Cardol triene inhibits dengue infectivity by targeting kl loops and preventing envelope fusion

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Dengue virus causes a global burden that specific chemotherapy has not been established. A previous report suggested that anacardic acid inhibited hepatitis C virus infection. Here, we explored structure activity relationship of anacardic acid, cardanol, and cardol homologues with anti-DENV cellular infectivities. Cardol triene showed the highest therapeutic index at 29.07 with the CC50 and EC50 of 207.30 ± 5.24 and 7.13 ± 0.72 µM, respectively. Moreover, we observed that the more unsaturated the hydrocarbon tail, the higher the CC50s in all head groups. High CC50s were also found in HepG-2, THP-1, and HEK-293 cell lines where cardol triene CC50s were 140.27 ± 8.44, 129.77 ± 12.08, and 92.80 ± 3.93 µM, respectively. Cardol triene expressed pan-dengue inhibition with the EC50s of 5.35 to 8.89 µM and kl loops of dengue envelope proteins were major targets. The strong binding energy at T48, E49, A50, P53, K128, V130, L135, M196, L198, Q200, W206, L207, I270, and L277 prevented cellular pH-dependent fusion. Zika virus kl loops were aligned in the closed position preventing cardol triene to bind and inhibit fusion and infectivity. This study showed for the first time that cardol triene had a potential for further development as anti-dengue inhibitors.

Dengue infection causes a burden over worldwide tropics with 390 million infections per year1. Of these statistics, Asia represented approximately 75% of the total reports2. Dengue virus belongs to the family Flaviviridae, genus Flavivirus, and consists of four serotype (DENV1-4). The infectious virion contains lipid envelope surrounding an icosahedral nucleocapsid. The nucleocapsid contains an 11 kilobases, single stranded, positive sense RNA. Envelope (E) protein is an immunogen and a major antigenic cross-reactive agent3. All serotypes of dengue viruses are transmitted by Aedes mosquitoes2. Severe clinical manifestations including plasma leakage, hemorrhage, and shock are caused by robust but incompetent immunological responses to secondary heterotypic infection4,5. Remarkably, the antibodies responding to those secondary heterotypic infection are unable to perform complete neutralization and the antibody-tagged viruses are opsonized by macrophage. The viral replication is enhanced inside the cell resulting in increasing the risk to severe dengue development4. Moreover, it was clearly showed that the correlation existed between high viral load, prolonged viremia and clinical severity6–8. Therefore, reducing the viral load should be a direct means to treat acute dengue infection in order to reduce the progression to severe clinical outcomes.

Antiviral drug discovery and development have been substantially advanced within a few decades. Five active leads have entered the clinical trials9–13 but none of them has been launched as a licensed drug yet. Natural products derived from medicinal plants are rich source of potential antiviral inhibitors14. Phenolic lipids are amphiphilic molecules containing phenolic rings and long aliphatic chains. Recently, accumulated evidences suggested that phenolic lipids showed a broad spectrum antimicrobials15, and anti-cancer16 activities by disturbing the phospholipid membrane17 and inducing apoptosis16,18. One of the major sources of phenolic lipids was

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Cashew nut shell liquid (CNSL) existing as mixture of anacardic acid, cardanol, and cardol head groups and saturated (C15:0), monoene (C15:1), diene (C15:2), and triene (C15:3) tails. There is a previous study describing saturated anacardic acid (C15:0) as an inhibitor of hepatitis C virus by interfering with the viral entry, translation, replication, and release19. Dengue virus and hepatitis c virus are likewise members of the family Flaviviridae, therefore it is possible that phenolic lipids should also effectively inhibit dengue and Zika viruses. Since there is limited knowledge of phenolic lipids as antivirals, we aimed to characterize the efficacies of all homologues of CNSL-derived phenolic lipids for dengue virus inhibition. Moreover, the potential candidates would be studied for activities against dengue virus serotype 1–4 and Zika virus; as well as cytotoxicity to human-derived cell lines. The mechanism of drug action and molecular target would also be explored to bring insights into further lead optimization on potential targets.

Results
Preliminary results of phenolic lipids as DENV2 inhibitors. A previous report suggested that anacardic acid mixture extracted from leaves of Gingko biloba inhibited several steps of hepatitis C virus replication19. Hepatitis C virus is one of the members of the family Flaviviridae, therefore, it is likely that anacardic acids, cardanol, and cardol should inhibit dengue virus replication. We extracted phenolic lipids from cashew nut shell liquid (CNSL) and primarily characterized by the three head groups into anacardic acid, cardanol, and cardol (Fig. 1A). Each compound was first screened for druglikeness using two pharmacological characters and four toxicity risks listed in materials and methods. Results showed that trienes from all phenolic heads scored the highest, whereas dienes scored the lowest regarding from the irritant factor (Fig. 1B). Moreover, anacardic acid, cardanol

Figure 1. Chemical structure and computational screening. (A) Phenolic lipid structures consisting of three head groups a) anacardic acid, b) cardol, c) cardanol; and the C15 hydrocarbon tails R1) saturated, R2) monoene, R3) diene, and R4) triene. (B) Computational screening for drug likeness using Molecular Property Explorer and RTECS database.
and cardol mixtures were co-incubated with DENV2 infected Vero cells (M.O.I. of 0.1) at 10 µM and 25 µM for 72 h and the viral inhibition was quantified by plaque titration assay. This preliminary data suggested that DENV2 virions were inhibited at 97.95, 65.30, and 95.91% when incubated with anacardic acid, cardanol, and cardol at 10 µM, respectively (Fig. 2A); whereas the DENV2 virions were potently inhibited (> 99%) by all compounds at 25 µM (Fig. 2A). Moreover, the viability of Vero cells were accessed in the presence of each compound at 10 and 25 µM for 48 and 72 h (Fig. 2B-C) and cytotoxicity results were less than 20 percent in all experimental conditions.

Taken together, our preliminary data showed that anacardic acid, cardanol, and cardol inhibited DENV2 infectivity similar to that of HCV previously reported19.

Cytotoxicity of phenolic lipid homologues.

Next, the compound mixture were further purified and synthesized into saturated (C15:0), monoene (C15:1), diene (C15:2), and triene (C15:3) as described in materials and methods. Toxicities were analyzed and results were calculated to percent viability and CC50 were analyzed using non-linear regression curve (Table 1). Interestingly, anacardic and cardol trienes, had the highest CC50s at 115.13 ± 14.12, and 207.3 ± 5.24 µM, comparing with other homologues from the same head group. As the hydrocarbon tail became more saturated, the CC50 gradually decreased until reaching the lowest value at saturation (C15:0). Cardanol triene was insoluble in DMSO, therefore the compound was excluded from this study. It could be inferred that the polyunsaturated hydrocarbon may involve in stabilizing the cell membrane. Moreover, CC50s of the mixtures tend to follow those of the saturated homologues suggesting that the presence of saturated homologues, regardless of the proportion, could dominate the cell membrane disruption. Moreover, we further explored the CC50 of cardol triene in other cell lines to verify its protective effect. Three human-derived cell lines (THP-1, HEK-293, and HepG2) were used to co-incubate with cardol triene and cardol mixture (73% diene and

Table 1. EC50, CC50, and TI of CNSL-derived phenolic lipids. 1Effective concentration of indicated phenolic lipid to DENV2 NGC (M.O.I. of 0.1) infected Vero cells. 2Cytotoxic concentration of indicated phenolic lipid to Vero cells. 3Therapeutic index (TI, CC50/EC50). *Insoluble. **Data represented means ± standard error of the means (SEM) from three independent experiments, unless otherwise indicated. °Data represented means ± standard error of the means (SEM) from two independent experiments due to limited compound availability.

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Results showed that CC50s of cardol triene were about 30–50 µM higher than cardol mixture in all cell lines (Fig. 3A–C). In conclusion, cardol triene showed a distinctive CC50 value comparing with other homologues towards four distinctive mammalian cell lines.

Efficacies of phenolic lipid homologues. Next, we studied the efficacies (EC50) of the phenolic lipid homologues against DENV2 infectivity in Vero cells. Briefly, compounds at 10 different concentrations were added to DENV2-infected Vero cells (M.O.I. of 0.1) during and post infection and the viral inhibition was determined by plaque titration. Efficacies of anacardic acids varied among homologues so that the structure-activity relationship was inconclusive. The most efficient anacardic acid was the saturated homologue and its result was similar to that of the mixture although only 18% of saturated anacardic acid was found in the total mixture. It is possible that the viral inhibition and cellular response might be predominately influenced by saturated anacardic acid regardless of the percent constituent. Noted that anacardic acids were major components of CNSL extracts.

Figure 3. CC50s and EC50s of cardol triene. (A–C) Non-linear regression curves and calculated cytotoxicity concentrations (CC50s) of cardol triene (bold line) and cardol mixture (dash line) in (A) HepG2, (B) THP-1, (C) HEK-293 cell lines were shown. (D–H) Non-linear regression curves and calculated effective concentrations (EC50s) of cardol triene in Vero cells infected with (D) DENV2 (E) DENV1 (F) DENV3 (G) DENV4 and (H) ZIKV were shown. Graphs represented one of the independent experiments and values indicated means ± standard error of the means (SEM) from three independent experiments.
and all homologues were purified directly from CNSL itself. However, CC_{50} of anacardic acids were generally lower than those of cardols (Table 1), therefore they were not chosen for further characterization.

Naturally derived cardanol mixture consisted of monoene, diene, and triene hydrocarbon species. The saturated cardanol was later synthesized by hydrogenation. Overall, the efficacy of cardanol species were less potent than those of anacardic acids and cardols (Table 1). One of the homologues, cardanol triene, was insoluble in DMSO as previously described, therefore it was excluded from the study. Efficacy of cardanol mixture fell between those of the monoene and diene, suggesting that both homologues contributed to the viral inhibition. Interestingly, hydrogenation of cardanol potentiated the efficacy (EC_{50}) to 5.89 ± 2.83 μM. However, cardanols were the most cytoxic of all head groups, thus resulted in the least TI values. The group were then excluded from further investigations.

Last, cardol mixture and homologues were analyzed for their efficacies using the DENV2 infected cells previously described. Cardol mixture was the most potent DENV2 inhibitor beyond any cardiol homologue with the EC_{50} of 3.24 ± 0.51 μM, and therapeutic index (TI) of 18.63 (Table 1), suggesting a synergistic effect from cardol diene and triene residing in the mixture. Analysis of the homologue showed that increasing the tail unsaturation correlated with increasing the viral inhibition. Cardol triene was the most effective at EC_{50} of 7.13 ± 0.72 μM, and therapeutic index (TI) of 29.07 (Table 1, Fig. 3D). Considering the TI values, the triene homologues showed the broadest spectrum suggesting strong candidates for further investigation. Moreover, the cell morphology under the treatment of DENV2 (M.O.I. of 1 and 5) and cardols were observed at 2, 24, and 48 h after infection and no apparent cytotoxicity were observed in any condition (Supplementary 1). From these results, cardol triene was selected for subsequent studies to verify the drug mechanism.

We further explored whether cardol triene would exhibit broad spectrum inhibition against dengue virus serotype 1–4 and Zika virus. Efficacies against DENV1 (16007), DENV3 (16562), DENV4 (c0036), and ZIKV (sv0010/15) (Fig. 3E–H) suggested that cardol triene similarly inhibited all dengue viruses, but did not inhibit Zika virus. Since dengue and Zika viruses are closely related, their protein structures and their mechanisms of viral replication were highly similar. Therefore, the character of potential targets were supposed to be conserved exclusively within dengue virus, but not in Zika virus. Investigating for the discrepancies between mechanism of dengue and zika virus replication, as well as between the two viral proteins, could result in identification of the molecular targets.

Screening for possible targets of viral inhibition. Time of drug addition assay is a standard method to primarily identify the possible target of a newly characterized compound20. Based on the previous efficacy results, cardol triene was chosen as a representative of all CNSL-derived phenolic lipids. Cardol triene was prepared in DMSO and added to the final concentrations of 10 μM and 20 μM to DENV2 infected Vero cells (M.O.I. of 0.1). The compound was added to the cells at different time-points. DMSO alone was added to the infected cells as a mock treatment. After 72 h incubation, cell lysates and supernatants were collected for analysis of intracellular RNAs and infectious virions by RT-qPCR and plaque titration, respectively. Results showed that intracellular RNAs (Fig. 4A) and infectious virions (Fig. 4B) decreased immediately at 1 h and persisted until 48 h after infections suggesting that the compound globally inhibited the virus at early and late stages of the life cycle. Noted that the EC_{50} of cardol triene was 7.13 ± 0.72 μM; therefore, the viral replication should be suppressed beyond 50 percent by plaque titration under the treatment of cardol triene at both concentrations. From these results, we hypothesized that the phenolic compounds could bind to multiple targets at both early and late stages of the virus life cycle, similar to a previous report on HCV19.

Cardol triene bound to kl loops and inhibited DENV fusion. The early life cycle of DENV mainly involves attachment, entry, and fusion. Due to the fact that cardol triene did not inhibit ZIKV, the close relatives of DENV; it is most likely that the drug targets were present only in DENV proteins and absent in those of ZIKV. Dengue and Zika envelope (E) proteins were first tested using blind docking technique. The cardol triene was preliminarily docked into DENV and ZIKV E dimers in order to detect the differences of ligand binding sites (Fig. 5A). Pairwise structural alignment at β-OG pocket revealed 50% identity between DENV and ZIKV with z-score of 5.5 (Fig. 5A). The β-OG pocket consists of four protein fragments including 48–54, 128–135, 191–209, and 265–282 in DENV, and 48–54, 128–135, 197–215 and 270–288 in ZIKV. The major difference was indicated to open and closed positioning of the kl loop, 265–282 in DENV and 270–288 in ZIKV. Also, blind docking results suggested that cardol triene was able to bind exclusively into dengue β-OG pocket (Fig. 5A). Molecular docking by CDOCKER was also used to analyze other phenolic lipids interacting at the kl loop pockets of DENV E dimer (Fig. 5B). The interaction energy scored between −41.44 to −50.47 kcal/mol and a β-OG crystal ligand was listed at 44.43 kcal/mol.

MD simulation was setup using the initial cardol triene/DENV E complex structure adopted from the best orientation and interaction energy score calculated by CDOCKER. The structural dynamics at 300 ns-trajectories under 297 and 310 Kelvin were analyzed and results showed that the complex at 310 K (3.03 Å) was slightly more fluctuated than that of the 297 K (2.91 Å) (Supplementary 2). The binding free energy (ΔG bind) was calculated from the equilibrium phase which occurred after 40 ns at both conditions. Results were subsequently compared to previously described (FNSY)21 by SIE method (Fig. 5C). The binding free energy of cardol triene to the K and K’ sites were −9.41 ± 0.03, and −8.45 ± 0.03 kcal/mol at 297 K; and −9.29 ± 0.04, and −8.87 ± 0.07 kcal/mol at 310 K, respectively. Interestingly, the energy from cardol triene/DENV E complex were significantly higher than that of the previously reported fusion inhibitor, FNSY at −3.96 ± 0.03 and −4.30 ± 0.03 kcal/mol at K and K’ sites, respectively, with p-value = 0.002621. In addition to SIE, MM-GB/PSBS and QM/MM-GBSA (SCC-DFTB) methods were used to corroborate the binding energy of cardol triene and the kl hydrophobic pockets. The binding energy was achieved from 200 ns-trajectories and delicately computed at K and K’ sites at 297 and 310 K using previously described methods (Fig. 5D). Results showed that cardol triene consistently bound to K and K’ sites...
regardless of the binding free energy calculations or temperature conditions. The major reason that cardol triene contributed to a significantly tight binding was the additional hydrophobic interactions from the hydrocarbon chain and the kl hydrophobic pocket. Major binding residues were revealed and grouped into four regions as follows; 48–54 (T48, E49, A50, P53), 128–135 (K128, V130, L135), 191–207 (M196, L198, Q200, W206, L207), and 270–281 (I270, and L277) (Fig. 5E). Noted that there were multiple hydrophobic interactions at A50, L135, M196, L198, W206, L207, I270 and L277 residues (Fig. 5E). However, the major ligand-binding interaction was contributed by polar and charged residues at T48, E49, K128 and Q200 (Fig. 5E).

Since dengue envelope protein was the key factor to receptor-binding, endocytosis, and fusion, we asked the question whether the compound would mainly neutralize the virion, inhibit the receptor binding, or inhibit endosomal fusion. Cardol triene at 10 µM was then co-incubated with DENV2 before, during, and after infection to Vero cells (Fig. 6A). Results showed that the major inhibition of intracellular RNA and infectious virion was observed at post-infection at 87.00 ± 6.43% and 91.73 ± 4.53% (Fig. 6B), respectively. Therefore, the major target was most likely at a pH-dependent endosomal fusion. Moreover, moderate inhibition was also noticed in infectious virions during pre-incubation and co-infection, suggesting that the compound also moderately neutralized the virion and interfered with the receptor-binding. However, the major post-infection inhibition required further investigations.

Fusion inhibition study of DENV and ZIKV were used to confirm the findings. Briefly, cardol triene at 10 µM was added to DENV2 or ZIKV (Fig. 6C-D) infected C6/36 cells at the M.O.I. of 1 for 48 h before inducing acidic condition by MES. The 4G2 antibody was used as a positive control inhibiting both DENV2 and ZIKV fusion. Results showed that cardol triene exclusively inhibited dengue fusion as seen in the unfused cells and nuclei (Fig. 6C-D). ZIKV-induced fused cells and nuclei were present in cardol triene-treated samples (Fig. 6C-D). Therefore, we concluded that one of the targets of cardol triene, and other CNSL-derived phenolic lipids, was the kl loops of DENV E protein, thus preventing the conformational change of DENV dimer to trimer that was required to initiate pH-dependent fusion.

Cardol triene inhibited late stages of DENV replication. We analyzed the replication inhibition using BHK-21/DENV2 replicon cells stably expressed the non-structural proteins. Cardol triene at 10µM and 20µM were added to the replicon cells for 72h and the replication inhibition was accessed by RT-qPCR of DENV2 NS1. Ribavirin, a known inhibitor of flaviviral replication, was used as a positive inhibition control. We found 76.60 ± 7.59 and 89.44 ± 4.02 percent DENV2 RNA inhibitions from 10µM and 20µM cardol triene, respectively (Fig. 7A), comparable to those of ribavirin. The inhibitory effect at 20µM of cardol triene was closed to ribavirin.
that showed 98.42 ± 0.68 percent and 98.09 ± 0.72 percent at 10 µM and 20 µM, respectively. Replicon cell viability under cardol triene treatment was also analyzed using an MTS assay. Cardol triene treatment at 10 µM and 20 µM for 72 h did not have effect on cell viability (Fig. 7B). Therefore, we concluded that cardol triene inhibited DENV RNA replicon replication, potentially in dose-dependent manner.

Discussion

Phenolic lipids were reported as active ingredients of plants in the Family Anacardiaceae with versatile biological activities. Major components of phenolic lipids in this study were directly isolated from cashew nut shell liquid (CNSL) and synthesized by hydrogenation. The viral inhibition was analyzed and cardol triene was the most effective compound in this group considering from the highest CC50 and TI values. Cardol triene was also similarly effective against the other DENV but no inhibition was found in Zika virus. Moreover, cardol triene was observed with high CC50 values in three human-derived cell lines; HepG2, HEK-293, and THP-1. One of the molecular targets was the kl hydrophobic pockets that would undergo conformational change under acidic pH and performed endosomal fusion to release the viral genome. Our findings showed for the first time that CNSL-derived phenolic lipids had a potential for further development as anti-dengue inhibitors.

Since all homologues showed similar efficacies against DENV2 at the kl loops (Table 1, Fig. 5A), we then hypothesized that all CNSL-derived phenolic lipids shared the same molecular targets. The major target of cardol triene, and potentially to other phenolic lipids, was the kl loop of DENV E protein. This target was one of the hotspots in dengue drug discovery and a variety of ligands were reported as fusion inhibitors. Cardol triene was structurally similar to β-OG, the crystalized ligand, and similar binding moieties were identified. Additional structure-activity relationship (SAR) study should focus on optimizing the phenolic lipids into this binding pocket. A previous report on mannoside glycolipid conjugates suggested that the C24 hydrocarbon tail, either saturated or unsaturated, was the kl loop of DENV E protein. This target was one of the hotspots in dengue drug discovery and a variety of ligands were reported as fusion inhibitors. Cardol triene was structurally similar to β-OG, the crystalized ligand, and similar binding moieties were identified. Additional structure-activity relationship (SAR) study should focus on optimizing the phenolic lipids into this binding pocket. A previous report on mannoside glycolipid conjugates suggested that the C24 hydrocarbon tail, either saturated or unsaturated, was the kl loop of DENV E protein. This target was one of the hotspots in dengue drug discovery and a variety of ligands were reported as fusion inhibitors. Cardol triene was structurally similar to β-OG, the crystalized ligand, and similar binding moieties were identified. Additional structure-activity relationship (SAR) study should focus on optimizing the phenolic lipids into this binding pocket. A previous report on mannoside glycolipid conjugates suggested that the C24 hydrocarbon tail, either saturated or unsaturated, was the kl loop of DENV E protein. This target was one of the hotspots in dengue drug discovery and a variety of ligands were reported as fusion inhibitors. Cardol triene was structurally similar to β-OG, the crystalized ligand, and similar binding moieties were identified. Additional structure-activity relationship (SAR) study should focus on optimizing the phenolic lipids into this binding pocket. A previous report on mannoside glycolipid conjugates suggested that the C24 hydrocarbon tail, either saturated or unsaturated, was the kl loop of DENV E protein. This target was one of the hotspots in dengue drug discovery and a variety of ligands were reported as fusion inhibitors. Cardol triene was structurally similar to β-OG, the crystalized ligand, and similar binding moieties were identified. Additional structure-activity relationship (SAR) study should focus on optimizing the phenolic lipids into this binding pocket. A previous report on mannoside glycolipid conjugates suggested that the C24 hydrocarbon tail, either saturated or unsaturated, was the kl loop of DENV E protein. This target was one of the hotspots in dengue drug discovery and a variety of ligands were reported as fusion inhibitors.
However, cardol triene did not inhibit ZIKV although the virus was closely related to DENV. Based on the findings, we speculated that the similar efficacies of DENV and HCV might be a coincidence of the compound targeting different molecules. Indeed, the viral inhibition (EC_{50}) was a summary of hundreds of viral activities and virus-host interactions.

Cardol triene showed cytoprotective effects towards Vero, HepG2, THP-1, and HEK-293 cell lines with the CC_{50} of 207.3 ± 5.24, 140.27 ± 8.44, 129.77 ± 12.08, 92.8 ± 3.93 µM, respectively (Fig. 3A–C). The cytoprotective effect of cardol triene was also reported in normal human lung fibroblast cell line, GM07492A, with the 192.6 ± 6.0 µM. Cardol monoene, however, was reported with a cytotoxicity against the SW620 (CC_{50} of 14.18 ± 0.76 µM), KATO-III (CC_{50} of 19.06 ± 0.39 µM), HepG2 (CC_{50} of 2.23 ± 0.22 µM), Chago I (CC_{50} of 2.55 ± 0.18 µM) and BT474 (CC_{50} of 13.46 ± 0.14 µM) cell lines. Also, the saturated anacardic acid was cytotoxic against K562 and AGS cell lines with the CC_{50}s of 25.4 µM and 41.6 µM, respectively. Therefore, the saturated and monoene compounds were suitable for further development as anti-cancer chemotherapy. Moreover,
we noticed that the increasing unsaturated hydrocarbon tails positively correlated to the higher CC50 values (Table 1). Previous reports showed that the longer and the more unsaturated hydrocarbon tail related to higher affinity to either disruptive and stabilizing the lipid bilayer. It is possible that the increasing unsaturated hydrocarbon tail would facilitate the compound incorporation into lipid bilayer and stabilizing the bilayer by increasing the membrane fluidity. Moreover, among the three head groups, cardol showed the highest cytoprotection possibly because cardol had a rod-like shape with adequate hydrophobicity for insertion and stabilization into the hydrophobic parts of bilayer. Anacardic acids had a conical shape and more hydrophilic than cardol, therefore it was incorporated into the subsurface and introduced a constraint on lipid packing leading to earlier membrane disruption. We proposed that insertion and stabilization was a strategy of cardol triene to protect the cell membrane, thus tremendously increasing CC50 in all tested cell lines. In addition, a previous report suggested that the incorporation of phenolic lipids into lipid membrane could promote liposome fusion to the cell membrane and could activate the release of drug molecules from the interior of liposomes to the target cells. In our case, cardol triene itself also actively inhibited DENV envelope fusion; therefore, the compound could possibly be delivered into DENV-inhabiting endosome via liposome fusion, which would subsequently bind to kl hydrophobic pockets as previously described.

Our results showed that DENV replicon RNA was inhibited by cardol triene (Fig. 7). The replicon is a self-replicating viral RNA utilized as a screening for RNA replication inhibition. However, the exact molecular target had not been elucidated. Viral replication is driven and modulated by hundreds of viral and host factors including translation machineries, lipid biosynthesis, chaperones, etc. Previous reports showed that saturated anacardic acid inhibited host-derived histone acetyltransferase (HATs), inhibited IL-8 production and HCV replication. However, more evidence is required in order to conclude that dengue virus use the same inhibitory platform. Investigating the compounds with similar constructs can be further employed by similarity search and analysis of their biological activities could lead to the possible molecular target. Note that the inhibition was limited only to DENV so the targets would also be narrowed down to factors involving in DENV replication.

In conclusion, we demonstrated for the first time that phenolic lipids, especially cardol triene, were potential candidates for drug development. The compound inhibited all serotypes of DENV with good efficacy and mild cytotoxicity. One of the molecular targets were identified at kl loops of E protein at the early stage. The structure-activity relationship study of phenolic lipids and the kl loops should be further explored for potentials to be antiviral chemotherapy. Moreover, its selective efficacy towards DENV but not to ZIKV could become a crucial tool for structural and functional comparative studies of these viruses. Since this group of compounds are relatively new in antiviral drug discovery, a lot of knowledge is awaited to be explored and discovered. And we clearly showed in this research that cardol triene has strong potentials to become novel antivirals.

Materials and Methods

Extraction, Purification, and Identification of Anacardic acids, Cardanols, and Cardols. Phenolic lipids were extracted from cashew nut shell liquid (CNSL) (Anacardium occidentale), purified according to a previously reported protocol and characterized by 1H NMR (CDCl3) spectroscopy. The products were analyzed as anacardic acid, cardanol, and cardol, 84.2, and 0.1 percent, respectively. Briefly, 5% aqueous MeOH (300 mL) (Sigma Aldrich®, St. Louis, USA) was used to dissolve CNSL (50 g). Ca(OH)2 (50 g) (Sigma Aldrich®, St. Louis, USA) was gradually added to the reaction mixture while stirring at 50 °C for 5 h. The reaction was monitored by Kieselgel 60 PF254 silica-based thin layer chromatography (TLC) (Merck, New Jersey, USA). After 5 h reaction, dark brown calcium anacardate was precipitated which was then filtered and washed with MeOH before drying under vacuum evaporator. The product was suspended in HCl solution (Sigma Aldrich®, St. Louis, USA) and continuously stirred for 1 h before extraction with EtOAc (Sigma Aldrich®, St. Louis, USA). The organic layer was collected and washed with water, and subsequently dried over anhydrous Na2SO4 (Sigma Aldrich®, St. Louis, USA). Finally, the product was concentrated under reduced pressure to yield anacardic acid (42.1 g, 84% yield). The leftover filtrates from calcium anacardate precipitation was also evaporated under reduced pressure and subsequently extracted using EtOAc. The organic layer was recovered and dried over anhydrous Na2SO4 before continuous stirred for 1 h before extraction with EtOAc (Sigma Aldrich®, St. Louis, USA). Finally, the product was concentrated under reduced pressure to yield anacardic acid (42.1 g, 84% yield). The leftover filtrates from calcium anacardate precipitation was also evaporated under reduced pressure and subsequently extracted using EtOAc. The organic layer was recovered and dried over anhydrous Na2SO4 before
concentration under reduced pressure. The crude product was further purified by silica column with hexane/ EtOAc elution to cardanol (1.1 g, 2% yield) and cardol (319.1 mg, 0.1% yield). Each species was further separated into pure homologues of saturated (C15:0), monoene (C15:1), diene (C15:2), and triene (C15:3) using semi-preparative high performance liquid chromatography. The characteristics and signals of all compounds were shown as follows:

Anacardic acid C15:0 (6-pentadecylsalicylic acid): white solid (18% of total anacardic acids) 1H NMR (CDCl3) δH (ppm): 11.01 (1H, s), 7.36 (1H, t, J = 7.9 Hz), 6.87 (1H, dd, J = 8.3, 1.2 Hz), 6.78 (1H, dd, J = 7.5, 1.2 Hz), 2.98 (2H, t, J = 8.0 Hz), 1.59 (2H, m), 1.26 (19H, m), and 0.88 (3H, t, J = 6.8 Hz).

Anacardic acid C15:1 (6-[8(Z)]-pentadecenyl)salicylic acid): yellow liquid (31% of total anacardic acids) 1H NMR (CDCl3) δH (ppm): 11.04 (1H, s), 7.35 (1H, t, J = 7.9 Hz), 6.86 (1H, d, J = 8.3 Hz), 6.77 (1H, d, J = 7.5 Hz), 5.35 (2H, m, J = 4.8 Hz), 2.97 (2H, t, J = 7.9 Hz), 2.01 (4H, m), 1.60 (2H, m), 1.29 (16H, m), and 0.88 (3H, t, J = 6.5 Hz).

Anacardic acid C15:2 (6-[8(Z), 11(Z)]-pentadecadienyl)salicylic acid): yellow liquid (26% of total anacardic acids) 1H NMR (CDCl3) δH (ppm): 11.07 (1H, s), 7.35 (1H, t, J = 7.9 Hz), 6.86 (1H, d, J = 8.3 Hz), 6.77 (1H, d, J = 7.5 Hz), 5.37 (4H, m), 2.97 (2H, t, J = 7.9 Hz), 2.77 (2H, t, J = 6.3 Hz), 2.01 (4H, m), 1.33 (12H, m), and 0.90 (3H, t, J = 7.4 Hz).

Anacardic acid C15:3 (6-[8(Z), 11(Z), 14-pentadecatrienyl]salicylic acid): yellow liquid (25% of total anacardic acids) 1H NMR (CDCl3) δH (ppm): 11.09 (1H, s), 7.35 (1H, t, J = 7.9 Hz), 6.86 (1H, d, J = 8.3 Hz), 6.76 (1H, d, J = 7.5 Hz), 5.81 (1H, m), 5.39 (4H, m), 5.01 (2H, m), 2.97 (3H, t, J = 7.9 Hz), 2.79 (4H, dd, J = 13.7, 7.5 Hz), 2.05 (2H, m), 1.59 (2H, m), and 1.34 (8H, m).

Cardanol C15:1 (3-[8(Z)]-pentadecenyl)phenol): yellow liquid (31% of total cardanols) 1H NMR (CDCl3) δH (ppm): 7.14 (1H, t, J = 7.7 Hz), 6.75 (1H, d, J = 7.6 Hz), 6.65 (2H, d, J = 8.4 Hz), 5.35 (2H, m), 2.55 (2H, t, J = 7.8 Hz), 2.01 (4H, m), 1.59 (2H, m), 1.29 (16H, m), and 0.89 (3H, t, J = 6.5 Hz).

Cardanol C15:2 (3-[8(Z), 11(Z)]-pentadecadienyl)phenol): yellow liquid (46% of total cardanols) 1H NMR (CDCl3) δH (ppm): 7.14 (1H, t, J = 7.7 Hz), 6.75 (1H, d, J = 7.5 Hz), 6.65 (2H, d, J = 8.1 Hz), 5.36 (4H, m), 2.78 (2H, t, J = 6.3 Hz), 2.55 (2H, t, J = 7.8 Hz), 2.04 (4H, m), 1.58 (12H, m), and 0.91 (3H, t, J = 7.4 Hz).

Cardanol C15:3 (3-[8(Z), 11(Z), 14-pentadecatrienyl]phenol): yellow liquid (23% of total cardanols) 1H NMR (CDCl3) δH (ppm): 7.14 (1H, t, J = 7.6 Hz), 6.76 (1H, d, J = 7.6 Hz), 6.65 (2H, d, J = 7.9 Hz), 5.83 (1H, m), 5.40 (4H, m), 5.03 (2H, m), 2.82 (4H, dt, J = 16.1, 6.1 Hz), 2.56 (2H, t, J = 7.7 Hz), 2.04 (2H, m), 1.60 (2H, m), and 1.29 (8H, m).

Cardol C15:1 (5-[8(Z), 11(Z)-pentadecenyl]resorcinol): brown liquid (73% of total cardols) 1H NMR (CDCl3) δH (ppm): 6.24 (2H, s), 6.17 (1H, s), 5.36 (4H, m), J = 8.1, 4.8 Hz), 2.78 (2H, t, J = 6.3 Hz), 2.48 (2H, t, J = 7.8 Hz), 2.04 (4H, m), 1.56 (2H, m), 1.33 (8H, m), and 0.91 (3H, t, J = 7.4 Hz).

Cardol C15:2 (5-[8(Z), 11(Z)-pentadecadienyl]resorcinol): brown liquid (27% of total cardanols) 1H NMR (CDCl3) δH (ppm): 6.25 (2H, s), 6.17 (1H, s), 5.82 (1H, m), 5.39 (4H, m), 5.02 (2H, m), 2.80 (4H, dt, J = 14.1, 7.9 Hz), 2.46 (2H, t, J = 7.7 Hz), 2.04 (2H, m), 1.35 (2H, m), and 1.28 (8H, m).

In addition, the saturated homologues (C15:0) of cardanol and cardol were synthesized from hydrogenation reaction and their signals were shown as follows:

Cardanol C15:0: 1H NMR (CDCl3) δH (ppm): 7.15 (1H, t, J = 7.7 Hz), 6.77 (1H, d, J = 7.6 Hz), 6.68 (2H, d, J = 8.4 Hz), 5.91 (1H, s), 2.56 (2H, t, J = 7.9 Hz), 1.60 (2H, t, J = 6.6 Hz), 1.30 (24H, m), 0.91 (3H, t, J = 6.6 Hz). Cardol C15:0: 1H NMR (CDCl3) δH (ppm): 6.24 (2H, s), 6.18 (1H, s), 2.47 (2H, t, J = 7.8 Hz), 1.55 (2H, m), 1.25 (24H, m), 0.87 (3H, t, J = 6.6 Hz).

All compounds were stored as solid powder at room temperature. The compounds were prepared in dimethyl sulfoxide (DMSO) (Sigma Aldrich®, St. Louis, USA) to 50 mM stock solutions and stored as aliquots at −20°C until use.

Cells and viruses. Vero (ATCC® CCL-81), LLC/MK2 (ATCC® CCL-7), C6/36 (ATCC® CRL-1660), HEK-293 (ATCC® CRL-1573), HepG2 (ATCC® HB-8065), and THP-1 (ATCC® TIB-202) cells were maintained as previously described42. Also, DENV1 (16007), DENV2 (New Guinea C strain), DENV3 (16562), DENV4 (c0036), and Zika virus (SV0010/15) reference strains were propagated in Vero and C6/36 cells as previously described42.

Efficacy studies. The efficacies of anacardic acid, cardol, and cardanol mixtures were primarily studied using DENV2 infected Vero cells. Vero (5 × 104) cells were seeded into each well of 24-well plates. After an overnight incubation, cells were infected with DENV2 (M.O.I. of 0.1) and 10μM and 25μM phenolic lipids. DMSO at 1% was used as a mock treatment with 100% cells viability. Cell viability was assessed at designated time-points using CellTiter 96Aqueous One Solution Cell Proliferation Assay kit (Promega, Wisconsin, USA)
and analyzed by spectrophotometer ($A_{\text{260\text{nm}}}$) (VICTORTM X3, PerkinElmer, Massachusetts, USA). Results were calculated to percent cell death.

Cytotoxic concentrations (CC$_{50}$) were also studied using Vero cells. Briefly, the 10$^4$ cells were seeded into each well of 96-well plates. After an overnight incubation, phenolic lipids were serially diluted in DMSO to the final concentrations as follows: 25, 50, 100, 200, 300, 400, and 500 $\mu$M. Each concentration was performed in quadruplicates. DMSO at 1% was used as a mock treatment referring to 100% cell viability. Cells were incubated for 48 h after cell viability was accessed using CellTiter 96® Aqueous One Solution Cell Proliferation Assay kit (Promega, Wisconsin, USA). Non-linear regression analysis was used to determine cytotoxic concentrations (CC$_{50}$) of each experiment. Results were derived from three independent experiments and reported as means and standard error of the means (SEM).

**Time of drug addition study (TOA).** Vero (5 $\times$ 10$^4$) cells were seeded into 24-well plates as previously described. DENV2 (M.O.I. of 0.1) was diluted in ice-cold MEM with 1% fetal bovine serum and was allowed to adsorb on the cells for 1 h with continuous rocking. Cardol triene (10 $\mu$M) was prepared in DMSO and was mixed with DENV2 (M.O.I. of 1) before adsorption for 1 h (pre-attachment). Cardol triene (10 $\mu$M) was also mixed with DENV2 (M.O.I. of 1) and immediately applied to the cells (co-attachment). Last, cardol triene was added to the cells after 1 h DENV2 adsorption (post-attachment). The experiments were incubated for 2 days at 37°C. Intracellular viral RNA was determined from cell lysate using RT-qPCR and infectious virion was determined using plaque titration of the supernatant. DMSO was added to the experiment as a no-inhibition control. Means and standard error of the means (SEM) of three independent experiments were noted.

**Attachment inhibition study.** C6/36 (2 $\times$ 10$^4$) cells were seeded into 24-well plates before infection with DENV2 and ZIKV (M.O.I. of 1) as described. Cardol triene at 10 $\mu$M was introduced to the DENV2 and ZIKV infected cells during and after infection. DMSO was used as a 0% inhibition control and mouse anti-Flavivirus envelope protein antibody (4G2) was used as a 100% inhibition control. Cells were incubated at 28°C for 2 days before addition of 0.5 M 2-(N-morpholino) ethanesulfonic acid (MES) (pH 5.0) (Sigma Aldrich, St Louis, USA). Moreover, cells from selected wells subsequently underwent DAPI staining. Cell–cell fusion was observed under brightfield and fluorescence using an Eclipse TS100 Inverted Routine Microscope (Nikon, New York, USA).

**Fusion inhibition study.** BHK-21 (5 $\times$ 10$^4$) cells expressing DENV-2 replicon (BHK-21/DENV2) were seeded as previously described. Cardol triene at 10 $\mu$M and 20 $\mu$M, DMSO, and ribavirin (TargetMol, USA) were added to the replicon cells and incubated at 37°C for 3 days. Cells lysates were collected and viral RNA was quantified by RT-qPCR as previously described. Data were reported as percent inhibition compared with ribavirin. Three biological replicates were performed and results were means and SEM of all replicates.

**Replicon inhibition study.** BHK-21 (5 $\times$ 10$^4$) cells were seeded into 24-well plates before infection with DENV2 and ZIKV (M.O.I. of 1) as described. Cardol triene at 10 $\mu$M and 20 $\mu$M, DMSO, and ribavirin (TargetMol, USA) were added to the replicon cells and incubated at 37°C for 3 days. Cells lysates were collected and viral RNA was quantified by RT-qPCR as previously described. Data were reported as percent inhibition compared with ribavirin. Three biological replicates were performed and results were means and SEM of all replicates.

**Computational study.** Phenolic lipids were evaluated for druglikeness by scoring their pharmacological characters (cLogP and solubility) and toxicity risks (mutagenicity, tumorigenicity, irritating effect, and reproductive effect) using Molecular Property Explorer and RTECS database.

To refine the screening result and construct the initial complex structure, the 3D DENV envelope protein (PDB code: 1OKE) was obtained to calculate the interaction energy with phenolic lipids using CDOCKER software (Discovery Studio 2.5, Accelrys Inc., San Diego, USA). Each kl loop was assigned to 10 Å diameter according to n-octyl-beta-D-glucoside originally bound in the crystal structure. All phenolic lipids were calculated for partial charge distribution and docked into kl loop with 100 replicas with random conformation of ligand. The best score predicted by CDOCKER interaction energy was selected to construct an initial complex structure for molecular dynamics (MD) simulation using 3 different methods; Solvated Interaction Energy (SIE), Molecular Mechanics-generalized Born/Poisson Boltzmann solvent accessible surface area (MM-GB/PBSA), and Quantum mechanics/molecular mechanics generalized Born surface area (QM/MM-GBSA) with density functional theory-based tight binding (SCC-DFTB) methods.

The MD simulations were conducted for cardol triene/DENV E protein complex with two temperature systems; 297 K (23.85 °C) and 310 K (36.85 °C) implemented with the GPU accelerated pmemd.cuda program in the AMBER16 package program. Both 297 K and 310 K systems were simulated with explicit solvent TIP3P water molecule with 15 Å around protein surface. Periodic boundary condition was applied in this MD simulation. The protonation state of the ligand, cardol triene, was assigned by pKa calculation in MarvinSketch program and 2 atoms of Na$^+$ were added to neutralize the electrical charge of DENV E dimer using PDBePQR. The 1,500 steps of steepest descents (SD) followed by 1,500 steps of conjugated gradients (CG) in the SANDER module by AMBER 16 were used to minimize the interference of water molecules and hydrogen atoms. The systems were subsequently brought up to 297 K and 310 K within 200 ps and was stabilized at the designated temperature until 300 ns. The MD trajectories were collected every 0.2 ps during 200–300 ns production phase. The stability of the cardol triene/DENV E protein complex at equilibrium was analyzed for the global root-mean-square displacement (RMSD) of binding free energy using CPPTRAJ module. Snapshots, taken every 1 ns within 200–300
ns-MD trajectories, were used for calculating total binding free energy using SIE, MM/GB/PBSA, and QM/MM-GBSA (SCC-DFTB) methods. Snapshots were also used to identify the per-residue decomposition energy contribution by MM/GBSA method.

**Ethical Approval.** This project was approved by Ethical Review board, Faculty of Medicine, Chulalongkorn University (COE 021/2017 and IRB 375/60) as exempted in compliance with the International guidelines for human research protection as Declaration of Helsinki, The Belmont Report, CIOMS guideline, International Conference on Harmonization in Good Clinical Practice (ICH-GCP) and 45CFR 46.101(b).

**The level of bio-containment.** We performed all pathogen-related experiments in bio-containment level 2. BSL-2 standard operating procedure was performed under the regulation of ISO15189 and ISO15190.

**Data Availability Statement** All data supporting the findings can be found in the results and supplementary sections.

**References**

1. Bhatt, S. et al. The global distribution and burden of dengue. *Nature* **496**, 504–507, nature12060.
2. Murray, N. E., Quann, M. B. & Wilder-Smith, A. Epidemiology of dengue: past, present and future prospects. *Clinical epidemiology* **5**, 299–309, https://doi.org/10.2147/clep.s34440 (2013).
3. Marimuthu, P. & Ravinder, J. R. Trends in clinical trials of dengue vaccine. *Perspectives in clinical research* **7**, 161–164, https://doi.org/10.4103/2229-3485.192035 (2016).
4. Gubler, D. J. & Clark, G. G. Dengue/dengue hemorrhagic fever: the emergence of a global health problem. *Emerg Infect Dis* **1**, 55–57, https://doi.org/10.3201/eid0102.950204 (1995).
5. Guzman, M. G., Alvarez, M. & Halstead, S. B. Secondary infection as a risk factor for dengue hemorrhagic fever/dengue shock syndrome: an historical perspective and role of antibody-dependent enhancement of infection. *Archives of virology* **158**, 1445–1459, https://doi.org/10.1007/s00705-013-1645-3 (2013).
6. Solbrig, M. V. & Perng, G. C. Current neurological observations and complications of dengue virus infection. *Current neurology and neuroscience reports* **15**, 29, https://doi.org/10.1007/s11910-015-0550-4 (2015).
7. Wang, W. K. et al. High levels of plasma dengue viral load during defervescence in patients with dengue hemorrhagic fever: implications for pathogenesis. *Virology* **305**, 330–338 (2003).
8. Wang, W. K. et al. Slower rates of clearance of viral load and virus-containing immune complexes in patients with dengue hemorrhagic fever. *Clinical infectious diseases: an official publication of the Infectious Diseases Society of America* **43**, 1023–1030, https://doi.org/10.1086/507635 (2006).
9. Tricou, V. et al. A randomized controlled trial of chloroquine for the treatment of dengue in Vietnamese adults. *PLoS neglected tropical diseases* **4**, e785, https://doi.org/10.1371/journal.pntd.0000785 (2010).
10. Tam, D. T. et al. Effects of short-course oral corticosteroid therapy in early dengue infection in Vietnamese patients: a randomized, placebo-controlled trial. *Clinical infectious diseases: an official publication of the Infectious Diseases Society of America* **55**, 1216–1224, https://doi.org/10.1093/cid/ciu655 (2012).
11. Nguyen, N. M. et al. A randomized, double-blind placebo controlled trial of halapiravir, a polymerase inhibitor, in adult dengue patients. *The Journal of infectious diseases* **207**, 1442–1450, https://doi.org/10.1093/infdis/jiy470 (2013).
12. Low, J. G. et al. Efficacy and safety of celgosivir in patients with dengue fever (CELADEN): a phase Ib, randomised, double-blind, placebo-controlled, proof-of-concept trial. *The Lancet. Infectious diseases* **14**, 706–715, https://doi.org/10.1016/s1473-3099(14)70730-3 (2014).
13. Whitehorn, J. et al. Lovastatin for adult patients with dengue: protocol for a randomised controlled trial. *Trials* **13**, 203, https://doi.org/10.1186/1745-6215-13-203 (2012).
14. Teixeira, R. R. et al. Natural products as source of potential dengue antivirals. *Molecules (Basel, Switzerland)* **19**, 8151–8176, https://doi.org/10.3390/molecules19068151 (2014).
15. Hamad, F. B. & Mubofu, E. B. Potential Biological Applications of Bio-Based Anacardic Acids and Their Derivatives. *International Journal of Molecular Sciences* **16**, 8569–8590, https://doi.org/10.3390/ijms16088569 (2015).
16. Kustiawan, P. M. et al. Molecular mechanism of cardol, isolated from Trigona incisa stingless bee propolis, induced apoptosis in the SW620 colorectal cancer cell line. *BMC pharmacology & toxicology* **18**, 32, https://doi.org/10.1186/s12895-017-0139-4 (2017).
17. Stasiuk, M. & Kozubek, A. Membrane perturbing properties of natural phenolic and resorcinolic lipids. *FEBS letters* **582**, 3607–3613, https://doi.org/10.1002/jebs.2008.09.039 (2008).
18. Kustiawan, P. M. et al. Propolis from the Stingless Bee Trigona incisa from East Kalimantan, Indonesia. *Induces In Vitro Cytotoxicity and Apoptosis in Cancer Cell lines, Asian Pacific journal of cancer prevention: APJCP* **16**, 6581–6589 (2015).
19. Hundt, J., Li, Z. & Liu, Q. The inhibitory effects of anacardic acid on hepatitis C virus life cycle. *PloS one* **10**, e0117514, https://doi.org/10.1371/journal.pone.0117514 (2015).
20. Daelmans, D., Pauwels, R., De Clercq, E. & Panneconque, C. A time-of-drug addition approach to target identification of antiviral compounds. *Nature protocols* **6**, 925–933, https://doi.org/10.1038/nprot.2011.330 (2011).
21. Srivrangkul, P. et al. A novel flavanone derivative inhibits dengue virus fusion and infectivity. *Antiviral research* **151**, 27–38, https://doi.org/10.1016/j.antiviral.2018.01.010 (2018).
22. Wang, Y. G. et al. A small molecule dengue virus entry inhibitor. *Antimicrob Agents Chemother* **53**, 1823–1831, AAC.01148-08.
23. Tambunan, U. S., Zahroh, H., Parikesit, A. A., Idrus, S. & Kerami, D. Screening Analogs of beta-OG Pocket Binder as Fusion Inhibitor of Dengue Virus 2. *Drug target insights* **9**, 33–49, https://doi.org/10.4137/dti.s31566 (2015).
24. Jaday, S. S. et al. Design, synthesis and antiviral activity of a class of hybrid dengue virus E protein inhibitors. *Bioorganic & Medicinal Chemistry Letters* **25**, 1747–1752, https://doi.org/10.1016/j.bmcl.2015.02.059 (2015).
25. Ichiyama, K. et al. Sulfated polysaccharide, curdlan sulfate, efficiently prevents entry/fusion and restricts antibody-dependent enhancement of dengue virus infection in vitro: a possible candidate for clinical application. *PLoS neglected tropical diseases* **7**, e2188, https://doi.org/10.1371/journal.pntd.0002188 (2013).
26. Poh, M. K. et al. A small molecule fusion inhibitor of dengue virus. *Antiviral research* **84**, 260–266, S0166-3542(09)00482-3.
27. Randolph, V. B. & Stollar, V. Low pH-induced cell fusion in flavivirus-infected Aedes albopictus cell cultures. *J Gen Virol* **71(Pt 8)**, 1845–1850 (1990).
28. Summers, P. L., Cohen, W. H., Ruiz, M. M., Hase, T. & Eckels, K. H. Flaviviruses can mediate fusion from without in Aedes albopictus mosquito cell cultures. *Virus research* **12**, 383–392 (1989).
29. Rattanaburee, T. et al. Inhibition of dengue virus production and cytokine/chemokine expression by ribavirin and compound A. *Antiviral research* **124**, 83–92, https://doi.org/10.1016/j.antiviral.2015.10.005 (2015).
30. Przeworska, E., Gubernator, J. & Kozubek, A. Formation of liposomes by resorcinolic lipids, single-chain phenolic amphiphiles from Anacardium occidentale L. *Biochimica et biophysica acta* **1513**, 75–81 (2001).
Author Contributions

S.B., W.C. and T.R. conceived and designed the study. P.K. and T.S. equally performed most experiments. A.S. technically assisted most cell-based experiments and performed Fig. 7. K.H. and T.R. provided computational assistance and performed Figs 1B, 5, and Supplementary 2. W.C. purified, synthesized, and identified all compounds by NMR. S.B., P.K. and T.S. analyzed the data. S.B. wrote the original manuscript. All authors reviewed and edited the manuscript.

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Kustiawan, P. M., Puthong, S., Arung, E. T. & Chanchao, C. In vitro cytotoxicity of Indonesian stingless bee products against human cancer cell lines. Asian Pacific Journal of tropical biomedicine 4, 549–556, https://doi.org/10.2992/aphit.20141014-pbth-2013-0039 (2014).

Alam-Escamilla, D., Estrada-Muniz, E., Solis-Villegas, E., Elizondo, G. & Vega, L. Genotoxic and cytostatic effects of 6-pentadecyl salicylic anacardic acid in transformed cell lines and peripheral blood mononuclear cells. Mutation research. Genetic toxicology and environmental mutagenesis 777, 43–53, https://doi.org/10.1016/j.mrgentox.2014.11.008 (2015).

Stasiuk, M. & Kozubek, A. Biological activity of phenolic lipids. Cellular and molecular life sciences: CMLS 67, 841–860, https://doi.org/10.1007/s00018-009-0193-1 (2010).

Siwko, M. E., de Vries, A. H., Mark, A. E., Marrink, S. J. Disturb or stabilize? A molecular dynamics study of the effects of resorcinolic lipids on phospholipid bilayers. Biophysical journal 96, 3140–3153 (2009).

Cieslik-Boczula, K. & Koll, A. The effect of 3-pentadecylphenol on DPPC bilayers ATR-IR and 31P NMR studies. Biophysical chemistry 140, 51–56, https://doi.org/10.1016/j.bpc.2008.11.009 (2009).

Yasutake, T. et al. Anacardic acid, a histone transferase inhibitor, modulates LPS-induced IL-8 expression in a human alveolar epithelial cell line A549. F1000research 2, 78, https://doi.org/10.12688/f1000research.2-78.v1 (2013).

Masuoka, N., Tamsampaoloet, K., Chavasiri, W. & Kubo, I. Superoxide anion scavenging activity of alk(en)yl phenol compounds by using PMS-NADH system. Heliyon 2, https://doi.org/10.1016/j.heliyon.2016.e00169 (2016).

Suroengrit, A. et al. Halogenated Chrysins Inhibit Dengue and Zika Virus Infection. Scientific reports 7, 13696, https://doi.org/10.1038/s41598-018-04171-z (2017).

Boonyasuppayakorn, S. et al. Simplified dengue virus microwell plaque assay using an automated quantification program. Journal of virological methods 237, 25–31, https://doi.org/10.1016/j.viromet.2016.08.012 (2017).

Boonyasuppayakorn, S., Reichert, E. D., Manzano, M., Nagarajan, K. & Padmanabhan, R. Amodiaquine, an antimalarial drug, inhibits dengue virus type 2 replication and infectivity. Antiviral research 106, 125–134, https://doi.org/10.1016/j.antiviral.2014.03.014 (2014).

Lani, R. et al. Antiviral activity of salmeyrin against chukungunya virus. Scientific reports 5, 11421, https://doi.org/10.1038/srep11421 (2015).

Pollaristi, M. P. Overview on the Rule of Five. Current Protocols in Pharmacology 49, 12.9:11–12.9:18, https://doi.org/10.1002/0471141755.ph0912s49 (2010).

Modis, Y., Ogata, S., Clements, D. & Harrison, S. C. A ligand-binding pocket in the dengue virus envelope glycoprotein. Proceedings of the National Academy of Sciences of the United States of America 100, 6986–6991, https://doi.org/10.1073/pnas.0308213100 (2003).

Wu, G., Robertson, D. H., Brooks, C. L. & Vieth, M. Detailed analysis of grid-based molecular docking: A case study of COX-2—A CHARMM-based MD docking algorithm. Journal of Computational Chemistry 24, 1549–1562, https://doi.org/10.1002/jcc.10306 (2003).

Zhang, X. et al. Dengue structure differs at the temperatures of its human and mosquito hosts. Proceedings of the National Academy of Sciences of the United States of America 110, 6795–6799, https://doi.org/10.1073/pnas.1304300110 (2013).

Sion Campos, J. L. et al. Temperature-dependent folding allows stable dimerization of secretory and virus-associated E proteins of Dengue and Zika viruses in mammalian cells. Scientific reports 7, 966, https://doi.org/10.1038/s41598-017-01097-5 (2017).

Dixon, S. L. & Jurs, P. C. Estimation of pKa for organic oxacids using calculated atomic charges. Journal of Computational Chemistry 14, 1460–1467, https://doi.org/10.1002/jcc.14014208 (1993).

Csizmadia, F., Tsantili-Kakoulidou, A., Panderi, I. & Darvas, F. Prediction of distribution coefficient from structure. 1. Estimation method. Journal of Pharmaceutical Sciences 86, 865–871, https://doi.org/10.1021/ps960177b (2000).

Dolinsky, T. J., Nielsen, J. E., McCammon, J. A. & Baker, N. A. PDB2PQR: an automated pipeline for the setup of Poisson–Boltzmann electrostatics calculations. Nucleic Acids Research 32, W656–W667, https://doi.org/10.1093/nar/gkh381 (2004).

Mepreesrat, A. et al. Binding pattern of the long acting neuraminidase inhibitor laninamivir towards influenza A subtypes HSNI and pandemic H1N1. Journal of Molecular Graphics and Modelling 38, 148–154, https://doi.org/10.1016/j.jmgm.2012.06.007 (2012).

Nutho, B. et al. Binding mode and free energy prediction of fisetin/β-cyclodextrin inclusion complexes. Beilstein Journal of Organic Chemistry 10, 2789–2799, https://doi.org/10.3762/bjoc.10.296 (2014).

Sangphrek, W., Khuntawee, W., Wolchmann, P., Pongsawadi, P. & Rungrotmongkol, T. Enhanced stability of a naringin/β,6-dimethylβ-cyclodextrin inclusion complex: Molecular dynamics and free energy calculations based on MM- and QM-PBSA/GBSA. Journal of Molecular Graphics and Modelling 50, 10–15, https://doi.org/10.1016/j.jmgm.2014.03.001 (2014).

Nain, M. et al. Solved Interaction Energy (SIE) for Scoring Protein—Ligand Binding Affinities. 1. Exploring the Parameter Space. Journal of Chemical Information and Modeling 47, 122–133, https://doi.org/10.1021/ci040460v (2007).

Sulea, T. & Purismia, E. O. In Computational Drug Discovery and Design (ed Riccardo Baroni) 295–303 (Springer New York, 2012).

Kollman, P. A. et al. Calculating Structures and Free Energies of Complex Molecules: Combining Molecular Mechanics and Continuum Models. Accounts of Chemical Research 33, 889–897, https://doi.org/10.1021/ar000033 (2000).
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