

# How can researchers prepare to make the switch to open science?

Associate Professor Siouxsie Wiles  
University of Auckland

 @SiouxsieW

Who?



@SiouxsieW

[www.superbugslab.org](http://www.superbugslab.org)

Image credit:  
Maki Naro



Who?



## Bioluminescent Superbugs Lab

[www.superbugslab.org](http://www.superbugslab.org)

Image credit:  
Maki Naro

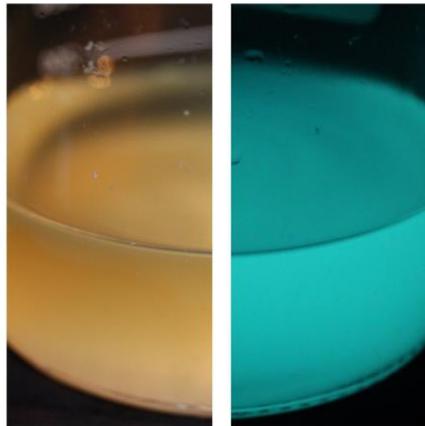


# Hunting for new antibiotics!

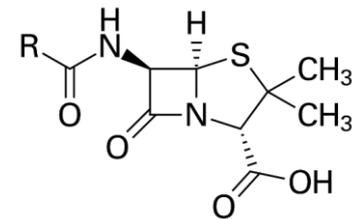
Dr Bevan Weir



\*Gross misrepresentation of the fungi in the ICMP...



Prof Brent Copp



\*Hopefully not just penicillin again and again...

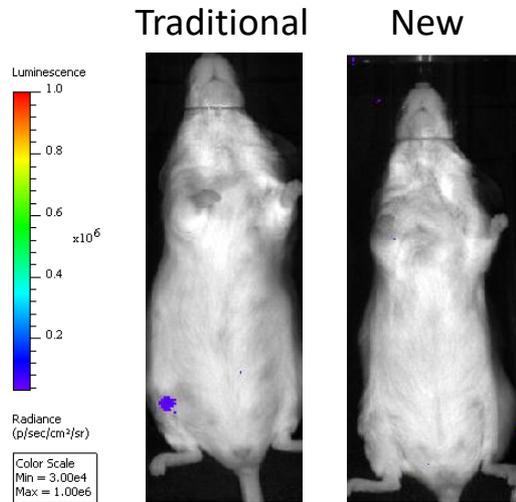


What?

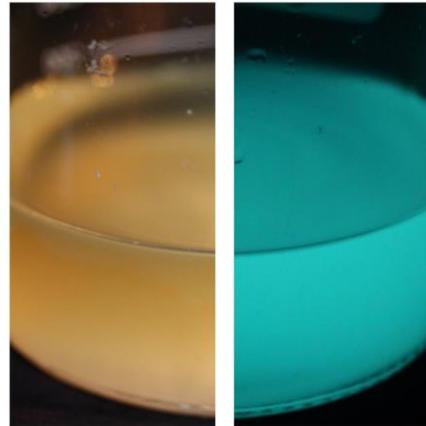
@SiouxsieW

# Making surgery safer?

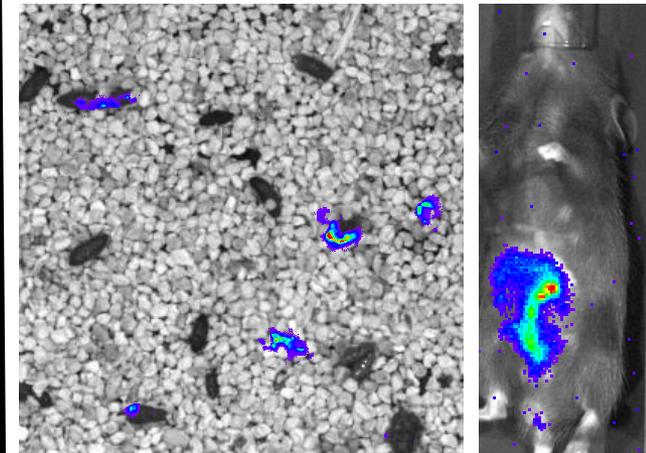
Dr Simon Young



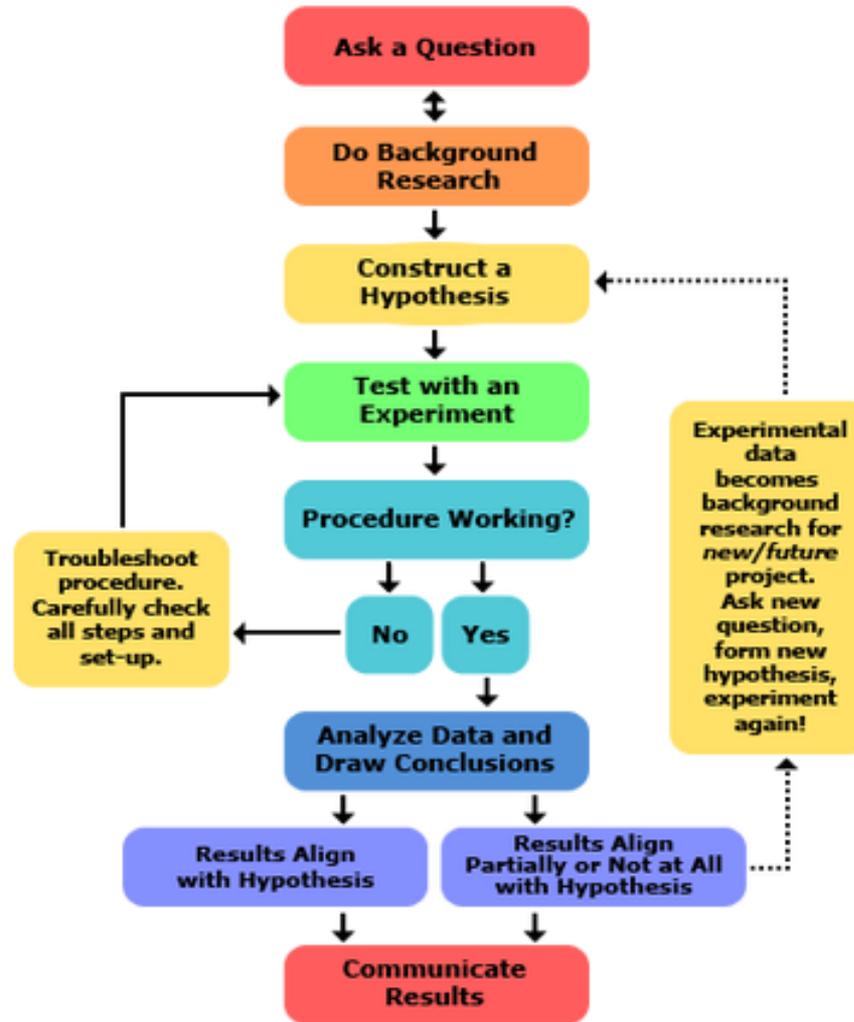
Young et al (2015). Clinical Orthopaedics and Related Research. 473(11):3573-84.



# What makes a microbe infectious?



# How we science!



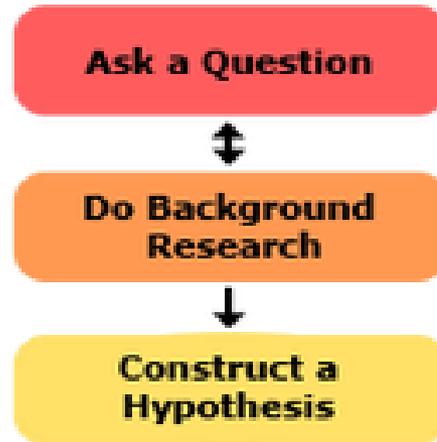
\$\$\$

How we science!



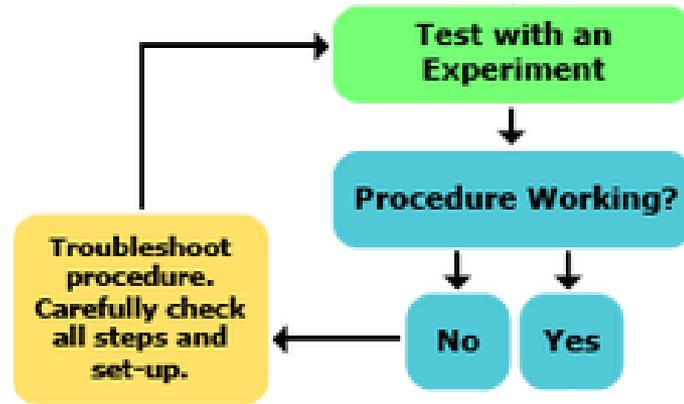
**Communicate  
Results**

How we science!



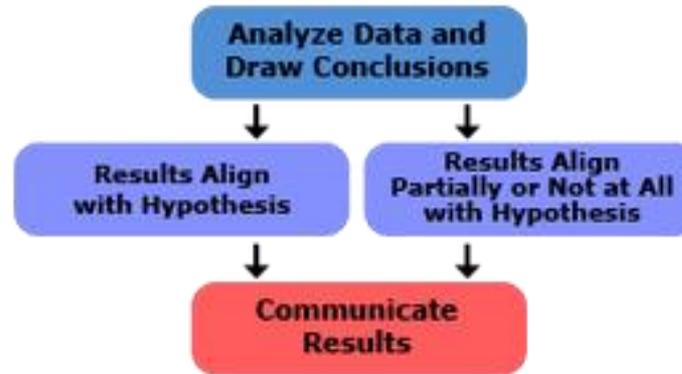
What's  
published/available?

How we science!



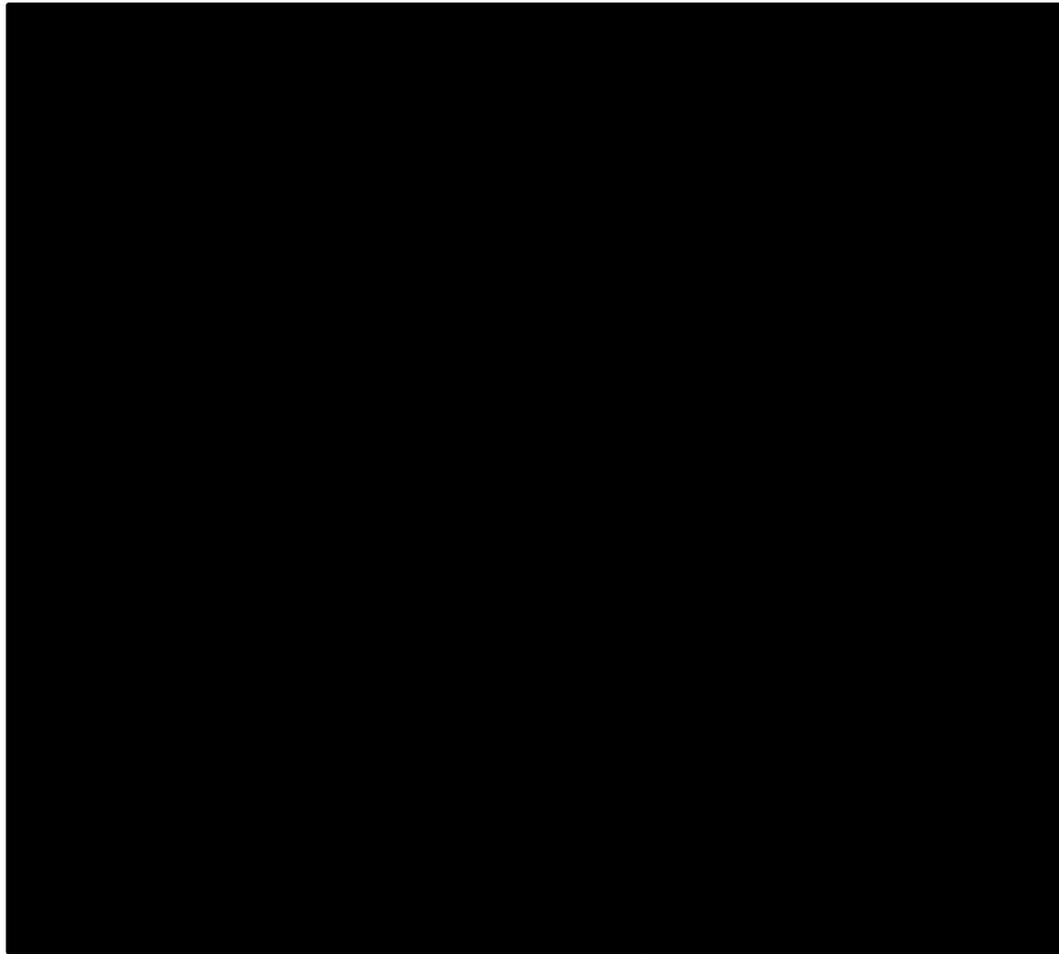
What's  
published/available?

How we science!



Warts and all?  
Completely documented?

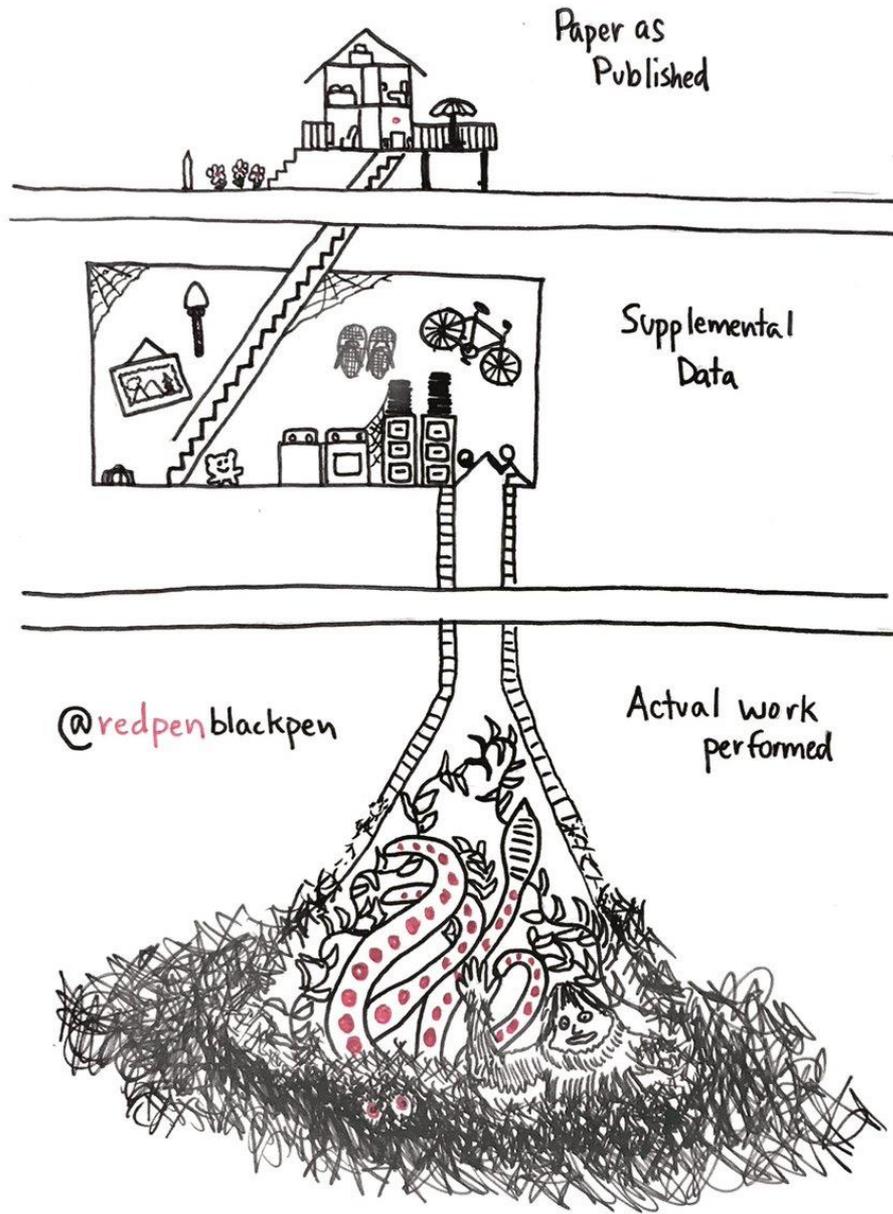
How we should science!



**Communicate  
Results**







**MY ATTEMPTS AT OPEN(ISH) SCIENCE...**

[ENTRIES](#)[PROTOCOLS](#)[ABOUT LAB BOOK](#)[BACK TO SBL](#)

## WELCOME!

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## RESEARCH

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[Evolution in Action \(27\)](#)



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## HANNAH 2/07/2015

Performed 16/06/2015 – 30/06/2015. A mouse competition experiment competing the ancestral ICC169 strain with the ancestral ICC180, N3, and W5 strains.

As before: 15 mL overnight cultures of ICC169 (3x), ICC180, N3 and W5 were spun down and resuspended in 1.5 mL sterile PBS (resulting in 10x concentrated cultures), and cultures were mixed in a 1:1 ratio as follows: ICC169 with ICC180, ICC169 with N3, and ICC169 with W5. 200  $\mu$ L of the mixtures were orally gavaged to groups of 6 C57BL/6 mice. For the ICC169/N3 group, the animals received 10  $\mu$ g/mL nalidixic acid in their drinking water for 1-day prior.

Mice received fresh drugs on Mondays, Wednesdays, and Fridays, and were imaged using an IVIS Kinetic (Caliper LifeScience) machine on Mondays and Fridays. Stools were taken and plated daily to measure relative ratios of the strains shed. At day 6 post-infection, mice were housed in a fresh cage for 1-day. At day 7, mice were re-housed and the 1-day old 'contaminated' cages were used to house naive animals (groups of 6) in order to follow natural transmission of the strains. These animals similarly received antibiotic (in the case of the ICC169/N3 group) in the drinking water and were monitored the same as the original animals.

Raw data: Bacteria shed from mice

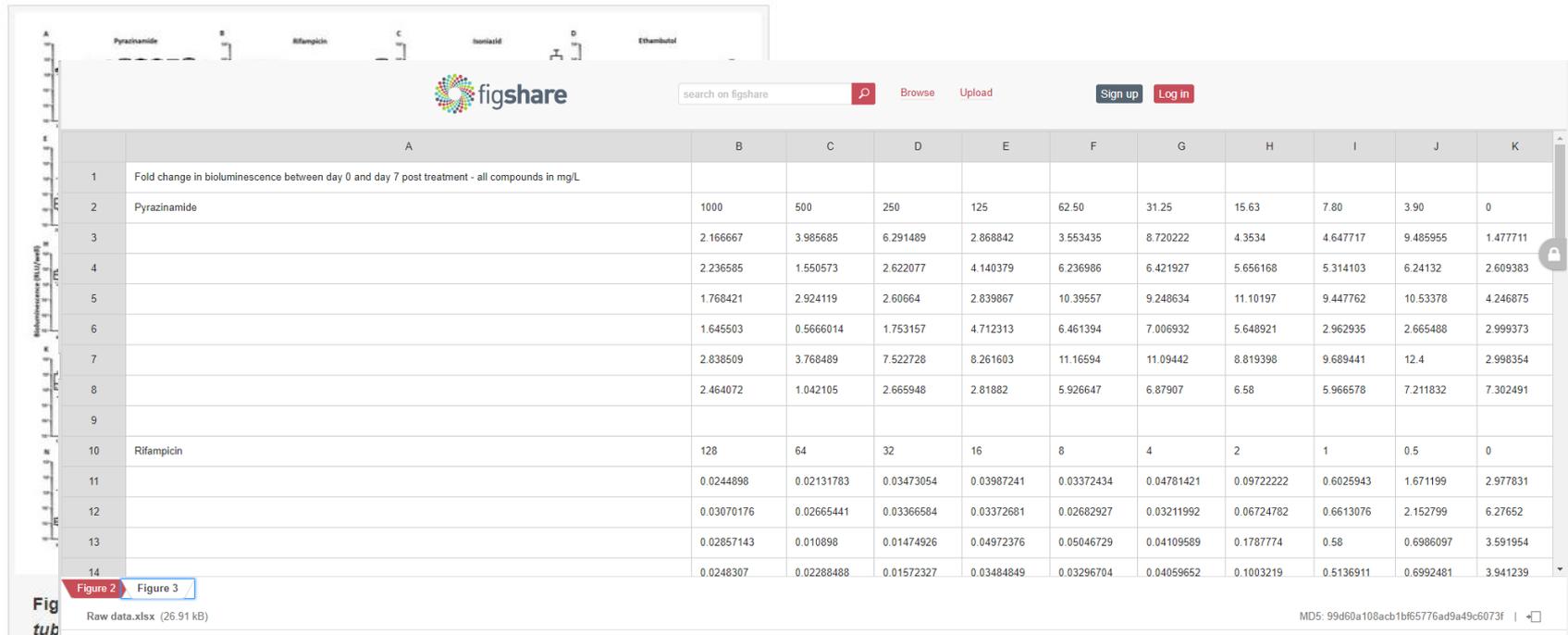
B	C	D	E	F	G	H	I	J	K	L	M	N	O	P	Q	R	S
			Nal plate (50 ug/mL nalidixic acid)					Km plate (50 ug/mL kanamycin)									
29-Jun	13 Days Post-infection	Mouse	Colonies Counted	Amount plated (uL)	Dilution	CFU/g/stool	Colonies Counted	Amount plated (uL)	Dilution	CFU/g/stool	ICC169	ICC180					
		21	9	14	9	10	1.00E+01	1.07E+05	5	8	6	10	1.00E+01	6.33E+04	4.33E+04	6.33E+04	
		22	32	24	19	10	1.00E+00	2.50E+04	22	19	24	10	1.00E+00	2.17E+04	3.33E+03	2.17E+04	
		23	33	19	35	10	1.00E+03	2.90E+07	17	29	25	10	1.00E+03	2.37E+07	5.33E+06	2.37E+07	
		24	24	31	27	10	1.00E+00	2.73E+04	20	16	17	10	1.00E+00	1.77E+04	9.67E+03	1.77E+04	
		25	29	20	21	10	1.00E+00	2.33E+04	22	18	14	10	1.00E+00	1.80E+04	5.33E+03	1.80E+04	
		26	5	13	9	10	1.00E+01	9.00E+04	8	4	8	10	1.00E+01	6.67E+04	2.33E+04	6.67E+04	
			Nal plate (50 ug/mL nalidixic acid)					Km plate (50 ug/mL kanamycin)									
29-Jun	13 Days Post-infection	Mouse	Colonies Counted	Amount plated (uL)	Dilution	CFU/g/stool	Colonies Counted	Amount plated (uL)	Dilution	CFU/g/stool	ICC169	ICC180					
		31	1	1	0	10	1.00E+00	6.67E+02	0	0	0	10	0.00E+00	0.00E+00	6.67E+02	0.00E+00	
		32	41	31	36	10	1.00E+02	3.60E+06	17	17	17	10	1.00E+00	1.70E+04	3.58E+06	1.70E+04	
		33	7	6	4	10	1.00E+00	5.67E+03	0	0	0	10	0.00E+00	0.00E+00	5.67E+03	0.00E+00	
		34	7	6	7	10	1.00E+00	6.67E+03	0	0	0	10	0.00E+00	0.00E+00	6.67E+03	0.00E+00	
		35				10		#DIV/0!				10		#DIV/0!	#DIV/0!	#DIV/0!	
		36	45	42	31	10	1.00E+01	3.93E+05	5	1	1	10	1.00E+00	2.33E+03	3.91E+05	2.33E+03	
			Nal plate (50 ug/mL nalidixic acid)					Km plate (50 ug/mL kanamycin)									
30-Jun	14 Days Post-infection	Mouse	Colonies Counted	Amount plated (uL)	Dilution	CFU/g/stool	Colonies Counted	Amount plated (uL)	Dilution	CFU/g/stool	ICC169	ICC180					
		11	3	1	3	10	1.00E+00	2.33E+03	0	1	3	10	1.00E+00	1.00E+03	1.00E+03	1.33E+03	
		12	57	52	46	10	1.00E+01	5.17E+05	18	23	26	10	1.00E+00	2.23E+04	4.94E+05	2.23E+04	
		13	11	11	6	10	1.00E+00	9.33E+03	0	0	0	10	0.00E+00	0.00E+00	9.33E+03	0.00E+00	
		14	27	14	19	10	1.00E+00	2.00E+04	14	11	12	10	1.00E+00	1.23E+04	7.67E+03	1.23E+04	
		15	1	1	0	10	1.00E+00	6.67E+02	0	0	0	10	0.00E+00	0.00E+00	6.67E+02	0.00E+00	
		16	0	0	0	10	0.00E+00	0.00E+00	0	0	0	10	0.00E+00	0.00E+00	0.00E+00	0.00E+00	
			Nal plate (50 ug/mL nalidixic acid)					Km plate (50 ug/mL kanamycin)									
30-Jun	14 Days Post-infection	Mouse	Colonies Counted	Amount plated (uL)	Dilution	CFU/g/stool	Colonies Counted	Amount plated (uL)	Dilution	CFU/g/stool	ICC169	ICC180					
		21	0	0	0	10	0.00E+00	0.00E+00	0	0	0	10	0.00E+00	0.00E+00	0.00E+00	0.00E+00	
		22	0	0	0	10	0.00E+00	0.00E+00	0	0	0	10	0.00E+00	0.00E+00	0.00E+00	0.00E+00	
		23	12	12	7	10	1.00E+02	1.03E+06	5	13	3	10	1.00E+02	7.00E+05	3.33E+05	7.00E+05	
		24	0	0	0	10	0.00E+00	0.00E+00	0	0	0	10	0.00E+00	0.00E+00	0.00E+00	0.00E+00	
		25	8	11	10	10	1.00E+00	9.67E+03	4	3	1	10	1.00E+00	2.67E+03	7.00E+03	2.67E+03	
		26	0	0	0	10	0.00E+00	0.00E+00	0	0	0	10	0.00E+00	0.00E+00	0.00E+00	0.00E+00	
			Nal plate (50 ug/mL nalidixic acid)					Km plate (50 ug/mL kanamycin)									
30-Jun	14 Days Post-infection	Mouse	Colonies Counted	Amount plated (uL)	Dilution	CFU/g/stool	Colonies Counted	Amount plated (uL)	Dilution	CFU/g/stool	ICC169	ICC180					
		31	7	5	16	10	1.00E+00	9.33E+03	0	0	0	10	0.00E+00	0.00E+00	9.33E+03	0.00E+00	
		32	56	68	72	10	1.00E+01	6.53E+05	37	38	46	10	1.00E+00	4.03E+04	6.13E+05	4.03E+04	
		33	1	2	1	10	1.00E+00	1.33E+03	0	0	0	10	0.00E+00	0.00E+00	1.33E+03	0.00E+00	
		34	1	1	0	10	1.00E+00	6.67E+02	0	0	0	10	0.00E+00	0.00E+00	6.67E+02	0.00E+00	
		35				10		#DIV/0!				10		#DIV/0!	#DIV/0!	#DIV/0!	
		36	7	0	5	10	1.00E+00	4.00E+03	0	0	0	10	0.00E+00	0.00E+00	4.00E+03	0.00E+00	

# Effect of common and experimental anti-tuberculosis treatments on *Mycobacterium tuberculosis* growing as biofilms

Related research

[Research article](#) [Microbiology](#) [Infectious Diseases](#)

James P. Dalton<sup>1,2,3</sup>, Benedict Uy<sup>1,2</sup>, Narisa Phummarin<sup>4</sup>, Brent R. Copp<sup>3,4</sup>, William A. Denny<sup>3,5</sup>, Simon Swift<sup>1</sup>, Siouxsie Wiles<sup>1,2,3</sup>



**LESSONS LEARNED...**

Open Access is not open science/research.....

Need to really think about how we document our entire research workflow (warts and all...)

If I was hit by a car tomorrow would anyone else be able to make sense of that documentation?

Raw data vs tidy data...



Image credit:  
Gravity Falls (<https://disneynow.go.com/shows/gravity-falls>)



Image credit:  
<https://www.deviantart.com/alicornofmagic/art/Collab-My-Very-Best-Friends-370402852>

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www.redbubble.com/people/siouxsiew



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