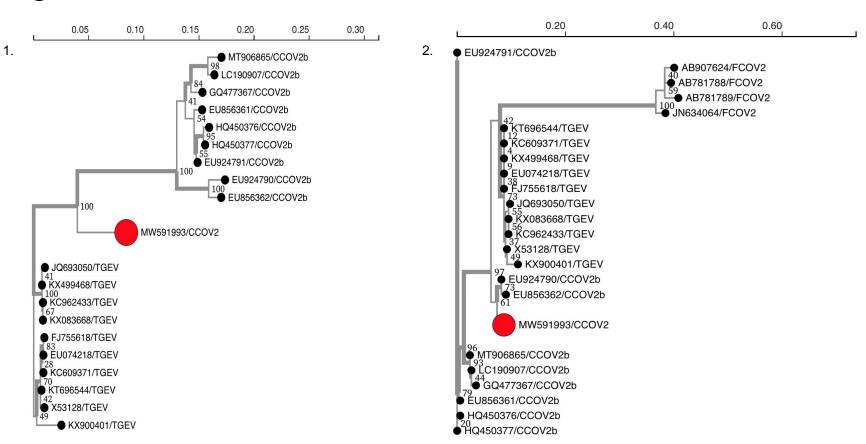
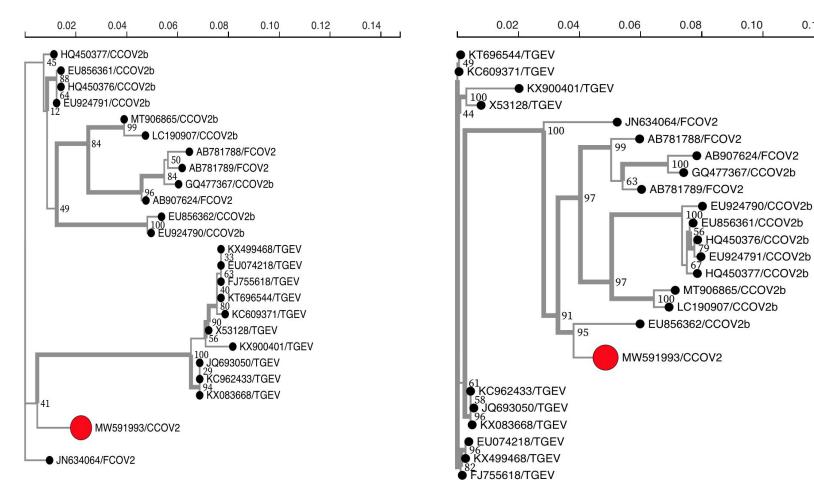
Table S1: Isolate details and sequence accessions for the included CoVs.

Alignment set	nment set Accession number Alpha-1 Type		Strain Name	Collection date	Location Malaysia	
1 & 2	MW591993.1	CCoV-HuPN-2018	-HuPN-2018 CCoV-HuPN-2018			
1 & 2	EU856361.1	CCoV2b	341/05	Dec 2005	Italy	
1 & 2	EU924791.1	CCoV2b	119-08	Mar 2008	Italy	
1 & 2	HQ450377.1	CCoV2b	68-09	2009	Greece	
1 & 2	HQ450376.1	CCoV2b	66-09	2009	Greece	
1 & 2	EU924790.1	CCoV2b	430-07	Oct 2007	Italy	
1 & 2	EU856362.1	CCoV2b	174-06	Mar 2006	Italy	
1 & 2	MT906865.1	CCoV2b	2020-7	2020	United Kingdom	
1 & 2	GQ477367.1	GQ477367.1 CCoV2b CCOV-NTU336-2008		Nov 2008	Taiwan	
1 & 2	LC190907.1	CCoV2b	CCOV-Dog-HCM47-2015	June 2015	Vietnam	
1 & 2	KX900401.1	TGEV	TGEV/USA/Tennessee144/2008	15 Apr 2008	USA - Tennessee	
1 & 2	KC609371.1	TGEV	NA	21 May 2013	China	
1 & 2	FJ755618.2	TGEV	H16	1973	China	
1 & 2	KT696544.1	TGEV	JS2012	2012	China	
1 & 2	EU074218.2	TGEV	Attenuated H	NA	China	
1 & 2	X53128.1	TGEV	TGEV FD772-70	NA	China	
1 & 2	KX499468.1	TGEV	TGEV AHHF	22 Dec 2015	China	
1 & 2	KC962433.1	TGEV	TGEV HX	5 May 2012	China	
1 & 2	JQ693050.1	TGEV	NA	NA	South Korea	
1 & 2	KX083668.1	TGEV	NA	2015	China	
2	AB781789.1	FCOV2	KUK-H-L	NA	Japan	
2	AB781788.1	FCOV2	M91-267	NA	Japan	
2	AB907624.1	FCOV2	Tokyo/cat/130627	NA	Tokyo, Japan	
2	JN634064.1	FCOV2	WSU-79-1683	1979	USA - Washingto	



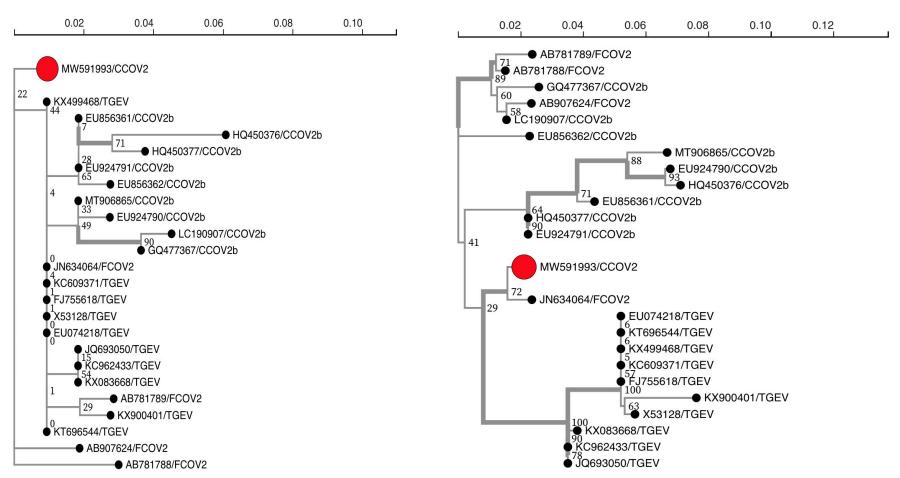
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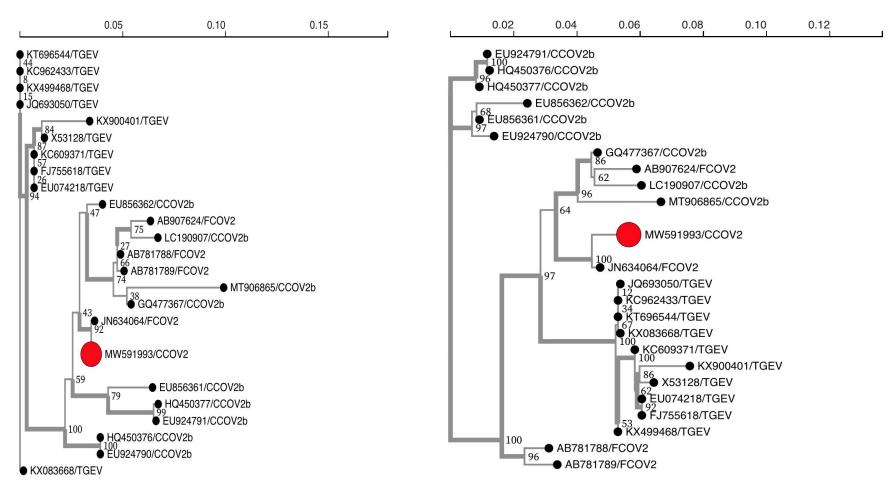
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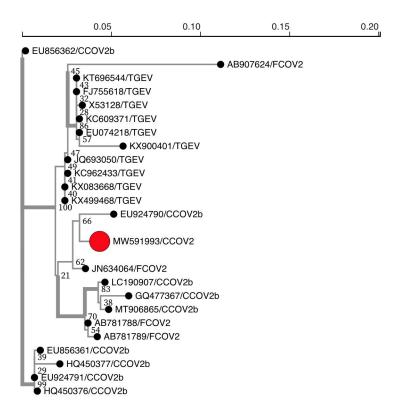


Figure S1. Phylogenetic trees for each GARD fragment inferred using RaxML. 1. Is the first non-recombinant fragment from alignment set I and the further trees correspond to non-recombinant fragments from alignment set II. Tree 2 corresponds to non-recombinant fragment 2 as depicted in the Fig. 1 gene map; tree 3 corresponds to non-recombinant fragment 3, and so on. These trees represent the phylogenetic incongruencies between the different non-recombinant fragments.

Table S2: Positive selection statistics from MEME and FEL

HuPN-2018 sites	FIPV mapped sites	MEME Non-synonymous substitution rate	MEME Synonymous substitution rate	FEL Non-synonymous substitution rate	FEL Synonymous substitution rate	MEME p-value	FEL p-value	
13	13	3.58	0	NA	NA	0.04	NA	
110	130	30.94	0	29.432	0	0.02	0.02	
124	144	3.87	0	NA	NA	0.02	NA	
248	262	2.9	0	2.784	0	0.02	0.04	
367	384	4.8	0	4.817	0	0.04	0.05	
575	603	11.17	0	NA	NA	0.02	NA	
529	549	9.23	0	9.46	0	NA	0.05	
698	715	11.26	0	NA	NA	0.05	NA	
619	637	12.48	0	12.47	0	0.03	0.02	
589	619	10.84	0	NA	NA	0.04	NA	
1227	1248	27.49	0	NA	NA	0.04	NA	
1296	1311	38.37	0	37.386	0	0.01	0.01	
1233	1252	3.3	0	3.281	0	0.04	0.04	

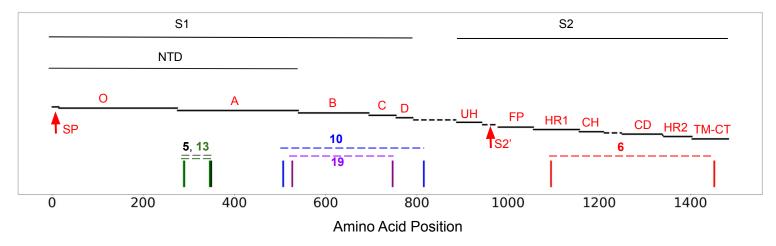


Figure S2. RDP5 results with supported recombination events (event boundaries outlined) that implicate CCoV-HuPn-2018, positioned along a spike gene map. A complete, enumerated list of the RDP5 recombination events, involving all sequences, can be found in Table S3. Event 5 (black) involves a proposed FCoV2 recombinant, with genetic donors: CCoV-HuPn-2018 and a different FCoV2. Event 13 (green) extensively overlaps with 5, as well as with GARD fragment 2 (Fig. 1), where CCoV-HuPn-2018 is a proposed recombinant, with genetic donors: FCoV2 from event 13, as well as a TGEV. Event 10 (cyan) proposes the same FCoV2 sequence from 5 and 13 as the recombinant, with genetic donors: CCoV-HuPn-2018 and a TGEV. Event 19 (purple) falls within 10, with CCoV-HuPn-2018 as the proposed recombinant and the genetic donors: a CCoV2b and a TGEV. Event 6 (red) proposes a CCoV2b as the recombinant, with genetic donors: CCoV-HuPn-2018 and another CCoV2b.

Table S3: RDP5 Recombination Results

bio kays								
 It is possible that this apparent recombination signal could have been caused by an evolutionary 								
process offer than recombination.								
= The actual breakpoint position is undetermined (it was root likely either overprinted by a subsequent								
recombination event or off the edges of the analysed sequence fragments).								
= The recombinant sequence may have been misidentified jone of the identified parents might be the recombinant								
or Parent - Parent contributing the smaller fraction of sequence.								
or Parent - Parent contributing the larger fraction of sequence.								
nown - Only one parent and a recombinant need be in the alignment for a recombination event to be detectable.								
The sequence field as unknown was used to infer the existance of a missing parental sequence.								
« No significant P-value was recorded for this recombination event using the particular method in question.								
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	_	_	Desipont Postors					
	-	in Alignment		In Recombinant Sequence		Passane to ABTRITIOS_1_FCOV2_SEKUK_H_L_INA_SPELIS_CATUS_CUAPAN		
embination Event Number	Number In PICP File	Baren	End	Begin	End	Manus	Box1	Pacombinant Sequencers
tion the contraction		1 179			2290	1704		HQHS6876_1_CCOVSb_1666_66_NA_NDOG_cGREECE_co
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process other than recombination.								
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ABBOTECH 1 PCOVZ SITOKYO CAT 130827 INA INFELIS CAT

HQ480377_1_CCC0v2b_9888_09_NA_FEXCG_cGREECE_cod2009

EU856981_1_CCOV2b_st941_05_INA_HDOG_clTALY_cooDEC BURGITRE 1 CCCV2b st119 08 INA NDOG cITALY codMAR HQ450378_1_CCCV2b_st89_09_INA_NDOG_cGREECE_cod2009 ADBOTROM 1 PCOVO SITORYO CAT 130807 INA HEBUS CAT

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HQ450377_1_CCC0V2b_sit8_09_INA_hDOG_cGREECE_cod2009

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EU886962_1_CCCV25_9174_96_INA_NDOG_0ITALY_codMAR

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MANSCHOOL 1 COOKS SINA ICCOV HURS 2018 SHOMD CMALAY

A8781789_1_FCOV2_BIKUK_H_L_INA_RFELIS_CATUS_GJAPAN

MW591993 1 CCOV2 IENA ICCOV HUPI 2018 IHDMO IMALAY

Uninown (ABB07824_1_FCOV2_SETOKYO_CAT_180827_INA_RFELIS_CAT) INS

Unknown (EU824791_1_CCCV2b_st110_68_INA_NDGG_clTALY_codMAR_) INS

AB781789_1_FCOV2_stKUK_H_L_INA_NFELIS_CATUS_cJAPAN

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Unknown(AB781788_1_FCOV2_99991_287_INA_FFELIS_CATUS_GJAPAN)

9 04/9994014979ME-02 NF

6.077888015894E-04.NS

0.0000000008179 1.29024890729025E-07 NS

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Detection Methods 2.54929441781989E-21 2.89904252019586E-19 1.88842061271346E-21 1.80241825590646E-12 1.11712519923247E-12 3.85788931003041E-19 NS NS 3.99149434799133E-27 A TANAHAMAN IN TONON MANAGEMENT AND A TANAHAMAN IN TONON MANAGEMENT AND A STANDARD MANAGEMENT AN 1.00000000000100E-10 1.01010040130001E-10 2.0400010030700E-10 3.01017339024100E-14 2.07102103000000E-00 8.49132400300000E-10 NS NS 1.47000002270140E-13

1.74095030001779E-05 8.39077111901959E-04 M²

8.83742902309869E-03 3.76911218069698E-04 NS

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3.485773323854304E-03. 1.415369253113477E-03.165

4.76176649908222E-03 4.10096299581509E-02 6.80442619455942E-19 NS NS NS

1.18918131310852E-03 5.2541093519971E-03 1.83969470554917E-03 NS

4.83472913678042E-02 8.68564366541979E-06 1.16862403256412E-06 NG

mumo Lamp 2000 8.42976000758612E-06 6.57764854284350E-12 NS NS 1.47650662275146E-13 1.615477377519446-06 7.490091293771936-12 NS NS 2.595406644411456-09 2.56815477303156E-06 8.52400660331166E-13 1.34314511108633E-10 1.77766291254701E-11 2.50404885152511E-16 NG NS 1.47650662275146E-12 1.13700574314452E-04 1.82373320519041E-08 2.72418943094248E-02 NS

0.0080075189019E-05 0.0076180227046E-10 2.0077776201688E-12 0.5886404347002E-00 2.828100075559E-06 1.88070874100544E-18 NS NS 4.98800249015477E-06

0.000000017955 2.930545594508405-10 2.36129641144729E-02 6.76594627581509E-03 2.3715029253094E-07 NS NS 1.39269509157434E-04 NS NS 3.19280417075829E-09

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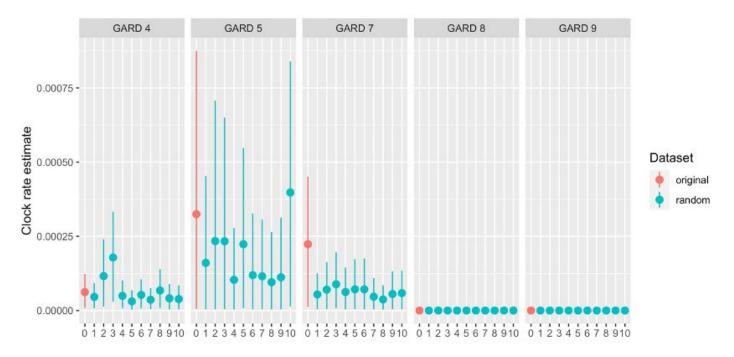


Figure S3. Mean and 95% HPD clock rate estimates for the original dataset and ten datasets with random dates are shown for the five GARD fragments with evidence of temporal signal on the root-tip regression analysis. Only GARD 7 had a mean clock rate estimate above the 95% HPD of the randomized datasets, indicating the presence of a temporal signal.

Table S4: Root-tip-regression results for each GARD Partition

GARD partition	Correlation coefficient	R ²	
1	0.083	0.0070	
2	-0.046	0.0021	
3	-0.11	0.013	
4	0.20	0.040	
5	0.20	0.039	
6	0.088	0.0077	
7	0.36	0.13	
8	0.30	0.089	
9	0.21	0.046	

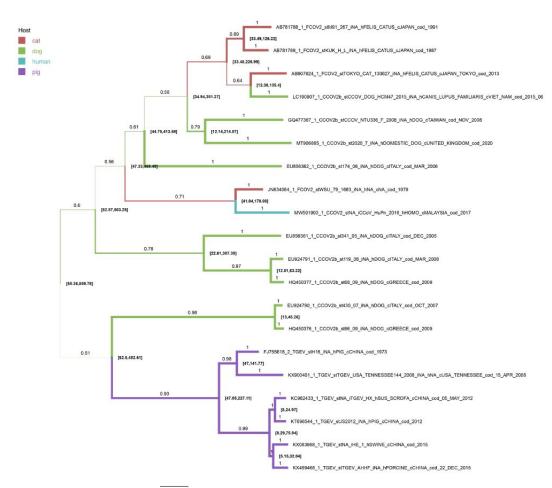


Figure S4. Ancestral host reconstruction and divergence time estimates. Branches are colored by inferred host species. The branch width is proportional to the posterior probability of host assignment (also labeled on branches). Internal nodes are labeled with the divergence time 95% HPD in years from the most recent sample date, 2017. CCoV-HuPn-2018 diverged from FCoV2 JN634064

between 40 and 170 years ago, with a median date estimate of 1957.

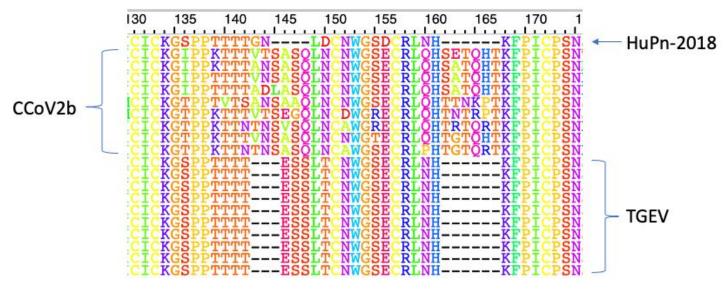


Figure S5. Set I sequence alignment highlighting the region involved in the proposed sialic acid binding, identified in the Krempl et al. (1997) mutation experiments.