Species-specific variation in RELA underlies differences in NF-B activity

Citation for published version:

Digital Object Identifier (DOI):
10.1128/jvi.00331-11

Link:
Link to publication record in Edinburgh Research Explorer

Document Version:
Publisher's PDF, also known as Version of record

Published in:
Journal of Virology

General rights
Copyright for the publications made accessible via the Edinburgh Research Explorer is retained by the author(s) and / or other copyright owners and it is a condition of accessing these publications that users recognise and abide by the legal requirements associated with these rights.

Take down policy
The University of Edinburgh has made every reasonable effort to ensure that Edinburgh Research Explorer content complies with UK legislation. If you believe that the public display of this file breaches copyright please contact openaccess@ed.ac.uk providing details, and we will remove access to the work immediately and investigate your claim.
Species-Specific Variation in RELA Underlies Differences in NF-κB Activity: a Potential Role in African Swine Fever Pathogenesis

Christopher J. Palgrave,1,2,3† Linzi Gilmour,2,3‡ C. Stewart Lowden,3 Simon G. Lillico,2
Martha A. Mellencamp,4§ and C. Bruce A. Whitelaw2∗

Veterinary Pathology Unit, Division of Veterinary Clinical Sciences, Royal (Dick) School of Veterinary Studies, University of Edinburgh, Easter Bush Veterinary Centre, Roslin, Midlothian EH25 9RG, United Kingdom1; Division of Developmental Biology, The Roslin Institute and Royal (Dick) School of Veterinary Studies, University of Edinburgh, Easter Bush Campus, Roslin, Midlothian EH25 9RG, United Kingdom2; Veterinary Health Research Pty Ltd., Trevenna Rd., West Armidale, NSW 2350, Australia3; and Genus/PIC, 100 Bluegrass Commons Blvd., Hendersonville, Tennessee 370754

Received 17 February 2011/Accepted 15 March 2011

African swine fever virus (ASFV) is a highly infectious disease of domestic pigs, with virulent isolates causing a rapidly fatal hemorrhagic fever. In contrast, the porcine species endogenous to Africa tolerate infection. The ability of the virus to persist in one host while killing another genetically related host implies that disease severity may be, in part, modulated by host genetic variation. To complement transcription profiling approaches to identify the underlying genetic variation in the host response to ASFV, we have taken a candidate gene approach based on known signaling pathways that interact with the virus-encoded immunomodulatory protein A238L. We report the sequencing of these genes from different pig species and the identification and initial in vitro characterization of polymorphic variation in RELA (p65; v-rel reticuloendotheliosis viral oncogene homolog A), the major component of the NF-κB transcription factor. Warthog RELA and domestic pig RELA differ at three amino acids. Transient cell transfection assays indicate that this variation is reflected in reduced NF-κB activity in vitro for warthog RELA but not for domestic pig RELA. Induction assays indicate that warthog RELA and domestic pig RELA are elevated essentially to the same extent. Finally, mutational studies indicate that the S531P site conveys the majority of the functional variation between warthog RELA and domestic pig RELA. We propose that the variation in RELA identified between the warthog and domestic pig has the potential to underlie the difference between tolerance and rapid death upon ASFV infection.

African swine fever virus (ASFV) is a pathogen of the Suidae (domestic and wild pig species), which may be transmitted directly or via an arthropod vector in the form of Ornthodoros ticks (35). ASFV is highly infectious, with virulent isolates causing an acute, rapidly fatal hemorrhagic fever in domestic pigs (Sus scrofa) (10, 34). This is thought, in part, to be the result of a proinflammatory cytokine storm driven by infected macrophages (9, 15, 16, 42–44, 56). Initiation of a systemic inflammatory response results in severe hematological and vascular perturbations, ultimately leading to cardiovascular collapse in a manner not dissimilar to septic shock (3, 17, 18, 20, 21, 40, 52, 53). In addition to hemorrhage, severe widespread apoptosis of infected macrophages and uninfected lymphocytes is a prominent feature of this disease; this is also likely related to markedly elevated proinflammatory cytokine levels (19, 33, 37, 43). In comparison to the severe disease which occurs in domestic pigs, in its natural hosts, warthogs (Phacochoerus sp.) and bushpigs (Potamochoerus sp.), ASF is subclinical and persistent (2, 32, 50). ASFV is notifiable to the World Organization for Animal Health (OIE), placing it in the highest category of infectious animal pathogens. It exhibits remarkable potential for transboundary spread, and outbreaks in domestic pig populations have a serious socioeconomic impact worldwide. Furthermore, ASF is considered to be the major limiting factor to pig production in Africa (34). ASFV is a large, double-stranded DNA virus and the only member of the Asfarviridae family (12), suggesting that it may carry novel genes that are not carried by other virus families. Furthermore, the ability of the virus to persist in one host while killing another genetically related host alludes to the possibility that disease severity may, in part, be modulated by host genetic variation.

Several candidate ASFV-encoded immune modulatory factors have been identified, including homologues of CD2 (8-DR/CD2v) (5, 6, 41), IAP (A224L) (31, 39), Bcl-2 (A179L; 5-EL) (1, 7, 8, 30), and IκBα (A238L; 5-EL) (36, 49). Of these, A238L shares 40% sequence homology and 20% identify with domestic pig IκBα (NFκBIA) and substitutes for NFκBIA by binding to the RELA (p65; v-rel reticuloendotheliosis viral oncogene homolog A) subunit of NF-κB. Thus, A238L reduces the ability of NF-κB to be activated (36, 49). In addition to inhibiting host NF-κB, A238L also suppresses calcineurin phosphatase activation of NFAT signaling by the following two mechanisms: direct binding to calcineurin phosphatase 3, β isoform (PPP3CB), and binding to the immunophilin carrier cyclophilin A (PPIA) in a manner similar to that of the immunosuppressive drug cyclosporine A (28, 29).

Various groups have initiated transcription profiling of host genes implicated in ASFV infection (15, 16, 42–44, 56). These
studies identified numerous upregulated host genes, but to date, all are limited to analysis in domestic pig cells. In this study, we take a complementary approach to this question by the testing of variation in targeted candidate genes. Clearly A238L represents a novel and versatile immunoregulatory mechanism by which ASFV can inhibit both the NF-κB and NFAT signaling pathways (11, 28, 29, 36, 49). We therefore consider the three A238L target proteins, RELA, PPP3CB, and PPIA, and the two proteins it mimics, NFKBIA and NFATC1, as candidates for the genetic variation between pig species which may contribute to species-specific responses to ASFV infection. We now report the sequencing of these genes from different pig species and identification and initial in vitro characterization of polymorphic variation in one of them.

MATERIALS AND METHODS

mRNA isolation, cDNA synthesis, and DNA sequencing. Whole blood (5 ml) was collected into EDTA from a domestic pig (Sus scrofa, commercial pig; United Kingdom), common warthog (Phacochoerus africanus; Rotterdam Zoo, Holland), and babirusa (Babyrousa babyrussa; Marwell Zoo, United Kingdom) and transferred immediately into DNA/RNA stabilization reagent for blood/ bone marrow (Roche Diagnostics). This was processed using a mRNA isolation kit for white blood cells (Roche Diagnostics). Initially, cDNA libraries were synthesized using the SMART RACE (rapid amplification of cDNA ends) kit for white blood cells (Roche Diagnostics). Initially, cDNA libraries were synthesized using the SMART RACE (rapid amplification of cDNA ends) kit for white blood cells (Roche Diagnostics). cDNA amplification kit (Clontech) to enable partial sequencing and design of species-specific primers (Table 1) for synthesis of individual cDNAs using proof-reading PCR using a 1-μl sample in a 25-μl PCR mixture consisting of 20 pmol of each primer in 2 μl MgCl2 and 2 μl deoxynucleotide triphosphates (dNTPs) with 0.7 U High-Fidelity DNA polymerase (Roche Diagnostics). The PCR cycling conditions were optimized for each gene (data not shown). PCR products were resolved on a 1% agarose gel, excised with a scalpel blade, and transferred immediately into DNA/RNA stabilization reagent for blood/ bone marrow (Roche Diagnostics). This was processed using a mRNA isolation kit for white blood cells (Roche Diagnostics). Initially, cDNA libraries were synthesized using the SMART RACE (rapid amplification of cDNA ends) kit for white blood cells (Roche Diagnostics). cDNA amplification kit (Clontech) to enable partial sequencing and design of species-specific primers (Table 1) for synthesis of individual cDNAs using proof-reading PCR using a 1-μl sample in a 25-μl PCR mixture consisting of 20 pmol of each primer in 2 μl MgCl2 and 2 μl deoxynucleotide triphosphates (dNTPs) with 0.7 U High-Fidelity DNA polymerase (Roche Diagnostics). The PCR cycling conditions were optimized for each gene (data not shown).

Genomic DNA isolation and sequencing. Samples of skeletal muscle were collected into 20% dimethyl sulfoxide (DMSO)-saturated salt (NaCl) solution and stored at −70°C. DNA was extracted from 0.2 g muscle using the BACC2 extraction kit for blood and cell cultures (Nucleon Biosciences). For PCR, 50 ng genomic DNA was used as template in a 25-μl PCR mixture consisting of 20 pmol of each primer in 2 μM MgCl2 and 2 μM dNTPs, with 0.7 U High-Fidelity DNA polymerase (Roche Diagnostics). Plasmid construction. Restriction sites were introduced into RELA products from the above-described sequencing study by nested PCR; this enabled insertion of domestic pig and warthog RELA genotypes into the multiple cloning site of the pFLAG-CMV-4 vector (Sigma-Aldrich). Plasmid products were digested to 1:500 in sterile water, and 1 μl was used as a template in a 25-μl PCR mixture consisting of 20 pmol of each primer (forward HindIII [5’-CCA AGC TGG ACC TCT TCC CCC TCA TCT T-3'] and reverse NotI [5’-GGG CCG CCG CTT AGG AGC TGA TCA GAT GAC TCT TC-3’]) in 2 μM MgCl2 and 2 μM dNTPs, with 0.7 U High-Fidelity DNA polymerase (Roche Diagnostics). Restriction sites were underlined. Each ~1.6-kbp PCR product was resolved on a 1% agarose gel, excised with a scalpel blade, and transferred using the QIAQuick gel extraction kit (Qiagen). Following HindIII and NotI restriction digestion of the vector and RELA PCR products, the products were ligated into the open plasmid. These constructs allowed constitutive expression of warthog RELA and domestic pig RELA driven by the cytomegalovirus (CMV) promoter. In addition, an N-terminal eight-amino-acid FLAG sequence is incorporated into the protein, which is recognized by an anti-FLAG monoclonal antibody. Site-directed mutagenesis was preformed using a QuikChange II site-directed mutagenesis kit (Stratagene).

Plasmid transient transfection was performed using Lipofectamine2000 (Invitrogen) in ~1 × 105 cells/well of a 12-well plate in culture medium at 37°C and 5% CO2. COS-7 cells were cultured in Glasgow minimal essential medium (GMEM, Sigma), containing 10% fetal bovine serum (Invitrogen), 1% l-glutamine (Invitrogen), 1% sodium pyruvate (Invitrogen), 1% nonessential amino acids (Invitrogen), and 0.2% β-mercaptoethanol (Invitrogen). Mouse embryonic fibroblasts (MEFs) and RELA+ MEFs were cultured in Dulbecco’s modified Eagle medium (DMEM, Sigma), containing 10% fetal bovine serum (Invitrogen). Luciferase activity was determined in triplicate using the dual reporter assay system (Promega) and analyzed with Excel (Microsoft), with total protein measured with the bicinchoninic acid (BCA) protein assay kit ( Pierce).

The data presented are the means ± standard deviations; statistical significance was evaluated using the unpaired Student t test, with a difference between groups being considered statistically significant if the P value of the comparison was <0.05.

Western blotting. For Western blot analysis, denatured protein (10 μg) was run in Tris-glycine-SDS running buffer (National Diagnostics) on prestained NuPage 12% Tris-glycine gels (Invitrogen) before transfer to a nitrocellulose membrane (National Diagnostics). Proteins were visualized using anti-FLAG M2-peroxidase (horseradish peroxidase [HRP]) (Sigma)-conjugated primary antibody and mouse β-actin primary antibody with goat anti-mouse IgG HRP (Sigma)-conjugated secondary antibody and detected with Immobilon Western (Millipore) chemiluminescent HRP substrate.

RESULTS

Limited sequence variation in candidate porcine genes. cDNAs were produced for PPIA, NFKBIA, NFATC1 (regulatory domain), PPP3CB, and RELA. These were obtained from the following three pig species: the domestic pig (Sus scrofa), common warthog (Phacochoerus africanus), and babirusa (Babyrousa babyrussa). The domestic pig and warthog represent ASFV-susceptible and ASFV-tolerant species, respectively. The “outspecies” is the babirusa, which is considered to be the most ancient extant species of pig; its range is restricted to the island of Sulawesi in the Indonesian archipelago. It has no common ancestor with the domestic pig more recently than approximately 10 to 19 million years ago (38). It is not known how this pig species would respond regarding ASFV infection. We generated sequence data (deposited in the EMBL database) (Table 2) and aligned all sequences to those of the human homologues which we used as reference sequences. Limited sequence variation was observed in this study (data not shown), with the following genes displaying complete homology at the translated protein level between the domestic pig and warthog: PPIA, NFATC1 regulatory domain, and

<table>
<thead>
<tr>
<th>Gene/domain</th>
<th>Primer sequence (5’–3’)</th>
</tr>
</thead>
<tbody>
<tr>
<td>PPIA (cDNA)</td>
<td>ATC TNT CAG TGC TGC TCA GC</td>
</tr>
<tr>
<td>NFKBIA (cDNA)</td>
<td>AAG GAG CGG CTA CTT GAC G</td>
</tr>
<tr>
<td>NFKBIA (5’-end DNA)</td>
<td>CTC ATC GCA GGG AGG TTC TCT GC</td>
</tr>
<tr>
<td>NEATC1 (cDNA)</td>
<td>ATC TCA GCT GTT GGG TCA GC</td>
</tr>
<tr>
<td>PPP3CB (cDNA)</td>
<td>CCC ACA ACA TCG TTT GAC AT</td>
</tr>
<tr>
<td>RELA (cDNA)</td>
<td>GAC CTC TCC CCC ATC ATC TT</td>
</tr>
<tr>
<td>RELA transactivation domains (DNA)</td>
<td>GGA AGG GAC ACTGAC AGA GG</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Gene/domain</th>
<th>Sense</th>
<th>Antisense</th>
</tr>
</thead>
<tbody>
<tr>
<td>PPIA (cDNA)</td>
<td>CAG AAG GAA TGG TCT GAT GG</td>
<td></td>
</tr>
<tr>
<td>NFKBIA (cDNA)</td>
<td>CAT GGT CTT TTA GAC ACT TCT CA</td>
<td></td>
</tr>
<tr>
<td>NFKBIA (5’-end DNA)</td>
<td>TCC TCG TCC TTC ATG GAG TC</td>
<td></td>
</tr>
<tr>
<td>NEATC1 (cDNA)</td>
<td>AGT GAG GGT GAG TGG TCC AG</td>
<td></td>
</tr>
<tr>
<td>PPP3CB (cDNA)</td>
<td>ATG TGA GAG TCT CGG AGA AG</td>
<td></td>
</tr>
<tr>
<td>RELA (cDNA)</td>
<td>CCC CTG AGG AGC TGA TCT GA</td>
<td></td>
</tr>
<tr>
<td>RELA transactivation domains (DNA)</td>
<td>TCA GAA GGG CTG AGA AGT CC</td>
<td></td>
</tr>
</tbody>
</table>
NFkBIA. In the warthog, PPP3CB contained two insertions with respect to the domestic pig. However, these were located outwith the known functional domains. Furthermore, these correlate with splice variants described in human PPP3CB (22, 27). Only in the RELA subunit of NF-κB were potentially significant coding differences between the domestic pig and warthog identified (Fig. 1).

**Sequence of the porcine RELA gene.** The RELA open reading frame (ORF) is 1,662 nucleotides in the domestic pig and warthog and is slightly smaller at 1,556 nucleotides in the babirusa. These encode proteins which are 554 and 552 amino acids in length, respectively. Human RELA ORF and protein are the same lengths as the babirusa sequences. Due to insufficient primer binding sites in the 5′-untranslated region (UTR), we sequenced all but the first 6 nucleotides for all three pig species (Fig. 1).

Although the babirusa and human sequences are both 6 nucleotides shorter than the other pig sequences, the nucleotide deletions occur at different locations. The babirusa sequence has a single 6-nucleotide deletion, in relation to those of the other pig species, located between the Rel homology domain and the transactivation 2 domain. Human RELA has two 3-nucleotide deletions, one 20 nucleotides upstream from the babirusa deletion and the other in the transactivation 2 domain.

Of the 13 nucleotide differences between the domestic pig and warthog, only three are nonsynonymous and result in codon changes, T448A, S485P, and S531P. The threonine at 448 in the domestic pig occurs as an alanine in the warthog and babirusa and is absent in human RELA. The serine at 485 in the domestic pig sequence acts as an alanine in the warthog and babirusa and is absent in human RELA. The serine at 531 in the domestic pig is also found in babirusa and human but is a proline in warthog RELA. All three amino acid differences occur outside the Rel homology domain, with amino acids 448 and 485 located within transactivation domain 2 and amino acid 531 within transactivation domain 1. To confirm these sequence differences, a 268-nucleotide region was amplified directly from genomic DNA and sequenced for an additional 5
domestic pigs and 11 warthogs (domestic pig sequences, EMBL accession numbers FN424224 to FN424228; warthog sequences, EMBL accession numbers FN424229 to FN424239). All domestic pig sequences were identical to each other over this region, and similarly, all warthog sequences were identical to each other over this region.

**Predicted structural changes as a consequence of RELA sequence variation.** Given that even single amino acid changes can have significant effects on protein structure, we performed *in silico* analyses on the identified changes in RELA in an attempt to determine the structural consequences of the amino acid substitutions. At the three sites, the greater hydrophobicity was conferred by the threonine at 448 in the domestic pig sequence and the prolines at 385 and 531 in the warthog sequence. No difference in structural disorder was evident for domestic pig RELA and warthog pig RELA. To determine if the higher activity level of domestic pig RELA may not be compatible with survival in these cells; a similar scenario may have occurred during a study investigating phosphorylation at the equivalent site (23).

**Basal activity of porcine RELA variants.** To determine whether the identified sequence variation between domestic pig RELA and warthog RELA affects NF-κB activity, we established a cell transfection assay. Cells were transiently transfected in duplicate using a triple plasmid cotransfection strategy involving the following: (i) 1 μg expression vector for FLAG-tagged domestic pig or warthog RELA or an empty vector control (pFLAG-CMV4; Sigma Aldrich); (ii) 1 μg NF-κB reporter plasmid comprising 4 copies of the NF-κB consensus binding sequence, driving expression of firefly luciferase (pNFκB-Luc; BD Biosciences, Clontech); and (iii) 1 μg transfection control vector expressing Renilla luciferase driven off the herpes simplex virus (HSV) thymidine kinase (TK) promoter (pRL-TK; Promega). Cells were harvested at 24 h posttransfection, and a dual luciferase assay (firefly luciferase activity relative to Renilla luciferase activity) was performed on COS-7 cells after 24 h of transient transfection with 1 μg warthog (WH) or 1 μg domestic pig (DP) RELA. Activity is presented as the fold difference between RELA-induced NF-κB—luciferase activity normalized to cotransfected TK-Renilla luciferase activity relative to that of the empty vector. Error bars = standard deviations from the means. P < 0.05 between WH and DP. (C) Fold difference between NF-κB—luciferase activities in RELA−/− MEFs after 24 h of transient transfection with warthog (WH) and domestic pig (DP) RELA genes. Activity is presented as the fold difference between RELA-induced NF-κB—luciferase activity normalized to cotransfected TK-Renilla luciferase activity relative to that of the empty vector normalized to cotransfected TK-Renilla luciferase activity. Error bars = standard deviations from the means. P < 0.05 between WH and DP.

**Effect of RELA sequence variation on induced NF-κB activity.** To determine if the ability to induce NF-κB activity was altered by the different pig RELA proteins, we stimulated transiently transfected RELA−/− MEFs with known inducers of NF-κB signaling (9, 14, 15, 16, 42–44, 56). Induction with tumor necrosis factor alpha (TNF-α) nearly doubled (90% increase) warthog RELA activity, whereas domestic pig RELA was always greater than that of domestic pig RELA. It is possible that the higher activity level of domestic pig RELA may not be compatible with survival in these cells; a similar scenario may have occurred during a study investigating phosphorylation at the equivalent site (SS29) in human RELA (54).

**Identification of functional mutations in porcine RELA.** In our sequence data, we identified three amino acid differences between domestic pig RELA and warthog pig RELA. To determine if the *in vitro* difference in domestic pig and warthog
RELAs activities was a cumulative effect of these three differences, or due to one or other individual mutations, we generated versions of RELA carrying the following single-base changes: T448A, S485P, and S531P. Transfection of these RELA variants into RELA−/− MEFs demonstrated that the majority of the reduced basal activity observed for warthog RELA compared to that observed for domestic pig RELA was attributed to the S531P mutation (Fig. 4).

**DISCUSSION**

In this study, we have taken a candidate approach, identifying genes which may affect the severity of the host response to ASFV infection in the highly susceptible domestic pig and ASFV-tolerant warthog. The ASFV immunomodulatory factor A238L is known to interact with components of both NF-κB and NFAT host signaling pathways (11, 28, 29, 36, 49). We have sequenced five key factors in these pathways, including three A238L-targeted proteins, RELA, PPP3CB, and PPIA, and the two proteins it mimics, NFKBIA and NFATC1 (regulatory domain). Modest sequence differences have been identified at the cDNA (mRNA) level between the domestic pig and warthog; however, the majority of these are synonymous (silent) and do not alter the resulting amino acid sequences. We also sequenced the same genes from the ancient babirusa and observed a small number of amino acid differences between this species and the other two species of pig. These differences likely reflect the long evolutionary distance that exists between these species (38).

Despite the high degree of conservation observed between warthog and domestic pig PPP3CB, PPIA, NFKBIA, and NFATC1-translated protein sequences, significant variation was detected in RELA. RELA is the predominant member of the heterodimeric transcription factor NF-κB (47). Moreover, the sequence variation includes a phosphorylation site in transcription activation domain 1 (position 531), which is highly conserved across mammals and has been demonstrated to modulate the activity of human RELA (equivalent to SS529) (54, 55). The function of approximately one-third of all eukaryote proteins is controlled by phosphorylation; thus, the observed SS531P sequence variation represents an intriguing candidate regulator for the reduced pathology observed in African pigs infected with ASFV. We demonstrate that the genetic variation in the RELA sequence between the domestic pig and warthog is reflected in NF-κB activity in vitro, with warthog RELA displaying significantly reduced basal and induced NF-κB activity. We discuss three (not mutually exclusive) scenarios of how the genetic variation we have identified between the domestic pig and warthog may underlie the dramatic phenotypic difference in how these two pig species respond to ASFV.

First, as suggested by our in vitro assays, warthog RELA is inherently less active than the domestic pig RELA, exhibiting lower basal and induced levels. In a study of macrophage transcription profiles following ASFV infection in vitro, several factors within the NF-κB signaling pathway displayed elevated expression (i.e., NFKB1, NFKBIA); however, RELA expression was not observed to be altered (56). The impact of ASFV infection on the warthog RELA expression level is not known, since this study used only domestic pig macrophages. Therefore, although we have not determined whether the porcine RELA variants are differentially phosphorylated, it would appear that variation in domestic pig RELA activity may not be determined by altered expression levels. Alternatively, the warthog will have to function with reduced basal NF-κB activity, which would presumably indicate an adapted NF-κB-dependent transcriptome between the two species.

In the second scenario, phosphorylation of domestic pig RELA at S531 activates a set(s) of genes which are not activated by warthog RELA, as it lacks this phosphorylation site. Much is known about activation of the NF-κB pathway, with data coming primarily from studies of human RELA (47). NF-κB activity is regulated by two different mechanisms. The classical canonical pathway involves inhibitors (e.g., NFKBIA) that sequester this transcription factor in the cytoplasm until they are proteolytically degraded by the ubiquitin pathway (24). The alternative pathway revolves around posttranscriptional modifications, predominantly phosphorylation of RELA (46). At least 8 inducible phosphorylation sites have been identified in human RELA, which enable transcription of subsets of NF-κB-dependent genes (26, 45). One such site is S529, which is equivalent to domestic pig S531, suggesting that this site may perform a similar role. Indeed, differential expression of subsets of immune and inflammatory proteins as a result of S531

**FIG. 3.** Induction of RELA allelic variants. RELA−/− MEFs were transiently cotransfected with 1 μg warthog (clear box) or 1 μg domestic pig (filled box) RELA. Cells were treated with TNF-α (30 ng/ml), lipopolysaccharide (LPS; 10 μg/ml), phorbol-12-myristate 13-acetate (PMA; 20 mM), and hydrogen peroxide (10 μM) immediately after and for the duration of the transfection. Cells were harvested at 24 h. RELA-induced NF-κB–luciferase (1 μg) activity was normalized to cotransfected TK-Renilla luciferase (1 μg) activity and presented as the fold induction above nondrug-treated cells (relative value of 1 depicted by the line).

**FIG. 4.** Comparison of individual allelic RELA variation on NF-κB activity. Comparison of NF-κB–luciferase activity in RELA−/− MEFs after 24 h transient transfection with 1 μg warthog (WH) or 1 μg domestic pig (DP) RELA or 1 μg of RELA variants encoding the individual amino acid substitutions T448A, S485P, and S531P. Activity is presented as the fold difference between RELA-induced NF-κB–luciferase activity normalized to cotransfected TK-Renilla luciferase activity relative to that of the empty vector normalized to cotransfected TK-Renilla luciferase activity. Error bars = standard deviations from the means, *, P < 0.08 versus DP.
phosphorylation could play a role in determining ASFV pathogenesis in the domestic pig. Furthermore, phosphorylation not only controls specific transcription profiles but also can underlie the developmental timing in gene activation (13). It is tempting to speculate that if a similar mechanism is applied through the domestic pig S531 site, then it could also play a role in the gross physical differences that characterize the various pig species.

In the third scenario, we consider whether the S531P variation in RELA results in different outcomes of interaction with A238L during ASFV infection. Phosphorylation of human S529 (equivalent to domestic pig S531) is inhibited by the interaction of RELA with NFKBIA; only upon activation and degradation of NFKBIA can S529 phosphorylation occur (55). During ASFV infection, NFKBIA is degraded and replaced by A238L, which mimics NFKBIA but is not susceptible to proteolytic degradation (36, 49). As a result, in domestic pig cells infected with ASFV, S531 may not be exposed for phosphorylation. In comparison, warthogs lack this phosphorylation site; therefore, the ability of A238L to block phosphorylation at this site is irrelevant. The role of NFKBIA is to prevent nuclear translocation of NF-κB (24). We did not observe any gross temporal differences in nuclear transportation rates upon stimulation between cells expressing domestic pig RELA and warthog RELA; likewise, phosphorylation of human S529 also does not affect nuclear translocation (54). Furthermore, studies using recombinant ASFV lacking A238L indicate that neither nuclear import nor export of RELA is affected by this immune modulator (48). This suggests that the immunomodulatory functions of A238L are not the result of preventing nuclear translocation in a manner similar to that used by NFKBIA. Instead, A238L may inhibit NF-κB-mediated transcription by other mechanisms, for example, by preventing phosphorylation of RELA in domestic pigs, as discussed above. Whether such differences in how domestic pig RELA and warthog RELA interact with A238L exist will require further investigation. Likewise, elucidating how different mechanisms governing RELA activity and NF-κB-mediated transcription have evolved in these species, and determining the full extent of their functional implications on the immune system and wider transcriptome, will require additional study.

ASFV is highly infectious, with virulent isolates causing an acute, rapidly fatal hemorrhagic fever in domestic pigs (10, 34). In contrast, the porcine species endogenous to Africa tolerate the virus. Outbreaks have significant economic repercussions in addition to welfare concerns. In Africa, ASFV limits the use of the genetically improved breeding pig stock that is pervasive in the Eurasian landscape. Furthermore, ASF poses a constant threat to Europe and Asia, as documented by the list of outbreaks that have occurred over the last 30 to 40 years. As our world climate changes and the international movement of pork products continues to rise, this risk may well increase. No effective vaccine has been developed, so many look to genetic strategies to mitigate the geographical limits of pig breeding imposed by ASFV. We have demonstrated that a polymorphic RELA variant found in warthogs has the potential to underlie the difference between tolerance and rapid death upon infection with ASFV.

ACKNOWLEDGMENTS

This work was supported by the BBSRC (United Kingdom) and a Genus/PIC Postgraduate Studentship, C.J.P. was also funded by a Postgraduate Centenary Fellowship (University of Edinburgh). L.G. was funded through a Faraday-BBSRC CASE studentship.

REFERENCES


