EXP. NUMBER	EXPERIMENT/SUBJECT	DATE
NAME	LAB PARTNER	LOCKER/DESK NO. / COURSE & SECTION NO.

8/13/12
- 9c, 11c, 8a, 9d, 1a, 5a, 11a, 12a, 12b, 12c are working (all others were bleached)
  - made new liquid cultures of each

8/14/12
- plasmid preps of all working detectors ↑

8/15/12
- PCR of 10 Detectors

dH_2O	16.5	x 11 =	181.5λ
Buffer	2.5		27.5
$MgCl_2$	1.5		16.5
$dNTP_3$	1		11
VF_2	1		11
VR	1		11
Taq	.5		5.5
	24λ		

(1λ of DNA)

Gel = 8/16/12

Ldr	9c	12c	11c	12a

Gel 2 - 8/16/12

(empty)	1kb ladder	9d	8a	5a	1a	11a	12b

8/17/12
- started liquid cultures and made a transformation plate of the single ~~colonie~~ colony working detectors
- resuspended the Nitrate detectors in PBS (took before and after pictures)

- Gels = errors

SIGNATURE	DATE	WITNESS/TA	DATE