# **State Water Board DPR Pathogen Research**

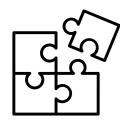
# Tools to Evaluate Microbial Risk, Plant Performance, and Reliability (DPR-1)

# Raw Wastewater Pathogen Monitoring (DPR-2)

Brian Pecson, P.I. for DPR-1 and DPR-2, Trussell Technologies







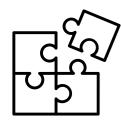
# How Much Pathogen Treatment?

#### Wastewater

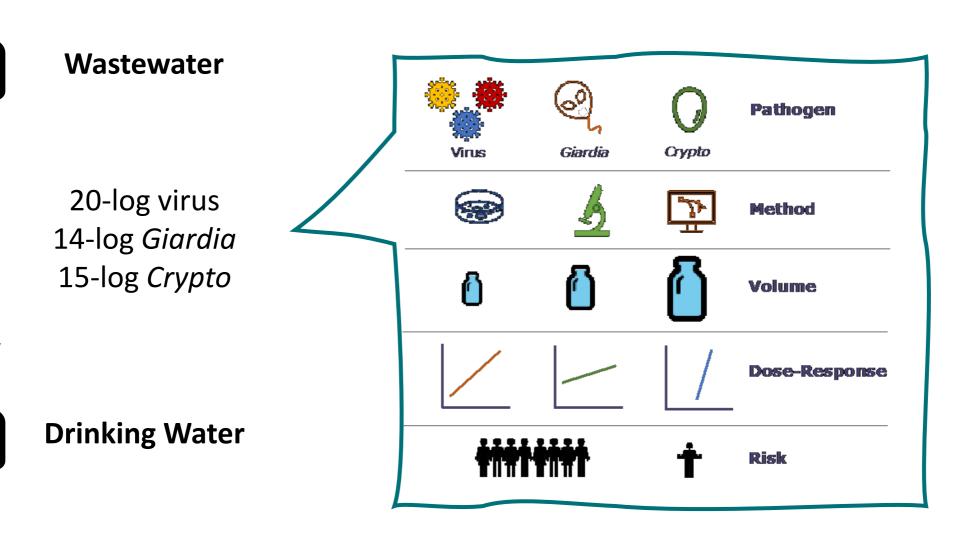
20-log virus 14-log *Giardia* 15-log *Crypto* 

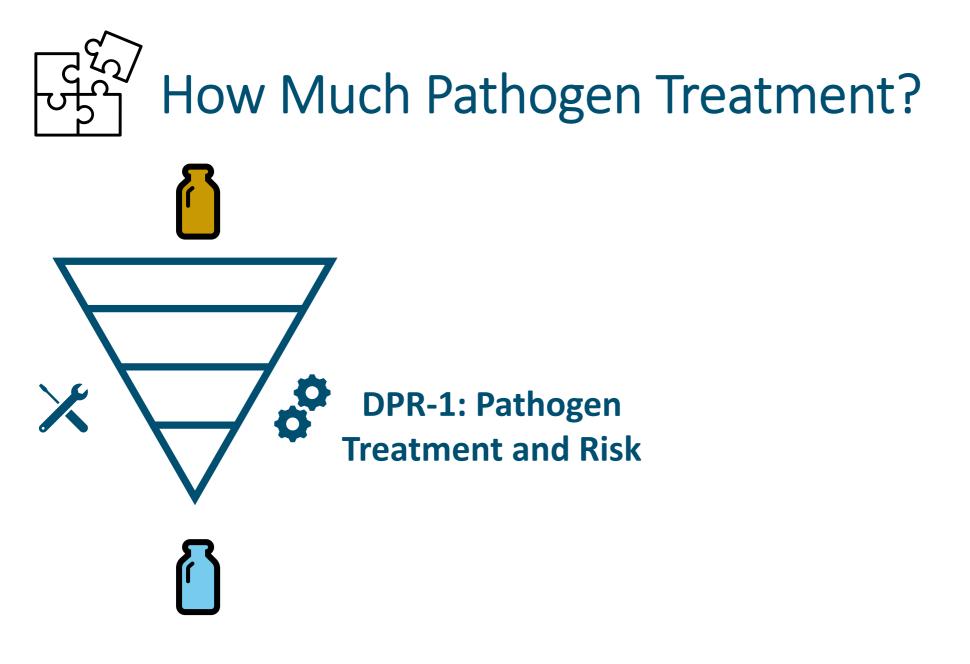


**Drinking Water** 



# How Much Pathogen Treatment?





## DPR-1 Technical Work Group and Research Team



Brian Pecson (chair) Trussell Technologies

Nick Ashbolt Southern Cross University



**Charles Haas** Drexel University



#### **Technical Work Group**

**Theresa Slifko** Metropolitan Water District

#### **Research Team**



Dan Gerrity SNWA



**Edmund Seto** University of Washington

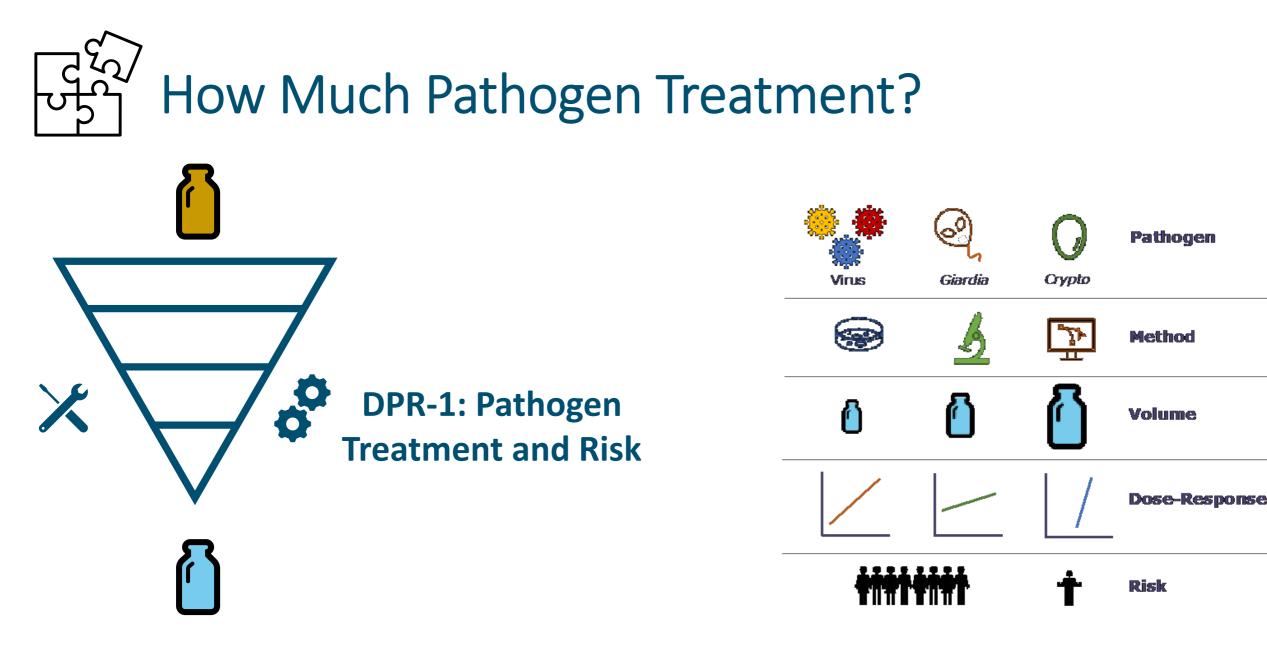
#### **DPR-1 Staff**

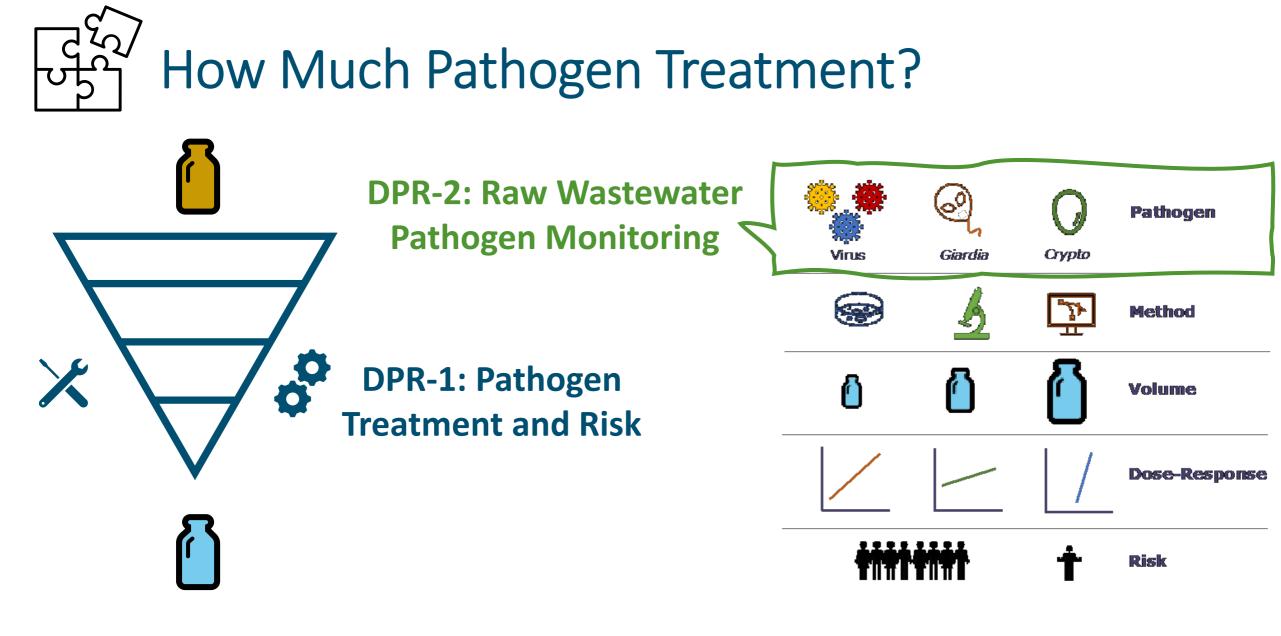


Anya Kaufmann Trussell Technologies

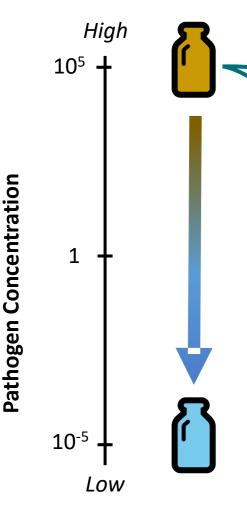


Adam Olivieri WRF/State Board Coordination





#### Motivation for Research



- Wastewater pathogen concentrations are key inputs
- Industry does not have sufficient high-quality data
- SOPs needed to address previous limitations

## **DPR-2** Technical Work Group



**George Di Giovanni** Metropolitan Water District



Menu Leddy Essential Environmental & Engineering Systems



Kara Nelson UC, Berkeley



Brian Pecson Trussell Technologies



**Channah Rock** University of Arizona



Theresa Slifko (chair) Metropolitan Water District



cel analytical, inc. water, wastewater, and soil laboratory services

DPR-2 Staff:



**Emily Darby** Trussell Technologies



Adam Olivieri WRF/State Board Coordination



Walt Jakubowski QA/QC Officer

DPR-2 Labs:



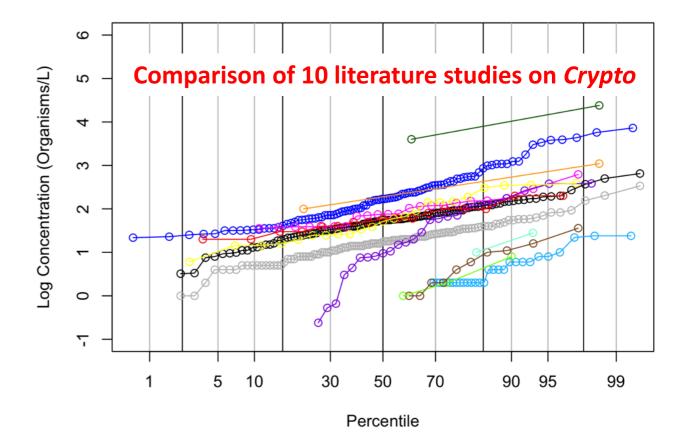
SCIENTIFIC METHODS

BIOLOGICAL CONSULTING SERVICES OF NORTH FLORIDA, INC.



## Literature Review

- Limited number of studies
- Low method sensitivity and high frequency of non-detects
- Recovery often not measured
- QA/QC often not strictly followed



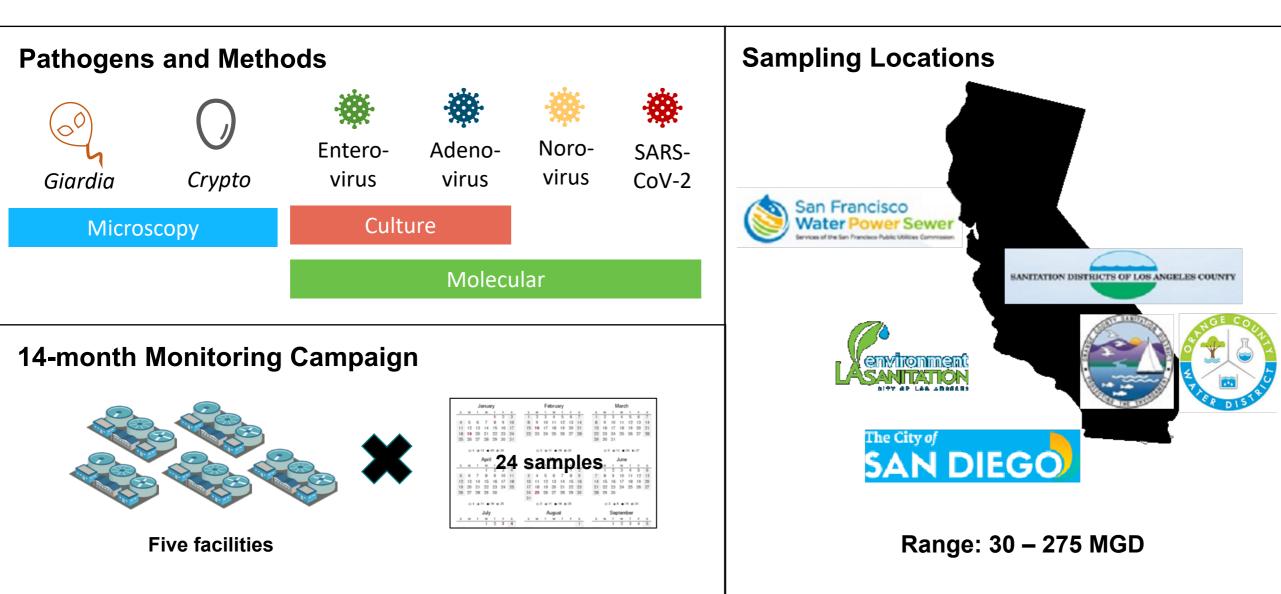
- Conclusions
  - Possible to measure pathogens in raw WW
  - -<u>Amount</u> and <u>quality</u> of data are insufficient

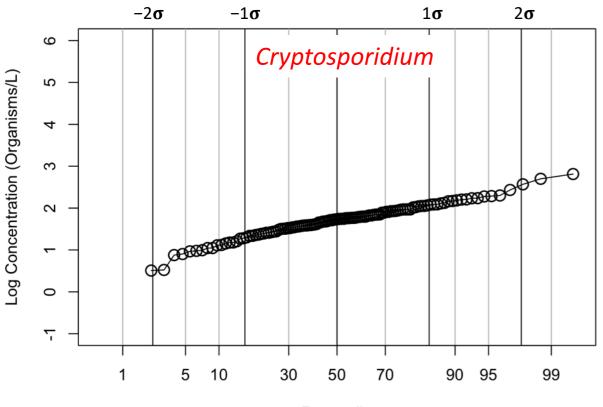
## Method Optimization

- Used to address limitations of past studies
  - Concentration method
  - Volume of sample to process
- Require strict QA/QC
  - Matrix spikes in 75% of all samples
  - Full set of controls
- Recommendation: use DPR-2 QAPP for future studies

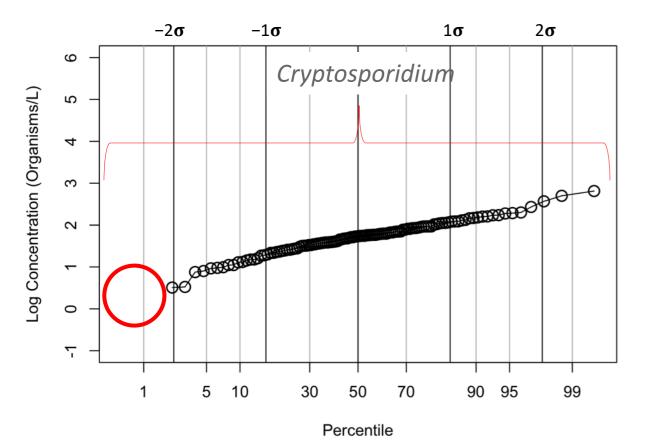
	QAPP Analytical Microbiology Supporting Version 4.0.	WRF Contract No: 4952 Date: 05.06.20
	Quality Assurance Project Pla	in
	Analytical Microbiology Services	
	Water Research Foundation Contract #4952	
	Prepared for:	
	The Water Research Foundation	
	Prepared by:	
<b>4</b> :::	<i>cel analytical, inc.</i> weter, westewater, and soil laboratory services	
	82 Mary Street Suite 2 San Francisco, CA 94103	
	Yeggie Dearborn Ph.D. Program Manager	
	Email: yeggie@celanalytical.com	
	ugust; October Version 1.0, Rev.01 November	
	Version 2.0, Rev.02 Version 2.0, Rev.03	Eabruary 2021
1	Version 3.0	February 2021

# Sampling campaign

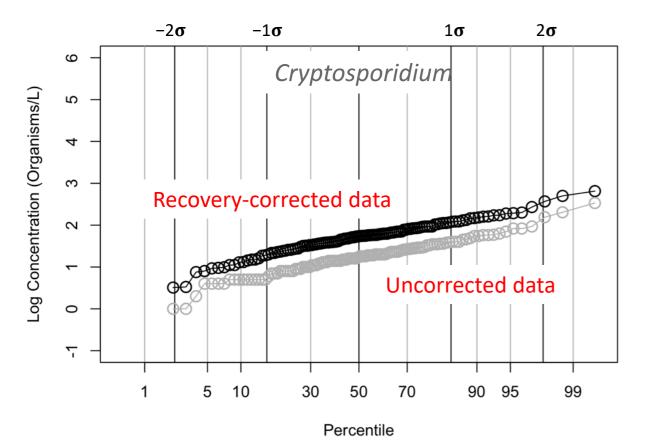




Percentile

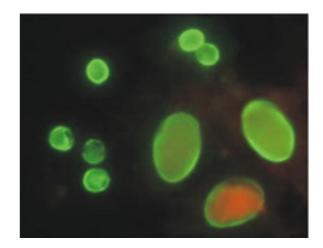


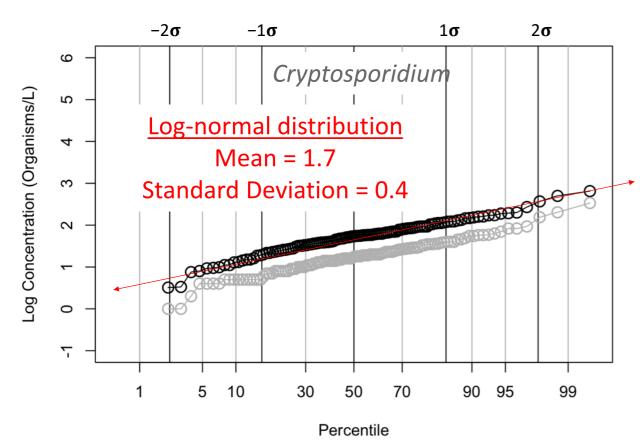
• High rate of <u>detects</u> across full range



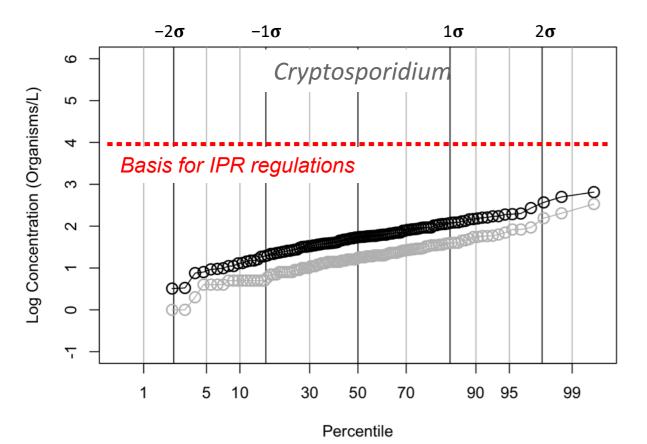
• High rate of <u>detects</u> across full range

- Matrix spikes used to <u>correct</u> for losses
- High <u>recovery efficiency</u>

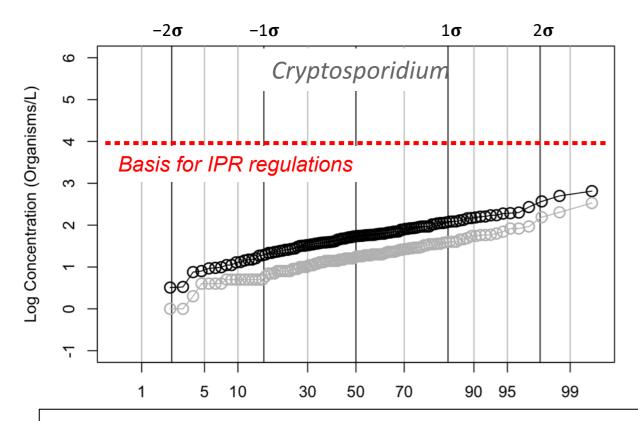




- High rate of <u>detects</u> across full range
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- Models estimate past measured range



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- Matrix spikes used to <u>correct</u> for losses
- High <u>recovery efficiency</u>
- <u>Models</u> estimate past measured range
- Allows for <u>comparison</u> with IPR regs

#### **Recommendation**:

Use modeled distributions for probabilistic assessments of treatment targets

## **Other Key Findings**

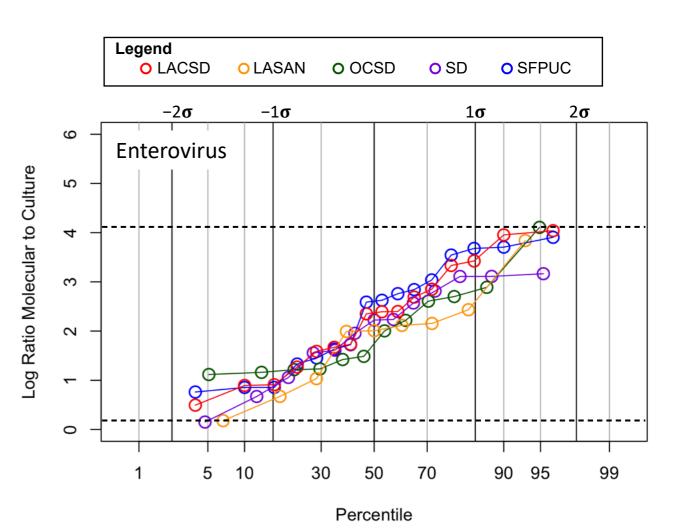
- Pathogen distributions similar across treatment plants
  - 94% of comparisons had no significant differences between facilities
- Minimal level of seasonality observed
  - Enterovirus higher in summer / adenovirus higher in winter
- No clear impact of COVID-19 on concentrations
  - Data collected before and after Stay-at-Home order showed minimal change
- Uncertainties associated with the use of molecular data

## Issues with the use of molecular data

 Genome copies (GC) not always associated with *infective* virus

• Difficult to link GC with infectivity

- DPR-2 ratios span orders of magnitude:
  - 10,000:1 to 1:1 (enterovirus)
  - 100,000:1 to 1:1 (adenovirus)

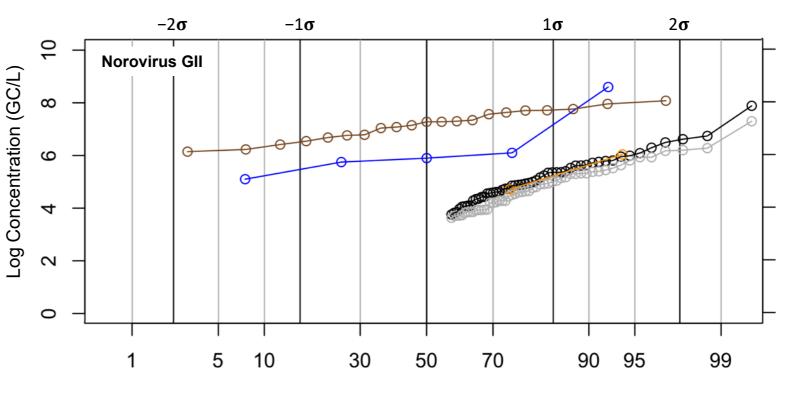


# When is this important?

Norovirus not culturable

 Dose-response function makes assumptions about "infectivity" of genome copies

 If we assume 1:1, then each GC is an infectious unit (IU)



Percentile

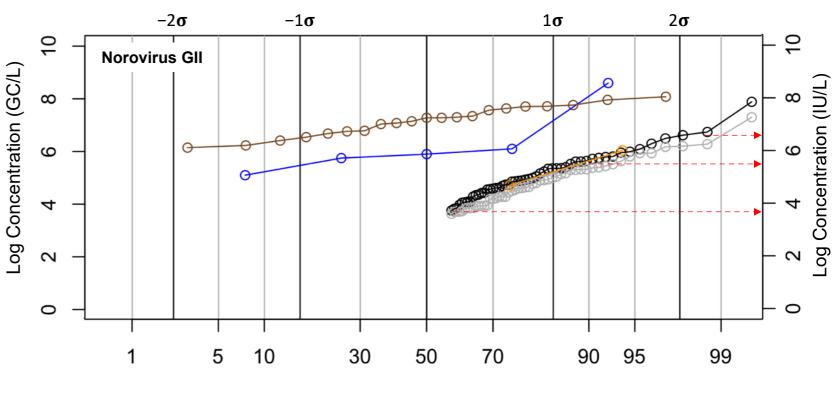


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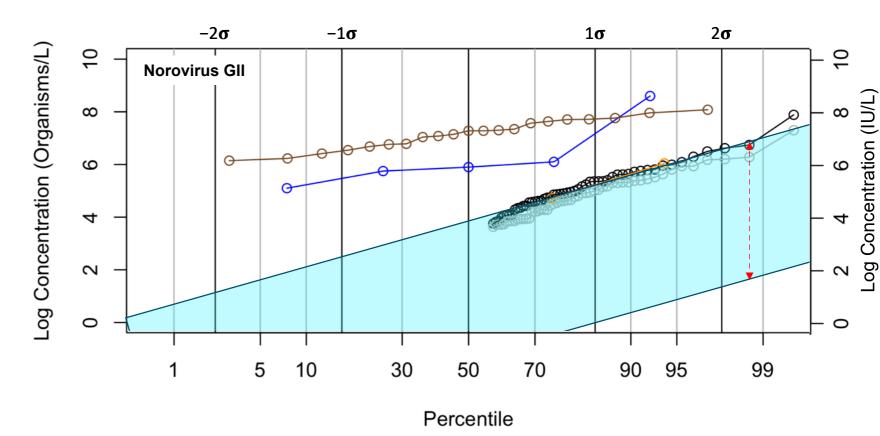
Percentile

Legend		
O This work - corrected data		O San Diego 2016
O This work - raw data	O Simmons & Xagoraraki 2011b	

#### Incorporate uncertainty in risk analyses

 DPR-1 Final Report shows how to incorporate molecular data into analysis

 Results in a "band" of potential values

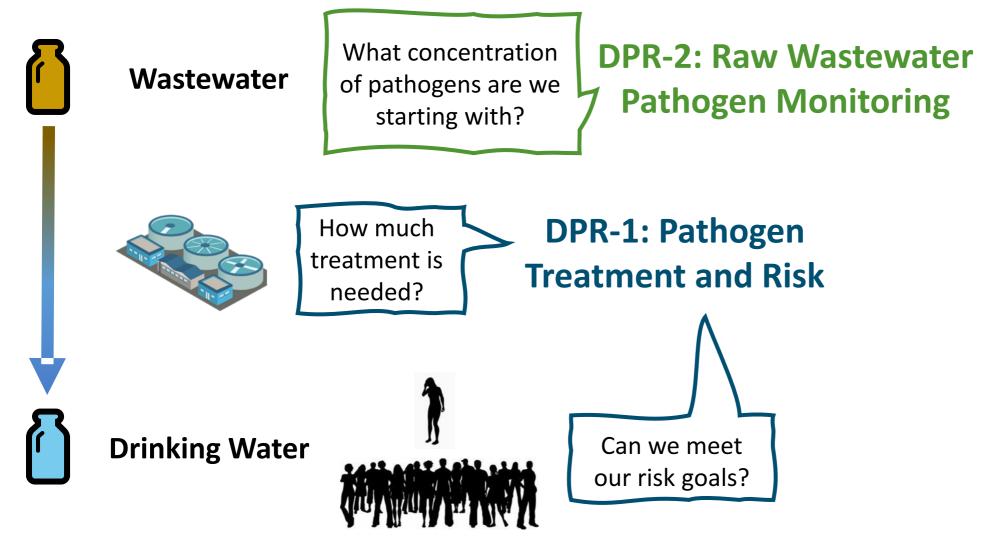


iego 2016
goraraki 2011b

#### **Recommendations for Regulatory Development**

- Use DPR-2 datasets as the raw wastewater inputs for QMRA
- Correct pathogen data for recovery using matrix spikes
- Use culture data to reduce uncertainties with molecular interpretation; follow TWG recommendations for the use of molecular data
- Model the DPR-2 distributions (and relevant literature) for use in probabilistic assessments
- Require DPR-2 QAPP/SOPs for future pathogen monitoring studies

## How Much Pathogen Treatment?



## **DPRisk Tool and Guidance Document**

#### DPRisk: QMRA Tool

DPRisk		Guidance Do
ersion 1.0.1 (11.05.2020) ponsored by: The Water Research Foundation opyright (C)2017 by The Water Research Foundation. ALL	Research FOUNDATION	Table of Conten
Introduction	Quantitative Microbial Risk Assessment and Probabilistic Assessment of Treatment Train Performance for Direct Potable Reuse Scenarios	List of Acronyms Project Definition a
Background	This tool is intended to facilitate quantitative microbial risk assessment (QMRA) and probabilistic assessment of treatment train	Historical Context . Overview of DPRisl
How to use the tool	performance (PATTP) for various direct potable reuse (DPR) scenarios. There are many possible analyses that you can conduct with this tool, including:	Step 1: Target Path
License	There are many possible analyses that you can conduct with this tool, including:	Step 2: Raw Waste
Litense	Developing a distribution of treatment train performance for different potential DPR treatment trains.	Step 3: Raw Waste
Model Specification	<ul> <li>Evaluating daily and annual risks of infection for multiple microbial pathogens for different potential DPR treatment trains.</li> <li>Comparing different DPR treatment trains in terms of treatment performance and risk.</li> </ul>	Step 4: Identifying
Raw Wastewater Pathogen Concentrations	Evaluating the impact of failures on treatment performance and risk.	Step 5: Assigning T
Treatment Train	The accompanying Guidance Document provides useful context for this tool, including: <ul> <li>The background motivation for the creation of the tool.</li> </ul>	Step 6: Treatment
	• The historical context for the use of PATTP and QMRA in DPR.	Step 7: Manageme
Treatment Failure	<ul> <li>The project process that resulted in this tool.</li> <li>Detailed descriptions of each step of the tool, including references for default assumptions.</li> </ul>	Step 8: Drinking W
Management Barriers	<ul> <li>Details on the computations implemented by the tool.</li> <li>Example case studies to help you get started with using the tool.</li> </ul>	Step 9: Pathogen D
Exposure	This tool was developed in the R statistical language.	Step 10: Risk Chard
Dose-Response		Final Tool Consider
Juse-Response		Case Study 1: QMR
Results		Case Study 2: QMR
PATTP Output		Case Study 3: QMR
QMRA Output		Conclusions
-		References
Summary of PATTP and QMRA Output		Appendix 1 – Sumn
Comparison of Risk Curves		Appendix 2 – Instal

#### **DPRisk: Guidance Document**

#### **Document for DPRisk**

#### nts

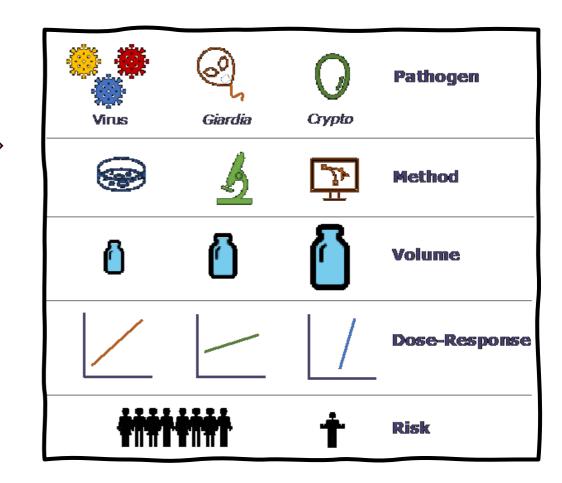
List of Acronyms
Project Definition and Background
Historical Context
Overview of DPRisk
Step 1: Target Pathogens (Hazard Identification)7
Step 2: Raw Wastewater Pathogen Datasets9
Step 3: Raw Wastewater Pathogen Distributions13
Step 4: Identifying Unit Processes for the Treatment Train
Step 5: Assigning Treatment Process Log Reduction Values
Step 6: Treatment Process Failure Framework
Step 7: Management Barriers (Blending, Dilution, and Die-off)
Step 8: Drinking Water Ingestion (Exposure Assessment)
Step 9: Pathogen Dose Response Models (Dose Response Assessment)
Step 10: Risk Characterization
Final Tool Considerations
Case Study 1: QMRA for Enterovirus in a Default DPR Scenario
Case Study 2: QMRA for Cryptosporidium in a FAT-Based DPR Scenario
Case Study 3: QMRA for Adenovirus in an FAT-Based DPR Scenario
Conclusions
References
Appendix 1 – Summary of Output File Headers
Appendix 2 – Installation of DPRisk on Shinyapps.io

#### Also: User Input Files for 3 Case Studies

#### **DPRisk Features**

#### **INPUTS:**

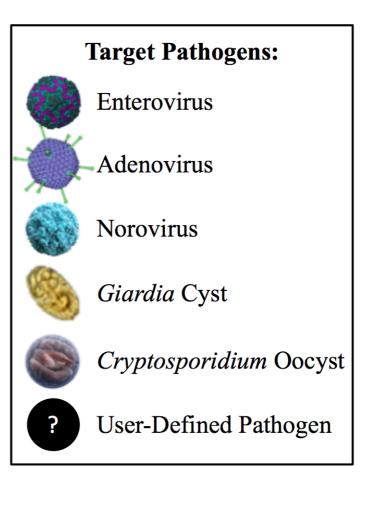
- Raw Wastewater Pathogen Concentrations
- Treatment Train
- Treatment Failure
- Exposure
- Dose Response



#### **DPRisk Features**

#### **INPUTS:**

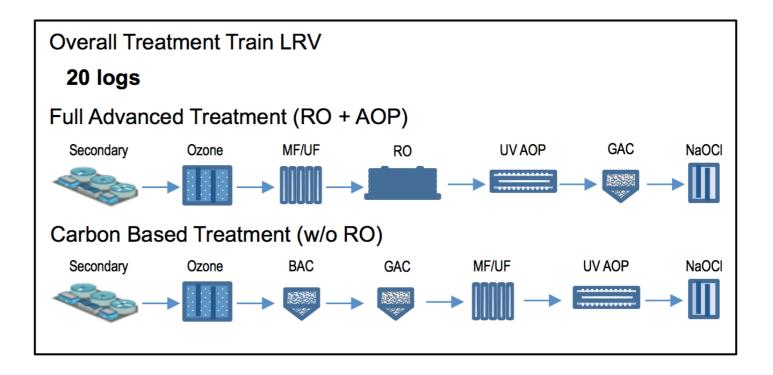
- Raw Wastewater Pathogen Concentrations
- Treatment Train
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#### **DPRisk Features**

#### **INPUTS:**

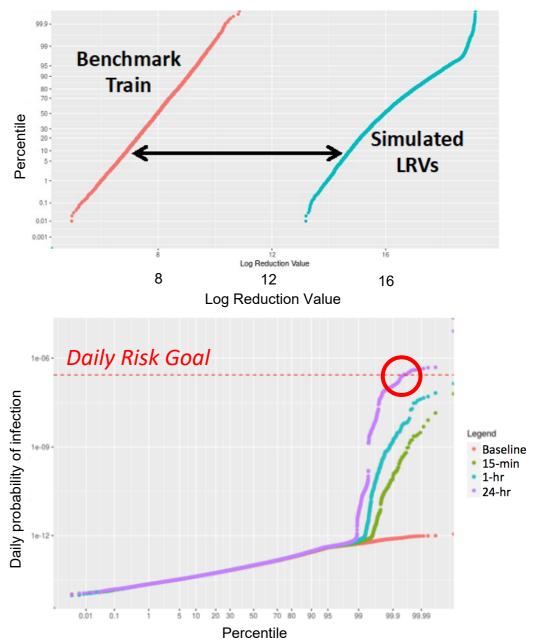
- Raw Wastewater Pathogen Concentrations
- Treatment Train
- Treatment Failure
- Exposure
- Dose Response



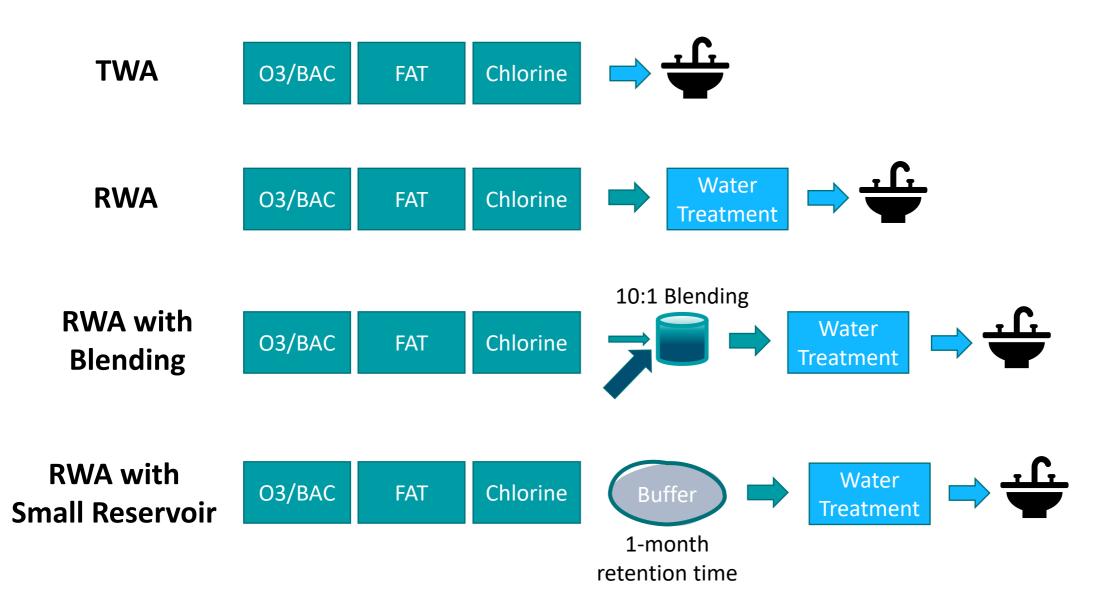
#### **DPRisk Outputs**

• Probabilistic Assessment of Treatment Train Performance (PATTP)

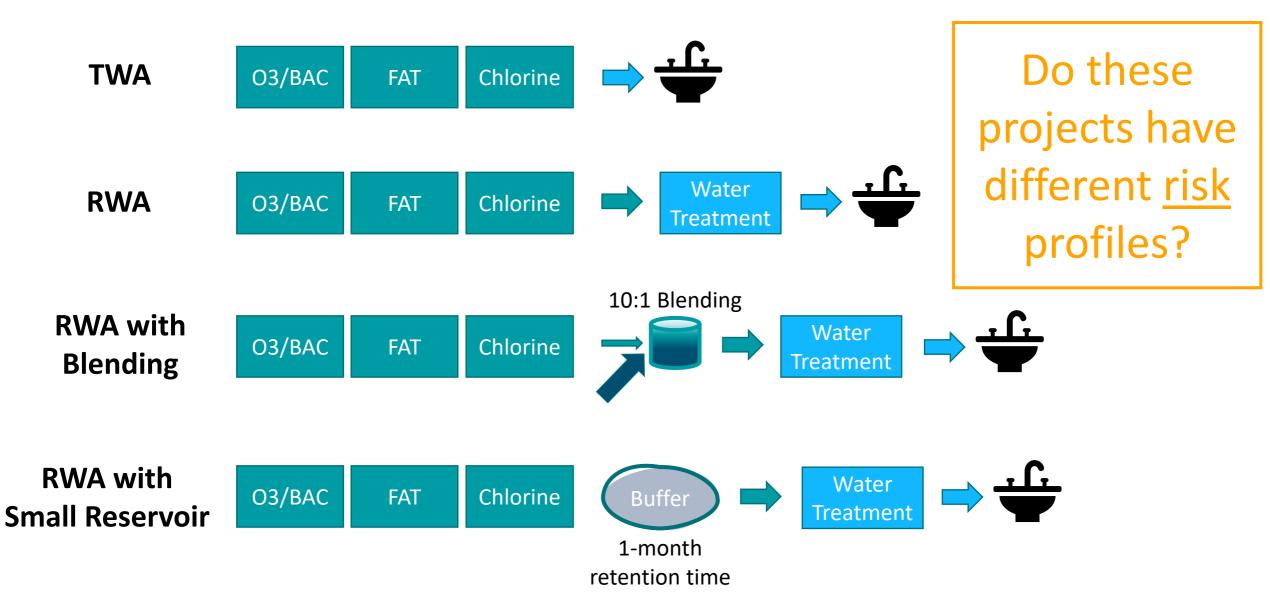
Quantitative Microbial Risk Assessment



#### Not All DPR Projects Are Alike

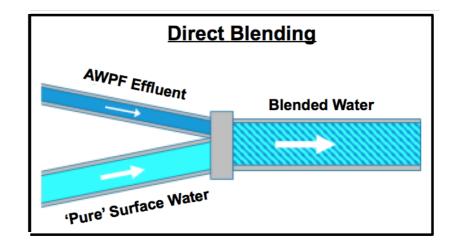


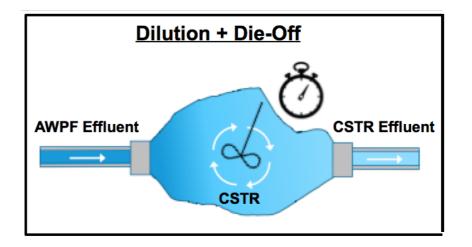
#### Not All DPR Projects Are Alike



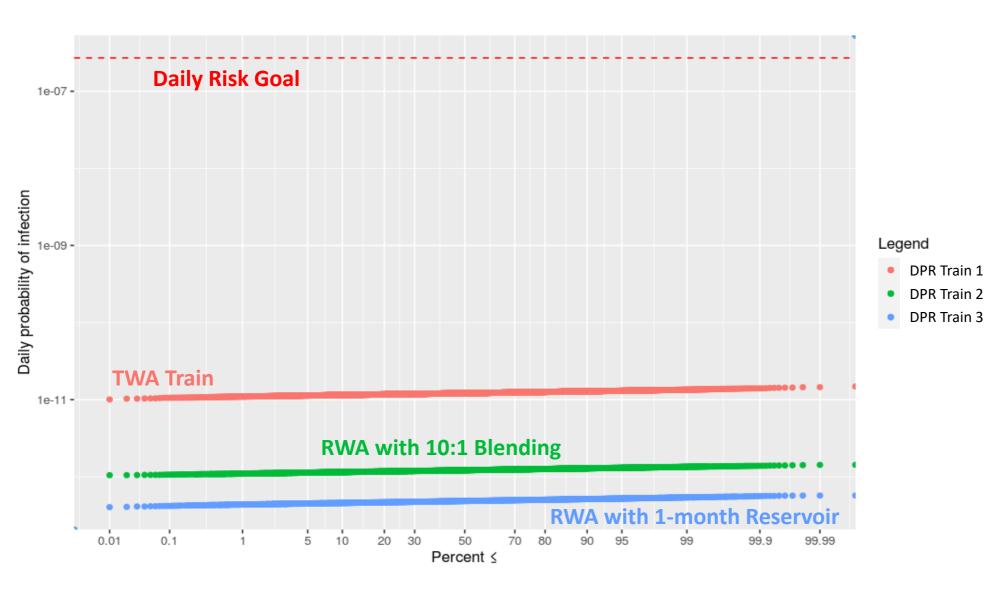
## **DPRisk Includes Management Barriers**

	Blending	
	Specify the log removal associated with blending.	
	Specify log removal for blending as:	
	Point estimate	
	Log Removal:	
	0	
DPRisk Inputs		
	Dilution	
	Specify the log removal associated with dilution.	
	Specify log removal for dilution as:	
	Point estimate	
	Log Removal:	
	0	

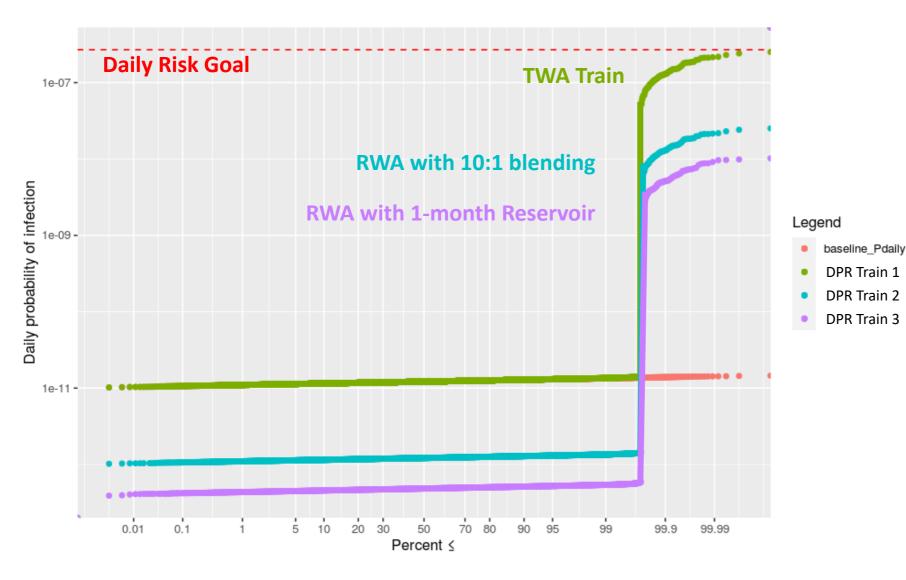




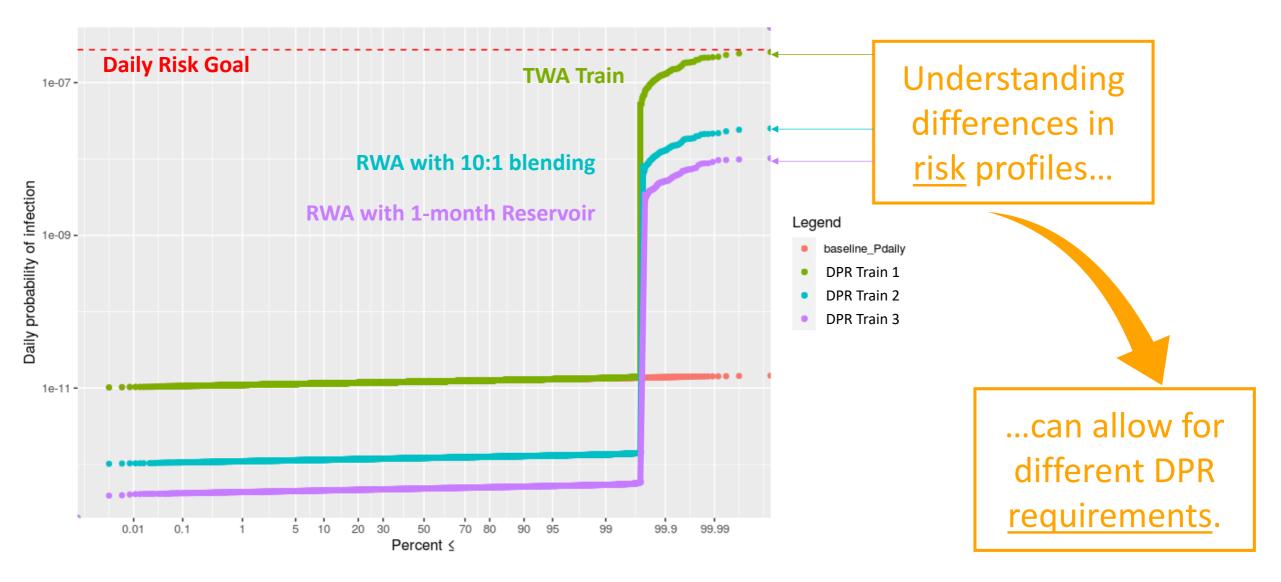
## Risk Profiles of RWA and TWA Trains (no failures)



#### Risk Profiles of RWA and TWA with Failure Analysis



#### Risk Profiles of RWA and TWA with Failure Analysis



#### Recommendations

• Select modeled distributions from DPR-2 as raw wastewater inputs

• Use DPRisk for *probabilistic* assessments of performance and risk

 Develop frameworks to incorporate the benefits of non-treatment (management) barriers in RWA and TWA

## Acknowledgements

- Anya Kaufmann, Trussell Technologies (DPR-1)
- Dan Gerrity, Southern Nevada Water Authority (DPR-1)
- Emily Darby, Trussell Technologies (DPR-2)

#### Questions?

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