Small-molecule inhibitors of the PDZ domain of Dishevelled proteins interrupt Wnt signalling

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Abstract

Dishevelled (Dvl) proteins are important regulators of the Wnt signalling pathway, interacting through their PDZ domains with the Wnt receptor Frizzled. Blocking the Dvl PDZ/Frizzled interaction represents a potential approach for cancer treatment, which stimulated the identification of small molecule inhibitors, among them the anti-inflammatory drug Sulindac and Ky-02327. Aiming to develop tighter binding compounds without side effects, we investigated structure-activity relationships of sulfonamides. X-ray crystallography showed high complementarity of anthranilic acid derivatives in the GLGF loop cavity and space for ligand growth towards the PDZ surface. Our best binding compound inhibits Wnt signalling in a dose-dependent manner as demonstrated by TOP-GFP assays (IC$_{50}$ ~50 µM), and Western blotting of β-catenin levels. Real-time PCR showed reduction in the expression of Wnt-specific genes. Our compound interacted with Dvl-1 PDZ (K$_D$=2.4 µM) stronger than Ky-02327 and may be developed into a lead compound interfering with the Wnt pathway.

KEYWORDS: Drug Design, NMR, PDZ, Frizzled, Wnt signalling

INTRODUCTION

Dishevelled (Dvl) proteins comprise 500 to 600 amino acids and contain three conserved domains: an N-terminal DIX (**Di**shevelled/A**x**in) domain (Schwarz-Romond 2007, Madrzak 2015), a central PDZ (**P**SD95/**D**lg1/**Z**O-1) domain (Doyle 1996, Ponting 1997), and a C-terminal DEP (**D**ishevelled/**E**gl-10/**P**leckstrin) domain (Wong 2000, Wallingford 2005). Dvl transduces Wnt signals from the membrane receptor Frizzled to downstream components *via* the interaction between Dvl PDZ and Frizzled (Wong 2003), thus it has been proposed as drug target (Klaus 2008, Holland 2013, Polakis 2012). Several studies identified internal peptides of the type (KTXXXW) as well as C-terminal peptides of the type (ΩΦGWF) in which Ω is any aromatic amino acid (F, W or Y) as Dvl PDZ targets (Lee 2009, Zhang 2009). Three Dvl homologues, Dvl-1, Dvl-2 and Dvl-3, have been identified in humans. Sequence identity is 88% between Dvl-3 PDZ and Dvl-1 PDZ and 96% between Dvl-3 PDZ and Dvl-2 PDZ (Supporting Information Figure S1). Dvl proteins are found to be upregulated in breast, colon, prostate, mesothelium, and lung cancers (Weeraratna 2002, Uematsu 2003, Bui 1997, Mizutani 2005).

PDZ domains appear in 440 copies spread over more than 260 proteins of the human proteome (Ponting 1997). They maintain relatively specific protein-protein interactions and are involved, for example, in signalling pathways, membrane trafficking and in the formation of cell-cell junctions (Zhang 2003, Fanning 1996, Kurakin 2007). Hence, they are potentially attractive drug targets (Rimbault 2019, Christensen 2020). PDZ domains consist of about 90 amino acids which fold into two α-helices and six β-strands exposing a distinct peptide-binding groove (Doyle 1996), Lee 2017). The conserved carboxylate-binding loop (GLGF loop, FLGI in Dvl-2 and -3, Figure 1) is essential for the formation of a hydrogen bonding network between the PDZ domain and PDZ-binding, C-terminal peptide motifs, in most cases coordinating the C-terminal carboxylate group of the interaction partner. In the respective complexes, the C-terminal residue of the ligand is referred to as P_0; subsequent residues towards the N-terminus are termed P_{-1}, P_{-2}, and P_{-3} etc. Previous studies have revealed that P_0 and P_{-2} are most critical for PDZ-ligand recognition (Songyang 1997, Schultz 1998).

PDZ domains are divided into at least three main classes on the basis of their amino acid preferences at these two sites: class I PDZ domains recognize the motif S/T-X-Φ-COOH (Φ is a hydrophobic residue and X any amino acid). Class II PDZ domains recognize the motif Φ-X-Φ-COOH, whereas class III

PDZ domains recognize the motif X-X-COOH. However, some PDZ domains do not fall into any of these specific classes (Pawson 2007, Sheng 2001, Zhang 2003). The Dvl PDZ domains, for example, recognize the internal sequence (KTXXXW) within the frizzled peptide 525(GKTLQSWRRFYH)536 ($K_D \sim 10\ \mu M$) (Wong 2003, Chandanamali 2009).

Due to their occurrence in important proteins, PDZ domains received early attention as drug targets, nicely summarized in Christensen 2019. There are several examples of Dvl PDZ inhibitors of peptide or peptidomimetic nature (eg. Hammond 2006, Haugaard-Kedstrom 2021), including peptide conjugates (eg. Qin 2021, Hegedus 2021), and on an organic, small-molecule basis. The latter approach is considered most beneficial in long term medical treatments of conditions like cancer or neurological disorders. NSC668036 (Shan 2005, Wang 2015) is a peptide-mimic compound which interferes with Wnt signalling at the Dvl level. Based on a computational pharmacophore model of NCS668036, additional compounds were later reported (Shan 2012). Known as first non-peptide inhibitor, the 1H-indole-5-carboxylic acid derivative FJ9 (Fujii 2007) showed therapeutic potential. Further examples including Sulindac (Lee 2009), 2-((3-(2-Phenylacetyl)amino)benzoyl)amino)benzoic acid (3289-8625, also called CalBioChem(CBC)-322338) (Grandy 2009, Hori 2018), N-benzoyl-2-amino-benzoic acid analogs (Hori 2018), phenoxyacetic acid analogs (Choi 2016), and Ethyl 5-hydroxy-1-(2-oxo-2-((2-(piperidin-1-yl)ethyl)amino)ethyl)-1H-indole-2-carboxylate (KY-02327) (Kim 2016) have been reported, with the latter showing the highest *in-vitro* affinity (8.3 µM) of all. Despite the existence of the abovementioned inhibitors of Dvl PDZ, the development of tighter-binding, non-peptidic small-compound modulators of the respective functions, binding with nanomolar affinity, is necessary and remains challenging. On this path, we explore optimal fits for the primary binding pocket by cycles of chemical synthesis and X-ray crystallography and further avenues for systematically growing ligands along the Dvl PDZ surface to provide SAR for the development of inhibitors in the low or medium nanomolar range. Nuclear magnetic resonance (NMR) spectroscopy was used to detect primary hits and for follow-up secondary screening. The ability of NMR to detect weak intermolecular interactions ($\mu M < K_D < mM$) make it an ideal screening tool for identifying and characterizing weakly binding fragments, to be optimized subsequently by chemical modification in order to improve binding (Zartler 2006, Shuker 1996, Zartler 2003). Besides NMR, the determination of X-ray crystal structures of selected

90   complexes was fundamental for further design of new compound structures with improved binding. In

91   the first round of screening, a library constructed after computational docking of candidates into the

92   peptide binding site of the Dvl PDZ domains were investigated, followed by secondary screening

93   utilizing a library of 120 compounds containing rhodanine or pyrrolidine-2,5-dione moieties.

94

95   **RESULTS AND DISCUSSION**

96   **PDZ targeted library design**

97   The PDZ targeted library was designed to cover all PDZ domains with available structure. For this, all

98   X-ray and NMR derived PDZ structures were retrieved from the PDB, clustered, and 6 selected centroids

99   were subjected to the virtual screening routine. The area considered is shown in Figure 1A, with the blue

100 sphere indicating the geometrical centre. The clustering of the PDZ domains was performed according

101 to the shapes of their binding sites, rather than backbone conformation. This approach accounts for the

102 importance of surface complementarity of protein-small molecule interactions and the critical

103 contribution of van der Waals interactions to the binding free energy. On another hand, PDZ domains

104 have evolved to recognize a carboxyl group that is mostly derived from the C-terminus of natively

105 binding proteins. Finally, the fact that PDZ can recognize internal motifs (Hillier 1999), including

106 KTXXXW of Frizzled-7 recognised by Dvl PDZ (Wong 2003, Chandanamali 2009), raises the question

107 of what are key binding contributions with PDZ domains: negative charge, hydrogen bonding or shape

108 complementarity (Harris 2003). For this reason, tangible compounds were preselected to have extensive

109 hydrophobic contacts as well as chemical groups that mimic the carboxylic group.

110 Virtual screening was performed with QXP, and the generated complexes were sequentially filtered with

111 a self-designed MultiFilter algorithm. From the resulting 1119 compounds a randomly selected set of

112 250 compounds was subjected to NMR validation.

113

114 **NMR Screening and development of compounds**

115 The results of virtual screening were checked experimentally by comparing 2D ^{1}H-^{15}N HSQC

116 (Heteronuclear Single Quantum Correlation) spectra of Dvl-3 PDZ in the absence and presence of the

117 compound to elucidate ligand-induced changes of chemical shifts. Chemical shift perturbation

118   differences (ΔCSP, representing the average of the three strongest shifting cross peaks according to

119   equation 1) were evaluated in cases where the residues responding strongest are inside the area defined

120   by Figure 1A. The responses were classified into: (i) inactive compounds (ΔCSP < 0.02); (ii) very weak

121   interactions (0.02 ≤ ΔCSP ≤ 0.05); (iii) weak interactions (0.05 < ΔCSP ≤ 0.1); (iv) intermediate

122   interactions (0.1 < ΔCSP ≤ 0.2); (v): strong interactions (0.2 < ΔCSP ≤ 0.5) and (vi) very strong

123   interactions (ΔCSP > 0.5). In most cases, the signals of residues S263, V287 and R320 (Figure 1A)

124   within the conserved binding site were most strongly perturbed (Supporting Information Figure S2).

125   With the ΔCSP of 0.12 ppm, the isoleucine-derived compound **1** ((2,3-dihydrobenzo[b][1,4]dioxin-6-

126   yl)sulfonyl)-L-isoleucine containing a sulfonamide moiety was detected initially as one of the best "hits"

127   according to chemical shift changes. The sulfonamide is a well-known moiety in drug discovery

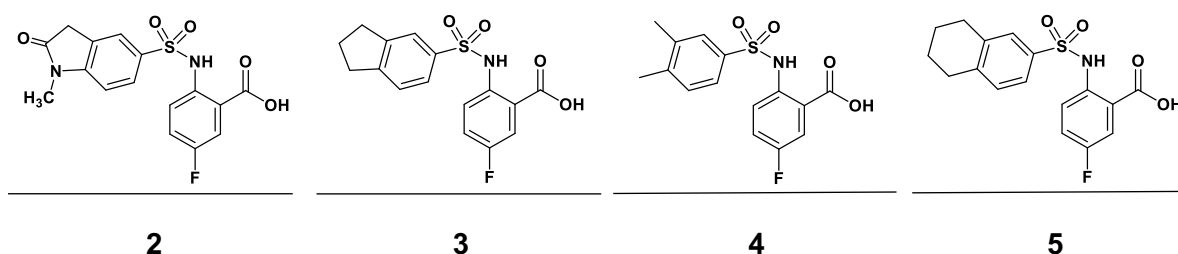128   (Mathvink 1999, Wu 1999, Sleight 1998 O'Brien 2000, Tellew 2003).

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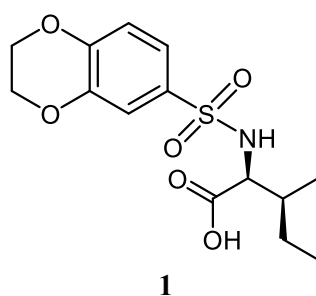
130
131   **1**

132   Upon NMR titration experiments for compound **1** (Supporting Information Figure S2) with Dvl-3 PDZ,

133   the largest chemical shift perturbations were observed for S263 in strand βB and R320 in helix αB of

134   Dvl-3 PDZ, confirming the conserved binding site.

135

136   **Scheme 1**: Compounds **2**, **3**, **4**, **5**

137   By comparing the binding of several sulfonamide compounds in a secondary screening event and mak-

138   ing use of our in-house library, four new compounds (**2**, **3**, **4**, **5**) that induced chemical shift perturbations

139   larger than 0.2 ppm were found (for binding constants see Table 1) and considered further as reasonably

140   strong binders. The similarity of the structures led us to define Scheme 2 as a scaffold for further refine-

141   ments. Sulfonamides were considered more drug-like, and hence followed up at higher priority than

142   other hits. We realised that our four new compounds had different moieties at R_2 in combination with a

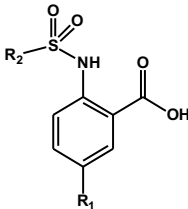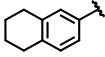143   small R_1 (fluorine). A decrease of binding was observed with decreasing size of R_2.
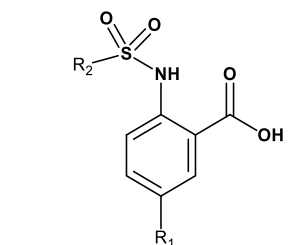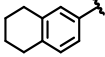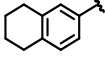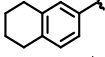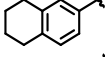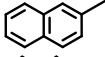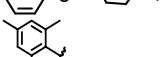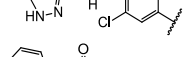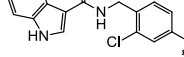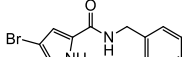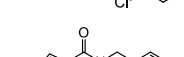
144   **Scheme 2**: Basic fragment for further synthesis

145   In order to assess the importance of the aryl group at R_2 for complex formation, it was replaced by a

146   methyl group as substituent to yield compound **6**, which showed a drastic decrease of binding (Table 1).

147   Compounds **3**, **4**, and **5** did not distinguish between the Dvl-3 PDZ and Dvl-1 PDZ. In order to obtain

148   detailed insight into the binding mode of these compounds, crystal structures of Dvl-3 PDZ in complex

149   with compounds **3**, **5** and **6** were determined (Figure 1). For compound **3** the crystal structure revealed

150   two complexes within the crystallographic asymmetric unit (AU) at 1.43 Å resolution. Both show the

151   anthranilic acid with the attached fluorine pointing into the hydrophobic binding pocket (Figure 1B and

152   Supporting Information Figure S3A), while the carboxyl group forms a hydrogen-bond network with

153   amide residues of the carboxylate binding loop, in particular strand βB (Figure 1B) and specifically with

154   residues I262, G261 and L260. The two sulfonamide oxygen atoms form hydrogen bonds with R320

155   and H324 (weak) of helix αB for only one complex in the AU. The aromatic aryl group

156   (tetrahydronaphtalene) attached to the sulfonamide is involved in hydrophobic interactions with F259

157   (Supporting Information Figure S3B). The 1.6-Å complex structure with compound **5** (4 molecules per

158   AU) exhibits a comparable binding mode as found for compound **3** with a hydrogen-bond network

159   involving the carboxyl group and the amides of I262, G261, L260, and of the sulfonamide to H324

160   (Figure 1C). No hydrogen bond was observed to R320 in all four molecules of the AU, but small

161   variations of the aryl moiety relative to F259 (Supporting Information Figure S3C). The crystals of the

162   complex with **6** show two molecules in the AU (Figure 1D). The sulfonamide is bound by H324 in both

complexes (Supporting Information Figure S3D). However, compound **6** bound only in the mM range as compared to **3** and **5**, which obviously results from the missing aromatic rings.

ID	R$_1$	R$_2$	(K$_D$, μM) Dvl-3PDZ	(K$_D$, μM) Dvl-1 PDZ
2	F		nd	237.6 ± 38.5^{NMR}
3	F		80.6 ± 6.1^{NMR}	112.7 ± 25.9^{NMR}
4	F		83.9 ± 7.8^{NMR}	114.4 ± 9.8^{NMR}
5	F		140.6 ± 14.1^{NMR}	160.1 ± 14.6^{NMR}
6	F	CH$_3$	> 1000^{ITC}	-
7	Br		20.6 ± 2.4^{NMR}	18.2 ± 2.4^{NMR}
8	CF$_3$		17.4 ± 0.5^{ITC}	24.5 ± 1.5^{ITC}
9	Cl		41.1 ± 3.1^{NMR}	45.6 ± 4.5^{NMR}
10	CH$_3$		62.5 ± 4.7^{NMR}	60.5 ± 5.3^{NMR}
11	Br		13.8^{ITC}	119.9^{ITC}
12	Br		58.5^{ITC}	nd
13	Br		7.2^{ITC}	213.2^{ITC}
14	Br		58.1 ± 2.1^{ITC}	nd
15	CF$_3$		52.9 ± 1.7^{ITC}	nd
16	CF$_3$		59.1 ± 1.5^{ITC}	nd
17	CF$_3$		49.5^{ITC}	nd
18	CH$_3$		9.4 ± 0.6^{ITC}	2.4 ± 0.2^{ITC}
19	CH$_3$		21.8 ± 1.7^{ITC}	8.0 ± 0.5^{ITC}
20	CH$_3$		9.8 ± 0.3^{ITC}	4.7 ± 0.3^{ITC}
21	CH$_3$		12.5 ± 0.5^{ITC}	4.7 ± 0.2^{ITC}

Table 1. Binding constants K$_D$ (μM) of Dvl-3 PDZ and Dvl-1 PDZ for compounds 3 – 21 derived by ITC or NMR if not specified. The K$_D$ values determined by NMR are reported as means ± standard deviations of measurements evalutating shifts of cross peaks of at least six residues influenced upon binding of the ligand. The K$_D$ values (1/K$_A$) determined by ITC were obtained as fits to a one-site binding model (n in the range of 0.95-1.2) with K$_D$ errors obtained by ΔK$_A$/K$_A^2$.
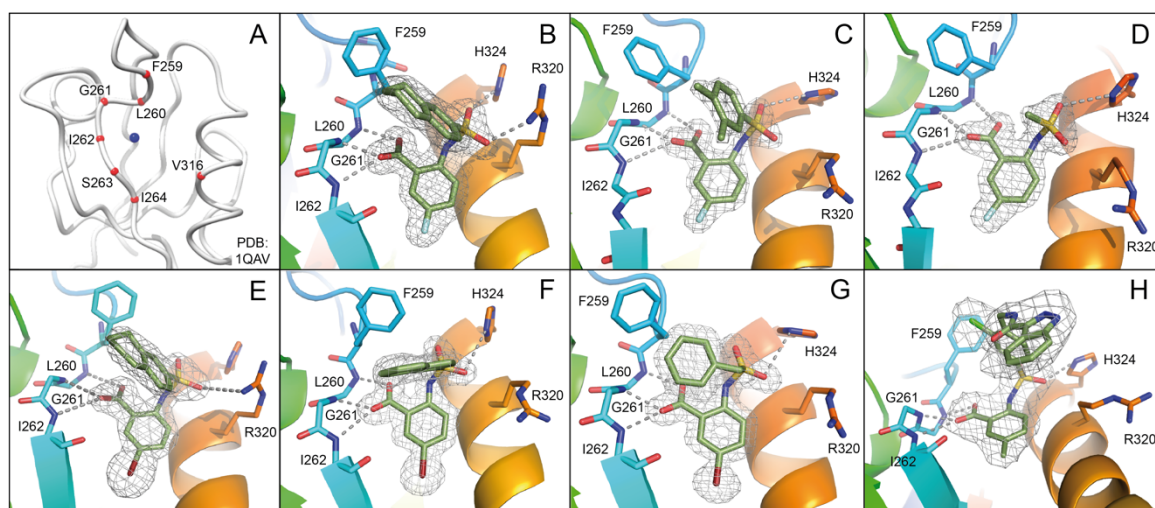
Figure 1: **A** Definition of PDZ binding site. The center of the binding site (blue sphere) is defined as the geometric center of Cα atoms (red spheres) of 7 residues (typed in red) defined by multiple sequence alignment. **B-H** Magnified views into crystal structures of various compounds bound to the Dvl-3 PDZ domain. The 2Fo-Fc electron density around the compounds is shown at 1σ contour level, and the dotted lines indicate formed hydrogen bonds. In the bound compounds covalent bonds to carbon atoms are shown as green sticks. Important residues involved in compound binding are labelled and displayed in atom colours (carbons blue or dark yellow). **B-D** show compound **3**, **5** and **6** respectively. All compounds in **B-D** contain fluorine (light blue) in para position to the amine. **E-G** represents the bound compounds **7**, **11** and **12**, respectively. All have bromine (dark red) in para position to the amine. **H** shows compound **18** within the binding site. The accession codes of the structures B-H are 6ZBQ, 6ZBZ, 6ZC3, 6ZC4, 6ZC6, 6ZC7 and 6ZC8, respectively.

To further explore the importance of the fluorine site inside the hydrophobic pocket, substitutions by bromine, chlorine, methyl and trifluoromethyl were chosen. In fact, the methyl group has a similar vdW radius as the CF_3 group. Iodine was not considered a good candidate since it increases molecular weight substantially and the compounds may be chemically less stable, in particular in biological assays. Taking into account that compound **6** did not bind because of the missing aromatic ring at the R_2 position, our overall strategy was to increase the aromatic ring at R_2 while finding a good fit for R_1, keeping an eye on the molecular weight to enable further compound modifications that fulfil key properties as defined by Lipinski (Lipinski 2000, Lipinski 1997). Our preference to continue exploration at the R_1 position of the aromatic ring in Scheme 1 was inspired by the absence of hits with other substitutions in the secondary screening event and the initial X-ray structures that showed a hydrophobic pocket available for substituents in this position while other sites at the aromatic ring would include steric hindrance. Therefore, compounds **7-17** were obtained and were classified in three different groups to derive structure activity relationships (SAR). The compounds **7-10** in group 1 contain different R_1 (Br, CF_3, Cl, CH_3) but the same moiety (tetrahydronaphthalene) at R_2. As expected, binding could be further improved by displacement of the fluorine with elements exhibiting larger van der Waals (vdW) radii.

8

199   Indeed, the K_D decreased stepwise and the best fit was observed for compound **8** containing a

200   trifluoromethyl group (K_D = 17.4 μM for Dvl-3 PDZ and 24.5 μM for Dvl-1 PDZ). The different

201   substituents at the R_1 position contribute to an increased binding affinity in the following order: F < Cl

202   < Br < CF_3 (compound **3** < **9** < **7** < **8**, respectively). Compound **10** with a methyl group at the R_1 position

203   showed only marginally improved binding, although the methyl group has a similar vdW radius as the

204   CF_3 group of compound **8**. The difference in binding results most likely from their different

205   hydrophobicity.

206   The 1.85-Å crystal structure of the Dvl-3 PDZ domain with compound **7** (K_D = 20.6 μM for Dvl-3 PDZ

207   and 18.2 μM for Dvl-1 PDZ) showed an identical hydrogen-bond network involving the amide groups

208   of residues I262, G261 and L260 of the carboxyl binding loop as seen for all other complex structures

209   reported here (Figure 1E). Only one hydrogen bond between the sulfonamide and R320 was found in

210   addition for one of the two Dvl-3 PDZ molecules per AU. H324 of Dvl-3 PDZ was not addressed by the

211   sulfonamide as seen previously. The bromine at position R_1 points into the hydrophobic pocket, similar

212   as the fluorine in the complex structure with compound **3**. The two complexes in the AU show significant

213   variations in the positions of the tetrahydronaphtalene rings as well as for the side chain of F259 and

214   R320 (Supporting Information Figure S3E).

215   Following the analysis of the complex involving compound **7**, the binding characteristics of the group-

216   2 compounds (**11-14**) were investigated. They contain bromine as R_1 and different substituents at the R_2

217   position to assess the importance of π-π stacking interactions involving F259. K_D values of 7.2 μM for

218   compound **13** and 13.8 μM for compound **11** were found with respect to the interaction with Dvl-3 PDZ.

219   Crystal structures of Dvl-3 PDZ in complex with compound **11** (1.58 Å resolution, 1 molecule per AU)

220   and **12** (1.48 Å, 2 molecules per AU) revealed very similar binding as observed in the crystal structures

221   with compounds **3** and **7**. The aromatic rings at R_2 show hydrophobic interactions to F259, but not a

222   classical π-π stacking as expected. Nevertheless, the tighter binding of compound **11** could be explained

223   by the larger aromatic substituent at the R_2 position compared to compound **12**. Both complex structures

224   show also non-specifically bound ligands in crystal contacts (Supporting Information Figure S3H,

225   Supporting Information Tables S2 and S3). The additional ligand molecules in both complex structures

226   can be explained as a crystallographic artefact, which is verified with ITC experiments that indicate 1:1

stoichiometries in both cases (Figure S5). With respect to the selectivity of the tested compounds we observed a 6 to 30-fold stronger binding of compounds **7**, **9**, **11** and **13** to Dvl-3 PDZ as compared to Dvl-1 PDZ. These differences are related to the different sequences at the end of αB. Most importantly, H324 is replaced by a serine residue in the Dvl-1 PDZ domain.

The group-3 compounds **(15-17)** contain a trifluoromethyl at position R_1 and were tested to investigate a cooperative role of this moiety with various substituents at position R_2. All compounds bind weaker to Dvl-1 and Dvl-3 than compound **8** which contains tetrahydronaphthalene at the R_1 position, revealing its important role in the interaction.

Further modifications towards higher affinity and reduced toxicity

Possible cytotoxic effects of compounds **3**, **7**, **8, 9**, and **10** were evaluated in cell viability assays using HEK293 cells (Supporting Information Figure S4). These compounds were selected due to different substituents at the R_1, including halogens. Cell viability was measured 24h after treatment with the individual compounds, and half maximal inhibitory concentrations (EC_{50}) were calculated for each compound. The compounds exhibited EC_{50} values in the range of 61-131 μM (Supporting Information Figure S4A). Compounds **3** and **10** that contained fluorine or methyl group substituents at R_2, respectively, were the least toxic, while compound **7**, containing bromine, was the most toxic. The results from crystallography, modelling studies and of the cell proliferation assays led us to further investigate compounds **18-21** that contain a methyl group at the R_1 position and different substituents as R_2. In this way, we aimed to develop both potent and less toxic, cell permeable inhibitors. All compounds showed strong interactions as indicated by chemical shift perturbation values between 0.30 to 0.34 ppm (Supporting Information Table S1). The binding constants were evaluated by ITC whereby compound **18** (K_D = 9.4 μM for Dvl-3 PDZ and 2.4 μM for Dvl-1 PDZ) appeared to be most potent. Compound **18** contains a pyrazole ring which is considered as an important biologically active heterocyclic moiety (Lv 2010). Compounds **20** (K_D = 9.8 μM for Dvl-3 PDZ and 4.7 μM for Dvl-1 PDZ) and **21** (K_D = 12.5 μM for Dvl-3 PDZ and 4.7 μM for Dvl-1 PDZ) contain pyrrole rings. Their binding constants almost have the same value despite the different substituents (bromine or chlorine) at the pyrrole rings. The binding of compounds **18-21** to both Dvl PDZ domains is mainly enthalpy-driven as

indicated in Table **2**, with a slightly stronger effect for Dvl-1 PDZ than for Dvl-3 PDZ. To our surprise, the crystal structure of Dvl-3 PDZ in complex with compound **18** shows the pyrazole substituent in the R_2 position orientated away from the binding pocket. Instead, a π-π stacking interaction with F259 was observed (Supporting Information Figure S3I). Cytotoxicity of **18-21** was determined *via* MTT assays (Mosmann 1983) that displayed viability up to concentrations above 150 µM (Supporting Information Figure S4B).

Compound	Dvl-3 PDZ				Dvl-1 PDZ			
	K_D (µM)	ΔH (kcal/mol)	TΔS (kcal/mol)	ΔG (kcal/mol)	K_D (µM)	ΔH (kcal/mol)	TΔS (kcal/mol)	ΔG (kcal/mol)
18	9.4 ± 0.6	-8.0	-1.2	-6.8	2.4 ± 0.2	-12.2	-4.7	-7.5
19	21.0 ± 1.7	-5.9	0.4	-5.5	8.0 ± 0.5	-7.3	-0.3	-7.0
20	9.8 ± 0.3	-10.4	-3.6	-6.8	4.7 ± 0.3	-9.4	-2.2	-7.2
21	12.5 ± 0.5	-5.9	0.7	-6.8	4.7 ± 0.2	-8.5	-1.5	-7.0
NPL-1011	79.7 ± 53.3							
Sulindac	8.3 ± 2.5							
CBC-322338/ 3289-8625	> 400 µM							
NSC668036	> 400 µM							
Ky-02327					8.3 ± 0.8[16g]			
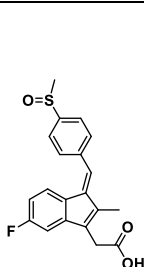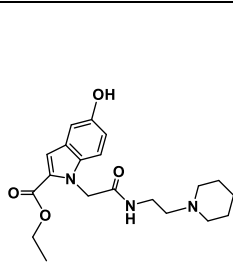
NPL-1011   **Sulindac**   **Ky-02327**   **NSC668036**   **CBC-322338 / 3289-8625**
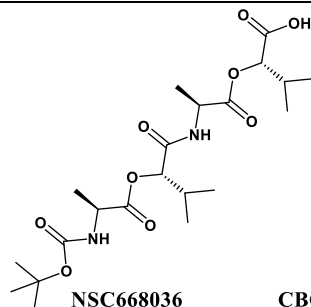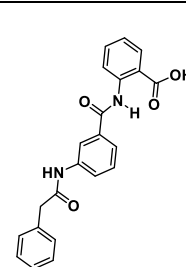
Table 2: Isothermal titration calorimetric data for the reaction between Dvl-3 PDZ, Dvl-1 PDZ and our compounds **18**, **19**, **20** and **21** respectively. Compounds NPL-1011 (Hori 2018), and Sulindac (Lee 2009), CBC-322338/3289-8625 (Grandy 2009, Hori 2018), and NSC668036 (Shan 2005), for more thermodynamic parameters see Supporting Information Figure S7. For Ky-02327 the value from literature is included.

Comparison to reported Dvl PDZ-binding molecules

Our compounds bind to Dvl-3 with a K_D better than 10 µM, and slightly tighter to Dvl-1, see Table 2, with **18** showing a K_D of 2.4 µM and chemical shift changes indicating binding to the canonical binding site (Figure 1A). For comparison, four compounds of those shown in Supporting Information Figure S6 were assayed by ITC (Supporting Information Figure S7) regarding their affinity to Dvl-3 PDZ. Ky-02327 was already determined to bind with a K_D of 8.3 ± 0.8 µM (Kim 2016) to Dvl-1 PDZ. Our first interest was oriented towards sulfonamides. Hori et al (Hori 2018) have recently reported 3-({3-[(2-carboxyphenyl)sulfamoyl]phenyl}sulfamoyl)benzoic acid (NPL-1011) binding to Dvl-1 PDZ via the

279   detection of chemical shift changes, and further sulfonamide compounds that showed smaller effects,

280   indicating weaker binding. We examined the binding constant of NPL-1011 which possesses two

281   sulfonamide moieties by ITC and found a value of 79.7 ± 53.3 µM, see Table 2. For further comparisons,

282   we assayed also CBC-322338/3289-8625, Sulindac and NSC668036 by ITC. Surprisingly, CBC-

283   322338/3289-8625 showed very low affinity, with a K_D above 400 µM in our ITC assay, in line with

284   the value found by Hori et al (Hori 2018) (954 +/- 403 µM). We also applied an NMR shift assay (Figure

285   S8), yielding a ΔCSP around 0.1. Based on NMR and ITC studies, the binding affinity of CBC-

286   322338/3289-8625 to DVL-3 seems to be less than 50 micromolar (comparing the CSPs from the NMR

287   assay with those of our other compounds listed in Table S1 and the respective binding constants in Table

288   1, and considering also the weak heat development in our ITC assay) which was larger than the originally

289   reported value (10.6 +/- 1.7) (Grandy 2009) that was obtained with a different method. Concerning non-

290   sulfonamide compounds, a K_D of 8.3 ± 2.5 µM was detected for Sulindac, while NSC668036 (Shan

291   2005) did not show high-affinity binding. These results are largely in agreement with literature. In all

292   cases, compounds were tested for purity after K_D measurements (see Supporting Information Figures

293   S9A-D).

294

295   **Selectivity testing using a set of selected PDZ domains**

296   Compounds **18**, **20** and **21** were tested towards other PDZ domains for selectivity. The set included

297   PSD95-PDZ 2 and 3, Shank-3, α-syntrophin, and AF-6 PDZ. According to the determined chemical

298   shift perturbations (Supporting Information Table S4), our compounds show no or very weak

299   interactions with the selected PDZ domains (0.05 < ΔCSP ≤ 0.1) ppm. These findings led to the

300   conclusion that our compounds show considerable selectivity towards Dvl PDZ domains. This

301   selectivity might be due to a unique feature of Dvl PDZ where R320 (Dvl-3 PDZ) or R322 (Dvl-1 PDZ)

302   are crucial for interactions, explaining selectivity with respect to other PDZ domains. In addition, the

303   large hydrophobic cavity for the side chain of the C-terminal residue of the interacting peptide is

304   occupied by a large moiety in case of compounds **18**, **20** and **21** which might not be accommodated in

305   most other PDZ domains.

306

307   **Dvl inhibitors antagonize canonical Wnt signalling and Wnt-related properties of cancer cells**

308 Taking advantage of a lentivirus that encodes GFP in a β-catenin/TCF-dependent fashion (TOP-GFP,

309 SABiosciences), a stable HEK293 reporter cell line was established to evaluate the inhibitory effect of

310 compounds **18**, **20** and **21** on canonical Wnt signalling activity. TOP-GFP expression in this cell line

311 was induced by the ligand Wnt3a, which directly activates the Frizzled-Dishevelled complex and

312 protects β-catenin from degradation by the destruction complex (Figure 2A). Remarkably, all three

313 compounds inhibited Wnt signalling induced by Wnt3a in a dose-dependent manner (Figure 2B),

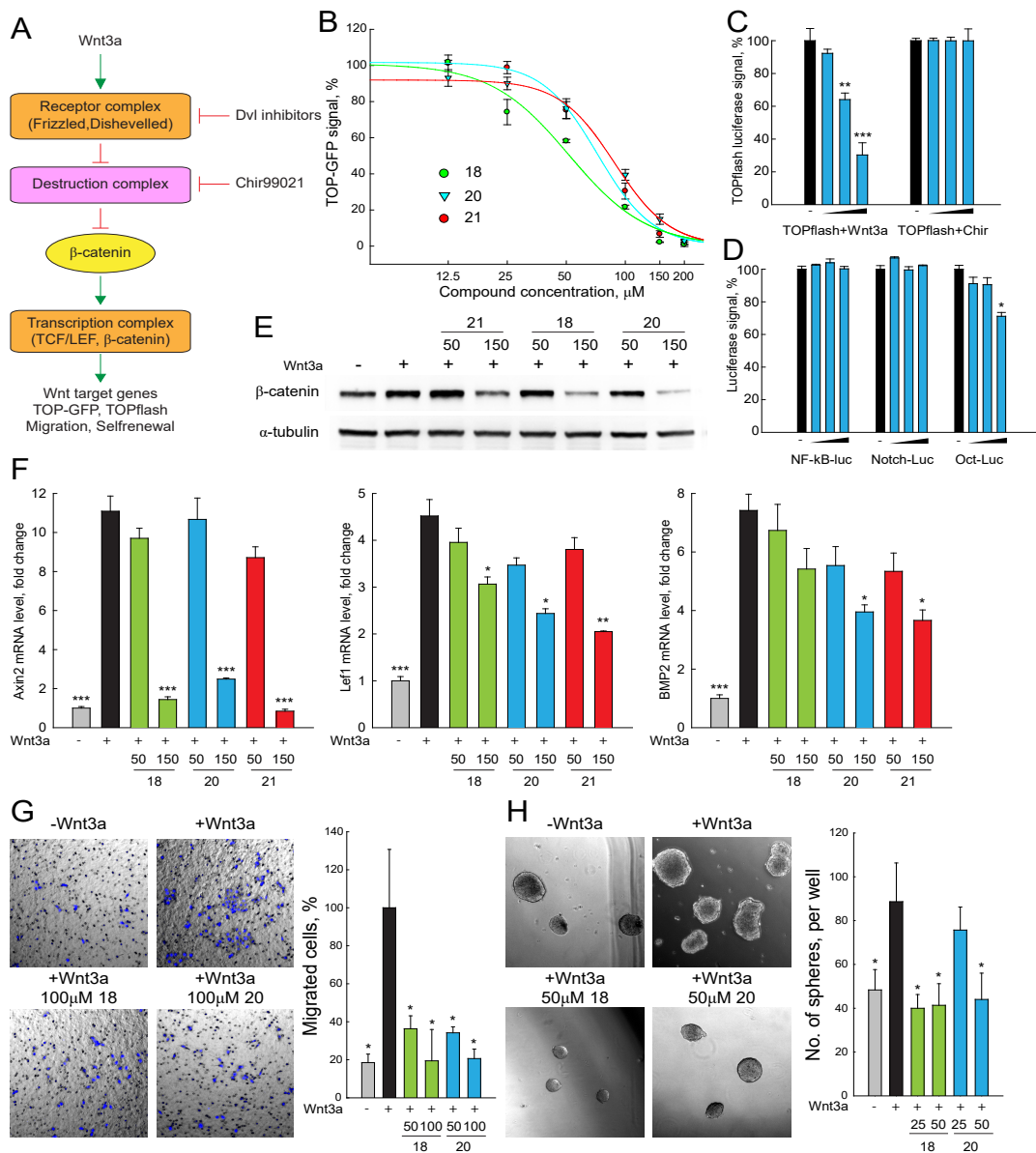314 yielding IC_{50} values between 50-80 μM.

Figure 2

315
316 **Figure 2.** DVL inhibitors antagonize Wnt signalling and Wnt related properties of cancer cells induced by Wnt3a. A. Scheme
317 of Wnt signalling pathway. Important components of the Wnt signalling pathway are schematically presented. Wnt3a treatment

13

318 increases the transcription of Wnt targets, enhances signals of TOP-GFP and TOPflash assays, and promotes Wnt related
319 biological properties of cancer cells. **B.** TOP-GFP reporter assays were performed with HEK293 reporter cell line. Compound
320 **18**, **20** and **21** inhibited Wnt3a induced Wnt activation in dose-dependent manner with IC_{50} of 50~75 μM. **C&D.** TOPflash
321 assays stimulated with Chir99021 and reporter assays of other pathway were used to evaluate the specificity of compound **20**.
322 Compound **20** specifically inhibited Wnt3a induced Wnt activation, and had no or mild effect on Chir99021 induced Wnt
323 activation and other signalling pathways including NF-κB, Notch and Oct4. **E.** β-Catenin protein levels were detected with
324 Western blotting in Hela cells. Compound **18**, **20** and **21** (150 μM) inhibited accumulation of β-catenin in Hela cells treated
325 with Wnt3a. **F.** The mRNA levels of Wnt target genes (Axin2, Lef1 and Bmp2) in Hela cells were measured with quantitative
326 real-time PCR. Compounds **18**, **20** and **21** (150 μM) reduced the transcription of Wnt target genes that are enhanced by Wnt3a
327 treatment in Hela cells. **G.** Cell migration of SW480 cells after Wnt3a treatment was assessed by transwell assays. Compounds
328 **18** and **20** (50~100 μM) reduced the migration of SW480 cells enhanced by Wnt3a. **H.** SW480 cells were cultured in serum-
329 free non-adherent condition to evaluate the self-renewal property enhanced by Wnt3a treatment. Compound **18** and **20** (25~50
330 μM) reduced sphere formation of SW480 cells that was enhanced by Wnt3a treatment. For all tests, three independent
331 biological replicas were performed and error bars represent standard deviations. P-values were calculated from T-test. *: P <
332 0.05; **: P < 0.01; ***: P < 0.001.

333
334

335 To further evaluate the specificity of our Dvl inhibitors, the conventional TOPflash (Molenaar 1996)

336 and other luciferase reporter assays were performed. In Hela cells, **20** inhibited TOP-luciferase signals

337 stimulated by Wnt3a but not by CHIR99021(Sineva 2010), a compound that activates Wnt signalling

338 downstream of Dvl (Figure 2A, C). Compound **20** had no significant inhibitory effects in reporter assays

339 that measure the activity of other signalling systems, e.g., NF-κB-luciferase stimulated by recombinant

340 TNFα, Notch-luciferase stimulated by the overexpression of the Notch intracellular domain, or the Oct-

341 luciferase assay that is stimulated by overexpression of Oct4 (SABiosciences, Figure 2D). These results

342 strongly indicate that **20** is specific for canonical Wnt signalling at the upstream level.

343 Increased β-catenin protein level is a hallmark of active Wnt signalling (Kishida 1999). Once β-catenin

344 is accumulated in the cytoplasm, it can translocate into the nucleus and activate the transcription of Wnt

345 target genes by interacting with transcription factors of the TCF/LEF family (Figure 2A) (Behrens

346 1996). In Hela cells, all three Dvl inhibitors blocked the increase of production of β-catenin by Wnt3a

347 in dose-dependent manners, as seen by Western blotting (Figure 2E). Increased mRNA levels of the

348 Wnt target genes Axin2, Lef1 and Bmp2 (Riese 1997, Jho 2002, Lewis 2010) were induced by Wnt3a

349 treatment, as measured by qRT-PCR, and these increases were reduced by compounds **18**, **20** and **21**

350 (Figure 2F). These results demonstrate that compounds **18**, **20** and **21** inhibit Wnt signalling as indicated

351 by reduced accumulation of β-catenin and low expression of typical Wnt target genes.

352 Canonical Wnt signalling contributes to cancer progression by inducing high motility and invasion of

353 cancer cells while retaining the self-renewal property of cancer initiating cells (Fritzmann 2009, Sack

354 2011, Vermeulen 2010, Malanchi 2008). In particular, cancer initiating cells are propagated and

355 enriched in non-adherent sphere culture, demonstrating the self-renewal capacity of the stem cells

356 (Kanwar 2010, Fan 2011). To investigate the potential value of the Dvl inhibitors for interfering with

357 these Wnt-related properties of cancer cells, the subline SW480WL was derived from the SW480 colon

358 cancer cell line, which exhibits a low level of endogenous Wnt activity (Fang 2012). The cell migration

359 and self-renewal properties of SW480WL cells were enhanced by Wnt3a treatment, as revealed by

360 transwell and sphere formation assays (Figure 2G, H). Compounds **18** and **20** prevented increased cell

361 migration and sphere formation. These results indicate that our Dvl inhibitors may be developed into

362 lead compounds that interfere with Wnt signalling.

363

364 **CONCLUSIONS**

365 In the present work, small molecules that bind to Dvl PDZ in the one-digit micromolar range with

366 considerable selectivity have been developed by an extensive structure-based design approach. With

367 regards to the affinity determined by ITC, compound **18** binds to Dvl-1 and Dvl-3 in vitro with K_D values

368 of 2.4 and 9.4, respectively, comparing very well with known ligands. X-ray structures of Dvl-3 PDZ

369 complexes with selected compounds provided insight into crucial interactions and served as the basis

370 for the design of tight binding compounds with reduced toxicity. The structural investigations revealed

371 that these compounds form hydrogen bonds with the amide groups of residues L260, G261 and I262 in

372 the PDZ-domain loop and the side chains of residues H324 and R320. Finally, the chosen methodology,

373 virtual screening followed by a two-stage NMR based screening, X-ray crystallography, and chemical

374 synthesis is an excellent path towards bioactive interaction partners. Our best compounds effectively

375 inhibited the canonical Wnt signalling pathway in a selective manner and could be developed for further

376 studies.

377

378 **Experimental Section**

379 **Clustering binding sites and selection of representative PDZ domains**

380 Three-dimensional structures of PDZ domains were retrieved from the PDB (Berman 2000). At the time

381 of the study from a total of 266 PDB files, 126 were NMR solution structures and 140 derived from X-

382 ray diffraction studies. The structures belong to 163 PDZ domains of 117 different proteins from 11

383 organisms. Files which contain more than one 3D conformation for a domain (up to 50 for NMR-derived

384 data) were split into separate structures and considered independently. The total number of unique 3D

385 structures was 2,708.

386 Amino acid sequences of PDZ domains were aligned using Clustal Omega software (Sievers 2011).

387 Based on the alignment, for each structure, residues which form the binding site (strand βB and helix

388 αB) were determined (Supporting Information Figure S8). The centre of the binding site was defined as

389 a geometric centre of Cα atoms of 7 residues (6 residues from the βB strand and the second residue from

390 the αB helix). Such bias toward the βB strand was made to cover sites occupied by residues in -1 and -

391 3 positions.

392 The triangulated solvent accessible surface for each PDZ structure was built using MSMS software

393 (Sanner 1996) with a spherical probe radius of 1.4 Å and vertex density 10 Å^{-1}. The largest connected

394 set of surface vertices within 9 Å from the centre of the binding site was used to construct shape-based

395 numerical descriptors. The descriptors are 508-dimensional vectors of non-negative integer numbers

396 and were built using a shape distributions approach (Osada 2002). In total 10 (Pawson 2007) vertex

397 triplets were selected randomly, each forming a triangle. Triangles which had a side longer than 16 Å

398 were discarded. Triangle sides were distributed into 16 length bins, each 1 Å wide, covering lengths

399 from 0 to 16 Å. A combination of three sorted side lengths, each belonging to one of 16 distance bins,

400 defines one of 508 categories of the triangles. The number of triangles of each category was calculated,

401 resulting in a 508-dimensional vector which is used as a numerical descriptor of the binding pocket

402 shape. For further operations with descriptors, Euclidian metric was introduced. Shape descriptors were

403 distributed into 6 clusters using k-means algorithm (Jain 1988). For each cluster, a centroid structure

404 was defined as the one, whose descriptor is the closest to mean descriptor for the cluster. The centroid

405 structures (2O2T#B.pdb, 1VA8#3.pdb, 2DLU#01.pdb, 1UHP#8.pdb, 2OS6#8.pdb, 3LNX#A.pdb) were

406 used for docking.

407

408 **PDZ targeted library design**

409 Screening collection by Enamine Ltd. (Chuprina 2010) containing a total of 1,195,395 drug-like

410 compounds was used as the primary source of small molecules. Natural ligand of PDZ is the C-terminus

411 of a peptide with carboxylic group making extensive hydrogen bond network with the "ΦGΦ" motif.

412   Since the carboxyl provides either of negative charge and hydrogen bond acceptor, we want our ligands

413   to retain at least one of these features. Therefore, we pre-filtered the stock library to bear chemical

414   groups which have negatively charged and/or hydrogen bond acceptor functionality. In total 65,288

415   compounds were selected for the virtual screening study. The selected 6 centroids of PDZ domains and

416   the prepared compound set were subjected to high-throughput docking using the QXP/Flo software

417   (McMartin 1997). Complexes were generated with 100 steps of sdock + routine, and 10 conformations

418   per complex were saved.

419   Processing of docking poses started with filtering by contact term *Cntc* from the QXP/Flo scoring

420   function. Entries with Cntc < 45 were discarded, which removed complexes with weak geometries of

421   bound ligands. The remained filtering was performed with the in-house MultiFilter program that allows

422   flexible geometry-based filtering. We applied two algorithms, *nearest-atom* filter and *hydrogen-bond*

423   filter. The former filters complexes by distance from a given protein atom to the nearest heavy ligand

424   atom, while in the latter, filtering is based upon the number of hydrogen bonds calculated for a given

425   complex geometry. With the *nearest-atom* routine we selected compounds that filled the P_0 pocket and

426   sterically mimicked binding of a peptide carboxylic group. Peptide group hydrogens of the "ΦGΦ" motif

427   and atoms forming the hydrophobic pocket were used for that. With the *hydrogen-bond* filter we selected

428   compounds that formed extensive hydrogen bonding with the PDZ domain. Both these properties might

429   have larger impact on binding rather than negative charge (Harris 2003). Details on atoms used for

430   filtering and thresholds for *hydrogen-bond* filtering, as well as the resulting number of compounds, are

431   provided in Supporting Information Table S5. Compounds from complexes which passed through these

432   filters were incorporated into a targeted library for the PDZ-domain family. The final library contained

433   1119 compounds in total.

434

435   **Screening of compounds**

436   Two-dimensional ^{1}H-^{15}N HSQC spectra were used to screen a library of 212 compounds designed by

437   the company Enamine for PDZ domains. 50 μM of ^{15}N-labeled protein samples were prepared in a 20

438   mM sodium phosphate buffer, containing 50 mM sodium chloride, 0.02% (w/v) NaN_3, at pH 7.4. Stock

439   solutions of small molecules were prepared in DMSO-*d6* at a concentration of 160 mM. A ^{1}H-^{15}N HSQC

440   spectrum of Dvl PDZ was acquired at 300 K with 5% DMSO-*d6* in the absence of ligand as reference

441   spectrum. Mixtures of 16 compounds were added to ^{15}N-labeled Dvl PDZ at 8-fold molar excess each.

442   The final concentration of DMSO-*d6* in the protein-ligand solutions was 5%. Spectra were acquired

443   with 8 scans and 256 points in the indirect dimension. Compound binding was deduced if the resonance

444   position of a cross-peak was significantly shifted compared to the reference spectrum. The active

445   compound was obtained through successive deconvolution. Experiments were recorded on a Bruker

446   DRX600 spectrometer equipped with a triple-resonance cryoprobe. The preparation of samples was

447   done automatically by a Tecan Genesis RSP 150 pipetting robot. Spectra were analysed using the

448   programs TOPSPIN and SPARKY.[47]

449

450   **Synthesis of compounds**

451   All reagents and starting materials were purchased from Sigma-Aldrich Chemie GmbH, ABCR GmbH

452   & Co.KG, alfa Aesar GmbH & Co.KG or Acros Organics and used without further purification. All air

453   or moisture-sensitive reactions were carried out under dry nitrogen using standard Schlenk techniques.

454   Solvents were removed by evaporation on a Heidolph Laborota 4000 with vacuum provided by a PC

455   3001 Vaccubrand pump. Thin-layer chromatography (TLC) was performed on plastic-backed plates pre-

456   coated with silica gel 60 F_{254} (0.2 mm). Visualization was achieved under an ultraviolet (UV) lamp (254

457   and 366 nm). Flash chromatography was performed using J.T Baker silica gel 60 (30-63 µm). Analytical

458   high-performance liquid chromatography (HPLC) was performed on a Shimadzu LC-20 (degasser

459   DGU-20A3, controller CBM-20A, autosampler SIL-20A) with a DAD-UV detector (SPD-M20A),

460   using a reverse-phase C18 column (Nucleodur 100-5, 5 µM, 250 mm x 4 mm, Macherey-Nagel, Düren,

461   Germany). Separation of compounds by preparative HPLC was performed on a Shimadzu LC-8A

462   system equipped with a UV detector (SPD-M20A), using a semi-preparative C18 column (Nucleodur

463   100-5, 5 µM, 250 mm x 10 mm, Macherey-Nagel) or preparative C18 column (Nucleodur 100-5, 5 µM,

464   250 mm x 21 mm, Macherey-Nagel). The detection wavelength was 254 nm. Gradients of acetonitrile-

465   water with 0.1% TFA were used for elution at flow rates of 1 mL/min, 8 mL/min, and 14 mL/min on

466   the analytical, semi-preparative and preparative columns respectively. Melting points (mp) were

467   determined with Stuart Melting Point Apparatus SMP3 and are not corrected. Mass spectra were

468 recorded on a 4000Q TRAP LC/MS/MS/ System for AB Applied Biosystems MDS SCIEX. NMR

469 spectra were recorded on a Bruker AV300 spectrometer instrument operating at 300 MHz for proton

470 frequency using DMSO-*d6* solutions. Chemical shifts were quoted relative to the residual DMSO peak

471 (^{1}H: δ = 2.50 ppm, ^{13}C: δ = 39.52 ppm). Coupling constants (J) are given in Hertz (Hz). Splitting patterns

472 are indicated as follows: singlet (s), doublet (d), triplet (t), quartet (q), multiple (m), broad (b). Purity of

473 each compound used for biological testing was $\geq$95% unless otherwise noted. The purity check of known

474 inhibitors purchased for comparison with our compounds are found in Supporting Information Figure

475 S9.

476

477 **Synthesis of compounds 8, 11 – 17**

479

R$_1$ = CF$_3$, Br, Cl

R$_2$ =
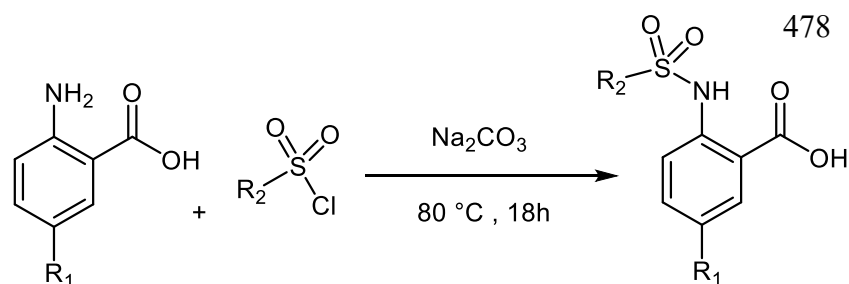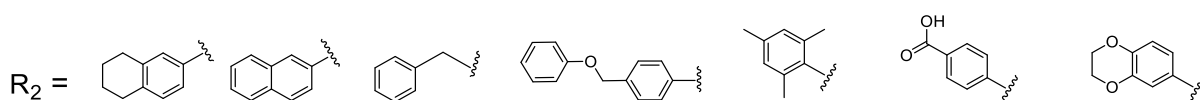
480 **Scheme 3**: Synthesis of compounds 8, 11 - 17

481 To a solution of anthranilic acid substituted with the appropriate R$_1$ (1.32 mmol) and sodium carbonate

482 (3.17 mmol) in water (2 mL) at 80 °C, the sulfonyl chloride (1.58 mmol) substituted with the appropriate

483 R$_2$ was added over a period of 5 minutes. The stirring continued for 18 h at 80 °C. The reaction mixture

484 was cooled to room temperature and acidified with 6 N HCl, and the resulting solid precipitate was

485 filtered, washed with water and dried to give the crude product. The final product was obtained by

486 preparative HPLC (Puranik 2008).

487

488 **2-(5,6,7,8-tetrahydronaphthalene-2-sulfonamido)- 5- (trifluoromethyl) benzoic acid (8)**

(0.52 g, 74% yield) **^{1}H-NMR** (300 MHz, DMSO-d6): δ = 11.77 [s, 1H, CO*OH*], 8.13 [s, 1H, *NH*],

7.85 [d, $^3J_{6,4}$ = 2.1 Hz, 1H, 6-H$_{Ar}$] 7.62 [d, $^4J_{1',3'}$ = 2.1Hz , 1H , 1'-H$_{Ar}$] 7.53 [dd, $^3J_{4,3}$ = 7.1 Hz, $^4J_{4,6}$ =

2.1 Hz, 4-H$_{Ar}$] 7.36 [dd, $^3J_{3'4'}$ = 7.5 Hz, $^4J_{3',1'}$ = 2.4 Hz, 1H, 3'-H$_{Ar}$] 7.15 [d, $^3J_{4',3'}$ = 7.5Hz, 1H, 4'-H$_{Ar}$]

, 6.90 [d, $^3J_{3,4}$ = 7.1Hz, 1H , 3-H$_{Ar}$] 2.73 (m, 4H, CH$_2$); 1.6 (m, 4H, CH$_2$);-**^{13}C-NMR** (75 MHz, DMSO-

d6): δ = 169.1(C, C$_{Ar}$-8], 152.7(C, C$_{Ar}$-2), 143.8 (C, C$_{Ar}$-4a'), 138.7(C, C$_{Ar}$-2'), 135.9 (C, C$_{Ar}$-8a'),

130.4(CH, C$_{Ar}$-4), 128.7 (CH, C$_{Ar}$-6), 127.5 (CH, C$_{Ar}$-1'), 124.0 (CH, C$_{Ar}$-4'), 121.6 (C, C-6), 118.2 (C,

C$_{Ar}$-5), 116.9 (C, C$_{Ar}$-3), 29.0 (CH$_2$, C-8'), 28.8 (CH$_2$, C-5'), 22.3 (CH$_2$, C-6'), 22.2 (CH$_2$, C-7'); mp:

177°C; MS (ESI) *m*/z:calcd. for C$_{18}$H$_{16}$F$_3$NO$_4$S, 399; found 400 [M+H]$^+$.

5-bromo-2-(naphthalene-2-sulfonamido) benzoic acid (11)

(0.13 g, 67% yield) **^{1}H-NMR** (300 MHz, DMSO-d6): δ = 10.2 [s, 1H, *COOH*], 9.8 [s, 1H, *NH*] 8.59 [d,

$^4J_{1',3'}$ = 1.4 Hz, 1 H, 1'-H$_{Ar}$], 8.17 [d, $^3J_{8',7'}$ = 7.8 Hz, 1 H, 8'-H$_{Ar}$], 8.10 [d, $^3J_{4'3'}$ = 8.8 Hz, 1 H, 4'-H$_{Ar}$],

8.02 [d, $^3J_{5',6'}$ = 7.8 Hz, 1 H, 5'-H$_{Ar}$], 7.93 [d, $^4J_{6,4}$ = 2.4 Hz, 1 H, 6-H$_{Ar}$], 7.77 [dd, $^3J_{3',4'}$ = 8.8 Hz, $^4J_{3',1'}$

= 1.4Hz ,1 H, 3'-H$_{Ar}$], 7.72 – 7.65 [m, 3 H, 4-H$_{Ar}$, 6'-H$_{Ar}$, 7'-H$_{Ar}$] , 7.51 [d, $^3J_{3,4}$ = 8.9 Hz, 1 H, 3-H$_{Ar}$]. −

^{13}C-NMR (75 MHz, DMSO-d6): δ = 168.2 (C, C-7) ,138.8 (C, C$_{Ar}$-2),136.8 (CH, C$_{Ar}$-4),135.3 (C, C$_{Ar}$-

4a'),134.4 (C, C$_{Ar}$-8a') ,133.4 (CH, C$_{Ar}$-6) ,131.4 (CH, C$_{Ar}$-6'), 129.3 (CH, C$_{Ar}$-4') ,128.5 (CH, C$_{Ar}$-

8'),127.8 (2xCH, C$_{Ar}$-5', C$_{Ar}$-7') 121.6 (CH, C$_{Ar}$-3'),120.6 (CH, C$_{Ar}$-3),119.0(C, C$_{Ar}$-1) ,114.9 (C, C$_{Ar}$-

5). Mp: 199°C; (ESI) *m*/z: calcd.for C$_{17}$H$_{11}$BrNO$_4$S$^-$; 403.9560: found   403.9613 [M-H]$^-$.

5-bromo-2-(phenylmethylsulfonamido)benzoic acid (12)

 (0.07g, 42% yield) **^{1}H-NMR** (300 MHz, DMSO-d6): δ = 10.57 [s, 1 H, COO*H*], 8.05 [d, $^4J_{6,4}$ = 2.4

Hz, 1 H, 6-H$_{Ar}$], 7.75 [dd, $^3J_{4,3}$ = 8.9 Hz, $^4J_{4,6}$ = 2.4Hz ,1 H, H-4$_{Ar}$], 7.49 [d,$^3J_{3,4}$ = 8.9 Hz,1 H, 3-H$_{Ar}$],

7.33 – 7.28 [m, 3 H, 3'-H$_{Ar}$, 5'-H$_{Ar}$], 7.23 – 7.20 [m, 2 H, 4'-H$_{Ar}$], 5.75 [s, 1 H, N*H*], 4.72 [s, 2 H, 1'-

H] **^{13}C-NMR** (75 MHz, DMSO-d6): δ = 168.3 (C, C-7) , 139.9 (C, C$_{Ar}$-2), 137(CH, C$_{Ar}$-4), 133.4 (CH,

C$_{Ar}$-6), 130.7 (CH, C$_{Ar}$-3') , 128.6 (C, C$_{Ar}$-2') , 128.4 (CH, C$_{Ar}$-5') , 128.3 (CH, C$_{Ar}$-4') , 119.5 (CH,

C$_{Ar}$-3) , 117.5 (C, C$_{Ar}$-1)   , 113.9 (C, C$_{Ar}$-5) , 57.4 (CH$_2$, C-1'). Mp: 216°C; (ESI) *m*/z: calcd.for

C$_{14}$H$_{11}$BrNO$_4$S$^-$ 367.9860; found 367.9878 [M-H]$^-$.

5-bromo-2-(4-(phenoxymethyl)phenylsulfonamido)benzoic acid (13)

(0.6 g, 29% yield) 1H-NMR (300 MHz, DMSO-d6): δ = 7.97 [d, $^4J_{6,4}$ = 2.4 Hz, 1 H, 6-H$_{Ar}$), 7.85 (d,

$^3J_{2',3'}$ = 8.3 Hz, 2 H, 3'-H$_{Ar}$), 7.73 [dd, $^3J_{4,3}$ = 8.9 Hz, $^4J_{4,6}$ = 2.4Hz , 1 H,4-H$_{Ar}$], 7.63 [d, $^3J_{2',3'}$ = 8.3 Hz, 2

H, 2′-H$_{Ar}$], 7.47 [d, $^3J_{3,4}$ = 8.9 Hz, 1 H, 3-H$_{Ar}$], 7.29 [dd, $^3J_{3'',2''}$ = $^3J_{3'',4''}$=7.3 Hz, 2 H, 3''-H$_{Ar}$], 7.00 – 6.92

[m, 3 H, 4''-H$_{Ar}$, 2''-H$_{Ar}$], 5.17 [s, H, 5′-H]. − **13C-NMR** (75 MHz, DMSO-d6): δ = 168.2 (C, C-7) ,

157.9 (C, C$_{Ar}$-1'') , 143.2 (C, C$_{Ar}$-4′), 138.8 (C, C$_{Ar}$-2), 137.5 (C, C$_{Ar}$-1′) , 136.9 (CH, C$_{Ar}$-4) 133.5 (CH,

C$_{Ar}$-6), 129.4(CH, C$_{Ar}$-3''), 128.1(CH, C$_{Ar}$-2′),127.0 (CH, C$_{Ar}$-3′) , 120.9 (CH, C$_{Ar}$-4'') , 120.5 (CH, C$_{Ar}$-

3) , 119.0 (C, C$_{Ar}$-1) , 114.9(CH, C$_{Ar}$-5), 114.7 (CH, C$_{Ar}$-2''), 68.0 (CH$_2$, C-5′) Mp: 175°C; (ESI) m/z:

calcd for C20H15BrNO5S- 459.9860 found 459.9878 [M-H]-.

5-bromo-2-(2,4,6-trimethylphenylsulfoamido)benzoic acid (14)

(0.6 g, 78% yield) **^{1}H-NMR** (300 MHz, DMSO-d$_6$): δ = 11.77 [s,1H, *COOH*], 9.98 [s, 1H, *NH*], 7.68

[d, $^3J_{6,4}$ = 7.4 Hz, 1H, 6-H$_{Ar}$], 7.51[dd, $^3J_{4,3}$ =7.1Hz, $^4J_{4,6}$ =7.4 Hz, 1H 4-H$_{Ar}$], 7.17 [d, 2H, 4′-H$_{Ar}$, 6′-

H$_{Ar}$], 7.14 [d, $^3J_{3,4}$= 1H,3-H$_{Ar}$] , 2.56 [s, 6H, CH$_3$, 9′-H, 7′-H], 2.21 [s, 3H, CH$_3$, 8′-H];-**13C-NMR** (300

MHz, DMSO-d$_6$): δ = 168.8 (C, C-7), 143.3 (C, C$_{Ar}$-2), 139.5 (C, C$_{Ar}$-2′), 139.0 and 139.0 (2xC, C$_{Ar}$-

3′, C$_{Ar}$-1′) 137.3 (CH, C$_{Ar}$-4), 134.0 (CH, C$_{Ar}$-6′), 133.0 (CH, C$_{Ar}$-6), 132.5 and 132.5 (2XCH, C$_{Ar}$-4′,

C$_{Ar}$-6′) 119.1(CH, C$_{Ar}$-3), 117.9(C, C$_{Ar}$-5), 114.3 (C, C$_{Ar}$-1), 22.5 and 22.5 (2 x CH$_3$, C-7′, C-9′) 20.7

(CH$_3$, C-8′) ;   mp: 185; MS (ESI): m/z 399 [M+H]+.

2-(4-acetylphenylsulfoamido)-5-(trifluoromethyl)benzoic acid (15)

 (0.4 g, 63% yield) **^{1}H-NMR** (300 MHz, DMSO-d6): δ = 12.28 [s, 1H, *COOH*]; 12.10 [s, 1H, *NH*], 8.11

[d, $^4J_{6,4}$= 2.5 Hz, 1H, 6-H$_{Ar}$], 8.08 [d, $^3J_{3'2'}$ = 7.5 Hz, 2H, 3′-H$_{Ar}$], 7.86 [dd, $^4J_{4,6}$ = 2.5 Hz, $^3J_{4,3}$ = 7.3Hz,

1H, 4-H$_{Ar}$], 7.64 [d, $^3J_{4,3}$ = 7.3 Hz, 1H, 3-H$_{Ar}$] , 7.56 [dd $^3J_{2',3'}$ = 7.5Hz, $^4J_{2',6'}$ =2.3Hz, 2H, 2′-H$_{Ar}$, 6′-

H$_{Ar}$] 7.22 [dd , $^3J_{3',2'}$ = 7.5Hz , $^4J_{3',5'}$ = 2.1Hz , 2H, 3′-H$_{Ar}$, 5′-H$_{Ar}$] 2.50 [s, 3H, CH$_3$, 8′-H]; - **^{13}C-NMR**

(75 MHz, DMSO-d6): δ = 197.9 (C, C-7′), 169.1(C, C-8), 151.8 (C, C$_{Ar}$-2) 143.5 (C, C$_{Ar}$-1′), 142.5 (C,

C$_{Ar}$-4′), 140.6 (CH, C$_{Ar}$-4), 131.4 (CH, C$_{Ar}$-7), 129.6 (2XCH, C$_{Ar}$-3′, C$_{Ar}$-5′), 128.6 (2xCH, C$_{Ar}$-2′, C$_{Ar}$-

6′), 127.6 (C, C$_{Ar}$-6), 123.0 (C, C-$_{Ar}$-5), 118.7 (CH, C$_{Ar}$-3), 27.3 (CH$_3$, C-8′); mp: 170°C; MS (ESI) *m*/z

: calcd. for C$_{16}$H$_{12}$F$_3$NO$_5$S. 387; found 388 [M+H]$^+$.

2-(2,3-dihydrobenzo[*b*][1,4]dioxine-6-sulfonamido)-5-(trifluoromethyl)benzoic acid (16)

(0.4 g, 65% yield) **^{1}H-NMR** (300 MHz, DMSO-d6): δ = 11.48 [s, 1H, *COOH*], 8.13[s, 1H, *NH*] , 7.89

[d, $^4J_{6,4}$ = 3,9 Hz , 1H , 6-H$_{Ar}$] 7.66 [dd, $^3J_{4,3}$ = 7.2 Hz, $^4J_{4,6}$ = 4.3 Hz, 1H, 4-H$_{Ar}$],

7.23 [d, $^3J_{4,3}$ = 7,2 Hz 1H, 3-H$_{Ar}$],   7.11 [dd, $^3J_{2',3'}$ = 7.3Hz , $^4J_{2',8'}$= 3.2Hz , 1H, 2′-HAr] 6.95 [d, $^4J_{2',8'}$=

3.2 Hz, 1H , 8′-H$_{Ar}$] 4.23 – 4.31 [m, 4H, 5′-H, 6′-H]; - **^{13}C-NMR** (75-MHz, DMSO-d6): δ = 168.9(C,

C-8), 148.3(C, C-4′), 143.8 (C, C-2), 143.5 (C, C-7′) , 131.3 (C, C-1′), 130.8 (CH, C-4), 128.6(CH, C-6), 125.7 (C, C-7), 122.1(C, C-5), 120.9(CH, C-2′), 118.3 (CH, C-3), 118.1(CH, C-3′), 116.8 (CH, C-8′) , 64.7(CH$_2$, C-5′) 64.3 (CH$_2$, C-6′); mp: 178°C; MS (ESI) m/z: calcd. for C$_{16}$H$_{12}$F$_3$NO$_6$S. 403; found 404 [M+H]$^+$.

5-(trifluoromethyl)-2-(2,4,6-trimethylphenylsulfoamido)benzoic acid (17)

(0.38 g, 62% yield) **^{1}H-NMR** (300 MHz, DMSO-d6): δ = 12.28 [s, 1H, *COOH*], 11.60 [s, 1H, *NH*], 8.15 [d, $^4J_{6,4}$ = 4.3 Hz, 1H, 6-H$_{Ar}$] 7.92 [dd, $^3J_{4,3}$ =7.9 Hz, $^4J_{4,6}$= 2.1 Hz, 1H,4-H$_{Ar}$] 7.87 [d, $^4J_{6′,4′}$= 1.9 Hz, 2H, 4′-H$_{Ar}$, 6′-H$_{Ar}$], 7.48 [d, $^3J_{3,4}$ = 7.9 Hz, 1H, 3-H$_{Ar}$] , 2.60 [s, 6H, CH$_3$, 9′-H, 7′-H], 2.23 [s, 3H, CH$_3$, 8′-H]; - **^{13}C-NMR** (75 MHz, DMSO-d6): δ = 169.3 (C, C-7), 154.2 (C, C-2), 143.6 (C, C-2′) , 139.1 and 139.1 (2xC, C-1′, C-3′) 132.9 (C, C-5′), 132.5 (CH, C-4), 131.5 and 131.5 (2xCH, C-4′, C-6′) , 130.1(CH, C-6), 128.7 (C, C-8), 122.5 (C, C-5), 117.0 (CH, C-3), 109.0 (C, C-1) ,22.4 and 22.4 (2xCH$_3$,C-7′, C-9′), 20.8 (CH$_3$, C-8′) ; mp:184°C; MS (ESI) m/z: calcd. for C$_{17}$H$_{16}$F$_3$NO$_4$S; 387: found 388 [M+H]$^+$.

18, 19, 20, and **21** were purchased from Enamine, Kiev, Ukraine as pure compounds (see also Table S6, Supporting Information).

Determination of ligand binding and binding constant by NMR

50 μM of ^{15}N-labeled protein samples were prepared in a 20 mM sodium phosphate buffer containing 50 mM sodium chloride, 0.02 % (w/v) NaN$_3$, at pH 7.4. Stock solutions of small molecules were prepared in DMSO-*d6* at a concentration of 160 mM. A ^{1}H-^{15}N HSQC spectrum of Dvl PDZ was acquired at 300 K with 5% DMSO-*d6* in the absence of ligand as reference spectrum. Mixtures of 16 compounds were added to ^{15}N-labeled Dvl PDZ at 8-fold molar excess each. The final concentration of DMSO-*d6* in the protein-ligand solutions was 5%. Spectra were acquired with 8 scans and 256 points in the indirect dimension.

Binding was deduced if the resonance position of a cross-peak was significantly shifted compared to the reference spectrum. The active compound was obtained through successive deconvolution. Experiments were recorded on a Bruker DRX600 spectrometer equipped with a triple-resonance cryoprobe. The

574 preparation of samples was done automatically by a Tecan Genesis RSP 150 pipetting robot. Spectra

575 were analysed using the programs TOPSPIN and SPARKY.

576 Chemical shift perturbations were obtained by comparing the ^{1}H-^{15}N backbone resonances of protein

577 alone to those of protein-ligand complex. The mean shift difference ($\Delta\delta$ in ppm) was calculated using

578 the equation 1 (Garrett 1997, Bertini 2011).

579
$$\Delta\delta = \sqrt{\frac{[\Delta\delta_H]^2 + [\Delta\delta_N/5]^2}{2}} \qquad (Eq.\,1)$$

580 Here $\Delta\delta_N$ and $\Delta\delta_H$ are the amide nitrogen and amide proton chemical shift differences between the free

581 and the bound states of the protein. In order to estimate binding constants, titration experiments

582 monitored by NMR were done. A series of ^{1}H-^{15}N HSQC were recorded as a function of ligand

583 concentration. Residues showing a continuous chemical shift change and for which the intensity

584 remained strong were classified as being in fast exchange. The dissociation binding constant was

585 estimated by fitting the observed chemical shift change to equation 2 (Shuker 1996, Hajduk 1997).

586
$$\frac{\Delta\delta}{\Delta\delta_{max}}$$

587
$$= \frac{([L_T] + [P_T] + K_D) - \sqrt{([L_T] + [P_T] + K_D)^2 - 4[L_T] \cdot [P_T]}}{2[P_T]} (Eq.\,2)$$

588

589 $\Delta\delta$ is the observed protein amide chemical shift change at a given compound concentration and $\Delta\delta_{max}$

590 the maximum chemical shift change at saturation. [L_T] the total concentration of the compound, and [P_T]

591 the total concentration of the protein. K_D is the equilibrium dissociation constant. The K_D values are

592 reported as means ± standard deviations of at least six residues influenced upon binding of the ligand.

593

594 **Determination of binding constant by Isothermal Titration Calorimetry (ITC)**

595 Isothermal Titration Calorimetry (ITC) experiments were performed using a VP-ITC system

596 (MicroCal). Protein in 20 mM Hepes buffer, 50 mM NaCl, pH 7.4, was centrifuged and degassed before

597 the experiment. A 200 µM ligand solution containing 2% DMSO was injected 30 times in 10 µL aliquots

598 at 120 s intervals with a stirring speed of 1000 rpm into a 1.4 mL sample cell containing the Dvl PDZ

599   domain at a concentration of 20 μM at 25 °C. Control experiment was initially determined by titrating

600   ligand into buffer at same conditions. Titration of ligand into buffer yielded negligible heats.

601   Thermodynamic properties and binding constants were determined by fitting the data with a nonlinear

602   least-squares routine using a single-site binding model with Origin for ITC v.7.2 (Microcal).

603

604   **Protein expression**

605   PDZ domains of human AF6 (P55196-2, residues 985–1086) and murine α1-syntrophin (Q61234,

606   residues 81–164) were cloned into pGEX-6P-2 (Amersham Biosciences, Freiburg, Germany) and

607   pGAT2 (European Molecular Biology Laboratory, Heidelberg, Germany), respectively. Proteins were

608   expressed in *E. coli* BL21 (DE3) cells and purified as previously described (Boisguerin 2004). For the

609   cloning of the Dvl-1 PDZ domain (O14640, residues 245–338), IMAGp958J151157Q (ImaGenes) was

610   used as template. V250 is exchanged to isoleucine as in human Dvl-3 or murine Dvl-1. The C-terminal

611   C338 of the domain was exchanged by serine. Via cloning in pET46EK/LIC, a coding sequence for a

612   TEV (Tobacco Etch Virus) protease cleavage site was introduced. The resulting plasmid pDVL1 was

613   transformed in *E. coli* BL21 (DE3). Expression on two-fold M9 minimal medium with 0.5 g/L ^{15}N

614   NH_4Cl as sole nitrogen source in shaking culture was done at 25 °C overnight with 1 mM IPTG. A yield

615   of 25 mg of pure Dvl-1 was obtained from 1 L culture after IMAC, TEV protease cleavage, a second

616   IMAC, and gel filtration (Superdex 75). The protein domain Dvl-1_245–338 was supplied for NMR in

617   20 mM phosphate buffer, pH 7.4, 50 mM NaCl.

618   The production of Dvl-3 (Q92997 residues 243-336), mShank3 (Q4ACU6, residues 637-744) PDZ

619   domains and the 3 PDZ domains of PSD95 was described by Saupe et al (Saupe 2011).

620

621   **Crystallization and X-ray diffraction**

622   The His-tagged cleaved human Dvl-3 PDZ domain was concentrated to 12-20 mg/mL in the presence

623   of a 5-fold molar excess of compound **3**, **5**, **6**, **7**, **11** and **12**. Crystals of all complexes were grown at

624   room temperature by the sitting drop vapour-diffusion method. 200 nL Dvl-3/compound solution was

625   mixed with an equal volume of reservoir solution using the Gryphon (Formulatrix) pipetting robot.

626   Crystals of all complexes were grown to their final size within 4 to 14 days. The Dvl-3 PDZ domain

627 crystallized in complex with compound **3**and **7** in crystallization condition 30% PEG 8000, 0.2 M

628 ammonium sulphate, 0.1 M MES pH 6.5; with compound **5** in 30% PEG 8000, 0.1 M MES pH 5.5; with

629 compound **6** in 1.2 M ammonium sulphate, 0.1 M citric acid pH 5.0; with compound **12** in 32% PEG

630 8000, 0.2 M ammonium sulphate, 0.1 M Na-cacodylate pH 6.0; with compound **11** in 1 M ammonium

631 sulphate, 1% PEG 3350, 0.1 M Bis-Tris pH 5.5; with compound **12** in 1.26 M sodium phosphate, 0.14

632 M potassium phosphate and with compound **18** in 1.5 M ammonium sulphate, 12% glycerol, 0.1 M Tris-

633 HCl pH 8.5. The crystals were cryoprotected if necessary, for data collection by soaking for few seconds

634 in precipitant solution containing 20% (v/v) glycerol and subsequently frozen in liquid nitrogen.

635 Diffraction data were collected at 100 K at beamline BL14.1 at the synchrotron-radiation source BESSY,

636 Helmholtz-Zentrum Berlin and processed with XDS.

637

638 **Structure determination and refinement**

639 Phases for the Dvl-3 PDZ domain in complex with compound 3 were obtained by molecular replacement

640 with PHASER (McCoy 2007) using the *Xenopus laevis* Dishevelled PDZ domain structure (PDB code

641 2F0A) as a starting model. The reasonable crystal packing and electron density allowed further model

642 and compound building using the program COOT (Emsley 2004) with iterative refinement with

643 REFMAC (Murshudov 1997). All further complex structures were obtained in the same way but using

644 the final refined compound free Dvl-3-PDZ structure as model for molecular replacement. The

645 Ramachandran statistics were analysed by Molprobity (Chen 2010) for all complexes and all

646 crystallographic statistics are given in Supporting Information Tables S2 and S3. Figures were prepared

647 with PyMol. Atomic coordinates and structure factor amplitudes for DVL-3 PDZ domain in complex

648 with compound **3**, **5**, **6**, **7**, **11**, **12**, **18** were deposited in the Protein Data Bank with accession codes

649 6ZBQ, 6ZBZ, 6ZC3, 6ZC4, 6ZC6, 6ZC7 and 6ZC8, respectively.

650

651 **MTT assay**

652 HEK293 cells were plated on a 96-well plate and treated with different concentrations of Dvl inhibitors.

653 After 24 h treatment, 20 µl of MTT solution (5 mg/mL) was added into each well. After 2 h incubation,

654   cell culture medium was replaced with 50 µL DMSO, and the signal of the purple formazan, produced

655   by living cells, was measured by a plate reader.

656

657   **TOP-GFP reporter assay**

658   The lentivirus particle (CCS-018L, SABiosciences) encoding GFP under the control of a basal promoter

659   element (TATA box) joined to tandem repeats of a consensus TCF/LEF binding site was transfected

660   into HEK293 cells. Stable cells were selected by puromycin (2 µg/mL) treatment. Wnt signalling

661   activity indicated by GFP intensity was measured by flow cytometry after 24 h incubation with

662   recombinant mouse Wnt3a (100 ng/mL) or GSK3 inhibitor CHIR99021 (3 µM) in the presence of Dvl

663   inhibitors.

664

665   **Luciferase reporter assays**

666   Plasmids encoding a firefly luciferase reporter gene under the control of different responsive elements

667   were transfected into Hela cells with a pRL-SV40 normalization reporter plasmid using the

668   Lipofectamine 2000 (Invitrogen). After desired treatment, cells were harvested in the passive lysis buffer

669   (Promega), and 15 µL cell lysate were transferred to 96-well LumiNunc plates (Thermo Scientific).

670   Firefly luciferase and Renilla luciferase were detected with the D-luciferin buffer (75 mM Hepes, 4 mM

671    $MgSO_4$, 20 mM DTT, 100 µM EDTA, 0.5 mM ATP, 135 µM Coenzyme A and 100 µM D-Luciferin

672   sodium salt, pH 8.0) and the coelenterazine buffer (15 mM Na4PPi, 7.5 mM NaAc, 10 mM CDTA, 400

673   mM Na_2SO_4, 25 µM APMBT and 1.1 µM coelenterazine, pH 5.0) respectively using the CentroXS

674   LB960 lumiometer (Berthold Technologies).

675

676   **Immunoblotting**

677   To assess the β-catenin accumulation in Hela cells, cells were treated with Wnt3a in the presence of Dvl

678   inhibitors for 24 h and lysed in RIPA buffer (50 mM Tris, pH 8.0, 1% NP-40, 0.5% deoxycholate, 0.1%

679   SDS, 150 mM NaCl). Equal amounts of protein were loaded on a SDS-PAGE. Separated proteins were

680   blotted onto PVDF membranes for immunoblot analysis using anti-β-catenin antibody (610154, BD).

681   HRP-conjugated anti-mouse antibody (715-035-150, Jackson ImmunoResearch laboratories) was used

682 for secondary detection with Western lightning chemiluminescence reagent plus (PerkinElmer) and

683 Vilber Lourmat imaging system SL-3.

684

685 **qRT-PCR analysis**

686 To measure the Wnt target accumulation at mRNA level, Hela cells were treated with Wnt3a in the

687 presence of Dvl inhibitors for 24 h. mRNA was extracted according to the standard TRIzol® protocol

688 (Invitrogen) and reverse-transcribed using random primers (Invitrogen) and M-MLV reverse

689 transcriptase (Promega). The qRT-PCR was performed in iQ5 Multicolor Real-Time PCR Detection

690 System (Bio-Rad) using SYBR® Green (Thermo Scientific) and gene-specific primer pairs of Bmp2,

691 Axin2, Lef1 and β-actin (endogenous control).

692

693 **Migration assay**

694 Cell motility was assessed using 24-well transwell (pore diameter: 8 μm, Corning). SW480WL cells

695 were seeded in the upper chamber in serum free DMEM with 0.1% BSA; 20% serum was supplemented

696 to medium in the lower chamber. After incubation with Wnt3a in the presence of Dvl inhibitors for 24

697 h, nonmigrant cells were scraped off using a cotton swab; the migrated cells on the filters were stained

698 with DAPI, photographed and counted.

699

700 **Colon sphere culture**

701 SW480WL cells were trypsinised into single cells, seeded on 24-well cell culture plates precoated with

702 250 μl polyhema (12 mg/mL in 95% ethanol, Sigma) per well, and incubated with Wnt3a in the presence

703 of Dvl inhibitors in the sphere culture medium (F12 : DMEM 1 : 1, 1X B-27 supplement, 20 ng/mL

704 EGF, 20 ng/mL FGF, 0.5% methylcellulose) for 10 days. Numbers of spheres were then counted under

705 the microscope.

706

707 **Notes**

708 The authors declare no competing financial interest.

709

ABBREVIATIONS USED

NMR, nuclear magnetic resonance; HSQC, Heteronuclear Single Quantum Correlation;

AU, asymmetric unit; SAR, derive structure activity relationships; vdW, van der Waals;

ITC, Isothermal titration calorimetry; PDZ, PSD95/Disc large/Zonula occludens 1);

Dvl, Dishevelled; PPI, protein-protein interactions; PDB, Protein Data Bank;

CSP, chemical shift perturbation; GFP, green fluorescent protein; DMSO, dimethyl sulfoxide;

PEG, polyethylene glycol; RNA, ribonucleic acid; mRNA, messenger RNA;

qRT-PCR, quantitative real-time polymerase chain reaction; DMEM Dulbecco's modified Eagle's medium ; BSA, bovine serum albumin;

ASSOCIATED CONTENT

Accession Codes

Atomic coordinates and structure factor amplitudes for DVL3 PDZ domain in complex with compound **3**, **5**, **6**, **7**, **11**, **12**, **18** were deposited in the Protein Data Bank with accession codes 6ZBQ, 6ZBZ, 6ZC3, 6ZC4, 6ZC6, 6ZC7 and 6ZC8, respectively. Authors will release the atomic coordinates and experimental data upon article publication.

Supporting information

1. Structure-based alignment of the amino acid sequences of Dvl-1,2,3 PDZ ; PSD95-PDZ-1,2,3 ; Af-6 and Syn PDZ domains. (S.2)

2. 1H-15N HSQC spectra of Dvl-3 PDZ domain alone and in the presence of varying concentrations of compound 3. (S.3)

738   **3.** Detailed views of diverse compounds bound to the Dvl-3 PDZ domain. (S.4)

739   **4.** Cell viability assays of compounds 3, 7,8, 9, 10, (A) and 18, 20, 21 (B). (S.5)

740   **5.** ITC binding assays of compound 18 with Dvl-3 PDZ (A) and with Dvl-1 PDZ (B). (S.5)

741   **6.** Structures of selected compounds used for comparison to our compounds. (S.6)

742   **7.** ITC data of selected compounds used for comparison to our compounds. (S.7)

743   **8.** NMR binding assay with compound 322338/3289-8625. (S.8)

744   **9.** Purity check of compounds. (S.9)

745   –   Purity check of NPL-1011 compound. (S.9)

746   –   Purity check of Sulindac compound. (S.10)

747   –   Purity check of CalBioChem-322338 compound. (S.11)

748   –   Purity check of NSC668036 compound. (S.12)

749   –   LCMS of intermediate compound 8. (S.13)

750   –   LCMS of intermediate compound 14. (S.13)

751   **10.** Chemical shift perturbation values of Dvl-3 PDZ and Dvl-1 PDZ for compounds (3-21). (S.14)

752   **11.** Data collection and refinement statistics of compounds 3, 5, 6, 7. (S.15)

753   **12.** Data collection and refinement statistics of compounds 11, 12, 18. (S.16)

754   **13.** Selectivity of ligands derived from chemical shift perturbation of compounds tested at other PDZ

755   domains. (S.17)

756   **14.** Details of Multifilter routines. (S.17)

757   **15.** Smiles codes and Compounds ID. (S.18)

758   **16.** NMR characterization of synthesized compounds (8, 11, 13, 14, 15, 16, 17). (S.21)

759

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