

# The Ecological Niche of bat Viruses Closely Related to SARS-CoV-2, as Inferred from Phylogeographic Analyses of Rhinolophus Species

Alexandre HASSANIN (✉ [alexandre.hassanin@mnhn.fr](mailto:alexandre.hassanin@mnhn.fr))

Sorbonne University

Vuong Tan Tu

Vietnam Academy of Science and Technology

Manon Curaudeau

Sorbonne University

Gabor Csorba

Hungarian Natural History Museum

---

## Research Article

**Keywords:** coronavirus, horseshoe bats, reservoir hosts, Indochina, China, immunity

**Posted Date:** April 26th, 2021

**DOI:** <https://doi.org/10.21203/rs.3.rs-418434/v1>

**License:**  This work is licensed under a Creative Commons Attribution 4.0 International License.

[Read Full License](#)

---

**Version of Record:** A version of this preprint was published at Scientific Reports on July 12th, 2021. See the published version at <https://doi.org/10.1038/s41598-021-93738-z>.

**The ecological niche of bat viruses closely related to SARS-CoV-2, as inferred from** 1  
**phylogeographic analyses of *Rhinolophus* species** 2

Alexandre Hassanin<sup>1°\*</sup>, Vuong Tan Tu<sup>2°</sup>, Manon Curaudeau<sup>3</sup>, and Gabor Csorba<sup>4</sup> 3  
4

<sup>1</sup> Institut de Systématique, Évolution, Biodiversité (ISYEB), Sorbonne Université, MNHN, CNRS, 6  
EPHE, UA, Paris; [alexandre.hassanin@mnhn.fr](mailto:alexandre.hassanin@mnhn.fr) 7

<sup>2</sup> Institute of Ecology and Biological Resources, Vietnam Academy of Science and Technology, 8  
No. 18, Hoang Quoc Viet Road, Cau Giay District, Hanoi, Vietnam; [tuvuongtan@gmail.com](mailto:tuvuongtan@gmail.com) 9

<sup>3</sup> Institut de Systématique, Évolution, Biodiversité (ISYEB), Sorbonne Université, MNHN, CNRS, 10  
EPHE, UA, Paris; [manon.curaudeau1@mnhn.fr](mailto:manon.curaudeau1@mnhn.fr) 11

<sup>4</sup> Department of Zoology, Hungarian Natural History Museum, Baross u. 13, Budapest H-1088, 12  
Hungary; [csorba.gabor@nhmus.hu](mailto:csorba.gabor@nhmus.hu) 13

<sup>°</sup> co-first authors 14

\* correspondence: [alexandre.hassanin@mnhn.fr](mailto:alexandre.hassanin@mnhn.fr); Tel.: +33 1 40 79 56 93 15

16

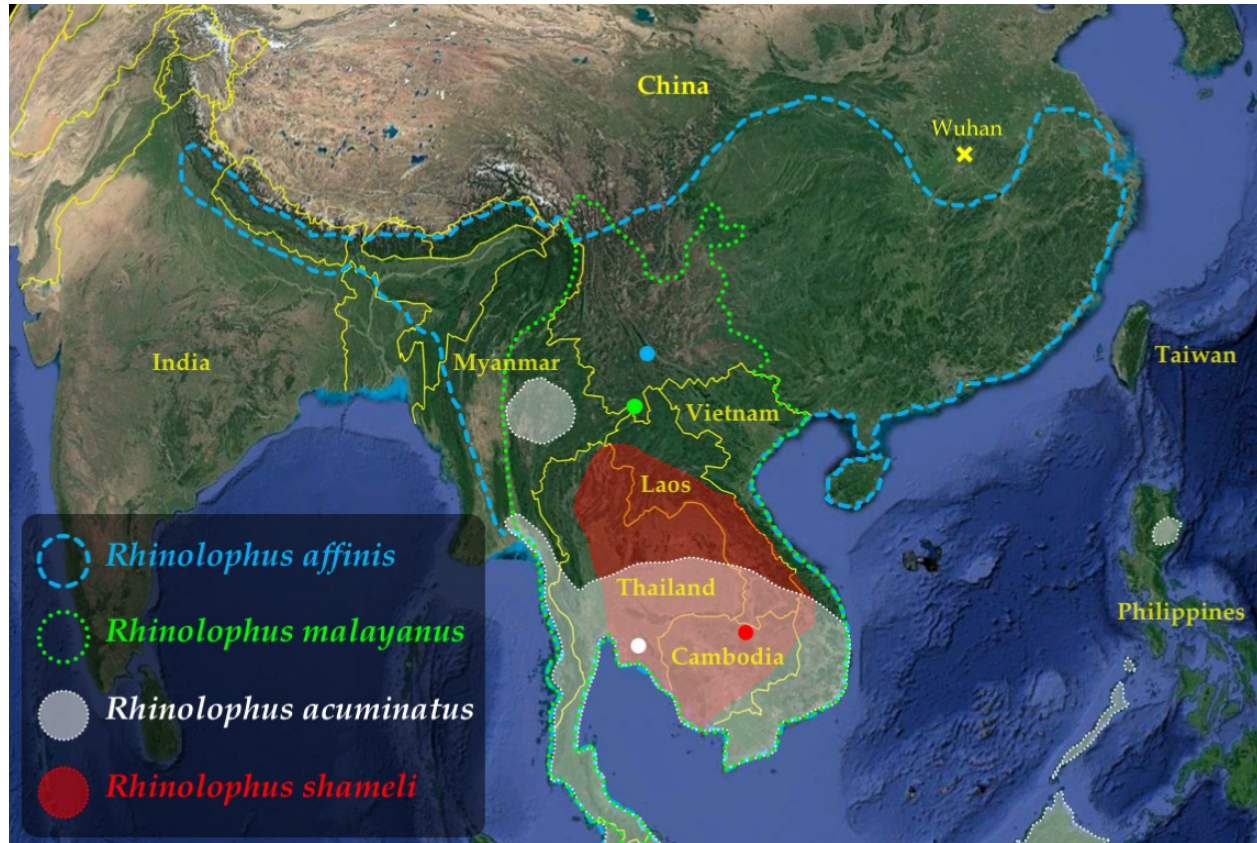
<b>Abstract</b>	17
	18
To date, viruses closely related to SARS-CoV-2 have been reported in four bat species:	19
<i>Rhinolophus acuminatus</i> , <i>Rhinolophus affinis</i> , <i>Rhinolophus malayanus</i> , and <i>Rhinolophus shameli</i> .	20
Here, we analysed 343 sequences of the mitochondrial cytochrome c oxidase subunit 1 gene ( <i>COI</i> )	21
from georeferenced bats of the four <i>Rhinolophus</i> species identified as reservoirs of viruses closely	22
related to SARS-CoV-2. Haplotype networks were constructed in order to investigate patterns of	23
genetic diversity among bat populations of Southeast Asia and China. No strong geographic	24
structure was found for the four <i>Rhinolophus</i> species, suggesting high dispersal capacity. The	25
ecological niche of bat viruses closely related to SARS-CoV-2 was predicted using the four	26
localities in which bat viruses were recently discovered and the localities where bats showed the	27
same <i>COI</i> haplotypes than virus-positive bats. The ecological niche of bat viruses related to SARS-	28
CoV was deduced from the localities where bat viruses were previously detected. The results show	29
that the ecological niche of bat viruses related to SARS-CoV2 includes several regions of mainland	30
Southeast Asia whereas that of bat viruses related to SARS-CoV is mainly restricted to China. In	31
agreement with these results, human populations in Laos, Vietnam, Cambodia, and Thailand appear	32
to be much less affected by the Covid-19 pandemic than other countries of Southeast Asia. In the	33
climatic transitional zone between the two ecological niches (southern Yunnan, northern Laos,	34
northern Vietnam), genomic recombination between highly divergent viruses is more likely to	35
occur. Since recombinant viruses can threaten the benefit of vaccination campaigns, these regions	36
should be under surveillance.	37
	38
<b>Keywords:</b> coronavirus; horseshoe bats; reservoir hosts; Indochina; China; immunity.	39

## Introduction

The Severe Acute Respiratory Syndrome coronavirus 2 (SARS-CoV-2) emerged in December 2019 in Wuhan (China) [1]. After 16 months, the coronavirus disease 2019 (Covid-19) pandemic has affected more than 137 million of people around the world, claiming over 2.95 million lives [2]. The origin of SARS-CoV-2 remains enigmatic and many hypotheses have been advanced to explain the first animal-to-human transmission [3].

Within the family Coronaviridae, the subgenus *Sarbecovirus* includes two human viruses, SARS-CoV-2 and SARS-CoV (which was responsible for the SARS epidemic in 2002-2004) [4]. The genomes of these two viruses share only 80% of nucleotide identity, and whole-genome phylogenies have shown that they belong to two divergent lineages [1, 5-7], which we refer to herein as SARS-CoV related coronaviruses (SCoVrCs) and SARS-CoV-2 related coronaviruses (SCoV2rCs). Most SCoVrCs were discovered in bats collected in China after the SARS epidemic, and the great majority were found in horseshoe bat species of the genus *Rhinolophus* (family Rhinolophidae), suggesting that this taxon is the natural reservoir host of sarbecoviruses [8]. More recently, several viruses showing between 96.2 and 91.8 % of genome identity with SARS-CoV-2 were identified from fecal samples of four horseshoe bat species: the RaTG13 virus was isolated from a *R. affinis* bat collected in Yunnan in 2013 [1]; the RmYN02 virus was found in a *R. malayanus* bat sampled in Yunnan in 2019 [5]; two variants of the same virus RshSTT200 (other variant: RshSTT182) were detected in two *R. shameli* bats caught in northern Cambodia in 2010 [6]; and five variants of the same virus RacCS203 (other variants: RacCS224, RacCS253, RacCS264, and RacCS271) were sequenced from five *R. acuminatus* bats collected in eastern Thailand in 2020 [7]. The bat species *R. acuminatus* and *R. shameli* are endemic to Southeast Asia whereas the two other bat species, *R. affinis* and *R. malayanus*, are distributed in both Southeast Asia and China (Fig. 1), suggesting that Southeast Asia is the main region where bat SCoV2rCs diversified. In addition, these recent data confirmed that the genus *Rhinolophus* is the natural reservoir host of all sarbecoviruses [3,8]. Note that this hypothesis was already corroborated by the

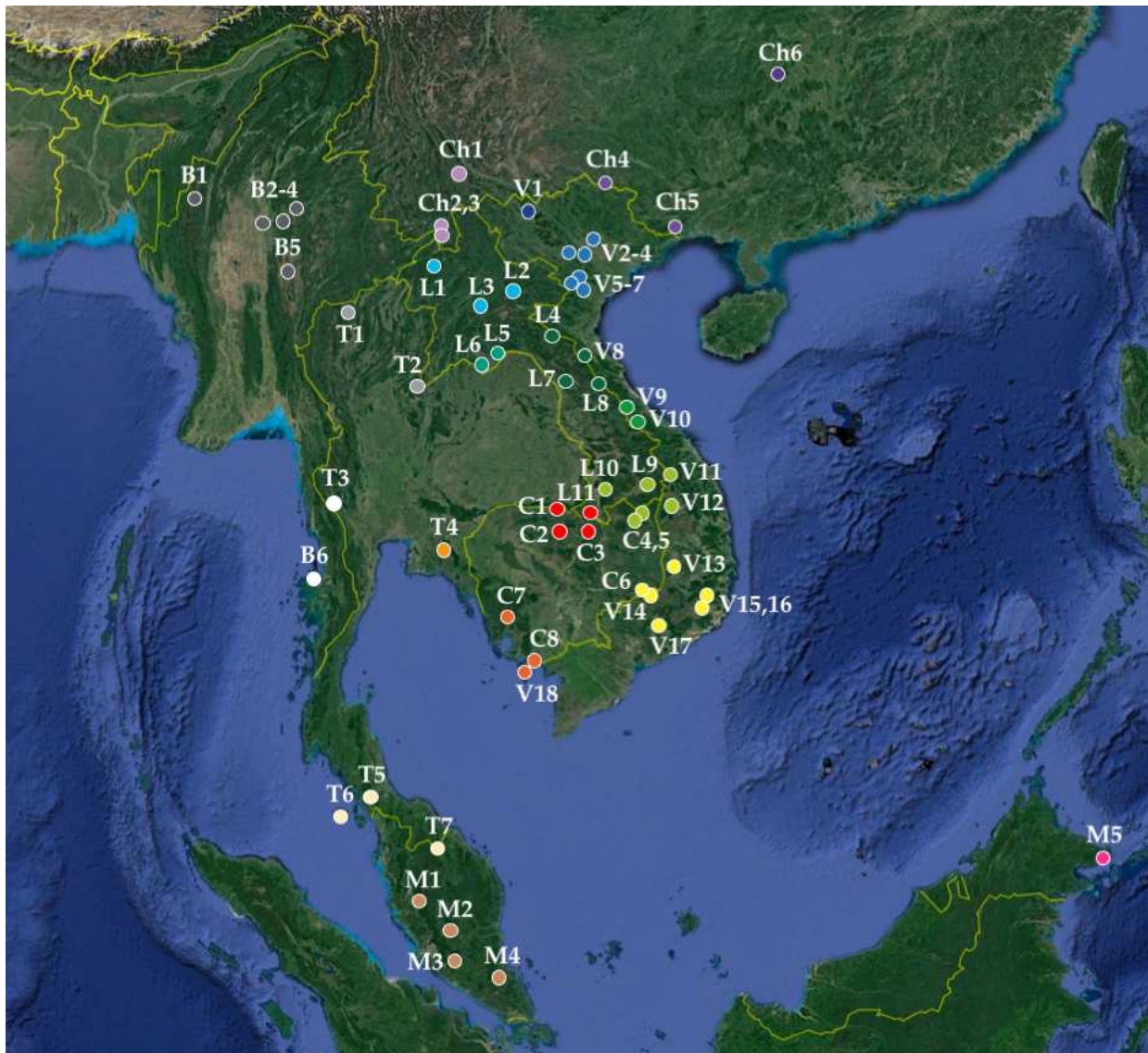
discovery of two divergent sarbecovirus genomes in horseshoe bat species collected out of Asia: 66  
one in *Rhinolophus blasii* from Bulgaria (BM48-31) [9] and another in an unidentified *Rhinolophus* 67  
species from Kenya (BtKY72) [10]. 68



**Figure 1. Geographic distribution of the four *Rhinolophus* species found positive for viruses closely related to SARS-CoV-2 in southern China and Southeast Asia.** Map from Google Earth US Dept of State Geographer © 2020 Google – Image Landsat / Copernicus - Data SIO, NOAA, U.S. Navy, NGA, GEBCO. For each of the four *Rhinolophus* species, the geographic distribution was extracted from the IUCN website [11]. The coloured dots show the four geographic locations where bats found positive for SCoV2rCs were collected. 69  
70  
71  
72  
73  
74  
75  
76

Since SCoV2rCs have been circulating in horseshoe bats for many decades [12], it is important to 77  
study population genetic structure of bats found positive for these sarbecoviruses in order to 78  
evidence their dispersal capacity in China and Southeast Asia. In this report, the phylogeography 79  
of the four species *R. acuminatus*, *R. affinis*, *R. malayanus* and *R. shameli* was therefore examined 80  
using the mitochondrial cytochrome c oxidase subunit 1 gene (CO1) from 343 georeferenced bats 81  
collected in 62 localities of Southeast Asia and China (Fig. 2). For each of the four species, 82  
haplotype networks were constructed to investigate geographic patterns of genetic diversity among 83

bat populations. The results of these analyses were used to select specific location coordinates to 84  
predict the ecological niche of bat SCoV2rCs. 85

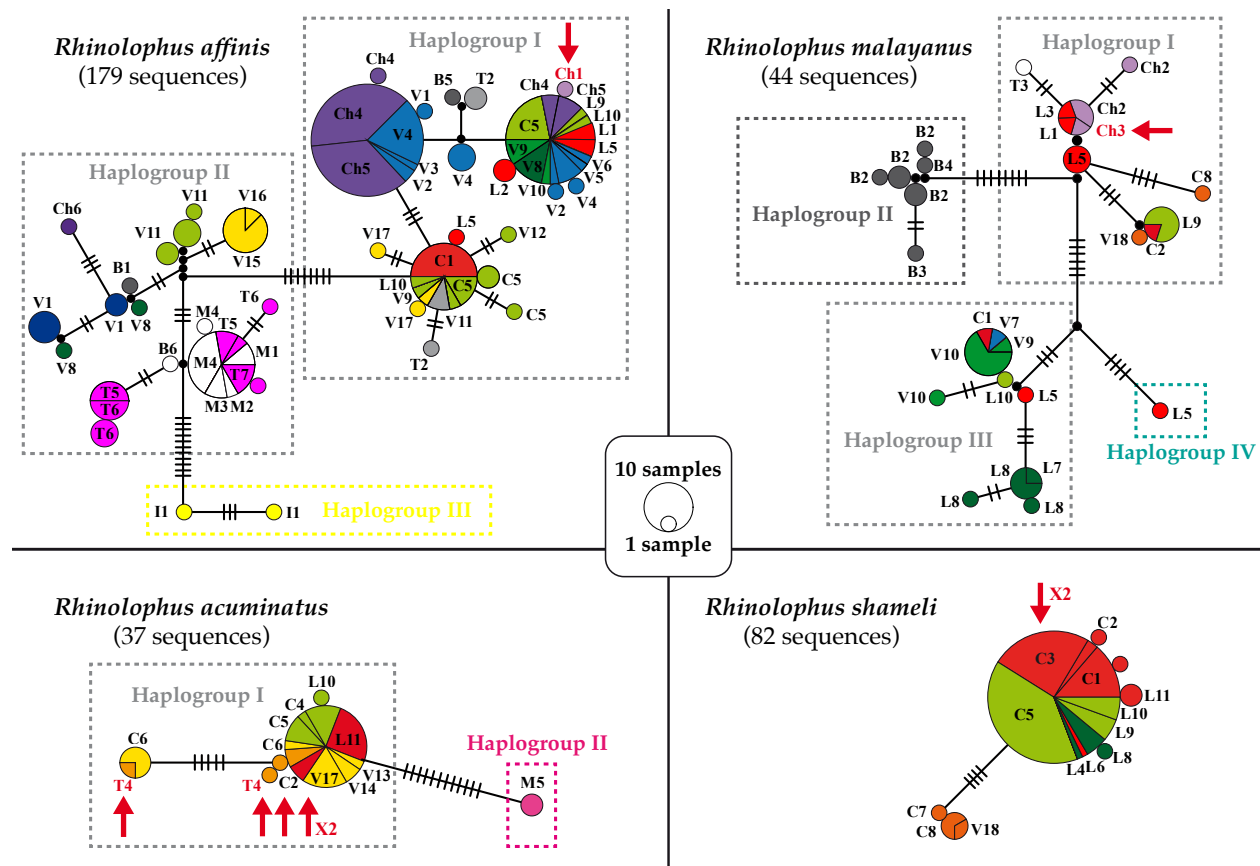


**Figure 2. Geographic localities analysed in this study.** The codes used for the countries are the 87  
following: B (Myanmar), C (Cambodia), Ch (China), I (Indonesia), L (Laos), M (Malaysia), T 88  
(Thailand), and V (Vietnam). Different regions were highlighted using colours. Map from Google 89  
Earth US Dept of State Geographer © 2020 Google – Image Landsat / Copernicus - Data SIO, 90  
NOAA, U.S. Navy, NGA, GEBCO. The locality names and GPS coordinates are provided in online 91  
supplementary table S1. 92  
93

## Results and Discussion 94

**Genetic analyses of *Rhinolophus* species identified as reservoirs of viruses closely related to SARS-CoV-2** 95  
96

Until now, SCoV2rCs have been found in four bat species of the genus *Rhinolophus*: *R. acuminatus*, *R. affinis*, *R. malayanus*, and *R. shameli*. The haplotype networks constructed using *COI* sequences of these four species are shown in Fig. 3. A star-like genetic pattern, characterized by one dominant haplotype and several satellite haplotypes was found for the two bat species endemic to Southeast Asia, i.e. *R. acuminatus* and *R. shameli*. 97  
98  
99  
100  
101



**Figure 3. Haplotype networks based on *COI* sequences of the four *Rhinolophus* species found positive for viruses closely related to SARS-CoV-2 (SCoV2rCs).** The networks were constructed with the median joining method available in PopART 1.5 [13]. The codes used for the countries are the following: B (Myanmar), C (Cambodia), Ch (China), I (Indonesia), L (Laos), M (Malaysia), T (Thailand), and V (Vietnam). Colours indicate the geographic origin of haplotypes according to Fig. 2 (see online supplementary table S1). The circles indicate haplotypes separated by at least one mutation. The black lines on the branches show the number of mutations  $\geq 2$ . Black circles represent missing haplotypes. Circle size is proportional to the number of haplotypes. Haplogroups separated by more than seven mutations (pairwise nucleotide distances  $> 1\%$ ) are highlighted by dotted lines. The red arrows show the positions of the nine bats found positive for SCoV2rCs. In the network of *R. acuminatus*, the most common haplotype (named Rac1 in online supplementary table S1) was found in northern Cambodia, southern Laos, eastern Thailand and southern Vietnam, indicating recent gene flow among these populations. Since a virus related to 102  
103  
104  
105  
106  
107  
108  
109  
110  
111  
112  
113  
114  
115

SARS-CoV-2 (91.8% of genome identity), named RacCS203, was detected in five *R. acuminatus* 116  
bats caught in eastern Thailand in June 2020 [6], the genetic pattern obtained for this species 117  
suggests that viruses closely related to RacCS203 may have circulated in most southern regions of 118  
mainland Southeast Asia. In contrast, *R. acuminatus* bats collected in Borneo (M5) showed a 119  
divergent haplotype (separated by 12 mutations; haplogroup II), suggesting that the South China 120  
Sea between mainland Southeast Asia and Borneo constitutes a barrier to gene flow. Isolated 121  
populations of *R. acuminatus* described in northern Myanmar, Indonesia (Java and Sumatra) and 122  
the Philippines [14] should be further studied. 123

The network of *R. shameli* shows a typical star-like pattern, the most common haplotype (named 124  
Rsh1 in online supplementary table S1) being detected in northern Cambodia and Laos. Since a 125  
virus related to SARS-CoV-2 (93.1% of genome identity), named RshSTT200, was recently 126  
discovered in two *R. shameli* bats collected in northern Cambodia in December 2010 [7], the 127  
genetic pattern obtained for this species suggests that viruses closely related to RshSTT200 may 128  
have circulated, at least in the zone between northern Cambodia and central Laos. The bats sampled 129  
south to the Tonle Sap lake (n = 4; southern Cambodia and Vietnamese island of Phu Quoc) were 130  
found to be genetically isolated from northern populations (four mutations). However, further 131  
sampling in the south is required to confirm this result, as it may reveal *COI* sequences identical 132  
to the haplotypes detected in the north. 133

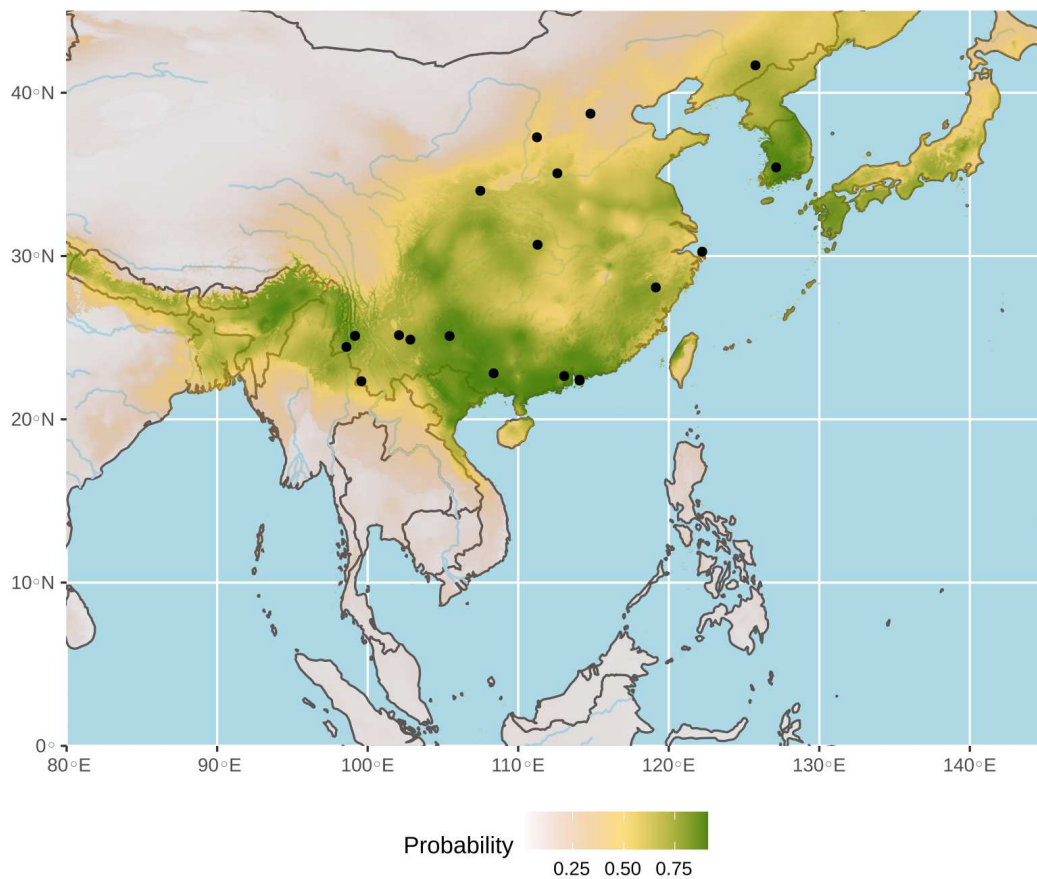
For the two species distributed in both China and Southeast Asia, i.e. *R. affinis* and *R. malayanus*, 134  
the genetic patterns are more complex with different haplogroups showing more than 1% of 135  
nucleotide divergence. In the network of *R. affinis*, there are three major haplogroups (named I, II 136  
and III in Fig. 3) separated by a minimum of seven mutations. The results are therefore in agreement 137  
with those previously published using *COI* and D-loop mitochondrial sequences [15]. The *COI* 138  
haplotypes detected in the localities sampled in southern China (ch1, ch4, ch5) are distantly related 139  
to the single haplotype available for central China (ch6), but they are also found in Laos, northern 140  
and central Vietnam, northern Thailand and northeastern Myanmar. This result suggests recent 141



gene flow between populations from southern Yunnan and those from northern mainland Southeast Asia. Since a virus related to SARS-CoV-2 (96.2% of genome identity), named RaTG13, was detected in one *R. affinis* bat captured in southern Yunnan in 2013 [1], the genetic pattern obtained for this species suggests that viruses closely related to RaTG13 may have circulated in the zone comprising southern Yunnan and northern mainland Southeast Asia. In the network of *R. malayanus*, there are four major haplogroups (named I, II, III and IV in Fig. 3) separated by a minimum of seven mutations. The *COI* haplotypes detected in the localities sampled in southern China (ch2 and ch3) were also found in northern Laos (L1 and L3), suggesting recent gene flow between populations from these two countries. Since a virus related to SARS-CoV-2 (93.7% of genome identity), named RmYN02, was recently isolated from one *R. malayanus* bat collected in southern Yunnan in June 2019 [5], the genetic pattern obtained for this species suggests that viruses closely related to RmYN02 may have circulated, at least between southern Yunnan and northern Laos. In contrast, the bats sampled in Myanmar were found to be genetically isolated from other geographic populations (haplogroup II in Fig. 3).

**Two different ecological niches for bat viruses related to either SARS-CoV or SARS-CoV-2** Sarbecoviruses have been already detected in a wide diversity of horseshoe bat species collected in Asia, Africa and Europe [5-10]. Therefore, there is no doubt that *Rhinolophus* species constitute the natural reservoir host of all sarbecoviruses [3,8]. The genus *Rhinolophus* currently includes between 92 [11] and 109 [16] insectivorous species that inhabit temperate and tropical regions of the Old World, with a higher biodiversity in Asia (63-68 out of the 92-109 described species) than in Africa (34-38 species), Europe (5 species) and Oceania (5 species). Although some *Rhinolophus* species are solitary, most of them are gregarious and live in large colonies or small groups generally in caves and hollow trees, but also in burrows, tunnels, abandoned mines, and old buildings [11,16]. However, they prefer large caves with total darkness, where temperatures are stable and less affected by diurnal and seasonal climatic variations. Importantly, all *Rhinolophus* species in

which sarbecoviruses were detected in previous studies [1, 5-9; 17] are cave species that form small groups or colonies (up to several hundreds) [11,18,19].

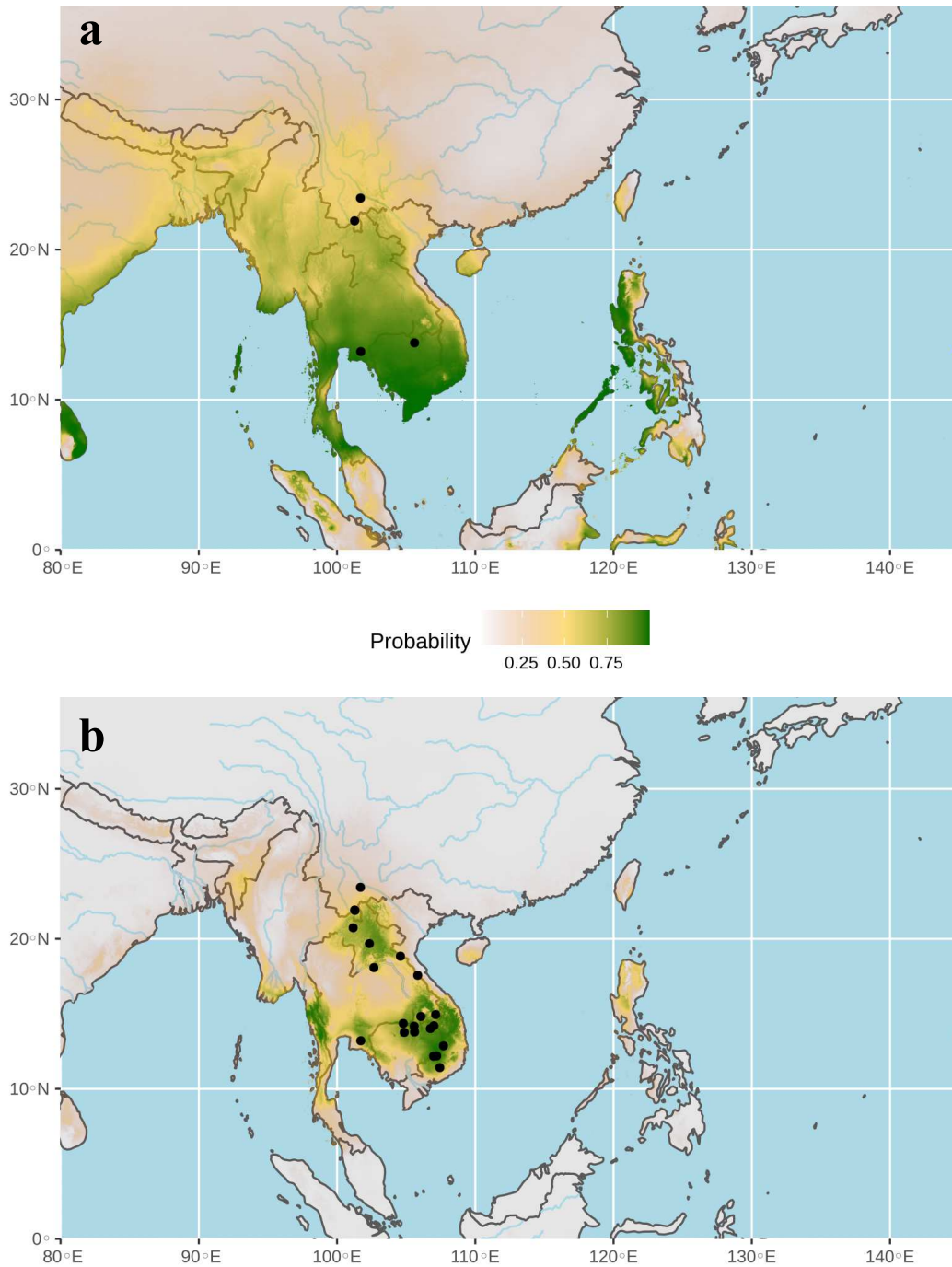


**Figure 4. Ecological niche of bat viruses related to SARS-CoV (SCoVrCs).** The geographic distribution of suitable environments was predicted using the Maxent algorithm in ENMTtools (see Methods for details). AUC = 0.81. Black circles indicate localities used to build the distribution model (see geographic coordinates in online supplementary table S2).

In China, many SCoVrCs were previously detected in several horseshoe bat species, including *Rhinolophus sinicus*, *Rhinolophus ferrumequinum* (currently *R. nippon*) [16], *Rhinolophus macrotis* (currently *R. episcopus*) [16], *Rhinolophus pearsoni*, and *Rhinolophus pusillus*, and it has been shown that they circulate not only among conspecific bats from the same colony, but also between bat species inhabiting the same caves [17,20,21]. The ecological niche predicted for bat SCoVrCs using a data set of 19 points (see online supplementary table S2) is shown in Fig. 4. The AUC was 0.81. The value was > 95% CI null-model's AUCs (0.68), indicating that the model performs significantly better than a random model (see online supplementary Fig. S1). The highest probabilities of occurrence (highlighted in green in Fig. 4) were found in Nepal, Bhutan,

Bangladesh, northeastern India, northern Myanmar, northern Vietnam, most regions of China south 185  
of the Yellow River, Taiwan, North and South Korea, and southern Japan. 186  
187

In Southeast Asia and southern China, SCoV2rCs have currently been found in four *Rhinolophus* 188  
species (*R. acuminatus*, *R. affinis*, *R. malayanus* and *R. shameli*) [1,6-8], but the greatest diversity 189  
of horseshoe bat species in mainland Southeast Asia (between 28 and 36 species) [11,16] suggests 190  
that many sarbecoviruses will be discovered soon. Despite the limited data currently available on 191  
SCoV2rCs, several arguments support that bat intraspecific and interspecific transmissions also 192  
occur with SCoV2rCs. Firstly, recent genomic studies have revealed that SCoV2rCs circulate and 193  
evolve among horseshoe bats of the same colony, as five very similar genomes (nucleotide 194  
distances between 0.03% and 0.10%) were sequenced from five *R. acuminatus* bats collected from 195  
the same colony in eastern Thailand [6], and as two genomes differing at only three nucleotide 196  
positions (distance = 0.01%) were sequenced from two *R. shameli* bats collected at the same cave 197  
entrance on the same night [7]. Secondly, the discovery of four viruses closely related to SARS- 198  
CoV-2 (between 96.2 and 91.8 % of genome identity) in four different species of *Rhinolophus* is a 199  
strong evidence that interspecific transmission occurred several times in the past. As detailed in 200  
online supplementary Table S1, these species were collected together in several localities of 201  
Cambodia (three species in C1, C2, and C5; two species in C8), Laos (four species in L10; three 202  
species in L9; two species in L1, L5, L8, L11), and Vietnam (two species in V10, V9, V17, V18). 203  
These data corroborate previous studies suggesting that sarbecoviruses can be transmitted, at least 204  
occasionally, between *Rhinolophus* species sharing the same caves. 205



**Figure 5. Ecological niches of bat viruses closely related to SARS-CoV-2 (SCoV2rCs) predicted using 4 points (a) (AUC = 0.58) and 21 points (b) (AUC = 0.96).** The geographic distributions of suitable environments were predicted using the Maxent algorithm in ENMTools (see Methods for details). Black circles indicate localities used to build the distribution model (see geographic coordinates in online supplementary table S1).

206  
207  
208  
209  
210  
211  
212

The ecological niche of bat SCoV2rCs was firstly predicted using the four localities where bat viruses were previously detected [1,6-8] (Fig. 5a). The highest probabilities of occurrence (highlighted in green in Fig. 5a) were found in Southeast Asia rather than in China. However, the

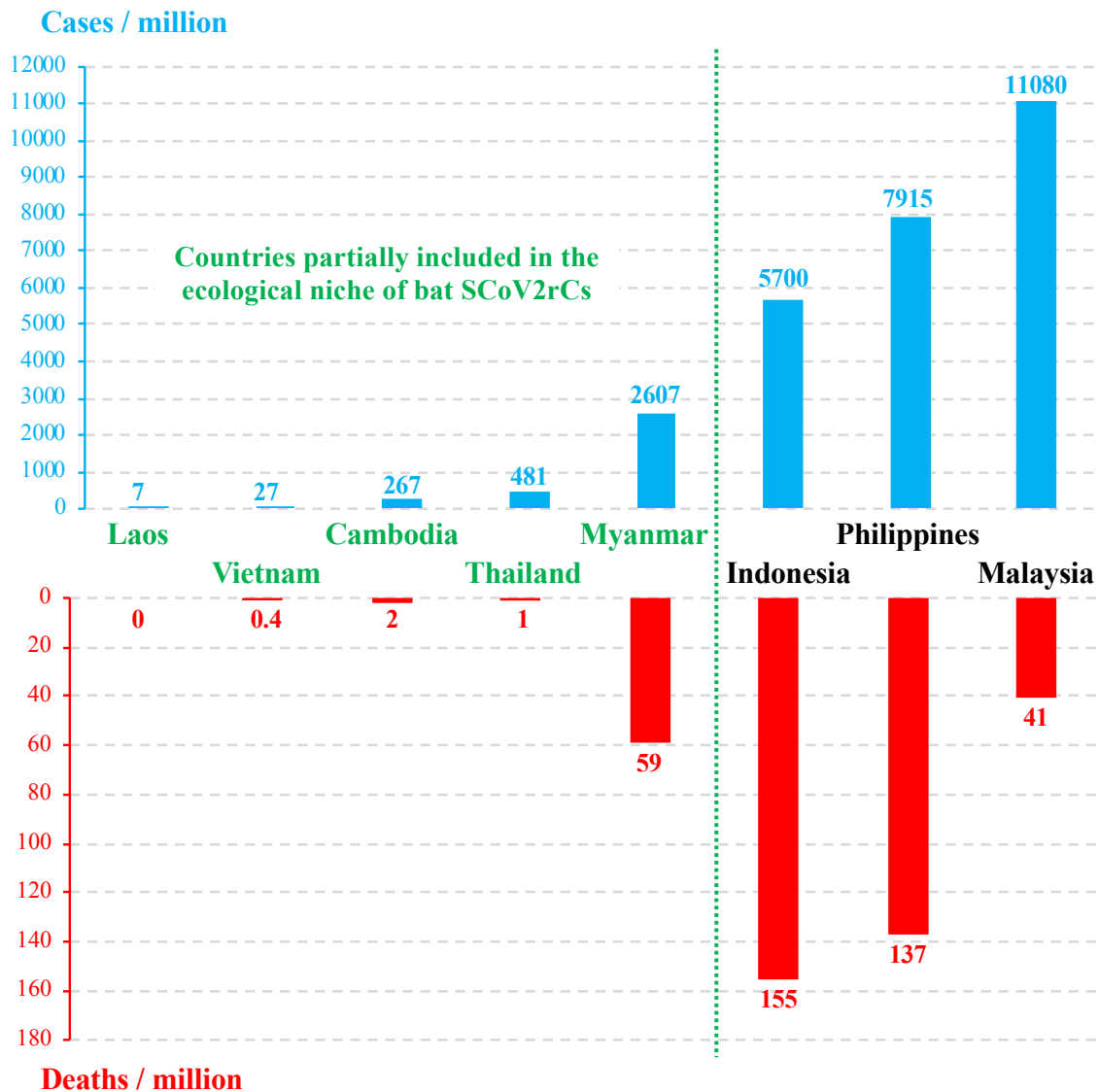
213  
214  
215

AUC was only 0.58, and the value was < 95% CI null-model's AUCs (0.74), indicating that the model was not statically supported at a significance level of 0.05 (see on line supplementary Fig. S2). As expected, these preliminary results confirmed that more than four records are needed to increase the accuracy of the distribution model [22]. For that reason, we used a genetic approach to determine geographic localities where bat SCoV2rCs are more likely to be found. The *COI* sequences of the nine bats in which a SCoV2rC was detected are shown by red arrows in Fig. 3. For *R. affinis*, the *COI* haplotype sequenced for the bat found positive for a SCoV2rC in southern Yunnan (site named Ch1 in Fig. 2 and 3) was not found in other sampled localities. For the three other bat species found positive for SCoV2rCs, identical *COI* sequences were detected in bats from 17 other geographic localities (see online supplementary table S1). For *R. acuminatus*, the four *COI* haplotypes sequenced for the bats found positive for SCoV2rCs in eastern Thailand (site named T4 in Fig. 2 and 3) were also found in four localities in Cambodia (C2, C4, C5, and C6), two localities in southern Laos (L10 and L11), and three localities in southern Vietnam (V13, V14, and V17). The results indicate high connectivity among *R. acuminatus* populations from eastern Thailand, Cambodia, southern Laos and southern Vietnam. For *R. malayanus*, the *COI* haplotype sequenced for the bat found positive for a SCoV2rC in southern Yunnan (site named Ch3 in Fig. 2 and 3) was also found in another locality in southern Yunnan (Ch2) and two localities in northern Laos (L1 and L3). The results indicate high connectivity among *R. malayanus* populations from southern China and northern Laos. For *R. shameli*, the single *COI* haplotype sequenced for the two bats found positive for SCoV2rCs in northern Cambodia (site named C3 in Fig. 2 and 3) was also found in three other localities in Cambodia (C1, C2, and C5) and five localities in Laos (L4, L6, L8, L9, and L10). The results indicate high connectivity among *R. shameli* populations from Cambodia and Laos. Based on these genetic data, the ecological niche of bat SCoV2rCs was predicted using 21 records corresponding to the four localities where bat viruses were previously detected [1,6-8] and the 17 localities where bats showed the same *COI* haplotype than virus-positive bats (data set B: 21 points; see online supplementary table S1 for details). The AUC was

0.96. The value was > 95% CI null-model's AUCs (0.81), indicating that the model performs significantly better than a random model (see online supplementary Fig. S3). The areas showing the highest probabilities of occurrence (highlighted in green in Fig. 5b) include southern Yunnan, northern and southern Laos, northwestern and southern Vietnam, northeastern and southwestern Cambodia, eastern, northern and western Thailand, and southern Myanmar.

Our results show that bat SCoVrCs and SCoV2rCs have different ecological niches: that of SCoVrCs covers mainly China and several adjacent countries and extends to latitudes between 18° and 43°N, whereas that of SCoV2rCs covers northern mainland Southeast Asia and extends to latitudes between 10° and 24°N. Most *Rhinolophus* species involved in the ecological niche of SCoVrCs have to hibernate in winter when insect populations become significantly less abundant. This is not the case for most *Rhinolophus* species involved in the ecological niche of SCoV2rCs. Since this ecological difference may be crucial for the dynamics of viral transmission among bat populations, it needs to be further studied through comparative field surveys in different regions of China and Southeast Asia.

The ecological niches of SCoVrCs and SCoV2rCs slightly overlap in the zone including southern Yunnan, northern Laos, and northern Vietnam (Fig. 4 and Fig. 5b). This zone corresponds to the northern edge of tropical monsoon climate [23]. Highly divergent sarbecoviruses of the two main lineages SCoVrCs and SCoV2rCs are expected to be found in sympatry in this area. This is confirmed by the discovery of both SCoVrCs and SCoV2rCs in horseshoe bats collected in southern Yunnan [1,6,21]. Collectively, these data suggest that genomic recombination between viruses of the two divergent lineages are more likely to occur in bats roosting, at least seasonally, in the caves of these regions. Since highly recombinant viruses can threaten the benefit of vaccination campaigns, southern Yunnan, northern Laos, and northern Vietnam should be the targets of closer surveillance.



**Figure 6. Number of Covid-19 patients per million inhabitants (in blue) and deaths per million inhabitants (in red) for the different countries of Southeast Asia.** Data extracted from the Worldometers website [2] on April 12, 2021.

269  
270  
271  
272  
273

Chinese researchers have actively sought sarbecoviruses in all Chinese provinces after the 2002- 274  
2004 SARS outbreak. They found many bat SCoVrCs [16,20,21] but only two SCoV2rCs [1,5] 275  
and both of them were discovered in southern Yunnan, the Chinese province bordering Southeast 276  
Asia. The ecological niches predicted herein for bat sarbecoviruses suggest that SCoVrCs are 277  
dominant in China (Fig. 4) while SCoV2rCs are present mostly in Southeast Asia (Fig. 5). This 278  
means that viruses similar to SARS-CoV-2 have been circulating for several decades throughout 279

Southeast Asia, and that different species of bats have exchanged these viruses in the caves they inhabit. The data available on human cases and deaths caused by the Covid-19 pandemic support the hypothesis that the cradle of diversification of bat SCoV2rCs is mainland Southeast Asia, and in particular the areas highlighted in green in Fig. 5b. Indeed, human populations in Cambodia, Laos, Thailand, and Vietnam appear to be much less affected by the Covid-19 pandemic than other countries of the region, such as Indonesia, Malaysia, Myanmar, and the Philippines (Fig. 6). This suggests that some human populations of Cambodia, Laos, Thailand, and Vietnam, in particular rural populations living in contact with wild animals for several generations, have a better immunity against SCoV2rCs because they have been regularly contaminated by bats and/or infected secondary hosts such as pangolins.

#### **Pangolins contaminated by bats in Southeast Asia**

Apart from bats, the Sunda pangolin (*Manis javanica*) and Chinese pangolin (*Manis pentadactyla*) are the only wild animals in which viruses related to SARS-CoV-2 have been found so far. However, these discoveries were made in a rather special context, that of pangolin trafficking. Several sick pangolins were seized by Chinese customs in Yunnan province in 2017 (unpublished data), in Guangxi province in 2017-2018 [24] and in Guangdong province in 2019 [25]. Even if the viruses sequenced in pangolins are not that close to SARS-CoV-2 (one was 85% identical and the other 90%), they indicate that at least two sarbecoviruses could have been imported into China well before the emergence of Covid-19 epidemic. Indeed, it has been shown that Sunda pangolins collected from different Southeast Asian regions have contaminated each other while in captivity on Chinese territory [3]. The question remained on how the Sunda pangolins became infected initially. Could it have been in their natural Southeast Asian environment, before being captured? The discovery of two new viruses close to SARS-CoV-2 in bats from Cambodia and Thailand supports this hypothesis, as *Rhinolophus* bats and pangolins can meet, at least occasionally, in



forests of Southeast Asia, possibly in caves, tree hollows or burrows. Further substantiating this hypothesis, the geographic distribution of *Manis javanica* [26] overlaps the ecological niche here predicted for bat SCoV2rCs (Fig. 5), and SARS-CoV-2 neutralizing antibodies have been recently detected in a Sunda pangolin collected in eastern Thailand [6]. Collectively, these data strengthen the hypothesis that pangolin trafficking is responsible for multiple exports of viruses related to SARS-CoV-2 to China [3].

## Methods

### DNA extraction and sequencing

One hundred and forty-four tissue samples of morphologically identified bats of *R. acuminatus* (n = 10), *R. affinis* (n = 57), *R. malayanus* (n = 14), and *R. shameli* (n = 63) were specially analysed for this study. In the field, bats were captured with mist nets and harp-traps and handled according to guidelines and recommendations of the American Society of Mammologists [27]. They were measured, photographed and identified by the authors. Tissue samples were taken from the chest muscles of voucher specimens or from the patagium (biopsy punches; 2 mm diameter) of released bats. Samples were preserved in 95% ethanol.

Total DNA was extracted using QIAGEN DNeasy Tissue Kit (Qiagen, Germany) in accordance with the manufacturer's instructions. The barcode fragment of the *COI* gene (657 bp) was amplified and sequenced using the primers UTyr and C1L705 [28]. PCR amplifications of the *COI* gene were performed as previously published [29]. PCR products were purified using ExoSAP Kit (GE Healthcare, UK) and sequenced using the Sanger method on an ABI 3730 automatic sequencer at the Centre National de Séquençage (Genoscope) in Evry (France). Haplotypes were assembled with forward and reverse electropherograms using Sequencher 5.1 (Gene Codes Corporation, Ann Arbor, MI, USA). No gaps and stop codons were found in the *COI* sequences after translation into amino-acids. Sequences generated for this study were deposited in the GenBank database (accession numbers MW712891-MW713034) (see online supplementary table S1).

331

**Analyses of CO1 sequences**

332

Our sequences were aligned with 199 additional *COI* sequences downloaded from GenBank. Note 333

that the *COI* sequences of seven bats found positive for viruses closely related to SARS-CoV-2 334

[1,5,6] were assembled on Geneious® 10.2.2 (Biomatters Ltd., Auckland, New Zealand) by 335

mapping available SRA data to a *COI* reference. Sequences were aligned using AliView 1.22 [30]. 336

Our final *COI* alignments contain 37 sequences for *R. acuminatus*, 44 sequences for *R. malayanus*, 337

82 sequences for *R. shameli*, and 180 sequences for *R. affinis*. These four alignments were analysed 338

in PopART 1.5 [13] to construct haplotype networks using the median joining method with equal 339

weights for all mutations. The 62 localities where bats were sampled are shown in the map of Fig. 340

2 and their geographic coordinates are detailed in online supplementary table S1. 341

342

**Prediction of ecological niches**

343

For bat SCoVrCs, the ecological niche was inferred using GPS data collected for viruses published 344

during the last two decades. The list of the 19 available geographic records is provided in online 345

supplementary table S2. For bat SCoV2rCs, the ecological niche was initially predicted using the 346

four geographic localities where viruses were previously detected [1,5-7]: two in Yunnan, one in 347

northern Cambodia, and one in eastern Thailand (data set A). However, the use of only four records 348

is questionable since Van Proosdij et al. [22] have estimated that a minimum of 13 records is 349

required to develop accurate distribution models for widespread taxa. For that reason, we used a 350

genetic approach to increase the number of geographic records. Since the detection of identical 351

*COI* sequences in different bat populations is indicative of recent dispersal events of females, we 352

also selected the 17 geographic records where bats showed the same *COI* haplotypes than virus- 353  
positive bats (data set B: 21 points; see online supplementary table S1). 354

For each of the three data sets (bat SCoVrCs; data sets A and B for bat SCoV2rCs), the 19 355  
bioclimatic variables available in the WorldClim database [31] were studied for an area 356  
corresponding to the minimum and maximum latitudes and longitudes of the selected points (19 357  
points for bat SCoVrCs; 4 and 21 points, respectively for the SCoV2rCs data sets A and B) and the 358  
caret R package [32] was used to determine the least correlated variables ( $|r| < 0.7$ ) [33]. For bat 359  
SCoVrCs, the following five predictor bioclimatic variables were retained: Bio3 (isothermality), 360  
Bio4 (temperature seasonality), Bio5 (maximum temperature of the warmest month), Bio15 361  
(precipitation seasonality), and Bio18 (precipitation of the warmest quarter). For data set A, the 362  
following seven predictor bioclimatic variables were retained: Bio3, Bio7 (temperature annual 363  
range), Bio10 (mean temperature of the warmest quarter), Bio13 (precipitation of the wettest 364  
month), Bio14 (precipitation of driest month), Bio15, and Bio18. For data set B, the following 365  
seven predictor bioclimatic variables were selected: Bio2 (mean diurnal range), Bio3, Bio7, Bio10, 366  
Bio13, Bio15, Bio17 (precipitation of the driest quarter), and Bio18. Ecological niche modelling 367  
was performed with the MaxEnt algorithm using ENMTools in R [34]. The MaxEnt approach was 368  
chosen for its ability to work with presence-only data sets and to produce results with a low sample 369  
size [35]. The area under the curve (AUC) of the receiver operating characteristic plot was used as 370  
a first measure of model accuracy, a value of 0.5 indicating model accuracy not better than random, 371  
and a value of 1 indicating perfect model fit [36,37]. To test for sampling bias, the distribution 372  
model using all selected localities was tested against a null model developed by 1000 times drawing 373  
an equal number of random points from the entire study area [37]. The position of the AUC value 374

was tested against the 95% confidence interval (CI) of the 1000 AUC values of the null-models. If 375  
the AUC value is  $\geq$  95% CI null-model's AUCs, the model is considered performing significantly 376  
better than a random model [37]. 377

## References 379

1. Zhou, P. *et al.* A pneumonia outbreak associated with a new coronavirus of probable bat origin. 380  
*Nature* **579**, 270-273; 10.1038/s41586-020-2012-7 (2020). 381
2. COVID-19 Coronavirus pandemic. <https://www.worldometers.info/coronavirus/>. downloaded 382  
on April 12, 2021. 383
3. Hassanin, A., Grandcolas, P. & Veron, G. Covid-19: natural or anthropic origin? *Mammalia* 384  
**85**, 1-7; 10.1515/mammalia-2020-0044 (2021). 385
4. Walker, P.J. *et al.* Changes to virus taxonomy and the statutes ratified by the international 386  
committee on taxonomy of viruses. *Arch Virol.* **165**, 2737-2748; 10.1007/s00705-020-04752- 387  
x (2020). 388
5. Zhou, H. *et al.* A novel bat coronavirus closely related to SARS-CoV-2 contains natural 389  
insertions at the S1/S2 cleavage site of the spike protein. *Curr. Biol.* **30**, 3896; 390  
10.1016/j.cub.2020.09.030 (2020). 391
6. Wacharapluesadee, S. *et al.* Evidence for SARS-CoV-2 related coronaviruses circulating in 392  
bats and pangolins in Southeast Asia. *Nat. Commun.* **12**, 972; 10.1038/s41467-021-21240-1 393  
(2021). 394
7. Hul, V. *et al.* A novel SARS-CoV-2 related coronavirus in bats from Cambodia. 395  
10.1101/2021.01.26.428212 (submitted). 396
8. Li, W. *et al.* Bats are natural reservoirs of SARS-like coronaviruses. *Science* **310**, 676-679; 397  
10.1126/science.1118391 (2005). 398

9. Drexler, J.F. *et al.* Genomic characterization of severe acute respiratory syndrome-related coronavirus in European bats and classification of coronaviruses based on partial RNA-dependent RNA polymerase gene sequences. *J. Virol.* **84**, 11336-11349; 10.1128/JVI.00650-10 (2010). 399  
400  
401  
402
10. Tao, Y. & Tong, S. Complete genome sequence of a severe acute respiratory syndrome-related coronavirus from Kenyan bats. *Microbiol. Resour. Announc.* **8**, e00548-19; 10.1128/MRA.00548-19 (2019). 403  
404  
405
11. The IUCN Red List of Threatened Species. Version 2020-3. <https://www.iucnredlist.org>. Downloaded on 23 February 2021. 406  
407
12. Boni, M.F. *et al.* Evolutionary origins of the SARS-CoV-2 sarbecovirus lineage responsible for the COVID-19 pandemic. *Nat. Microbiol.* **5**, 1408–1417; 10.1038/s41564-020-0771-4 (2020). 408  
409  
410
13. Leigh, J.W. & Bryant, D. PopART: Full-feature software for haplotype network construction. *Methods Ecol. Evol.* **6**, 1110-1116; 10.1111/2041-210X.12410 (2015). 411  
412
14. Thong, V.D., Thanh, H.T., Soisook, P. & Csorba, G. *Rhinolophus acuminatus*. The IUCN Red List of Threatened Species e.T19520A21974227; 10.2305/IUCN.UK.2019-3.RLTS.T19520A21974227.en. (2019). 413  
414  
415
15. Ith, S. *et al.* Taxonomic implications of geographical variation in *Rhinolophus affinis* (Chiroptera: Rhinolophidae) in mainland Southeast Asia. *Zool. Stud.* **54**, e31; 10.1186/s40555-015-0109-8 (2015). 416  
417  
418
16. Burgin, C.J., Wilson, D.E., Mittermeier, R.A., Rylands, A.B., Lacher, T.E., Sechrest, W. Illustrated Checklist of the Mammals of the World. Vol. 2. Lynx Edicions, Barcelona (2020). 419  
420
17. Fan, Y., Zhao, K., Shi, Z. L. & Zhou, P. Bat coronaviruses in China. *Viruses* **11**, 210; 10.3390/v11030210 (2019). 421  
422
18. Francis, C. Field Guide to the Mammals of South-east Asia. Bloomsbury, London, UK. (2019). 423

19. Smith, A.T. & Xie, Y. A Guide to the Mammals of China. Princeton University Press, 424  
Princeton, New Jersey (2008). 425
20. Han, Y. *et al.* Identification of diverse bat alphacoronaviruses and betacoronaviruses in China 426  
provides new insights into the evolution and origin of coronavirus-related diseases. *Front.* 427  
*Microbiol.* **10**, 1900; 10.3389/fmicb.2019.01900 (2019). 428
21. Hu, D. *et al.* Genomic characterization and infectivity of a novel SARS-like coronavirus in 429  
Chinese bats. *Emerg. Microbes Infect.*, **7**, 154; 10.1038/s41426-018-0155-5 (2018). 430
22. van Proosdij, A.S.J., Sosef, M.S.M., Wieringa, J.J. & Raes N. Minimum required number of 431  
specimen records to develop accurate species distribution models. *Ecography* **39**, 542-552; 432  
10.5061/dryad.8sb8v (2016). 433
23. Song, Y., Achberger, C. & Linderholm, H.W. Rain-season trends in precipitation and their 434  
effect in different climate regions of China during 1961–2008. *Environ. Res. Lett.* **6**, 034025; 435  
10.1088/1748-9326/6/3/034025 (2011). 436
24. Lam, T.T. *et al.* Identifying SARS-CoV-2-related coronaviruses in Malayan pangolins. *Nature* 437  
**583**, 282-285; 10.1038/s41586-020-2169-0 (2020). 438
25. Liu, P., Chen, W. & Chen, J.P. Viral metagenomics revealed Sendai virus and coronavirus 439  
infection of Malayan pangolins (*Manis javanica*). *Viruses* **11**, 979; 10.3390/v11110979 440  
(2019). 441
26. Challender, D. *et al.* *Manis javanica*. The IUCN Red List of Threatened Species 2019: 442  
e.T12763A123584856; 10.2305/IUCN.UK.2019-3.RLTS.T12763A123584856.en. (2019). 443
27. Sikes, R.S., Gannon W.L., and the animal care and use committee of the American Society of 444  
Mammalogists. Guidelines of the American Society of Mammalogists for the use of wild 445  
mammals in research. *J. Mammal.* **92**, 235–253; 10.1644/10-MAMM-F-355.1 (2011). 446
28. Hassanin, A. *et al.* Pattern and timing of diversification of Cetartiodactyla (Mammalia, 447  
Laurasiatheria), as revealed by a comprehensive analysis of mitochondrial genomes. *C.R. biol.* 448  
**335**, 32–50; 10.1016/j.crv.2011.11.002 (2012). 449

29. Tu, V.T. *et al.* Integrative taxonomy of the *Rhinolophus macrotis* complex (Chiroptera, Rhinolophidae) in Vietnam and nearby regions. *J. Zool. Syst. Evol. Res.* **55**, 177-198; 10.1111/jzs.12169 (2017).
30. Larsson, A. AliView: a fast and lightweight alignment viewer and editor for large data sets. *Bioinformatics* **30**, 3276-3278; 10.1093/bioinformatics/btu531 (2014).
31. Hijmans, R.J., Cameron, S.E., Parra, J.L., Jones, P.G. & Jarvis, A. Very high resolution interpolated climate surfaces for global land areas. *Int. J. Climatol.* **25**, 1965-1978; 10.1002/joc.1276 (2005).
32. Kuhn, M. Building predictive models in R using the caret package. *J. Stat. Softw.* **28**, 1-26; 10.18637/jss.v028.i05 (2008).
33. Dormann, C.F. *et al.* Collinearity: a review of methods to deal with it and a simulation study evaluating their performance. *Ecography* **36**, 27-46; 10.1111/j.1600-0587.2012.07348.x (2013).
34. Warren, D.L. *et al.* ENMTools 1.0: an R package for comparative ecological biogeography. *Ecography* **44**, 504-511; 10.1111/ecog.05485 (2021).
35. Wisz, M.S. *et al.* Effects of sample size on the performance of species distribution models. *Divers. Distrib.* **14**, 763-773; 10.1111/j.1472-4642.2008.00482.x (2008).
36. Fielding, A.H. & Bell, J.F. A review of methods for the assessment of prediction errors in conservation presence/absence models. *Environ. Conserv.* **24**, 38-49; 10.1017/S0376892997000088 (1997).
37. Raes, N. & ter Steege, H. A null-model for significance testing of presence-only species distribution models. *Ecography* **30**, 727-736; 10.1111/j.2007.0906-7590.05041.x (2007).

## Acknowledgements

476

In Cambodia, we would like to acknowledge T.D. Vong, V. Hul and S. In for their support during 477  
the field surveys, and C. Garnero-Morena, G. Diffloth, B. Kilian, A. Beschaouch and N. Hayashi- 478  
Denis for their assistance. The field mission was organized by UNESCO under the permissions of 479  
the National Authority for Preah Vihear, the President, His Excellency Mr. Chuch Phoeurn and the 480  
Director of the Preah Vihear temple, Prof. Hang Soth. In Laos, we would like to thank N. Beevang, 481  
J. Fuchs, E. Ifticene, and S. Xaythany for their support during the field surveys, and C. Bear Khue, 482  
F. Bouvier, C. Hatten, R. Humphrey, A.M. Ohler, P. Rousseau, Mr. Tipavanh, M. Thoummabouth, 483  
K. Valakone, Y. Varelides, and A.G. Verdier for their assistance. The fieldwork was supported and 484  
organized by the UNESCO and « La Maison du Patrimoine » (World Heritage House) of Luang 485  
Prabang, and the Phongsaly Forest Conservation and Rural Development Project, a Lao-European 486  
cooperation. Survey permission and authorization for tissue samples collecting were granted by the 487  
Ministry of Agriculture and Forestry, Department of Livestock and Fisheries. In Vietnam, we 488  
would like to acknowledge P.D. Tien (IEBR), D.Q. Thang and N.X. Nghia (Ngoc Linh NR) and 489  
N.T. Son (Vu Quang NP) for their support during the field surveys. We are grateful to L.X. Canh, 490  
T.H. Thai, N.V. Sinh and other colleagues of the IEBR, Hanoi for administrative assistance. The 491  
field research was done under the permissions of the People's Committees of numerous provinces 492  
and the Vietnamese Ministry of Agriculture and Rural Development (Vietnam Administration of 493  
Forestry). The fieldwork was also supported by the "Société des Amis du Muséum et du Jardin des 494  
Plantes" and the National Research, Development and Innovation Fund of Hungary NKFIH 495  
KH130360. M.C. is funded by the SCOR Corporate foundation for Science. We are grateful to C. 496  
Bonillo, C. Ferreira, J. Lambourdière, and J. Utge (UMS 2700, MNHN) for their technical 497  
assistance. We would like to thank Huw Jones and Anne Ropiquet for helpful comments on the 498  
first version of the manuscript. This research was funded by the AAP RA-COVID-19, grant number 499  
ANR-21-CO12-0002, the CNRS, the MNHN, the INRA and the CEA (Genoscope). 500

501



<b>Author Contributions</b>	502
A.H. wrote the main manuscript text; V.T.T. and A.H. sequenced the CO1 barcodes; V.T.T. and	503
A.H. prepared table S1; A.H. prepared figures 1, 2, 3, and 6, and table S2; M.C. and A.H. prepared	504
figures 4, 5, and S1-S3; A.H, G.C., and V.T.T. collected bats in different localities of Southeast	505
Asia. All authors reviewed the manuscript.	506
	507
<b>Competing interests</b>	508
The author(s) declare no competing interests.	509
	510
<b>Data Availability:</b> DNA sequences generated for this study were deposited in the GenBank	511
database (accession numbers MW712891-MW713034).	512
	513
<b>Ethical statement:</b> Ethical review and approval were not available for our study because the field	514
missions were carried out between 2004 and 2011, i.e., before the creation of the ethical committee	515
at the Muséum national d’Histoire naturelle. However, the field studies were carried out in	516
compliance with the ARRIVE guidelines.	517

## Figures

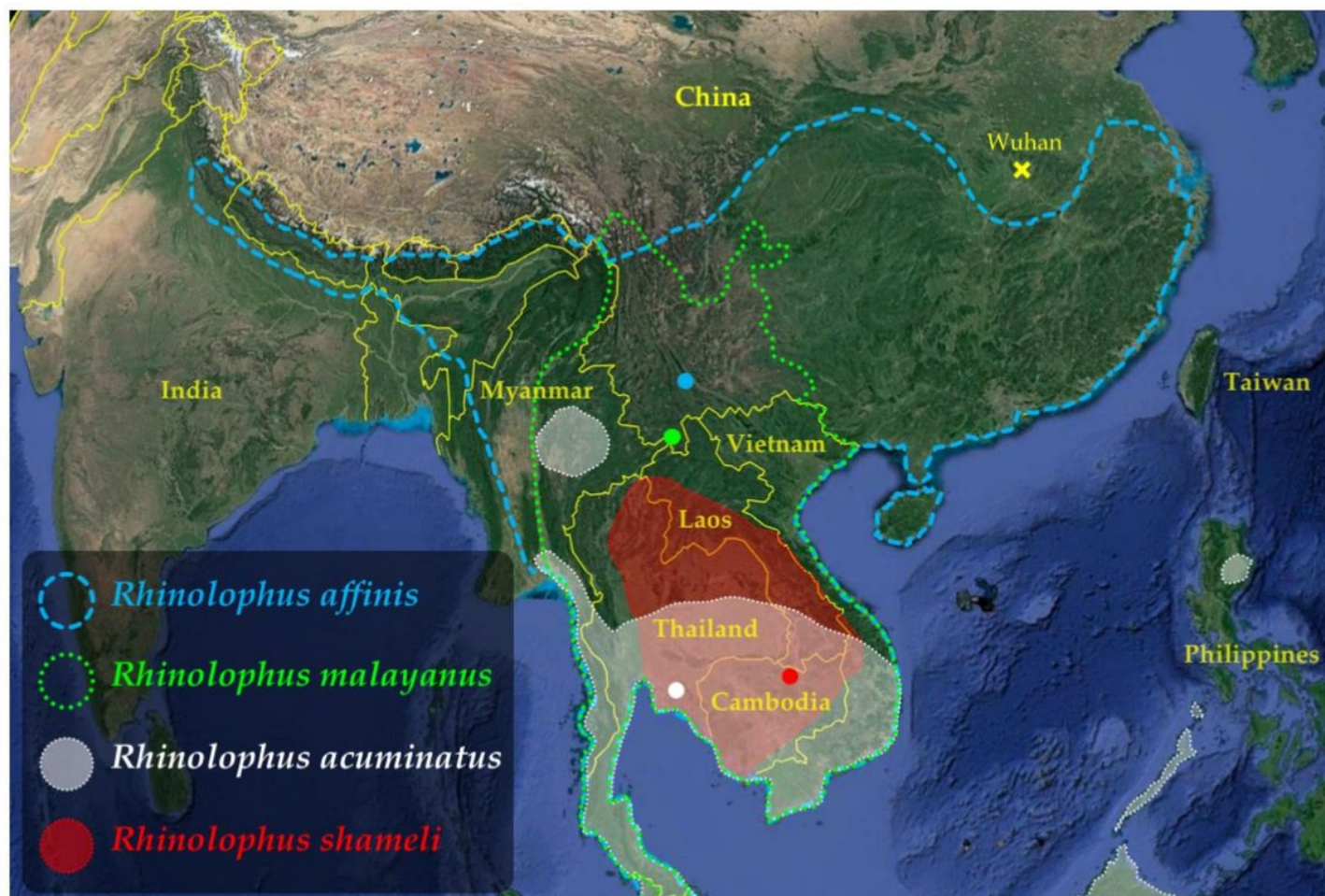
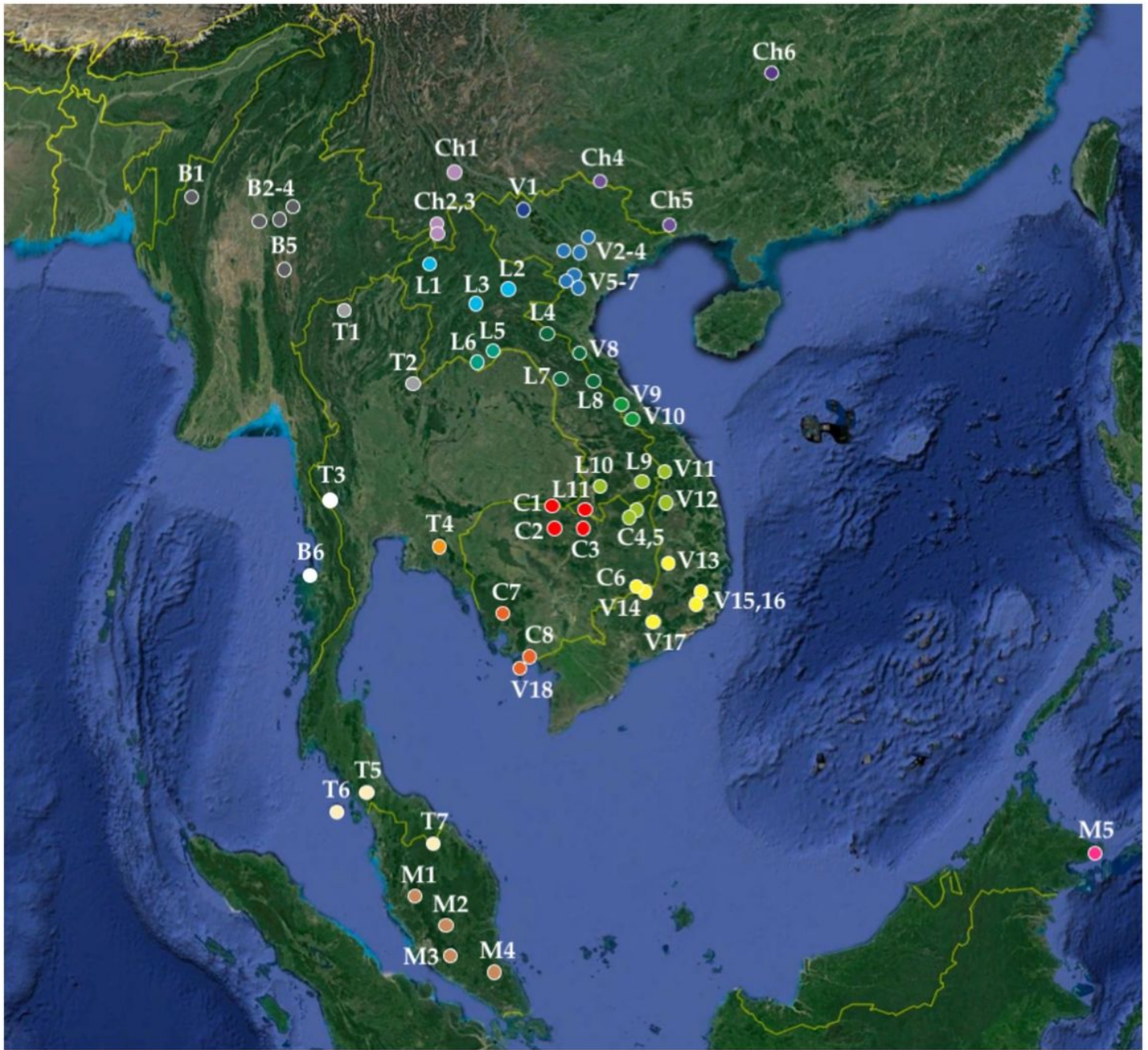


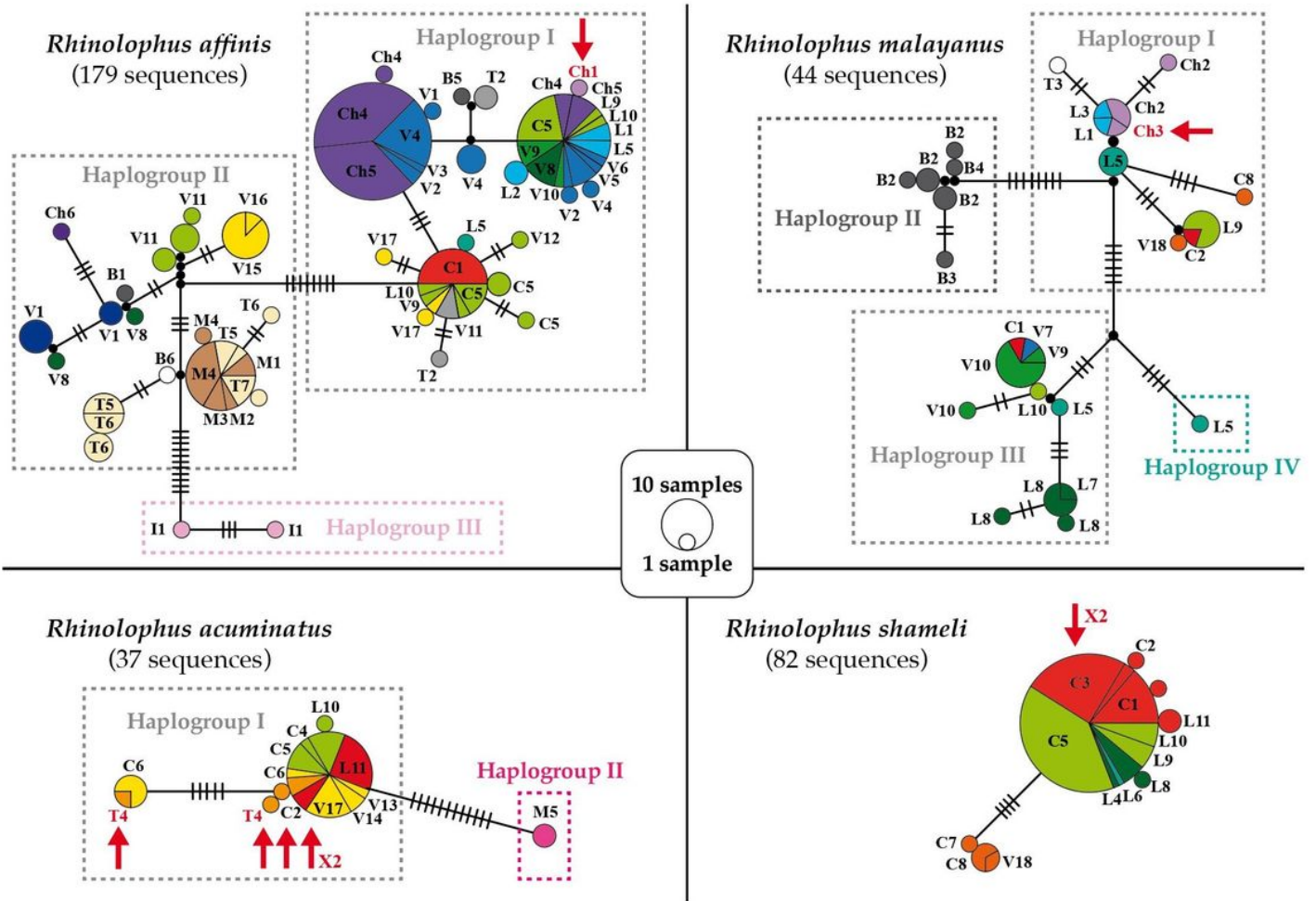
Figure 1

Geographic distribution of the four *Rhinolophus* species found positive for viruses closely related to SARS-CoV-2 in southern China and Southeast Asia. Map from Google Earth US Dept of State Geographer © 2020 Google – Image Landsat / Copernicus - Data SIO, NOAA, U.S. Navy, NGA, GEBCO. For each of the four *Rhinolophus* species, the geographic distribution was extracted from the IUCN website [11]. The coloured dots show the four geographic locations where bats found positive for SCoV2rCs were collected. Note: The designations employed and the presentation of the material on this map do not imply the expression of any opinion whatsoever on the part of Research Square concerning the legal status of any country, territory, city or area or of its authorities, or concerning the delimitation of its frontiers or boundaries. This map has been provided by the authors.



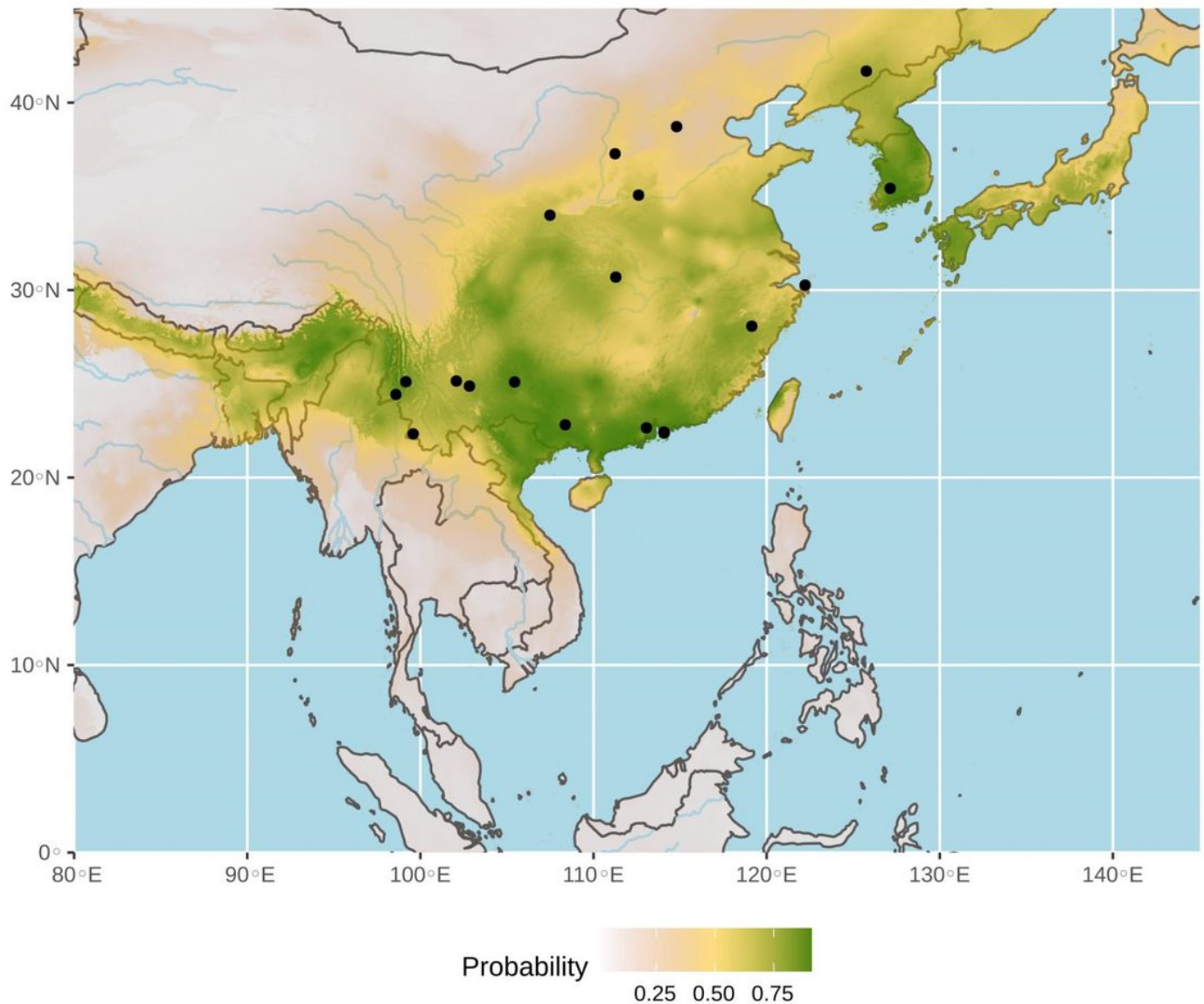
**Figure 2**

Geographic localities analysed in this study. The codes used for the countries are the following: B (Myanmar), C (Cambodia), Ch (China), I (Indonesia), L (Laos), M (Malaysia), T (Thailand), and V (Vietnam). Different regions were highlighted using colours. Map from Google Earth US Dept of State Geographer © 2020 Google – Image Landsat / Copernicus - Data SIO, NOAA, U.S. Navy, NGA, GEBCO. The locality names and GPS coordinates are provided in online supplementary table S1 Note: The designations employed and the presentation of the material on this map do not imply the expression of any opinion whatsoever on the part of Research Square concerning the legal status of any country, territory, city or area or of its authorities, or concerning the delimitation of its frontiers or boundaries. This map has been provided by the authors.



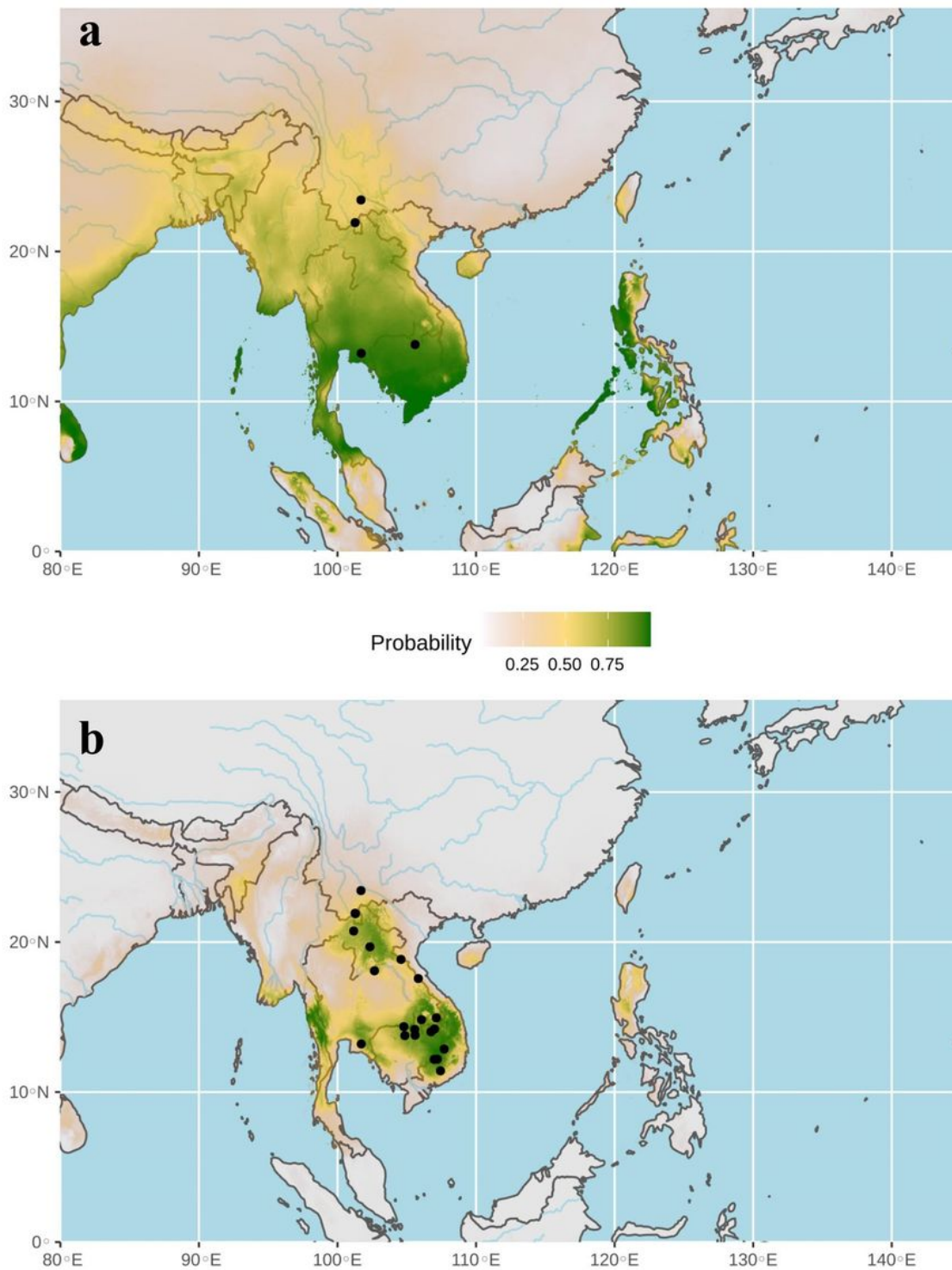
**Figure 3**

Haplotype networks based on CO1 sequences of the four *Rhinolophus* species found positive for viruses closely related to SARS-CoV-2 (SCoV2rCs). The networks were constructed with the median joining method available in PopART 1.5 [13]. The codes used for the countries are the following: B (Myanmar), C (Cambodia), Ch (China), I (Indonesia), L (Laos), M (Malaysia), T (Thailand), and V (Vietnam). Colours indicate the geographic origin of haplotypes according to Fig. 2 (see online supplementary table S1). The circles indicate haplotypes separated by at least one mutation. The black lines on the branches show the number of mutations  $\geq 2$ . Black circles represent missing haplotypes. Circle size is proportional to the number of haplotypes. Haplogroups separated by more than seven mutations (pairwise nucleotide distances  $> 1\%$ ) are highlighted by dotted lines. The red arrows show the positions of the nine bats found positive for SCoV2rCs.



**Figure 4**

Ecological niche of bat viruses related to SARS-CoV (SCoVrCs). The geographic distribution of suitable environments was predicted using the Maxent algorithm in ENMTools (see Methods for details). AUC = 0.81. Black circles indicate localities used to build the distribution model (see geographic coordinates in online supplementary table S2). Note: The designations employed and the presentation of the material on this map do not imply the expression of any opinion whatsoever on the part of Research Square concerning the legal status of any country, territory, city or area or of its authorities, or concerning the delimitation of its frontiers or boundaries. This map has been provided by the authors.



**Figure 5**

Ecological niches of bat viruses closely related to SARS-CoV-2 (SCoV2rCs) predicted using 4 points (a) (AUC = 0.58) and 21 points (b) (AUC = 0.96). The geographic distributions of suitable environments were predicted using the Maxent algorithm in ENMTools (see Methods for details). Black circles indicate localities used to build the distribution model (see geographic coordinates in online supplementary table S1). Note: The designations employed and the presentation of the material on this map do not imply the

expression of any opinion whatsoever on the part of Research Square concerning the legal status of any country, territory, city or area or of its authorities, or concerning the delimitation of its frontiers or boundaries. This map has been provided by the authors.

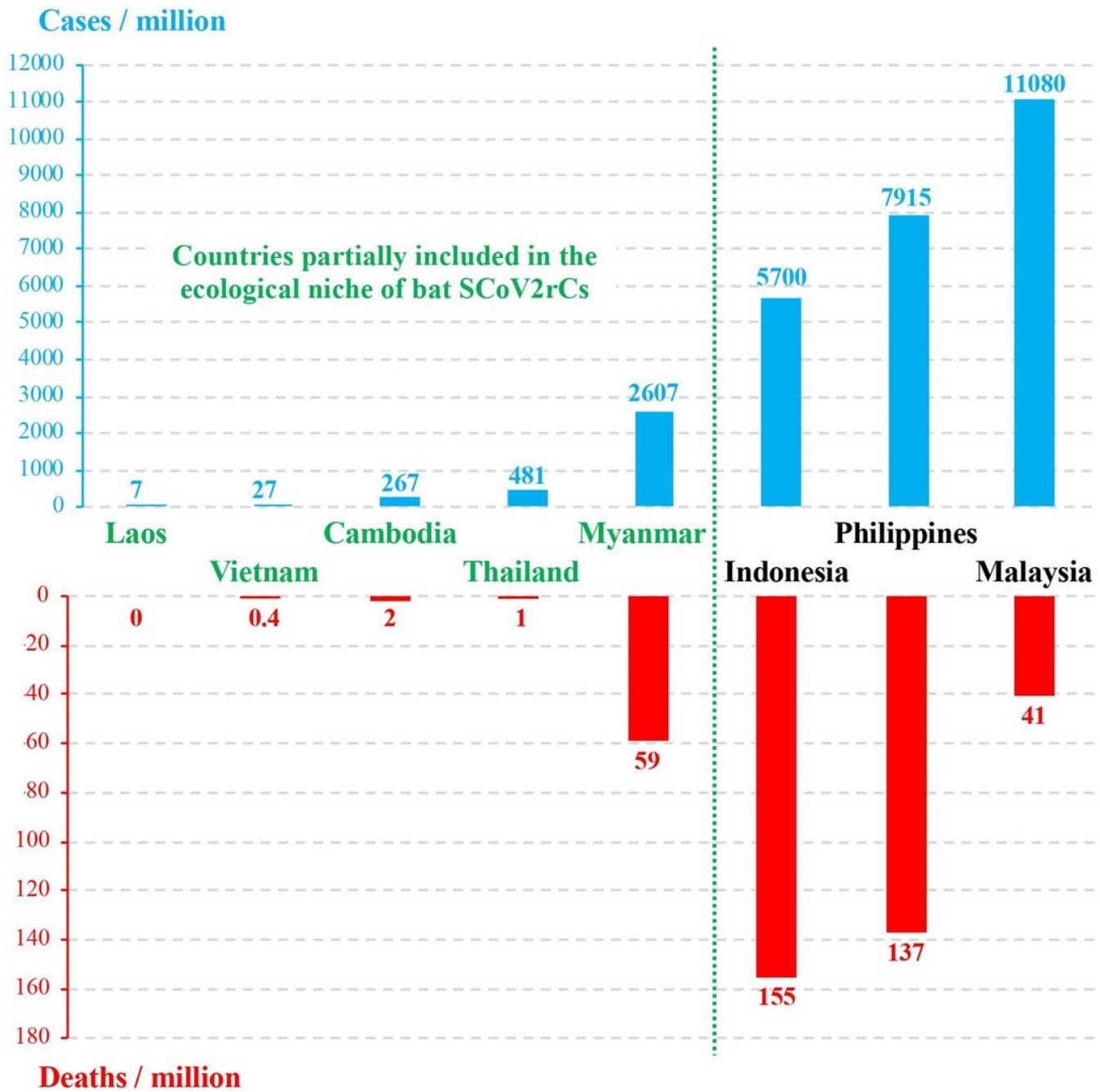


Figure 6

Number of Covid-19 patients per million inhabitants (in blue) and deaths per million inhabitants (in red) for the different countries of Southeast Asia. Data extracted from the Worldometers website [2] on April 12, 2021

## Supplementary Files

This is a list of supplementary files associated with this preprint. Click to download.

- [SupplementaryInformation.pdf](#)