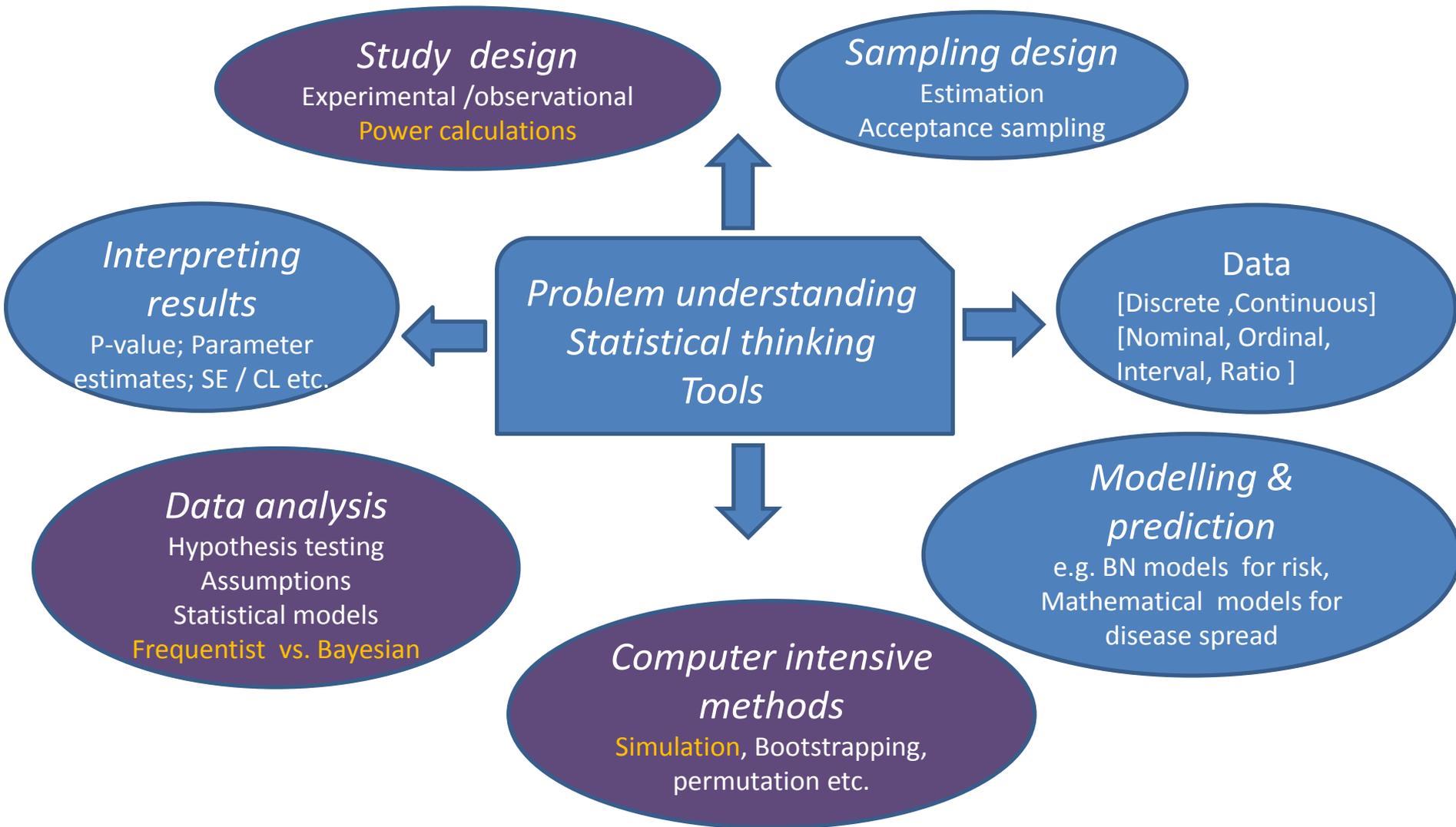


Mind map outline



Power Analysis

What is Statistical Power?

- Statistical Power is the probability of rejecting the Null Hypothesis if it is false

e.g. The probability of getting a statistically significant treatment effect in your experiment if a true treatment effect exists.

- Formally Power = $1 - \beta$ where β is the type II error.

- It depends on a number of factors: α –level, magnitude of the variation from the relevant sources, sample size, experimental design, and statistical test used.

Power Analysis

Why is power important?

- A prospective power analysis helps to decide the appropriate sample size for a trial.
- For many clinical trials a power of 80% or 90% is often required for ethical approval
- Retrospective or Post-hoc power analyses are less useful (and more controversial).

Power Analysis

How to calculate power

- Analytically

e.g. For the difference in proportions with equal sized samples and $\alpha=0.05$:

$$Z_{1-\beta} = \sqrt{\frac{n(p_1 - p_2)}{2\bar{p}(1-\bar{p})}} - 1.96$$

- Most common statistical software packages include a power calculation module. There are also specialised programs available for the more complicated designs e.g. G*Power, Russ Lenth's Power and Sample size Java applet.

Power Analysis

How to calculate power

- Through simulation e.g.

```
res.prop.test<-rep(0,1000)
n<-110
for(i in 1:1000){
  p1<-rbinom(1,n,.4)
  p2<-rbinom(1,n,.6)
  res.prop.test[i]<-prop.test(c(p1,p2),c(n,n))$p.value
}
sum(res.prop.test<0.05)/1000
[1] 0.818
```

(Analytical power=0.813)

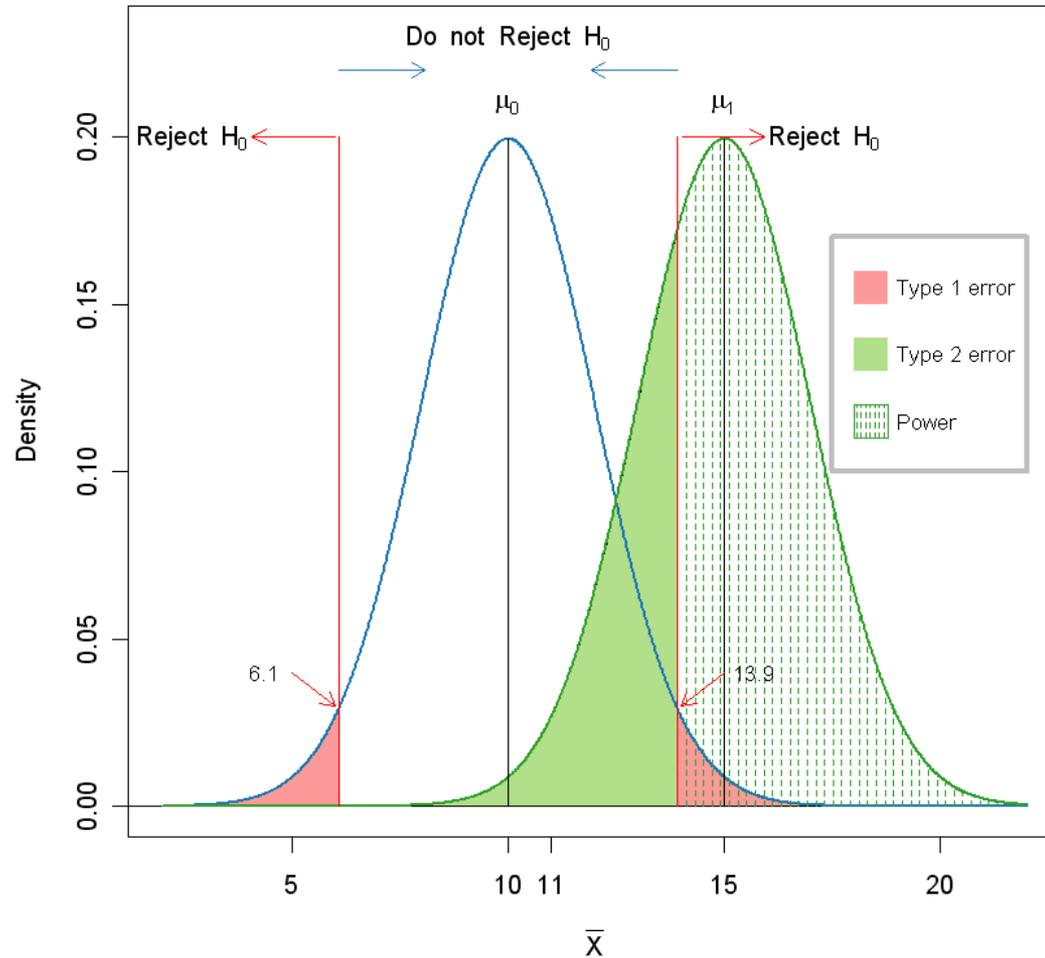
- Can be very useful when the design or model is non standard and there is no analytical method easily available.

Power Analysis

Factors that influence power:

- The lower the α -level the lower the power.
- Power increases with sample size.
- The larger the minimum detectable difference the greater the power.
- The experimental design and model can influence the power
e.g. Paired or two sample t-test

Power \uparrow as the difference to be detected \uparrow



Power Analysis

<http://homepage.stat.uiowa.edu/~rlenth/Power/>



Java applets for power and sample size

Select the analysis to be used in your study:

- CI for one proportion
- Test of one proportion
- Test comparing two proportions
- CI for one mean
- One-sample t test (or paired t)
- Two-sample t test (pooled or Satterthwaite)
- Linear regression
- Balanced ANOVA (any model)**
- Two variances (F test)
- R-square (multiple correlation)
- Generic chi-square test
- Generic Poisson test
- Online tables of common distributions
- Pilot study

Run selection

Note: These require a web browser capable of running Java applets. See www.oracle.com/technetwork/java/. Due to a compatibility bug, many plug-ins size the applet window.

Please read this comment

I receive quite a few questions that start with something like this:
"I'm not much of a stats person, but I tried [details...] -- am I doing it right?"
Please compare this with:
"I don't know much about heart surgery, but my wife is suffering from ... and I plan to operate ... can you advise me?"
Folks, just because you can plug numbers into a program doesn't change the fact that if you don't know what you're doing, you're almost guaranteed to get meaningless results -- if not dangerously misleading ones. Statistics really is like rocket science; it isn't easy, even to us who have studied it for a long time. Anybody who think it's easy surely lacks a deep enough knowledge to understand why it isn't! If your scientific integrity matters, and statistics is a mystery to you, then you need expert help. Find a statistician in your company or at a nearby university, and talk to her face-to-face if possible. It may well cost money. It's worth it.

If you're blocked by a security setting

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Discussion group

Please read this comment

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Discussion group

Bayesian Statistics

- Frequentist hypothesis tests use the p-value as evidence against the Null hypothesis. It does not always mean that the alternative hypothesis is more likely.

An example: Suppose we have a test for PSA-X with sensitivity of 100% and specificity 99.9%. If we take a single sample at random and get a positive test result, then what is the probability that PSA-X is present?

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Answer: We need more information! We need to know the probability of PSA-X occurring.

- If $\Pr(\text{PSA-X})=0.5$ then

$$\Pr(\text{PSAX} | T^+) = \frac{\Pr(T^+ | \text{PSAX}) \times \Pr(\text{PSAX})}{\Pr(T^+)} = \frac{1 \times 0.5}{1 \times 0.5 + 0.001 \times 0.5} = 0.999$$

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- However if $\Pr(\text{PSA-X})=1/1001$ then

$$\Pr(\text{PSA-X} | T^+) = \frac{1 \times \frac{1}{1001}}{1 \times \frac{1}{1001} + 0.001 \times \frac{1000}{1001}} = \frac{\frac{1}{1001}}{\frac{1}{1001} + \frac{1}{1001}} = 0.5$$

Bayesian Statistics

- Frequentists base their findings on the data and do not incorporate prior knowledge.
- In the previous example the probability of a positive test if PSA-X is not present is 0.1% and so a standard Hypothesis test would reject the Null.
- Bayesian statistics aims to use prior knowledge in addition to the data to make inferences about the unknown parameters using Bayes' Theorem

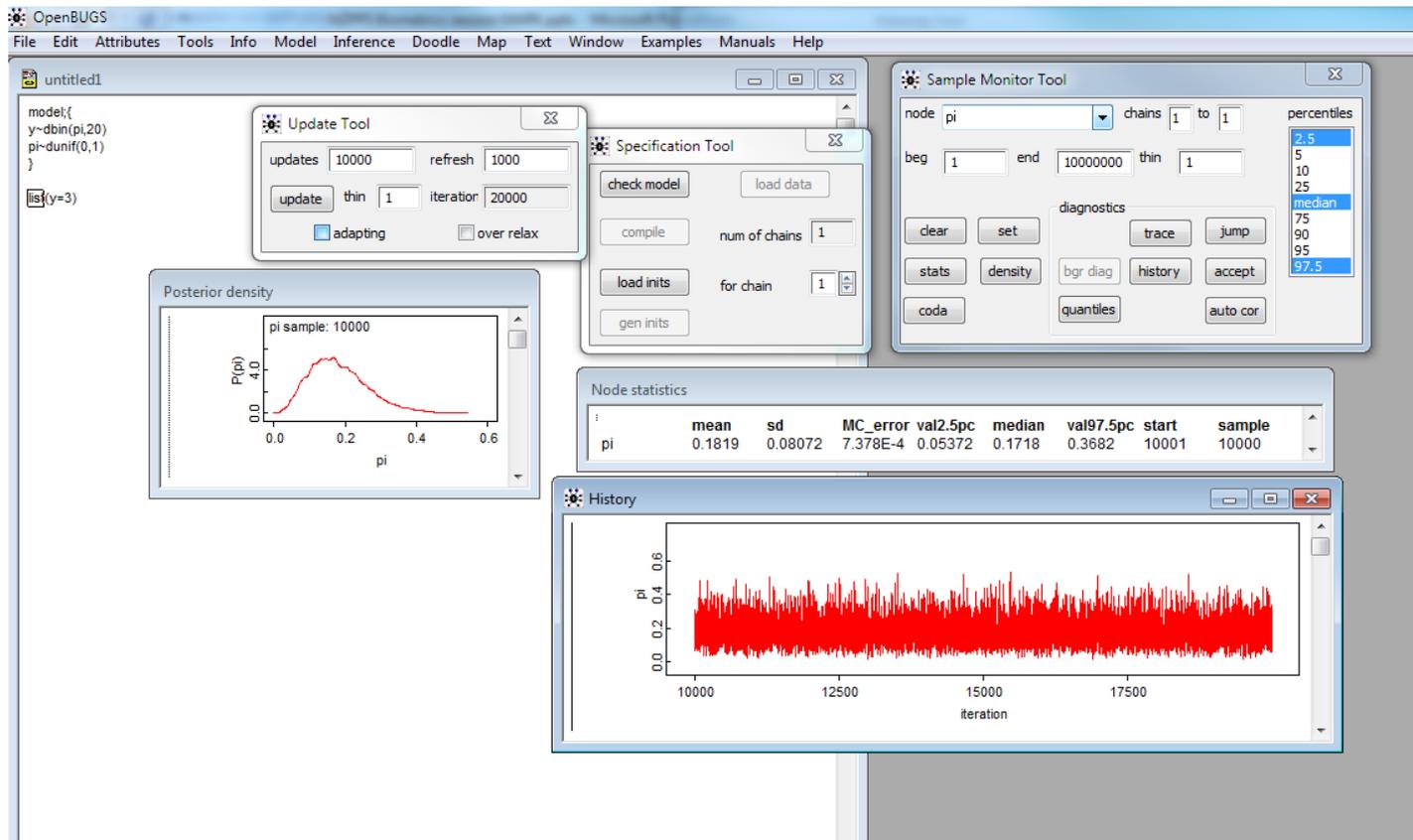
$$\Pr(\theta | data) = \frac{\Pr(data | \theta) \times \Pr(\theta)}{\int \Pr(data | \theta) \times \Pr(\theta) d\theta}$$

- Often the integral is intractable and so MCMC methods are used to sample from the posterior distribution using the relationship

$$Posterior \propto likelihood \times prior$$

Bayesian Statistics

- Once the model is set up a large number of observations are sampled from the posterior distribution.
- Since the earlier samples may be influenced by the starting values it is common practice to ignore the first batch (burn-in) of initial values.
- Base inferences on the posterior sample e.g. Median, mean, credible intervals...



Bayesian Statistics

Criticisms

- Using prior knowledge introduces a subjective element to the analysis. Flat priors and sensitivity analyses are some methods used to address these concerns.
- With large samples the data will dominate and the Bayesian results should be similar to the frequentist equivalent.
- Can be overly complex and difficult to get the model to converge. More advanced algorithms have helped in this area.

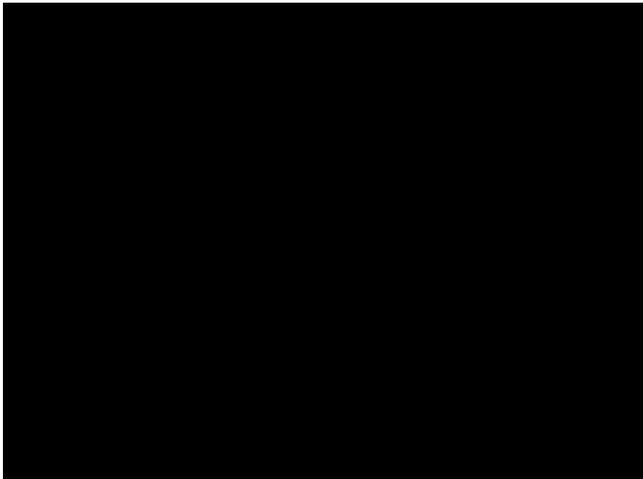
Advantages

- Can produce more intuitive results e.g. Credible intervals vs. Confidence intervals, posterior predictive distribution.
- Doesn't rely on large sample approximations
- Forces you to think about the model and the sources of variation

Bayesian Statistics

<http://twiecki.github.io/blog/2014/01/02/visualizing-mcmc/>

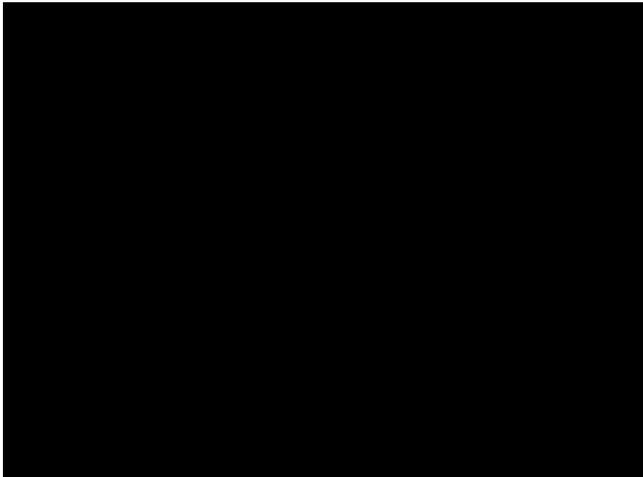
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Step Method

- 1) Metropolis-Hastings Algorithm
- 2) Gibbs sampling
- 3) NUTS (Hamiltonian Monte Carlo)

2



3

