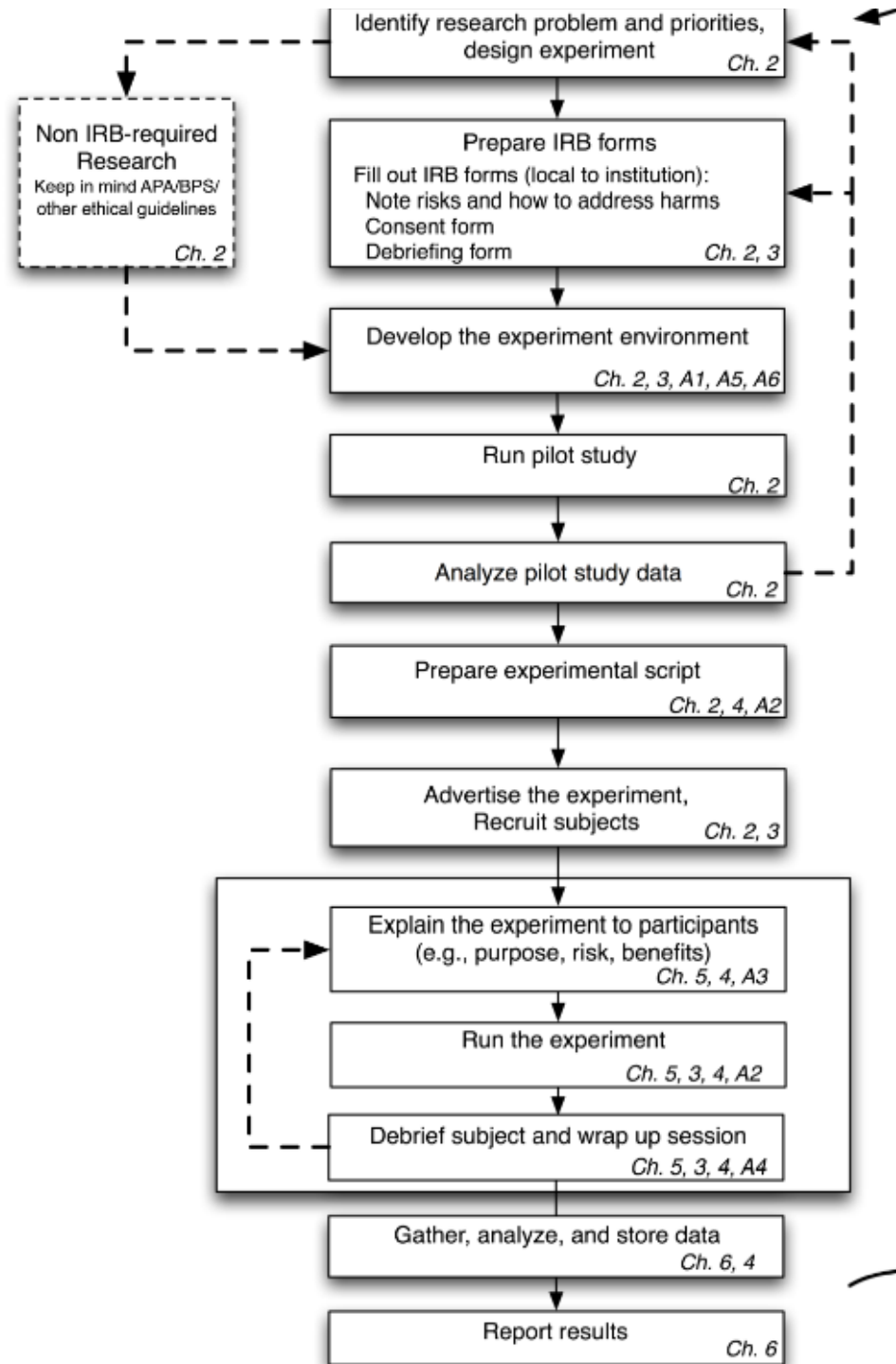
Partially sighted and blind users

HRI

Experimental Process Overview, linear

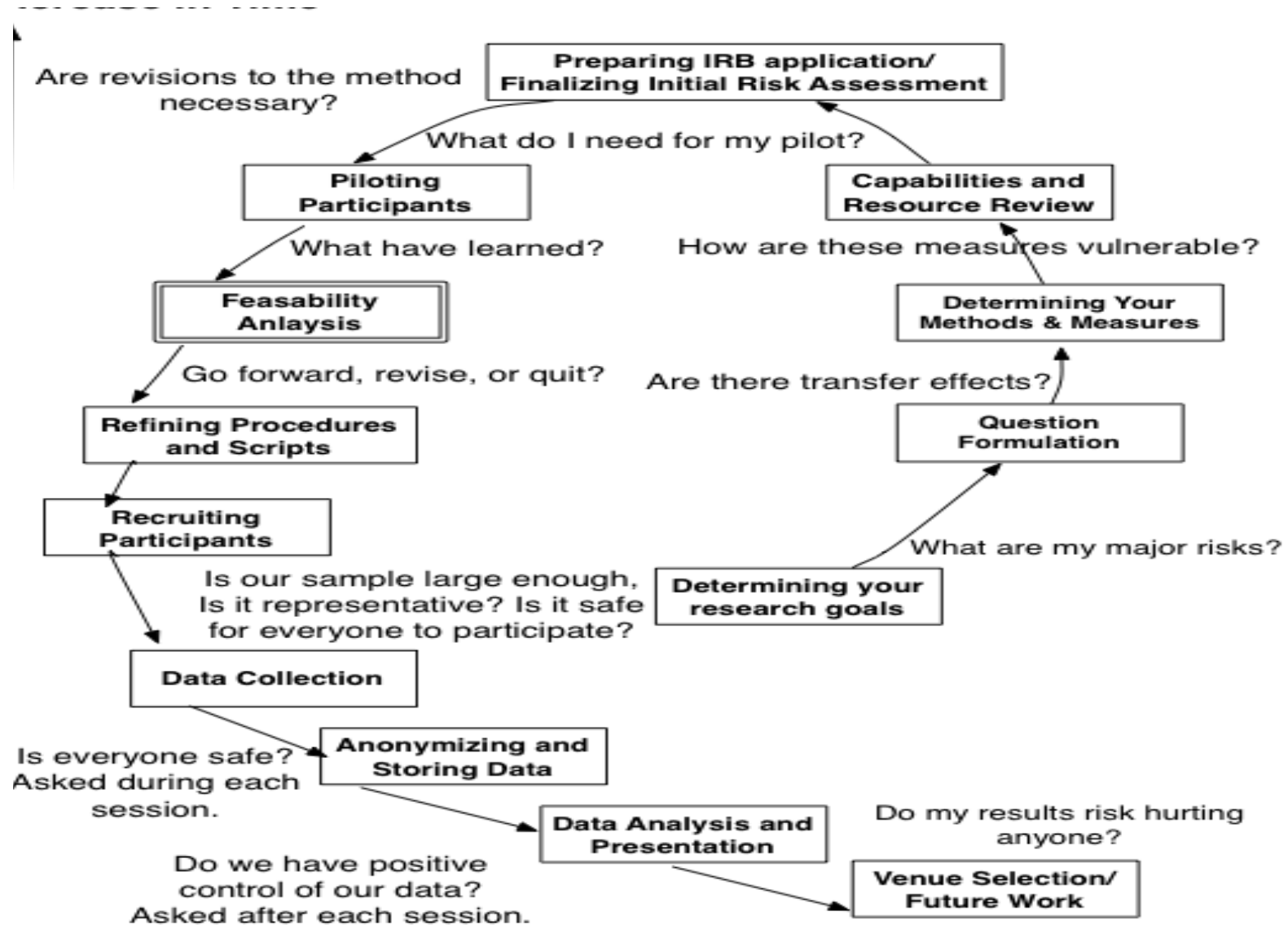
(TB, p. 11)

An iterative,
and often over-
lapping process



Experimental Process Overview

Risk Driven, more spiral (TB, p. 4)

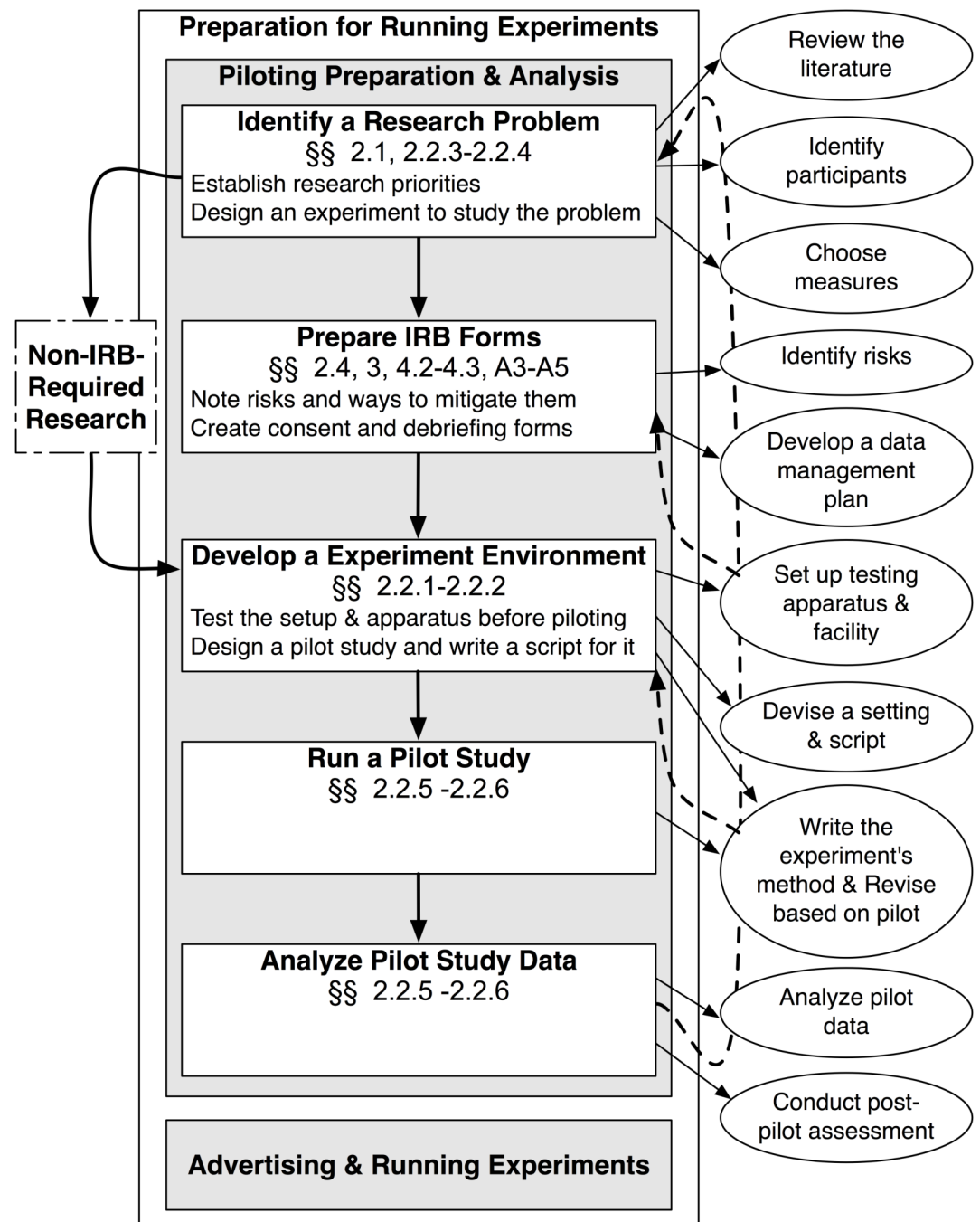


Summary: Lessons so Far

- More steps than I thought
- Iterative and risk-driven (if you pay attention)
- A process but not a set process
- Studies will overlap each other and inspire each other
- It is useful to have the RAs/Es pay attention
 - Ss suddenly 'get it'
 - Ss don't get some aspect
 - Ss comments
 - Ss 'cheat' somehow

Preparation for an experiment (TB, p. 14)

Experiments are driven by their questions and shaped by the methods available to explore those questions and existing results/lessons in that area



What studies need IRB?

■ In the US

- if not publishing no IRB (but, be careful), includes class projects
- If only authors are Ss, no IRB
- If only published / publicly available data, no IRB but IRB has to ok this (!)
- Else, IRB
- Blood, sexual history, etc. are high-risk,=> full IRB

■ Outside US

- Depends, UK used to do IRB only on high-risk studies
- Can you tell me?

■ In all cases, worth having someone check your work

IRB Forms

- Used to check your work
- May be worth being clear and concise
- Also check with example forms for language
- Draft for the PI

Summary: Piloting

- Write out method
- Used to check your work
- Use a script,
Step 1, start program, Step 2 “Welcome to...”
- Start local, e.g., YOU, and then officemate, and then move further and further away
- Mount a scratch monkey
- Check your apparatus and data gathering and use of data
- Consider/reconsider, number of Ss to run
 - Previous studies
 - Power analyses (Cohen for Ss; Ritter et al. for models)
 - Why not prefer large effects?

Ethical Challenges Associated with the Experimental Process

Ethical problems can be decreased by deliberate proactive action.

A couple of bad examples and then a general view

Assessing & Addressing Ethical Risks

Sources of Risk

Recruiting Participants

§§ 3.2, 2.3-2.4

Issues regarding equal access to the study
Issues regarding compensation

Conducting Studies

§§ 3.4, 3.5, 3.11

Location risks
Task related risks/coercion of participants

Sensitive Data

§§ 3.6, 6.1

Identifying information or data misuse
Data loss

Plagiarism & Fraud

§§ 3.6-3.7, 6.3

Formal and informal misattribution
Fraud in response to pressure or data loss

Conflicts of Interest

§§ 3.9

Sponsor or institutional conflicts of interest
Local conflicts of interest

Authorship and data ownership

§§ 3.10, 6.4

Conflicts over authorship credit
Conflicts over data ownership

Understand your sample population

Ensure fair compensation & access

Describe the task sufficiently but no more to participants

Perform a risk assessment & address risks point-by-point

Enact and follow a data management plan

Know: what is plagiarism or fraud, & what is a contribution

Place yourself to succeed

Address potential conflicts of interest in your risk strategy

Communicate with your colleagues often and early

The Monster Study: Wendell Johnson's Stuttering Study (1939)



AP

- Evaluated the effect of external valuations on stuttering
 - interrupting vs. non-interrupting conditions
- Studied 22 orphans ranging in age from 5-15 years old, grouping them into 5 fluency categories
- Resulted in long-term developmental and psychological harm, with \$925,000 awarded to six of the participants in 2007
- Avoid manipulations that can harm people

Jesse Gelsinger (1981-1999)



- Included in a bio-medical intervention study to replace a missing participant despite testing positive for high ammonia levels
- The informed consent agreement failed to disclose either known adverse drug effects or the death of two monkeys in animal trials.
- A profound conflict-of-interest existed
- Avoid conflict of interests
- Cases like this give rise to the need for IRBs

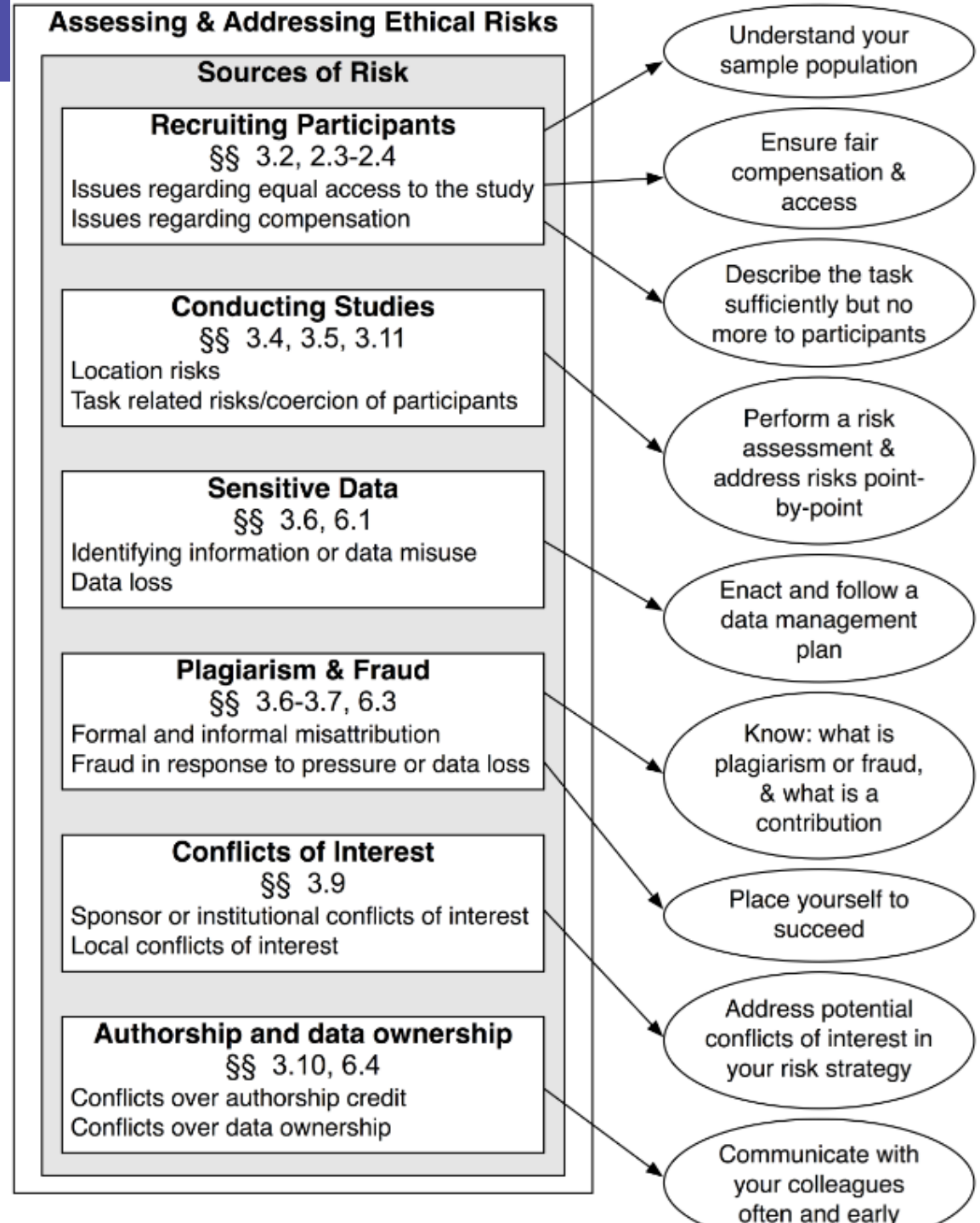
A HCI Study Gone Wrong (circa 2008)



- No informed consent
- No privacy grantees or data management plan
- “You have no friends.”
Yes, a student researcher felt compelled to inform a participant and the S’s teachers and Dean of this fact.

Ethical Challenges Associated with the Experimental Process

Ethical problems can be decreased by deliberate proactive action.



Summary:

How to avoid ethical problems

- Recruit fairly
- Look out for your Ss
- Anonymise data at the beginning of each session by using subject IDs, not names
- Have a plan for surprising data (e.g., high BP)
- Communicate early and relatively often about publication plans and data ownership
- Some argue that you have an obligation to use the data you gather

Challenges to Validity: Constraints on your study

Or: alternative
hypothesis for
results (TB, p. 21)

Challenges to
validity can be
anticipated and
mitigated.

Assessing & Addressing Risks to Validity

Risks to Internal Validity

Insufficient Sample/Effect Size
§§ 4.2.1 & 2.7

Experimenter & Participant Effects
§§ 4.2.2-4.2.3

Demand Characteristics
§ 4.2.4

Equipment and Setup Effects
§ 4.2.5

Failures to Perform the Task
§ 4.2.6

Risks to External Validity

Poor Task Fidelity
§ 4.3.1

Poor Sample Representativeness
§ 4.3.2

Use rules-of-thumb

Perform power
analysis

Make & observe
protocols

Use a double-blind
design if applicable

Randomize when
possible

Keep the setup the
same

Practice directing
participants

Ensure
psychological fidelity

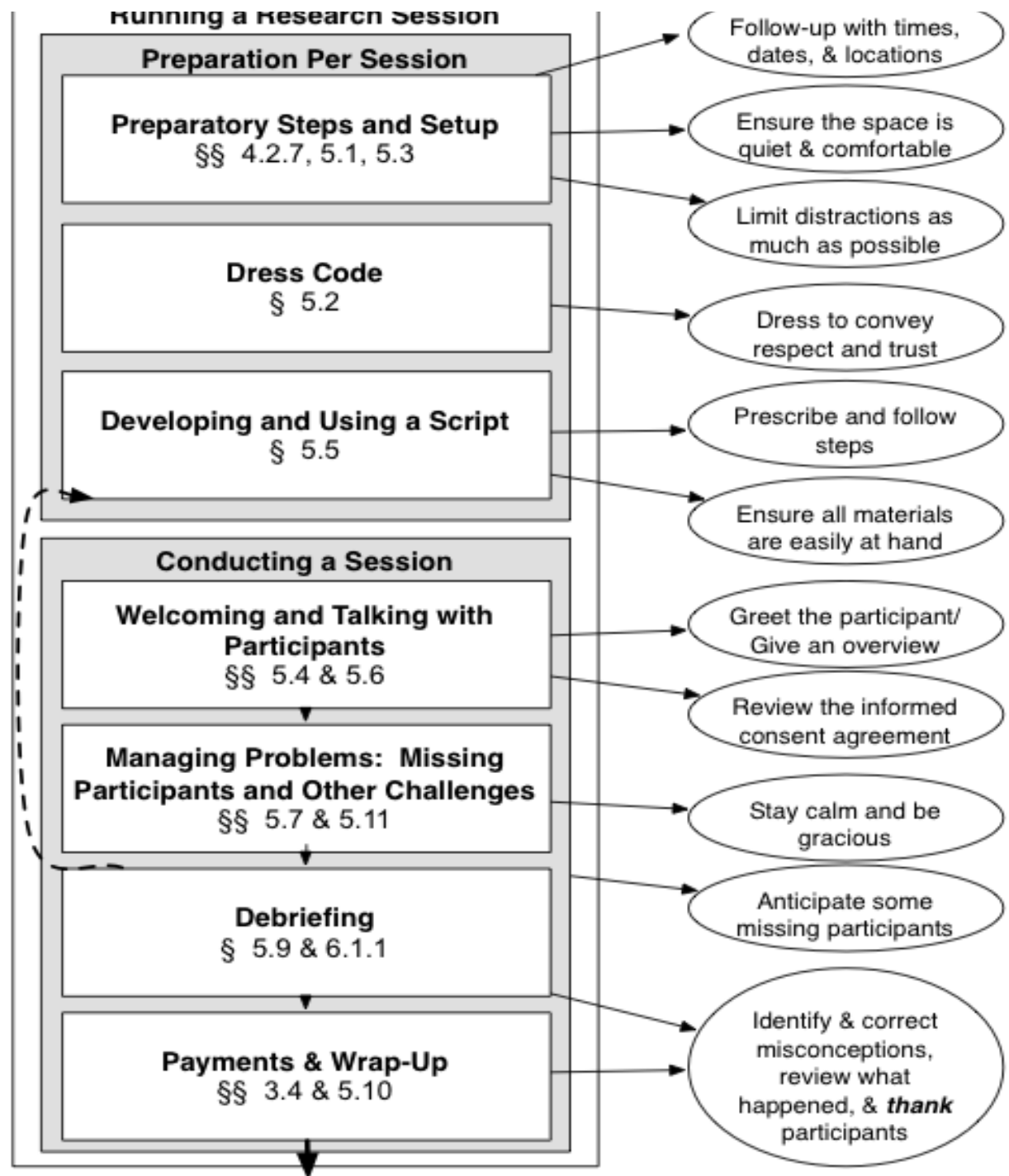
Establish what is
relevant to the study
population

Avoid inadvertent
selection bias

Conducting an Experiment

(TB, p. 24-25)

Success in
execution is
directly correlated
to careful
preparation



Summary: Running a session

- Use of piloting means no surprises (except for the data!)
- Script keeps treatment the same, it includes session set up
- Keep eyes open while running for further insights
- Anonymise data as soon as possible

Concluding an Experiment and Reporting Your Results

(TB, p.27)

Debrief, debrief, debrief!

Concluding a Study and Relaying Results

Data Care & Backup
§§ 6.1 & 3.6

Analyzing Data & Reporting Results

Documenting Data Analyses
§§ 6.2.1 & 2.2.4

Using Descriptive & Inferential Statistics
§ 6.2.2

Planned vs. Exploratory Data Analysis
§ 6.2.3

Displaying Data
§ 6.2.4

Communicating Your Results
§§ 6.3

Keep raw data as a backup

Record all data transformations

Try numerous measures

Think about what you are aggregating

Don't be afraid to do additional analyses

Explore graphing your data

Consider your writing outlet

Summary: Concluding an Experiment and Reporting Your Results

■ Concluding a session

- Finish with the subject (thank, debrief, check paperwork)
- Check the data was collected and saved
- Comment on the data if anomalies

■ Data care, security and privacy

- Anonymizing removes nearly all IDs

■ Back up data (daily, weekly)

■ Data analysis

- Not how, but note how (document and keep track of)
- Know your data if you are the RA that analyses
- Save the analyses, time is not important, space is not important, the insights and results are important
- Aside: we prefer regression
- Aside: we prefer individual analysis

Ch 6.5 Communicating your results

- Start with a target in mind
(if you can)
- Work to larger publications
(workshop, conf, journal, book)
- Rewrite, rewrite, rewrite
(the book is draft 49 [mar12], now 53)

Exercise: setting up space [iff time]

- (a) Describe your space with your partner for your next study
- (b) Does it match the description pp. 32-33?
- (c) How could you improve it?
- (d) Should you improve it?

Ch. 7 Afterward

- Appropriate behavior with subjects
- Insights
- Repeatability
- Reportability

Summary 1 of tutorial:

Relooking at failure: What constitutes a failure?

- Someone got hurt.
- After committing significant resources, the study was never completed.
- We have learned nothing new because our data is not repeatable or generalizable.
- We have failed to communicate our results or their significance to anyone else.

Sources of Failure?

- Why did someone get hurt?
 - *We failed to do a risk assessment.*
 - *Being prepared for unanticipated problems.*
 - *We failed to screen participants properly.*
 - *We failed to either develop or follow procedures, either experimental procedures or data management procedures.*
 - *We did not anticipate or mitigate situational risks either in our experimental setting or outside of it that hurt our participants.*
 - *We ignored additional insights we could have learned from the participants through observation or debriefing.*
 - *Others?*

Sources of Failure?

■ Why we were unable to complete the study?

- *We were overly ambitious, perhaps because we failed to fit the research question or methods to the problem at hand.*
- *We ran out of time.*
- *We ran out of resources or lacked them in the first place.*
- *We lacked the people, either participants or staff, or trained staff.*

(experiments appear to have less risk than modeling)

Sources of Failure?

- Why we were unable to reproduce our results or generalize them?
 - *We failed to use the same experimental procedures or test under the same conditions for each S.*
 - *We failed to achieve an adequate sample size or sufficient degree of representativeness in our sample.*
 - *Our task fidelity was poor. We failed to construct an experimental task that was analogous with respect to its key points.*

Sources of Failure?

- Why have we been unable to report our results or communicate their significance?
 - *We failed to properly catalog or backup our data.*
 - *We failed to write as we went. We no longer remember some of the critical early details.*
 - *We made poor data analysis or display choices.*
 - *We failed to identify a venue early, or understand who we should consider our audience.*

How do we avoid failure?

- We recognize that running a study is an incremental risk-driven process, similar in some respects to spiral development (Boehm & Hansen, 2001; Pew & Mavor, 2007).
- To be successful, we need to:
 - Formulate a research question that meets our research goals
 - Have a theory of transfer effects that minimizes risks associated with confounding variables, and enables us to conserve time and resources.
 - Pilot studies and study components
 - Be candid in our risk assessments and be willing to adapt and refine.

Summary 2 of Tutorial

- There are steps to running a study separate from design and analysis
- These are practical, hands-on, implicit knowledge
- They are informed by previous studies
- To be successful, we need to:
 - Formulate a research question that meets our research goals
 - Pilot studies and study components
 - Be candid in our risk assessments and be willing to adapt and refine
 - Be aware of alternative hypotheses, and avoid what we can and control what we cannot avoid
 - Plan for reporting results early





If you will teach this....

- Full book available shortly from Sage
- Slides available as ppt or pdf
- Workbook available as pdf

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acs.ist.psu.edu/papers

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