

Gain, Preservation, and Loss of a Group 1a Coronavirus Accessory Glycoprotein[∇]

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Coronaviruses are positive-strand RNA viruses of extraordinary genetic complexity and diversity. In addition to a common set of genes for replicase and structural proteins, each coronavirus may carry multiple group-specific genes apparently acquired through relatively recent heterologous recombination events. Here we describe an accessory gene, ORF3, unique to canine coronavirus type I (CCoV-I) and characterize its product, glycoprotein gp3. Whereas ORF3 is conserved in CCoV-I, only remnants remain in CCoV-II and CCoV-II-derived porcine and feline coronaviruses. Our findings provide insight into the evolutionary history of coronavirus group 1a and into the dynamics of gain and loss of accessory genes.

Coronaviruses (CoVs), enveloped positive-strand RNA viruses of human clinical and veterinary relevance, are exceptional in terms of genetic complexity and variety. With genome sizes of ~30 kb, they are the largest RNA viruses known thus far (17, 42). One major contributing factor to CoV diversity is high-frequency RNA recombination (1, 28, 33). New sero- and biotypes have arisen from homologous RNA recombination, i.e., the exchange of corresponding sequences among related CoVs (3, 21, 25, 27, 29, 43), while heterologous RNA recombination events with noncoronaviral donor RNAs have led to the acquisition of novel genes (31, 44, 56).

All CoVs have a similar genome organization with a common set of five genes arranged in a conserved order (10, 12). The polymerase gene, occupying the 5'-most 70% of the genome, encodes the replicase polyproteins from which up to 16 mature products are derived as well as an unknown number of functional processing intermediates (58). Downstream of the polymerase gene and expressed through a 3'-coterminal nested set of subgenomic (sg) mRNAs are the genes for the structural proteins S, E, M, and N (12). In addition, each CoV may possess up to seven group-specific "accessory" genes that are also expressed from sg mRNAs (34). In most cases, the functions of the accessory gene products are not known, and in general, they are not essential for replication in cultured cells (6, 36, 41, 53–55). Quite the opposite, their expression might decrease viral fitness *in vitro*, and mutants with inactivated accessory genes readily become selected during serial passage (22, 30, 45, 51). In field strains, however, accessory genes as a rule are maintained (13, 22, 43), and their loss—either through spontaneous mutation (37) or by design via reversed genet-

ics—generally causes loss of virulence in the natural host (11, 20, 36).

CoVs can be divided into three main phylogenetic groups (16). Canine coronaviruses (CCoVs), common enteric pathogens of dogs (8, 47), belong to subgroup 1a together with feline coronaviruses types I and II (FCoV-I and FCoV-II, respectively) and transmissible gastroenteritis virus (TGEV) of swine (16, 18). Like FCoV, CCoV-I and CCoV-II, respectively) sharing ~90% sequence identity in most of their genome (A. Lorusso, N. Decaro, C. Buonavoglia, and R. J. de Groot, unpublished data). In the coding region for the S ectodomain, however, sequence identity is only 56%.

The evolutionary history of CoV group 1a is not completely understood, but it apparently entailed multiple homologous and heterologous recombination events. The available data suggest that CCoV-I and FCoV-I arose by linear descent from a common ancestor and that recombination of CCoV-I with an unknown CoV led to acquisition of a new S gene, thus giving rise to CCoV-II (Lorusso et al., unpublished). In turn, CCoV-II strains donated this S gene and flanking sequences in recombinational exchanges to FCoV-II strains, leading to the independent emergence of FCoV-II strains (21). TGEV also appears to be of CCoV-II origin; in phylogenetic analyses of the genomic region downstream of the S gene, TGEV consistently clusters with extant CCoV-II field strains (7).

To date, the FCoVs and CCoVs described share the same complement of accessory genes, three of which (the "ORF3abc cluster") are located between the S and E genes (10, 14, 24, 46, 50, 52). The two remaining ones, ORF7a and ORF7b, are located at the 3' end of the genome (9, 24, 50) (Fig. 1a). In TGEV and related porcine CoVs, ORF3b is inactivated and ORF7b is lacking. Here we describe a novel functional accessory gene unique to CCoV-I and discuss the implications of our findings for our understanding of the evolutionary history of group 1a CoVs.

ORF3, a functional accessory gene unique to CCoV-I. During sequence analysis of CCoV-I variant Elmo/02 (39), we discovered, immediately downstream of the S gene, a 624-

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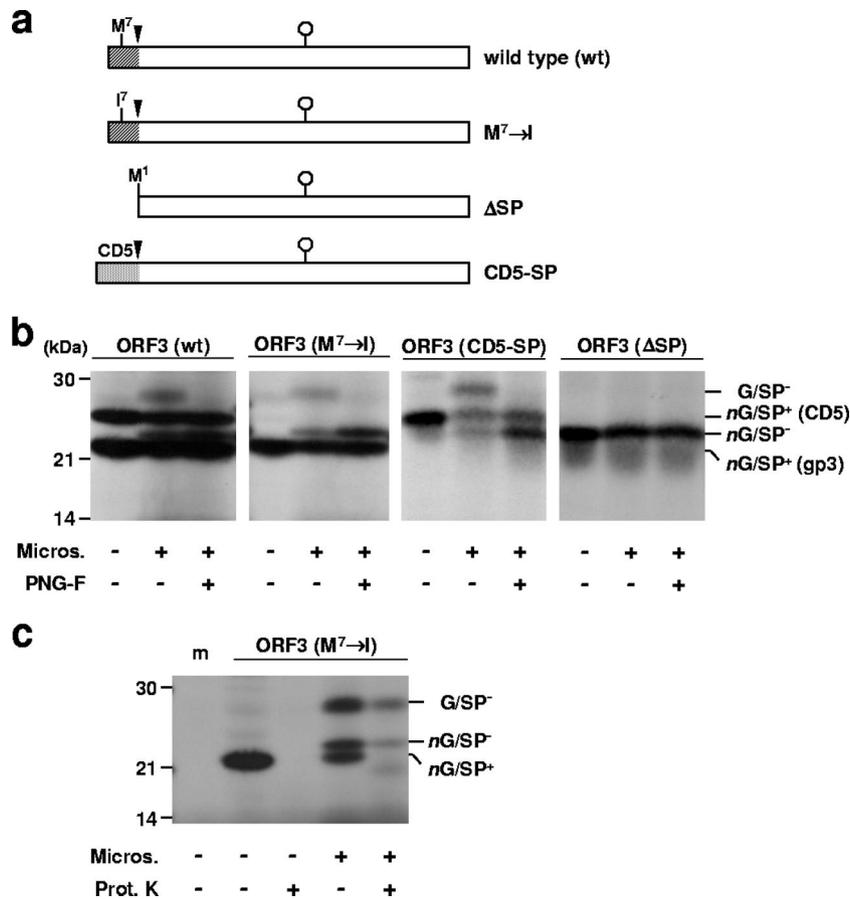


FIG. 2. ORF3 encodes a glycoprotein with a cleavable N-terminal signal sequence. (a) Linear representation of the wild-type ORF3 protein and derivatives expressed from pTUG-31-based expression plasmids. The signal sequences of the ORF3 product and of CD5 are indicated by shading. Signalase cleavage sites (black arrowheads) and potential N-glycosylation sites (white lollipop) are indicated. (b) In vitro translation of wild-type (wt) ORF3 and derivatives. Translations were performed either in the absence (-) or presence (+) of dog pancreas microsomes (Micros.). Prior to SDS-polyacrylamide gel electrophoresis (PAGE) analysis, translation products were treated with PNGase F (PNG-F) (+) or left untreated (-). The positions and masses (in kilodaltons) of proteins from the molecular size markers are shown at the left. The positions of the various products are shown at the right as follows: G/SP⁻, glycosylated product without signal peptide; nG/SP⁺ (CD5), nonglycosylated product with CD5 signal peptide; nG/SP⁻, nonglycosylated product without signal peptide; nG/SP⁺ (gp3), nonglycosylated product with gp3 signal peptide. (c) Proteinase K protection assay. ORF3 derivative M⁷→I was translated either in the absence (-) or presence (+) of microsomes (Micros.) and treated with proteinase K (Prot. K) (+) or mock treated (-) prior to SDS-PAGE analysis. The positions of the various products are shown on the right as described above for panel b; the positions of molecular size standards and their masses (in kilodaltons) are shown on the left. A sample from a translation reaction supplemented with water instead of expression plasmid was included as a negative control (m).

cloned downstream of the T7 RNA polymerase promoter (Fig. 2a). Translation of the wild-type gene in the absence of microsomes yielded not one but two products with molecular weights of 22,000 (22K) and 25K [Fig. 2b, panel ORF3 (wt), leftmost lane]. Scrutiny of the Elmo/02 ORF3 sequence revealed an AUG at codon position seven (Fig. 2a). Arguing that internal initiation of translation at this site might give rise to an additional, smaller product, we replaced Met⁷ by Ile (note that in many naturally occurring CCoV-I strains, Ile is found at this position [Fig. 1b]). Upon expression of this mutant in the absence of microsomes, only a single protein species was found. Surprisingly, however, it was not the 25K product but the faster migrating 22K product [Fig. 2b, panel ORF3 (M⁷→I), leftmost lane]. Apparently, the intact signal peptide of the ORF3 protein causes aberrant migration in sodium dodecyl sulfate (SDS)-polyacrylamide gels. Indeed, expression of a mutant with the ORF3 signal peptide replaced by that of

CD5 yielded a single protein species of 25K [Fig. 2b, panel ORF3 (CD5-SP), leftmost lane], in accordance with its calculated molecular mass (25.7 kDa).

Translation of wild-type ORF3 and derivatives in the presence of microsomes consistently yielded two additional products of 28K and 23K [Fig. 2b, panels ORF3 (wt), ORF3 (M⁷→I), and ORF3 (CD5-SP), middle lanes) that were fully protected from digestion by proteinase K (Fig. 2c) and hence appeared to be contained within the microsomal lumen. The 28K product is N glycosylated; treatment of translation products with endoglycosidase PNGase (Promega) resulted in loss of this protein species with a concomitant increase in the amount of the 23K product; the latter comigrated with mutant ΔSP that lacks a signal peptide [Fig. 2b, panel ORF3 (ΔSP)]. The combined findings conclusively show that CCoV-I ORF3 codes for a 28K glycoprotein (gp3) with a cleavable N-terminal signal sequence; hydrophobicity

plots did not reveal additional transmembrane regions (data not shown), indicating that gp3 is either a secretory or peripheral membrane protein.

ORF3 remnants in CCoV-II and in CCoV-II-related viruses.

Comparative sequence analyses suggest that the horizontal gene transfer that resulted in the CCoV-I/II split-up was restricted to the coding sequences for the signal peptide and ectodomain of S and should have left ORF3 intact (not shown). Under the assumption that CCoV-I represents the parental biotype and CCoV-II its recombinant offspring, the latter must have lost ORF3 subsequently. Indeed, close inspection of the intergenic regions (IGRs) separating the S and ORF3a genes in CCoV-II variants (4, 7, 24, 32, 40), in FCoV-II strains 79-1146 (NCBI accession number AY994055) and 79-1683 (NCBI accession number Y13921) (the S gene and downstream sequences are of CCoV-II origin in this strain; 21), and in TGEV (57) revealed remnants of ORF3 and/or its preceding IGR (Fig. 1c and d). In FCoV-I strains C1Je and Black (14, 46), however, there is no trace of ORF3, and the S gene and ORF3a are separated by a very short 11-nt IGR that except for the TRS bears little similarity to the IGRs preceding ORF3 or ORF3a in CCoV. Presumably, ORF3 was acquired after type I FCoV and CCoV diverged, although the possibility that ORF3 was already present in their common ancestor and then lost completely in the FCoV lineage cannot be excluded (Fig. 3). In any case, our findings suggest that while gp3 is important during CCoV-I infection, it became obsolete in CCoV-II.

Is gp3 advantageous only in combination with CCoV-I S protein? The function of gp3 is not known, but based upon its biochemical properties, it may act either in the infected cell within the compartments of the exocytotic route or in the extracellular milieu. Given that apart from ORF3 the main difference between CCoV-I and CCoV-II strains lies in the type of S proteins they carry, it would seem that the function of gp3 is in some way connected to the function of S, i.e., gp3 apparently provides an advantage only in combination with a type I spike protein. Conceivably, gp3 may be involved in the biogenesis of CCoV-I S or required for S-mediated attachment or fusion during entry. There is the alternative possibility, however, that gp3 is advantageous to the virus only in certain types of host cells or tissues correlating with the cell tropism conferred by S. Accumulating evidence suggests that FCoV-I and -II, and hence by extension CCoV-I and -II, recognize different receptors, which may well translate to a difference in host cell preference (2, 15, 23).

ORF3 would not be the sole example of an accessory gene lost after a tropism change. ORF3c is conserved among group 1 CoVs, yet it is inactivated in FCoV variants that cause feline infectious peritonitis; loss of expression seemingly correlates with a shift from enteric to systemic infection and in host cell tropism from enterocytes to monocytes (49). In severe acute respiratory syndrome CoV, loss of nsP8 may have been the consequence of cross-species transmission and adaptation to the human host (19, 35). In the case of TGEV, adaptation of CCoV-II to swine apparently was accompanied by inactivation of ORF3b and loss of ORF7b. Clearly, further studies of CoV accessory proteins are warranted, as these studies will not only broaden our under-

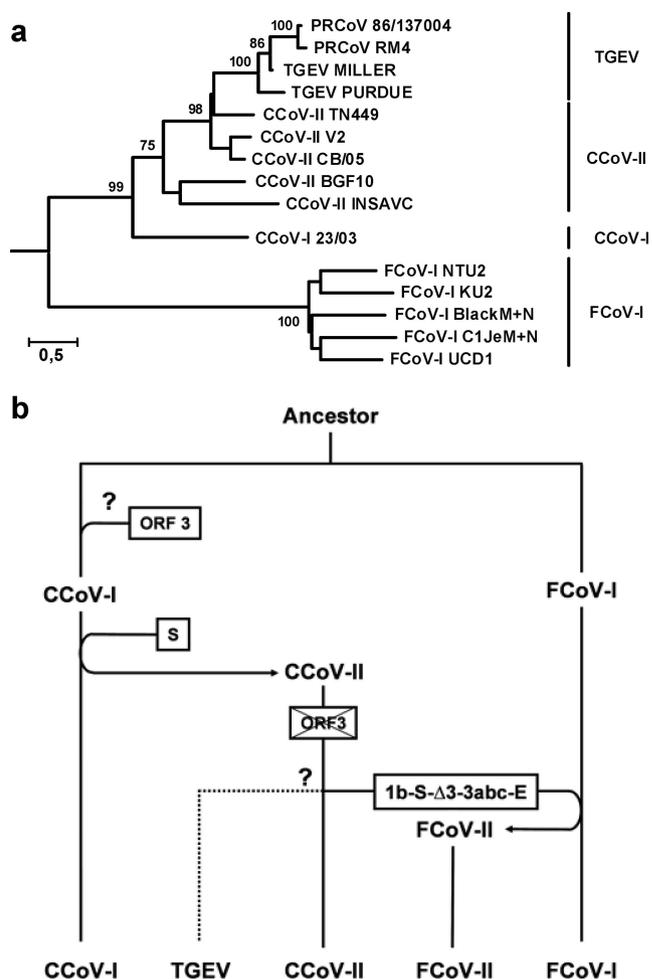


FIG. 3. Hypothetical scenario for the evolution of CoV cluster 1a. (a) Rooted neighbor-joining tree inferred from multiple amino acid sequence alignments of the M and N proteins, illustrating the evolutionary relationships between members of phylogroup 1a. Human CoV 229e served as an outgroup. Support from bifurcations from 100 bootstraps is indicated. PRCoV, porcine respiratory CoV. (b) CCoV-I and FCoV-I apparently arose from a common ancestor by linear descent. As these viruses diverged, several distinct RNA recombination events led to the emergence of CCoV-II, FCoV-II, and TGEV. Details are explained in the text. Question marks indicate steps that are not yet completely understood.

standing of coronavirus host adaptation and speciation but may also open new avenues to antiviral intervention.

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