Introduction to Bayesian A/B testing

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Agenda

- Setting the scene
- Priors and probability distributions
- Running a Bayesian A/B test
- Posterior analysis
- Summary and some final remarks
- Thank you!
- When to stop a Bayesian A/B test?
- Extras
- Game of chances



Setting the scene



Randomised Control Trials (RCTs)

A simplistic example:

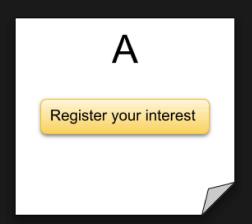
- Users are assigned at random to two groups, A and B, with equal probability.
- Let A be our control group and B be our treatment group.

We want to know what effect our treatment has.



Hypothetical scenario

- A button on a landing page that takes users to a sign up form.
- At present, the button is labelled "Register your interest".
- Test whether changing it to "Get started" will result in an increased click-through rate (CTR).
- "Get started" was suggested by an experienced and skilled UX designer.







Priors and probability distributions

The key to speeding up your experiment



Prior knowledge and beliefs

Before running an experiment, we form opinions and gather evidence such as:

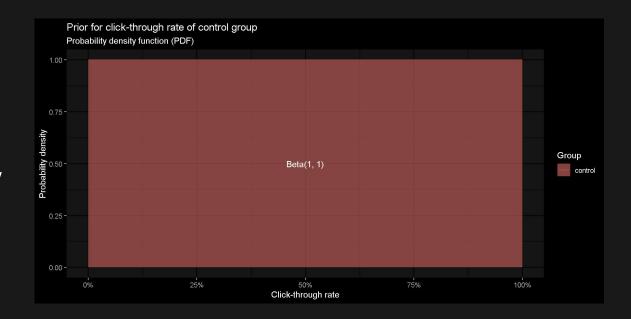
- The baseline click-through rate of the button (with its current label) and knowledge of any outside variables that affects click-through rate, e.g. seasonality
- Effects we have seen from similar previous experiments
- Qualitative research, such as usability tests, focus groups, and surveys that are related to the test
- Opinions (including critical) from stakeholders and experts



Priors are probability distributions

Express prior beliefs about the click-through rate of the control group using a **probability distribution**.

Here's an example of an extremely uninformative prior – a uniform prior that says any range of click-through rate is as probable as any other equally wide range, i.e. naive.

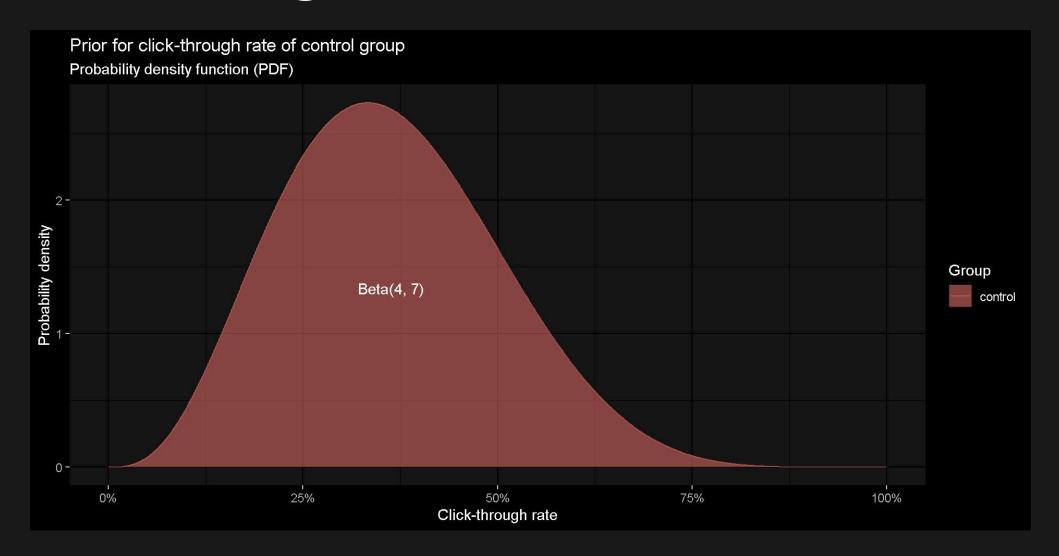




The **Beta distribution** is a **probability density function (PDF)** with two **shape parameters**: B(shape1, shape2). It's used to describe proportions, such as click-through rate.



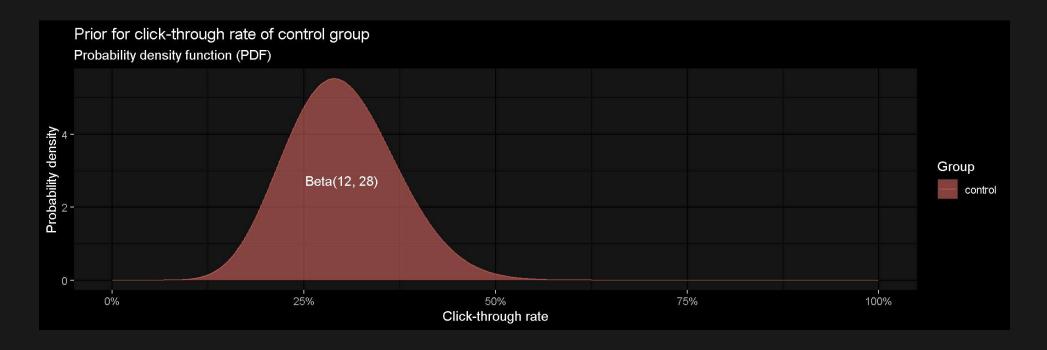
Something a little more informative



As the curve narrows, notice that the shape parameters of the Beta distribution increase.



Something even more informative

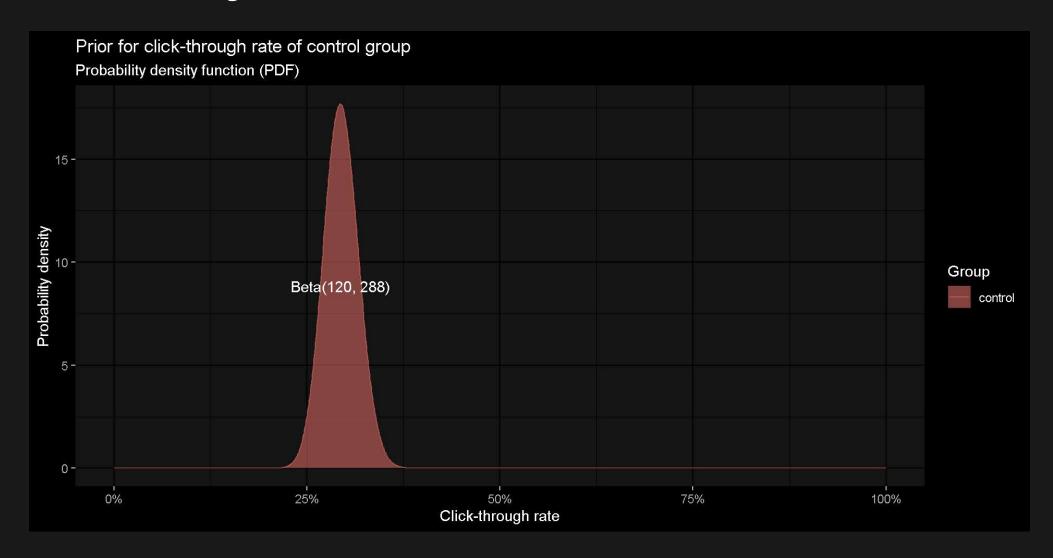


The shape parameters (shape1 and shape2) of the Beta distribution can be considered counts of successes and failures, respectively. The mean probability of success (i.e. average click-through rate) can be calculated by this formula:

$$\frac{shape1}{shape1+shape2}$$



Let's say we've settled on this:



The more confident we are about our beliefs, the narrower the curve.

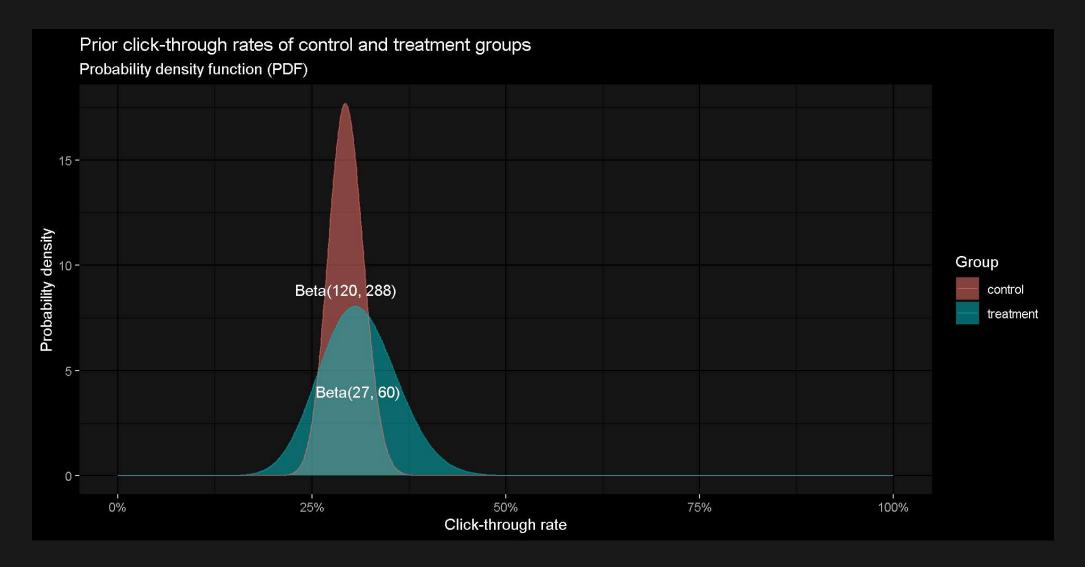


What about the treatment group?

- We expect that the click-through rates of the treatment and control groups will be correlated.
- We're unsure about how correlated they will be, but we're not expecting a dramatic difference.
- We're more confident than not that the treatment will be an improvement, but we're open to other possibilities.
- We don't want to bias the experiment results in favour of treatment or control, or towards a conclusion of there being a difference or no difference.



We've settled on these priors:





Running a Bayesian A/B test



Prior agreement

- Agreement must be reached on the priors before collecting and analysing data from the experiment.
- Once the priors are agreed to and locked in, we can start the experiment.

Here's a summary of the priors we have chosen:

group	shape1	shape2	mean	sd
control	120	288	29.4%	2.3 p.p.
treatment	27	60	31.0%	4.9 p.p.



Let's run an experiment

We'll generate some fake data to mimic a real experiment.

It'll be rigged though, as we'll already know the click-through rates for control and treatment, which are:

• Control: 32%

• Treatment: 35%

That's a relative uplift of 9.4%.

If we're successful at applying Bayesian inference then we should hope (but can't guarantee due to randomness) that the results somewhat match with these expected CTRs.



The next day

Let's pretend that on average 150 users enter our experiment each day, and we've received the following data from day 1:

group	total_users	clicked	not_clicked	CTR
control	86	32	54	37.2%
treatment	81	26	55	32.1%



Let's now incorporate our priors

Posteriors represent your updated beliefs once you've incorporated experiment data with your priors. Like priors, posteriors represent your beliefs about the metric of interest, which in our case is click-through rate.

For each experiment group, we derive our posterior shape parameters through simple arithmetic addition:

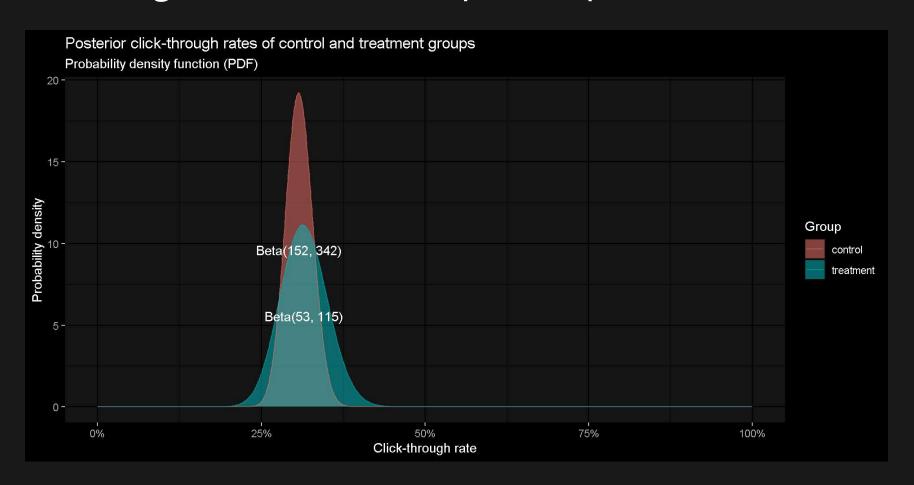
- Increment the first shape parameter by the count users who had clicked
- Increment the second shape parameter by the count users who **didn't click**

count	control	treatment
prior_shape1	120	27
clicked	32	26
posterior_shape1	152	53
count	control	treatment
count prior_shape2	control 288	treatment 60



Posterior distribution of each group

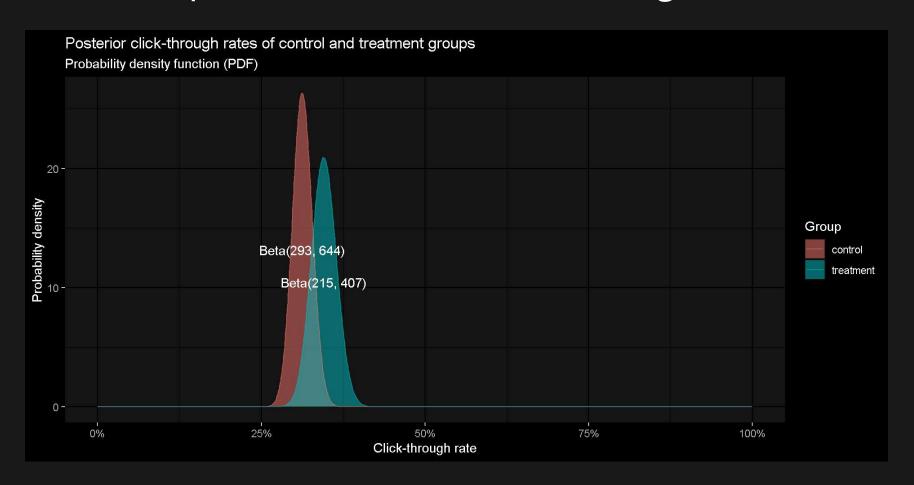
We have now updated our beliefs. These posteriors can now be thought of as our new updated priors.





Another six days later...

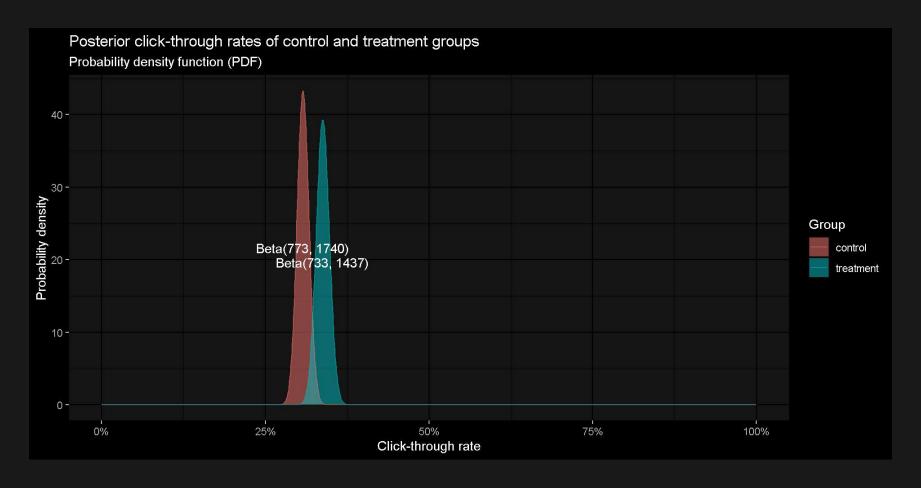
We've collected more data, so let's again update our priors to form new posteriors for the click-through rates of each group.





Another three weeks later...

We've now observed a total sample of 4,188 users and a decision is made to end the experiment.





Posterior analysis

Statistical inferences using the posterior distributions



Monte Carlo simulation

Let's draw a very large quantity of random samples from our posterior distributions to make inferences about the experiment.

This is called Monte Carlo simulation – named after a casino.



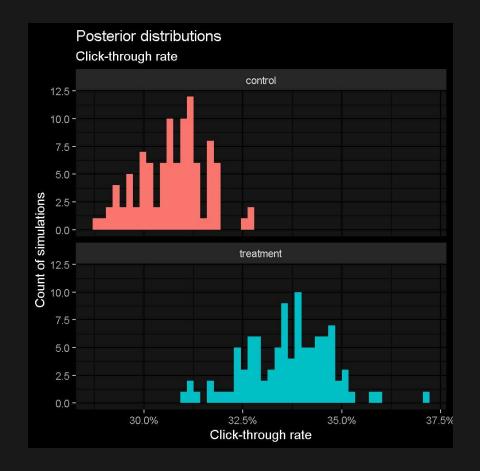


100 simulations

Let's start slowly by drawing 100 random samples from our distributions and plot them using histograms...

Here's some of our Monte Carlo samples:

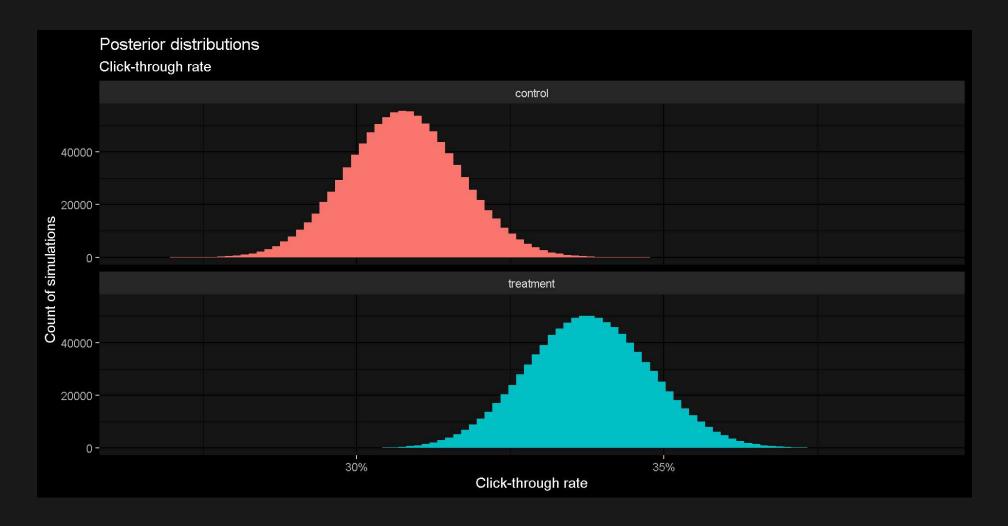
control	treatment	uplift	beats_control
0.316	0.340	0.074	TRUE
0.317	0.358	0.131	TRUE
0.310	0.347	0.119	TRUE
0.294	0.328	0.117	TRUE
0.318	0.341	0.075	TRUE
0.301	0.347	0.154	TRUE
0.309	0.346	0.118	TRUE





Let's now beef it up a bit...

We'll now draw 1,000,000 samples...





We can now make some inferences

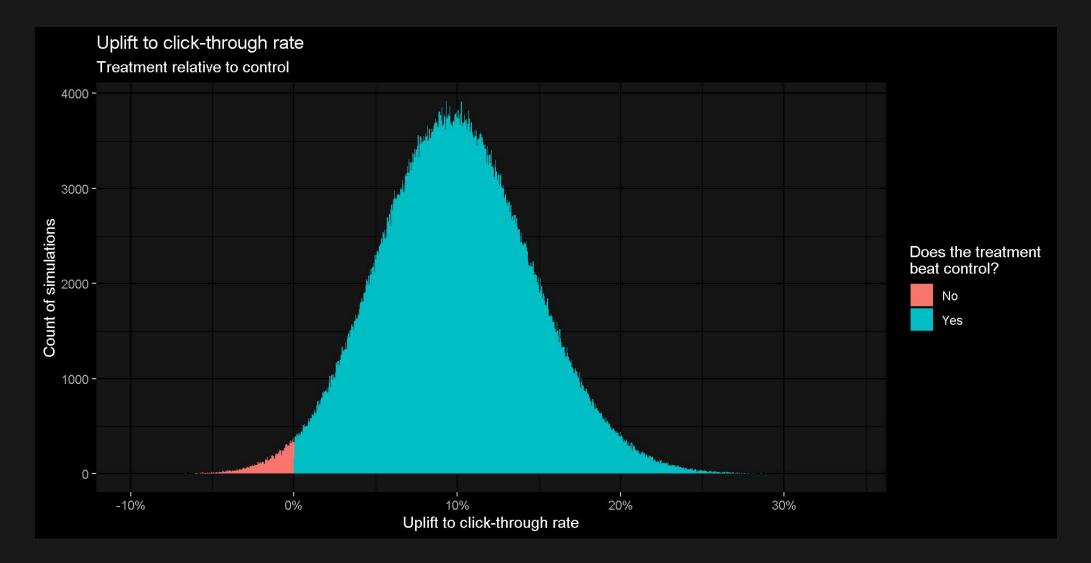
A summary of our 1,000,000 posterior samples for click-through rate:

```
control
          treatment
                                   uplift
                                                 beats control
                      :0.2902
                               Min.
                                                 Mode : logical
Min.
      :0.2656
               Min.
                                      :-0.09909
1st Ou.:0.3014
              1st Ou.:0.3309
                               1st Ou.: 0.06710
                                                 FALSE: 13815
Median : 0.3075
               Median :0.3377
                               Median : 0.09819
                                                 TRUE :986185
Mean :0.3076
              Mean :0.3378
                              Mean : 0.09914
3rd Qu.:0.3138
              3rd Qu.:0.3446
                             3rd Ou.: 0.12999
Max. :0.3525
               Max. :0.3924
                               Max. : 0.34028
```

- How do these compare to our theoretical CTRs of 32% for control and 35% for treatment, and uplift of 9.4%?
- What is the posterior probability that the CTR of the treatment is greater than that of control? Answer: 98.62%



Posterior distribution of the CTR uplift





Summary and some final remarks



Before starting an experiment

Gather prior knowledge and articulate beliefs:

- Establish a baseline what do you know about the control group?
- What do you expect the effect of the treatment to be? How sure are you?

Express those beliefs and knowledge as distributions - these are your priors for your control and treatment groups.



Running the experiment

- Start the experiment, gather data, and update your priors to form posteriors about the metric of interest
- Draw inferences by running a large number of Monte Carlo simulations using the posterior distributions
- Know when to end the experiment try to plan for this ahead of running the experiment



Thank you!

Further topics that might interest you:

- Bayesian Generalised Linear Models to better isolate the effect of the treatment from other predictors.
- Survival Analysis, such as Kaplan Meier, to analyse lagged conversion outcomes.
 - These slides and simulations were produced in RStudio using Quarto. Download the source code and slides at: https://github.com/jdeboer/measurecamp2022





When to stop a Bayesian A/B test?



If using uninformative priors...

If your original priors are uninformative or too weak, then you face the same risks as with frequentist experiments.

Perform **power analysis** ahead of running the experiment. This is to determine the required sample size before any inferences are made.

Before commencing the experiment, decide on:

- The minimum detectable effect size
- The accepted false positive rate
- The accepted false negative rate



If using informative priors...

If your priors are relatively informative and chosen carefully, then this can reduce the chances of false positives and negatives. But:

- Be careful not to bias the results of the experiment.
- Power analysis is still recommended in order to gauge the worse case scenario for how long the experiment might run.

Bayesian inference, with informative priors, can make it possible to end an experiment early.



If deciding to end early...

Ask yourself:

- Has the experiment run for at least a couple of cycles?
 (e.g. at least two full weeks)
- Have the results stabilised? Is there a clear winner?
- Could it be worth running longer to learn more?
- What are the risks of continuing or ending now? What if the results you see are just a fluke and are therefore misguiding you? What is the impact of making the wrong choice? What are the chances?



Extras



Bayes theorem

$$P(B \cap A) = P(A \cap B)$$

$$P(B) \times P(A|B) = P(A) \times P(B|A)$$

$$P(A|B) = rac{P(B|A)P(A)}{P(B)}$$

$$f(heta|data) = rac{f(data| heta)f(heta)}{f(data)}$$

 $Posterior \propto \mathcal{L}(\theta|data) imes prior$



Thomas Bayes - 1701 – 1761



Some useful formulas

Let α and β represent the first and second shape parameters of the Beta distribution, respectively.

The mean of this distribution is: $\mu = rac{lpha}{lpha + eta}$

The standard deviation is:
$$\sigma = \sqrt{\frac{lpha eta}{(lpha + eta)^2(lpha + eta + 1)}}$$

Through substitution and rearrangement, you can determine α and β from μ and σ .

$$v=rac{\mu(1-\mu)}{\sigma^2}-1$$
 $lpha=\mu v$ $eta=(1-\mu)v$

This way, you can determine the shape parameters based on centrality and spread.



Game of chances



To play this game:

- There is one host and at least 2 contestants.
- The host will need a uniform random number generator.



Instructions

- 1. The host secretly picks a number, Y, between 0% and 100% and writes it down.
- 2. The host will then secretly generate a random number, X, again between 0% and 100%:
 - If X is less than Y then the host will mark it as a 'success', otherwise as a 'failure', using a tally board that's visible to all contestants.
- 3. The objective of the game is for contestants to estimate Y by asking the host to perform step 2 as many times as they need (to a reasonable limit). The closer their guess is, the better. However, each contestant can only call out their guess once.
- 4. Once two contestants have called out what they believe Y is, then the game ends and the host reveals the true answer for Y.
 - The first contestant to call out their guess wins the game if they are within 5 points of Y. If not, then the contestant whose guess is closest to Y wins.



Lessons of the game

- As the number of successes and failures increases, you get closer to knowing what Y is.
- The compromise each contestant makes between speed and certainty will influence who wins.



Tip

Try a modification to the game where Y is a number related to a topic that the audience will have some prior knowledge about. This way they can incorporate their prior expectations when making a guess.

