

Synthesis of Skeletal Analogues of Saxitoxin Derivatives and Evaluation of Their Inhibitory Activity on Sodium Ion Channels Na_v1.4 and Na_v1.5**

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Abstract: Skeletal analogues of saxitoxin (STX) that possess a fused-type tricyclic ring system, designated FD-STX, were synthesized as candidate sodium ion channel modulators. Three kinds of FD-STX derivatives **4a–c** with different substitution at C13 were synthesized, and their inhibitory activity on sodium ion channels was examined by means of cell-based assay. (–)-FD-

STX (**4a**) and (–)-FD-dcSTX (**4b**), which showed moderate inhibitory activity, were further evaluated by the use of the patch-clamp method in cells

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that expressed Na_v1.4 (a tetrodotoxin-sensitive sodium channel subtype) and Na_v1.5 (a tetrodotoxin-resistant sodium channel subtype). These compounds showed moderate inhibitory activity towards Na_v1.4, and weaker inhibitory activity towards Na_v1.5. Uniquely, however, the inhibition of Na_v1.5 by (–)-FD-dcSTX (**4b**) was “irreversible”.

Introduction

Voltage-gated sodium channels (Na_vCh) are transmembrane proteins that provide inward current carried by sodium ions, and they contribute to the control of membrane excitability, as well as the propagation of action potentials along axons.^[1] Sodium channels within neurons are composed of a single α subunit, which forms the voltage-sensing and ion-selective pore, and one or more auxiliary β subunits, which are proposed to serve a number of functions, including modulation of α -subunit function and targeting/anchoring the channels at specific sites in the plasma membrane.^[2] To date, nine

genes that encode α subunits (Na_v1.1–Na_v1.9) and four genes that encode β subunits have been identified. The α subunits can be classified into two groups on the basis of their sensitivity to tetrodotoxin (TTX), the pufferfish toxin. There are six TTX-sensitive (TTX-s) α subunits (i.e., Na_v1.1–Na_v1.4, Na_v1.6, and Na_v1.7) and three TTX-resistant (TTX-r) α subunits (i.e., Na_v1.5, Na_v1.8, and Na_v1.9), which are only blocked by high concentrations of TTX.^[3,4] Recently, a related protein (Na_xX) has been recognized as a tenth member of the group.^[5] These subtypes have quite similar structures, but the expression of the α subunits is strongly cell-type- and tissue-specific. Therefore, each of these subtypes is believed to have unique properties. Indeed, mutation studies have thrown some light on the functions of these channels. For example, mutation of Na_v1.4 is associated with primary periodic paralysis,^[6] whereas mutation of Na_v1.5 leads to cardiovascular syndromes.^[7] Consequently, these channels are considered to be good candidates for new drug targets. In this context, Na_vCh-subtype-selective small-molecular modulators from natural sources have been explored, as well as synthetic compounds. Synthetic studies have led to some subtype-selective compounds.^[8] For example, A-803467 is an Na_v1.8-selective blocker developed by an Abbott research group^[8c,9] for possible application in pain control. However, more selective modulators with a variety of action mechanisms are still of great interest.

Saxitoxin (STX) (**1a**) is a naturally occurring shellfish neurotoxin that blocks ion influx to TTX-s Na_vChs.^[10] STX (**1a**) is an indispensable tool for studying Na_vCh-related electrophysiology, and its derivatives are also promising candidates for Na_vCh-subtype-selective modulators. STX (**1a**) and its analogues are believed to bind to the P-loop region of the ion-selective filter in Na_vCh in a similar manner to

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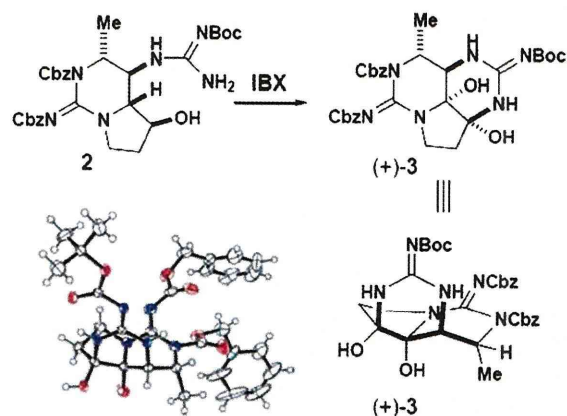
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[**] Na_vCh denotes voltage-gated sodium channels. Na_v1.4 is a tetrodotoxin-sensitive sodium channel subtype; Na_v1.5 is a tetrodotoxin-resistant sodium channel subtype.

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TTX (i.e., in a reversible manner).^[11] Recently, a binding model of STX with the P-loop domain was proposed based on molecular docking studies, and it was suggested that the two guanidinium groups, the carbamoyl group at C13 and the hydrated ketone at C12, have critical roles in the binding.^[12] On the other hand, most naturally occurring STX analogues isolated to date are functionalized at C11, C12, C13, N1, and N7, and these modifications have a significant influence on the Na_vCh-inhibitory activity.^[13] Thus, structure–activity relationship (SAR) studies of STX have been focused on those positions,^[14] especially C13, with the aim of developing Na_vCh-subtype-selective modulators.

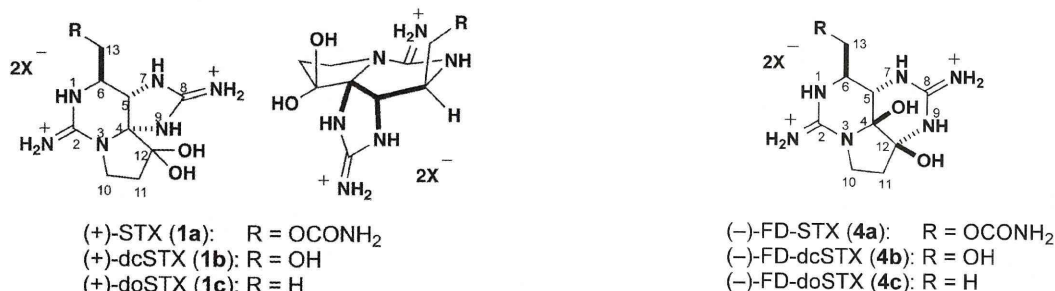
We have recently reported a total synthesis of STX (**1a**; see Scheme 1) and some of its naturally occurring derivatives.^[15] In the course of our synthetic studies of these compounds, we unexpectedly obtained a fused-type tricyclic compound (+)-**3** during the oxidation of **2** with 2-iodoxybenzoic acid (IBX; Scheme 2).^[15a] An X-ray analysis showed compound **3** to have a conical shape similar to that of STX (see Scheme 1), and compound **3** was expected to be able to access the channel conical vestibule in a similar manner to STX. Thus, this fused-type tricyclic skeleton can be regarded as a new skeletal analogue of STX, and we named it FD-STX. The FD-STX structure has almost the same functional groups as STX at positions that interact with Na_vCh, although the angles and/or directions of functional groups are slightly different from those of STX.^[16] We considered that these small structural differences of FD-STX compared with STX might represent an opportunity to acquire new subtype



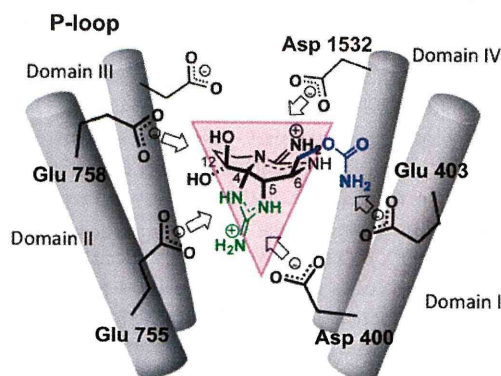
Scheme 2. Synthesis of fused-type tricyclic compound (+)-**3**.

selectivity and/or might result in a distinctive interaction mode of FD-STX with Na_vChs.

In this report, we describe the synthesis of FD-STX derivatives (–)-**4a–c** functionalized at C13, which correspond to the skeletal analogues of (+)-STX (**1a**), (+)-dcSTX (**1b**), and (+)-doSTX (**1c**), respectively (Scheme 3). The Na_vCh-inhibitory activity of these FD-STX derivatives was initially examined by cell-based assay, and (–)-FD-STX (**4a**) and (–)-FD-dcSTX (**4b**) were further evaluated by means of the patch-clamp method in cells that express Na_v1.4 or Na_v1.5. The results are of particular interest because no SAR studies that focus on the STX skeleton itself have been reported so far.



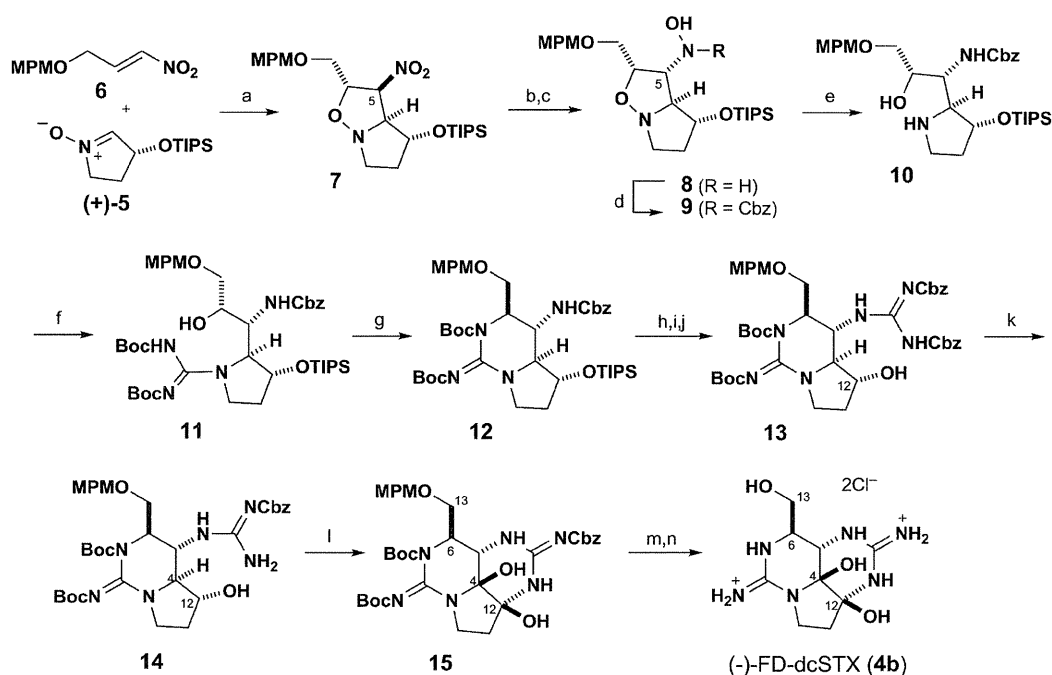
Scheme 3. Structures of (–)-FD-STXs **4a–c**, skeletal analogues of (+)-STXs **1a–c**.



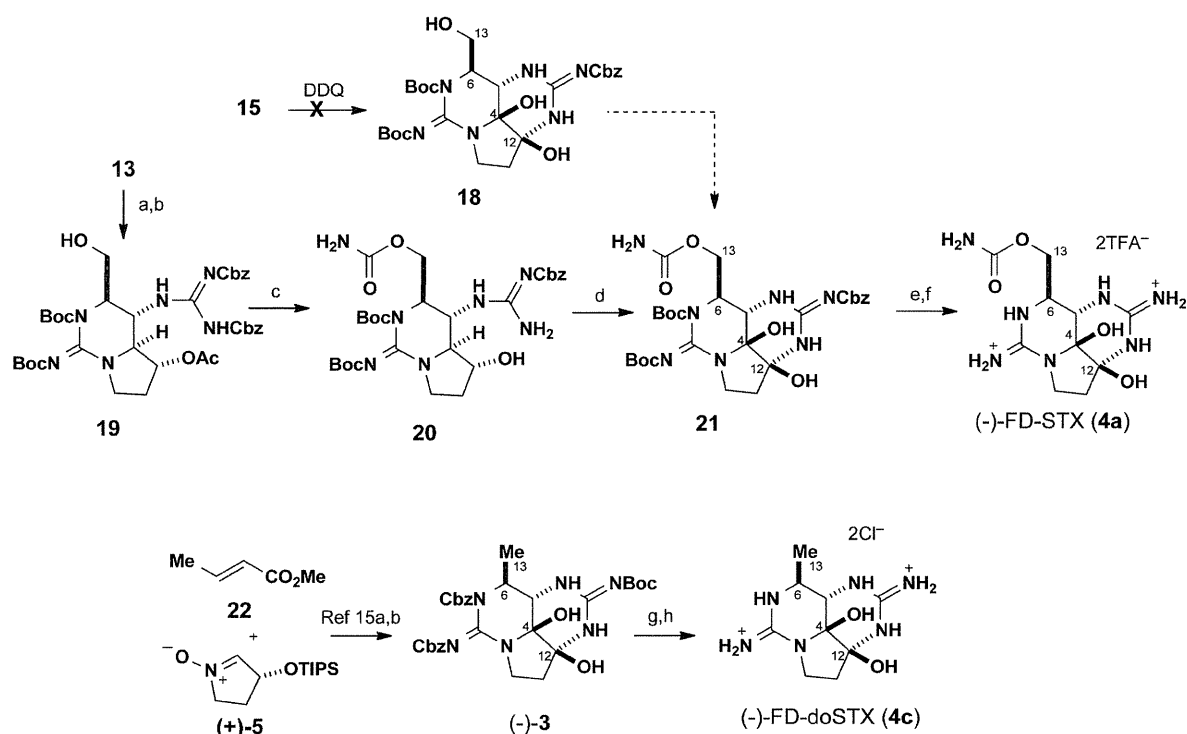
Scheme 1. Structures of saxitoxin (STX) and its derivatives, and a model of their binding to the sodium ion channel.^[12c]

Results and Discussion

Synthesis of (–)-FD-STX and its derivatives (4a–c): Scheme 4 and Scheme 5 illustrate the synthesis of **4a–c**. We first focused on the synthesis of (–)-FD-dcSTX (**4b**) (Scheme 4).^[15] 1,3-Dipolar cycloaddition reaction between chiral nitrene **5** and nitro alkene **6** at 40 °C gave the *endo* adduct of isoxazolidine **7** as a single diastereomer. The stereochemistry at C5 was epimerized with 1,8-diazabicyclo[5.4.0]undec-7-ene (DBU) in dichloromethane at 0 °C (95:5), and selective reduction of the nitro group was carried out with zinc powder in acetic acid to give hydroxylamine **8**



Scheme 4. Synthesis of (-)-FD-dcSTX (**4b**): a) 40°C; b) DBU, CH₂Cl₂, -40°C; c) AcOH, then Zn, -40°C, 72% (3 steps); d) CbzCl, K₂CO₃, THF/H₂O=3:1, 0°C, 83%; e) NaOAc, TiCl₃/HCl, Zn, CH₂Cl₂/MeOH=2:1, -50°C; f) BocN=C(SMe)NH₂ (**16**), HgCl₂, Et₃N, DMF, 82% (2 steps); g) ClCH₂SO₂Cl, *i*Pr₂EtN, CH₂Cl₂, 87%; h) TBAF, THF, 0°C, 93%; i) H₂, Pd(OH)₂/C, MeOH; j) CbzN=C(SMe)NHCbz (**17**), HgCl₂, Et₃N, DMF, 85% (2 steps); k) NaOMe, THF/MeOH, 0°C, 65%; l) IBX, DMSO, 70°C, 24%; m) H₂, Pd(OH)₂/C, MeOH; n) 3 N HCl, 78% (2 steps).



Scheme 5. Synthesis of (-)-FD-STX (**4a**): a) Ac₂O, pyridine (py), 50°C, 96%; b) DDQ, CH₂Cl₂, H₂O, 94%; c) trichloroacetyl isocyanate, CH₂Cl₂, then K₂CO₃, MeOH, 34%; d) IBX, DMSO, 70°C, 20%; e) H₂, Pd(OH)₂/C, MeOH; f) TFA, CH₂Cl₂, 82% (2 steps). Synthesis of (-)-FD-doSTX (**4c**): g) TFA, CH₂Cl₂; h) H₂, Pd/C, MeOH, 92% (2 steps).

in 72 % yield from (+)-**5**. The amino group of hydroxylamine in **8** was protected with a carbobenzyloxy (Cbz) group using Cbz chloride in the presence of potassium carbonate (83 %), then the two N–O bonds in **9** were reduced with TiCl_3 –NaOAc and zinc powder in aqueous MeOH–HCl to give diamine **10**. Next, a *tert*-butoxycarbonyl (Boc)-protected guanidine group was introduced with bis(Boc)-2-methyl-2-thiopseudourea (**16**) in the presence of mercury(II) chloride and triethyl amine to give guanidine **11** in 82 % yield from **8**. Cyclization of guanidine in **11** was carried out by treatment with monochloromethanesulfonyl chloride and diisopropylethylamine to give **12** in 87 % yield, and this product was subsequently converted into **13** in three steps: 1) deprotection of the triisopropylsilyl (TIPS) group with tetra-*n*-butylammonium fluoride (TBAF) in tetrahydrofuran (THF) at 0 °C, 2) deprotection of the Cbz group with hydrogen in the presence of Pearlman's catalyst,^[17] and 3) guanylation with bis(Cbz)-2-methyl-2-thiopseudourea (**17**) in the presence of mercury(II) chloride in 79 %.^[18] One of the Cbz groups of **13** was removed with sodium methoxide in THF/methanol to give **14** in 65 % yield.^[19] Construction of the tricyclic fused-type guanidine was carried out by oxidation of alcohol **14** with IBX (4 equiv) in DMSO at 70 °C to give **15** in 24 % yield. In this reaction, oxidation of the alcohol at C12 takes place first, then the C4 position is oxidized through the enamine–IBX complex.^[15] Finally, (–)-FD-dcSTX (**4b**) was obtained by deprotection of the Cbz and Boc groups on guanidines by hydrogenolysis in the presence of Pearlman's catalyst, followed by treatment with 3N HCl in 78 % yield from **15** (i.e., two steps).

The synthesis of (–)-FD-STX (**4a**) is illustrated in Scheme 5. Synthesis of (–)-FD-STX (**4a**) was attempted firstly from the tricyclic compound **15** by deprotection of the *p*-methoxybenzyl (MPM) group with 2,3-dichloro-5,6-dicyano-1,4-benzoquinone (DDQ). However, the desired reaction did not occur, and decomposition of the starting material was observed. The difficulty of this deprotection reaction presumably arises from steric effects at C13. Thus, functionalization at C13 was carried out before construction of the tricyclic FD-STX ring system. After protection of the hydroxyl group of bis-guanidine **13** at C12 with acetic anhydride, the MPM group in the resulting compound was successfully removed with DDQ to give alcohol **19** in 90 % yield (two steps). A carbamoyl group was introduced at C13 by treating the alcohol **19** with trichloroacetyl isocyanate, followed by hydrolysis with potassium carbonate in MeOH, thereby affording carbamate **20** in 34 % yield. Then, the fused-type tricyclic skeleton was constructed under IBX oxidation conditions to give **21** in 20 % yield. Finally, deprotection of the Cbz and Boc groups was carried out under the same conditions as described for the synthesis of **4b** to give (–)-FD-STX (**4a**) in 82 % yield (two steps).

(–)-FD-doSTX (**4c**) was synthesized by deprotection of fused-type tricyclic guanidine (–)-**3**, which was obtained from methyl crotonate (**22**) and the chiral nitrone (+)-**5** as previously reported (Scheme 5).^[15a,b]

Evaluation of Na_vCh -inhibitory activity of (–)-FD-STXs

4a–c: Next, the Na_vCh -inhibitory activity of **4a–c** was evaluated in terms of inhibition of the cytotoxicity of ouabain and veratridine in mouse neuroblastoma cell line Neuro-2a.^[20] In this cell-culture assay, Neuro-2a is treated with a sodium channel activator, veratridine, in the presence of ouabain, an inhibitor of Na^+/K^+ ATPase. This serves to block sodium ion efflux, and decreases the cell viability. Na_vCh inhibition by TTX, STX, and related compounds antagonizes this effect, and rescues the cells in a dose-dependent manner. The results obtained with **4a–c** are summarized in Table 1. (–)-FD-STX (**4a**) and (–)-FD-dcSTX (**4b**) showed a concentration-dependent inhibitory effect on the cytotoxicity of ouabain and veratridine, and their IC_{50} values were determined to be (7.3 ± 2.7) and (15 ± 5.4) μM , respectively (synthetic (+)-dcSTX (**1b**),^[15e] used as a control, showed an IC_{50} value of (82 ± 17) nM). On the other hand, (–)-FD-doSTX (**4c**) showed no inhibitory activity even at concentrations over 500 μM . Since (–)-FD-STX (**4a**) and (–)-FD-dcSTX (**4b**) were inhibitory, these compounds were further evaluated by means of the patch-clamp method with cells that express $\text{Na}_v1.4$ and $\text{Na}_v1.5$, which are representative members of the TTX-s and TTX-r channel subtypes, respectively.

Table 1. Na_vCh -inhibitory activity of FD-STXs **4a–c** in Neuro-2a cells.^[21]

Compounds	IC_{50} (mean \pm SD)	<i>n</i>
(–)-FD-STX (4a)	(7.3 ± 2.7) μM	3
(–)-FD-dcSTX (4b)	(15 ± 5.4) μM	3
(–)-FD-doSTX (4c)	> 500 μM	3
(+)-dc-STX (1b)	(82 ± 17) nM	3

The sodium ion channel α subunits of the rat skeletal muscle channel $\text{rNa}_v1.4$ and rat heart channel $\text{rNa}_v1.5$ were each transiently transfected into HEK293 cells, and electrophysiological recording was performed with a whole-cell patch clamp. In this experiment, the sodium ion channels were activated by applying 10-millisecond voltage steps from a holding potential of -120 mV every 5 s, which induces peak sodium currents (I_{Na}).^[22,23] Figure 1 shows time courses of changes in I_{Na} , the effects of inhibitors **4a** and **4b**, and the recovery upon washout of these compounds. In the case of $\text{Na}_v1.4$, sodium ion current was decreased by the addition of 10 μM (–)-FD-STX (**4a**) or (–)-FD-dcSTX (**4b**), and reached an equilibrium peak current ($I_{\text{inhibitor}}$ **4a** or **4b**). Recovery of sodium ion current was observed upon washout of the inhibitor, which indicates that the ligands were removed from the channel (Figure 1a and b). This is a typical reversible inhibition, like that of TTX and STX.^[24] The IC_{50} values of **4a** and **4b** were (3.8 ± 0.9) and (16 ± 8.6) μM , respectively (Table 2),^[25] and paralleled the results of the cell-based assay. In the case of $\text{Na}_v1.5$, the sodium ion current was decreased by the addition of 1 mM **4a** or **4b**, but recovery of the current following washout was observed only in the case of (–)-FD-STX (**4a**) (Figure 1c); the IC_{50} value of **4a** was (118 ± 8.1) μM (Table 2).^[25] On the other hand, recov-

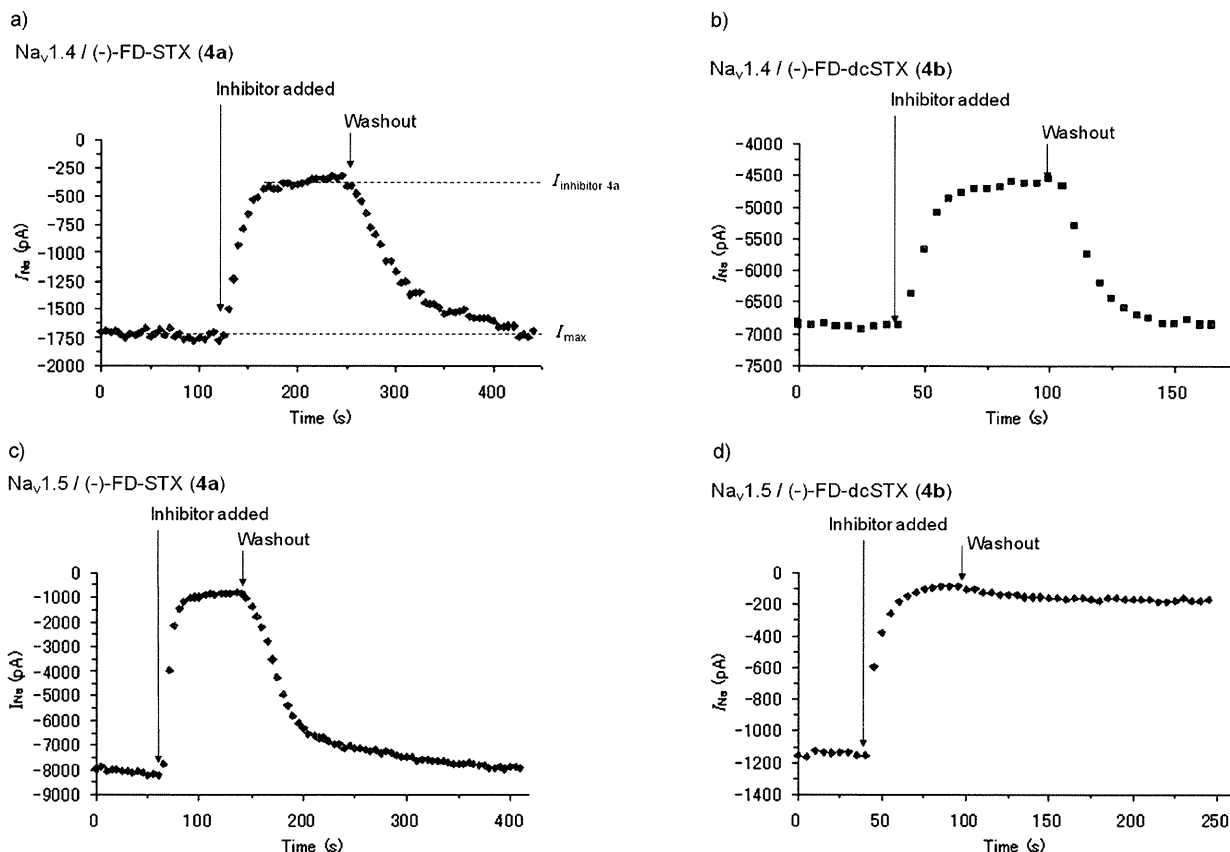


Figure 1. Time courses of inhibition by (–)-FD-STXs and recovery following washout. a) Effect of (–)-FD-STX (**4a**) (10 μM) on rNa_v1.4. b) Effect of (–)-FD-dcSTX (**4b**) (10 μM) on rNa_v1.4. c) Effect of (–)-FD-STX (**4a**) (1 mM) on rNa_v1.5. d) Effect of (–)-FD-dcSTX (**4b**) (1 mM) on rNa_v1.5.

Table 2. Inhibitory activity of (–)-FD-STXs **4a** and **4b** towards Na_v1.4 and Na_v1.5.

Compounds	IC ₅₀ (Na _v 1.4) (mean \pm SD)	<i>n</i>	IC ₅₀ (Na _v 1.5) (mean \pm SD)	<i>n</i>
(–)-FD-STX (4a)	(3.8 \pm 0.9) μM	3	(118 \pm 8.1) μM	3
(–)-FD-dcSTX (4b)	(16 \pm 8.6) μM	4	(182 \pm 16) μM	4

ery of the sodium ion current did not occur when (–)-FD-dcSTX (**4b**) was washed out (Figure 1d), thus indicating that the inhibition of Na_v1.5 by (–)-FD-dcSTX (**4b**) was “irreversible.” Although the inhibitory action of **4b** against Na_v1.5 was quite weak, with an IC₅₀ value of (182 \pm 16) μM (Table 2), this is the first observation of “irreversible” inhibition of Na_v1.5 by any STX- or TTX-related molecule. The interaction mechanism of **4b** and Na_v1.5 has not yet been clarified, but **4b** may be a good lead compound for the development of Na_vCh-subtype-selective modulators.

Conclusion

In summary, we have synthesized a novel skeletal analogue of saxitoxin, FD-STX. Three FD-STX derivatives **4a–c**, which differ in substitution at C13, were synthesized, and

their inhibitory effect on sodium ion channels was evaluated by means of cell-based assay and the patch-clamp method. (–)-FD-STX (**4a**) and (–)-FD-dcSTX (**4b**) showed IC₅₀ values of (7.3 \pm 2.7) and (15 \pm 5.4) μM , respectively, in cell-based assay. These compounds inhibited Na_v1.4 (a representative tetrodotoxin-sensitive Na_vCh subtype) and Na_v1.5 (a representative tetrodotoxin-resistant Na_vCh subtype) in cells that express those channels. Uniquely, (–)-FD-dcSTX (**4b**) showed “irreversible” inhibition of Na_v1.5; it is the first STX- or TTX-related molecule to do so. Further work to elucidate the mechanism of the “irreversible” interaction between (–)-FD-dcSTX (**4b**) and Na_v1.5 is in progress.

Experimental Section

General: Flash chromatography was performed on silica gel 60 (spherical, particle size 40–100 μm ; Kanto), Chromatorex NH (particle size 75–150 μm ; Fuji Silysia). Optical rotations were measured using a JASCO P-2200 polarimeter with the sodium D line. ¹H and ¹³C NMR spectra were recorded using a JEOL JNM ECX 400 or JNM-ECA 500 spectrometer. The spectra are referenced internally according to residual solvent signals of CDCl₃ (¹H NMR, δ =7.26 ppm; ¹³C NMR, δ =77.0 ppm); CD₃OD (¹H NMR, δ =3.30 ppm; ¹³C NMR, δ =49.0 ppm); or 4% CD₃CO₂D₂O (¹H NMR, δ =4.79 ppm (D₂O); ¹³C NMR, δ =179.0 ppm

(CD₃CO₂D)). Mass spectra were recorded using a JEOL JMS-T100X spectrometer in ESI-MS mode with methanol as solvent.

Isoxazolidine 8: A mixture of nitron (+)-**5** (10.4 g, 40.5 mmol) and nitro-olefin **6** (9.49 g, 42.5 mmol) was stirred for 30 min at 40 °C without any solvents. The reaction mixture was diluted with CH₂Cl₂ (150 mL) and cooled to -40 °C. DBU (6.28 mL, 44.6 mmol) was added to the solution, and the resulting mixture was stirred for 1 h at the same temperature. AcOH (11.6 mL, 203 mmol) and Zn powder (freshly activated with diluted aqueous HCl, 10.6 g, 162 mmol) were added to the reaction mixture, and the resulting mixture was warmed to 0 °C over 5 h. The reaction mixture was then filtered through a pad of Celite, and the filtrates were diluted with CH₂Cl₂ (150 mL) and washed with H₂O and brine. The combined organic layer was dried over MgSO₄, filtered, and concentrated in vacuo. The residue was purified by column chromatography on silica gel (hexane/EtOAc=15:1 to 5:1) to give hydroxylamine **8** (13.5 g, 72 %). Spectral data for **8**: [α]_D²⁵ = +1.3 (*c* = 2.3 in CHCl₃); ¹H NMR (400 MHz, CDCl₃): δ = 7.25 (d, *J* = 8.2 Hz, 2H), 6.86 (d, *J* = 8.2 Hz, 2H), 5.47 (br, 1H), 4.48 (s, 2H), 4.23 (dt, *J* = 2.3, 5.0 Hz, 1H), 4.12 (q, *J* = 6.0 Hz, 1H), 3.79 (s, 3H), 3.72 (m, 2H), 3.58 (m, 2H), 3.3 (m, 2H), 2.06 (m, 1H), 1.73 (m, 1H), 1.04 ppm (m, 21H); ¹³C NMR (100 MHz, CDCl₃): δ = 159.3, 129.6, 129.5, 113.8, 77.9, 77.8, 77.4, 73.4, 71.9, 67.9, 55.8, 55.2, 34.4, 17.9, 12.0 ppm; HRMS (ESI): *m/z* calcd for C₂₄H₄₂N₂O₅SiNa: 489.2761 [*M*+Na⁺]; found: 489.2758.

Isoxazolidine 9: CbzCl (4.01 mL, 28.1 mmol) was added to a solution of 5 α -hydroxylamine **8** (10.1 g, 21.6 mmol) and K₂CO₃ (15.0 g, 108 mmol) in THF/H₂O (1:1, 60 mL) at 0 °C. After stirring for 10 min at 0 °C, methanol (150 mL) was added to the solution, and resulting mixture was warmed to room temperature. The reaction mixture was then concentrated in vacuo, and the resulting material was diluted with EtOAc/H₂O and extracted with EtOAc. The combined organic layer was dried over MgSO₄, filtered, and concentrated in vacuo. The residue was purified by column chromatography on silica gel (hexane/EtOAc=8:1 to 4:1) to give *N*-Cbz-hydroxylamine **9** (10.7 g, 83 %). Spectral data for isoxazolidine **9**: [α]_D²⁵ = +4.5 (*c* = 2.9 in CHCl₃); ¹H NMR (400 MHz, CDCl₃): δ = 7.35 (m, 5H), 7.19 (d, *J* = 8.7 Hz, 2H), 6.79 (d, *J* = 8.7 Hz, 2H), 5.20 (d, *J* = 12.4 Hz, 1H), 5.15 (d, *J* = 12.4 Hz, 1H), 4.90 (dd, *J* = 4.6, 5.0 Hz, 1H), 4.44 (d, *J* = 11.0 Hz, 1H), 4.40 (d, *J* = 11.0 Hz, 1H), 4.34 (m, 1H), 4.12 (dt, *J* = 3.2, 6.0 Hz, 1H), 3.97 (dd, *J* = 2.8, 3.2 Hz, 1H), 3.83 (dd, *J* = 5.5, 11.0 Hz, 1H), 3.77 (s, 3H), 3.63 (dd, *J* = 3.2, 11.0 Hz, 1H), 3.40 (ddd, *J* = 6.0, 12.6, 9.6 Hz, 1H), 3.16 (ddd, *J* = 4.1, 6.9, 12.6 Hz, 1H) 2.02 (m, 1H), 1.75 (m, 1H), 1.01 ppm (m, 21H); ¹³C NMR (100 MHz, CDCl₃): δ = 159.5, 156.0, 136.0, 129.8, 128.5, 128.2, 113.9, 77.4, 76.3, 75.2, 73.9, 67.8, 67.1, 66.7, 55.2, 33.9, 17.9, 17.8, 11.9 ppm; HRMS (ESI): *m/z* calcd for C₂₂H₄₈N₂O₅SiNa: 623.3129 [*M*+Na⁺]; found: 623.3137.

Guanidine 11: TiCl₃/HCl (20 % aq, 14.0 mL, 21.5 mmol; Kanto Chemical) was added to a solution of *N*-Cbz hydroxylamine **9** (10.7 g, 17.9 mmol), NaOAc (4.40 g, 53.7 mmol), and freshly activated Zn powder (3.51 g, 53.7 mmol) in CH₂Cl₂/MeOH (2:1, 100 mL) at -50 °C under Ar atmosphere. After stirring for 24 h, the reaction was neutralized by an excess amount of Na₂CO₃ solid at -50 °C. Then the dark-green suspension was warmed to room temperature and vigorously stirred under O₂ atmosphere for 30 min. The light-gray suspension was extracted with EtOAc, and the extracts were washed with water and brine. The organic layer was dried over MgSO₄, filtered, and concentrated in vacuo to give diamine **10** (10.8 g). HgCl₂ (4.37 mg, 16.1 mmol) was added to a solution of crude diamine **10** (10.3 g), bis(Boc)-2-methyl-2-thiopseudourea (**16**) (4.68 g, 16.1 mmol), and Et₃N (7.48 mL, 53.7 mmol) in DMF (40 mL) at room temperature under Ar atmosphere. After stirring for 1 h, the reaction mixture was diluted with EtOAc (150 mL) and filtered through a pad of Celite. The filtrates were washed with H₂O and brine. The organic layer was dried over MgSO₄, filtered, and concentrated in vacuo. The residue was purified by column chromatography on silica gel (hexane/EtOAc=15:1 to 6:1) to give guanidine **11** (12.1 g, 82 %, two steps). Spectral data for guanidine **11**: [α]_D²⁵ = +0.6 (*c* = 2.6 in CHCl₃); ¹H NMR (500 MHz, CDCl₃): δ = 10.31 (s, 1H), 7.33 (m, 5H), 7.21 (d, *J* = 8.6 Hz, 2H), 6.83 (d, *J* = 8.6 Hz, 2H), 5.69 (br, 1H), 5.44 (d, *J* = 9.7 Hz, 1H), 5.09 (d, *J* = 12.0 Hz, 1H), 5.04 (d, *J* = 12.0 Hz, 1H), 4.47 (d, *J* = 2.9 Hz, 1H), 4.42 (d, *J* = 12.0 Hz, 1H), 4.39 (d, *J* = 9.7 Hz, 1H), 4.33 (d, *J* = 12.0 Hz,

1H), 3.92 (d, *J* = 10.9 Hz, 1H), 3.89 (dd, *J* = 6.3, 6.9 Hz, 1H), 3.77 (s, 3H), 3.59 (dt, *J* = 8.0, 10.9 Hz, 1H), 3.50 (m, 2H), 3.31 (dd, *J* = 6.9, 9.7 Hz, 1H), 2.35 (m, 1H), 1.96 (dd, *J* = 6.9, 13.8 Hz, 1H), 1.49 (s, 9H), 1.43 (s, 9H), 0.97 ppm (m, 21H); ¹³C NMR (125 MHz, CDCl₃): δ = 161.3, 159.1, 157.6, 156.5, 150.2, 136.4, 130.2, 129.6, 128.4, 128.0, 127.9, 113.6, 82.3, 79.5, 73.0, 72.5, 69.8, 66.8, 55.2, 53.0, 46.8, 32.1, 28.1, 28.0, 17.8, 12.0 ppm; HRMS (ESI): *m/z* calcd for C₄₃H₆₈N₄O₁₀SiNa: 851.4569 [*M*+Na⁺]; found: 851.4602.

Cyclic guanidine 12: Chloromethanesulfonyl chloride (5.72 mL, 47.6 mmol) was added to a solution of guanidine **11** (9.90 g, 11.9 mmol) and *i*Pr₂NEt (12.3 mL, 71.4 mmol) in CH₂Cl₂ (60 mL) at 0 °C under an N₂ atmosphere. After stirring for 12 h, the reaction was quenched with saturated aqueous NaHCO₃, and the organic layer was extracted with CH₂Cl₂. The combined organic layer was dried over MgSO₄, filtered, and concentrated in vacuo. The residue was purified by column chromatography on silica gel (hexane/EtOAc=10:1 to 8:1) to give cyclic guanidine **12** (8.35 mg, 87 %). Spectral data for cyclic guanidine **12**: [α]_D²⁵ = +70.9 (*c* = 2.0 in CHCl₃); ¹H NMR (400 MHz, CDCl₃): δ = 7.33 (m, 5H), 7.20 (d, *J* = 8.24 Hz, 2H), 6.82 (d, *J* = 8.7 Hz, 2H), 5.09 (d, *J* = 12.2 Hz, 1H), 5.05 (br, 1H), 4.99 (d, *J* = 12.2 Hz, 1H), 4.45 (d, *J* = 10.3 Hz, 1H), 4.42 (m, 1H), 4.29 (d, *J* = 10.3 Hz, 1H), 4.25 (m, 1H), 3.99 (m, 1H), 3.80 (m, 1H), 3.78 (s=3H), 3.71 (dd, *J* = 3.7, 10.1 Hz, 1H), 3.58 (m, 1H), 3.38 (dd, *J* = 7.3, 9.6 Hz, 1H), 3.18 (d, *J* = 11.0 Hz, 1H), 1.76 (m, 2H), 1.49 (s, 9H), 1.48 (s, 9H), 0.98 ppm (m, 21H); ¹³C NMR (100 MHz, CDCl₃): δ = 159.6, 159.0, 155.6, 151.6, 135.9, 130.4, 129.6, 128.6, 128.3, 128.2, 113.4, 82.6, 78.1, 74.1, 72.9, 71.4, 68.6, 67.2, 59.4, 55.2, 54.5, 46.0, 32.6, 28.5, 28.1, 17.9, 12.0 ppm; HRMS (ESI): *m/z* calcd for C₄₃H₆₇N₄O₅Si: 811.4677 [*M*+H⁺]; found: 811.4678.

Bis(guanidine) 13: TBAF (3.23 g, 12.4 mmol) was added to a solution of cyclic guanidine **12** (8.35 g, 10.3 mmol) in a solution of THF (50 mL) at 0 °C. After stirring for 1 h, the reaction mixture was quenched with saturated aqueous NH₄Cl and extracted with EtOAc. The organic layer was dried over MgSO₄, filtered, and concentrated in vacuo. The residue was purified by column chromatography on silica gel (hexane/EtOAc; 3:1 to EtOAc) to give alcohol (6.27 g, 93 %). Pd(OH)₂/C (5 %, 33.3 mg) was added to a solution of alcohol (665 mg 1.02 mmol) in methanol (15 mL), and the mixture was vigorously stirred under an H₂ atmosphere (balloon) at room temperature for 3 h. The reaction mixture was filtered through a pad of Celite, and the filtrates were concentrated in vacuo to give amine (538 mg). HgCl₂ (331 mg, 1.22 mmol), Et₃N (425 μ L, 3.05 mmol), and 1,3-bis(Cbz)-2-methyl-2-thiopseudourea (**17**) (437 mg, 1.22 mmol) were added to a solution of amine (538 mg) in DMF (10 mL) at room temperature for 2 h. The reaction mixture was diluted with EtOAc, and the resulting solution was filtered through a pad of Celite. The filtrates were washed with H₂O and brine twice, and the organic layer was dried over MgSO₄, filtered, and concentrated in vacuo. The residue was purified by column chromatography on Chromatorex NH (hexane/EtOAc=3:2 to 1:2) to give bis(guanidine) **13** (718.2 mg, 85 %, two steps). Spectral data for bis(guanidine) **13**: [α]_D²⁵ = +122.8 (*c* = 0.46 in MeOH); ¹H NMR (400 MHz, CDCl₃): δ = 11.59 (s, 1H), 8.97 (d, *J* = 6.41 Hz, 1H), 7.35 (m, 10H), 7.22 (d, *J* = 8.42 Hz, 2H), 6.82 (d, *J* = 8.42 Hz, 2H), 5.48 (brs, 1H), 5.16 (dd, *J* = 11.91, 17.40 Hz, 2H), 5.13 (dd, *J* = 12.36, 45.34 Hz, 2H), 4.62 (m, 1H), 4.51 (t, *J* = 5.50, 1H), 4.39 (dd, *J* = 11.45, 27.93 Hz, 2H), 3.90 (dd, *J* = 7.33, 19.00 Hz, 1H), 3.77 (dd, *J* = 4.12, 9.62 Hz, 1H), 3.73 (s, 3H), 3.68 (m, 1H), 3.47 (dd, *J* = 8.24, 9.62 Hz, 1H), 3.17 (dd, *J* = 4.12, 7.79 Hz, 1H), 3.09 (m, 1H), 2.01 (m, 1H), 1.67 (m, 1H), 1.53 (s, 9H), 1.49 ppm (s, 9H); ¹³C NMR (100 MHz, CDCl₃): δ = 162.73, 159.61, 155.96, 153.58, 136.42, 134.40, 130.17, 129.56, 129.09, 128.83, 128.67, 128.18, 127.87, 113.93, 83.86, 78.85, 74.93, 73.33, 69.34, 68.79, 68.69, 67.43, 55.36, 54.95, 45.17, 30.13, 28.53, 28.06 ppm; HRMS (ESI): *m/z* calcd for C₄₃H₅₅N₆O₁₁: 831.3928 [*M*+H⁺]; found: 831.3896.

Mono-Cbz-guanidine 14: NaOMe (9.0 mg, 0.167 mmol) was added to a solution of bis(guanidine) **13** (139 mg, 0.167 mmol) in THF/MeOH (1:1, 4 mL) at 0 °C under an N₂ atmosphere, and the mixture was stirred for 5 h. Saturated aqueous NH₄Cl was added to the reaction mixture, and organic layer was extracted with EtOAc. The extracts were washed with brine, dried over MgSO₄, filtered, and concentrated in vacuo. The residue was purified by column chromatography on Chromatorex NH (hexane/

EtOAc=3:1 to 1:2) to give mono-Cbz-guanidine **14** (75.7 mg, 65%). Spectral data for mono-Cbz-guanidine **14**: $[\alpha]_D^{25} = +202.3$ ($c = 1.0$ in MeOH); $^1\text{H NMR}$ (500 MHz, CD_3OD): $\delta = 7.17$ (m, 5H), 7.08 (d, $J = 8.70$ Hz, 2H), 6.67 (d, $J = 8.70$ Hz, 2H), 4.98 (dd, $J = 12.82, 39.38$ Hz, 2H), 4.32 (d, $J = 10.53$ Hz, 1H), 4.19 (m, 2H), 4.10 (d, $J = 3.21$ Hz, 1H), 3.67 (m, 3H), 3.63 (s, 3H), 3.41 (m, 1H), 3.20 (m, 2H), 1.75 (m, 1H), 1.61 (m, 1H), 1.38 (m, 18H); $^{13}\text{C NMR}$ (125 MHz, CD_3OD): $\delta = 162.07, 160.11, 159.89, 152.55, 138.36, 129.96, 128.86, 128.17, 128.03, 113.94, 83.39, 78.93, 73.32, 73.02, 71.32, 67.93, 60.84, 55.11, 49.34, 46.76, 32.09, 28.42, 27.87$ ppm; HRMS (ESI): m/z calcd for $\text{C}_{35}\text{H}_{49}\text{N}_6\text{O}_9$: 697.3561 $[\text{M}+\text{H}^+]$; found: 697.3557.

Fused-type bicyclic guanidine 15: IBX (117 mg, 0.416 mmol) was added to a solution of mono-Cbz-guanidine **14** (48 mg, 0.0694 mmol) in DMSO (1 mL), and the resulting suspension was stirred for 10 min at room temperature. Then the reaction mixture was warmed to 70 °C and stirred for 6 h. The reaction was quenched with 10% aqueous $\text{Na}_2\text{S}_2\text{O}_3$ and saturated aqueous NaHCO_3 , and the solution was diluted with EtOAc. The organic layer was dried over MgSO_4 , filtered, and concentrated in vacuo. The residue was purified by column chromatography on Chromatorex NH (hexane/EtOAc; 1:2 to pure EtOAc) to give fused-type bicyclic guanidine **15** (11.8 mg, 24%). Spectral data for fused-type bicyclic guanidine **15**: $[\alpha]_D^{25} = +16.9$ ($c = 0.47$ in MeOH); $^1\text{H NMR}$ (500 MHz, CD_3OD): $\delta = 7.20$ (m, 7H), 6.79 (d, $J = 8.59$ Hz, 2H), 4.94 (d, $J = 1.72$ Hz, 2H), 4.61 (m, 1H), 4.51 (d, $J = 10.88$ Hz, 1H), 4.39 (d, $J = 10.88$ Hz, 1H), 4.00 (d, $J = 1.72$ Hz, 1H), 3.90 (dd, $J = 7.45, 9.74$ Hz, 1H), 3.72 (m, 1H), 3.68 (s, 3H), 3.52 (m, 1H), 3.44 (m, 1H), 2.27 (m, 1H), 2.08 (m, 1H), 1.34 ppm (m, 18H); $^{13}\text{C NMR}$ (125 MHz, CD_3OD): $\delta = 160.95, 160.37, 153.10, 150.41, 138.66, 131.02, 130.75, 129.38, 128.86, 114.79, 86.42, 85.32, 82.96, 80.05, 74.15, 69.82, 67.48, 59.99, 55.69, 45.08, 33.10, 28.77, 28.27$ ppm; HRMS (ESI): m/z calcd for $\text{C}_{35}\text{H}_{47}\text{N}_6\text{O}_{10}$: 711.3353 $[\text{M}+\text{H}^+]$; found: 711.3400.

(–)-**FD-dcSTX (4b)**: $\text{Pd}(\text{OH})_2/\text{C}$ (2.5 mg) was added to a solution of **15** (4.8 mg, 0.00676 mmol) in methanol (500 μL), and the suspension was vigorously stirred under an H_2 atmosphere (balloon) at room temperature. After 30 min, the reaction mixture was filtered through a pad of Celite. The filtrates were concentrated in vacuo to give bis-Boc-bis-aminal (3.1 mg). HCl (3N, 500 μL) was added to the resulting bis-Boc-bis-aminal at 0 °C. After stirring for 1 h at room temperature, the reaction mixture was concentrated in vacuo to give FD-dcSTX (**4b**) as a 2HCl salt (1.7 mg, 78%). Spectral data for FD-dcSTX (**4b**): $[\alpha]_D^{25} = -63.9$ ($c = 0.11$ in MeOH); $^1\text{H NMR}$ (400 MHz, D_2O): $\delta = 4.06$ (s, 1H), 3.82 (m, 3H), 3.70 (t, $J = 10.07$ Hz, 1H), 3.48 (dd, $J = 10.07, 17.40$ Hz, 2H), 2.50 (m, 1H), 2.34 ppm (m, 1H); $^{13}\text{C NMR}$ (100 MHz, 4% $\text{CD}_3\text{CO}_2\text{D}$ in D_2O): $\delta = 152.91, 152.16, 85.53, 81.49, 62.52, 55.24, 45.78, 43.00, 31.44$ ppm; HRMS (ESI): m/z calcd for $\text{C}_9\text{H}_{17}\text{N}_6\text{O}_3$: 257.1362 $[\text{M}+\text{H}^+]$; found: 257.1319.

Alcohol 19: Ac_2O (2 mL) and pyridine (2 mL) were added to bis-guanidine **13** (230 mg, 0.277 mmol) at room temperature. The reaction mixture was warmed to 50 °C and stirred for a further 3 h, then cooled to room temperature, quenched with toluene, and concentrated in vacuo. The resulting residue was purified by column chromatography on Chromatorex NH (hexane/EtOAc; 6:1 to 3:1) to give acetate (245 mg, 96%). DDQ (377 mg, 1.66 mmol) was added to a solution of acetate (144 mg, 0.166 mmol) in $\text{CH}_2\text{Cl}_2/\text{H}_2\text{O}$ (2:1, 4.5 mL) at room temperature. The reaction mixture was stirred for 5 h, saturated aqueous NaHCO_3 was added, and the resulting solution was extracted with EtOAc three times. The combined organic layer was dried over MgSO_4 , filtered, and concentrated in vacuo. The residue was purified by column chromatography on Chromatorex NH (hexane/EtOAc; 6:1 to 1:1) to give alcohol **19** (117 mg, 94%). Spectral data for alcohol **19**: $[\alpha]_D^{25} = +171.7$ ($c = 0.3$ in MeOH); $^1\text{H NMR}$ (400 MHz, CDCl_3): $\delta = 11.76$ (s, 1H), 8.65 (d, $J = 8.70$ Hz, 1H), 7.36 (m, 10H), 5.23 (dd, $J = 11.91, 20.15$ Hz, 2H), 5.11 (dd, $J = 12.82, 16.94$ Hz, 2H), 4.48 (m, 1H), 4.08 (m, 1H), 4.00 (d, $J = 11.91$ Hz, 1H), 3.90 (t, $J = 8.70$ Hz, 1H), 3.65 (m, 3H), 3.52 (t, $J = 9.63$ Hz, 1H), 2.05 (m, 5H), 1.49 ppm (m, 18H); $^{13}\text{C NMR}$ (100 MHz, CDCl_3): $\delta = 170.83, 163.55, 156.44, 153.82, 136.63, 134.44, 128.82, 128.59, 128.52, 127.87, 83.51, 80.11, 75.41, 68.63, 67.33, 64.91, 51.36, 46.46, 28.77, 28.38, 28.20, 21.16$ ppm; HRMS (ESI): m/z calcd for $\text{C}_{37}\text{H}_{49}\text{N}_6\text{O}_{11}$: 753.3459 $[\text{M}+\text{H}^+]$; found: 753.3496.

Carbamate 20: Trichloroacetyl isocyanate (19 μL , 0.16 mmol) was added to a solution of alcohol **19** (60 mg, 0.080 mmol) in CH_2Cl_2 (1 mL) at 0 °C under an N_2 atmosphere. The reaction was stirred for 1 h. Then K_2CO_3 and methanol were added at 0 °C and warmed to room temperature. The reaction mixture was stirred for 3 h, quenched with H_2O , and extracted with EtOAc twice. The combined organic layer was dried over MgSO_4 , filtered, and concentrated in vacuo. The residue was purified by preparative TLC (EtOAc, four times) to give carbamate **20** (16.6 mg, 34%). Spectral data for carbamate **20**: $[\alpha]_D^{25} = +52.3$ ($c = 0.32$ in MeOH); $^1\text{H NMR}$ (400 MHz, CD_3OD): $\delta = 7.30$ (m, 5H), 5.04 (dd, $J = 3.66, 12.82$ Hz, 2H), 4.25 (m, 2H), 3.58 (m, 4H), 3.28 (m, 2H), 1.84 (m, 1H), 1.69 (m, 1H), 1.41 ppm (m, 18H); $^{13}\text{C NMR}$ (100 MHz, CD_3OD): $\delta = 162.6, 160.2, 159.1, 154.6, 152.8, 138.8, 129.4, 128.6, 84.2, 79.7, 73.3, 68.3, 64.8, 59.8, 55.6, 47.2, 45.5, 32.8, 28.8, 28.7$ ppm; HRMS (ESI): m/z calcd for $\text{C}_{28}\text{H}_{42}\text{N}_7\text{O}_9$: 620.3044 $[\text{M}+\text{H}^+]$; found: 620.3074.

Fused-type bicyclic guanidine 21: IBX (62 mg, 0.223 mmol) was added to a solution of **20** (23 mg, 0.0371 mmol) in DMSO (1 mL), and the resulting suspension was stirred for 10 min at room temperature. Then, the reaction mixture was warmed to 70 °C and stirred for 6 h. The reaction was quenched with 10% aqueous $\text{Na}_2\text{S}_2\text{O}_3$ and saturated aqueous NaHCO_3 , and the solution was diluted with EtOAc. The organic layer was dried over MgSO_4 , filtered, and concentrated in vacuo. The residue was purified by column chromatography on Chromatorex NH (hexane/EtOAc; 1:8 to EtOAc/MeOH; 10:1) to give **21** (4.6 mg, 20%). Spectral data for fused-type bicyclic guanidine **21**: $[\alpha]_D^{25} = -7.4$ ($c = 0.1$ in MeOH); $^1\text{H NMR}$ (400 MHz, CD_3OD): $\delta = 7.34$ (m, 5H), 5.02 (d, $J = 1.37$ Hz, 2H), 4.53 (dd, $J = 5.5, 11.9$ Hz, 2H), 4.28 (dd, $J = 8.24, 11.91$ Hz, 2H), 4.02 (d, $J = 3.96$ Hz, 1H), 3.63 (m, 3H), 2.37 (m, 1H), 2.19 (m, 1H), 1.48 (m, 9H), 1.44 ppm (m, 9H); $^{13}\text{C NMR}$ (100 MHz, CD_3OD): $\delta = 164.5, 160.6, 159.3, 153.0, 149.5, 138.6, 130.0, 129.5, 86.9, 86.2, 83.9, 80.8, 68.1, 65.3, 59.8, 45.8, 33.5, 29.3, 28.9$ ppm; HRMS (ESI): m/z calcd for $\text{C}_{28}\text{H}_{40}\text{N}_7\text{O}_{10}$: 634.2836 $[\text{M}+\text{H}^+]$; found: 634.2884.

(–)-**FD-STX (4a)**: $\text{Pd}(\text{OH})_2/\text{C}$ (3.0 mg) was added to a solution of **21** (7.0 mg, 0.011 mmol) in methanol (600 μL) under an N_2 atmosphere, and the suspension was vigorously stirred under an H_2 atmosphere (balloon) at room temperature. After 7 h, the reaction mixture was filtered through a pad of Celite. The filtrates were concentrated in vacuo to give bis-Boc-bis-aminal (5.3 mg). TFA (500 μL) was added to the resulting bis-Boc-bis-aminal (5.3 mg) in CH_2Cl_2 (1 mL) at room temperature. Methanol was added to the reaction mixture, and it was concentrated in vacuo to give FD-STX (**4a**) as a 2TFA salt (4.8 mg, 82%). Spectral data for FD-STX (**4a**): $[\alpha]_D^{25} = -34.9$ ($c = 0.32$ in MeOH); $^1\text{H NMR}$ (400 MHz, D_2O): $\delta = 4.39$ (dd, $J = 8.70, 10.99$ Hz, 1H), 4.17 (dd, $J = 4.58, 10.99$ Hz, 1H), 4.05 (s, 1H), 3.95 (dd, $J = 4.58, 8.70$ Hz, 1H), 3.68 (t, $J = 10.7$ Hz, 1H), 3.45 (m, 1H), 2.47 (m, 1H), 2.33 ppm (dd, $J = 7.79, 13.28$ Hz, 1H); $^{13}\text{C NMR}$ (100 MHz, 4% $\text{CD}_3\text{CO}_2\text{D}$ in D_2O): $\delta = 159.1, 153.0, 152.1, 85.6, 81.7, 65.5, 52.5, 45.3, 43.2, 31.4$ ppm; HRMS (ESI): m/z calcd for $\text{C}_{10}\text{H}_{18}\text{N}_7\text{O}_4$: 300.1420 $[\text{M}+\text{H}^+]$; found: 300.1390.

(–)-**FD-doSTX (4c)**: TFA (0.5 mL) was added to a solution of (–)-**3** (12.1 mg, 0.0199 mmol) in CH_2Cl_2 (1 mL) at 0 °C. After being stirred for 1 h at room temperature, the reaction mixture was concentrated under reduced pressure to give bis-Cbz-bis-aminal. Pd/C (catalytic amount) was added to a solution of bis-Cbz-bis-aminal in methanol (1 mL), and the reaction mixture was stirred at room temperature under an atmosphere of hydrogen (balloon). After 4 h, the reaction mixture was filtered through a pad of Celite, and the filtrates were acidified with 10% HCl and concentrated in vacuo to give FD-doSTX (**4c**) (5.7 mg, 92%, two steps). Spectral data for **4c**: $[\alpha]_D^{20} = -62.7$ ($c = 0.9$ in MeOH); $^1\text{H NMR}$ (400 MHz, D_2O): $\delta = 3.88$ (d, $J = 1.0$ Hz, 1H), 3.75 (dd, $J = 1.0, 7.1$ Hz, 1H), 3.63 (dd, $J = 9.6, 9.6$ Hz, 1H), 3.42 (ddd, $J = 7.6, 9.6, 10.1$ Hz, 1H), 2.43 (ddd, $J = 9.6, 10.1, 13.6$ Hz, 1H), 2.27 (dd, $J = 7.6, 13.6$ Hz, 1H), 1.38 ppm (d, $J = 7.1$ Hz, 3H); $^{13}\text{C NMR}$ (100 MHz, CD_3OD): $\delta = 152.4, 151.4, 85.2, 81.8, 49.4, 48.4, 42.6, 30.9, 19.3$ ppm; HRMS (ESI): m/z calcd for $\text{C}_9\text{H}_{17}\text{N}_6\text{O}_2$: 241.1413 $[\text{M}+\text{H}^+]$; found: 241.1434.

Cell culture: HEK293 was grown in Dulbecco's modified Eagle's medium (DMEM; Life Technologies, Grand Island, NY) supplemented with 10% fetal bovine serum (30 units; BioWhittaker, Walkersville, MD) and penicillin-streptomycin (30 $\mu\text{g mL}^{-1}$; Gibco, Grand Island, NY), re-

spectively, at 37°C under 5% CO₂ in air. Neuro-2a was grown in RPMI1640 (Sigma, St Louis, MO) supplemented with 10% fetal bovine serum (50 units) and penicillin-streptomycin (50 µg mL⁻¹), respectively.

Inhibitory activity to cytotoxicity of ouabain and veratridine: Two days' worth of cultured cells (Neuro-2a) were harvested with a trypsin/EDTA solution (0.5%/0.2%) (Sigma) in phosphate buffer saline (PBS), and cells were suspended at a density of 5×10^5 cells mL⁻¹ in growth medium. This suspension (100 µL) was plated into each well of 96-well plates and cultured at 37°C for 24 h. The test medium (100 µL), which contained FD-STXs at various concentrations with 1:3 serial dilutions, and ouabain (200 µM) (Sigma) and veratridine (100 µM) (Sigma) were poured into each well. After incubation at 37°C for 24 h, the number of viable cells was determined using Cell Counting Kit-8 (Dojindo, Kumamoto, Japan). Cell Counting Kit-8 solution (5 µL) was added to each well, and the mixture was incubated at 37°C for 3 h, and then absorbance at 450/655 nm was measured using a microplate reader (Model 550, Bio-Rad, Hercules, CA). Three wells were used for one concentration of FD-STXs, and the same experiments were performed three times for each compound. The cell viability of the well that contained the test medium in the presence of only ouabain and veratridine (in the absence of FD-STXs) was taken as 0%, and that of the well that contained the medium in the absence of all toxins was taken as 100%. The IC₅₀ were determined from the Hill plot.

Transient transfection: One day before transfection, HEK293 cells were plated in a 60 mm dish (10^6 cells dish⁻¹). Complexes for transfection were prepared as follows: DNA (1 µg) was diluted in Buffer EC (150 µL), the DNA condensation buffer for the Effectene (Hilden, Germany). Next, enhancer (8 µL) from the kit was added to the solution of DNA. After 2 min, the above medium (150 µL) and Effectene Transfection Reagent kit (25 µL; Hilden, Germany) were mixed and incubated for 5 min. Then, the medium from 50%-confluent cells was removed and replaced with fresh medium (3 mL). The mixed complexes with fresh medium (1 mL) were added to the dish, which was swirled by rocking the plate. Incubation was continued at 37°C under humidified 5% CO₂ in air for at least 4 h. The medium was removed and replaced with fresh medium (4 mL). The transfected cells were used for electrophysiological experiments within 24–72 h after transfection.

Electrophysiological recording: Macroscopic sodium ion currents from the transfected cells were measured using a whole-cell patch-clamp method using an Axopatch 200B amplifier (Axon Instruments; Foster City, CA, USA); more than 80% of the series resistance was compensated to minimize voltage errors. The bath solution contained: NaCl (70 mM), *N*-methyl-D-glucamine (67 mM), CaCl₂ (1 mM), MgCl₂ (1.5 mM), glucose (10 mM), and HEPES (5 mM) (pH 7.4). The pipette solution contained: CsF (70 mM), CsCl (60 mM), NaF (12 mM), EGTA (5 mM), and HEPES (5 mM) (pH 7.4). Whole-cell patch pipettes with a resistance of 1.5–2 MΩ were used to achieve optimum voltage control. Recordings were started 10 min after establishing a whole-cell recording configuration. Whole-cell membrane currents were digitized at a sampling rate of 50–100 kHz using a 12-bit analogue-to-digital converter (DigiData 1321A interface; Axon Instruments), controlled by pClamp software (version 8; Axon Instruments). All experiments were conducted at room temperature (22–24°C). Data are presented as mean ± SD along with the number of observations (*n*), unless otherwise stated.

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- [21] See the Supporting Information for calculation of IC₅₀ values.
- [22] The voltage that provided a peak current (maximum active potential) for rNa_v1.4 and rNa_v1.5 was –10 mV.
- [23] The holding potentials of Na_v1.4 and Na_v1.5 are –100 and –120 mV, respectively.
- [24] Under the same conditions, (+)-dc-STX (**1b**) showed inhibitory activities against Na_v1.4 and Na_v1.5 with IC₅₀ values of (29 ± 1.8) nM and (1.2 ± 0.1) μM, respectively. Compound (+)-dc-STX (**1b**) interacted reversibly with both channels.
- [25] The IC₅₀ values were determined according to the following equation; $IC_{50} = [inhibitor\ 4] (I_{inhibitor4}/I_{max}) / (1 - (I_{inhibitor4}/I_{max}))$, in which I_{max} is the peak current under control conditions and $I_{inhibitor4}$ is the equilibrium peak current in the presence of an inhibitor **4**; J. L. Penzotti, H. A. Fozzard, G. M. Lipkind, S. C. Dudley, Jr., *Biophys. J.* **1998**, *75*, 2647–2657.

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Correlation Analyses on Binding Affinity of Sialic Acid Analogues and Anti-Influenza Drugs with Human Neuraminidase Using *ab Initio* MO Calculations on Their Complex Structures – LERE-QSAR Analysis (IV)

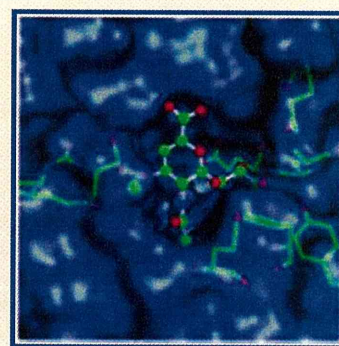
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S Supporting Information

ABSTRACT: We carried out full *ab initio* fragment molecular orbital (FMO) calculations for complexes comprising human neuraminidase-2 (hNEU2) and sialic acid analogues including anti-influenza drugs zanamivir (Relenza) and oseltamivir (Tamiflu) in order to examine the variation in the observed inhibitory activity toward hNEU2 at the atomic and electronic levels. We recently proposed the LERE (linear expression by representative energy terms)-QSAR (quantitative structure–activity relationship) procedure. LERE-QSAR analysis quantitatively revealed that the complex formation is driven by hydrogen-bonding and electrostatic interaction of hNEU2 with sialic acid analogues. The most potent inhibitory activity, that of zanamivir, is attributable to the strong electrostatic interaction of a positively charged guanidino group in zanamivir with negatively charged amino acid residues in hNEU2. After we confirmed that the variation in the observed inhibitory activity among sialic acid analogues is excellently reproducible with the LERE-QSAR equation, we examined the reason for the remarkable difference between the inhibitory potencies of oseltamivir as to hNEU2 and influenza A virus neuraminidase-1 (N1-NA). Several amino acid residues in close contact with a positively charged amino group in oseltamivir are different between hNEU2 and N1-NA. FMO-IFIE (interfragment interaction energy) analysis showed that the difference in amino acid residues causes a remarkably large difference between the overall interaction energies of oseltamivir with hNEU2 and N1-NA. The current results will be useful for the development of new anti-influenza drugs with high selectivity and without the risk of adverse side effects.



1. INTRODUCTION

Neuraminidases (NAs, also called sialidases) are ubiquitous exoglycosylases that hydrolyze the terminal sialic acids of glycoproteins, glycopeptides, gangliosides, oligosaccharides, and polysaccharides and have been proved to play important roles in various biological processes through regulation of cellular sialic acid contents. NAs are widely distributed in nature, from viruses, and microorganisms such as bacteria, protozoa, and fungi to higher animals and humans.¹ Structural and functional studies on NAs have been widely reported so far. Most of them were on influenza virus NAs,^{2–4} a smaller number on human neuraminidases (hNEUs) having also been reported.^{5–8} The NA inhibitor first developed, 2-deoxy-2,3-didehydro-*N*-acetyl-neuraminic acid (Neu5Ac2en, DANA),⁹ was designed as a putative transition state analogue of sialic acid in the late 1960s. Although the inhibition by DANA is not selective for influenza virus NAs,^{10,11} DANA shows moderate inhibitory activity toward a variety of NAs, suggesting that amino acid residues in the active sites of NAs are highly conserved among a number of species including influenza virus and human NAs.

hNEUs are classified into four types,¹² the lysosomal neuraminidase (hNEU1),¹³ cytosolic neuraminidase (hNEU2),¹⁴

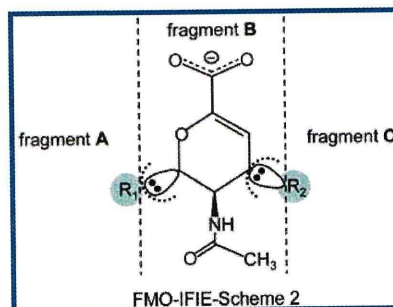
plasma membrane-associated neuraminidase (hNEU3),¹⁵ and lysosomal/mitochondrial membrane-associated neuraminidase (hNEU4),¹⁶ according to several characteristics, such as their subcellular distributions, enzymatic properties, and substrate specificities. They are involved in various cellular functions such as cell proliferation, differentiation, and apoptosis.¹⁷ hNEU1 is essential for the lysosomal catabolism of sialylated glycoconjugates,^{18–20} hNEU2 is able to recognize not only sialosyl linkages but also saccharide units close to sialic acids and the supramolecular organization of such as gangliosides, hNEU3 is expected to be involved in cell surface functions by modulating gangliosides, and hNEU4 is active against a broad range of sialylated glycoconjugates and has been implicated in the catabolism of glycolipids, but the details are not known at present. The crystallographic structure of hNEU2²¹ is the only available one among those of the four hNEUs.

As mentioned previously, DANA inhibits influenza virus NAs moderately but is not used as an anti-influenza drug because of its practically low potency. Through structure-based approaches

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Table 1. Chemical Structure and Inhibitory Potency of Compounds 1–5 against hNEU2



compound no.	name	R ₁ (fragment A)	R ₂ (fragment C)	hNEU2 log K _i ^a
1	IEM ^b	OCH ₂ CH(CH ₃) ₂	OH	-3.06
2	HEM ^c	OCH ₂ CH ₂ CH ₂ (OH ¹)	OH	-3.13
3	DEM ^d	OCH ₂ CH(OH ¹)CH ₂ (OH ²)	OH	-2.85
4	DANA ^e	CH(OH ¹)CH(OH ²)CH ₂ (OH ³)	OH	-3.85
5	zanamivir	CH(OH ¹)CH(OH ²)CH ₂ (OH ³)	NHC(=NH ₂ ⁺)NH ₂	-4.77

^a Taken from ref 33. ^b Isobutyl ether DANA mimetic. ^c 3-Hydroxypropyl ether DANA mimetic. ^d 2,3-Dihydroxypropyl ether DANA mimetic. ^e 2-Deoxy-2,3-didehydro-*N*-acetyl-neuraminic acid.

and the structure–activity relationships (SAR) of analogues of DANA, four anti-influenza drugs have been developed so far: zanamivir (Relenza),²² oseltamivir (Tamiflu),²³ peramivir (Rapiacta),²⁴ and laninamivir (Inavir).²⁵ These drugs have the potential to affect the activities of endogenous hNEUs, which can recognize similar substrates. Indeed, zanamivir and oseltamivir have been reported to inhibit hNEUs at the micromolar level or more, while they exhibit inhibitory activity in the nanomolar range against influenza virus NAs.²⁶ Over a thousand adverse events have been reported so far to have occurred with the use of oseltamivir, including a small number of rare but severe neuropsychiatric adverse events, especially in children.²⁷ Therefore, it is of great importance to elucidate the differences in sensitivity to these anti-influenza drugs among human and influenza virus NAs from the aspect of the inhibitory interaction mechanism at the atomic and electronic levels.

We previously reported correlation analyses of the interaction energies of influenza A virus neuraminidase-1 (N1-NA) with a series of eight sialic acid analogues including oseltamivir involving ab initio FMO (fragment molecular orbital)-IFIE (interfragment interaction energy) calculations at the Møller–Plesset (MP2) level.²⁸ Our previous study revealed that the dispersion and/or hydrophobic interaction energies of the alkoxy side chain in each ligand with several amino acid residues in N1-NA, which surrounds the former, govern the overall free-energy change in the binding interaction. Recently, we proposed a novel quantitative structure–activity relationship (QSAR) procedure called linear expression by representative energy terms (LERE) involving molecular calculations such as ab initio FMO and generalized Born/surface area (GB/SA) ones.^{29,30} The proposed LERE-QSAR procedure was demonstrated to provide detailed quantitative information as to the ligand–protein interaction of carbonic anhydrase with a series of substituted benzenesulfonamides at the atomic and electronic levels, which cannot be obtained with any other procedures.

There are a small number of compounds of which the inhibitory potency against hNEU2 have been measured and reported.^{26,31–34}

In this work, we constructed complexes of hNEU2 with six sialic acid analogues exhibiting relatively potent inhibitory activity, zanamivir, oseltamivir, DANA, and three analogues of DANA, by using molecular mechanics calculations and then carried out ab initio FMO and continuum solvation model calculations for each complex structure. Subsequently, we performed LERE-QSAR analysis and quantitatively examined the binding mechanism at the atomic and electronic levels. Finally, we discuss the observed difference between the sensitivities of oseltamivir to hNEU2 and N1-NA in terms of structure and interaction energy.

2. METHODS

2.1. Compound Set. Table 1 shows the chemical structures of four sialic acid analogues [IEM (compound 1), HEM, (2), DEM (3), and DANA (4)] and an anti-influenza drug, zanamivir (5), along with their inhibitory constants K_i against hNEU2.³³ Compounds 1–4 have a hydroxyl group as a common substituent (R₂, fragment C) and variable substituents (R₁, fragment A), hydroxyl groups existing in compounds 2–4. Compounds 4 and 5 have an identical substituent R₁, but compound 5 has a guanidino group (NHC(=NH₂⁺)NH₂) in fragment C, instead of the hydroxyl one of compounds 1–4. Thus, a total of five compounds with the common skeletal structure of 3-acetamido-3,4-dihydro-2*H*-pyran-6-carboxylic acid (fragment B) were subjected to analyses. Another anti-influenza drug, oseltamivir (compound 6), was added to the above compound set in Section 3.2.

2.2. Modeling of Complex Structures. The X-ray structures of complexes of hNEU2 with compounds 1 (PDB code: 2F11, complex 1), 2 (2F12, complex 2), 3 (2F13, complex 3), 4 (DANA, 1VCU, complex 4), and 5 (zanamivir, 2F0Z, complex 5) are all available. The structure of hNEU2 varies slightly among the crystallographic structures of complexes 1–5. The initial geometries of complexes 1–3 and 5 were taken from the crystallographic structure of complex 4 having a minimum number of missing residues, by replacing DANA in complex 4 with each of compounds 1–3 and 5. Loops A (Ile103–Thr122) and B

(Val265–Ser275) in hNEU2 are located in the vicinity of fragment A in compound 4. Loops A and B in the crystallographic structure of complex 5 occupy very close positions to those in complex 4, respectively. However, loops A and B in the crystallographic structure of complex 2 take on significantly different positions from those in complexes 4 and 5. Although several amino acid residues in loop A are not found (missing residues) in the crystallographic structures of complexes 1 and 3, the atomic positions of the corresponding amino acid residues other than the missing ones are nearly the same among the crystallographic structures of complexes 1, 2, and 3. Therefore, the initial geometries of complexes 1–3 were reconstructed by replacing loops A and B in the crystallographic structure of complex 4 with those in the crystallographic one of complex 2.

In our previous work,²⁸ we showed that two crystal water molecules inside the active site of N1-NA play an important role in the binding interaction: ligands having guanidino and amino groups in fragment C bind with only one water molecule (W1) and with two molecules (W1 and W2), respectively. The X-ray structures of nine complexes of hNEU2 with ligands (PDB codes: 2F11, 2F12, 2F13, 2F25 (dimer), 2F27 (dimer), and 1VCU (dimer)) were carefully examined. The root-mean-square deviation values for the atomic positions of W1 and W2 are both within 0.44 Å, indicating that their positions were nearly fixed. W1 and W2 for complexes 1–4 and W1 for complex 5 were placed at the respective average positions determined by the above procedure. Five and four sodium ions (Na⁺) were added to neutralize the total –5 and –4 charge of the complexes of hNEU2 with negatively charged ligands (compounds 1–4) and with neutral ones (compounds 5 and 6), respectively. Each complex was then solvated in a truncated octahedral box of TIP3P waters extending 12 Å from hNEU2.

Three-step energy minimizations were performed as follows: (i) hydrogen atoms, (ii) water molecules and counterions, and (iii) the entire system were sequentially relaxed for 2000, 6000, and 6000 steps, respectively. During these minimizations, the atomic positions of Glu111, Tyr179, Tyr181, Glu218, and Gln270, which could be involved in electrostatic and/or hydrogen-bonding interactions with fragment A in compounds 1–5, were constrained with a harmonic potential of 10 kcal/mol/Å². The finally obtained energy minimized complex structures were used in the following calculations. All molecular mechanics calculations were carried out using the AMBER10 package³⁵ with parameters of parm99³⁶ and general AMBER force field (GAFF)³⁷ for hNEU2 and each compound, respectively. The X-ray structures of compounds 1–5 bound with hNEU2 were subjected to partial geometry optimization (bond lengths and angles), fixing dihedral angles to the corresponding X-ray ones with HF/6-31G* calculations (Gaussian03 program³⁸). Similarly, that of compound 6 bound with N1-NA was geometry optimized. Then, partial atomic charges in each compound were determined according to the restrained electrostatic potential (RESP) fitting procedure^{39–41} with HF/6-31G* calculations. The atomic charges in compounds 1–6 are consistent with those of atoms in amino acid residues, because the latter charges were determined with the same procedure in the Amber (parm99) force field generally used for proteins.^{36,37} Atom names, atom types, and partial atomic charges of compounds 1–6 are listed in Table S1 (Supporting Information). The summation of partial atomic charges q within each fragment [$\sum q_i(X)$ (summation over atoms in fragment X), X = A, B, and C] for zanamivir

(compound 5) and oseltamivir (compound 6) is similar with that reported by Udommaneehanakit et al.⁴¹

2.3. LERE Formalism. We recently proposed LERE analysis involving molecular calculations such as the ab initio FMO and GB/SA ones and reported LERE-QSAR analyses of a series of substituted benzenesulfonamides with bovine carbonic anhydrase.^{29,30}

A basic assumption when dissecting free-energy terms is that they are all additive^{28,42,43}

$$\Delta G_{\text{obs}} = \Delta G_{\text{bind}} + \Delta G_{\text{sol}} + \Delta G_{\text{dis}} + \Delta G_{\text{others}} \quad (I)$$

ΔG_{obs} on the left-hand side of eq I is the overall free-energy change obtained from the observed inhibitory constant K_i : $\Delta G_{\text{obs}} = 2.303 RT \log K_i$ ($T = 310$ K). ΔG_{bind} , ΔG_{sol} , and ΔG_{dis} in eq I are the intrinsic interaction energy between an inhibitor and protein, the solvation free-energy change associated with complex formation, and the dissociation free-energy change of an inhibitor, respectively. ΔG_{others} , which represents the sum of free-energy terms such as the deformation energies of a protein and ligand other than ΔG_{bind} , ΔG_{sol} , and ΔG_{dis} , is assumed to be linear with the sum of representative free-energy terms ΔG_{bind} , ΔG_{sol} , and ΔG_{dis} in eq I, and ΔG_{others} is expected to act as a penalty term ($\beta < 0$), i.e., LERE approximation

$$\Delta G_{\text{others}} = \beta(\Delta G_{\text{bind}} + \Delta G_{\text{sol}} + \Delta G_{\text{dis}}) + \text{const (LERE approximation)} \quad (II)$$

During complex formation between compounds 1–5 and hNEU2, ΔG_{dis} does not need to be considered in the current case, because a carboxylate group common to compounds 1–5 and a guanidino one in compound 5 take on nearly complete ionized forms [COO[–] ($pK_a \sim 2.4$) and NHC(=NH₂⁺)NH₂ ($pK_a \sim 13$), respectively] before and after complex formation at experimental pH = 5.6.^{33,44}

ΔG_{obs} very probably exhibits a ubiquitous energetic relation on complex formation between a protein and a series of ligands having a similar structure, the entropy–enthalpy compensation rule^{45–48}

$$T\Delta S_{\text{obs}} = \alpha\Delta H_{\text{obs}} + \text{const (entropy–enthalpy compensation)} \quad (IIa)$$

The value of α ($\alpha > 0$) on complex formation between a sialic acid analogue and hNEU2 is supposed to be within the range of 0.70 ± 0.20 , judging from the results of isothermal titration calorimetry (ITC) experiments on complex formation of various sugars and sugar mimetics with their binding proteins.^{49–52} Frederick et al.⁵³ showed that a change in the protein conformational entropy (ΔS_{bind} , which includes changes in the vibrational, rotational, and translational entropies) is linearly correlated with a change in overall entropy on the formation of complexes of calmodulin with its binding peptides, using nuclear magnetic resonance (NMR) relaxation methods. They proposed the possibility that ΔS_{bind} makes a major contribution to the overall binding entropy change. In the case where the complex formation is mostly driven by the electrostatic interaction energy between a charged ligand and protein, the intrinsic binding enthalpy term ΔH_{bind} and the polar component of solvation free-energy change $\Delta G_{\text{sol}}^{\text{pol}}$, calculated with continuum solvation models such as the generalized Born (GB) and Poisson–Boltzmann (PB) ones, are probably major contributors to the overall free-energy changes. Avbelj et al.⁵⁴ recently showed that the observed solvation (hydration) free-energy changes of the polar parts in several dipeptides consist mostly of the enthalpy term arising

from the electrostatic interaction, the energy of which can be well reproduced with PB type calculation. Directly from the original Born equation: $\Delta G_{\text{sol}}^{\text{pol}} = -(q^2/2r)(1 - 1/\epsilon)$, the enthalpy change is given by the expression: $\Delta H_{\text{sol}} = \Delta G_{\text{sol}}^{\text{pol}} - T(\partial G_{\text{sol}}^{\text{pol}}/\partial T)_P = -(q^2/2r)[1 - 1/\epsilon - (T/\epsilon^2)(\partial\epsilon/\partial T)_P]$, where r , q , ϵ , and T are the cavity radius formed by the ion in a particular solvent, ion charge, dielectric constant, and temperature, respectively.^{55,56} The reported value of $[(T/\epsilon)(\partial\epsilon/\partial T)_P]$ for water ($\epsilon = 78.4$) at 298 K is -1.357 ,⁵⁵ giving 0.983 for the ratio between $\Delta G_{\text{sol}}^{\text{pol}}$ and ΔH_{sol} ($\Delta G_{\text{sol}}^{\text{pol}}/\Delta H_{\text{sol}} = 0.983$). The ratio close to unity suggests that ΔH_{sol} is generally replaceable by $\Delta G_{\text{sol}}^{\text{pol}}$ with satisfactory accuracy. We showed the possibility that the energy term (E_{sol}) calculated with the ab initio MO-continuum solvation model (COSMO) corresponds to the effective enthalpic energy term associated with the partition equilibrium between the aqueous and organic solvent phases rather than the overall solvation free-energy term, when we analyzed more than 200 log $P_{\text{sol/w}}$ values.⁵⁷

ΔH_{bind} can be replaced by ΔE_{bind} , because of the lack of significant volume and pressure changes in solution. ΔE_{bind} consists of $\Delta E_{\text{bind}}^{\text{HF}}$ and E^{disp} , which are the intrinsic binding energy at the Hartree–Fock (HF) level and the dispersion (van der Waals) energy, respectively. ΔG_{sol} consists of $\Delta G_{\text{sol}}^{\text{pol}}$ and the nonpolar free-energy change $\Delta G_{\text{sol}}^{\text{nonpol}}$. $\Delta E_{\text{bind}}^{\text{HF}}$ and $\Delta G_{\text{sol}}^{\text{pol}}$ are considered to be “electrostatic energy terms”⁵⁸ involved in the electrostatic interactions among a ligand, protein, and solvent medium. E^{disp} and $\Delta G_{\text{sol}}^{\text{nonpol}}$ are considered to be “local energy terms”, which are effective only at a short distance. We reported that $\Delta G_{\text{sol}}^{\text{pol}}$ shows an excellent anticorrelation with $\Delta E_{\text{bind}}^{\text{HF}}$ during complex formation between carbonic anhydrase and a series of substituted benzenesulfonamides.^{29,30} We assumed that the sum of the electrostatic energy terms represents the effective overall enthalpic change, i.e., the effective binding enthalpy change in a solvent, $\Delta H_{\text{bind}}^{\text{sol}} = \Delta E_{\text{bind}}^{\text{HF}} + \Delta G_{\text{sol}}^{\text{pol}}$, on the condition that contributions from the local energy terms to $\Delta H_{\text{bind}}^{\text{sol}}$ are significantly smaller than those from the electrostatic ones. This condition is equivalent to that the statistical variances of E^{disp} and $\Delta G_{\text{sol}}^{\text{nonpol}}$ are negligibly smaller than those of $\Delta E_{\text{bind}}^{\text{HF}}$ and $\Delta G_{\text{sol}}^{\text{pol}}$ in a correlation equation. The entropy–enthalpy compensation is now expressed as

$$T\Delta S_{\text{bind}}^{\text{sol}} = \alpha(\Delta E_{\text{bind}}^{\text{HF}} + \Delta G_{\text{sol}}^{\text{pol}}) + \text{const} \quad (\text{III})$$

Assuming eq III, the effective overall free-energy change $\Delta G_{\text{bind}}^{\text{sol}}$ is expressed as $(1 - \alpha)(\Delta E_{\text{bind}}^{\text{HF}} + \Delta G_{\text{sol}}^{\text{pol}}) + \text{const}$. Although the protein conformational entropic term ($T\Delta S_{\text{bind}}$) is often estimated using normal mode and molecular dynamics calculations, the calculated one is known to exhibit significant large uncertainties.^{59–61} Alternatively, eqs I, II, and III yield eq IV without estimation of $T\Delta S_{\text{bind}}$ ^{62,63} directly

$$\Delta G_{\text{obs}} = (1 + \beta)(1 - \alpha)(\Delta E_{\text{bind}}^{\text{HF}} + \Delta G_{\text{sol}}^{\text{pol}}) + \text{const} \quad (\text{IV})$$

Equation IV appears to take a similar form to the solvated interaction energy (SIE) model proposed by Naïm et al.⁶⁴ Although they introduced several adjusting parameters such as the effective interior dielectric constant in order to adjust the calculated overall free-energy change to the observed one, eq IV has only two scaling factors, α and β , which effectively and naturally consider the entropy–enthalpy compensation and penalty effects, respectively. It is obvious but noteworthy that the α and β values depend on a set of a series of ligands and their target proteins. Also, it should be noted that the above procedure is probably applicable

to a congeneric series of ligands having the same skeleton but probably not to a set containing structurally diverse ones.

2.4. Calculation of Representative Energy Terms. $\Delta E_{\text{bind}}^{\text{HF}}$ ($= E^{\text{HF}}(\text{complex}) - [E^{\text{HF}}(\text{protein}) + E^{\text{HF}}(\text{ligand})]$) and E^{disp} ($= \Delta E_{\text{bind}}^{\text{MP2}} - \Delta E_{\text{bind}}^{\text{HF}}$) were calculated using ab initio fragment molecular orbital (FMO) calculations with one residue (including W1 and W2) per FMO-fragment partition (FMO-IFIE-Scheme 1)^{65–68} at the HF and MP2/6-31G levels, respectively, using the ABINIT-MP program.^{69,70} Our previous studies involving FMO calculations^{28–30,71,72} demonstrated that interfragment interaction energy (IFIE) analysis can provide valuable information independent from that obtained with the LERE-QSAR one. In the present work, the structures of compounds 1–5 were further divided into three FMO-fragments A, B, and C (FMO-IFIE-Scheme 2, shown in Table 1),^{28,73–75} in order to clarify the contribution of each fragment to the overall interaction energy. Amino acid residues, which are closely located in fragments A, B, and C, are denoted by pockets A, B, and C, respectively. IFIE^{HF} (fragment X, Y) denotes the IFIE value between X (fragment X in a ligand) and Y (amino acid residue Y in a protein) at the HF level, and $\Sigma\text{IFIE}^{\text{HF}}$ (fragment X, pocket Z) represents summation of IFIE^{HF} (fragment X, Y) over all the amino acid residues Y in pocket Z. As expected,^{28,29,71,72} $\Delta E_{\text{bind}}^{\text{HF}}$ is almost completely linear with $\Sigma\Sigma\text{IFIE}^{\text{HF}}$ (fragment X, pocket Z) (summation over X, Z = A, B, and C) for compounds 1–5 ($r = 0.999$). This fact guarantees that the total intrinsic binding energy can be decomposed into local fragment–pocket interaction energies. The solvation free-energy terms $\Delta G_{\text{sol}}^{\text{pol}}$ and $\Delta G_{\text{sol}}^{\text{nonpol}}$ were calculated with the molecular dynamics (MD)-PB/SA (Poisson–Boltzmann/surface area) and MD-GB/SA (generalized Born/surface area) procedures^{76,77} included in the AMBER10 package. These terms were obtained as ensemble averages of snapshot structures in the MD trajectory, $\langle G_{\text{sol}}^{\text{pol}}(\text{PB}) \rangle$, $\langle G_{\text{sol}}^{\text{pol}}(\text{GB}) \rangle$, and $\langle G_{\text{sol}}^{\text{nonpol}} \rangle$. It is necessary to average G_{sol} , because $G_{\text{sol}}^{\text{pol}}$ is supposed to be sensitive to subtle conformational changes occurring in domains other than the binding one.^{62,63,78,79} Short-time MD with 200 ps of a production run at 300 K was performed to refine each complex structure (structure annealing), and snapshot structures were collected for the last 150 ps trajectory at 3.0 ps intervals. We confirmed that $\langle \Delta G_{\text{sol}}^{\text{pol}}(\text{PB}) \rangle$ and $\langle \Delta G_{\text{sol}}^{\text{pol}}(\text{GB}) \rangle$ were not significantly changed during a longer simulation time, by performing 1 and 10 ns simulations for (1) the complex of hNEU2 with compound 4 and (2) that of N1-NA with compound 6, respectively. The absolute differences of $\langle \Delta G_{\text{sol}}^{\text{pol}} \rangle$ between the 200 ps and longer time simulations are less than 1.4 and 3.3 kcal/mol for (1) and (2), respectively.

3. RESULTS AND DISCUSSION

3.1. LERE-QSAR Analysis of Complex Formation of hNEU2 with Sialic Acid Analogues and Zanamivir. Table 2 lists the observed and calculated overall free-energy changes along with the electrostatic interaction energy terms $\Delta E_{\text{bind}}^{\text{HF}}$ and $\langle \Delta G_{\text{sol}}^{\text{pol}} \rangle$, and the local interaction energy terms E^{disp} and $\langle \Delta G_{\text{sol}}^{\text{nonpol}} \rangle$, of compounds 1–5 listed in Table 1. Figure 1 shows the variations in these energy terms. The variances of $\Delta E_{\text{bind}}^{\text{HF}}$ and $\langle \Delta G_{\text{sol}}^{\text{pol}}(\text{PB}) \rangle$, which mainly arise from the electrostatic interaction, are considerably large (3989 and 1967 kcal²/mol², respectively). On the other hand, those of E^{disp} and $\langle \Delta G_{\text{sol}}^{\text{nonpol}} \rangle$ are much smaller (5.06 and 0.01 kcal²/mol², respectively). Thus, the variation in overall free-energy change is governed by $\Delta E_{\text{bind}}^{\text{HF}}$ and $\langle \Delta G_{\text{sol}}^{\text{pol}}(\text{PB}) \rangle$, satisfying the condition for eq IV.

Table 2. Overall Free-Energy Change ΔG and Representative Energy Terms^a (hNEU2)

compound									
no.	name	ΔG_{obs}^b	ΔG_{calc}^c	ΔG_{calc}^d	$\Delta E_{\text{bind}}^{\text{HF}}$	$\langle \Delta G_{\text{sol}}^{\text{pol (PB)}} \rangle^e$	$\langle \Delta G_{\text{sol}}^{\text{pol (GB)}} \rangle^e$	E^{disp}	$\langle \Delta G_{\text{sol}}^{\text{nonpol}} \rangle^{e,f}$
1	IEM	-4.33	-4.03	-3.95	-78.27	79.45 (6.79)	67.62 (7.06)	-38.88	-5.00 (0.05)
2	HEM	-4.44	-4.44	-4.44	-85.77	78.82 (4.76)	67.35 (5.31)	-37.32	-4.99 (0.05)
3	DEM	-4.05	-4.30	-4.41	-98.06	93.87 (7.66)	80.17 (8.50)	-40.34	-5.08 (0.06)
4	DANA	-5.47	-5.65	-5.86	-154.28	123.44 (9.27)	113.31 (8.93)	-36.81	-4.87 (0.11)
variance ^g					889.36	327.25	351.25	1.92	0.01
5	zanamivir	-6.77	-6.62	-6.38	-247.22	197.14 (7.71)	197.98 (6.05)	-43.03	-5.22 (0.05)
variance ^g					3989.09	1967.09	2429.14	5.06	0.01

^a In kcal/mol. ^b $\Delta G_{\text{obs}} = 2.303 RT \log K_i$ ($T = 310 \text{ K}$). ^c Calculated from eq 1. ^d Calculated from eq 2. ^e Average value and standard deviation (in parentheses). ^f $\Delta G_{\text{sol}}^{\text{nonpol}} = \gamma \Delta \text{ASA}$, where ASA is the water accessible surface area and γ is taken to be $0.0072 \text{ kcal/mol}/\text{\AA}^2$. ^g In $\text{kcal}^2/\text{mol}^2$.

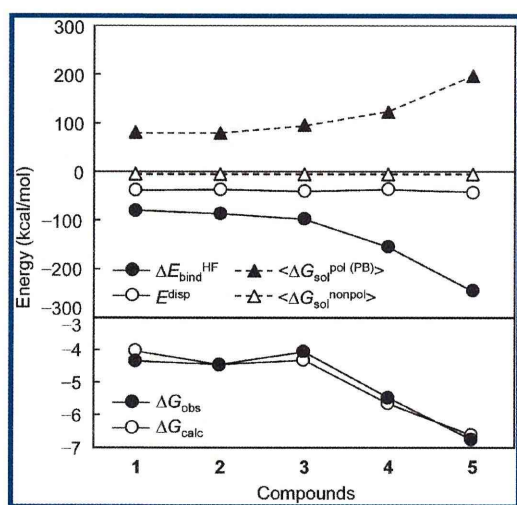


Figure 1. Variations of $\Delta E_{\text{bind}}^{\text{HF}}$, $\langle \Delta G_{\text{sol}}^{\text{pol (PB)}} \rangle$, E^{disp} , $\langle \Delta G_{\text{sol}}^{\text{nonpol}} \rangle$, ΔG_{obs} , and ΔG_{calc} values (hNEU2). ΔG_{calc} is calculated by eq 1.

The correlation coefficient between $\Delta E_{\text{bind}}^{\text{HF}}$ and $\langle \Delta G_{\text{sol}}^{\text{pol (PB)}} \rangle$ is -0.996 for compounds 1–5 (-0.988 for compounds 1–4). This means that these two energy terms are dependent on and compensate for each other. The variance of $\Delta E_{\text{bind}}^{\text{HF}}$ for compounds 1–5 is $\sim 200\%$ larger than that of $\langle \Delta G_{\text{sol}}^{\text{pol (PB)}} \rangle$ ($\sim 270\%$ for compounds 1–4), and the difference between $|\Delta E_{\text{bind}}^{\text{HF}}|$ and $\langle \Delta G_{\text{sol}}^{\text{pol}} \rangle$ is distinctly large for compounds 4 (DANA) and 5 (zanamivir). That is, the stabilization due to $\Delta E_{\text{bind}}^{\text{HF}}$ overwhelms the destabilization due to $\langle \Delta G_{\text{sol}}^{\text{pol}} \rangle$. The sum of $\Delta E_{\text{bind}}^{\text{HF}}$ and $\langle \Delta G_{\text{sol}}^{\text{pol}} \rangle$ is excellently correlated with ΔG_{obs} , as shown in eqs 1 and 2

$$\Delta G_{\text{obs}} = (1 + \beta)(1 - \alpha)(\Delta E_{\text{bind}}^{\text{HF}} + \langle \Delta G_{\text{sol}}^{\text{pol (PB)}} \rangle) - 4.09$$

$$n = 5, r = 0.979, s = 0.265, F = 68.3, (1 + \beta)(1 - \alpha) = 0.0505 \quad (1)$$

$$\Delta G_{\text{obs}} = (1 + \beta)(1 - \alpha)(\Delta E_{\text{bind}}^{\text{HF}} + \langle \Delta G_{\text{sol}}^{\text{pol (GB)}} \rangle) - 3.28$$

$$n = 5, r = 0.940, s = 0.439, F = 22.9, (1 + \beta)(1 - \alpha) = 0.0630 \quad (2)$$

Although the statistical quality of eq 1 using PB is slightly better than that of eq 2 using GB, it is probably difficult and mostly pointless to discuss the difference, because each of the two

computing procedures, PB and GB, involves several approximations and parameters. The correlation coefficient between $\langle \Delta G_{\text{sol}}^{\text{pol (PB)}} \rangle$ and $\langle \Delta G_{\text{sol}}^{\text{pol (GB)}} \rangle$ is 0.999, clearly showing that these two $\langle \Delta G_{\text{sol}}^{\text{pol}} \rangle$ values are equivalent in terms of statistical correlation, at least. When compound 5, which shows the largest $|\Delta E_{\text{bind}}^{\text{HF}}|$ and $|\langle \Delta G_{\text{sol}}^{\text{pol}} \rangle|$, is removed ($n = 4$), the statistical qualities of eqs 1 and 2 are poorer but still statistically significant ($r = 0.941$ and 0.914 , respectively). The absolute values of intercept in eqs 1 and 2 are small, 4.09 and 3.28 , respectively. The β values in eqs 1 and 2 are similar, -0.83 and -0.79 , respectively, when α is assumed to be 0.70 .^{49–52} These results indicate that the penalty energy term ΔG_{others} in eq 11 is nearly proportional to $(\Delta E_{\text{bind}}^{\text{HF}} + \langle \Delta G_{\text{sol}}^{\text{pol}} \rangle)$ and makes a negative contribution to the observed overall free-energy change ΔG_{obs} , as we expected. As noted above, there is an excellent anticorrelation between $\Delta E_{\text{bind}}^{\text{HF}}$ and $\langle \Delta G_{\text{sol}}^{\text{pol}} \rangle$, and the variation of the former overwhelms that of the latter in the current case.

We then attempted to determine the reason why compound 5 has the smallest $\Delta E_{\text{bind}}^{\text{HF}}$ value among compounds 1–5 in terms of structure and interaction. As shown in Table 2, the large variance in $\Delta E_{\text{bind}}^{\text{HF}}$ ($889 \text{ kcal}^2/\text{mol}^2$) among compounds 1–4 arises from differences in fragment A, because these compounds are different only in fragment A. We carefully examined $\Delta E_{\text{bind}}^{\text{HF}}$ arising from interaction of fragment A with pocket A in hNEU2. Figure 2 shows the IFIE^{HF} values of fragment A with amino acid residues in pocket A. Pocket A in hNEU2 consists of Glu111, Tyr179, Tyr181, Leu217, Glu218, and Gln270. The side chain of Glu218 is located in close to fragment A (R_1) but does not form a stable hydrogen-bond with hydroxyl groups in fragment A of compounds 2–4. The weak stabilization of IFIE^{HF} (fragment A, Glu218) commonly found in compounds 1–4 is probably due to the charge (electron) transfer (CT) from negatively charged Glu218 to neutral fragment A in compounds 1–4. In addition to the effect of CT, weak hydrogen-bonding/electrostatic interactions between hydroxyl groups on fragment A in compounds 2–4 and oxygen atoms in the side chain of Glu218 are possibly involved. The IFIE^{HF} (fragment A, pocket A) values for compounds 1 and 2 indicate that fragment A in these compounds does not interact with pocket A effectively. The negative IFIE^{HF} (fragment A/compound 3, Tyr179) value is due to a weak hydrogen-bonding interaction between the side chain of Tyr179 and the OH¹ group in fragment A (distance ($\text{H}^1\text{O} \cdots \text{HO} \text{---} \text{Tyr179}$) = 2.04 \AA). The OH¹ group attached to fragment A in compound 4 shows remarkably high attractive interaction energy with Glu111, because the OH¹ group and oxygen atoms in the side chain of Glu111 take on atomic positions optimum for effective

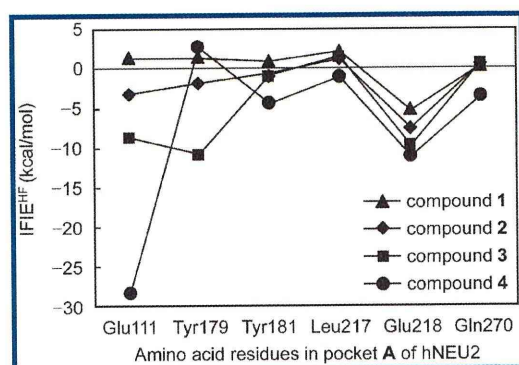


Figure 2. IFIE^{HF} values of fragment A in compounds 1–4 with amino acid residues in pocket A of hNEU2.

hydrogen-bonding and electrostatic interactions [distance ($H^1 \cdots O - Glu111$) = 1.54 Å, angle ($O - H^1 \cdots O - Glu111$) = 176 deg, and angle ($H^1 \cdots O - C_\delta - Glu111$) = 116 deg].

The variance analysis of the IFIE^{HF} value between residue X and fragment A in a ligand among compounds 1–4 (denoted as var.[IFIE^{HF} (fragment A, X)]) shows that var.[IFIE^{HF} (fragment A, Glu111)], var.[IFIE^{HF} (fragment A, Tyr179)], and var.[IFIE^{HF} (fragment A, Arg237)] are 128.7, 27.1, and 24.8 kcal²/mol², respectively, and that var.[IFIE^{HF} (fragment A, X)] values other than the above three ones are less than 5.5 kcal²/mol². This result clearly suggests that these three residues govern the overall variation in the electrostatic interaction energy among compounds 1–4. Glu111 is a key residue which interacts with the OH¹ group (fragment A) in compound 4 through the electrostatic and/or hydrogen-bonding interactions. In addition, Arg237 (pocket B) in close proximity to fragment A significantly contributes to the stabilization of the electrostatic interaction energy with compound 4, but not with compounds 1–3. In fact, IFIE^{HF} (fragment A, Arg237) of compound 4 is 10–13 kcal/mol smaller than those of compounds 1–3. This is because the OH² group in compound 4 only takes favorable position for the electrostatic interaction with the positively charged side chain of Arg237.

Fragment A in compound 4 undergoes the largest stabilization through the hydrogen-bonding and electrostatic interactions. Consequently, compound 4 exhibits the most potent inhibitory activity among compounds 1–4.

The above results quantitatively confirm that the hydrogen-bonding interaction energy of fragment A in compounds 1–4 with residues in pocket A mostly determines the observed overall free-energy change and suggest that the position of the hydroxyl group in fragment A is crucial for the optimum hydrogen-bond formation with residues in pocket A. Although the interaction energy of fragment A with amino acid residues in pocket A is not the most important contributor to the stabilization of the total binding energy, it should be noted that the structural factors that stabilize the total binding energy are different from those that govern the variation in the total binding energy among compounds 1–4.

Next, we examined the most potent compound 5 (zanamivir), which has the same skeletal structure (fragment B) as compounds 1–4 but has a positively charged guanidino group instead of a neutral hydroxyl group (fragment C). Notably, compounds 4 and 5 have the same fragment A as well as fragment B structure. Differences in the interaction energy between compounds 4 and 5 are unquestionably attributable to those in fragment C.

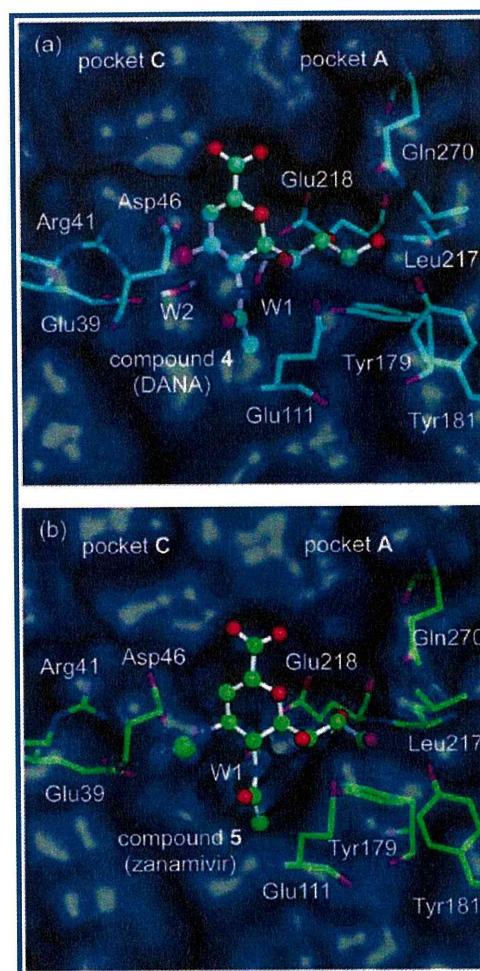


Figure 3. (a) Compound 4 (DANA) and (b) compound 5 (zanamivir) accommodated in the active site of hNEU2.

Figure 3 (a) and (b) shows the structures of compounds 4 and 5 bound at the active site of hNEU2, respectively. The guanidino group (fragment C) in compound 5 is involved in electrostatic interactions with Glu39 and Asp46 in hNEU2, and the corresponding hydroxyl group in compound 4 forms a hydrogen-bond with Arg41.

As can be seen in Figure 4, the IFIE^{HF} (fragment C/compound 5, Glu39) and IFIE^{HF} (fragment C/compound 5, Asp46) values represent large negative energies, and the IFIE^{HF} (fragment C/compound 5, Arg41) value is the positive energy due to the electrostatic repulsive interaction between the positively charged guanidino group and the side chain of Arg41. The relatively small negative IFIE^{HF} (fragment C/compound 4, Arg41) value represents the hydrogen-bonding interaction, which causes considerably weaker stabilization than electrostatic interactions of the guanidino group in compound 5 with Glu39 and Asp46. In total, $\Delta E_{\text{bind}}^{\text{HF}}$ of compound 5 is 93 kcal/mol more stable than that of compound 4.

3.2. Comparison of Inhibitory Activities of Anti-Influenza Drugs against hNEU2 and N1-NA. Although anti-influenza drugs zanamivir (compound 5) and oseltamivir (compound 6) exhibit excellent inhibitory activity toward various types of influenza virus NAs including N1-NA, their activity toward

hNEU2 is much lower. It is notable that compound **6** has a skeletal structure (fragment **B**) different from that in compounds **1–5**. Table 3 shows the ΔG_{obs} values of compounds **5** and **6** with hNEU2 and N1-NA. The large difference in ΔG_{obs} between compound **5** with hNEU2 and N1-NA as well as that between compound **6** with hNEU2 and N1-NA can be attributed to those between the binding interaction modes of hNEU2 and N1-NA with the two compounds.

The whole protein structures of hNEU2 and N1-NA take on a close topology: six bladed β -propeller folds commonly conserved among neuraminidases are arranged at topologically almost equivalent positions. Although the overall amino acid sequence homology between hNEU2 and N1-NA is considerably low (identity $\sim 16\%$, similarity $\sim 25\%$), several residues in their active sites seem to be conserved: an “arginine triad” (Arg21, 237, and 304 in hNEU2; Arg118, 292, and 371 in N1-NA) and tyrosine (Tyr334 in hNEU2; Tyr406 in N1-NA), which are essential for expression of their catalytic activities. We carefully compared the atomic positions of amino acid residues located in the active sites of the two PDB crystal structures, 1VCU (hNEU2)²¹ and 2HU4 (N1-NA).⁸² Each of the above corresponding pairs of residues takes on nearly the same position in hNEU2 and N1-NA (within 0.76 Å (C_{α} atoms)). Figure 5 (a) and (b) schematically

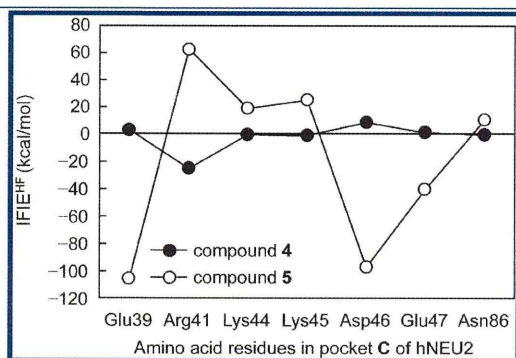


Figure 4. IFIE^{HF} values of fragment C in compounds **4** (DANA) and **5** (zanamivir) with amino acid residues in pocket C of hNEU2.

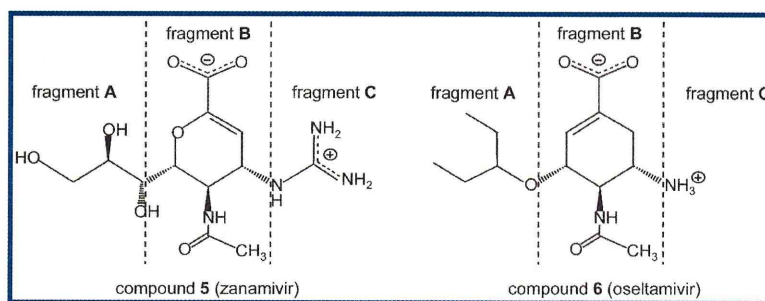
represents the interactions of compound **6** with hNEU2 and N1-NA, respectively.

In pocket **B**, one Tyr and three Arg residues (arginine triad) common in hNEU2 and N1-NA surround a negatively charged carboxylate group in fragment **B** of compound **6**, exhibiting a very close geometry.

A few amino acid residues in pocket **A** are different between hNEU2 and N1-NA. For example, a negatively charged Glu111 in hNEU2 is replaced by nonpolar Ile222 in N1-NA. Although fragment **A** in compound **5** as well as compound **4** form a hydrogen-bond with residues in pocket **A** of hNEU2, fragment **A** in compound **6** is not able to form a hydrogen-bond because it does not have a hydroxyl group. Both compounds **5** and **6** exhibit potent inhibitory potency as to N1-NA, probably because the hydrogen-bonding interaction between fragment **A** and pocket **A** in N1-NA is not dominant, but the dispersion type of interaction (E^{disp} and $\Delta G_{\text{sol}}^{\text{nonpol}}$) governs the interaction energy between fragment **A** and pocket **A** in N1-NA, as demonstrated in our previous study.²⁸

We then examined the differences between the IFIE^{HF} (fragment C (NH_3^+)/compound **6**, pocket C/hNEU2) and IFIE^{HF} (fragment C/compound **6**, pocket C/N1-NA) values. Figure 6 shows the IFIE^{HF} (fragment C/compound **6**, pocket C/hNEU2) and IFIE^{HF} (fragment C/compound **6**, pocket C/N1-NA) values. The $\Sigma \text{IFIE}^{\text{HF}}$ (fragment C, pocket C/hNEU2) and $\Sigma \text{IFIE}^{\text{HF}}$ (fragment C, pocket C/N1-NA) values are mostly due to those of fragment C with ionized residues: negatively charged Glu39 and Asp46 and positively charged Arg41 in hNEU2 and negatively charged Glu119, Asp151, and Glu227 and positively charged Arg156 in N1-NA. Asp46 in hNEU2 and Asp151 in N1-NA stabilize, and Arg41 in hNEU2 and Arg156 in N1-NA destabilize the interaction energy. The above stabilization and destabilization commonly occur in the two complexes. Ile22, Glu39, Met85, and Asn86 in pocket C of hNEU2 are replaced by Glu119, Leu134, Trp178, and Glu227 in N1-NA, respectively. Among the above replacements, that from neutral Ile22 in hNEU2 to negatively charged Glu119 in N1-NA makes the interaction energy of fragment C with N1-NA 105 kcal/mol more stable than that with hNEU2. Similarly, the replacement of Asn86 with Glu227 makes the interaction energy 54 kcal/mol

Table 3. Chemical Structure of Zanamivir and Oseltamivir along with Inhibitory Potency against hNEU2 and N1-NA



compound		ΔG_{obs}^a			
no.	name	R_1 (fragment A)	R_2 (fragment C)	hNEU2 ^b	N1-NA ^c
5	zanamivir	$\text{CH}(\text{OH}^1)\text{CH}(\text{OH}^2)\text{CH}_2(\text{OH}^3)$	$\text{NHC}(=\text{NH}_2^+)\text{NH}_2$	-6.77	-12.80
6	oseltamivir	$\text{OCH}(\text{CH}_2\text{CH}_3)_2$	NH_3^+	-3.26	-13.26

^a In kcal/mol. ^b $\Delta G_{\text{obs}} = 2.303 RT \log K_i$ ($T = 310 \text{ K}$). K_i value is taken from ref 33. ^c $\Delta G_{\text{obs}} = 2.303 RT \log (\text{IC}_{50})$ ($T = 310 \text{ K}$). IC_{50} value is taken from ref 81.

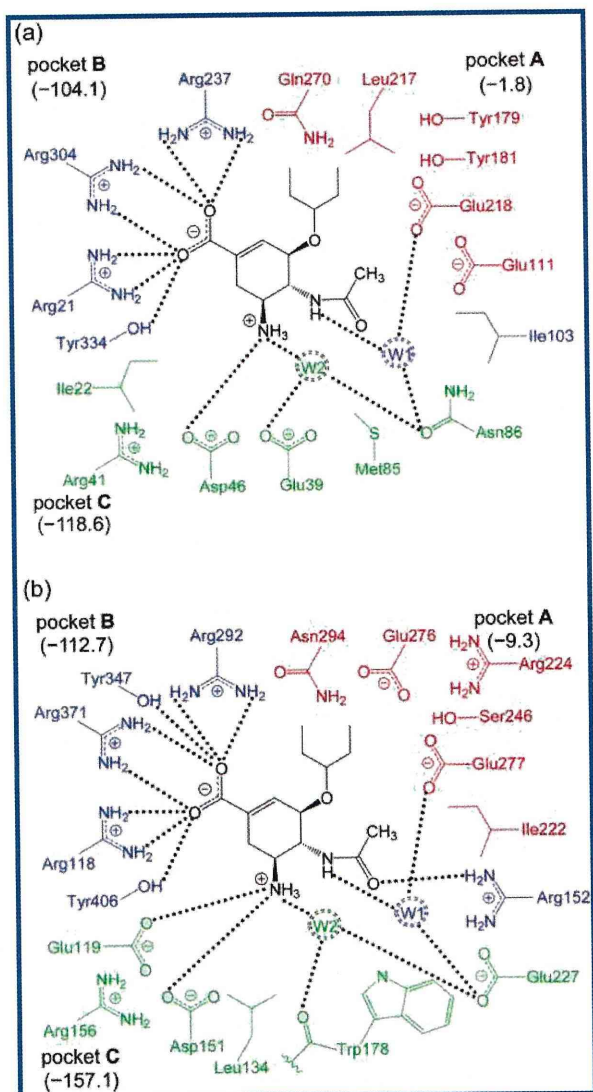


Figure 5. Interaction of compound 6 (oseltamivir) with amino acid residues in the active sites of (a) hNEU2 and (b) N1-NA. $\Sigma\text{IFIE}^{\text{HF}}$ (fragment X, pocket X: X = A (red), B (blue), and C (green)) values are shown in parentheses (kcal/mol). W1 and W2 represent water molecules in the active sites. A dotted line represents an attractive interaction.

more stable. Conversely, the replacement of negatively charged Glu39 with neutral Leu134 stabilizes the complex of hNEU2 with compound 6 by 70 kcal/mol. The energy differences $\Delta\Sigma\text{IFIE}^{\text{HF}}$ ($= \Sigma\text{IFIE}^{\text{HF}}$ (fragment C/compound 6, pocket C/hNEU2) $- \Sigma\text{IFIE}^{\text{HF}}$ (fragment C/compound 6, pocket C/N1-NA)) reaches a stabilization value of 38 kcal/mol.

In summary, compound 6 shows remarkably less inhibitory activity toward hNEU2 than that toward N1-NA, because of differences in pockets A and C between the two proteins.

We finally examined the difference between the inhibitory activities of compounds 5 and 6 with hNEU2. In our previous paper,²⁸ we showed that $\Sigma\text{IFIE}^{\text{HF}}$ (fragment C/compound 6, pocket C/N1-NA) is not far from $\Sigma\text{IFIE}^{\text{HF}}$ (fragment C in the compound where the amino group in compound 6 is replaced by a guanidino one, pocket C/N1-NA) (-157 and -145 kcal/mol, respectively). However, there is a significantly large difference

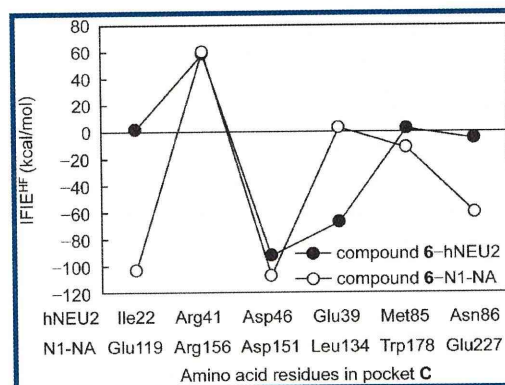


Figure 6. IFIE^{HF} values of fragment C in compound 6 (oseltamivir) with amino acid residues in pocket C of hNEU2 and N1-NA.

between $\Sigma\text{IFIE}^{\text{HF}}$ (fragment C (NH_3^+)/compound 6, pocket C/hNEU2) and $\Sigma\text{IFIE}^{\text{HF}}$ (fragment C ($\text{NHC}(=\text{NH}_2^+)\text{NH}_2$)/compound 5, pocket C/hNEU2). This difference between $\Sigma\text{IFIE}^{\text{HF}}$ (fragment C/compound 6, pocket C/hNEU2) and $\Sigma\text{IFIE}^{\text{HF}}$ (fragment C/compound 5, pocket C/hNEU2) (-119 and -149 , respectively) mostly arises from the interaction energy of fragment C with Glu39 [IFIE^{HF} (fragment C/compound 6, Glu39) and IFIE^{HF} (fragment C/compound 5, Glu39) are -68 (see Figure 6) and -106 (Figure 4) kcal/mol, respectively]. This is because the positively charged guanidino group in compound 5 is located closer to negatively charged Glu39 in hNEU2 than the amino group in compound 6. The nearest interatomic (heavy atoms) distances between Glu39 and fragment C are 2.76 and 4.48 Å for compounds 5 and 6, respectively. The above differences as well as those in the interactions between fragment A and pocket A make the inhibitory activity of compound 6 toward hNEU2 considerably less potent than that of compound 5.

4. SUMMARY AND CONCLUSION

In our previous study,²⁸ we showed that the variation in the inhibitory potency as to N1-NA among eight analogues (including oseltamivir) having alkoxy side chains ($\text{OC}_n\text{H}_{2n+1}$, $n = 0, 3, 4$, and 5) at R_1 (fragment A) is governed by the dispersion (E^{disp}) and/or hydrophobic interaction ($\Delta G_{\text{sol}}^{\text{nonpol}}$) energies. In the present study, we showed that the observed overall free-energy changes of DANA, three analogues of DANA, and an anti-influenza drug zanamivir, all of which have a common skeletal structure (fragment B), are linearly correlated with the calculated one using FMO and MD-PB(GB)/SA calculations (LERE-QSAR). Unlike the case reported in the previous study, the statistical variances of the local energy terms E^{disp} and $\langle \Delta G_{\text{sol}}^{\text{nonpol}} \rangle$ are both 3–6 orders of magnitude smaller than those of the electrostatic energy terms $\Delta E_{\text{bind}}^{\text{HF}}$ and $\langle \Delta G_{\text{sol}}^{\text{pol}} \rangle$. Because of penalty and entropy–enthalpy compensation effects, the coefficient of ($\Delta E_{\text{bind}}^{\text{HF}} + \langle \Delta G_{\text{sol}}^{\text{pol}} \rangle$) in the correlation equations eqs 1 and 2 is considerably smaller than unity (~ 0.05). There is an excellent anticorrelation between the electrostatic energy terms $\Delta E_{\text{bind}}^{\text{HF}}$ and $\langle \Delta G_{\text{sol}}^{\text{pol}} \rangle$ ($r = \sim -0.99$). Comparison of the variances of the two electrostatic energy terms showed that the contribution (stabilization) from the intrinsic binding energy term $\Delta E_{\text{bind}}^{\text{HF}}$ to the overall free-energy change overwhelms that (destabilization) from the polar solvation free-energy term $\langle \Delta G_{\text{sol}}^{\text{pol}} \rangle$. $\Delta E_{\text{bind}}^{\text{HF}}$ can be expressed as the sum of the FMO-IFIE terms for an inhibitor and hNEU2. FMO-IFIE analysis revealed that the

variation in the overall free-energy changes among four sialic acid analogues is determined by hydrogen-bonding interactions between fragment A in a sialic acid analogue and amino acid residues in pocket A in hNEU2. DANA and zanamivir, both of which have the same fragment A structure, show excellent inhibitory activities, because of the effective hydrogen-bond formation of a hydroxyl group in fragment A with Glu111 in hNEU2. The hydroxyl group in fragment C of DANA is replaced by a positively charged guanidino group in zanamivir. This replacement causes great stabilization of the interaction energy through the electrostatic interactions of the guanidino group with negatively charged Glu39 and Asp46, resulting in that zanamivir exhibits more potent activity than DANA.

Second, we determined the reason why an anti-influenza drug, oseltamivir, shows very weak inhibitory activity toward hNEU2, although it exhibits potent activity toward N1-NA. The observed free-energy change for complex formation of hNEU2 with oseltamivir is 10 kcal/mol larger (less stable) than that in the case of N1-NA. There are significant differences in amino acid residues in the active site between the two proteins: notably Ile22, Glu39, and Asn86 in hNEU2 are Glu119, Leu134, and Glu227 in N1-NA, respectively. These three residues surround the hydroxyl and guanidino groups in fragment C of DANA and zanamivir, respectively. IFIE analysis showed that these replacements cause a large difference in the free-energy change between the oseltamivir–hNEU2 and oseltamivir–N1-NA complexes. In addition to the difference in the interaction between fragment C and pocket C, fragment A in oseltamivir, unlike that in zanamivir, is not able to form hydrogen bonds with pocket A. These differences reasonably account for the inefficiency of oseltamivir as to hNEU2.

The present study could provide clues for understanding the neuropsychiatric side effects of oseltamivir, which are similar to the known symptoms of endogenous hNEUs related disorders. Li et al.⁸³ proposed that the hNEU2 variation caused by the Arg41Gln mutation could be associated with the adverse side effects of oseltamivir. They demonstrated that replacement of positively charged Arg41 with neutral Gln41 increases the sensitivity of hNEU2 to oseltamivir and reduces the catalytic activity of hNEU2. This observation is strongly supported by the present results, which suggest that the unfavorable electrostatic repulsive interaction between Arg41 and the positively charged amino group in oseltamivir can be abolished with the mutation from Arg41 to Gln41. In fact, IFIE^{HF} (fragment C, Gln41) is 67 kcal/mol more stable than IFIE^{HF} (fragment C, Arg41) (−9 and 58 kcal/mol, respectively).

The current and previous results²⁸ obtained with the LERE-QSAR analysis successfully reveal the quantitative contribution of each fragment in neuraminidases (hNEU2 and N1-NA) and sialic acid analogues including anti-influenza drugs to the overall interaction free-energy change. Consequently, the results bring a new and consistent understanding of the binding interaction mechanism, which cannot be obtained with any other procedures.

The present work will provide useful information for drug development and for the curing of influenza and other diseases in which neuraminidases are involved.

■ ASSOCIATED CONTENT

Supporting Information. Atom names, atom types, and partial atomic charges of compounds 1–6. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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