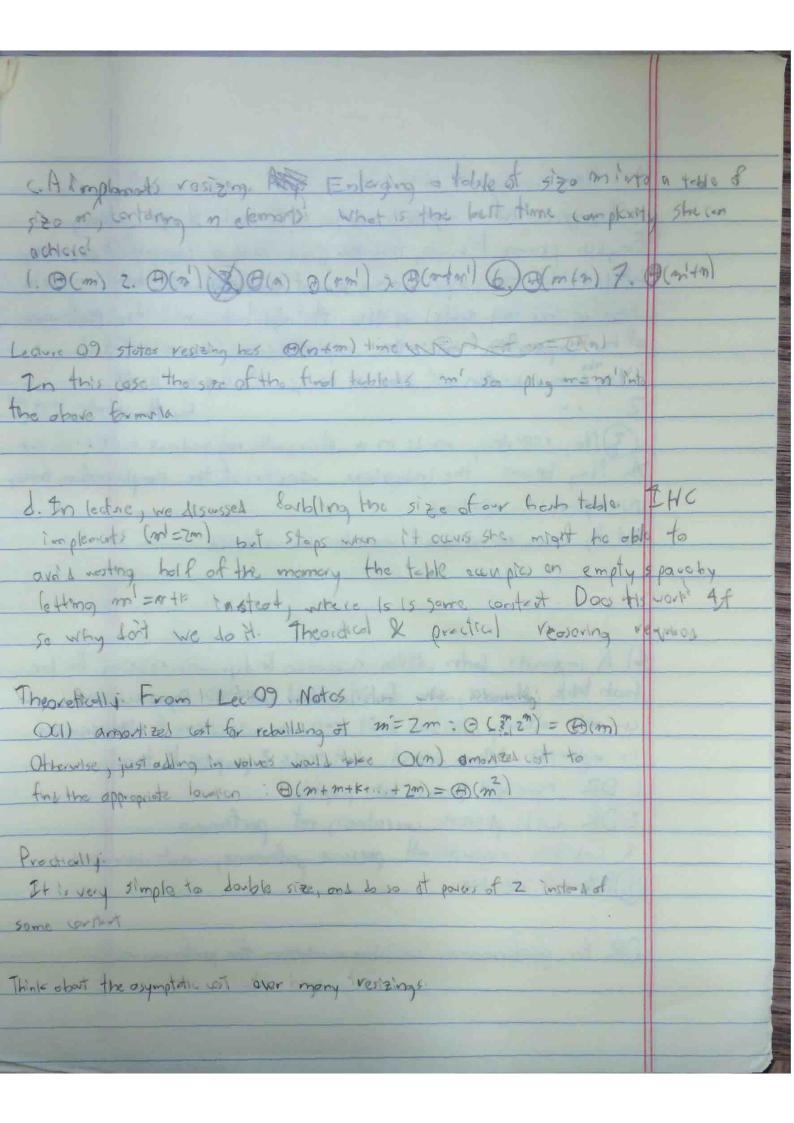
4-1 Wash functions & Low a. Imagine that an algorithm requires us to host things canding English phrase 15 mains that strong fire tared as servences of (hereto) APU doctats to simply use the rum of those character values Complibition size of box bosh table) as the strings bosh, will the performance of her implementation moter the expected volve shown in lecture Lyas sum a peraten will space out storage interely by levight by the wornders truly will (3) No, reorating words in a string will not preduce a afterest best \* No, because the inseparations consisten of the simple uniform texting USSUMPTION 18 Notated Not 4, hoshing does not depend on where other keys are hoshed. (b). A implanent's boton collister resolution & dynamic restring for har both teble. Howard, she doesn't want to age both Dynamic resizing co- word collison, collison don't wase convections issues & collision resolution is implemented when statement about trose 2 proposes is true? 1. DR alone will prosove both propose 2. DR WILL prosence correctness not performens 3. collisten resolution will presente performance, not correctues 4.) Bath are needed DR for performance, collision resolution for performance



42. v. Let's exemine the "membership testing" use uso which Adamst according desolber til Mary Inschools right ofter creation, and from mostly lookup.

Z. A maximized of every-milked inscriptions labeletis & lookups.

4. Alternating rounds of inscriptions be letters & lookups. It is described as "Creater one then verely changes" & many insertion, mostly Mary cells to has-key" => many look ups. b. Now largeline you here to prop a not function, sze, will from resolution statesy and so founts in order to make a book table pertectly suited to this use case alone. Aler tre statement that but describes are choices you might make. 1. A long min size on a growth rate of 2. 2. A small min size & growth rate of 2
3.) large min rize & growth rate of 4
4. small 6. The document recommends a law X => many slots por Key => kinge table size =) faster growth rete Mary keys inserted initially a start w/ a larger tolo

## 4-3 Matching DNA Sequences

The code and data used in this problem are available on the course website Please take a peak at the README. txt for some instructions.

Ben Bitdiblic has recently moved into the Kendall square area, which is full of biotechnology componies and their shiny, window-laden office buildings. While modeling their about himself, he is secretly their about himself, he is secretly Jealons, and so he sets out to earn one of his very own. To pick up the necessary geet cred, he begins experimenting with DNA-matching technologies. Ben would like to create mutants to do his bidding, and to get started, he'd like to know how closely related creatures are. If two sequences contain mostly the same subsequences in mostly the same place, then they're likely closely related; if they don't, they probably aren't.

For our purposes, we'll represent a DNA sample as a sequence of characters. These sequences are very long, so comparing subsequences of them quickly is important. We've provided code in Kfastsa. py that reads the fa files stering this data.

a. Let's start with subsequence Hashes, which returns all length-te subsequences and their hashes. (imprement your function as a generator).

See subsequence Hashes in draseq.py

b. Implement "Multidict" and verify that your work passes the simple sonity tests provided.

See class "Multidict" in drasequpy.

C. Now it's time to implement get Exact Submatches. I grave the parameter on for the time being; we'll get to that in the next part. Again, implementing this function as a generator is probably a good idea.

See get Exact Submatches in draseq. Py

d. The most significant reason why your solution is presently too slow to be useful is that you are hashing and inserting into your hash table of millions of elements, and then performing tens of millions of lookups into that hash table. Implement interval subsequence Hashes, which returns the same thing as 'subsequence Hashes' except that it hashes any one in m subsequences. Modify your implementation of get Exact submotions to have monly for sequence A.

See "interval subsequence Hashes" and "get Exact submotches" in draseq.py

e. Run comparisons between the two human samples (poternol & maternal) and between the poternol sample and each of the samples.

See composisons folder in /dist/