

Cite this: *RSC Adv.*, 2017, 7, 12391

Novel valproic aminophenol amides with enhanced glial cell viability effect†

Andrea Alpuche-García, Xochitl Dávila-González, Leticia Arregui* and Hiram I. Beltrán*

New valproic acid derivatives were synthesized by coupling valproyl chloride with *ortho*-aminophenols, resulting in seven *N*-(*ortho*-hydroxyphenyl)valproamides. These amides share similar structural characteristics and exhibit tuneable electronic and steric contributions either without particular substituents, or through the inclusion of electro-donating (–Me), electro-withdrawing (–NO₂) or pi electro-donating/sigma electro-withdrawing (–Cl) substituents at the aromatic ring. The identity of such derivatives was evidenced through spectroscopic characterization using FTIR, ¹H, ¹³C and HETCOR NMR, as well as by analyzing their melting points. In particular, for three derivatives it was feasible to determine their chemical structures in the crystal phase; all three behaved in a similar fashion and exhibited very similar conformations independent of the attached substituents. The base compound was found to exhibit 15.8 times more activity in C6 cells and 4.4 times more activity in U373 cells compared with VPA. In general, the parent compound, or those having –Me as a substituent, presented a greater effect on C6 cells than U373 cells. However, those with –NO₂ and –Cl substituents, as well as VPA, required similar doses for the IC₅₀ in both cell lines. Modification of the base compound with a –Me or –NO₂ substituent increased the effect on cell viability to ca. 20 times that of VPA in both C6 and U373, indicating that a larger structure causes an important enhancement in the inhibition of cell viability. In both cell lines, –Cl containing derivatives were the most active compounds. For these derivatives, an activity increase of ca. 59 and 47 times that of VPA was observed for C6 and U373 cells, respectively. An important perspective is that VPA analogues possessing an aromatic ring with a –Cl substituent may become central structures in the search for more potent pharmaceutical prototypes.

Received 3rd January 2017
Accepted 23rd January 2017

DOI: 10.1039/c7ra00048k

rsc.li/rsc-advances

Introduction

Valproic acid (VPA) is a fatty acid with anticonvulsant properties used in the treatment of epilepsy, and this drug also has anti-tumor activity due to its effects as a histone deacetylase inhibitor.¹ The occurrence of many types of cancer are generally accompanied by histone hypoacetylation.² Because of this, many new strategies to fight cancer are focused on the restoration of de-acetylation balance through the use of histone deacetylase inhibitors (iHDAC) such as VPA.^{3–6} Nevertheless, despite the promising effects offered by VPA against cancer, it causes side effects mainly related to blood and bone marrow toxicity, and high concentrations are required to achieve the desired pharmacological activity, mostly in the mM range.⁷

Glioblastoma is the most common and devastating primary tumour in adults. Epigenetic changes together with genetic modifications have been associated with its generation and

progression. VPA and other iHDAC have been explored for glioblastoma treatment alone or with chemotherapy or radiotherapy, and promising effects have been found. However, additional efforts to find more effective drugs are required, in particular those involving the testing of drugs in glioblastoma cell lines, given the previously reported evidence.⁸

In order to diminish or avoid the side effects associated with VPA, as well as to enhance its activity, a common and useful strategy has been the chemical modification of the parent compound. For instance, in 2005, Eyal *et al.*⁹ evaluated and compared the iHDAC of VPA and its constituent isomers, including valnoctic acid (VCA), propylisopropylacetic acid (PIA), diisopropylacetic acid (DIA), 2,2,3,3-tetramethylcyclopropylcarboxylic acid (TMCA), and VPA metabolites: 2-en-AVP and 4-en-AVP. The authors found that almost any chemical modification of the VPA structure results in the decay of its activity as iHDAC. Another strategy, developed in 2006 by Deubzer and co-workers,¹⁰ was the employment of VPA analogues with longer side chains, as well as the inclusion of alkenyl fragments in the structure. These changes caused multifunctional properties of such derivatives as iHDAC, showing interesting cell cycle modulation due to the induction

Departamento de Ciencias Naturales, DCNI, UAM Cuajimalpa, 05300, Ciudad de México, Mexico. E-mail: arregui@correo.cua.uam.mx; hbeltran@correo.cua.uam.mx

† Electronic supplementary information (ESI) available: CCDC no. 1525302 for **VA**₂, 1525303 for **VA**₃, and 1525301 for **VA**₆. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/c7ra00048k

of p21 expression, as well as low toxicity on CD34+ bone marrow cells.¹⁰ Another recent effort to enhance the beneficial effects of VPA derivatives was carried out using phosphovalproic acid, employed as iHDAC, which involved tracking its dominant molecular target STAT3. This compound includes ester functionality to anchor the VPA fragment to the diethyl (4-hydroxybenzyl) phosphate parent compound. This compound has shown synergistic behaviour in enhanced pancreatic cancer inhibition, compared with VPA.¹¹ Lastly, in 2009, the compound *N*-9-(2-hydroxy)ethoxymethylguanine disubstituted with VPA was patented¹² and in 2014 it was evaluated by Tarasenko and co-workers¹³ as an anticancer agent in various cell lines with promising results. In particular, ester and amide groups link this compound to the VPA moiety, indicating that these functionalities should work to generate new promising derivatives of VPA. Some other trials to find potent VPA analogues with enhanced iHDAC activity have also emerged lately.¹⁴

Thus, the current work aims to obtain and characterize seven new VPA derivatives, wherein the molecular design strategy is not to vary the valproic fragment, but instead the carboxylate moiety, through the formation of amidic functionalities. This change is accompanied by the inclusion of an aromatic ring containing different electro-withdrawing or electro-releasing substituents in order to modulate bioactivities.¹⁵ Finally, the new structures recover the OH fragment present in VPA, through the inclusion of a phenolic group. The results of this contribution are that the seven valproic amides exhibit an enhanced effect on cell viability compared with VPA in two glioma cell lines.

Experimental section

General information

Available solvents and reagents were used as received. 2-Propylpentanoic acid (VPA, CAT P6273-100ML), 2-amino-*p*-cresol (Aph2, CAT 144908-50G), 2-amino-5-methyl-phenol (Aph3, CAT 144916-10G), 2-amino-4-nitro-phenol (Aph4, CAT A70402-5G), 2-amino-5-nitro-phenol (Aph5, CAT 303585-25G), 2-amino-4-chloro-phenol (Aph6, CAT C44400-100G) and 2-amino-5-chloro-phenol (Aph7, CAT 552224-25G) were acquired from Sigma-Aldrich; 2-aminophenol (Aph1, CAT 09110-100G) was obtained from Fluka. The solvents employed were acetonitrile (CH₃CN, CAT 9011-03-4L), tetrahydrofuran (THF, CAT 9450-03-4L), ethyl acetate (AcOEt, CAT 75-05-8-2L) and hexane (Hex, CAT 930L) and were acquired from Reasol.

1D & 2D nuclear magnetic resonance spectroscopy (1D & 2D NMR)

NMR experiments were performed on a VARIAN Mercury 200-BB spectrometer and on an Anasazi spectrometer EFT-60. The ¹H and ¹³C chemical shifts [δ = ppm] were taken as relative to internal SiMe₄ (TMS, $\delta(^1\text{H}) = 0.00$ ppm, $\delta(^{13}\text{C}) = 0.00$ ppm), and the coupling constants were quoted in Hz. The employed solvent was CDCl₃, and just for the compounds AV4 and AV5, 0.1 mL of DMSO-*d*₆ was added to achieve full dissolution due to the presence of nitro groups.

Fourier transform infrared (FT-IR) spectra

The FT-IR spectra were recorded in the range of 4000–400 cm^{−1} on a Bruker Tensor-27 FT-IR spectrophotometer, using KBr pellets or KBr mixtures of each of the isolated samples. The samples were measured using the ATR technique.

Single crystal X-ray diffraction (X-ray)

X-ray diffraction data were acquired on an Enraf Nonius diffractometer FR590 Kappa-CCD ($\lambda_{\text{MoK}\alpha} = 0.71073$ Å, graphite monochromator, $T = 293$ K, using rotating image scan with CCD). Monocrystals were mounted in LINDEMAN tubes. Reflections were corrected for Lorentz and polarization effects. The SHELX-S and SHELX-L 2014 program suite¹⁶ was used for the first structural solution, as well as for the refinement and output data. These programs, as well as the molecular perspectives, were employed and obtained in the OLEX2 environment.¹⁷ All heavy atoms were found through electronic density differences in the Fourier maps and refined anisotropically. Some of the hydrogen atoms were also found through Fourier maps and refined isotropically, and the remnants were geometrically calculated and added for the final refinement.

Thin layer chromatography (TLC)

Thin layer chromatography (TLC) aluminium sheet plates of silica 60 F₂₅₄ of 0.2 mm width (Merck, KGaA) were employed to track the course of each reaction. The TLC separation of each compound was developed using a hexane/ethyl acetate 1 : 1 solvent mixture and the plates were revealed using a UV-vis lamp. In all obtained VA products, the appearance of single spots was indicative of their purity and also confirmed that the desired reaction had been achieved.

Synthetic methodology for compounds VA_{1–7}

Synthesis of VA₁. In a round flask, 0.5 g (3.5 mmol) of VPA was poured and dissolved in 40 mL of CH₃CN. Then, 0.68 mL (3.5 mmol) of SOCl₂ was added to yield the acid chloride of VPA *in situ*.¹⁸ This stage was carried out for 24 h at room temperature, keeping the system closed with a glass cap. Then 0.38 g (3.5 mmol) of Aph1 was predissolved in 5 mL of CH₃CN and 5 mL of triethylamine was added to the round flask drop by drop through an addition funnel, followed by magnetic stirring for an hour. The mixture was set to reflux at a temperature which was maintained for 24 h to finally yield the *N*-(2-hydroxyphenyl)-2-propylpentanamide (AV₁). The completion of the reaction was followed by FTIR carbonyl band displacement of the carboxylic acid, acyl chloride and finally secondary amide groups, as well as by thin layer chromatography of the reaction crudes. Volatiles were removed under reduced pressure, and 20 mL of THF was added to the remaining solids, thus keeping the triethylammonium chloride salt insoluble, which was separated through filtration. The soluble fraction was dried through reduced pressure distillation and the remnant solid was recrystallized in an ethyl acetate/hexane 1 : 4 solvent mixture. In this way it was feasible to obtain compound VA₁ in a yield of 64.8%. Similar to VA₁, the other six valproic amides (VA_{2–7}) were



prepared by just varying the corresponding *ortho*-aminophenol source. As stated in a very recent contribution, this synthetic strategy solely provided *N*-acylation of the carboxylate moiety, instead of *O*-acylation.¹⁹

***N*-(2-Hydroxyphenyl)-2-propylpentanamide (VA₁).** Light brown solid, yield 64.8%, mp 55–57 °C. FTIR ($\nu = \text{cm}^{-1}$, KBr): 3600–2600 (O–H), 3256 (N–H), 3077 (C_{sp^2} –H), 2957, 2929, 2873 (C_{sp^3} –H), 1626, 1603 (–C(=O)–N–), 1545, 1497 (–C(O)=N–), 1387, 1313, 1264, 772, 750, 722, 693. ^1H NMR ($\delta = \text{ppm}$, CDCl_3 , $J = \text{Hz}$): 8.95, 8.73 (OH, sb, 1H & NH, s, 1H), 7.44 (H3, d, $J_o = 7.25$, 1H), 7.02 (H5, t, $J_o = 7.25$, 1H), 6.95 (H4, t, $J_o = 7.25$, 1H), 6.83 (H6, d, $J_o = 7.25$, 1H), 2.46 (H8, q, $J = 6.42$, 1H), 1.49 (H9, m, 4H), 1.23 (H10, m, 4H), 0.92 (H11, t, $J = 6.42$, 6H). ^{13}C NMR ($\delta = \text{ppm}$, CDCl_3): 177.3 (C7, 1C), 148.2 (C1, 1C), 126.5 (C2, 1C), 126.0 (C5, 1C), 122.4 (C4, 1C), 120.4 (C3, 1C), 118.4 (C6, 1C), 48.0 (C8, 1C), 35.4 (C9, 2C), 20.8 (C10, 2C), 14.1 (C11, 2C).

***N*-(2-Hydroxy-5-methylphenyl)-2-propylpentanamide (VA₂).** Dark brown solid, yield 95.7%, mp 99–101 °C. FTIR ($\nu = \text{cm}^{-1}$, KBr): 3580–2400 (O–H), 3252 (N–H), 3221, 3033 (C_{sp^2} –H), 2957, 2927, 2872, 2861 (C_{sp^3} –H), 2660, 2341, 1630, 1605 (–C(=O)–N–), 1541, 1507 (–C(O)=N–), 1358, 1317, 1267, 1259, 1034, 1015, 949, 868, 821, 780, 766, 727, 713. NMR ^1H ($\delta = \text{ppm}$, CDCl_3 , $J = \text{Hz}$): 9.20, 8.94 (OH, sb, 1H & NH, s, 1H), 7.37 (H3, s, 1H), 6.83–6.81 (H5, H6, m, 2H), 2.48 (H8, q, $J = 7.3$, 1H), 2.24 (H12, s, 3H), 1.55 (H9, m, 4H), 1.36 (H10, m, 4H), 0.92 (H11, t, $J = 7.3$, 6H). NMR ^{13}C ($\delta = \text{ppm}$, CDCl_3): 175.9 (C7, 1C), 145.5 (C1, 1C), 128.8 (C4, 1C), 125.8 (C2, 1C), 125.8 (C5, 1C), 122.0 (C3, 1C), 117.2 (C6, 1C), 47.3 (C8, 1C), 35.2 (C9, 2C), 20.5 (C10, 2C), 20.4 (C12, 1C), 15.9 (C11, 2C). Suitable monocrystals for the X-ray diffraction analysis were obtained in a 4 : 1 hexane/ethyl acetate solvent mixture.

***N*-(2-Hydroxy-4-methylphenyl)-2-propylpentanamide (VA₃).** Light brown solid, yield 92.7%, mp 68–70 °C. FTIR ($\nu = \text{cm}^{-1}$, KBr): 3700–2600 (O–H), 3238 (N–H), 3202, 3046 (C_{sp^2} –H), 2959, 2929, 2875, 2858 (C_{sp^3} –H), 1639, 1611 (–C(=O)–N–), 1598, 1546, 1510 (–C(O)=N–), 1469, 1459, 1322, 943, 897, 872, 815, 796, 771, 760, 733. NMR ^1H ($\delta = \text{ppm}$, CDCl_3 , $J = \text{Hz}$): 8.90, 8.45 (OH, sb, 1H & NH, s, 1H), 7.16 (H3, d, $J_o = 7.02$, 1H), 6.82 (H6, s, 1H), 6.65 (H4, d, $J_o = 7.02$, 1H), 2.49 (H8, q, $J = 6.72$, 1H), 2.27 (H12, s, 3H), 1.48 (H9, m, 4H), 1.29 (H10, m, 4H), 0.92 (H11, t, $J = 6.72$, 6H). NMR ^{13}C ($\delta = \text{ppm}$, CDCl_3): 177.0 (C7, 1C), 148.4 (C1, 1C), 136.9 (C5, 1C), 123.4 (C2, 1C), 122.3 (C3, 1C), 121.1 (C4, 1C), 119.4 (C6, 1C), 47.9 (C8, 1C), 35.4 (C9, 2C), 20.8 (C12, 1C), 20.0 (C10, 2C), 14.2 (C11, 2C). Suitable monocrystals for the X-ray diffraction analysis were obtained in a 4 : 1 hexane/ethyl acetate solvent mixture.

***N*-(2-Hydroxy-5-nitrophenyl)-2-propylpentanamide (VA₄).** Light yellow solid, yield 24.6%, mp 85–87 °C. FTIR ($\nu = \text{cm}^{-1}$, KBr): 3500–2400 (O–H), 3416 (N–H), 3097 (C_{sp^2} –H), 2957, 2937, 2872 (C_{sp^3} –H), 2362, 2344, 1627, 1595 (–C(=O)–N–), 1539, 1503 (–C(O)=N–), 1465, 1461, 1427, 1400, 1341 (–N(=O)–O), 1327, 1268, 1244, 1197, 1167, 1153, 993, 878, 819, 781, 743. NMR ^1H ($\delta = \text{ppm}$, CDCl_3 , $J = \text{Hz}$): 9.22, 8.86 (OH, sb, 1H & H3, d, $J_m = 2.74$, 1H), 8.82 (NH, s, 1H), 7.88 (H5, dd, $J_o = 8.96$, $J_m = 2.74$, 1H), 6.98 (H6, d, $J_o = 8.96$, 1H), 2.52 (H8, q, $J = 7.14$, 1H), 1.50 (H9, m, 4H), 1.38 (H10, m, 4H), 0.92 (H11, t, $J = 7.14$, 6H). NMR ^{13}C ($\delta = \text{ppm}$, CDCl_3): 176.3 (C7, 1C), 153.4 (C1, 1C), 140.2 (C4, 1C),

126.7 (C2, 1C), 120.8 (C5, 1C), 116.8 (C3, 1C), 116.0 (C6, 1C), 47.8 (C8, 1C), 35.1 (C9, 2C), 20.6 (C10, 2C), 14.0 (C11, 2C).

***N*-(2-Hydroxy-4-nitrophenyl)-2-propylpentanamide (VA₅).** Orange solid, yield 30.6%, mp 161–163 °C. FTIR ($\nu = \text{cm}^{-1}$, KBr): 3600–2400 (O–H), 3408 (N–H), 3096 (C_{sp^2} –H), 2957, 2934, 2872 (C_{sp^3} –H), 2362, 2347, 1669, 1627 (–C(=O)–N–), 1592, 1506 (–C(O)=N–), 1465, 1424, 1400, 1346 (–N(=O)–O), 1327, 1268, 1247, 1194, 896, 875, 825, 880, 743. NMR ^1H ($\delta = \text{ppm}$, CDCl_3 , $J = \text{Hz}$): 9.07, 8.35 (OH, sb, 1H & NH, s, 1H), 8.20 (H3, d, $J_o = 8.88$, 1H), 7.76 (H4, d, $J_o = 8.88$, 1H), 7.63 (H6, s, 1H), 2.43 (H8, q, $J = 7.14$, 1H), 1.48 (H9, m, 4H), 1.36 (H2, m, 4H), 0.92 (H11, t, $J = 7.14$, 6H). NMR ^{13}C ($\delta = \text{ppm}$, CDCl_3): 175.0 (C7, 1C), 146.1 (C1, 1C), 142.5 (C5, 1C), 132.6 (C2, 1C), 118.7 (C3, 1C), 114.7 (C4, 1C), 109.5 (C6, 1C), 47.1 (C8, 1C), 34.5 (C9, 2C), 19.9 (C10, 2C), 13.4 (C11, 2C).

***N*-(2-Hydroxy-5-chlorophenyl)-2-propylpentanamide (VA₆).** Dark brown solid, yield 42.4%, mp 90–92 °C. FTIR ($\nu = \text{cm}^{-1}$, KBr): 3650–2400 (O–H), 3251 (N–H), 3185 (C_{sp^2} –H), 2958, 2932, 2876 (C_{sp^3} –H), 1630, 1593 (–C(=O)–N–), 1541, 1524 (–C(O)=N–), 1291, 867, 821 (C–Cl). NMR ^1H ($\delta = \text{ppm}$, CDCl_3): 8.17, 7.86 (OH, s, 1H & NH, s, 1H), 7.40 (H3, s, 1H), 6.95 (H5, d, 1H), 6.95 (H6, d, 1H), 2.35 (H8, q, $J = 6.60$, 1H), 1.45 (H9, m, 4H), 1.20 (H10, m, 4H), 0.92 (H11, t, $J = 6.60$, 6H). NMR ^{13}C ($\delta = \text{ppm}$, CDCl_3): 177.3 (C7, 1C), 147.0 (C1, 1C), 127.0 (C5, 1C), 126.3 (C2, 1C), 125.0 (C4, 1C), 121.9 (C3, 1C), 119.5 (C6, 1C), 48.3 (C8, 1C), 35.5 (C9, 2C), 21.0 (C10, 2C), 14.2 (C11, 2C). Suitable monocrystals for the X-ray diffraction analysis were obtained in a 4 : 1 hexane/ethyl acetate solvent mixture.

***N*-(2-Hydroxy-4-chlorophenyl)-2-propylpentanamide (VA₇).** Light brown solid, yield 35.7%, mp 125–127 °C. FTIR ($\nu = \text{cm}^{-1}$, KBr): 3560–2600 (O–H), 3297 (N–H), 3186 (C_{sp^2} –H), 2958, 2935, 2867 (C_{sp^3} –H), 1633, 1593 (–C(=O)–N–), 1531, 1495 (–C(O)=N–), 919, 896, 807 (C–Cl). NMR ^1H ($\delta = \text{ppm}$, CDCl_3 , $J = \text{Hz}$): 8.50, 7.95 (OH, s, 1H & NH, s, 1H), 7.18 (H3, d, $J_o = 8.40$, 1H), 7.00 (H6, d, $J_m = 2.04$, 1H), 6.80 (H4, dd, $J_o = 8.40$, $J_m = 2.04$, 1H), 2.36 (H8, q, $J = 6.60$, 1H), 1.47 (H9, m, 4H), 1.37 (H10, m, 4H), 0.92 (H11, t, $J = 6.60$, 6H). NMR ^{13}C ($\delta = \text{ppm}$, CDCl_3): 177.4 (C7, 1C), 149.1 (C1, 1C), 131.9 (C5, 1C), 124.8 (C2, 1C), 123.1 (C3, 1C), 120.6 (C4, 1C), 119.3 (C6, 1C), 48.4 (C8, 1C), 35.5 (C9, 2C), 21.0 (C10, 2C), 14.3 (C11, 2C).

Semiempirical determination of partition coefficient (log *P*)

A partition coefficient (log *P*) calculation was developed in Hyperchem v.8.0 computational chemistry software. This was performed for VPA and VA_{1–7}, starting from the X-ray structures obtained herein and through the replacement of particular functional groups and substituents for the whole series. Minimization was carried out with the PM3 semiempirical approach for these means and log *P* was computed for each compound at this level of theory.

Cell culture and drug treatment

Human U373 and rat C6 cells were cultured in Dulbecco's Modified Eagle's F12 Medium (DMEM-F12, Gibco, Thermo Fisher Scientific) with 10% fetal calf serum and 1% penicillin/streptomycin in a humidified atmosphere containing 5% CO₂.



at 37 °C. Due to drug hydrophobicity, the drug stocks were dissolved in Tween 80 (1 : 80 weight ratio) and then in DMSO (1 : 11 v/v). C6 and U373 cells were generously donated by Drs José Segovia and José Antonio Arias-Montaña, both from CINVESTAV México.

Cell viability

The cells were seeded at a density of 15 625 cells per cm² in a 96 well plate. The C6 cells were treated for 24 h with the drugs at different concentrations, while the U373 cells were treated for 48 h. Viability was determined using WST-1 reagent (Roche) and the absorbance was measured at 440 nm in a plate reader (Infinite M1000, Tecan). The cell viability was used to obtain the inhibitory concentration at 50% (IC₅₀) of the tested compounds in the C6 and U373 cells. At least three independent assays were performed, and in each assay triplicates were included. The IC₅₀ determination was done by non-linear logarithmic curve fitting using GraphPad Prism version 6.00 for Windows (GraphPad Software, La Jolla California USA, <http://www.graphpad.com>).

Statistical analysis

One-way ANOVA, followed by Dunnett's multiple comparisons test were performed also using GraphPad Prism v. 6.00.

Results and discussion

Synthetic proposal for valproic amides (VA)

In this research, the molecular modifications performed on the VPA parent were aimed on the acidic side, maintaining the valproyl moiety in an attempt to preserve the VPA biological activity. This is because acidic side modifications have been shown to cause a variety of effects that enhance the therapeutic capacity of VPA, *e.g.*, the simplest VPA amide, valproyl amide (VA), which despite having antiepileptic activity has been demonstrated to lack iHDAC activity.²⁰ Consequently, the exchange between acidic and amidic functionalities is not enough to produce the desired effect or even maintain it, because the iHDAC activity is suppressed for VA. Anyway, the preparation of other valproic amides is forthcoming, with the aim of preserving iHDAC activity as well as other particular biological properties like cell viability.

A common strategy in pharmaceutical design has been the inclusion of aminophenols to generate an amidic functionality, where the nitrogen atom forms a peptidic bond or derived groups, in order to form new chemical chimeras through molecular design.²¹ For example, in 2011, Yamazaki *et al.*²² patented a benzoxazole generated through aminophenols and intermediary amides. Their invention refers to a pharmaceutical composition containing this active ingredient to treat illnesses such as hyperlipidemia, atherosclerosis, diabetes and complications, inflammation, and heart disease. Another example, patented in 2013 by Casado-Centellas *et al.*,²³ is a catechol-*O*-methyltransferase inhibitor (iCOMT) that again is a benzoxazole derived from aminophenols and intermediary amides, employed in the prevention and treatment of amyloidosis. Moreover, one of the most commonly employed

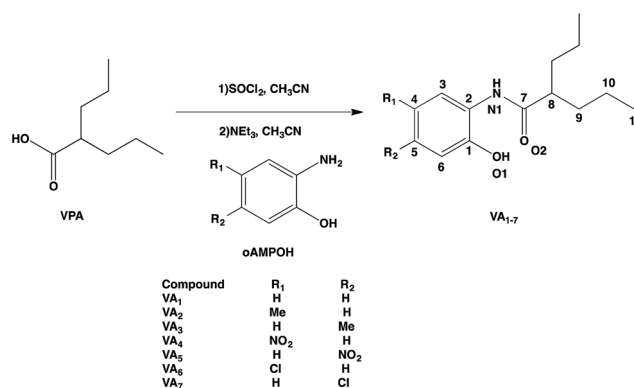
drugs is paracetamol or acetaminophen (APAP), which has analgesic properties and a chemical structure that contains a *para*-aminophenol moiety and an acetyl group attached as an amide to the aromatic ring.²⁴ Considering the latter, it is evident that the incorporation of fragments such as aminophenols in diverse structures provides special characteristics in drugs and prototypes, thus signifying a promising way to design new VPA derivatives.

Molecular design of VPA analogues of valproic amide (VA) containing aminophenols

If one considers the use of aminophenols to generate an amidic functionality, the nitrogen atom is employed for the peptidic bond, leaving the OH untouched and thus recovering this group that is present in the VPA. The incorporation of OH should be as close as possible to the amide functional group, and the *ortho*-aminophenol (oAMPOH) would provide this requisite, thus generating the desired parent compound that connects VPA with oAMPOH. With the inclusion of an aromatic ring in oAMPOH, it is also possible to include structural modifications with substituents that are electro-donating or electro-withdrawing in nature, in order to modulate a plausible biological response and determine what substitution enhanced the activity.²⁵ These variations include fragments with a slight effect on electronic density (–H), electro-donating methyl (–Me), electro-withdrawing nitro (–NO₂) and chlorine (–Cl), which has an electro-withdrawing nature but also has the capacity for pi retrodonation to the aromatic ring.

Synthesis of VA_{1–7}

To obtain the designed compounds, VPA was reacted with SOCl₂ in a 1 : 1 ratio in acetonitrile to yield valproyl chloride (VPCl), which was reacted again to achieve amide formation.¹⁸ Once the VPCl was formed, the corresponding oAMPOH_{1–7} was added to form a 1 : 4 mixture with triethylamine to neutralize the hydrochloric acid obtained during both synthetic steps. In this way, it was possible to generate the amidic bond that joined VPA with oAMPOH_{1–7} to obtain seven *N*-(2-hydroxyphenyl)-



Scheme 1 Synthetic procedure for obtaining VA_{1–7}. The numbering scheme employed was followed for the NMR and X-ray characterization.



valproamides (**VA**_{1–7}) (Scheme 1) in yields of 24.6–95.7%. In all cases, the completion of the reaction was followed by FTIR analysis, due to the change of the carbonyl band from acid to valproyl chloride and finally to valproyl amide. Thin layer chromatography of the crude products was also performed. All compounds were characterized using FTIR, ¹H, ¹³C and HETCOR NMR, and melting point analysis. Also, **VA**₂, **VA**₃ and **VA**₆ were characterized by their single crystal X-ray diffraction structures.

Spectroscopic and structural analysis of **VA**_{1–7}

In this section, the experimental and spectroscopic evidence for the obtained **VA** type compounds through Fourier transform infrared (FTIR) spectroscopy is presented. For the case of **VA**₁, a single band at 3256 cm^{−1} is characteristic of the N–H bond present in a secondary amide (R–NH–C=O). A very broad band is observed around 3600–2600 cm^{−1}, which is characteristic of a phenolic O–H bond. A band belonging to C_{sp}²–H from an aromatic ring at 3077 cm^{−1} and bands due to C_{sp}³–H at 2957, 2929 and 2873 cm^{−1} belonging to the valproyl moiety were identified. The bands of the amidic group appear doubled, and could be due to two possibilities of conformers. These bands appear since the amide and phenolic OH interact, (a) Ph–OH⋯O=C–NH and (b) Ph–(H)O⋯H–N–C=O, gives four amidic bands in 1626, 1603, 1545 and 1497 cm^{−1}. Analogously to **VA**₁, the FTIR band assignment was done for the remaining *N*-(2-hydroxyphenyl)-valproamides (**VA**_{2–7}). For **VA**₄ and **VA**₅, there were also NO₂ bands at 1400 and 1341–1346 cm^{−1}. Finally, the C–Cl bands appear at around 867 and 821 cm^{−1} for **VA**₆, and

896 and 807 cm^{−1} for **VA**₇ (see Experimental section for further details).

Compounds **VA**_{1–7} were also characterized using ¹H (Table 1), ¹³C (Table 2) and HETCOR (¹³C–¹H) NMR experiments. Multiplicities resulted according to their substitution patterns. The HETCOR spectra results were very useful to ascertain the unequivocal assignment of resonances in **VA**_{1–7}, both for the ¹H and ¹³C spectra. Quaternary carbons, secondary amide and phenol signals were confirmed as not having correlations in the HETCOR experiments. Protons H8, H9, H10 and H11, which belong to the valproyl moiety, appear in the ranges of 2.52–2.35 (H8), 1.55–1.45 (H9), 1.38–1.20 (H10) and 0.92 ppm (H11), these resonances are slightly influenced by the function of distance among aromatic and valproyl fragments as well as by the particular substituents at the aromatic ring. In the case of ¹³C NMR, the carbonyl signal C7 appears at 177.4–175.0 ppm, characteristic of a secondary amide with a double hydrocarbon chain, and is slightly influenced by aromatic substitution. In the case of C1, *ipso* to OH, the peak appears in the range of 153.4–145.5 ppm, clearly evidencing the *para* electro-withdrawing effect of the –NO₂ substituent in **VA**₄, as well as the –Cl substituent for **VA**₇, which is instead in the *meta* position. The C2 position, which is *ipso* to the amide, appears at 132.6–123.4 ppm, and is mostly affected by the *para* –NO₂ in **VA**₅. The remnant C3, C4, C5 and C6 positions appear at 123.1–116.8, 140.2–114.4, 142.5–120.8, and 119.5–109.5 ppm, respectively.

Moreover, out of the seven synthesized **VA** compounds, suitable monocrystals for three of them were obtained from ethyl acetate/hexane solvent mixtures. These cases were (crystalline systems and space groups) **VA**₃ (orthorhombic: *P*₂₁2₁2₁), **VA**₂ and **VA**₆ (both as monoclinic: *P*₂₁/*c*), for which crystallographic data were determined through single crystal X-ray diffraction and analysis. Refined data and structures are presented in Tables 3 and 4. Fig. 1 shows the molecular perspectives following the same numbering as that used in the spectroscopic assignment. Selected structural parameters are shown in Table 4. These are aimed to analyse the amidic junction between VPA and oAMPOH, including bond distances (Å), bond angles (°) and dihedral angles (°) for these three analogous structures **VA**₂, **VA**₃ and **VA**₆ (for which data will appear in this same order in this section). A very important observation is that aside from the attached substituent, these three amides are very similar in conformation. This behaviour clearly indicates a structural consensus depending on the

Table 1 ¹H NMR of compounds **VA**_{1–7}^a

Cmpd	OH	NH	H3	H4	H5	H6	H8	H9	H10	H11	H12
VA ₁	8.95	8.73	7.44	6.95	7.02	6.83	2.46	1.49	1.23	0.92	—
VA ₂	9.20	8.94	7.37	—	6.83	6.81	2.48	1.55	1.36	0.92	2.24
VA ₃	8.90	8.45	7.16	6.72	—	6.72	2.49	1.48	1.29	0.92	2.27
VA ₄	9.22	8.82	8.86	—	7.88	6.98	2.52	1.50	1.38	0.92	—
VA ₅	9.07	8.35	8.20	7.76	—	7.63	2.43	1.48	1.36	0.92	—
VA ₆	8.17	7.86	7.40	—	6.95	6.95	2.35	1.45	1.20	0.92	—
VA ₇	8.50	7.95	7.18	6.80	—	7.00	2.36	1.47	1.37	0.92	—

^a The numbering employed is shown in Scheme 1. Samples were dissolved in DMSO-*d*₆. δ = ppm.

Table 2 ¹³C NMR of compounds **VA**_{1–7}^a

Cmpd	C1	C2	C3	C4	C5	C6	C7	C8	C9	C10	C11	C12
VA ₁	148.2	126.5	120.4	122.4	126.0	118.4	177.3	48.0	35.4	20.8	14.1	—
VA ₂	145.5	125.8	122.0	128.8	125.8	117.2	175.9	47.3	35.2	20.5	13.9	20.4
VA ₃	148.4	123.4	122.3	121.1	136.9	119.4	177.0	47.9	35.4	20.0	14.2	20.8
VA ₄	153.4	126.7	116.8	140.2	120.8	116.0	176.3	47.8	35.1	20.6	14.0	—
VA ₅	146.1	132.6	118.7	114.7	142.5	109.5	175.0	47.1	34.5	19.9	13.4	—
VA ₆	147.0	126.3	121.4	125.0	127.0	119.5	177.3	48.3	35.5	21.0	14.2	—
VA ₇	149.6	124.8	123.1	120.6	131.9	119.3	177.4	48.4	35.5	21.0	14.3	—

^a The numbering employed is shown in Scheme 1. Samples were dissolved in DMSO-*d*₆. δ = ppm.



Table 3 Crystallographic data for compounds VA₂, VA₃ and VA₆^a

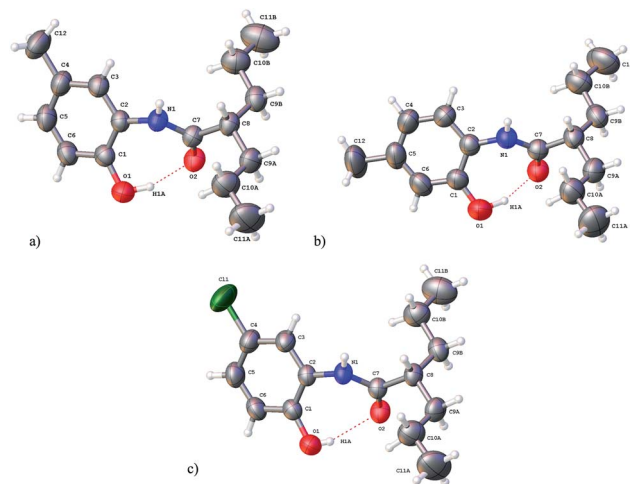
Compound	VA ₂	VA ₃	VA ₆
Molecular formula	C ₁₅ H ₂₃ NO ₂	C ₁₅ H ₂₃ NO ₂	C ₁₄ H ₂₀ ClNO ₂
Molecular weight [g mol ⁻¹]	249.34	249.34	269.76
Crystalline system	Monoclinic	Orthorhombic	Monoclinic
Space group	<i>P</i> 2 ₁ / <i>c</i>	<i>P</i> 2 ₁ 2 ₁ 2 ₁	<i>P</i> 2 ₁ / <i>c</i>
Cell dimensions			
<i>a</i> [Å]	8.9236(5)	8.5924(17)	9.0054(4)
<i>b</i> [Å]	8.7861(4)	12.938(3)	8.7908(4)
<i>c</i> [Å]	19.7125(10)	14.250(3)	19.4618(9)
β [°]	102.535(3)	90	102.377(3)
Volume [Å ³]	1508.69(13)	1584.1(6)	1504.88(12)
<i>Z</i>	4	4	4
ρ (calc.) [g cm ⁻³]	1.0977	1.0455	1.191
μ [mm ⁻¹]	0.072	0.069	0.249
2 θ range [°]	8.42 to 56.80	9.74 to 55.10	6.31 to 55.16
Parameters/restraints	207/30	218/30	222/18
Collected reflections	22 042	22 986	10 105
Independent reflections	3457	3584	3348
<i>R</i> (int.)	0.1127	0.0517	0.0294
GOOF	1.032	1.091	1.029
<i>R</i>	0.0681	0.0558	0.0551
<i>R</i> _w	0.1490	0.1151	0.1469

^a Numbers in parentheses are std. deviations.

characteristics of those attached moieties, and will be discussed as follows. For these three structures, there are clear hydrogen bond interactions between O1–H1a...O2–C7, the former

Table 4 Structural X-ray data for VA₂, VA₃ and VA₆^a

Compound	VA ₂	VA ₃	VA ₆
Bonding distances (Å)			
O1–H1a	0.91 (3)	0.99 (4)	0.67 (3)
O2...H1a	1.75 (3)	1.68 (4)	2.01 (3)
N1–H1	0.83 (2)	0.83 (3)	0.82 (2)
C7=O2	1.247 (2)	1.246 (3)	1.247 (2)
C7–N1	1.338 (3)	1.331 (3)	1.335 (2)
N1–C2	1.429 (3)	1.424 (3)	1.426 (2)
C2–C1	1.392 (3)	1.388 (4)	1.392 (2)
C1–O1	1.362 (3)	1.364 (3)	1.356 (2)
Bonding angles (°)			
O2–C7–N1	122.0 (2)	121.9 (2)	121.76 (15)
C7–N1–C2	127.56 (19)	127.6 (2)	127.13 (15)
N1–C2–C1	123.4 (2)	123.3 (2)	123.45 (16)
C2–C1–O1	123.4 (2)	122.9 (2)	123.60 (16)
H1a–O1–C1	105.7 (18)	105.0 (2)	107 (2)
H1–N1–C2	115.7 (13)	112.8 (18)	116.2 (14)
O1–H1a...O2	160 (3)	158 (4)	160 (3)
Dihedral angles (°)			
H1a–O1–C1–C2	–33.2 (18)	–39 (3)	–36 (3)
O1–H1a...O2–C7	65 (7)	41 (11)	58 (9)
H1–N1–C2–C1	–148.6 (15)	–144.9 (19)	–148.8 (15)
H1–N1–C7–O2	–176.2 (15)	–173 (2)	–175.8 (15)
N1–C7–O2–C8	178.5 (4)	–179.0 (4)	178.4 (4)
C1–C2–N1–C7	45.4 (3)	44.3 (4)	46.8 (3)

^a Numbers in parentheses are std. deviations.Fig. 1 X-ray diffraction structures for (a) VA₂, (b) VA₃ and (c) VA₆ (ellipsoids at 50% of probability).

belonging to the phenol and the latter being the carbonyl fragment, with bond distances of 1.75, 1.68 and 2.01 Å, where these values are smaller than the sum of the van der Waals radii (1.2 Å for hydrogen and 1.52 Å for oxygen).²⁶ The bonding angles (O1–H1a...O2) for this same interaction are 160, 158 and 160°, tending towards 180° due to the strong directionality between these two fragments. Meanwhile, the torsion angles (O1–H1a...O2–C7) present for this hydrogen bond are 65, 41 and 58°, evidencing a needed angularity to avoid steric hindrance between the aminophenol and valproyl moieties. This finding of hydrogen bonding correlates with the observation in the FTIR spectra, where four amidic bands instead of only two were identified, as is common in simpler amides. One pair may be due to the non-hydrogen bonded structure and the other may be due to the hydrogen-bonded counterpart.

For the covalent behaviour of the structures, the bond distances of O1–H1a and N1–H1 were 0.91, 0.99 and 0.67 Å for the former, and 0.83, 0.83 and 0.82 Å for the latter, showing bond contractions only for VA₆. In all cases, the double bonds between C7 and O2 are practically the same at 1.247, 1.246 and 1.247 Å, evidencing a negligible substituent effect through the structure. This last observation is reinforced, as C7–N1 also remains almost constant at 1.338, 1.331 and 1.35 Å for these compounds. As examples of the general structural behaviour, the remaining bond distances, bond angles and torsion angles almost behave the same for the amidic junction between VPA and oAMPOH. In particular, the amidic fragment is practically planar with dihedral angles for N1–C7–O2–C8 of 178.5, 179.0 and 178.4°. Meanwhile, the dihedral angles for the C1–C2–N1–C7 fragment are 45.4, 44.3 and 46.8°, clearly tending to minimize the steric hindrance among the VPA and oAMPOH moieties.

Cell viability of VA_{1–7}

For the cell viability tests, all the VA_{1–7} compounds were pre-dissolved in TWEEN 80 in 1 : 9 (w/w) ratios. This mixture was



reformulated using 1 : 11 (v/v) TWEEN 80/DMSO ratios due to the hydrophobicity of these derivatives, and the final concentrations were related to the respective VA compound. According to the employed ratios of TWEEN 80 and DMSO, the tested concentrations were below the toxicity levels stated for these substances in cell viability assays, and blank tests were also carried out.^{27,28} At the tested concentrations there were no toxic effects. For VPA and all VA₁₋₇ compounds, their cell viabilities in C6 (rat glioma) and U373 (human glioblastoma) lines were measured. The results show that all VA compounds presented more activity than VPA (see Table 5), hence indicating that the molecular design of these derivatives was clever in generating new molecules that fulfilled the desired goal. The cell viability of VPA was measured as the control, and the inhibitory concentration at 50% (IC₅₀) obtained for C6 cells (1179 ± 130.8 μM) and for U373 cells (936.5 ± 174.4 μM) was similar to that of previous reports,²⁹ requiring a millimolar concentration for 48 h. Meanwhile, for the VA₁₋₇ compounds, the required concentrations to achieve IC₅₀ ranged from 19.7 to 74.8 μM for C6, and 1.5 to 212.8 μM for U373 cells, which is comparable with other designed compounds aimed for cancer treatment.³⁰

Structure activity analysis of VA₁₋₇

VA₁ is the base compound and it was found to be 15.8 times more active than VPA in C6 cells (74.8 ± 8.4 μM) and 4.4 times more active than VPA in U373 cells (212.8 ± 12.3 μM), in the same range as other VPA amidic derivatives.³¹ In this case the molecular design gained credibility, due to the fact that this simple derivative augmented the biological response due to the recovery of the hydrogen-bonding group nearby, where it was originally placed in VPA. Furthermore, this modification not only recovers this interaction due to the presence of the aromatic ring, the biological effect is also enhanced by at least 4 times in U373 cells and almost 16 times in C6 cells. These results clearly show that the inclusion of an aromatic ring in this side of the parent molecule is a key method for developing other more potent derivatives. Therefore, the inclusion of steric and electronic modifiers of this aromatic ring, as is the case for VA₂₋₇, should provide insight into the types of preferred interactions in a given molecular receptor in the cell. In this trend,

the modification of the base compound VA₁ with an electro-donating –Me substituent in the *meta* or *para* position relative to the amide group indeed increased the cell viability of the base compound to 41.8 ± 8.4 μM for VA₂ (28.2 times more activity than VPA) and 55.8 ± 11.0 μM (21.1 times more activity than VPA) for VA₃ in C6 cells. In U373 cells, VA₂ and VA₃ also showed an increased effect on cell viability (151.8 ± 14.4 and 161.8 ± 11.2 μM). This result clearly indicates that a bigger structure causes an important enhancement in the cell viability inhibition. Then, the compounds with an electro-withdrawing group, such as the –NO₂ substituent at the *meta* or *para* position relative to the amide, presented a similar effect on the cell viability of C6 cells (67.9 ± 9.7 μM for VA₄ and 45.3 ± 12.3 μM for VA₅) as VA₂ and VA₃. The effect of these amides was more pronounced in U373 cells for the compounds with –NO₂ (64.9 ± 9.0 μM for VA₄ and 39.7 ± 4.4 μM for VA₅). Finally, for both cell lines, VA₆ and VA₇ were the most active compounds. They contain –Cl as the substituent, behaving in a pi electro-donating/sigma electro-withdrawing manner, placed in the *para* or *meta* position relative to the amide. For C6 cells, the IC₅₀ for VA₆ was 19.7 ± 4.7 μM and 21.8 ± 2.7 μM for VA₇; in U373 cells, the doses to obtain the IC₅₀ were 24.6 ± 2.2 μM for VA₆ and 15.5 ± 2.2 μM for VA₇. This is mainly due to the fact that a chlorine atom attached at an aromatic ring prevents a wide variety of chemical reactions, enhancing the possible interactions of the compound with biological receptors.³² Aromatic moieties possessing chlorine (and halogen substituents in general) are a key pharmacological feature¹⁵ justified because these fragments are present in an important amount of already tested and used pharmaceutical drugs and pesticides, and because of the plausible interactions that they present among them, such as R–Cl⋯C=O, R–Cl⋯H–R, R–Cl⋯Ar, R–Cl⋯Cl–R, *etc.*

Moreover, in order to track important physicochemical properties responsible for the biological findings, a semi-empirical (Hyperchem v.8.0) partition coefficient (log *P*) calculation was developed. In this line, log *P*_{VPA} was 2.61, log *P* for VA₁ was 3.39, for VA₂₋₃ was 3.85, for VA₄₋₅ was 3.34 and for VA₆₋₇ was 3.90. This calculation was carried out for each compound to track hydrophobic/hydrophilic ratios and correlate them with biologic trends. The results clearly evidence that the most hydrophobic ratio, resulting from the amide in VA₆₋₇, led to the most potent inhibition, and this was attained due to the –Cl substituent.

In general, VA₁, VA₂ and VA₃ presented a greater effect on C6 cells than U373 cells, but VA₄, VA₅, VA₆ and VPA required similar doses for the IC₅₀. From the analysis of this data we can conclude that the new compounds not only maintain the effect of VPA on the cell viability, but can also achieve the same viability with lower doses. Related references clearly state that amides of VPA do not undergo hydrolysis^{33,34} or further biotransformations, due to the steric hindrance of the valproyl moiety not only under chemical but also biological conditions. Therefore, this kind of compound should have a major effect on cell viability due to the newly obtained chemical structure and not due to the VPA metabolite. This demonstrates that the new base structure (with the neutral substituent) can be modified to increase the activity, and the best activity was found with –Cl, followed by –NO₂ and

Table 5 IC₅₀ of VA₁₋₇ measured by cell viability in glioma cell lines C6 and U373^a

Compound	Substituent*	C6 (μM)	U373 (μM)
VA ₁	–H	74.8 ± 8.4	212.8 ± 12.3
VA ₂	<i>m</i> -Me	41.8 ± 8.4	151.8 ± 14.4
VA ₃	<i>p</i> -Me	55.8 ± 11.0	161.8 ± 11.2
VA ₄	<i>m</i> -NO ₂	67.9 ± 9.7	64.9 ± 9.0
VA ₅	<i>p</i> -NO ₂	45.3 ± 12.3	39.7 ± 4.4
VA ₆	<i>m</i> -Cl	19.7 ± 4.7	24.6 ± 2.2
VA ₇	<i>p</i> -Cl	21.8 ± 2.7	15.5 ± 2.2
VPA	—	1179 ± 130.8	936.5 ± 174.4

^a Data represents median ± standard error of at least three independent experiments. Glioma cells were in contact with each compound (VA₁₋₇) in complete medium for 48 h. *Position relative to the amidic group.



finally with –Me in U373 cells. But in C6 cells, there was not a significant difference between –NO₂ and –Me groups. This may reflect different histone deacetylase expression or regulation in these cell lines and further work is under development on this issue.

Conclusions

Finally, seven designed valproic acid derivatives, *N*-(*ortho*-hydroxyphenyl)valproamides (VA_{1–7}), were obtained in moderate yields by coupling valproyl chloride with seven *ortho*-aminophenols. This led to a family of amides with similar structural characteristics but tuneable electronic and steric contributions, through the inclusion of electro-donating (–Me), electro-withdrawing (–NO₂) or pi electro-donating/sigma electro-withdrawing (–Cl) substituents at the aromatic ring. The identity of such derivatives was evidenced through spectroscopic characterization using FTIR, ¹H, ¹³C and HETCOR NMR, as well as by melting point analysis. In particular, for three of the derivatives (VA₂, VA₃ and VA₆) it was feasible to determine their chemical crystalline structures, in all cases behaving in a similar fashion and with very similar conformations independent of the attached substituents. In general, VA₁, VA₂ and VA₃ affected C6 cells more than U373 cells, but VA₄, VA₅, VA₆ and VPA required similar doses for the IC₅₀. From the analysis of this data we can conclude that the new compounds not only maintain the effect of VPA on cell viability, but also require lower doses. This demonstrates that the new base structure (with a neutral substituent) can be modified to increase the activity, and the best activity was found with –Cl, which was as expected due to the inclusion of this substituent in a huge amount of tested drugs. The hydrophobic effect was also an important variable in enhancing the potency of these derivatives, and the chlorine-bearing molecules produced the best results. In the biological trend –Cl is then followed by –NO₂ and finally by –Me in U373 cells. However, in C6 cells, there was not a significant difference between the –NO₂ and –Me groups, since they have almost the same hydrophobic/hydrophilic ratio according to the log *P* results. This may reflect different histone deacetylase expression or regulation in these cell lines, and work is under development on this issue. An important perspective is that VA₆ and VA₇ may become lead structures in the search for more potent pharmaceutical prototypes.

Acknowledgements

We acknowledge CONACyT projects 47310337 (L. Arregui) and 222872 (H. I. Beltrán), UAM and SEP-PROMEP for financial support. A. Alpuche-García and X. Dávila-González acknowledge university scholarships from CONACyT and UAM.

References

- 1 C. Wang, Z. Luan, Y. Yang, Z. Wang, Y. Cui and G. Gu, *Neurosci. Lett.*, 2011, **497**, 122–127.
- 2 S. Chateavieux, F. Morceau, M. Dicato and M. Diederich, *J. Biomed. Biotechnol.*, 2010, **2010**, 479364.
- 3 R. W. Johnstone, *Nat. Rev. Drug Discovery*, 2002, **1**, 287–299.
- 4 M. Dokmanovic, C. Clarke and P. A. Marks, *Mol. Cancer Res.*, 2007, **5**, 981–989.
- 5 D. Marchion and P. Münster, *Expert Rev. Anticancer Ther.*, 2007, **7**, 583–598.
- 6 M. Mottamal, S. Zheng, T. L. Huang and G. Wang, *Molecules*, 2015, **20**, 3898–3941.
- 7 A. V. Krauze, S. D. Myrehaug, M. G. Chang, D. J. Holdford, S. Smith, J. Shih, P. J. Tofilon, H. A. Fine and K. Camphausen, *Int. J. Radiat. Oncol., Biol., Phys.*, 2015, **92**, 986–992.
- 8 D. Thotala, R. M. Karvas, J. A. Engelbach, J. R. Garbow, A. N. Hallahan, T. A. DeWees, A. Laszlo and D. E. Hallahan, *Oncotarget*, 2015, vol. 6, pp. 35004–35022.
- 9 S. Eyal, B. Yagen, J. Shimshoni and M. Bialer, *Biochem. Pharmacol.*, 2005, **69**, 1501–1508.
- 10 H. Deubzer, B. Busche, G. Rönndahl, D. Eikel, M. Michaelis, J. Cinatl, S. Schulze, H. Nau and O. Witt, *Leuk. Res.*, 2006, **30**, 1167–1175.
- 11 G. G. Mackenzie, L. Huang, N. Alston, N. Ouyang, K. Vrankova, G. Mattheolabakis, P. P. Constantinides and B. Rigas, *PLoS One*, 2013, **8**, e61532.
- 12 C. Harmon and D. Myles, Valproic acid salts, *US Pat.*, WO2009142968 A2, 2009.
- 13 N. Tarasenko, S. M. Cutts, D. R. Phillips, G. Berkovitch-Luria, E. Bardugo-Nissim, M. Weitman, A. Nudelman and A. Rephaeli, *Biochem. Pharmacol.*, 2014, **88**, 158–168.
- 14 E. Perrino, G. Cappelletti, V. Tazzari, E. Giavini, P. D. Soldato and A. Sparatore, *Bioorg. Med. Chem. Lett.*, 2008, **18**, 1893–1897.
- 15 C. Bissantz, B. Kuhn and M. Stahl, *J. Med. Chem.*, 2010, **53**, 5061–5084.
- 16 G. M. Sheldrick, *Acta Crystallogr., Sect. C: Struct. Chem.*, 2015, **71**, 3–8.
- 17 O. V. Dolomanov, L. J. Bourhis, R. J. Gildea, J. A. K. Howard and H. Puschmann, *J. Appl. Crystallogr.*, 2009, **42**, 339–341.
- 18 E. Bechar and H. Astroug, *Arch. Pharm.*, 1997, **330**, 273–276.
- 19 N. Pariente-Cohen, M. Weitman, N. Tania, D. T. Major, H. E. Gottlieb, S. Hoz and A. Nudelman, *RSC Adv.*, 2015, **5**, 24038–24043.
- 20 C. U. Johannessen and S. I. Johannessen, *CNS Drug Rev.*, 2003, **9**, 199–216.
- 21 H. Andleeb, Y. Tehseen, S. J. Ali Shah, I. Khan, J. Iqbal and S. Hameed, *RSC Adv.*, 2016, **6**, 77688–77700.
- 22 Y. Yamazaki, T. Toma, M. Nishikawa, H. Ozawa, A. Okuda, K. Abe and S. Oda, Benzoxazole compound and pharmaceutical composition containing the compound, *Germany Pat.*, EP1433786 B1, 2011.
- 23 M. Casado Centellas, B. R. Insa, B. N. Reig and B. N. Gavalda, New therapy for transthyretin-associated amyloidosis, *Spain Pat.*, WO2013060668 A1, 2013.
- 24 J. A. Clements, R. C. Heading, W. S. Nimmo and L. F. Prescott, *Clin. Pharmacol. Ther. Pediatr. Perspect.*, 1978, **24**, 420–431.
- 25 J. Fu, K. Cheng, Z. M. Zhang, R. Q. Fang and H. L. Zhu, *Eur. J. Med. Chem.*, 2010, **45**, 2638–2643.
- 26 A. Bondi, *J. Phys. Chem.*, 1964, **68**, 441–451.



- 27 B. Arechabala, C. Coiffard, P. Rivalland, L. J. M. Coiffard and Y. D. Roeck-Holtzhauer, *J. Appl. Toxicol.*, 1999, **19**, 163–165.
- 28 Y. He, H. Liu, Z. Xie, Q. Liao, X. Lai and Z. Du, *Drug Dev. Ind. Pharm.*, 2014, **40**, 237–243.
- 29 J. A. Benitez, L. Arregui, G. Cabrera and J. Segovia, *Neuroscience*, 2008, **156**, 911–920.
- 30 V. Di Bussolo, E. C. Calvaresi, C. Granchi, L. Del Bino, I. Frau, M. C. Dasso Lang, T. Tuccinardi, M. Macchia, A. Martinelli, P. J. Hergenrother and F. Minutolo, *RSC Adv.*, 2015, **5**, 19944–19954.
- 31 M. Farooq, A. El-Faham, S. N. Khattab, A. M. Elkayal, M. F. Ibrahim, N. A. Taha, A. Baabbad, M. A. M. Wadaan and E. A. Hamed, *Asian Pac. J. Cancer Prev.*, 2014, **15**, 7785–7792.
- 32 H. Marquardt, *Angew. Chem.*, 1995, **107**, 1017.
- 33 M. Bialer, A. Haj-Yehia, K. Badir and S. Hadad, *Pharm. World Sci.*, 1994, **16**, 2–6.
- 34 S. Blotnik, F. Bergman and M. Bialer, *Drug Metab. Dispos.*, 1996, **24**, 560–564.

