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Maturation of dengue virus nonstructural protein 4B in monocytes enhances production of dengue hemorrhagic fever-associated chemokines and cytokines

James F. Kelley ^a, Pakieli H. Kaufusi ^{a,b}, Esther M. Volper ^a, Vivek R. Nerurkar ^{a,b,*}^a Department of Tropical Medicine, Medical Microbiology and Pharmacology, John A. Burns School of Medicine, University of Hawaii at Manoa, Honolulu, HI, 96813, USA^b Pacific Center for Emerging Infectious Diseases Research, John A. Burns School of Medicine, University of Hawaii at Manoa, Honolulu, HI, 96813, USA

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ABSTRACT

High levels of viremia and chemokines and cytokines underlie the progression of severe dengue disease. Dengue virus (DENV) preferentially infects peripheral blood monocytes, which secrete elevated levels of immunomodulators in patients with severe disease. Further, DENV nonstructural proteins (NS) are capable of modifying intracellular signaling, including interferon inhibition. We demonstrate that peak secretions of immunomodulators such as IL-6, IL-8, IP-10, TNF α or IFN γ in DENV-infected monocytes correlate with maximum virus production and NS4B and NS5 are primarily responsible for the induction of immunomodulators. Furthermore, we demonstrate that sequential NS4AB processing initiated by the viral protease NS2B3 (pro) and via the intermediate 2KNS4B significantly enhances immunomodulator induction. While the 2K-signal peptide is not essential for immunomodulator induction, it plays a synergistic role with NS4B. These data suggest that NS4B maturation is important during innate immune signaling in DENV-infected monocytes. Given similar NS4B topologies and polyprotein processing across flaviviruses, NS4B may be an attractive target for developing *Flavivirus*-wide therapeutic interventions.

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Introduction

Dengue virus (DENV) causes considerable risk to human health worldwide infecting an estimated 50–100 million people annually and causing explosive outbreaks with infection rates as high as 80–90% among individuals previously unexposed to the virus (World Health Organization, 1997). The incidence of dengue diseases has dramatically increased during the past two decades as the result of an expanding geographical distribution of the *Aedes* mosquito vector and increased human travel (Gubler, 2002; Gubler, 2006; World Health Organization, 1997). Over 90% of DENV infections are asymptomatic or result in self-limiting dengue fever (DF) cases that resolve without complications (Guzman and Isturiz, 2010; Guzman and Kouri, 2002). However, a subset of infected patients progresses to severe dengue hemorrhagic fever (DHF) and dengue shock syndrome (DSS), resulting in over a half-million hospitalizations each year worldwide (Gubler, 1998; World Health Organization, 2009). No definitive mechanisms explain the progression of DHF/DSS, which can be defined in part as bleeding and increased plasma leakage into the pleural cavities and peripheral tissue without morphological damage to the capillary endothelium (Chang et al., 1990; Srikiatkachorn et al., 2007). Clinical studies indicate that

patients who progress to severe disease demonstrate elevated viremia (Libraty et al., 2002; Murgue et al., 2000; Vaughn et al., 2000) and high levels of interleukin (IL)-6, IL-8 and tumor necrosis factor alpha (TNF α) in the bloodstream (Hober et al., 1993; Huang et al., 2000; Juffrie et al., 2001; Nguyen et al., 2004; Priyadarshini et al., 2010; Raghupathy et al., 1998). Similarly, peripheral blood monocytes from patients with DHF/DSS display elevated DENV antigen and increased expression of activation markers and production of immunomodulators (Durbin et al., 2008; Halstead, 1989; Kou et al., 2008), implicating monocytes as important cells during infection and severe disease pathogenesis. Moreover, DENV-infected primary monocytes secrete DHF/DSS-associated immunomodulators (Bosch et al., 2002; Chen and Wang, 2002).

DENV belongs to the family Flaviviridae and consists of four genetically distinct serotypes having a positive-sense, single-stranded RNA genome of approximately 11 kilobases (kb) in length. The RNA encodes for a polyprotein precursor that is co- and post-translationally processed into three structural (C, PrM and E) and seven nonstructural proteins (NS1, NS2A, NS2B, NS3, NS4A, NS4B and NS5) (Bell et al., 1985; Chambers et al., 1990). The NS are responsible for various enzymatic activities during replication, including the NS5 RNA-dependent RNA polymerase (RdRP) and methyltransferase activity required for viral RNA capping (Chung et al., 2010; Dong et al., 2010; Geiss et al., 2009; Selisko et al., 2010; Zou et al., 2011), the NS3 helicase and the NS2B3 protease (NS2B3pro) (Cahour et al., 1992; Falgout et al.,

* Corresponding author at: John A. Burns School of Medicine, University of Hawaii at Manoa, 651 Ilalo Street, BSB 325AA, Honolulu, HI, 96813, USA. Fax: +1 808 692 1984.
E-mail address: nerurkar@hawaii.edu (V.R. Nerurkar).

1991; Padmanabhan et al., 2006). Several NS, such as NS3, NS4B and NS5, interact as part of the viral replication complex facilitating transcription and translation of the viral genome (Harris et al., 2006; Khromykh et al., 1999a; Khromykh et al., 1999b; Mackenzie et al., 2007; Roosendaal et al., 2006; Westaway et al., 1997a; Westaway et al., 1997b). Further, accumulating evidence suggests that intrinsic DENV genetic characteristics within NS4B and NS5 are associated with severe disease outcomes (Leitmeyer et al., 1999).

DENV NS5 induces IL-8 transcription and protein secretion in human embryonic kidney cells, (Medin et al., 2005) and inhibits the interferon alpha (IFN α) response through binding and degradation of STAT2 (Ashour et al., 2009; Mazzon et al., 2009). Also, NS4B strongly inhibits the IFN transduction cascade by interfering with STAT1 phosphorylation (Munoz-Jordan et al., 2003); and sequential processing of NS4AB by viral and host proteases is required to initiate an IFN antagonistic function (Munoz-Jordan et al., 2005). Nonstructural protein-induced subversion of the host IFN response and induction of immunomodulators may simultaneously promote DENV survival while increasing the risk of severe disease outcomes. Based on the aforementioned data, to further understand the role of NS in dengue immunopathogenesis, we hypothesized that NS5 and maturation of NS4B expressed in monocytes would induce DHF-associated immunomodulators. In this report, we demonstrate that both NS5 and NS4B induce immunomodulators and that NS4B maturation via cleavage of the NS4AB polypeptide, in a 2KNS4B-dependent manner, significantly enhanced immunomodulator production in monocytes.

Results

Elevated secretion of immunomodulators from DENV-infected THP-1 monocytes corresponds with peak viral titers and copy numbers

To establish whether DENV-infected monocytes secrete DHF-associated immunomodulators, we infected THP-1 cells with DENV-2 New Guinea (NGC) strain and collected cells and culture supernatants each day for five consecutive days. Plaque assay and quantitative real-time polymerase chain reaction (qRT-PCR) data demonstrated that peak viral titers and copy numbers for both MOI-0.1 and -1 occurred on day 3 after infection; however, infection with MOI-1 resulted in approximately log 1.4 higher titer and log 0.5 higher viral RNA transcripts than infection with an MOI-0.1 on day 3 after infection (Fig. 1A). We determined by qRT-PCR that the peak induction of IL-8 and TNF α transcripts occurred at day 3 after infection (Supplemental Fig. 1); therefore, we assayed the THP-1 supernatants for immunomodulators at days 1 and 3 after infection. The Luminex $\text{\textcircled{R}}$ multiplex data corresponded with transcript data in that the THP-1 cells infected with an MOI-1 induced peak secretion of IL-6, IL-8, interferon gamma-induced protein 10 (IP-10) and TNF α at day 3 after infection (Fig. 1B). Levels of IL-6, IL-8, IP-10, and TNF α peaked at 52, 2000, 40 and 65 pg/mL, respectively. Also slight overall increases were observed in VEGF and IFN γ levels at day 3 after infection. THP-1 cells incubated with lipopolysaccharide (LPS) significantly produced all of the immunomodulators tested both at the transcript and protein levels (Supplemental Fig. 1 and Fig. 1B) whereas THP-1 cells mock-infected with UV-treated virus produced low expression levels similar to those observed in untreated THP-1 cells (Fig. 1B). Overall, these results demonstrated that DENV-2

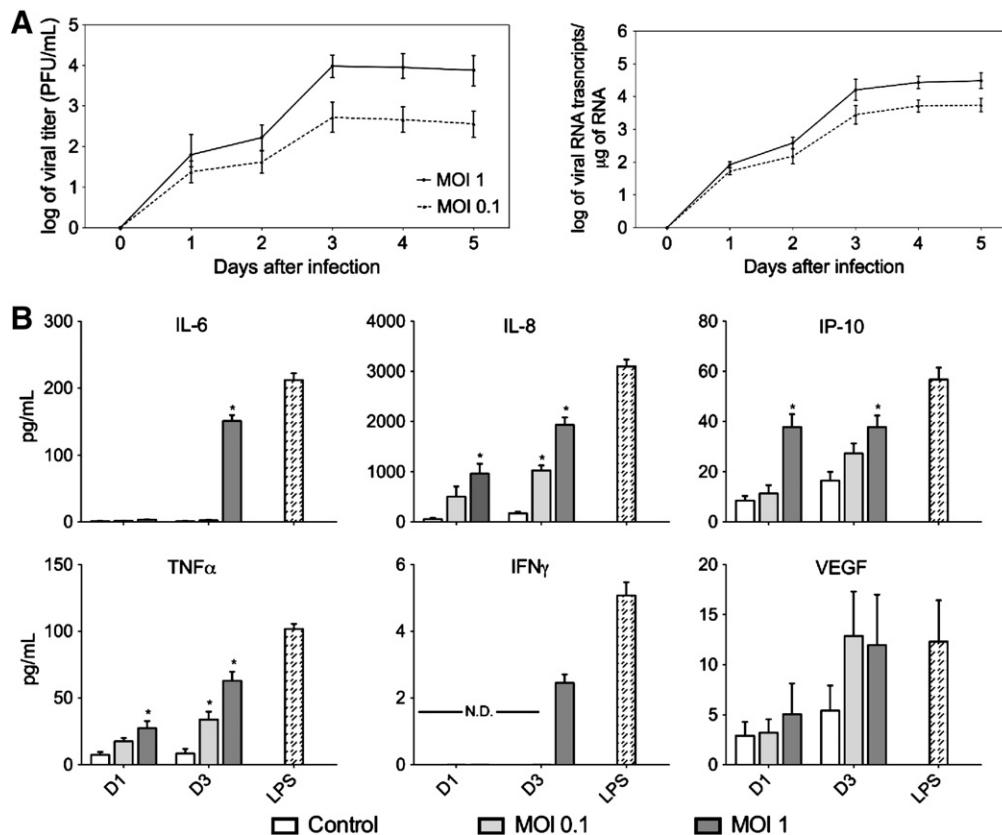


Fig. 1. Kinetics of chemokine and cytokine secretion in DENV-infected THP-1 cells. THP-1 cells were infected with DENV-2 NGC strain at MOI-0.1 and -1 and cells and supernatants were collected every 24 h after infection for five consecutive days. (A) Viral titers and RNA copy numbers were determined by plaque assay and qRT-PCR, respectively, on samples collected each day after infection (MOI-1, solid line; MOI-0.1, dotted line). (B) Chemokine and cytokine levels in mock- and DENV-infected THP-1 cell culture supernatants collected at days 1 and 3 after infection (mock-infected, white bars; DENV-infected MOI-0.1, light gray bars; DENV-infected MOI-1, dark gray bars; and LPS treatment, striped bars). Data points and bars represent the mean \pm SD of three independent experiments conducted in duplicate. (*) indicates statistical significance at $p < 0.05$ as compared to same day controls; N.D., not detected.

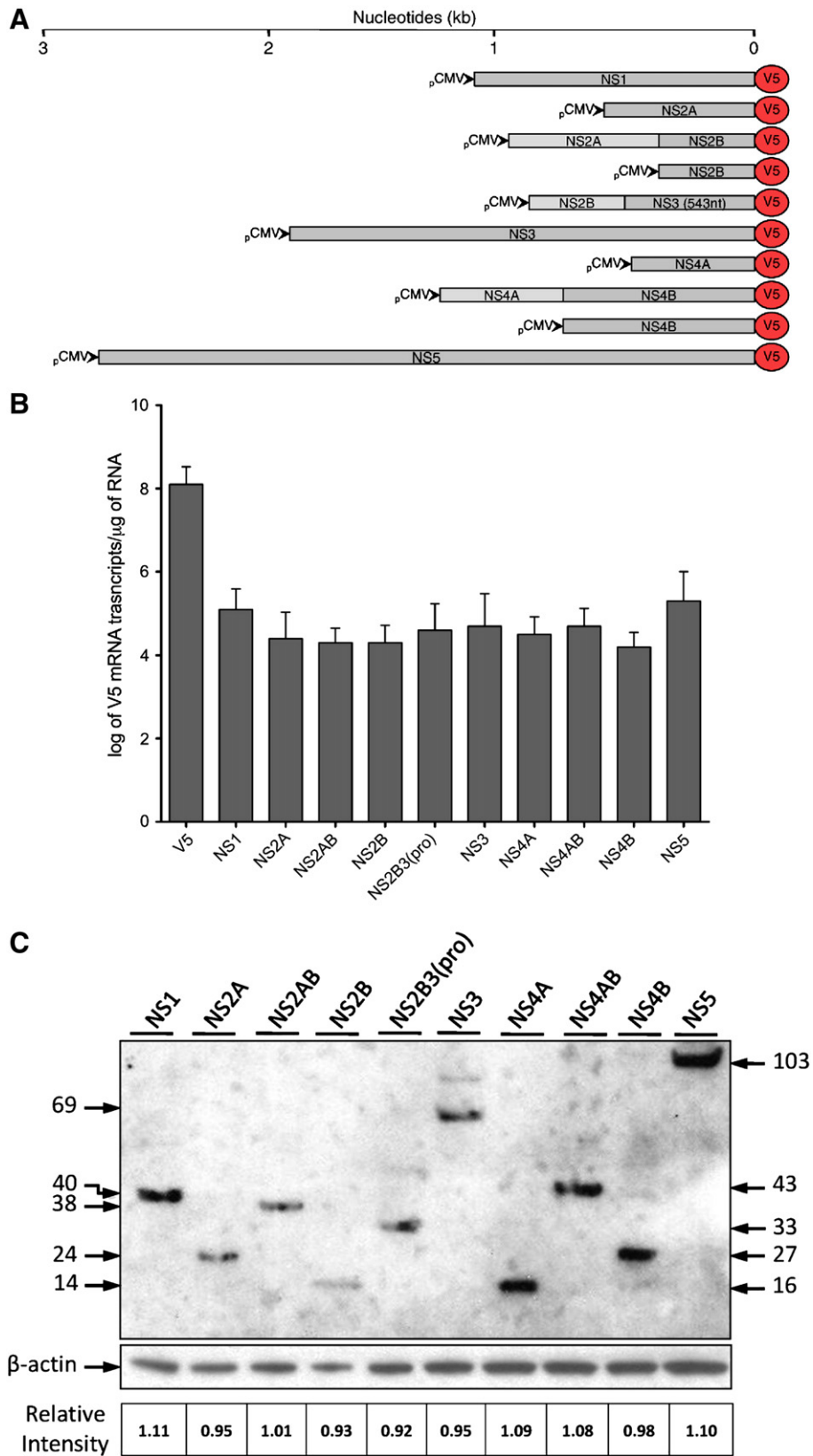


Fig. 2. Expression of nonstructural DENV-V5 fusion protein constructs in THP-1 cells. THP-1 cells were transfected with each DENV nonstructural gene plasmid and cell lysates were collected at 40 h after transfection for expression analysis by qRT-PCR and western blot. (A) Constructs and approximate nucleotide number (kb) of cloned viral genes. Each gene was constructed using a V5 reporter vector having the CMV promoter for expression in mammalian cells. (B) V5 copy numbers were determined for each expressing plasmid using qRT-PCR and reported as log of V5 mRNA transcripts per μ g of RNA. A standard curve was prepared by serially diluting known amount of V5 transcripts. (C) Western blot analysis of DENV-V5 proteins expression. Twenty microliter (approximately 50 μ g) of each cell lysate was loaded and electrophoresed on SDS-PAGE followed by immunoblotting with V5 antibody. Protein loading amounts were confirmed using β -actin and a relative intensity (RI) was calculated by dividing the absolute intensity of the DENV-V5 protein band by the absolute intensity of its corresponding β -actin band. The cell lysate loaded in each well is specified at the top and its corresponding molecular mass (kDa) is depicted at the left or right. Bars represent the mean \pm SD of three independent experiments conducted in duplicate. No significant differences ($p < 0.05$) were found across transcripts.

NGC-infected THP-1 cells produced DHF-associated immunomediators and that maximum transcript expression and protein secretion levels corresponded with peak viral titers and copy numbers. These data are consistent with other *in-vitro* data demonstrating that infected THP-1 cells secrete elevated chemokines and cytokines, which corresponds with increased viral titers (Chareonsirisuthigul et al., 2007; Hober et al., 1996; Ubol et al., 2008).

NS4B or NS5 expressed in THP-1 cells induces IL-6, IL-8 and IP-10

To evaluate the role of DENV NS on the observed induction of immunomediators, we constructed plasmids (Fig. 2A) with a commercially available V5 expression vector using primers designed from the wild-type NGC genome (Table 1). Following the manufacturer's optimization instructions for the Neon™ system, we determined that maximum expression levels of the V5 control plasmid or NS5-V5 plasmid and THP-1 cell viability (approximately 70% for all plasmids) occurred at 40 h after transfection (Supplemental Fig. 2). Therefore, successful expression of each DENV plasmid was confirmed at 40 h after transfection by immunofluorescence assay (data not shown), qRT-PCR and western blot (Figs. 2B–C). We conducted qRT-PCR using primers designed to detect plasmid expressing V5 mRNA (Table 2) and demonstrated similar mRNA expression levels, wherein approximately log 4.25 copies of transcripts were expressed by all DENV-V5 fusion plasmids with no significant difference between the minimum (NS4B) and maximum

(NS5) expression levels ($p=0.32$) (Fig. 2B). Also, we conducted western blot and demonstrated similar relative intensities (RIs), calculated as described in Materials and methods, for all DENV proteins, wherein NS1 reached a maximum RI of 1.11 and NS2B3 (pro) reached a minimum RI of 0.92 (Fig. 2C). Having established similar transcript and protein expression levels for all DENV plasmids at 40 h after electroporation, we screened transfected cells for immunomediator induction using qRT-PCR. We demonstrated that NS4B significantly increased IL-6 transcripts approximately 4-fold and IL-8 almost 9-fold relative to NS4AB (Fig. 3). Moreover, NS5 significantly increased IL-6 transcripts approximately 8-fold and IL-8 over 11-fold relative to NS4AB (Fig. 3). However, relative to NS4AB expression NS4B ($p=0.07$) or NS5 ($p=0.09$) did not significantly increase TNF α transcripts levels (Fig. 3).

Given that NS4B and NS5 induced DHF-associated immunomediator transcripts, we examined the culture supernatant for secreted immunomediators induced by NS4B, NS5 and the viral replication complex-associated proteins NS2B3(pro), NS3 and NS4AB. As an alternative method to confirm plasmid expression efficiencies, we conducted flow cytometry using a V5 antibody to detect each DENV-V5 fusion protein (Fig. 4A). We observed similar protein levels for all plasmids tested wherein a range of 12–18% of THP-1 cells expressed DENV-V5 fusion proteins (Fig. 4A). Having established similar NS-V5 transcript and protein expression levels by qRT-PCR, western blot and flow cytometry, we quantitated secreted immunomediator levels in the culture supernatants using

Table 1
Primers employed for the construction of DENV gene plasmids.

Construct and primer names ^a	Nucleotide position ^b	Primer sequence (5'–3') ^c	Tm (°C)	Amplicon size (bp)	PCR cycling conditions
NS1					
NS1-forward (F)	2422–2443	accaccatg GATAGTGGTTGCGTTGTGAGCT	65.9	1056	95 °C 5 min 1 cycle; 95 °C 45 s, 57 °C 45 s, 72 °C 1 min, 30 cycles; 72 °C 7 min 1 cycle
NS1-reverse (R)	3477–3449	GGCTGTGACCAAGGAGTTGACCAAAATCT	62.3		
NS2A					
NS2A-F	3478–3499	accaccatg GGACATGGGCAGATTGACAACCT	66.1	654	95 °C 5 min 1 cycle; 95 °C 30 s, 55 °C 30 s, 72 °C 45 s, 30 cycles; 72 °C 7 min 1 cycle
NS2A-R	4131–4102	CCTTTTCTTGTGGTTCTTGAAGGGTTGT	59.6		
NS2B					
NS2B-F	4132–4149	accaccatg ctAGCTGGCCACTAAATGAG	64.9	390	95 °C 5 min 1 cycle; 95 °C 30 s, 55 °C 30 s, 72 °C 45 s, 30 cycles; 72 °C 7 min 1 cycle
NS2B-R	4521–4995	CCGTTGTTTCTCACTCCACAGGTA	60.5		
NS2AB					
NS2A-F	3478–3499	accaccatg GGACATGGGCAGATTGACAACCT	66.1	1044	95 °C 5 min 1 cycle; 95 °C 45 s, 55 °C 45 s, 72 °C 1 min, 30 cycles; 72 °C 7 min 1 cycle
NS2B-R	4521–4995	CCGTTGTTTCTCACTCCACAGGTA	60.5		
NS2B3(pro)					
NS2B-F	4132–4149	accaccatg ctAGCTGGCCACTAAATGAG	64.9	933	95 °C 5 min 1 cycle; 95 °C 45 s, 53 °C 45 s, 72 °C 1 min, 30 cycles; 72 °C 7 min 1 cycle
NS3pro-R	5064–5035	GTCATCTCGATCTCTGGATTGTCTTCAAT	58.1		
NS3					
NS3-F	4522–4543	accaccatg GCTGGAGTATTGTGGGATGTCC	66.0	1854	95 °C 5 min 1 cycle; 95 °C 45 s, 55 °C 45 s, 72 °C 1 min, 30 cycles; 72 °C 7 min 1 cycle
NS3-R	6375–6348	CTTTCTCCAGCTGCAAACTCCTTGAAT	59.5		
NS4A					
NS4A-F	6376–6393	accaccatg ctTCCTGACCTGAACCTA	64.1	450	95 °C 5 min 1 cycle; 95 °C 30 s, 59 °C 30 s, 72 °C 45 s, 30 cycles; 72 °C 7 min 1 cycle
NS4A-R	6825–6800	TGCCATGGTGGCGGCCCACTGTGA	68.6		
NS4A(-2K)					
NS4A-F	6376–6393	accaccatg ctTCCTGACCTGAACCTA	64.1	381	95 °C 5 min 1 cycle; 95 °C 30 s, 59 °C 30 s, 72 °C 45 s, 30 cycles; 72 °C 7 min 1 cycle
NS4A(-2K)-R	6756–6731	TCTCTGTTTTCTGGTTCTGGAATAA	59.3		
NS4B					
NS4B-F	6826–6846	accaccatg ctAACGAGATGGGTTTCTGGAA	66.4	744	95 °C 5 min 1 cycle; 95 °C 45 s, 55 °C 45 s, 72 °C 1 min, 30 cycles; 72 °C 7 min 1 cycle
NS4B-R	7569–7540	CTTCTCGTGTGGTTGTGTCTTCATGAT	60.5		
NS4AB					
NS4A-F	6376–6393	accaccatg ctTCCTGACCTGAACCTA	64.1	1194	95 °C 5 min 1 cycle; 95 °C 45 s, 55 °C 45 s, 72 °C 1 min, 30 cycles; 72 °C 7 min 1 cycle
NS4B-R	7569–7540	CTTCTCGTGTGGTTGTGTCTTCATGAT	60.5		
NS4B-GFP-R	7569–7540	gCCTTCTCGTGTGGTTGTGTCTTCATGAT	60.5		
2KNS4B					
2KNS4B-F	6757–6777	accaccatg ctACACCCCAAGATAACCAATTG	64.4	813	95 °C 5 min 1 cycle; 95 °C 45 s, 55 °C 45 s, 72 °C 1 min, 30 cycles; 72 °C 7 min 1 cycle
NS4B-R	7569–7540	CTTCTCGTGTGGTTGTGTCTTCATGAT	60.5		
NS5					
NS5-F	7570–7591	accaccatg GGAATCGCAACATAGGAGAGA	65.2	2700	95 °C 5 min 1 cycle; 95 °C 1 min, 59 °C 60 s, 72 °C 1 min, 30 cycles; 72 °C 7 min 1 cycle
NS5-R	10,269–10,239	CCACAGGACTCTGCCTCTCTCTCTTTT	64.0		

^a Forward, sense sequence; reverse, antisense sequence.

^b Determined from NCBI accession number M29095 (DENV NGC strain).

^c Kozac sequence shown as bold lower-case letters, fill-in nucleotides for in-frame translation as lower case letters and DENV sequence as upper-case letters.

Table 2

Primers employed for the analysis of DENV and host genes by qRT-PCR.

Gene and primer names ^a	Primer sequence (5'–3')	Tm (°C)	Amplicon size (bp)
DENV2 capsid ^b			
DN-forward (F)	CAATATGCTGAAACGCGAGAG	55.8	167
DN-reverse (R)	CATCTATTAGAATCCCTGCT	58.3	
V5 epitope ^b			
V5-F	TCTGCAGATATCCAGCACAGT	55.8	87
V5-R	ACCGAGGAGAGGGTTAGGGAT	59.2	
GAPDH ^c			
GAPDH-F	AGTTAGCCGCATCTCTTTTTC	56.1	100
GAPDH-R	CAATACGACCAATCCGTTGAC	55.3	
IL-6 ^c			
IL6-F	CCAGGAGCCCAGCTATGAAC	57.8	63
IL6-R	CCCAGGGAGAAGGCAACTG	58.2	
IL-8 ^c			
IL8-F	GAAGTGAAGTATTGAGAGT	55.1	133
IL8-R	CTCTTCAAAACTTCTCCACA	56.6	
IP-10 ^c			
IP10-F	CCACGTGTTGAGATCATTGC	54.3	139
IP10-R	GCTCCCTCTCGTTTTAAGG	55.4	
TNF- α ^c			
TNF α -F	CCCAGGGACCTCTCTAATCA	55.9	79
TNF α -R	GCTACAGGCTTGCTACTCGG	58.6	

^a Forward, sense sequence; reverse, antisense sequence.^b Real-time PCR primers used to determine DENV2 NGC and V5 copy numbers.^c Real-time RT-PCR primers to amplify immunomediator genes.

the Luminex® technology. As expected, NS4B or NS5 significantly induced IL-6, IL-8, IP-10 and IFN γ when compared to cells expressing only the V5 epitope (Fig. 4B). Cells expressing NS4B secreted 57, 1100 and 48 pg/mL of IL-6, IL-8 and IP-10, respectively, while cells expressing NS5 secreted slightly higher levels of these immunomediators (Fig. 4B). NS4B and NS5 increased TNF α production but the difference was not significant, $p=0.11$ and 0.08 , respectively. However, DENV-infected or LPS-treated cells significantly induced TNF α (Fig. 1B) indicating an alternative route of induction. When we electroporated the THP-1 cells without plasmid DNA (pDNA) or the V5 vector control, we observed chemokine and cytokine expression levels similar to

endogenous THP-1 levels (Fig. 4B and data not shown). Cell viability reached approximately 70% at 40 h after treatment, including the no-plasmid transfected and LPS-treated control cells (Supplemental Fig. 2), further indicating that the differences observed were due to the expression of NS4B and NS5.

Maturation of NS4B via cleavage of NS4AB enhances IL-6 and IL-8 production

During replication, the DENV protease NS2B3(pro) interacts with the NS4AB polyprotein presumably as part of the viral replication complex. Previous work has demonstrated sequential processing of NS4AB wherein NS2B3(pro) cleaves NS4AB, releasing NS4A from NS4B (Cahour et al., 1992; Miller et al., 2006). We proposed that expressing both NS4AB and NS2B3(pro) in THP-1 cells would best mimic natural processing and maturation of NS4B and possibly affect the immunomediator induction pattern observed during the expression of individual viral proteins. To detect THP-1 cells expressing both NS4AB and NS2B3(pro), we constructed the NS4AB-GFP fusion protein plasmid (Fig. 5A) for co-expression with the NS2B3(pro)-V5 fusion protein. We demonstrated by flow cytometry that approximately 9% of electroporated THP-1 cells co-expressed NS4AB-GFP and NS2B3(pro)-V5, 23% expressed NS4AB-GFP and 12, 14 and 17% expressed NS4B, NS2B3(pro) and NS5, respectively, at 40 h after electroporation (Fig. 5B). To confirm the flow cytometry data and examine functional cleavage events, we conducted western blot using cells co-expressing NS4AB and the viral protease NS2B3(pro). As expected, we observed that the co-expression resulted in the cleavage of NS4AB which produced four distinct bands: NS4AB, NS2B3(pro), 2KNS4B and NS4B (Fig. 5C). Co-expression of NS4AB and NS2B3(pro) significantly enhanced the induction of IL-6 and IL-8 transcripts when compared to NS4B or NS5 alone, approximately doubling the relative fold change of IL-6 transcripts from 4- to 8-fold and IL-8 transcripts from 9- to 18-fold (Fig. 5D). Given that NS4B co-localizes with NS3 and NS5 during IFN antagonistic function and as part of the viral replication complex (Mazzon et al., 2008; Miller et al., 2006; Umareddy et al., 2006), we tested the induction potential of NS4B with NS3 or NS5 and demonstrated that the induction potential was unaltered relative to NS4B or NS5 alone for IL-6 or IL-8 (data not shown), further supporting our results that enhancement occurs primarily during NS4B maturation and polyprotein

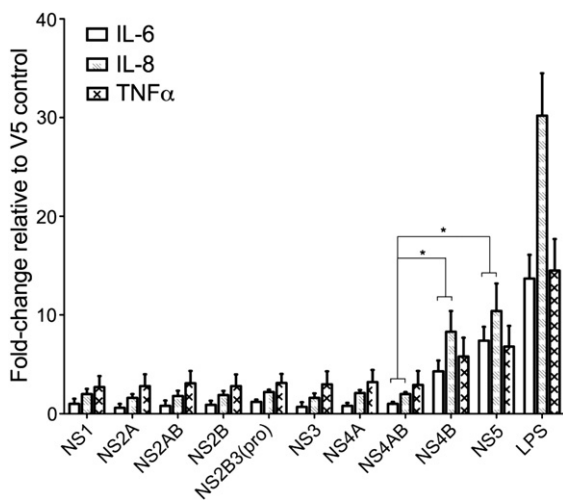


Fig. 3. DENV NS4B and NS5 induce IL-6, IL-8, and TNF α transcripts in THP-1 cells. THP-1 cells expressing individual DENV nonstructural genes were harvested at 40 h after transfection and total mRNA was isolated for immunomediator transcript analysis by qRT-PCR. The fold change induction of IL-6, IL-8, and TNF α transcripts was determined for each DENV gene relative to V5 controls. LPS was used as a positive control and GAPDH was used as the endogenous control. Bars represent the mean \pm SD of three independent experiments conducted in duplicate. (*) indicates statistical significance at $p < 0.05$ as compared to transcript levels induced by the NS4AB-V5 fusion protein.

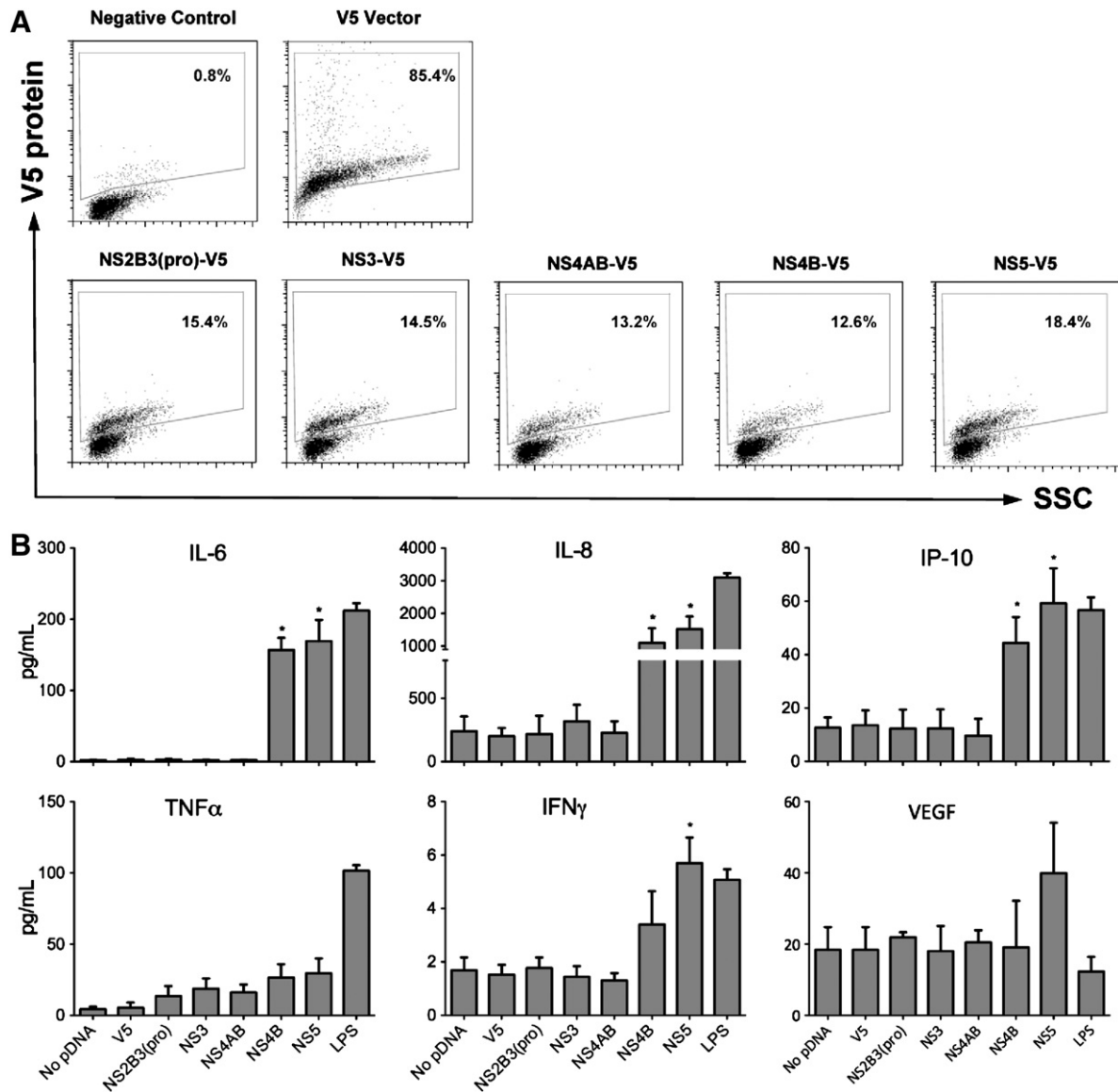


Fig. 4. DENV NS4B and NS5 induce DHF-associated immunomediators in THP-1 cells. THP-1 cells were electroporated with DENV NS and after 40 h, cells and supernatants were collected to measure expression efficiencies and immunomediator protein levels, respectively. (A) Percent of THP-1 cells expressing DENV-V5 fusion proteins as determined by flow cytometry. Cells were fixed, permeabilized and V5-expression proteins were detected using an anti-V5 primary antibody and Alexa Fluor® 594 secondary antibody. (B) Chemokine and cytokine levels (pg/mL) in the supernatants of THP-1 cells expressing NS at 40 h after electroporation and as determined by the Luminex® multiplex technology. Bars represent the mean \pm SD of three independent experiments conducted in duplicate and (*) indicates statistical significance at $p < 0.05$ as compared to vector controls.

cleavage events. Interestingly, the cleavage event did not significantly enhance the induction of IP-10, suggesting that multiple pathways may be responsible for inducing immunomediators. Although this result is unexpected, signal transduction crosstalk is regulated in a dynamic manner and may differ under homeostatic and pathologic conditions making it difficult to determine the exact mechanisms responsible for induction outcomes (Hu et al., 2007). However, taken together these results suggest that NS4AB expressed with NS2B3(pro) initiates sequential processing of NS4AB and maturation of NS4B (Miller et al., 2007; Miller et al., 2006), thereby enhancing the induction of immunomediators.

Expression of the 2K-signal peptide with NS4B is sufficient to enhance IL-6 and IL-8 induction

The DENV 2K-signal sequence is a 17-amino acid peptide linking NS4A with NS4B and has little known function. Having observed the cleavage product 2KNS4B in the co-transfected cells, we questioned whether the

2K-signal sequence was responsible for the observed enhanced induction of immunomediator transcripts. To address this question, we constructed the plasmids 2KNS4B and NS4A without 2K (-2K) (NS4A-2K) (Fig. 6A) and expressed them alone or in combination with other NS, particularly NS2B3(pro) due to its intrinsic enzymatic properties and involvement in the replication complex. By using qRT-PCR and western blot, we demonstrated the production of approximately log 4 copies of V5 mRNA transcripts by each plasmid or combination of plasmids (Fig. 6B) and cleavage of 2KNS4B, respectively (Fig. 6C). Importantly, 2KNS4B expression and cleavage, presumably by host proteases, significantly enhanced the induction of IL-6 and IL-8 transcripts (Fig. 6D). Expression of NS4A(-2K) with or without NS2B3(pro) did not induce transcripts of any of the immunomediators tested while NS4B expressing with NS2B3(pro) showed slightly reduced induction potential, suggesting that the viral protease may interfere with NS4B during the induction process (Fig. 6D). Overall, these data suggest that the induction of DHF-associated chemokines and cytokines by NS4B is enhanced when NS4AB is cleaved

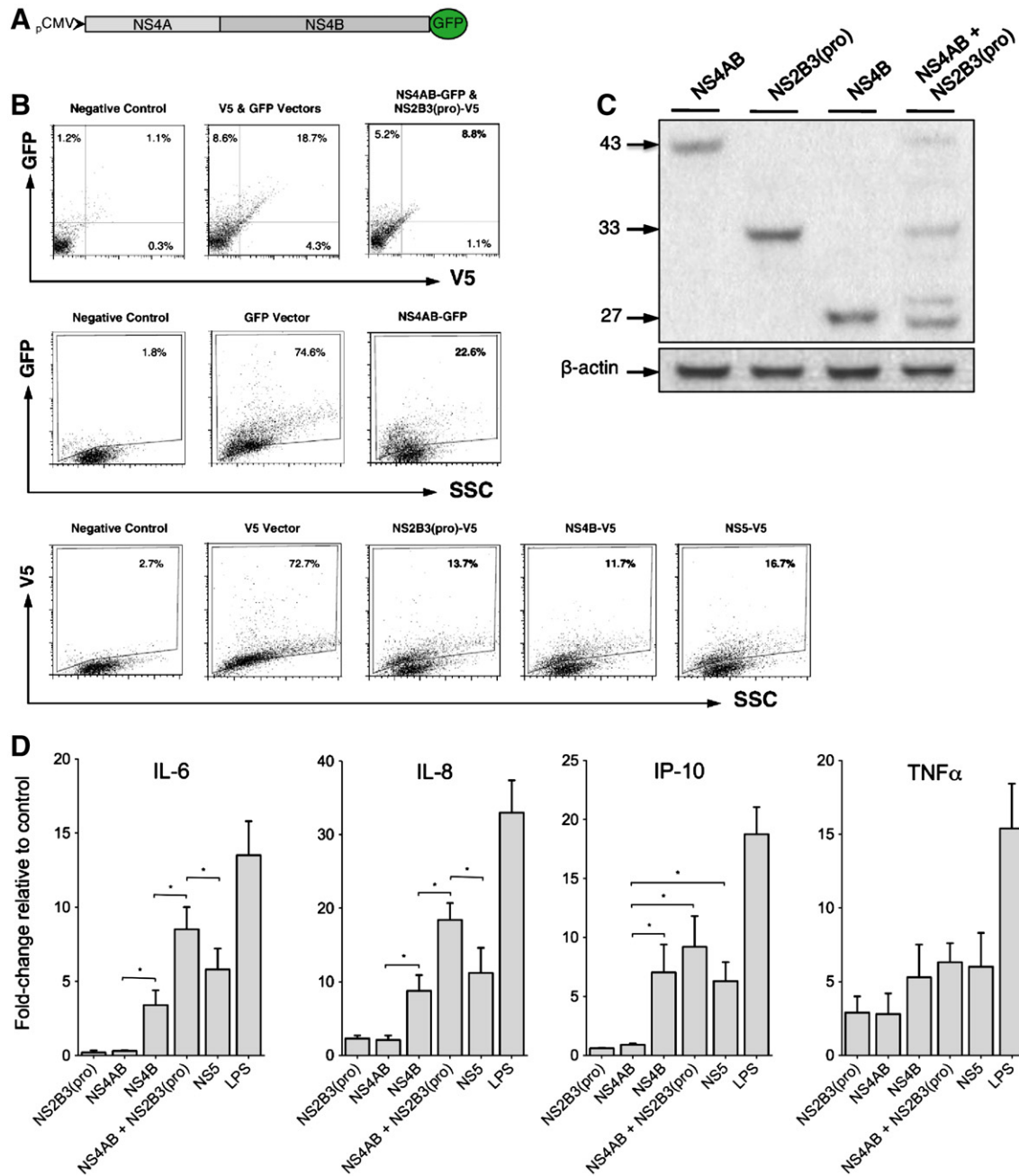


Fig. 5. Maturation of NS4B via processing of the polyprotein, NS4AB, enhances the production of IL-6 and IL-8. (A and B) The NS4AB gene was cloned into a GFP reporter vector and co-expressed with NS2B3(pro)-V5 fusion protein in THP-1 cells for detection by flow cytometry. The expression efficiencies were measured at 40 h after transfection and expressed as percent of THP-1 cells expressing DENV-V5-GFP, V5 or GFP fusion proteins. (C) Western blot analysis of THP-1 cells expressing DENV-V5 fusion proteins NS 4AB, 2B3(pro), 4B and both 4AB and 2B3(pro). β-actin was used as the endogenous control and estimated molecular weight markers (kDa) are shown on the left. (D) Induction of immunomediator transcripts in THP-1 cells relative to vector controls as determined by qRT-PCR. Bars represent mean ± SD of six experimental samples and (*) indicates statistical significance at $p < 0.05$.

by NS2B3(pro), specifically during 2KNS4B processing by host signalases and possibly during NS4B localization into the membrane structures (Miller et al., 2006, 2007; Roosendaal et al., 2006; Umareddy et al., 2006; Westaway et al., 1997a).

Discussion

It is well established that elevated levels of immunomediators detected in DHF/DSS patients can adversely alter endothelial cell integrity leading to plasma leakage and unresolved vascular permeability (Basu and Chaturvedi, 2008; Green and Rothman, 2006; Leong et al., 2007; Srikiatkachorn, 2009). However,

mechanisms underlying severe DENV immunopathogenesis remain elusive. The lack of an animal model to study dengue immunopathogenesis has led to extensive clinical and *in-vitro* studies suggesting that both host and viral factors influence elevated circulating immunomediators and severe disease outcomes (Kurane and Takasaki, 2001; Pang et al., 2007). DENV is capable of modulating intracellular signaling, such as the IFN response, or host inflammatory responses such as activation of the transcription factor NF-κB (Mogensen and Paludan, 2001a; Mogensen and Paludan, 2001b). DHF/DSS is presumed to occur when the virus outcompetes host antiviral responses and establishes infection while initiating a robust inflammatory response,

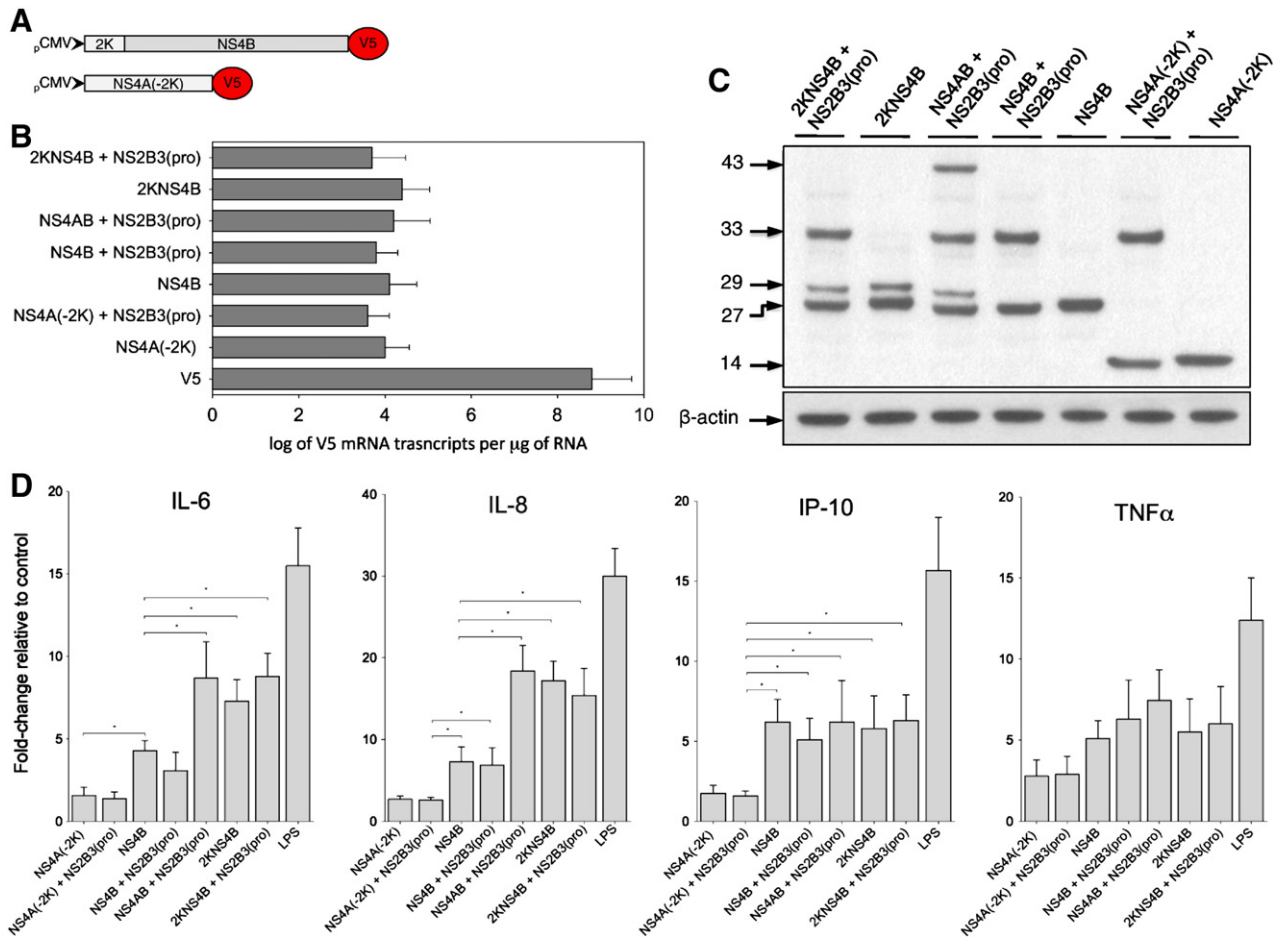


Fig. 6. NS4B with the 2K-signal sequence is sufficient to enhance the induction of immunomediators. (A) The DENV genes 2KNS4B and NS4A without the 2K-signal peptide, NS4A (-2K), were cloned in the V5 reporter vector. (B) Log of V5 mRNA transcripts in THP-1 cells expressing nonstructural plasmids at 40 h after transfection as determined by qRT-PCR. (C) Western blot analysis of THP-1 cells expressing DENV-V5 fusion proteins NS 2B3(pro) and 2K4B, 2K4B, 4AB and 2B3(pro), 2B3(pro) and 4B, 4B, 2B3(pro) and 4A(-2K), and 4A (-2K). Estimated molecular weights (kDa) are depicted on the left and β -actin was detected as the endogenous control. (D) Immunomediators' mRNA fold changes as compared to corresponding V5 controls; GAPDH was used as the endogenous control gene. Bars represent mean \pm SD of six independent experimental samples and (*) indicates statistical significance at $p < 0.05$.

which is critical for the progression of severe dengue disease. Although other investigators have examined DENV initiated IFN inhibition, very little is known regarding the mechanism by which the DENV proteins induce DHF/DSS-associated immunomediators. In this report, we demonstrate that maximum production of immunomediators from DENV-infected THP-1 cells corresponds with peak levels of secreted infectious virus and cellular viral RNA. Moreover, using a plasmid construct system that consistently expresses similar levels of viral gene transcripts and viral-V5 fusion proteins, we demonstrate that DENV NS4B and NS5 are potent inducers of DHF/DSS-associated immunomediators in THP-1 cells and that maturation of NS4B via host protease cleavage of 2KNS4B is a critical step during the observed induction.

Peak levels of chemokines and cytokines in DENV-infected THP-1 cells correspond with maximum virus yield

We demonstrate that DENV-infected THP-1 monocytes induce transcription and secretion of immunomediators such as IL-6, IL-8, IP-10, TNF α and IFN γ and that peak levels of these chemokine and cytokines correspond with an increased generation of infectious virus and viral RNA copy numbers. These data are consistent with previous reports demonstrating that secreted immunomediator levels directly correspond with the amount of virus production in

THP-1 cells (Chareonsirisuthigul et al., 2007; Ubol et al., 2008). Other investigators have demonstrated no infectious virus production from DENV-infected THP-1 cells using low passage clinical isolates and in the absence of enhancing antibodies (Diamond et al., 2000). This variation may be due to differences in virus passage, as high-passage virus isolates *in-vitro* may acquire dominant mutations that exhibit aberrant phenotypic characteristics not found in low-passage isolates (Rico-Hesse et al., 1998). Our data suggest that immunomediator induction occurs during virus replication when viral RNA copies are elevated. Moreover, others have demonstrated that the antiviral drug ribavirin inhibits DENV replication and IL-6 and IL-8 production in a dose dependent manner (Huang et al., 2000). Regardless, it remains difficult to pinpoint the precise immunomediator initiation step due to the dynamic characteristics of virus replication. It is possible that immunomediator induction corresponds with the initial translation of input positive-strand viral RNA, negative-strand viral RNA synthesis, viral protein production or packaging, and/or secretion of virions. Functional studies demonstrate that the expression of all NS in THP-1 cells induces the activation of NF κ B, the ubiquitous transcription factor involved in the rapid production of many chemokines and cytokines (Hershkovitz et al., 2008). Similarly, expression of NS or NS5 alone in human embryonic kidney cells induces IL-8 transcription and protein secretion (Medin

et al., 2005; Pryor et al., 2007), further suggesting that NS effectively modulates intracellular signaling and initiate immunomediator induction.

NS5 induces DHF-associated immunomediators in THP-1 monocytes

We demonstrate that NS5 induces high levels of IL-8 as well as IL-6, IP-10 and IFN γ transcripts and protein secretion in THP-1 cells. Moreover, it appears that NS5 triggers a rapid and sustained production of immunomediators as demonstrated by increased transcript and protein secretion at 24 and 72 h after DENV infection, and 20 and 40 h after transfection with NS5 constructs, respectively. NS5 has a potent induction potential as demonstrated by consistent and equivalent expression levels of all tested viral gene transcripts and V5 fusion proteins. Although we observed that NS5 localized to the nucleus and the cytoplasm at 20 and 40 h after transfection, most NS5 localized to the nucleus (data not shown) consistent with other reports (Pryor et al., 2007; Rawlinson et al., 2009).

The largest and most highly conserved protein among the flaviviruses, NS5 serves as the RNA-dependent RNA polymerase (RdRP) and methyltransferase (Chung et al., 2010; Dong et al., 2010; Egloff et al., 2007; Harris et al., 2006; Zhou et al., 2007; Zou et al., 2011). Other investigators have reported that DENV NS5 induces IL-8 via the activation of the CAAT/enhancer binding protein (c/EBP) and NF κ B (Medin et al., 2005) as it shuttles between nuclear and cytoplasmic compartments (Pryor et al., 2007; Rawlinson et al., 2009). Upon positive-sense RNA translation and polyprotein processing in the cytoplasm, soluble NS5 is thought to undergo phosphorylation allowing the host proteins importin- α/β to bind at two distinct nuclear localization sequences (NLS) and import NS5 into the nucleus (Pryor et al., 2007). While hyperphosphorylated NS5 localizes to the nucleus, its dephosphorylation elicits a conformational change, allowing exportin-1 to bind and translocate NS5 to the cytoplasm where it interacts with NS3 as part of the viral replication complex (Kapoor et al., 1995; Mackenzie et al., 2007; Rawlinson et al., 2009). Although the function of nuclear NS5 is unclear, greater nuclear accumulation of NS5 results in decreased IL-8 production and *vice versa* (Pryor et al., 2007; Rawlinson et al., 2009), suggesting that the observed induction of immunomediators occurs when NS5 is localized in the cytoplasm. Moreover, because the majority of NS5 localizes to the nucleus and only a relatively small amount of NS5 is required for virus replication in the cytoplasm (Grun and Brinton, 1987; Uchil and Satchidanandam, 2003), it is probable that very low levels of cytoplasmic NS5 is required to initiate immunomediator induction. Although the induction mechanisms by NS5 have not been defined, it is possible that NS5 triggers c/EBP and NF κ B signaling events during NS5/importin interactions in the cytoplasm. However, this theory remains to be tested. Recent studies demonstrate that NS5 interacts with NS4B in the cytoplasm, and inhibits IFN signaling by binding and preventing STAT2 phosphorylation (Ashour et al., 2009; Mazzon et al., 2009; Mazzon et al., 2008), further suggesting that viral/host protein interactions are important for modulating host signaling.

NS4B induces DHF-associated immunomediators in THP-1 monocytes

In screening the induction potential of DENV NS in monocytes, we demonstrate for the first time that NS4B induces the secretion of IL-6, IL-8, IP-10 and IFN γ . The potent NS4B induction pattern is a surprise given its highly hydrophobic nature and intimate integration within the ER membrane. Yet, we demonstrate that NS4B has an early and sustained induction potential as suggestive of its ability to induce IL-8 at 20 and 40 h after transfection. A highly hydrophobic protein, NS4B consists of three endoplasmic reticulum (ER) membrane-spanning segments and may serve as an anchor for the viral replication complex (Miller et al., 2007; Miller et al., 2006; Welsch

et al., 2007). NS4B can act as a potent IFN antagonist via interference and degradation of cellular STAT1 (Munoz-Jordan et al., 2003) which is synchronized with NS4B maturation via NS4AB cleavage by the viral protease, NS2B3(pro) (Munoz-Jordan, 2010; Munoz-Jordan et al., 2005). Interestingly, IFN- β -mediated inhibition of IL-8 expression requires the presence of STAT1 and STAT2 (Laver et al., 2008). Given that NS4B maturation inhibits IFN signaling via STAT1 degradation, we hypothesized that NS4B maturation via NS4AB proteolytic modification(s), including host signalase cleavage of 2KNS4B, would lead to increased levels of IL-8 and possibly other immunomediators. As such, we tested the induction potential of the processing events involved in NS4B maturation.

Maturation of NS4B via processing of 2KNS4B enhances the induction of immunomediators

During polyprotein processing, NS4AB is cleaved by the viral protease NS2B3(pro), releasing NS4A from NS4B having the 2K-signal peptide (2KNS4B) before a host signalase cleaves the 2KNS4B junction in the lumen of the ER to release mature NS4B and allowing for its integration into the ER membrane (Cahour et al., 1992; Lin et al., 1993; Miller et al., 2006). To mimic natural NS4B maturation via polyprotein processing, we co-expressed NS4AB with NS2B3(pro) and observed that polyprotein cleavage events initiated a more potent induction of IL-6, IL-8 and IP-10 transcripts than NS4B or NS5 alone. The enhancement appears to be due to NS4AB processing or maturation and localization of NS4B in the ER, not increased levels of NS expression as demonstrated by similar expression levels of viral protein and gene transcripts. Moreover, our data suggest that the 2K-signal peptide is not required for NS4B induction, yet post-translational proteolytic cleavage events including host signalase cleavage of the 2K-peptide from NS4B is primarily responsible for the enhancement.

The 2K-signal peptide appears to play an important role during a variety of intracellular alterations, including anchoring of NS4B to the ER membrane (Lin et al., 1993). Proteolytic removal of the 2K-peptide from NS4A during DENV replication induces membrane alterations that may harbor the viral replication complex (Miller et al., 2007) and the West Nile virus (WNV) 2K-peptide is important for cytoplasmic rearrangements, foci development and Golgi trafficking (Roosendaal et al., 2006). Moreover, IFN inhibition by NS4B appears to require the presence of the 2K-signal peptide for at least three flaviviruses, DENV, WNV and yellow fever virus (YFV) (Munoz-Jordan et al., 2005). Interestingly, the initial 125 amino acid residues of DENV 2KNS4B, predicted to be located in the cytoplasm between the first and second transmembrane domains, are required for IFN antagonism (Munoz-Jordan et al., 2005), while the YFV 2K-peptide may be important for the translocation of cleaved NS4B to the lumen side of the ER (Chambers et al., 1989).

Given that IFN- β -mediated inhibition of IL-8 production requires the presence of STAT1 and STAT2 (Laver et al., 2008), and NS5 or processing of 2KNS4B by host proteases is synchronized with IFN inhibition via STAT1 and STAT2 degradation, it will be interesting to determine whether both IFN inhibition and immunomediator induction occur in the presence of mature NS4B as intracellular STAT1 and STAT2 diminish. Overall, we demonstrate that the viral protease has no intrinsic induction potential alone, yet it cleaves NS4AB and generates NS4A(-2K) and 2KNS4B, which is then cleaved by a host protease, perhaps allowing NS4B to reorient across the ER and initiate IFN inhibition and immunomediator induction.

Conclusions

Effective host defenses against DENV infection include a robust IFN response and a rapid production of chemokines and cytokines

important for immune signaling. Importantly, DENV and other flaviviruses are capable of inhibiting host IFN signaling to establish productive infections and concurrently promote an overproduction of immunomediators associated with severe disease. In this report, we describe DENV-associated immunomediator induction patterns in the monocytic cell line, THP-1. We have demonstrated that THP-1 cells sustain excellent viability after electroporation, are permissive to DENV infection, and thus are an ideal model to examine early events after infection or expression of DENV NS proteins. We also demonstrated that NS4B maturation events, specifically cleavage of 2KNS4B by host proteases, induce significantly higher levels of immunomediators than NS4B or NS5 alone. Therefore, it appears that maturation of NS4B is important for both IFN inhibition and immunomediator induction. Given the shared phenotype of IFN antagonism across the virus family, it is quite possible that IFN inhibition and immunomediator induction patterns induced during NS4B maturation are similar for other flaviviruses. Even though *Flavivirus*-wide NS4B sequences show divergent and sometimes negligible genotypic resemblance (Lundin et al., 2003; Qu et al., 2001), predicted flavivirus NS4B structural topologies, including the multiple transmembrane and cytoplasmic regions, are strikingly similar as well as NS4AB cleavage events. Reduced flavivirus protein expression or polyprotein processing during infection may dampen virus-induced IFN inhibition and the overproduction of potentially deleterious immunomediators, minimizing the risk of severe disease. As such, NS4B may very well be an attractive target for the development of *Flavivirus*-wide therapeutic interventions.

Materials and methods

Virus and cell culture

DENV-2 New Guinea C (NGC) strain was obtained from Dr. Duane Gubler at the University of Hawaii at Manoa. A virus stock was produced by passaging virus twice in C6/36 cells. Given the extensive use of the THP-1 cell line by others examining DENV pathogenesis and due to their permissibility to DENV infection (Chareonsirisuthigul et al., 2007; Chen et al., 2009; Hershkovitz et al., 2008; Hober et al., 1996; Ubol et al., 2008), we chose to work with THP-1 cells in this study. The THP-1 cells were obtained from the American Type Culture Collection (ATCC, Rockwell, MD) and were cultured in RPMI-1640 (ATCC) supplemented with 1% penicillin/streptomycin, 10% heat-inactivated fetal bovine serum (FBS) (Gibco Labs, Grand Island, NY) and 2-mercaptoethanol to a final concentration of 0.05 mM; cells were incubated at 37 °C in a 5% CO₂ atmosphere. The Vero cells (monkey kidney epithelial cells) (ATCC) were maintained in M199 and supplemented with 1% penicillin/streptomycin and 10% FBS.

Infection of THP-1 cells

For infection experiments, 1×10^6 THP-1 cells were infected with DENV at MOI-0.1 or -1. After 1.5 h at 37 °C and 5% CO₂, the cells were washed and further cultured with fresh growth media. UV-inactivated DENV was generated using previously published protocols (Anderson et al., 1997; Verma et al., 2009). Briefly, DENV was diluted in 500 μ L PBS in a 35-mm culture plate and exposed to UV radiation using a Stratalink 2400 device (Stratagene) for 10 min. Inactivation of virus infectivity was verified by plaque assay using Vero cells. Mock-infected control THP-1 cells were infected with UV-inactivated DENV and for positive controls, THP-1 cells were incubated with 1 μ g/mL of LPS for 1 h, washed and incubated with fresh growth media until collection at 24 h. Every 24 h, cells and supernatants were collected while remaining cells were replenished with fresh growth media. Infectious virus released from infected cells was confirmed by plaque assay on Vero cells, as described below.

Plaque assay

To determine the amount of infectious virus released from DENV-infected THP-1 cells, plaque assay was conducted using Vero cell monolayers as described previously (Lambeth et al., 2005). Briefly, 2.5×10^5 Vero cells per well were seeded in 6-well culture plates (Corning, Lowell, MA) and incubated for 2 to 3 days until confluent. Supernatants from DENV-infected THP-1 cells were serially diluted using 10-fold dilutions in Dulbecco's Modified Eagles Medium (DMEM) (Gibco Labs) supplemented with 10% FBS and 100 μ L of each dilution was added to each well of the Vero cells followed by incubation at 37 °C and 5% CO₂ for 1 h with rocking every 15 min. Three milliliters of primary nutrient agar containing 1% SeaKem® LE agarose (Lonza, Walkersville, MD) was added to each well and the plates were incubated for 5 days. Three milliliters of secondary nutrient agar (primary agar containing 1% neutral red) was added to each well and the plates were incubated for an additional 2 days before counting plaques and calculating viral titers (Lambeth et al., 2005). Titers were expressed as plaque forming units (PFU) per mL.

Construction of plasmids

We employed standard molecular biology techniques to clone the DENV genes (Sambrook and Russell, 2001). Each primer listed in Table 1 was designed from the DENV-2 NGC reference genome, NCBI accession number M29095. All forward primers contain the Kozac sequence and ATG start codon (Table 1). The PCR step was conducted using AmpliTaq Gold® DNA polymerase (Applied Biosystems, Carlsbad, CA) with either the first-strand cDNA template (iScript® cDNA synthesis kit, Bio-Rad Inc., Hercules, CA) synthesized from the wild-type DENV-2 NGC RNA extracted from stock viral supernatant using the QIAamp viral RNA mini kit (Qiagen, Hilden, Germany) or the DENV-2 NGC DNA infectious clone (kindly provided by Barry Falgout, Food and Drug Administration) (Polo et al., 1997). Following the manufacturer's protocol, PCR products were cloned into either the pcDNA3.1/V5-His-TOPO® or the pcDNA3.1/CT-GFP-TOPO® vector (Invitrogen, Carlsbad, CA) having the CMV promoter for mammalian expression and detection. Proper gene orientation and identity were confirmed by DNA sequencing (Greenwood Molecular Biology Facility, University of Hawaii) and sequence analysis was conducted using Sequencer® 4.10.1 (Gene Codes Corporation, Ann Arbor, MI). Selected plasmids were isolated using Cesium Chloride ethidium bromide equilibrium centrifugation as previously described (Sambrook and Russell, 2001).

Transfections

In order to express the DENV-V5 fusion proteins, we transfected each construct into THP-1 cells using the Neon™ Transfection System (Invitrogen) after performing a series of optimization protocols as specified by the manufacturer. Briefly, 5×10^5 cells were electroporated with 1 μ g of plasmid DNA using 1250 V and 40 ms for 1 pulse. Cells were directly added to growth media without penicillin/streptomycin and immediately incubated at 37 °C and 5% CO₂. The supernatants and cells were harvested at 40 h after electroporation and used for the assays described below.

Flow cytometry

Intracellular expression of DENV-V5 or -GFP fusion proteins was detected at 40 h after transfection using a Guava EasyCyte flow cytometer (Guava, Hayward, CA). Cells expressing DENV-V5 proteins were washed, fixed and permeabilized (Fix & Perm reagents, Invitrogen) for intracellular labeling with a mouse monoclonal V5 antibody (Invitrogen) diluted 1:500 in PBS and a secondary antibody against mouse IgG coupled with Alexa Fluor® 594 antibody

(Invitrogen). Similarly, cells expressing DENV-GFP were washed, fixed and resuspended in PBS for flow cytometry. Data were analyzed using FlowJo software version 4.3 and expressed as percent cells expressing DENV protein.

Quantitative real-time RT-PCR (qRT-PCR)

Total cellular RNA was extracted from DENV-infected-, pDNA-transfected- and control-THP-1 cells harvested at various time points using the RNeasy® Plus kit with RNase-Free DNase (Qiagen) as per manufacturer protocol. cDNA was synthesized using 1 µg of RNA using the Bio-Rad iScript® kit in a 20 µL reaction volume. Bio-Rad iCycler iQ™ Multicolor Real-Time PCR Detection System was employed to conduct qRT-PCR for quantitation of DENV and V5 copy numbers using Bio-Rad iQ™ SYBR® Green Supermix, 2 µL of 1:3 diluted cDNA, and 10 pmol each of forward and reverse primers (Table 2) in a final reaction volume of 20 µL. Thermal cycling reactions for both DENV and V5 amplifications were initiated with a denaturing step of 4 min at 95 °C, followed by 40 cycles of 95 °C (30 s) and 55 °C (30 s for DENV and 15 s for V5). A standard curve was developed from 10-fold serial dilutions of linear DENV or V5 gene having known concentrations to quantitate the dynamic range of detection of 10¹ to 10⁸ copies per µg of RNA. Host cellular gene changes relative to the glyceraldehyde-3-phosphate dehydrogenase (GAPDH) housekeeping gene were determined as previously described (Verma et al., 2006). Primers used to measure cellular gene changes are listed in Table 2.

Cytokine quantitation

IL-6, IL-8, IP-10, TNFα, VEGF and IFNγ levels were measured in the supernatants of DENV-infected and pDNA-transfected THP-1 cells using a Milliplex human cytokine and chemokine 6-plex immunoassay kit (Millipore Corp., Billerica, MA) together with the Luminex® 100™ System (Luminex, Austin, TX) to determine mean fluorescent intensities (MFI) as recommended by the manufacturer. Protein concentrations were calculated from MFI data using 10-fold serially diluted standards and Bead View analysis software version 1.0.4 (Millipore). The minimum detectable concentrations were 0.4 pg/mL for IL-6, 0.3 pg/mL for IL-8, 1.3 pg/mL for IP-10, 0.2 pg/mL for TNFα, 10.1 pg/mL for VEGF and 0.4 pg/mL for IFNγ.

Western blot

Total cellular protein extracts were prepared from THP-1 cells at 40 h after electroporation with DENV-V5 fusion plasmids. Cells were washed once with cold PBS and extracted with 200 µL of M-PER mammalian protein extraction buffer (Thermo Scientific, Rockford, IL) or NP-40 detergent buffer containing EDTA-free complete protease inhibitor cocktail (Roche, Indianapolis, IN). Either 20 µL or 50 µg of total protein was fractionated on a 4–12% gradient SDS polyacrylamide gel using the Mini-Protean II (Bio-Rad, Hercules, CA) and then transferred onto a 0.2 µm nitrocellulose filter (Bio-Rad Laboratories) as previously described (Verma et al., 2006). Nonspecific binding sites were blocked using 5% FBS in 1× Tris buffered saline with 0.1% Tween and membranes were incubated at 4 °C overnight with primary V5 or β-actin antibodies (dilution 1:1000) followed by incubation with secondary antibodies conjugated to HRP (dilution 1:5000) at room temperature for 1 h. Protein was detected with enhanced chemiluminescence (Amersham ECL, GE Healthcare Limited, Buckinghamshire, UK) using Amersham ECL Hyperfilm (Kodak, Rochester, NY). To determine the relative intensity (RI) of protein bands, the absolute intensity of the DENV-V5 protein band was divided by the absolute intensity of its corresponding β-actin band. Absolute intensities were calculated using Photoshop by multiplying the given pixel value and mean intensity of selected bands as previously described (Luhtala and Parker, 2009).

Cell viability assay

To determine the cell viability, 1 × 10⁶ THP-1 cells were transfected with plasmid DNA or incubated with 1 µg/mL of LPS for 1 h, washed and incubated with fresh growth media until collection at 24 h. Transfected cells were collected at 40 h for measurement of cell viability. Cell viability was measured using the CellTiter96 AQueous One Solution Cell Proliferation Assay (Promega, Madison, WI) kit as per the manufacturers' protocol (Verma et al., 2008).

Statistical analysis

Statistical tests, including paired and unpaired Student's t-tests, were conducted for qRT-PCR and cytokine and chemokine immunoassay using GraphPad InStat version 5.0 (GraphPad Software, San Diego, CA). Values were expressed as mean ± SD of three independent observations. *P* values of <0.05 were considered significant.

Supplementary materials related to this article can be found online at doi:10.1016/j.virol.2011.07.006.

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