

Five-week warning of COVID-19 peaks prior to the Omicron surge in Detroit, Michigan using wastewater surveillance

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Overview



Introduction & research objectives

Lag time of wastewater signals preceding clinical cases

Methods & experiments

Prediction of COVID-19 cases based on wastewater data

Conclusions & future work

Introduction – Wastewater surveillance 🦷 MICHIGAN STATE UNIVERSITY



COVID-19 infections (SARS-CoV-2)





Wastewater treatment plant sampling

- 1. Provide early warnings of upcoming disease incidences
- 2. Predict upcoming disease incidences through modeling based on WBE data



COVID-19 cases collected for the Detroit metropolitan area

Virus concentration, RNA extraction, and RT-ddPCR



Wastewater viral concentrations data (e.g. SARS-CoV-2)

Objectives



Lag time for early warning

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Sampling



Interceptor tributary area



 Untreated wastewater samples were collected from three main interceptors: Detroit River Interceptor (DRI), North Interceptor-East Arm (NIEA), and Oakwood-Northwest-Wayne County Interceptor (ONWI) in southeast Michigan.
We used NanoCeram column filters based on the EPA Virus Adsorption-Elution (VIRADEL) method to collect wastewater.

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Sampling onsite

Sampling



Sampling apparatus



Sampling mechanism



Sample processing

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Vortex during RNA extraction Droplet generator fo RT-ddPCR Thermocycler for RT-ddPCR

Droplet reader for RTddPCR

Collection of COVID-19 cases



Study Area: Three Largest Counties in Michigan

COVID-19 cases collected for the City of Detroit, as well as Wayne, Macomb, and Oakland counties

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Results

Genomic copies/L



Total N1 and N2 gene concentrations in gc/L for the three interceptors and total confirmed COVID-19 cases in the study areas (gray area)

Correlations

Pearson's correlation between N1 and N2 gene concentrations (in different units) and total COVID-19 cases in city of Detroit, as well as Wayne, Macomb, and Oakland counties with lag times

Lag time	Unit	N1 vs. total cases	N2 vs. total cases	
	gc/l	0.27	0.28	
3 weeks	gc/d	0.11	0.11	
	gc/l of sanitary flow	0.10	0.09	
	gc/l	0.51	0.52	
4 weeks	gc/d	0.29	0.26	
	gc/l of sanitary flow	0.28	0.25	
gc/l		0.62	0.64	
5 weeks	gc/d	0.34	0.31	
	gc/l of sanitary flow	0.33	0.30	

Lag time

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	Lag time	Model	Equation (N1, gc/L)		Pearson r
	3 weeks	veeksLinear $y_{t(3)} = 0.015 x_{t(3)} + 534.96$ Autoregression $y_{t(3)} - 0.2928y_{t(3)-1} - 0.3741 y_{t(3)-2} = 0.015(x_{t(3)} - 0.2928x_{t(3)-1} - 0.3741x_{t(3)-2}) + 534.96$		7.22	0.26
				135.65	0.07
	Autoregression+ time effect $y_{t(3)} = 1052 - 1.7923t(3) - 0.044dx_{t(3)}$ $y_{t(3)}*= y_{t(3)} + 0.5857y_{t(3)-1}$		$y_{t(3)} = 1052 - 1.7923t(3) - 0.044dx_{t(3)}$ $y_{t(3)} = y_{t(3)} + 0.5857y_{t(3)-1}$	10.18	0.76
Ve Autore		Vector Autoregression	$y_{t(3)} = 1.30y_{t(3)-1} + 0.12 y_{t(3)-2} - 0.49 x_{t(3)-1} - 0.01x_{t(3)-2} - 73.46$	8.32	0.72
	4 weeks	Linear	$y_{t(4)} = 0.29 x_{t(4)} + 227.04$	7.26	0.51
		Autoregression	$y_{t(4)}-0.1706y_{t(4)-1} - 0.2799 y_{t(4)-2} = 0.29(x_{t(4)}-0.2354x_{t(4)-1} - 0.2799x_{t(4)-2}) + 227.04$	182.92	0.50
		Autoregression+ time effect	$y_{t(4)} = 1730 + 6.78t(4) + 0.06dx_{t(4)}$ $y_{t(4)} = y_{t(4)} + 0.59y_{t(4)-1}$	7.50	0.92
		Vector Autoregression	$y_{t(4)} = 1.42y_{t(4)-1} + 0.20 \ y_{t(4)-2} - 0.61 \ x_{t(4)-1} + 0.004x_{t(4)-2} + 109.89$	8.00	0.86
[5 weeks	Linear	$y_{t(5)} = 0.35 x_{t(5)} + 93.13$	1.83	0.62
Lag	time	Autoregression	$y_{t(5)}+0.2362y_{t(5)-1}-0.0785 y_{t(5)-2}=0.35(0.2362x_{t(5)-1}-0.0785x_{t(5)-2})+93.13$	105.81	0.67
		Autoregression+ time effect	$y_{t(5)} = 1337 + 20.20t(5) - 0.011 dx_{t(5)}$ $y_{t(5)}^* = y_{t(5)} + 0.64y_{t(5)-1}$	1.47	0.95
		Vector Autoregression	$y_{t(5)} = 1.54 y_{t(5)\text{-}1}$ -0.02 $y_{t(5)\text{-}2}$ -0.68 $x_{t(5)\text{-}1}$ -0.04 $x_{t(5)\text{-}2} - 191.18$	0.35	0.96

Modeling results of N1 gene in gc/L for September 2020 to August 2021

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Best prediction models based on (a) N1 gene concentrations (gc/L) and (b) N2 gene concentrations (gc/L) with a 5-week lag time

Conclusions



Publication





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Thank you!

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