

Surveillance of SARS-CoV-2 in the environment and animal samples of the Huanan Seafood Market

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4 Running title:

- 5 Prevalence of SARS-CoV-2 in Huanan Seafood Market
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Abstract

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Emerging in December 2019, coronavirus disease 2019 (COVID-19) eventually became a pandemic and has posed a tremendous threat to global public health. However, the origins of SARS-CoV-2, the causative agent of COVID-19, remain to be determined. It has reported that a certain number of the early case clusters had a contact history with Huanan Seafood Market. Therefore, surveillance of SARS-CoV-2 within the market is of vital importance. Herein, we presented the SARS-CoV-2 detection results of 1380 samples collected from the environment and the animals within the market in early 2020. By SARS-CoV-2-specific RT-qPCR, 73 environmental samples tested positive for SARS-CoV-2 and three live viruses were successfully isolated. The viruses from the market shared nucleotide identity of 99.980% to 99.993% with the human isolate HCoV/Wuhan/IVDC-HB-01. In contrast, no virus was detected in the animal swabs covering 18 species of animals in the market. The SARS-COV-2 nucleic acids in the positive environmental samples showed significant correlation of abundance of *Homo sapiens* with SARS-CoV-2. In summary, this study provided convincing evidence of the prevalence of SARS-CoV-2 in the Huanan Seafood Market during the early stage of COVID-19 outbreak.

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Keywords:

- 61 COVID-19, SARS-CoV-2, Huanan Seafood Market, zoonotic origin, cold chain,
- 62 high-throughput sequencing, virus isolation

Infections with novel human coronavirus 2019 (HCoV-19) (1, 2), named as severe acute respiratory syndrome coronavirus 2 (SARS-CoV-2) by International Committee on Taxonomy of Viruses (ICTV) (3), can result in coronavirus disease 2019 (COVID-19), characterized by various clinical outcomes from asymptomatic infections to severe pneumonia and even death (4-6). Globally, as of September 10th 2021, there have been 223,022,538 confirmed cases of COVID-19, including 4,602,882 deaths. No sign hints the pandemic has declined. (covid19.who.int).

Early human cases with COVID-19 were first reported in late December 2019 from Wuhan, China with pneumonia of unknown etiology (PUE), and a majority of them were found to be linked to Huanan Seafood Market (HSM) in Wuhan (6, 7), where various animal meats and exotic seafoods were available for purchase. Therefore, HSM has been suspected to be the source of the COVID-19 pandemic (8). However, the possibility that some individuals outside the market had brought the virus into the market through human-to-human transmission or cold chain could not be ruled out, considering some of the early cases without epidemiological link to the market (7, 9). In addition, recent studies from different countries suggest that SARS-CoV-2 circulation preceded the initial detection of cases by weeks. Some of the suspected positive samples were detected even earlier than the first case in Wuhan, suggesting that potential circulation of the virus in other regions may have been missed(10-14).

Considering the findings that SARS-CoV-2 had high similarities with a few coronaviruses derived from bats in Asian countries including China, Japan, Cambodia, Thailand and Laos, scientists have proposed that bats might be the original source of SARS-CoV-2 (1, 15-20). However, it is yet a mystery whether another animal might act as an intermediate host to have facilitated the virus spillover between bats and humans (21). One of such important findings was the discovery of SARS-CoV-2 related coronaviruses from pangolins that harbored highly similar receptor binding domain with that of SARS-CoV-2 (22-24). Despite that pangolins might be involved

in the ecology of coronaviruses, whether the pangolins are the intermediate hosts is still fairly controversial. A recent study documented the animal species in HSM between May 2017 and November 2019 and noted that no pangolins or bats were traded (25). Thus far, the origins of SARS-CoV-2 (26, 27) and the role of HSM in the origins and spread of SARS-CoV-2 remain poorly understood.

The HSM is located in the Jianghan District, a downtown of Wuhan, and is approximately 800 m far from Hankou Railway Station, a major railway travel hub. It occupies >50,000 m², with 678 stalls located close to each other in an extremely crowded condition (Fig. 1A). The market is separated into two zones, eastern and western zone, with seafood and animals mainly sold in the western zone and livestock meat in the eastern zone. Among the 678 stalls of the market, 10 domesticated wildlife animal-selling stalls (1.5%) were identified, located in the south-western corner of western zone (8/10) and the north-western corner of eastern zone (2/10), respectively (Fig. 1A). According to sales records, during late December 2019, animals or animal products were sold in these 10 animal stalls and animals included snakes, avian species (chickens, ducks, gooses, pheasants and doves), Sika deer, badgers, rabbits, bamboo rats, porcupines, hedgehogs, salamanders, giant salamanders, bay crocodiles and Siamese crocodiles, etc., among which snakes, salamanders and crocodiles were traded as live animals.

The market was closed in the morning of January 1st, 2020 shortly after the identification of the PUE. At the same time, in order to investigate the potential introduction of SARS-CoV-2 into the market, Chinese Center for Disease Control and Prevention (China CDC) dispatched an epidemiological team, together with experts from Hubei Provincial CDC and Wuhan Municipal CDC, to the HSM to collect environmental samples in the early morning of January 1st, 2020 (Fig. 1B). From January 1st, 2020 to March 2nd, 2020, a total of 923 environmental samples from different locations within and around this market and 457 animal samples including

animal bodies, stray animals and their feces were collected, with some stray animals sampled until March 30th (Extended Data Table 1), which may reflect the profile of SARS-CoV-2 contamination in the market during the early phase of the outbreak. After the closure of the stalls in the market, the outside surface of the rolling shutter doors of the stalls and the corridors was disinfected (with 1% bleach mixed with water) throughout January and February 2020. The goods inside the stalls were completely cleared and disinfected until early March 2020.

Out of the 923 environmental samples collected in and around the market, 73 were found to be positive for SARS-CoV-2 with positive rate of 7.9% through the nucleic acid test (NAT), with Cycle threshold (*Ct*) values of real-time polymerase chain reaction (PCR) ranging from 23.9 to 41.7 (Table 1). Among the 828 samples inside the HSM, 64 samples (7.7%) were positive. For the 14 samples from warehouses related to the HSM, five tested positive. Among the 51 sewerage wells in the surrounding areas, three tested positive (Table 1). Notably, one out of the 30 environmental samples from other markets in Wuhan tested positive.

For these 64 SARS-CoV-2 positive samples inside HSM, 87.5% (56/64) were collected in the western zone of the market, in particular streets from no. 1 to 8, with 71.4% (40/56) positive samples identified herein (Fig. 1A). Of the 110 samples collected from sewers or sewerage wells in the market, 24 samples were positive for SARS-CoV-2 nucleic acid. All the four sewerage wells in the market tested positive. This suggested that either contaminated sewage may have played a role in the cluster of cases in the market or that the infected people in the market contaminated the sewage.

The merchants' activities were assessed against the NAT results of the environmental samples. The sampling covered 19.8% (134/678) of the vendors in the market (95% confidence interval (CI): 16.8-23.0%). Of the positive samples, 44 were distributed

among 21 vendors in the market, 19 of whom were located in the west zone and the remaining two located in the east area (Fig. 1A). Some vendors sold more than one product type, leading to differences in the denominators. While the results provide some indication of association of cases with different products, but no significant differences were observed between different vendors, including cold-chain products (18.4%, 16/87, 95% CI: 10.9-28.1%), aquatic products (17.8%, 13/73, 95% CI: 9.8-28.5%), seafood products (11%, 6/56: 95% CI: 4-21.9%), poultry (22%, 8/37: 95% CI: 9.8-38.2%), livestock (14%, 5/36: 95% CI: 4.7-29.5%), wildlife products (11%, 1/9: 95% CI: 0.3-48.2%) and vegetables (25%, 2/8: 95% CI: 3.2-65%) (Extended Data Fig. 1). These results suggested that SARS-CoV-2 might have been circulating in the market, especially the western zone, for a period of time in December 2019, leading to an extensive distribution of the virus within the market, which might have been facilitated by the crowded buyers and the contaminated environment.

The 457 animal samples mainly collected between January 1st and March 2nd, 2020 included 188 individuals belonging to 18 species (with some stray animals sampled until March 30th) (Table 2). The sources of the samples include unsold goods kept in refrigerators and freezers in the stalls of HSM, and goods kept in warehouses and refrigerators related to the HSM. Samples from stray animals in the market were also collected, i.e. swab samples from 10 stray cats, 27 cat feces, one dog, one weasel, and 10 rats. All the 457 animal samples tested negative for SARS-CoV-2 nucleic acid, suggesting that the animal infections with SARS-CoV-2 might be rare in the market.

To determine whether there was live virus in the HSM, we inoculated 27 SARS-CoV-2 positive environmental samples collected on January 1st, 2020 into cell lines, including Vero E6 and Huh7.5 cells. Cytopathic effects (CPE) were observed 3 days post inoculation with sample F13 on Vero E6 cells. CPE was also observed 5 days post inoculation on Huh7.5 cells. We performed RT-qPCR on the supernatant of

the inoculated Vero E6 and Huh7.5 cells. The supernatants were found to be SARS-CoV-2 positive, with Ct values of 15.11 (Vero E6) and 14.44 (Huh7.5) for the ORF1ab gene, and 13.98 (Vero E6) and 13.59 (Huh7.5) for the N gene, respectively. We further used transmission electron microscopy to observe the culture supernatant and ultra-thin section of Vero E6 cells 5 days post inoculation. The electron micrographs showed that virus particles were present in both the supernatant and the cells. Negative-stained virus particles were generally spherical, pleomorphic and 60-140 nm in diameter. Spike protrusions were observed around the particles in a crown (corona) shape. In the ultra-thin cultured cell sections, a group of virus particles could be seen outside the cells, and sheets of virus particles could also be observed inside the cells (Fig. 2). Totally, live viruses were isolated from samples F13, F54, and B5, which were the only three samples with Ct values <30 in the NAT. F54 and B5 were swab samples of the ground and F13 were swab samples of the wall. Notably, samples F13 and F54 were from the stalls with confirmed patients. All the results of successful virus isolation and the Ct values of the original samples revealed the existence of live SARS-CoV-2 with high titers in the environment of HSM.

We further performed high-throughput sequencing on the Miseq platform and successfully obtained seven complete SARS-CoV-2 genome sequences, including three sequences from three environmental samples (F13, F54, and A20), and four sequences from cell supernatants of F13, F54, and B5 (Fig. 3). The genome sequences of two environmental samples, F13 and F54, were found to be highly identical to the reference strain HCoV/Wuhan/IVDC-HB-01 (WH01, sequence identity of 99.993%) and completely identical to the human stain Wuhan-Hu-1 (GenBank: NC_045512) (Fig. 3A). The genome sequence of the isolated virus from environmental sample F54 had two more synonymous mutations compared to HCoV/Wuhan/IVDC-HB-01, with sequence identity of 99.987% (Fig. 3A). Therefore, the SARS-CoV-2 sequences from environmental samples were highly similar to the clinical strains obtained during the early stage of the COVID-19 outbreak.

Previously, SARS-CoV-2 has been proposed to be classified into two major genotypes based on the two highly linked single nucleotide polymorphisms (SNPs): S (8782T and 28144C) and L (8782C and 28144T), with S most likely to be the ancestral lineage, because all of the SARS-CoV-2 related coronaviruses from bats and pangolins possessed 8782T and 28144C (28). Phylogenetic analysis revealed that most of the environmental strains belong to the L lineage and they fall into the basal positions of the L lineage (Fig. 3B). However, the environmental sample A20 fell within the S lineage in the tree (Fig. 3B, Extended Data Table 2).

To explore the potential origins of the SARS-COV-2, we conducted RNA-seq analysis using 27 SARS-CoV-2 positive environmental samples collected on January 1st, 2020 from the HSM. We analyzed the correlation of SARS-CoV-2 and the abundance of other species. The abundance of *Homo sapiens* showed the correlation to SARS-CoV-2 (Figure 4), which highly suggests the SARS-CoV-2 might have derived from *Homo sapiens* in the HSM. No animals were concluded, implying that no animal host of SARS-CoV-2 can be deduced.

In summary, SARS-CoV-2 RNA was detected in stalls in the western zone of HSM, suggesting the prevalence of SARS-CoV-2 in the market. Thus, the market might have acted as an amplifier due to the high number of visitors every day, causing many initially identified infection clusters in the early stage of the outbreak as indicated in the Report of WHO-convened global study of origins of SARS-CoV-2 (10). In addition, live SARS-CoV-2 viruses also existed in the environmental samples. However, no SARS-CoV-2 was detected in the animal samples from the market. Definitely, more work involving international coordination is needed to investigate the real origins of SARS-CoV-2(10), especially considering the SARS-CoV-2 positive results of samples collected in 2019 in retrospective studies of different countries (29, 30). Surveillance of wild animals using a viromic approach should be

enhanced to explore the natural and intermediate hosts for SARS-CoV-2 (8, 31), which would help to prevent future potential pandemics caused by animal-derived coronaviruses or alike.

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248 Author contributions

- 249 The study was designed by W.J.L, G.F.G. and G-Z.W. Samples were collected by
- 250 W.J.L., W.L, Z.J., X.H., J.W., F.W., G.W., K.Q., R.G., J.Z., M.L. and W.X.
- Experiment and data analysis were performed by W.J.L., P.L., Z.J., X.H., L.L., W.S.,
- 252 Y.T., S.Z., X.Z., J.W., F.W., G.W., Y.G., Z.X., Y.Z., J.S., Jing Z., W.Z., W-T.Z., B.Y.,
- J.S., M.Y., W-M.Z., Y.B., K.C., D.W., W.T., J.H. and W-B.X. The manuscript was
- written by W.J.L., P.L., W.S., Y.T., G.W., G.F.G. and G-Z.W.

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Competing interest declaration

No competing interest exists.

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333 Figure legends

- Figure 1. The distribution of the positive environmental samples in Huanan
- 335 **Seafood Market.**
- 336 A. As the place of the early cluster of COVID-19 patients, the Huanan Seafood
- Market is separated into eastern and western zones with the Xinhua Rd. between them.
- To detect for the presence of SARS-CoV-2 RNA, reverse transcription, quantitative
- polymerase chain reaction (RT-qPCR) was performed. The locations of the positive
- samples were marked in the map of the market within orange, while the location of
- the samples that the live viruses were isolated from were labeled with red. The map
- also shows locations of stalls where domesticated wildlife products were sold. B.
- 343 Timeline of environmental and animal samples collected within and around Huanan
- 344 Seafood Market.

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- Figure 2. The SARS-CoV-2 virus isolation from environmental samples of
- 347 Huanan Seafood Market.
- 348 The electron micrographs of the SARS-CoV-2 viruses isolated from the
- 349 environmental samples in Huanan Seafood Market. To determine whether
- 350 SARS-CoV-2 particles could be visualized from the cell supernatant and lysate, we
- used transmission electron microscopy to observe the culture supernatant and
- 352 ultra-thin section cells based from both VeroE6 and Huh7.5 cells. The electron
- micrographs showed that virus particles were present in both the supernatant (A, B)
- and the cells (C, D). Negative-stained virus particles were generally spherical,
- 355 pleomorphic and 60-140 nm in diameter. Spike protrusions were observed around the
- particles in a crown (corona) shape (A, B). In ultra-thin cultured cell sections, a group
- of virus particles can be seen outside the cell (C), and sheets of virus particles can also
- be observed inside the cells (D).
- Figure 3. The genomic analysis of SARS-CoV-2 sequenced from Huanan Seafood
- 360 Market.

Figure 4. Clustering of environmental samples in Huanan Seafood Market.

A, Correlation of SARS-COV-2 with other species in SARS-COV-2 positive cases.

The abundance of SARS-COV-2 and other species was used for analysis. Both Pearson's and Spearman's correlation values were plotted. B, Correlation of the abundance of SARS-COV-2 and *Homo sapiens*.

Tables
 Table 1. Information of the positive environmental samples from Huanan Seafood
 Market.

No.	Sample type	Sampling date	PCR	Ct	PCR target	NGS
E61	Ground	2020/1/1	+	36.04	ORF1ab/N	NA
A101	Surface of the door	2020/1/1	+	36.82	ORF1ab/N	NA
A14	Surface of packing bag for hairtail	2020/1/1	+	36.42	ORF1ab/N	NA
A15	Surface of the door	2020/1/1	+	35.51	ORF1ab/N	NA
A18	Shoe covers and soles	2020/1/1	+	33.79	ORF1ab/N	NA
A2	Ground	2020/1/1	+	35.07	ORF1ab/N	+
A20	Gloves	2020/1/1	+	32.48	ORF1ab/N	+
A33	Garbage truck	2020/1/1	+	34.46	ORF1ab/N	NA
A55	Ground	2020/1/1	+	34.84	ORF1ab/N	+
A61	Ground	2020/1/1	+	32.04	ORF1ab/N	NA
A63	Ground	2020/1/1	+	34.43	ORF1ab/N	+
A87	Surface of the door	2020/1/1	+	36.94	ORF1ab/N	NA
A88	Ground	2020/1/1	+	36.69	ORF1ab/N	NA
A90	Ground	2020/1/1	+	33.14	ORF1ab/N	NA
A96	Ground	2020/1/1	+	33.97	ORF1ab/N	NA
B17	Scale	2020/1/1	+	34.16	ORF1ab/N	NA
B5*	Ground	2020/1/1	+	29.32	ORF1ab/N	+
D32	Surface of a cart	2020/1/1	+	33.72	ORF1ab/N	NA
E48	Blood on the ground in front of the door	2020/1/1	+	35.93	ORF1ab/N	+
E7	Styrofoam desk in front of the door	2020/1/1	+	36.44	ORF1ab/N	+
F100	Ground	2020/1/1	+	34.72	ORF1ab/N	+
F13*	Surface of the wall	2020/1/1	+	23.85	ORF1ab/N	+
F33	Roller shutter	2020/1/1	+	34.13	ORF1ab/N	NA
F46	Ground	2020/1/1	+	31.8	ORF1ab/N	+
F54*	Ground	2020/1/1	+	25.8	ORF1ab/N	+
F98	Ground	2020/1/1	+	34	ORF1ab/N	+
G93	Sewage	2020/1/1	+	33.23	ORF1ab/N	NA
Q37	Inner surface of the freezer	2020/1/12	-	\	ORF1ab/N	+
Q61	Cart1	2020/1/12	-	\	ORF1ab/N	+
Q64	Cart2	2020/1/12	+	+	ORF1ab/N	+
Q68	Ground	2020/1/12	+	+	ORF1ab/N	+
Q69	Feather removal machine	2020/1/12	+	+	ORF1ab/N	+
Q70	Iron container in inner room	2020/1/12	-	\	ORF1ab/N	+
1-27-33	Water drain	2020/1/27	+	36	ORF1ab	NA
1-27-37	Water drain	2020/1/27	+	35	ORF1ab	NA
1-29-4	Water drain	2020/1/29	+	36	ORF1ab	NA
1-29-8	Water drain	2020/1/29	+	37	ORF1ab	NA
8-25-D	Ground inside the stalls	2020/2/3	+	35.9	ORF1ab	+
8-25-M1	Ground inside the stalls	2020/2/3	+	36.5	ORF1ab	+
WS-1-1	West sewage well 1	2020/2/5	+	36.00	ORF1ab	NA
WS-1-2	West sewage well 2	2020/2/5	+	38.00	ORF1ab	NA
WS-1-3	West sewage well 3	2020/2/5	+	34.01	ORF1ab	NA
WS-1-4	West sewage well 4	2020/2/5	+	38.00	ORF1ab	NA
WS-2-1	West sewage well 1	2020/2/5	+	37.30	ORF1ab	NA
WS-2-2	West sewage well 2	2020/2/5	+	37.65	ORF1ab	NA
WS-2-3	West sewage well 2	2020/2/5	+	36.72	ORF1ab	NA
WS-3-2	West sewage well 3	2020/2/5	+	37.60	ORF1ab	NA
WS-4-2	West sewage well 4	2020/2/5	+	37.10	ORF1ab	NA
zong-1	Water drain	2020/2/9	+	34.94	ORF1ab	NA
w-6-29-33	Water drain	2020/2/15	+	37.97	ORF1ab	NA
w-zong-1	Water drain	2020/2/15	+	36.54	ORF1ab	NA
EWS-2#-2	East sewage well 2	2020/2/15	+	35.32	ORF1ab	NA
EWS-3#-2	West sewage well 3	2020/2/15	+	36.05	ORF1ab	NA
WWS-1#	West sewage well 1	2020/2/15	+	34.44	ORF1ab	NA
WWS-1#-2	West sewage well 1	2020/2/15	+	33.63	ORF1ab	NA

WWS-1#-3	West sewage well 1	2020/2/15	+	33.58	ORF1ab	NA
WWS-2#-2	West sewage well 2	2020/2/15	+	37.44	ORF1ab	NA
WWS-2#-3	West sewage well 2	2020/2/15	+	36.88	ORF1ab	NA
wws-1#-0	West sewage well 1	2020/2/15	+	33.75	ORF1ab	NA
W-8-25-D1	Ground inside the stalls	2020/2/15	+	36.77	ORF1ab	NA
W-8-25-D2	Ground inside the stalls	2020/2/15	+	33.91	ORF1ab	NA
W-8-25-L	Container	2020/2/15	+	34.58	ORF1ab	NA
W-8-25-L2	Container	2020/2/15	+	37.16	ORF1ab	NA
C8	Wall inside the stalls	2020/2/20	+	32.59/-/37.34	RDRP/N/E	NA
SJ-D	Storehouse ground	2020/2/22	+	39.39/39.25/35.48	RDRP/N/E	NA
SJ-CS	Storehouse weight scale	2020/2/22	+	40.21/40.1/36.37	RDRP/N/E	NA
SJ-L3	Storehouse wire fence	2020/2/22	+	41.77/41.62/37.61	RDRP/N/E	NA
RLC-4	Storehouse bag surfaces	2020/2/22	+	41.71/41.53/37.31	RDRP/N/E	NA
RLC-3	Storehouse bag surfaces	2020/2/22	+	36.18/36.05/32.36	RDRP/N/E	NA

Note: Four positive samples were not included in this table. One sample was collected from other market in Wuhan and 3 collected from sewerage wells in surrounding areas.

Table 2. The animal samples collected in the Huanan Seafood Market.

Species	Animal number	Sample number	RT-PCR positive	Remarks
			number	
Rabbit/Hares	52	104	0	
Stray cat	27	80 ^a	0	Including faeces
Snake	40	80	0	
Hedgehog	16	67	0	
Muntjac	6	18	0	
Dog	7	17	0	Including one
				stray dog
Badger	6	16	0	
Bamboo rat	6	15	0	
Rat	10	12	0	Captured around
				the market
Pig	NA^c	6 ^b	0	
Chicken	5	5	0	
Chinese giant	3	5	0	
salamander				
Crocodile	2	4	0	
Wild boar	2	4	0	
Soft-shelled	2	3	0	
turtle				
Weasel	1	2	0	Captured around
				the market
Fish	2	2	0	
Sheep	1	1	0	
Others	NAc	16	0	
Rabbit/Hares	52	104	0	
Total	188	457	0	

^a Six of the cats were from the Huanan Seafood Market.

³⁷⁶ b Other markets.

^{377 °} Not applicable.

Extended data methods

Sample collection

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Huanan Seafood Market (HSM) was closed on early morning of January 1st 2020 and at the same time, China CDC began collecting environmental and animal samples. Staff from China CDC entered the market about 30 times before the market's final clean-up on 2 March 2020, with some stray animals sampled outside the market until March 30th. Environmental samples in the HSM were collected to represent exhaustively as possible, from a wide diversity of surfaces, animals and products (Table 1 and 2) according to different sampling principles, as described in detail in the Joint Report of WHO-convened Global Study of Origins of SARS-CoV-2: China Part (10).The principles and ranges of in-market sampling covered: (1) environmental samples from stalls related to early cases; (2) environmental samples from doors and floors of all stalls in the blocks where the early cases were located; (3) environmental samples in the east wing of the market were collected according to blocks; (4) transport carts, trash cans and similar objects; (5) environmental samples from stalls that sold livestock, poultry, farmed wildlife (also called "domesticated wildlife" or "domesticated wildlife products" in this report); (6) samples of sewage and silt from drainage channels and sewerage wells; (7) stray cats, rats and other potential vector animals in the market; (8) animal products and other commodity samples kept in the cold storages and refrigerators in the market; (9) the market's ventilation and air-conditioning system; and (10) public toilets, public activity rooms and other places where people gathered in the market. The investigators used full personal protective equipment during the sampling in the market. Commercial products of swabs and virus preservation solution were used for the sampling (Disposable Virus Sampling Tube, V5-S-25, Shen Zhen Zi Jian Biotechnology Co., Ltd., Shenzhen, China). For environmental samples, sampling swabs were applied to smear the floors, walls or surfaces of objects and then preserved them in virus preservation solution.

For animal samples, depending on the type of animal and whether it was alive or frozen, pharyngeal, anal, body surface and body cavity swabs or tissue samples were collected for nucleic acid testing (NAT). Generally, for alive animal and frozen full bodies, three samples, including pharyngeal, anal, and body surface swabs were collected for each animal individuals. And for animal bodies after "bai tiao" disposing (remaining parts of poultry or livestock after removal of hair and viscera), the body cavity swabs were collected.

Sewage (silt) samples were collected by the use of virus sampling swabs to probe into the silt at the bottom of drainage channels in the market. Sewage and silt samples were preserved in virus preservation solution. For the sewage well, a container was used to take a silt-water mixture from a location near the bottom of the well, and an appropriate amount of sample was collected by using virus sampling swabs and then preserved in virus preservation solution.

Nucleic acid extraction and SARS-CoV-2 real-time PCR assay

A virus nucleic acid extraction kit (Xi'an Tianlong) was used to extract viral nucleic acid from samples using an automated nucleic acid extraction instrument according to the manufacturer's instructions. Real-time (RT) PCR was performed on extracted nucleic acid samples with a SARS-CoV-2 nucleic acid assay kit. The reagent brands include BioGerm (40/38, cycle number/cut-off value, the same as below), DAAN (45/40), and BGI (40/38).

Virus isolations

Samples positive for SARS-CoV-2 were cultured in Vero E6 and Huh7.5 cells. The cell lines were inoculated with positive samples and three blind passages were performed for each sample. The culture supernatant and cell pellet of each passage were harvested for RT PCR. The morphology of viral particles in the cell sections and the supernatant were observed by transmission electron microscope (TEM).

Metagenomic sequencing

Metagenomic sequencing was conducted at Wuhan BGI. Nucleic acid was extracted using Qiagen's viral RNA microextraction kit and human nucleic acid was removed

using an enrichment kit to improve the sensitivity of viral RNA detection. Extracted RNA was reverse transcribed into cDNA and segmented into 150-200 bp by enzyme digestion. After repair, fitting, purification, PCR amplification and purification, sample concentration was assayed and SE50+10 sequencing performed by DNBSEQ-T7, and an average output of more than 200 million reads was obtained. Sequencing data were compared with those in a SARS-CoV-2 database to determine whether the samples contained coronavirus sequences.

Identification of the species abundances

The species abundances were obtained from the RNA-seq datasets of the environmental samples. The raw sequenced reads were mapped to the NCBI Nucleotide database using the kraken2 software (confidence 0.05) (32). The matrix of species was obtained by using the pavian algorithm (33). The correlation value between SARS-COV-2 and other species were examined by the Pearson's and Spearman's correlation coefficient method.

Extended data figure legends/tables.

Extended Data Figure S1. Positive environmental samples associated with different products in the Huanan Seafood Market.

Dots represent the percentage of positive environmental samples associated with each product. Bars represent 95% confidence intervals for the binomials in the text above. Note that the CI for some products (e.g. vegetables, farmed wildlife) have broad error bars that are likely due to the low number of vendors for these categories in the market. Nine of the 10 vendors selling farmed wildlife have been sampled.

Extended Data Table 1. Overview of environmental sample sampling and testing in the Huanan Seafood Market.

	Number	of	Number of positive	Number of	
	samples		samples by RT-PCR	isolated viruses	
Huanan Seafood Market	718		40	3	
Warehouses related to the Huanan Seafood Market	14		5		
Other markets in Wuhan*	30		1		
Drainage system in the Huanan Seafood Market	110		24		
Sewerage wells in surrounding areas	51		3		
Total	923		73	3	

Extended Data Table 2. Summary of number of reads mapped to positions 8782 and 28144 in different samples.

Position -	8782				28144				
	A	T	С	G	A	T	С	G	
A101	0	0	0	0	0	0	0	0	
A14	0	0	0	0	0	0	0	0	
A15	0	0	0	0	0	0	0	0	
A18	0	0	0	0	0	0	0	0	
A20	7	359	1	0	1	5	1596	5	
A2	0	0	0	0	0	0	0	0	
A33	0	0	2	0	0	0	1	0	
A55	0	0	0	0	0	0	0	0	
A61	0	0	2	0	0	0	1	0	
A63	0	0	0	0	0	0	0	0	
A87	0	0	0	0	0	1	0	0	

^{*}The other markets were Dongxihu market and Huanggang Center market.

A88	0	0	0	0	0	2	0	0
A90	0	0	0	0	0	0	0	0
A96	0	0	0	0	0	0	0	0
B17	0	0	0	0	0	0	0	0
B5	0	0	86	0	0	85	0	0
D32	0	0	0	0	0	0	0	0
E48	0	0	0	0	0	0	0	0
E61	0	0	0	0	0	0	0	0
E7	0	0	0	0	0	0	0	0
F100	0	0	0	0	0	0	0	0
F13	5	10	1874	3	0	1669	0	0
F33	0	0	0	0	0	0	0	0
F46	0	0	1	0	0	0	0	0
F54	1	1	106	0	1	39	0	0
F98	0	0	20	0	0	3	0	0
G93	0	0	0	0	0	0	0	0

Figures





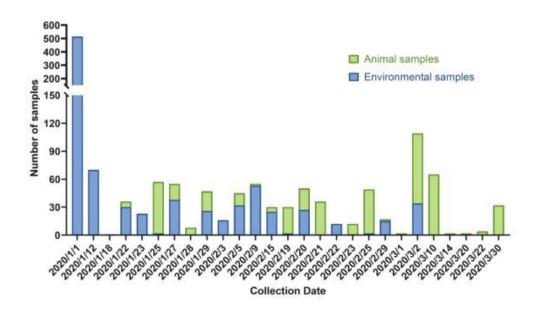


Figure 1

The distribution of the positive environmental samples in Huanan Seafood Market. A. As the place of the early cluster of COVID-19 patients, the Huanan Seafood Market is separated into eastern and western zones with the Xinhua Rd. between them. To detect for the presence of SARS-CoV-2 RNA, reverse

transcription, quantitative polymerase chain reaction (RT-qPCR) was performed. The locations of the positive samples were marked in the map of the market within orange, while the location of the samples that the live viruses were isolated from were labeled with red. The map also shows locations of stalls where domesticated wildlife products were sold. B. Timeline of environmental and animal samples collected within and around Huanan Seafood Market

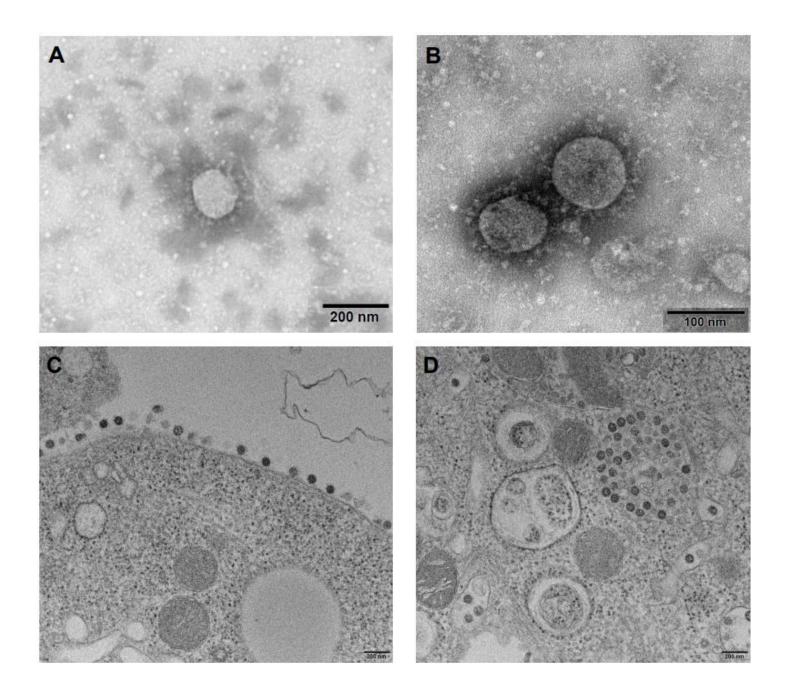


Figure 2

The SARS-CoV-2 virus isolation from environmental samples of Huanan Seafood Market. The electron micrographs of the SARS-CoV-2 viruses isolated from the environmental samples in Huanan Seafood Market. To determine whether SARS-CoV-2 particles could be visualized from the cell supernatant and

lysate, we used transmission electron microscopy to observe the culture supernatant and ultra-thin section cells based from both VeroE6 and Huh7.5 cells. The electron micrographs showed that virus particles were present in both the supernatant (A, B) and the cells (C, D). Negative-stained virus particles were generally spherical, pleomorphic and 60-140 nm in diameter. Spike protrusions were observed around the particles in a crown (corona) shape (A, B). In ultra-thin cultured cell sections, a group of virus particles can be seen outside the cell (C), and sheets of virus particles canalso be observed inside the cells (D).

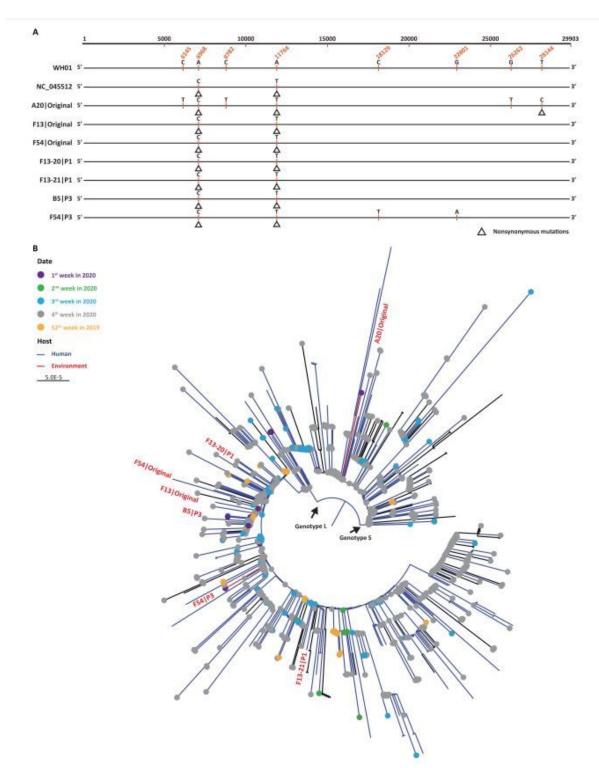


Figure 3

The genomic analysis of SARS-CoV-2 sequenced from Huanan Seafood Market.

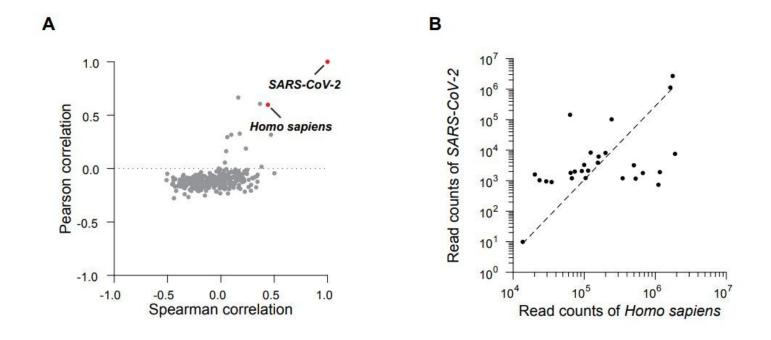


Figure 4

Clustering of environmental samples in Huanan Seafood Market. A, Correlation of SARS-COV-2 with other species in SARS-COV-2 positive cases. The abundance of SARS-COV-2 and other species was used for analysis. Both Pearson's and Spearman's correlation values were plotted. B, Correlation of the abundance of SARS-COV-2 and Homo sapiens.