

CORONASTEP Report 76 (Week 22) SARS-CoV-2 Sewage Surveillance in Luxembourg

Summary

This report 76 presents the results of SARS-CoV-2 contamination of wastewater at the entrance of the 13 wastewater treatment plants analysed during the week 22 of 2021.

The SARS-CoV-2 RNA flux measured in wastewater treatment plants shows a moderate national prevalence of the virus. The flux has around 1 to 2 x 10¹¹ RNA copies per day per 100,000 equivalent-inhabitants, with a continuing downward trend for several weeks. The SARS-CoV-2 contamination of wastewater seems to be stabilized at this moderate level since the three last sampling dates.

At the individual treatment plant scale, the trend is similar with several treatment plants having a SARS-CoV-2 flux near the detection limit of the test, or even negative.

Table 1 – National level of SARS-CoV-2 contamination of wastewaters in Luxembourg.



Dark green: negative samples for SARS-CoV-2 gene E (-), Green to red: positive samples for SARS-CoV-2 gene E. The intensity of the color is related to the national SARS-CoV-2 flux (RNA copies / day / 100 000 equivalent inhabitants).

National Contamination	Week	National	JOOM
Level		Level	V
	Week 49-2		Week 3
	Week 50-1		Week 7
	Week 50-2		Week 9
	Week 51-1		Week 11
	Week 51-2		Week 14
	Week 51-2		Week 15
	Week 52		Week 16
	Week 53		Week 17
	Week UI-1		Week 18
	Week 02-2		Week 19
	Week 02-2		Week 20
	Week 03-1		Week 21
	Week 03-2		Week 22
	Week 04-1		Week 23
	Week 04-2		Week 24
	Week 05-1		Week 25
	Week 06-1		Week 26
	Week 06-2		Week 27
	Week 07-1		Week 28
	Week 07-2		Week 29
	Week 08-1		Week 30
	Week 08-2		Week 31
	Week 09-1		Week 32
	Week 09-2		Week 33
	Week 10-1		Week 34
	Week 10-2		Week 35
	Week 11-1		Week 36
	Week 11-2		Week 37
	Week 12-1		Week 38
	Week 12-2		Week 39
	Week 13-1		Week 40
	Week 13-2		Week 41
	Week 14-2		Week 43
	Week 15-1		Week 44-1
	Week 15-2		Week 44-2
	Week 16-1		Week 45-1
	Week 16-2		Week 45-2
	Week 17-1		Week 45-3
	Week 17-2		Week 46-1
	Week 18-1		Week 46-2
	Week 18-2		Week 46-3
	Week 19		Week 47-1
	Week 20-1		Week 47-2
	Week 20-2		Week 48-1
	Week 21		Week 48-2
	Week 22-1		Week 48-3
	VCCN 22-2		Week 49-1



Figure 1a – RT-qPCR quantification time-course monitoring of SARS-CoV-2 (E gene) in Luxembourgish wastewater samples from December 2019 to May 2021. Grey squares: daily-confirmed cases for Luxembourgish residents (https://data.public.lu/fr/datasets/donnees-covid19/), Blue dots: cumulative SARS-CoV-2 flux (RNA copies / day / 100 000 equivalent inhabitants).

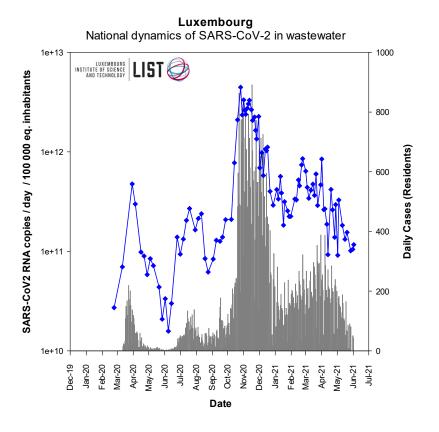


Figure 1b – Close-up of Figure 1a showing results from September 1st on.

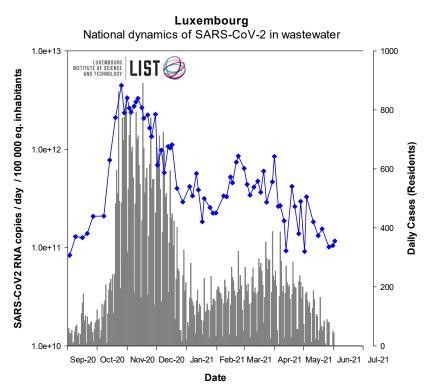




Table 2 - Level of SARS-CoV-2 contamination of each analyzed wastewater treatment plant in Luxembourg during the second wave. BEG: Beggen, BET: Bettembourg, SCH: Schifflange, BLE: Bleesbruck, MER: Mersch, PET: Pétange, HES: Hespérange, ECG: Echternach, UEB: Uebersyren, GRE: Grevenmacher, TRO: Troisvierges, BOE: Boevange sur Attert, WIL: Wiltz



Dark green: negative samples for SARS-CoV-2 gene E (-), Green to red: positive samples for SARS-CoV-2 gene E. The intensity of the color is related to the RT-qPCR signal (Ct values) Grey boxes: no data

																				2	2021	<u> </u>																			
WWTP	Week 01-1	Week 01-2	Week 02-1	Week 02-2	Week 03-1	Week 03-2	Week 04-1	Week 04-2	Week 05-1	Week 06-1	Week 06-2	Week 07-1	Week 07-2	Week 08-1	Week 08-2	Week 09-1	Week 09-2	Week 10-1	Week 10-2	Week 11-1	Week 11-2	Week 12-1	Week 12-2	Week 13-1	Week 13-2	Week 14-1	Week 14-2	Week 15-1	Week 15-2	Week 16-1	Week 16-2	Week 17-1	Week 17-2	Week 18-1	Week 18-2	Week 19	Week 20-1	Week 20-2	Week 21	Week 22-1	Week 22-2
BEG																																									
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SCH																																									
BLE																																									
MER																																									
PET																																									
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Figure 2a – RT-qPCR quantification time-course monitoring of SARS-CoV-2 (E gene) in the four most impacted wastewater treatment plants from March 2020 to June 2021. Grey squares: daily-confirmed cases for the contributory area of each wastewater treatment plant, dots: SARS-CoV-2 flux (RNA copies / day / 10 000 equivalent inhabitants).

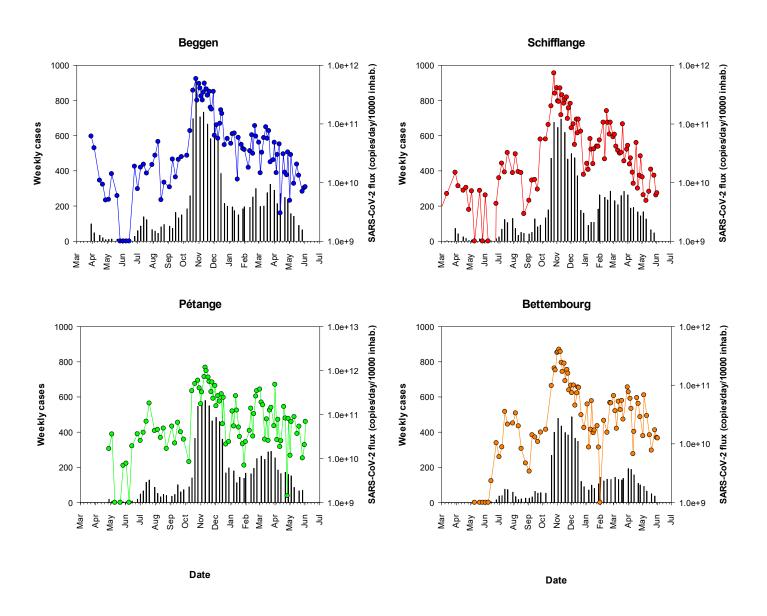




Figure 2b – Close-up of Figure 2a showing results from September 1st, 2020 on.

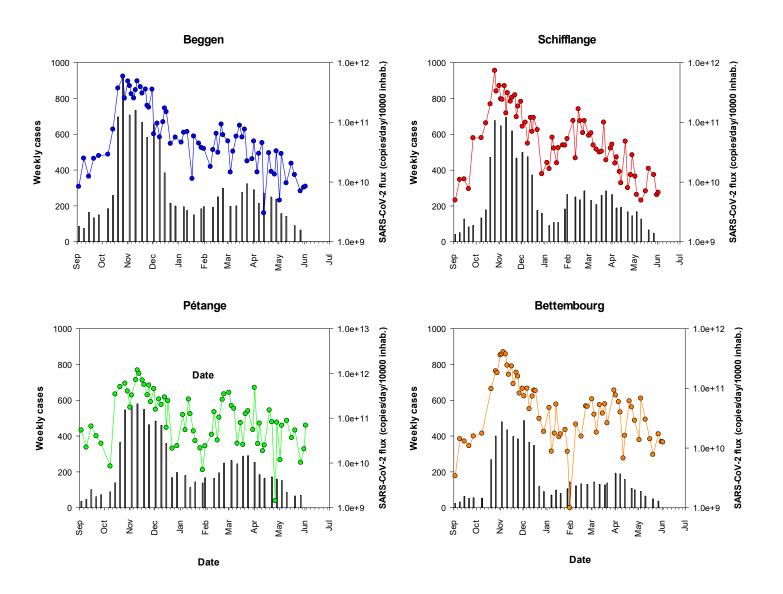
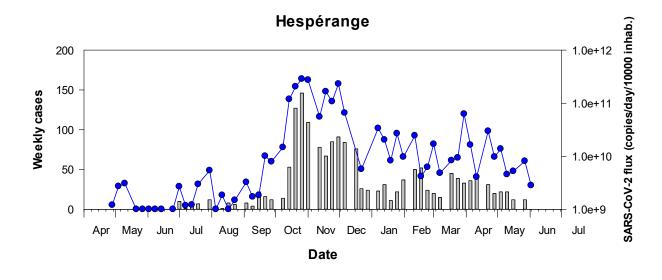
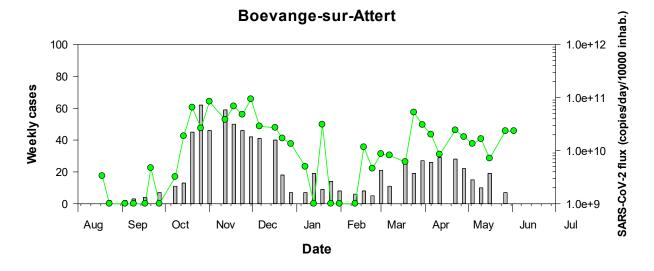




Figure 3a – RT-qPCR quantification time-course monitoring of SARS-CoV-2 (E gene) in Hespérange, Mersch and Boevange-sur-Attert wastewater treatment plants from March 2020 to June 2021. Grey squares: daily-confirmed cases for the contributory area of each wastewater treatment plant, dots: SARS-CoV-2 flux (RNA copies / day / 10 000 equivalent inhabitants).





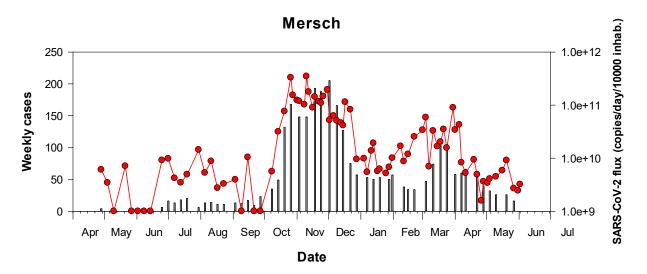


Figure 3b – Close-up of Figure 3a showing results from September 1st, 2020 on.

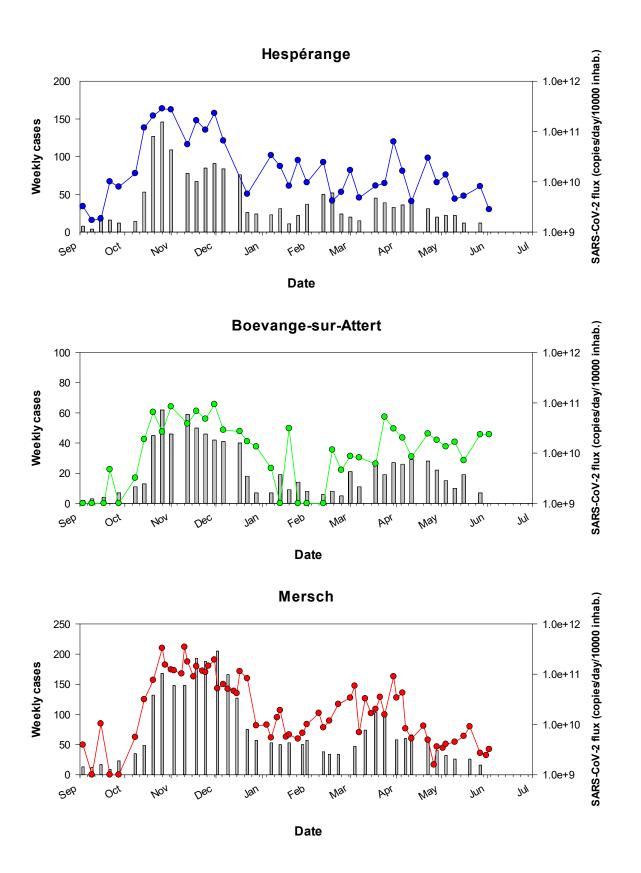
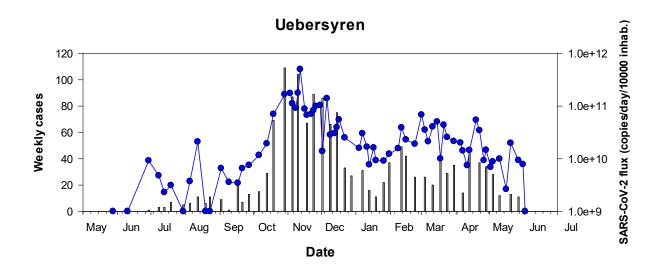
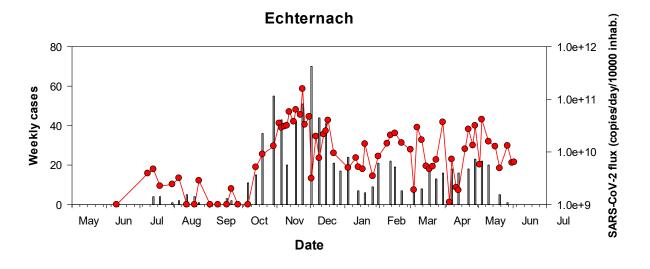




Figure 4a – RT-qPCR quantification time-course monitoring of SARS-CoV-2 (E gene) in SIDEST wastewater treatment plants from March 2020 to June 2021. Grey squares: daily-confirmed cases for the contributory area of each wastewater treatment plant, dots: SARS-CoV-2 flux (RNA copies / day / 10 000 equivalent inhabitants).





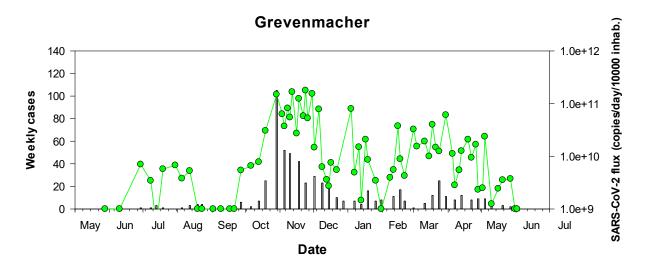
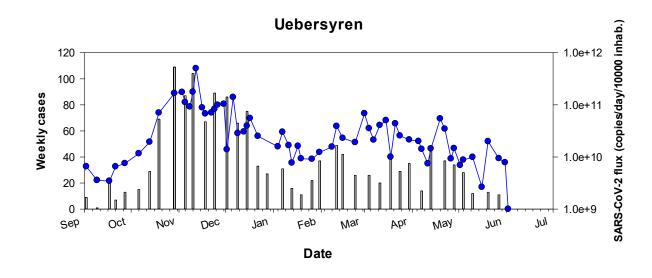
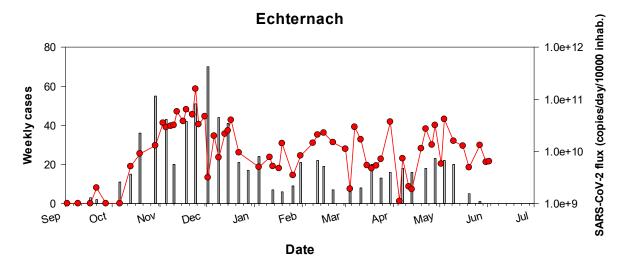


Figure 4b – Close-up of Figure 4a showing results from September 1st, 2021 on.





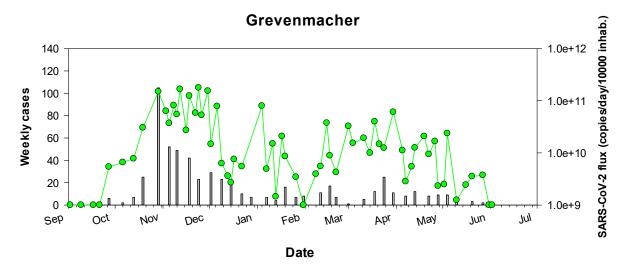
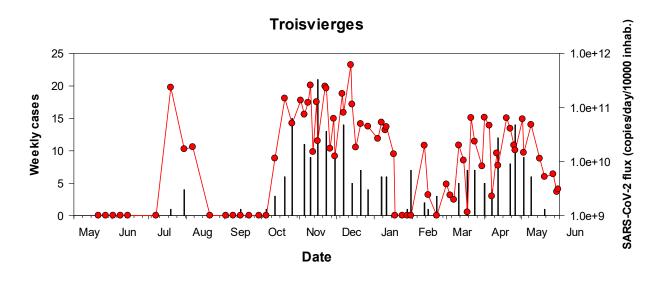
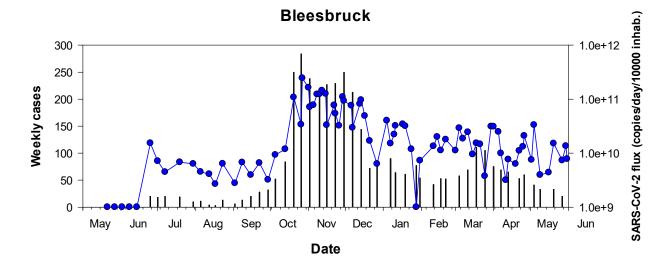




Figure 5a – RT-qPCR quantification time-course monitoring of SARS-CoV-2 (E gene) in SIDEN wastewater treatment plants from March 2020 to June 2021. Grey squares: daily-confirmed cases for the contributory area of each wastewater treatment plant, dots: SARS-CoV-2 flux (RNA copies / day / 10 000 equivalent inhabitants)





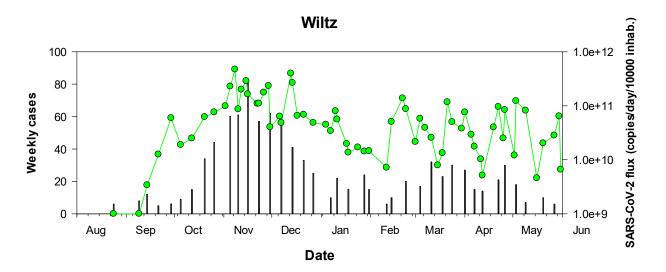


Figure 5b – Close-up of Figure 5a showing results from September 1st, 2021 on.

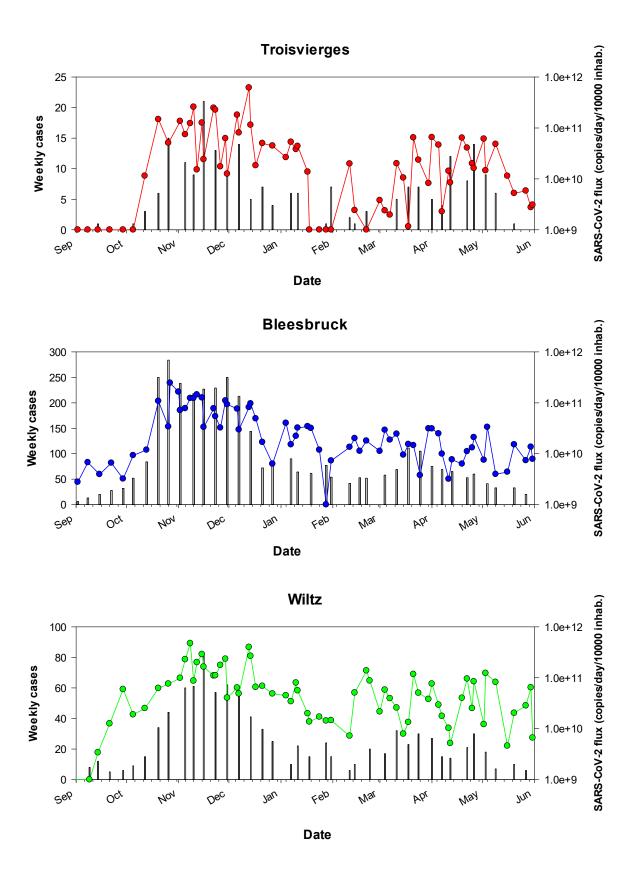




Table 3- Timing of sewage sampling since the beginning of the CORONASTEP study

		20	19																20	020																								2	2021								
WWTP		Week 41 Week 43	Week 46 Week 51	Week 3	Week 7	Week 11	Week 14	Week 15 Week 16	* *	Week 18			Week 22	Week 23 Week 24	Week 25	Week 26	Week 28	Week 29				Week 34 Week 35	Week 36	Week 37	Week 38 Week 39	Week 40	Week 41	Week 42			Week 46 Week 47		Week 49					Week 02 Week 03	Week 04	Week 05	Week 06 Week 07	Week 08	Week 09	Week 10 Week 11	Week 12		Week 14 Week 15		Week 17	Week 18			Week 22
Beggen	139731						1	1 1	1	1	1 1	1	1	1 1	1	1 1	1	1	1 1	1	1	1 1	1	1 1	1 1	1	1 1	1 1	2	3	3 2	3	2	2 3	1	1	2	1 2	2	2	2 2	2	2	2 2	2	2	2 2	2 2	2	2 1	. 2	1	2
Bettembourg	53606										1	1	1	1 1	1	1 1	1	1	1 1	1	1	1 1	1	1 1	1 1	1	1	1	2	3	3 2	3	2 :	2 3	1	1	2	2 2	2	2	1 2	2	2	2 2	2	2	2 2	2 2	1	2 1	. 2	1	2
Schifflange	68143	1 1	1 1	1	1 1	1	1	1 1	. 1	1	1 1	1	1	1 1	1	1 1	1	1	1 1	1	1	1 1	1	1 1	1 1	1	1 1	1 1	2	3	3 2	3	2	2 3	1	1	2 :	2 2	2	2	2 2	2	2	2 2	2	2	2 2	2 2	2	2 1	. 2	1	2 1
Bleesbrück	30930											1	1	1 1	1	1 1	1		1 1	1	1	1 1	1	1 1	1 1	1	1 1	1 1	2	3	3 2	3	2	2 3	1	1	2	2 2	2	2	1 2	2	2	2 2	2	2	2 2	2 2	2	2 1	. 2	1	2
Mersch	30473									1	1 1	1	1	1 1	1	1 1	1	1	1 1	1	1	1 1	1	1 1	1 1	1	1 1	1 1	2	2	3 2	3	2	2 3	1	1	2	2 2	2	2	2 2	2	2	2 2	2	2	2 1	1 2	2	2 1	. 2	1	2
Pétange	59481	1 1	1 1	1	1 1	1				1	1 1	1	1	1 1	1	1 1	1	1	1 1	1	1	1 1	1	1 1	1 1	1	1 1	1 1	2	2	3 2	3	2	2 3	1	1	2	2 2	2	2	2 2	2	2	2 2	2	2	2 2	2 2	2	2 1	. 2	1	2
Hespérange	15479									1	1 1	1	1	1 1	1	1 1	1	1	1 1	1	1	1 1	1	1 1	1 1	1	1 1	1 1	1	1	1 1	2	1	1 1	1	0	1	1 1	1	1	1 1	1	1	1 1	1	1	1 1	1 1	1	1 1	. 1	1	1
Echternach	7499													1			1	1	1 1	1	1	1 1	1	1 1	1 1	1	1 1	1 1	1	2	3 2	3	2	2 3	1	0	1 :	2 2	1	2	2 2	2	2	2 2	2	1	2 2	2 2	2	2 1	. 2	1	2
Uebersyren	18600												1	1		1	1	1	1 1	1	1	1 1	1	1 1	1 1	1	1 1	1 1	1	2	3 2	3	2	2 3	1	0	2	2 2	1	2	2 2	1	2	2 2	2	1	2 2	2 2	2	2 1	. 2	1	2
Grevenmacher	9835												1	1		1	1	1	1 1	1	1	1 1	1	1 1	1 1	1	1 1	1 1	1	2	3 2	3	2	2 3	1	0	2	2 2	1	2	2 2	2	2	1 2	2	1	2 2	2 2	2	2 1	. 2	1	2
Troisvierges	3411											1	1	1 1	1		1		1 1	1		1	1	1 1	1 1	1	1 1	1 1	1	2	3 2	3	2	2 3	1	1	2	2 2	2	2	1 2	2	2	2 2	2	2	2 2	2 2	2	2 1	. 2	1	2
Boevange sur Attert	1170																					1 1	1	1 1	1 1	1	1 1	1 1	1	1	1 1	2	1	1 1	1	1	1 :	1 1	1	1	1 1	1	1	1 1	. 1	1	1 1	1 1	1	1 1	. 1	1	1
Wiltz	6944																					1		1 1	1 1	1	1 1	1 1	1	2	3 2	3	2	2 3	1	1	2	2 2	2	2	1 2	2	2	2 2	2	2	2 2	2 2	2	2 1	. 2	1	2
Total	445302	2 2	2 2	2	2 2	2	2	2 2	2	5	5 6	8	10	8 11	8	9 7	11	9 1	1 11	11	10 1	2 12	12	13 1	3 13	13	13 1	2 13	19	28 3	35 24	37	24 2	24 35	13	9 :	23 2	3 24	21	24 2	20 24	23	24	23 24	1 24	21	24 23	3 24	23	24 1	3 24	13	24 1



Materials and Methods

Sewage samples

From March 2020 to June 2021, up to thirteen wastewater treatment plants (WWTPs) were sampled at their inlet according to the planning presented in Table 3. The operators of the WWTPs collected a 24-h composite sample according to their routine sampling procedure. Composite sample was stored at 4°C until sample processing.

Sample processing

The samples were transported to the laboratory at 4°C and viral RNA was isolated on the day of sampling. Larger particles (debris, bacteria) were removed from the samples by centrifugation at 2,400 x g for 20 min at 4°C. A volume of 120 mL of supernatant was filtered through Amicon® Plus-15 centrifugal ultrafilter with a cut-off of 10 kDa (Millipore) by centrifugation at 3,220 x g for 25 min at 4°C. The resulting concentrate was collected and 140 μ L of each concentrate was then processed to extract viral RNA using the QIAamp Viral RNA mini kit (Qiagen) according to the manufacturer's protocol. Elution of RNA was done in 60 μ L of elution buffer.

Real-time One-Step RT-PCR

Samples were screened for the presence of *Sarbecovirus* (*Coronaviridae*, *Betacoronaviruses*) and/or SARS-CoV-2 virus RNA by two distinct real-time one-step RT-PCR assays, trageting the E gene (Envelope small membrane protein) and the N gene (nucleoprotein). The E gene real-time RT-PCR can detect *Sarbecoviruses*, i.e. SARS-CoV, SARS-CoV-2 and closely related bat viruses. In the context of the COVID19 pandemic, it can be assumed that only SARS-CoV-2 strains will be detected by this assay given that SARS-CoV virus has been eradicated and other bat viruses do not commonly circulate in the human population. The E gene assay is adapted from Corman et al. [17]. The N gene real-time RT-PCR assay (N1 assay) specifically detects SARS-CoV-2 virus. It is adapted from the CDC protocol¹. The two primers/probe sets are presented in Table 3. The RT-qPCR protocols and reagents were all provided by the LIH.

Table 4 – RT-qPCR primer-probe sets

Target	Primer name	Primer sequence (5' to 3')	References
E gene	E_Sarbeco_F1	5-ACAGGTACGTTAATAGTTAATAGCGT-3	Corman et al.,
	E_Sarbeco_R2	5-ATATTGCAGCAGTACGCACACA-3	2020
	E_Sarbeco_P1	5'-FAM-ACACTAGCCATCCTTACTGCGCTTCG-BHQ1	
N gene	2019-nCoV_N1_Fw	5'-GAC CCC AAA ATC AGC GAA AT-3'	CDC, 2019
	2019-nCoV_N1_Rv	5'-TCT GGT TAC TGC CAG TTG AAT CTG-3'	
	2019-nCoV_N1 Probe	5'-FAM-ACC CCG CAT TAC GTT TGG TGG ACC-BHQ1-3'	

Each reaction contained 5 μ L of RNA template, 5 μ L of TaqPath 1-step RT-qPCR MasterMix (A15299, Life Technologies), 0.5 μ L of each primer (20 μ M) and probe (5 μ M) and the reaction volume was adjusted to a final volume of 20 μ L with molecular biology grade water. Thermal cycling reactions were carried out at 50 °C for 15 min, followed by 95 °C for 2 min and 45 cycles of 95 °C for 3 sec and 58 °C (E gene) or 55 °C (N gene) for 30 sec using a Viia7 Real-Time PCR Detection System (Life Technologies). Reactions were considered positive (limit of detection – LOD) if the cycle threshold (Ct value) was below 40 cycles.

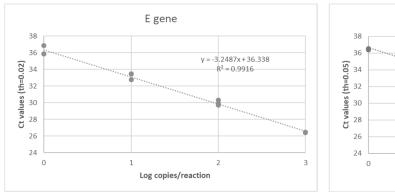
¹ https://www.cdc.gov/coronavirus/2019-ncov/downloads/rt-pcr-panel-primer-probes.pdf



Controls

A non-target RNA fragment commercially available (VetMAX™ Xeno™ IPC and VetMAX™ Xeno™ IPC Assay, ThermoFischer Scientific) was added to the viral RNA extract from sewage concentrates as an internal positive control (IPC). This IPC-RNA is used to control the performance of the RT-qPCR (E gene) and to detect the presence of RT-qPCR inhibitors.

Viral RNA copies quantification of both targeting genes in wastewater samples was performed using RT-qPCR standard curves generated using EDX SARS-CoV-2 Standard (Biorad). This standard is manufactured with synthetic RNA transcripts containing 5 targets (E, N, S, ORF1a, and RdRP genes of SARS-CoV-2, 200,000 copies/mL each). Using such a standard, the limits of quantification (LOQ) of both RT-qPCR assays were estimated to 1 RNA copy per reaction (Figure 6).



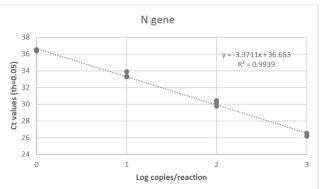


Figure 6 – RT-qPCR standard curves established for both target genes (E gene and N gene) of SARS-CoV-2 using a commercially available standard (Biorad).

Data interpretation

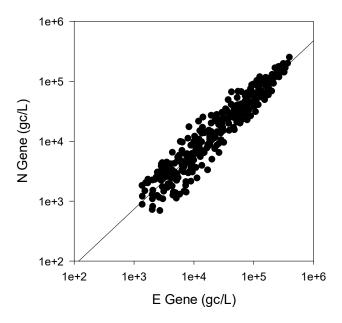
A sample is declared positive for the presence of SARS-CoV-2 if both targets (E and N gene) are detected with Ct values less than or equal to the LOQ. If only one target is detected or if target genes are detected with Ct values between the LOD and the LOQ, samples are reported as presumptive positive (+/-). A sample is declared negative when no target genes are detected (Ct values superior to the LOD).

In case of presumptive positive, sample is tested again using another RT-qPCR detection assay (Allplex 2019-nCoV Assay, Seegene). This commercially available detection kit is a multiplex real-time RT-PCR assay for simultaneous detection of three target genes of SARS-CoV-2 in a single tube. The assay is designed to detect RdRP and N genes specific for SARS-CoV-2, and E gene specific for all *Sarbecovirus* including SARS-CoV-2.

As shown in Figure 7, a highly significant correlation (Pearson Correlation, R^2 =0.964, p = 5.979.10⁻²⁴) was obtained between the SARS-CoV-2 RNA concentrations estimated using the E gene and the N gene, respectively. Therefore, only the E gene results were presented in this report.



Figure 7 - Relationship between the SARS-CoV-2 RNA concentration (RNA copies / L of wastewater) estimated by the both distinct RT-qPCR systems targeting the E and N gene, respectively (n=415),



Acknowledgments

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