

Figure 1: Problem 2

1 Problem 2 - Monte Carlo calculation of an arbitrarily drawn polygon

I was amused by the shape of this polygon. I generated points in the interval x = (0, 2), y = (0, 0.5) for the simple reason that its area is unity. Using the shapely package, it was simple to check whether a point were inside the polygon or not. I plotted the results, with the corresponding area in figure 1. The area (0.166) also seems appropriate upon visual inspection.

2 Problem 3 - Conditional probabilities for Slightly Evil Inc.

This section was by far the most tricky to get right.

2.1 Answers

My answers are contained in the output from my code:

Problem 3a:

Current aggregate defective rate: 0.03275

If defective, a PM is 18.32% likely to come from the facility A2 If defective, a PM is 23.66% likely to come from the facility A5 Problem 3b:

defective new_defective

A1	0.020	0.022143			
A2	0.040	0.051667			
A3	0.100	0.155000			
A4	0.035	0.038750			
A5	0.031	0.031000			
Problem 3c:					
facility	defective	new_defective			
A1	0.020	0.022037			
A2	0.040	0.059500			
A3	0.100	0.119000			
A4	0.035	0.074375			
A5	0.022	0.023800			
A6	0.092	0.180303			
A7	0.120	0.313158			
A8	0.070	0.070000			
A9	0.110	0.180303			
A10	0.020	0.297500			
) A11	0.070	0.396667			
A12	0.060	0.270455			
	A2 A3 A4 A5 Problem 3c facility A1 A2 A3 A4 A5 A6 A7 A8 A9 A10	A2 0.040 A3 0.100 A4 0.035 A5 0.031 Problem 3c: facility defective A1 0.020 A2 0.040 A3 0.100 A4 0.035 A5 0.022 A6 0.092 A7 0.120 A8 0.070 A9 0.110 A10 0.020 A11 0.070			

0.099

0.082

2.2 Explanation

A13

A14

12

13

facility

2.2.1 Problem 3a

Finding the chance a defective pacemaker is from A2 facility calls for the use of Bayes' Theorem:

0.396667 0.743750

$$P(A2|D) = \frac{P(D|A2) * P(A2)}{P(D)}$$

Applying the already found values:

$$P(A2|D) = \frac{0.04 * 0.15}{0.03275} = 18.32\%$$

Using the same formula, for all facilities, we find that:

$$P(A5|D) = \frac{0.031 * 0.25}{0.03275} = 23.66\%$$

Which is higher than all the others. It makes intuitive sense, too, as the only other candidate, facility A4, still produces fewer defective devices than

A5:

$$P(A4|D) = \frac{0.035 * 0.2}{0.03275} = 22.93\%$$

2.2.2 Problem 3b and 3c

Again, Bayes' Theorem shows itself handy.

In order to make all relative defective rates the same (20%) we need to adjust all but one of the defective rates up. (Analytically, it might make sense to increase the error rate for A5, as we know that it it the maximum.)

Doing so, I used the following formula to calculate the new defective rates from each factory:

$$P(D|A_i)_{new,j} = \frac{P(A_j)}{P(A_i)}P(D|A_i)_{old}$$

for all factories A_i , where j indicates which factory's defective rate remains constant.

At this point, I only accepted the set of new defective rates where

$$P(D|A_i)_{new} \ge P(D|A_i)_{old}, \forall A_i$$

Practically, this solution was scalable to be implemented on problem 3c as well.

A realization that would have made the solution easier I realized after finishing this, that an alternative solution would be to start with the relative defective rate (1.0/N) where N is the number of factories) and from there calculate the defective rates from there. Nonwithstanding, I didn't implement this solution, as I believe that the original solution was sufficient (and scalable) enough to work for the whole problem.

3 Problem 4 - Seawater

3.0.1 Problem 4a

Numerical and graphical results are contained in figure 2.

Sea Surface Water Temperatures 1.0 1997 KDE: Distribution of 100 λ s $\int_{-2}^{+4} P = 1.00,$ Gauss Mu 281.261 +/-12.770 $\int_{-2}^{0} P = 0.19$ 2017 KDE: Gauss Sigma 269 +/-0.8 - $\int_{-2}^{+4} P = 1.00,$ $\int_{-2}^{0} P = 0.22$ 10 0.6 5 1000 MC samples 0.4 300 250 0.2 0.0 0 1 3 4 -2 Temperature (C)

Figure 2: Problem 4. After generating one lambda, I was surprised when the value was several orders of magnitude larger than 1. I therefore generated 100 lambdas and plotted the distribution, which seems to fit nicely with a gaussian distribution. The distribution of the values as a gaussian with mean 285 and sigma 22 demonstrates that the values of lambda can be estimated to fall in the interval (241,329) with 95% confidence. We can therefore directly conclude that because our lambdas are significantly larger than one, we can not under any significance level reject the null hypothesis. The data generated from the 1997 KDE is definitely not compatible with the 2007 KDE PDF.

4 Problem 5 - Particles

4.1 Problem 5a and 5b

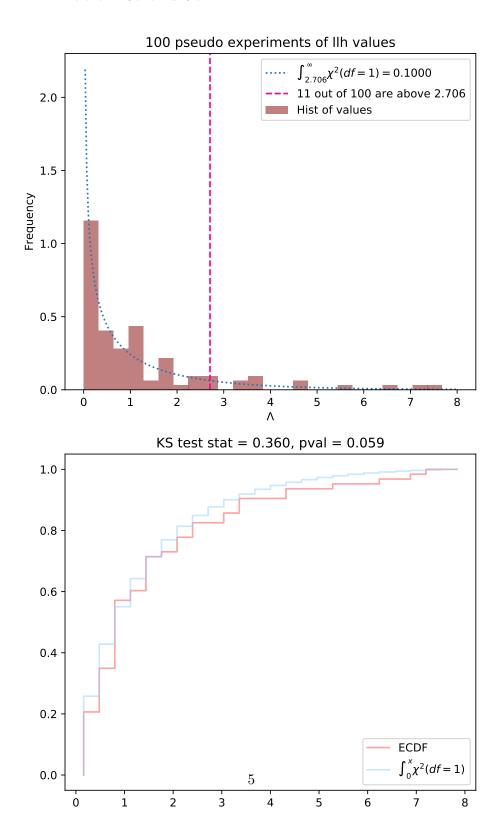


Figure 3: Problem 5

4.1.1 Problem 5b

Too see whether the distribution is actually chi squared distributed, I did a KS test as demonstrated in figure 2. Immidiately, I could see that the distributions lined up rather well. The p value close to 0.05 is also a good indicator that the values are chi square distributed, as one would expect if the samples are from the same distribution. As can also be seen in the figure, the number of pseudo-experiments with a test statistic over 2.706 is also comparable (11 observed vs 10 expected) to what one would expect from a chi square distributed test statistic.

4.2 problem 5c

Using all 20000 events for a LLH test, I get the following values for the likelihoods.

For the null hypothesis H_0 : b = 1.0, I get an LLH of $LLH_0 = -29653.08$, for the best fit parameters $\sigma_t = 0.618$, b = 1.0.

For the test hypothesis $H_1: b \neq 1.0$, I get an LLH of $LLH_1 = -29652.74$, for the best fit parameters $\sigma_t = 0.619, b = 0.993$.

Hence,

$$\Lambda = -2 * (LLH_0 - LLH_1) = 1.358$$

Which gives us a chi square integrated value

$$\int_{1.358}^{\infty} \chi^2(df = 1) = 0.244$$

Where we have 1 degree of freedom between the two models (one free parameter).

In order to reject a hypothesis at a 3 sigma threshold, the result needs to have a test statistic (chi square) value higher than 9. (p = 0.0027) As this model does not have this, we can NOT reject the null hypothesis.

I was surprised that I found such a large difference in the two approaches to finding a lambda, and tried to part 5a and 5b with RAN-DOM chunks of data, instead of splitting periodically. However, I found no significant changes between the randomly sampled chunks and the periodically split chunks. Thus, I can conclude that the different λ values are not due to a designed way of making the data chunks, and that the ordering of the points are uninmportant. The file in question is attached as p5_llhratios_random.pdf.

In order to get a bit wiser on these results, I plotted the two results in figure 4, from which we can clearly see that the two functions are substantially equivalent.

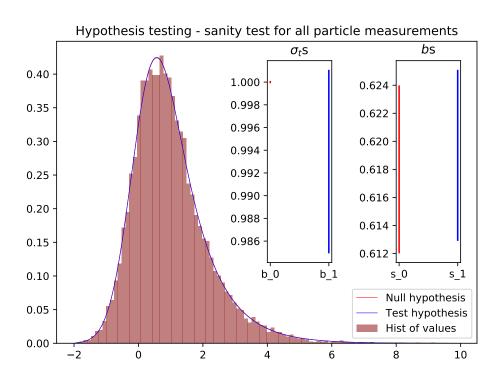


Figure 4: Visualization of the null hypothesis vs test hypothesis for problem 5c. Subplots show the 1 sigma symmetric error returned from minuit, except for b_0 , which was fixed. Clearly, the two are not distinguishable using conventional chi-by-eye techniques.