## CS250/EE387 - LECTURE 8 - APPLICATIONS of CONCATENATED RS CODES



## Why might we care about this?

1. Image processing and signal processing.



- · Most natural images/signals are sparse(ish) in some basis (or write some dictionary). · So if we can acquire that image/signal by just measuring linear combinations of it ond storing those, we can save time and space.
- Z. Streaming algorithms:

Consider a data stream:

 $X_1, X_2, X_3, \dots, X_t, \dots \in \text{some universe U of size } N$ 

You are interested in the frequency counts  $f_i = #times i \in U$  showed up.

But you don't want to store the vector  $f \in \mathbb{R}^n$ , especially if only a few items show up often.

Instead, keep a SKETCH  $\square$  =  $\square$  sketch.

Uf is spone (ish) When a new item anives, you can update the sketch by adding the appropriate column of  $\overline{\mathcal{P}}$ .

So this is exactly the same as synchronic decoding, except over Rinstead of F.

ANOTHER SYNTACTICALLY SIMILAR PROBLEM: GROUP TESTING.

Let 
$$B = \{0, 1\}$$
, with the operations "+" = V (aka, OR) and "\*" =  $\Lambda$  (ata AND).  

$$M \begin{cases} Pooling metrix \\ \overline{D} \in \mathcal{B} \end{cases} = \begin{bmatrix} & \overline{D} \cdot x \in \mathcal{B}^{m} = \text{"test outcomes}" \\ & \overline{D} \in \mathcal{B} \end{bmatrix} \\ & & \\$$

· Unfortunately, s << n of them are poisoned, but we don't know which.

W

- You want to decide BY TOMORROW which pots of coffee are poisoned (so that you can drink the rest).
   While using as few lab rats as passible. (so that the biology dept. doesn't notice...)
   The idea is to POOL the samples of coffee:
- n pots:

· IF a labrat drinks from ANY puisoned coffee pot, they become sick.\*\*

\* \* Not THAT sick. No animals were harmed in the making of these lecture notes.



So that's the picture we had before.

The PROBLEM is to recover x, the indicator vector of poisoned pots, and the GOAL is to minimize the number of lab rats borrowed "from the biologists.

MORE SERIOUSLY, this problem is usually motivated as follows:

- During WWII, the problem was indroduced for testing US soldiers for syphilis.

Soldiers ←→ coffee pots blood sample ←> coffee sample syphilis tests ←> lab rats

- Nowadays, for high-throughput screening.

Civilians ↔ coffeepots DNA samples ↔ coffee sample genetic tests ↔ lab rats

Tests are expensive, and not many soldiers/civilians are sick, so we'd like to use 25 few tests as possible.

So both GROUP TESTING and COMPRESSED SENSING are syntactically very similar b STNDROME DECUDING, it's just that they happen over B, Ror C, and FE, respectively.

The different algebraic and geometric structures make these problems very different. However, ideas from one are often useful in others.

Today, we'll see how RS codes can be used to make good GROUP TESTING matrices.  
DEF. A pooling matrix 
$$\overline{\Phi} \in \mathcal{B}^{m \times N}$$
 is d-disjunct if for any set  $\Lambda \in \mathbb{N}$  of sized, and  
any  $j \in \mathbb{N}$  is d-disjunct if for any set  $\Lambda \in \mathbb{N}$  of sized, and  
 $\overline{\Phi}_{ij} = 1$  and  $\overline{\Phi}_{il} = 0$   $\forall l \in \Lambda$   
Picture:  
 $i \rightarrow 0 \ 0 \ 0 \ 1$   
 $\Lambda$   $j$   
This is a good thing b/c if  $\Lambda$  we the true set of positives (aka, poisoned colleapoils),  
Then

$$i \rightarrow 0 \circ 0 1 = 0 \leftarrow i$$
  
 $\Lambda j = 0 \leftarrow i$   
 $I =$ 

which gives a j's status as not - puisoned. THM. If  $\overline{D}$  is d-disjunct, then as a pooling design it can identify up to d positive items.

Moreover, there is an algorithm that runs in time  $O(m \cdot N)$  to identify the d positives.

Pf. The algorithm is :

for each je [N]:
 if all the tests that j participates in are positive:
 l label j as positive.
 else j is not positive.
 L

Why does this work? Suppose that  $\Lambda$  is the true positive set and  $j \notin \Lambda$ . Then the def of d-disjunctness says that some test  $z \in \mathbb{E}m \ J$  which j participates in will come up negative, so the alg will label j "NOT POSITIVE."

So the goal is to come up with d-clisjunct-matrices  $\overline{\Phi} \in \mathbb{B}^{m \times N}$  so that m is as small as possible.

 BEST CONSTRUCTIONS KNOWN:
  $M = O(d^2 \log_d^2(N))$   $[Kautz - Singleton'(A] - we'll see this body (based on RS cades)]

 <math>m = O(d^2 \log(N))$  A random multix dues this - or check cut

 [Porat - Rothschild'08] for an explicit construction. (also based on coding theory).

ALGORITHMS: If m= O(d²log(N)), there's an EXPLICIT construction w/ SUBLINEAR TIME algorithm. [Ngo-Porat-Rudra 11(?)] (Also based on coding theory). We'll see some faster algs later in the course.

Today: A construction with 
$$m = O(d^2 \log^2_{d}(N))$$
.

IDEA: We'd like all these columns to be kind of far apart ,, let's use codewords?

Let 
$$C = RS_q(n,k)$$
, let  $N = q^k$ ,  $m = q \cdot n$ . Consider the matrix formed by:  
 $n = \sqrt{1 - 1} \sqrt{1 -$ 

$$i\in Ch_{2}$$

Now replace each symbol de Fg w/ a vector of length q.

$$\alpha_1 \longleftrightarrow \begin{pmatrix} 1 \\ 0 \\ \vdots \\ 0 \end{pmatrix}, \quad \alpha_2 \longleftrightarrow \begin{pmatrix} 0 \\ 1 \\ 0 \\ \vdots \\ 0 \end{pmatrix}, \dots, \quad \alpha_q \longleftrightarrow \begin{pmatrix} 0 \\ 0 \\ \vdots \\ 1 \end{pmatrix} \quad \text{where } F_q = \{ \alpha_{1,j-1} \alpha_q \}.$$

This results in a matrix W



The first column of  $\Lambda$  agrees w/c in at most n-dist(C) places: If these ones in the picture. The second column of  $\Lambda$  agrees w/c (and not w/ the first col) in  $\leq n - dist(C)$  places. If there are  $\cdot$  etc.

Altogether, there are at most  $|\Lambda| \cdot (n - dist(C))$  positions of C that are agreed with by SOME column of  $\Lambda$ .

By our guarantee on dist(C), 
$$|\Lambda|(n-dist(C)) < d(n-n(\frac{d+1}{d})) = n$$
.  
So there is at least one position that is not agreed with?

Let's instantiale this with 
$$RS_q(Ft_q, q, k)$$
, so  $n = q$ ,  $dist(C) = q-k+1$   
Setting  $dist(C) = n(\frac{d-1}{d}) + 1 = q(\frac{d-1}{d}) + 1$ , we get  $k = \lfloor b/d \rfloor$ .  
Then our matrix is:  
 $m = n_q \cdot q^2$   
 $\begin{cases} u^2 \\ u \\ u \\ u \\ v = q^k - q^{UU1} \end{cases}$   
Thus we choose  $q = \sqrt{m}$ , which implies  $\log_q(N) = \left\lfloor \frac{\sqrt{m}}{d} \right\rfloor$  ata,  $\sqrt{m} \approx d\log_q(N)$ .  
Then  $m \approx d^2 \log^2(N)$  which implies  $m = O\left(\frac{d^2 \log^2(N)}{\log^2(1)}\right)$ ,  
 $3^{SC}$  claimed.  
 $\begin{cases} Quick Note About Compressed SEMSING.$   
A very similar construction can be used to get deterministic compressed sensing medices.  
For those who know the lingo, this EXACT SAME construction is an S-RIP matrix.  
 $\overline{Q} \in [R^{n,N}]$  with  $m = O\left(3^{1}\log^2(N)\right)$ .  
And you can do slightly belier if you replace  $F_{\overline{P}}$  with up th roots of unity.  
 $\begin{bmatrix} See Cherceptch's * Cuding-Theoretic Methods for Sparse Recovery" for lots more! ]$ 

## QUESTIONS TO PONDER

- ① Can you come up with a recovery scheme for this group testing matrix that runs in time poly( dlog(N)) [in particular, sublinear in n?]
- 2) Can you make a group testing scheme using the semantic similarity to syndrome decoding? (Rather than the scheme we saw, which used a different connection to coding theory)