Primers name	Primer sequences	PCR condition
IAV-Primer set 1	Fwd: 5'-ATGAGTCTTCTAACCGAGGTC-3'	95°C for 5 min
(982 bp)	Rev: 5'-GTCAGCATAGAGCTGGAGTAA-3'	Ļ
		[95°C for 30 sec, 52°C for 30 sec, 72°C
		for 1 min] x 25, 24 times (Figure 2A, 2C)
		Ļ
		72 °C for 10 min
IAV-Primer set 2	Fwd: 5'-AAGACCAATCCTGTCACCTC-3'	95°C for 5 min
(253 bp)	Rev: 5'-CAGTTGTATGGGCCTCATATAC-3'	↓
		[95°C for 30 sec, 52°C for 30 sec, 72°C
		for 1 min] x 22 times
		Ļ
		72 °C for 10 min
IAV-Primer set 3	Fwd: 5'-ACAGATTGCTGACTCCCA-3'	95°C for 5 min
(320 bp)	Rev: 5'-TGATCCTCTCGCTATTGCC-3'	\downarrow
		[95°C for 30 sec, 52°C for 30 sec, 72°C
		for 1 min] x 22 times
		Ļ
		72 °C for 10 min
FCV-Primer set	Fwd: 5'-TCCACACTAGCGTCAACTGG-3'	95°C for 5 min
(264 bp)	Rev: 5'-GACGAGCGTCAAACAGAACA-3'	Ļ
		[95°C for 30 sec, 49°C for 30 sec, 72°C
		for 1 min] x 32 times
		Ļ
		72 °C for 10 min
MNV-F1 and -R1	Fwd: 5'-GCCATGCATGGTGAAAAG-3'	95°C for 5 min
[1] (721 bp)	Rev: 5'-CATGCARACCAGGCGCATAG-3'	↓
		[95°C for 30 sec, 49°C for 30 sec, 72°C
		for 1 min] x 33 times
		↓

Table S1 The sequence of primers used in this study and each PCR condition.

		72 °C for 10 min
NIID_2019-nCoV	Fwd: 5'-AAATTTTGGGGGACCAGGAAC-3'	95°C for 5 min
_N_F2 and R2	Rev: 5'-TGGCAGCTGTGTAGGTCAAC-3'	Ļ
(158 bp)		[95°C for 30 sec, 49°C for 30 sec, 72°C
		for 1 min] x 30 times
		Ļ
		72 °C for 10 min



Figure S1. Scheme of the sample extraction and isolation.



Figure S2. Evaluation of the MNV-inactivating activity of 25 μ g/ml SS-derived fractions. The SS-derived fractions and DMSO control were added to solutions containing MNV followed by incubation at 25°C for 48 h. Viral titer was then evaluated. Aqu: Aqueous extract.

Figure S3



Compound 8

HC

н

но





он

ОН

ΟН

OH

он



Figure S3. Chemical structures of compounds isolated from *SS*. (A–C) Compounds isolated from (A) Fr 1C, (B) Fr 1D, and (C) Fr 1E.

Figure S4

Electron microscopy (6.25 μm²)



Figure S4. Electron microscopic images of IAV under low magnification (see Figure 3). The IAV virions featured in Figure 3 were evaluated using transmission electron microscopy under low magnification. The panels to the left and right include representative images of DMSO- and Fr 1C-treated virion particles, respectively in 6.25 μ m² fields.



E Target: SARS-CoV-2, Viral titer related to Fig 5C



Figure S5. The viral titers of FCV, MNV, and SARS-CoV-2 treated with *SS*-derived fractions (see Figures 4 and 5).

(A–E) Viral titer of the viral mixtures in (A) Figure 4A, (B) Figure 4B, (C) Figure 4C, (D) Figure 4D, and (E) Figure 5C. Aqu: Aqueous extract.

Reference

 Kitajima, M.; Oka, T.; Tohya, Y.; Katayama, H.; Takeda, N.; Katayama, K. Development of a broadly reactive nested reverse transcription-PCR assay to detect murine noroviruses, and investigation of the prevalence of murine noroviruses in laboratory mice in Japan. *Microbiol. Immunol.* 2009, 53, 531-534, doi:10.1111/j.1348-0421.2009.00152.x.