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Large-scale synthesis of Notum inhibitor 1-(2,4-dichloro-3-(trifluoromethyl)-phenyl)-1*H*-1,2,3-triazole (ARUK3001185) employing a modified Sakai reaction as the key step[†]

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1-Phenyl-1*H*-1,2,3-triazole **1** (ARUK3001185) was prepared on large scale from aniline **4** by application of both (1) a copper catalyzed azide–alkyne cycloaddition (CuAAC) with (trimethylsilyl)acetylene, and (2) a Clark modification of the Sakai reaction. The one-pot Sakai–Clark method with (MeO)₂CHCH=NNHTos (**2b**) proved to be superior as it was operationally simple, metal-free, and avoided the use of aryl azide **7**. The Sakai–Clark method has been reliably performed on large scale to produce >100 g of **1** in good efficiency and high purity.

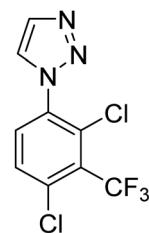
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Introduction

Notum is a carboxylesterase enzyme that suppresses Wnt signaling¹ through deacylation of an essential palmitoleate group on Wnt proteins.^{2,3} There is a growing understanding of the role Notum plays in human disease such as osteoporosis⁴ and colorectal cancer⁵ supporting the need to discover improved inhibitors.^{6–9} Recently, Willis *et al.* reported ARUK3001185 (**1**) as a potent, selective and brain penetrant inhibitor of Notum suitable for oral delivery in rodent models of disease.¹⁰ We are investigating the role of Notum in modulating Wnt signaling in Alzheimer's disease¹¹ and required multigram quantities of Notum inhibitor **1** for use in target validation and preclinical toxicology studies.


1: ARUK3001185

1-Aryl-1*H*-1,2,3-triazoles are an important structural motif in bioactive molecules and drugs¹¹ that has led to a number of synthetic methods for their construction.¹² These methods include the direct reaction of 1,2,3-triazole with a suitable aryl halide (ArBr, ArI) promoted by metal catalyzed cross-coupling. However, these tend to give mixtures of 1- and 2-aryl isomers through poor selectivity that limits its utility.^{13,14} Methods are preferred where the specific creation of the 1-aryl triazole can be controlled by the appropriate choice of reagents/synthons with defined and predictable reactivity. Copper(I) catalyzed azide–alkyne cycloaddition (CuAAC) promotes a [3 + 2] cyclisation where the three contiguous N atoms are provided by an aryl azide synthon (Ar–N₁–N₂–N₃). Aryl azides are readily available from anilines, *via* the diazonium salt, and so this approach has become popular, especially when combined with a suitably protected alkyne (*e.g.* TMS–C≡CH, C₄–C₅ synthon) that can be conveniently removed when necessary.

In 1986 Sakai *et al.* described a new method to prepare 1,2,3-triazoles from primary amines and α,α -dichloro tosylhydrazones **2a** under mild conditions as a metal-free alternative to azide–alkyne cycloaddition.¹⁵ The Sakai reaction was probably under-utilized until 2012 when Westermann *et al.* explored the scope and limitations of the reaction.¹⁶ The outcome and

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[†] Electronic supplementary information (ESI) available: Spectroscopic and analytical data for **1** (DSC, ¹H NMR, ¹³C and DEPT NMR, ¹⁹F NMR, HPLC, LCMS), **4** (¹H NMR, ¹⁹F NMR, HPLC) and **9** (¹H NMR, ¹⁹F NMR, HPLC, LCMS). Table of pilot reactions for the chlorination of 2-chloro-3-(trifluoromethyl) aniline (**3**) with *N*-chlorosuccinimide in various solvents. See <https://doi.org/10.1039/d2ra05132j>



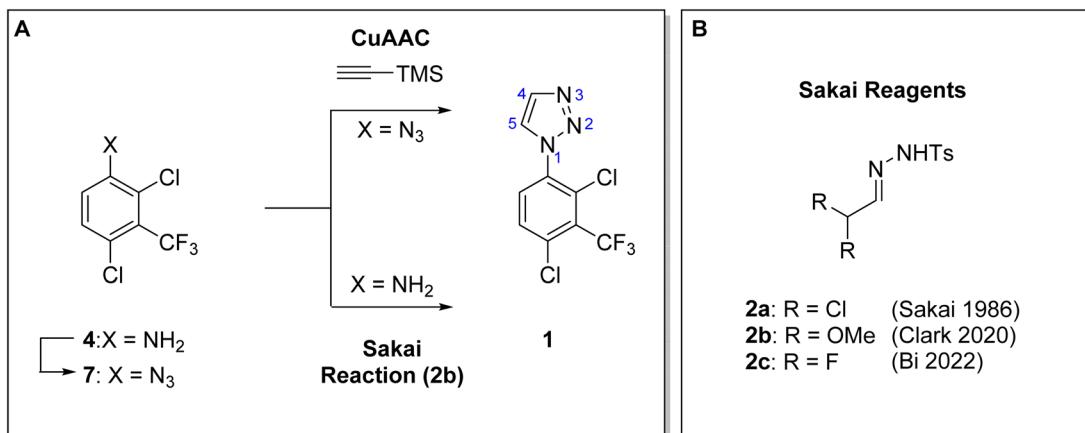


Fig. 1 (A). Synthesis of 1-aryl-1*H*-1,2,3-triazole **1** using both a CuAAC method and a modified Sakai reagent from aniline **4**. (B) α,α -disubstituted tosylhydrazones, Sakai reagents (**2**).

regioselectivity of the reaction is reliable in that the product triazole of the [1 + 4] cyclisation is created from the aniline (Ar-N₁) and tosylhydrazone (C₅-C₄-N₃-N₂).

Recently, Clark has reported a convenient and operationally simple modification of the Sakai reaction that produces 1-aryl triazoles with a range of anilines under mildly acidic conditions.^{17,18} The one-pot reaction of 2,2-dimethoxyacetaldehyde ((MeO)₂CHCHO) with *p*-toluene sulfonylhydrazide (TsNHNNH₂) in MeOH at RT for 2 h provides the α,α -dimethoxy tosylhydrazone **2b** *in situ*, and then addition of the aniline and AcOH with warming to 75 °C for 16 h produces the 1-aryl triazole in good yield.¹⁸ Further modifications of the Sakai reaction were presented by Bi using a base-promoted annulation of α,α -difluoro tosylhydrazone **2c** with a wide scope of amines under mild conditions.¹⁹ Proposed reaction mechanisms for these transformations have been reported.^{18,19}

Here, we disclose the application of both a CuAAC method and a modified Sakai reagent for the large-scale synthesis of aryl triazole **1** from aniline **4** (Fig. 1).

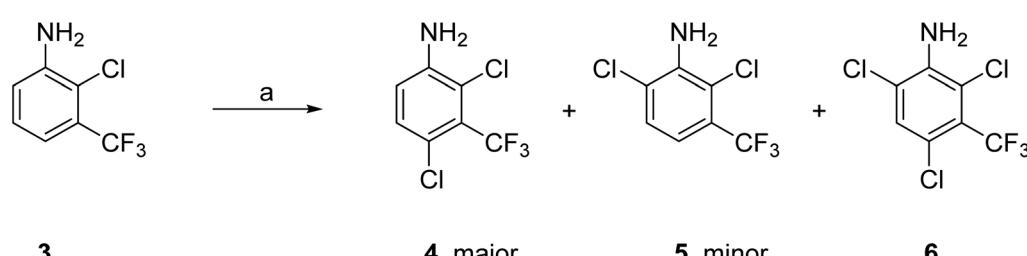
Results and discussion

These two complementary methods to prepare **1** both required 2,4-dichloro-3-(trifluoromethyl)aniline (**4**) as the starting material which provided an element of efficiency and economy to this approach. Aniline **4** was either purchased from

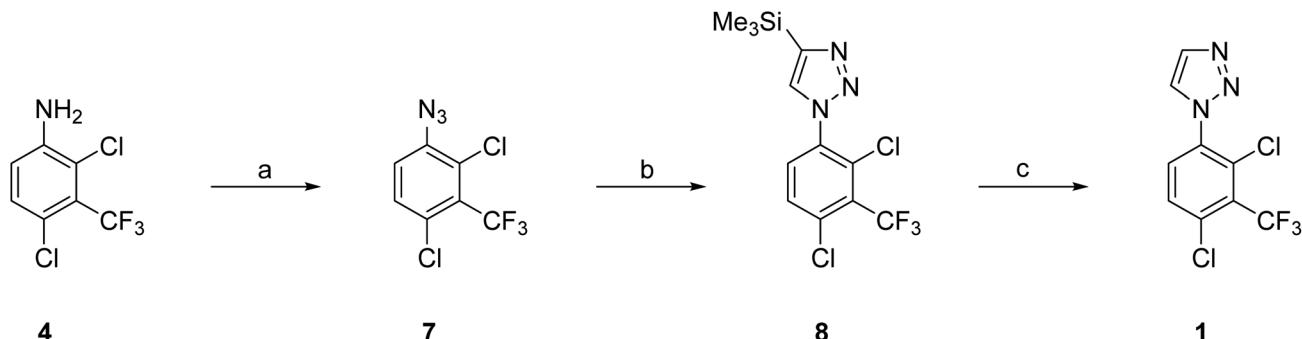
commercial suppliers or prepared by standard chlorination of 2-chloro-3-(trifluoromethyl)aniline (**3**) with *N*-chlorosuccinimide (NCS) as described below (Scheme 1).

Pilot reactions with **3** and NCS (1.1 equiv.) in a variety of solvents showed that the choice of solvent had a significant effect on both the conversion to **4** and the ratio of products (**4**–**6**) (Table S1†). NCS (1.1 equiv.) in MeCN gave the highest conversion to **4** with the lowest formation of **5** and **6**. Treatment of **3** (50 g scale) with NCS (1.1 equiv.) in MeCN at 60 °C for 16 h gave the required 4-chlorinated aniline **4** as the major product in 56% isolated yield. Analysis of the crude product mixture showed that the 6-chloro regio isomer **5** was also produced as a minor product along with small amounts of **6** through double chlorination at both the 4- and 6-positions (**3** : **4** : **5** : **6**, 1 : 72 : 19 : 6). The reaction was sequentially scaled up to 3 × 200 g of **3** without issue; the distribution of products was maintained as were the isolated yields of **4** (44%). Aniline **4** was purified by chromatography to give high purity material for use in the next step. As the need for quantities of **4** increased, it became economically more viable to prepare **4** from **3** (rather than purchase **4**), as **3** is significantly less expensive than **4** per unit cost.

Our efforts then focused on the development of a reliable synthesis of 50 g of 1-aryl-1*H*-1,2,3-triazole **1**. This multigram synthesis of **1** using the CuAAC method essentially followed the discovery route but with additional precautions to minimize the



Scheme 1 Synthesis of 2,4-dichloro-3-(trifluoromethyl)aniline (**4**). Reagents and conditions: (a) NCS (1.1 equiv.), MeCN, 60 °C, 16 h; ratio of products **4** : **5** : **6**, 72 : 19 : 6; **4** isolated yield 56%.



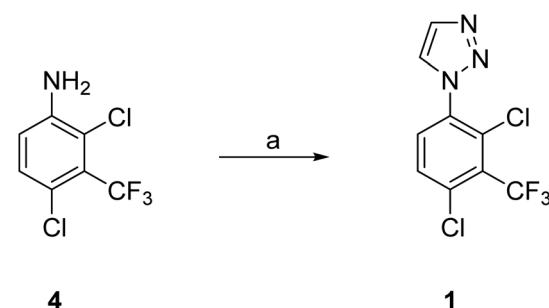
Scheme 2 First generation multigram synthesis of **1** using CuAAC method. Reagents and conditions: (a) (i) NaNO₂ (4.8 equiv.), CF₃CO₂H, 0 °C, 0.5 h → RT, 1.5 h; (ii) NaN₃ (4.8 equiv.), H₂O, 0 °C → RT, 16 h; (iii) NaOH, 0 °C; (b) HC≡CSiMe₃ (2.0 equiv.), sodium L-ascorbate (0.95 equiv.), CuSO₄·5H₂O (0.5 equiv.), MeOH-tBuOH-H₂O, RT, 16 h; (c) K₂CO₃ (2 equiv.), MeOH, RT, 3 d; 3-step overall total yield 65 g, 53%.

risk of azide **7** on larger scale (Scheme 2).¹⁰ In brief, reaction of aniline **4** with NaNO₂ in TFA gave the corresponding diazonium salt and then reaction with NaN₃ gave the required azide **7**. Copper catalyzed azide–alkyne cycloaddition of **7** with TMS-acetylene afforded a mixture of mostly 4-TMS triazole **8**, along with **1** (8 : 1, 5 : 2), which was treated with K₂CO₃ in MeOH to complete the proto-desilylation to give **1**.

Azides are potentially explosive reagents and should be treated with **caution** (see Experimental section for details). Differential scanning calorimetry (DSC) analysis of azide **7** established 42 °C as the maximum safe handling temperature for its preparation and use.¹⁰ In addition, azide **7** was maintained in solution throughout with solvent swapping as necessary. Reactions were performed batchwise (5 × 20 g of **4**) to limit the quantity of sodium azide used per reaction, and were carried through the 3-step sequence without purification until the end. Purification by chromatography gave **1** (13.5–15.8 g, 55–64%). These five batches were then pooled for a final purification to give a homogenous, crystalline product **1** (65 g, 53%) of very high purity.

Although successful in meeting its objective of producing >50 g of high purity **1**, the CuAAC method was inefficient through repetition of the 3-step procedure and would be scale limited without a more detailed understanding of the energetics of the azide **7** and the overall process. An improved procedure was required for scale-up to 100 g of **1** and larger quantities.

The Sakai reaction was an attractive alternative because of its simplicity through N₁-aryl specificity of the construction of the 1,2,3-triazole, readily available reagents, precedent for a toleration of wide variety of substituents on the aryl ring, and because it has been used to make representative bioactive molecules on gram scale.^{17–19} A pilot reaction with **4** (334 mg) using α,α -dimethoxy tosylhydrazone **2b** in MeOH at 75 °C for 16 h following the published conditions¹⁸ gave the required triazole **1** in 65% yield demonstrating the feasibility of this approach. Switching the solvent to THF improved the yield to 76%, whereas the use of DMF as solvent only gave recovered aniline. A few exploratory reactions of α,α -difluoro tosylhydrazone **2c** with **4** by the two reported methods (NaH/EtOAc or EtNiPr₂/MeOH) failed to produce **1** despite the reported success with similar



Scheme 3 Second generation multigram synthesis of **1** using Sakai–Clark method. Reagents and conditions: (a) (i) TsNNH₂ (2.0 equiv.), (MeO)₂CHCHO (3.3 equiv.), THF, RT, 2 h; (ii) **4** (1.0 equiv.), AcOH (3.0 equiv.), THF, 75 °C, 5 h, 30%.

anilines.¹⁹ Hence **2b** in THF was selected as the preferred reagent and solvent respectively for the larger scale procedures.

Three large scale reactions were then performed in parallel (**4**: 25 g and 2 × 140 g) and the crude products combined for purification. A mixture of TsNNH₂ and (MeO)₂CHCHO in THF was stirred at 20 °C for 2 h to create the α,α -dimethoxy tosylhydrazone **2b** *in situ* and then addition of aniline **4** and AcOH in THF with warming to 75 °C for 5 h produced crude 1-aryl triazole **1** (Scheme 3). The reaction mixture was cooled to RT, washed with brine and concentrated. The combined products from these three reactions was purified by column chromatography followed by preparative HPLC. The appropriate fractions were combined and partially concentrated, which prompted crystallisation. This approach proved to be very effective in producing **1** (113 g, 30%) of very high purity (99.8% pure by HPLC), although the overall yield was somewhat lower than the pilot reactions.²⁰ This chemistry, and these yields, have not been optimized and a more systematic approach to this process would be justified as the next step.

Conclusion

A one-pot synthesis of Notum inhibitor ARUK3001185 (**1**) has been developed enabling the delivery of multi-gram quantities of **1** for use in target validation and preclinical toxicology



studies. 1-Phenyl-1*H*-1,2,3-triazole **1** was prepared from aniline **4** by a modification of the Sakai reagent. The one-pot Sakai-Clark method with α,α -dimethoxy tosylhydrazone **2b** proved to be superior to the classical CuAAC method as it was operationally simple, metal-free, and avoided the use of aryl azide **7**. This method has been reliably performed on large scale to produce >100 g of **1** in good efficiency and high purity.

Experimental section

General information

Unless preparative details are provided, all reagents and solvents were purchased from commercial suppliers and used without further purification. Petroleum ether refers to fraction bp 60–90 °C. Thin-layer chromatography (TLC) was carried out on aluminum-backed Silica gel 60 F₂₅₄ plates. Organic solvent layers were routinely dried with anhydrous Na₂SO₄ or MgSO₄ and concentrated under reduced pressure using a Büchi rotary evaporator. Compound purification by column chromatography was performed using either a Biotage Isolera using prepacked Biotage Sfär silica cartridges, or silica gel packed columns. ¹H, ¹³C and ¹⁹F NMR spectra were recorded in deuterated ($\geq 99.5\%$) solvents on a Bruker Avance III 400 with ultrashield magnet and B-ACS-60 autosampler. Chemical shifts (δ) are reported as parts per million (ppm), coupling constants (J) are reported in Hz, and signal multiplicities are reported as singlet (s), doublet (d), doublet of doublets (dd), quartet (q), or multiplet (m).

Liquid chromatography-mass spectrometry (LCMS) analysis was performed on an Agilent 1260 infinity HPLC with Agilent 6130 single quadrupole MS in electrospray mode. Method 1: column, Phenomenex Kinetex XB-C₁₈, 50 × 4.6 mm, 2.6 μ m; column temperature, 40 °C; flow rate, 2 mL min⁻¹. Method 2: column, Luna C₁₈, 50 × 2.0 mm, 5 μ m; column temperature, 40 °C; flow rate, 1 mL min⁻¹. Solvent system and elution profiles, see Table 1.

Analytical high performance liquid chromatography (HPLC) was performed on either an Agilent 1260 (method 1) or Shimadzu LC-20AB (method 2): column, XBridge C₁₈, 50 × 2.1 mm, 5 μ m; column temperature, 40 °C; flow rate, 0.8–1.2 mL min; detection at λ 220 and 254 nm; solvent system and elution profile, see Table 2.

Table 1 LCMS solvent and elution profiles

Time (minutes)	% Aqueous (A) (0.1% formic acid in water)	% Organic (B) (100% acetonitrile)
Method 1		
0	95	5
1.37	2	98
1.83	95	5
2.25 end		
Method 2		
0	95	5
3.00	5	95
4.00	95	5
4.50 end		

Table 2 Analytical HPLC solvent and elution profiles

Time (minutes)	% Aqueous (A) (10 mM NH ₄ HCO ₃ in water)	% Organic (B) (100% acetonitrile)	Flow rate (mL min ⁻¹)
Method 1			
0.00	90	10	0.8
4.00	20	80	0.8
4.90	90	10	1.2
6.00 end			
Method 2			
0.00	90	10	0.8
4.50	20	80	0.8
5.40	90	10	1.2
6.00 end			

Table 3 Prep-HPLC solvent and elution profiles

Time (minutes)	% Aqueous (A) (0.09% TFA in water)	% Organic (B) (100% acetonitrile)
0	60	40
20	30	70
30	2	98
35 end		

Preparative high performance liquid chromatography (Prep-HPLC) was performed on Shimadzu LC-8A: column, Phenomenex Luna c18 250 × 100 mm, 10 μ m; column temperature, 40 °C; flow rate, 250 mL min; detection at λ 220 and 254 nm; solvent system and elution profile, see Table 3.

Chemical reactions were frequently performed in parallel batches and then combined for purification for expediency. Chemical yields refer to isolated and purified products unless otherwise stated. Purity of compounds was evaluated by NMR spectroscopy, HPLC and/or LCMS analysis. All compounds had purity $\geq 95\%$.

Full spectroscopic and analytical data for **1** have been reported.¹⁰ Additional spectroscopic and analytical data for **1** from this work are presented in the ESI† (Fig. S1–S6†).

2,4-Dichloro-3-(trifluoromethyl)aniline (4). 2,4-Dichloro-3-(trifluoromethyl)aniline (**4**) was either purchased from Activate Scientific (Ely, U. K.; cat. No. AS111481) or prepared by chlorination of 2-chloro-3-(trifluoromethyl)aniline (**3**) (Jiangsu Aikon, China; cat no. AK0033VU) as described below. Three reactions were performed in parallel (**3**: 3 × 200 g) and then combined for purification.

N-Chlorosuccinimide (150.0 g, 1.12 mol, 1.1 equiv.) was added to a stirred solution of 2-chloro-3-(trifluoromethyl)aniline (**3**) (200.0 g, 1.02 mol) in MeCN (2.4 L) at 20 °C, and the mixture was then heated at 60 °C for 16 h. The mixture was cooled to RT, concentrated, and the residue triturated with EtOAc (1.0 L).

The combined EtOAc filtrates (*ca* 3 L) from these three reactions was concentrated, and the crude product purified by column chromatography on silica gel (1–10% EtOAc in

petroleum ether) to afford **4** as brown oil (310 g, 1.35 mol, 44%) containing some residual EtOAc (*ca.* 0.15 equiv.) by ^1H NMR.

^1H NMR (400 MHz, CDCl_3) δ 7.19 (d, $J = 8.8$ Hz, 1H), 6.84 (d, $J = 8.9$ Hz, 1H), 4.28 (br s, 2H); ^{19}F NMR (376 MHz, CDCl_3) -55.71; HPLC (method 1): R_t 3.84 minutes, 99.6%.

1-(2,4-Dichloro-3-(trifluoromethyl)phenyl)-1*H*-1,2,3-triazole
(1) by CuAAC method.¹⁰ *Caution. Use of azides*. Azides are potentially explosive reagents and therefore all solutions containing azides were concentrated under reduced pressure behind a blast shield at RT. Furthermore, sodium azide reacts with acids to generate gaseous HN_3 , which is a toxic gas. All reactions were conducted behind a blast shield under a flow of nitrogen with exit gasses passed through two inline Drechsel bottles, the first empty and the second containing fresh NaOH, prior to release into ventilated fume hood.

Azide **7** was maintained in solution throughout its preparation and use, with solvent swapping as necessary. Solutions of **7** should not be evaporated to dryness. Differential scanning calorimetry (DSC) analysis of **7** established 42 °C as the maximum safe handling temperature.¹⁰

Reactions were performed batchwise (5×20 g of **4**) to limit the quantity of sodium azide used per reaction, and carried through the 3-step sequence without purification until the end. A representative experiment is presented.

Step 1. 2,4-Dichloro-3-(trifluoromethyl)aniline (**4**) (20.0 g, 87.0 mmol, 1.0 equiv.) was dissolved in 2,2,2-trifluoroacetic acid (400 mL) under nitrogen and the mixture was cooled to 0 °C with efficient stirring. Sodium nitrite (7.30 mol L^{-1} , 56.5 mL, 412 mmol, 4.8 equiv.) in water was added dropwise over 90 min, whilst maintaining the reaction mixture at *ca.* 0 °C throughout. The mixture was stirred at 0 °C for an additional 30 min., the ice bath was removed, and mixture allowed to warm slowly for 1 h (final temp. 5–10 °C). The solution of the diazonium salt was re-cooled to 0 °C, and sodium azide (3.85 mol L^{-1} , 107 mL, 412 mmol, 4.8 equiv.) in water was added dropwise over *ca.* 90 min., keeping the reaction temperature below 0 °C throughout. The reaction mixture was left in the ice bath and allowed to warm slowly to RT overnight. The solution of azide **7** in TFA-water was cooled to 0 °C and basified with the dropwise addition of aqueous sodium hydroxide (5.00 mol L^{-1} , 1045 mL, 5.23 mol) over *ca.* 5 h whilst maintaining the temperature at 0 °C. The mixture was warmed to RT, diluted with brine (500 mL) and extracted into Et_2O (3×700 mL). The combined organic extracts were dried (MgSO_4) and partially concentrated *in vacuo* (water bath set to 24 °C) to a volume of *ca.* 250 mL. *tert*-BuOH (200 mL) was then added, and the mixture evaporated until final traces of Et_2O have been removed to give 1-azido-2,4-dichloro-3-(trifluoromethyl)benzene (**7**) as solution in *tert*-BuOH (119 g total weight), which was used directly in the next step.

^1H NMR (400 MHz, CDCl_3) δ 7.45 (dd, $J = 8.8, 0.8$ Hz, 1H), 7.26 (dd, $J = 8.7, 0.7$ Hz, 1H); LCMS (method 1): R_t 1.97 minutes, 100%.

Step 2. Azide **7** as solution in *tert*-BuOH was dissolved in MeOH (200 mL) under nitrogen at RT. Copper sulfate pentahydrate (0.400 mol L^{-1} , 102 mL, 40.9 mmol, 0.47 equiv.) in water, sodium-L-ascorbate (1.01 mol L^{-1} , 81.8 mL, 82.6 mmol, 0.95 equiv.) in water and trimethylsilylacetylene (17.1 g,

174 mmol, 2.0 equiv.) were sequentially added, and the mixture stirred at RT overnight. TLC (20% EtOAc in hexane) showed no azide remaining. The mixture was extracted with CH_2Cl_2 (3 × 200 mL), dried (MgSO_4) and evaporated to give mostly 1-(2,4-dichloro-3-(trifluoromethyl)phenyl)-4-trimethylsilyl-1*H*-1,2,3-triazole (**8**) along with triazole **1** (33 g, **8** : **1**, 5 : 2), which was used directly in the next step to complete the desilylation.

^1H NMR (400 MHz, CDCl_3) δ 7.88 (s, 1H), 7.66–7.59 (m, 2H), 0.39 (s, 9H); LCMS (method 1): **1** R_t 1.70 minutes, 26% and **8** R_t 1.98 minutes, 68%.

Step 3. Crude 4-TMS triazole **8** was dissolved in MeOH (550 mL), potassium carbonate (24.0 g, 174 mmol, 2.0 equiv.) was added, and the mixture stirred at RT under nitrogen for 3 d. The reaction mixture was carefully concentrated under reduced pressure to remove the MeOH, and the residue suspended in CH_2Cl_2 (300 mL). This solution was filtered through a pad of silica gel and the silica pad was then washed with 20% MeOH in CH_2Cl_2 to ensure all product **1** was collected. The filtrate was evaporated, and the crude product (*ca.* 32 g) (range 28–35 g) purified by (1) column chromatography (0–100% CH_2Cl_2 in hexane), (2) column chromatography (0–20% EtOAc in hexane) and (3) triturated in hexane to give **1** as a white solid (14.7 g, 59%) (range 13.5–15.8 g, 55–64%).

Five batches of **1** were combined (73.4 g) and purified by column chromatography (15–20% EtOAc in hexane) to give **1** (65.3 g, 53%) as a crystalline, white solid of high purity.

Mp 107.2–108.6 °C; ^1H NMR (400 MHz, CDCl_3) δ 7.96 (d, $J = 1.1$ Hz, 1H), 7.90 (d, $J = 1.1$ Hz, 1H), 7.70–7.61 (m, 2H); ^{13}C NMR (101 MHz, CDCl_3) δ 136.37, 135.77, 133.93, 131.54, 131.01, 130.83, 127.99 (q, $J = 30.95$ Hz), 126.04, 122.04 (q, $J = 276.76$ Hz); LCMS (method 1): R_t 1.69 minutes, 100%, MS^+ *m/z* 282.0 (100%).

1-(2,4-Dichloro-3-(trifluoromethyl)phenyl)-1*H*-1,2,3-triazole

(1) by Sakai–Clark method

Small scale pilot reactions. THF (3.0 mL) was added to 4-methylbenzenesulfonohydrazide (250 mg, 1.34 mmol, 1.0 equiv.) in a thick-walled reaction vial with stir bar and the vial was sealed with a crimped septum cap. An aqueous solution of dimethoxyacetaldehyde (0.22 mL, 60% in H_2O , 1.48 mmol, 1.1 equiv.) was added *via* syringe and the mixture stirred at RT for 2 h. Separately, 2,4-dichloro-3-(trifluoromethyl)aniline (**4**) (334 mg, 1.48 mmol, 1.1 equiv.) and AcOH (80 μL , 1.34 mmol, 1.0 equiv.) was dissolved in THF (0.5 mL), and this solution was added to *via* syringe. The mixture was then heated at 75 °C for 18 h in a DrySyn block. TLC (30% EtOAc in cyclohexane) showed no aniline remaining. The reaction mixture was cooled to RT, concentrated and the residue partitioned between CH_2Cl_2 (30 mL) and water (20 mL) with vigorous mixing for 5 minutes. The 2-phase mixture was passed through a phase separator and the organic phase concentrated. The crude product was purified by chromatography on a 25 g Biotage SNAP KP Si cartridge (2–45% EtOAc in cyclohexane) to give **1** (286 mg, 1.01 mmol, 76%) as an off-white solid.

Three large scale reactions were then performed in parallel (4: 25 g and 2 × 140 g) and combined for purification.

25 g scale. A mixture of 4-methylbenzenesulfonohydrazide (40.5 g, 0.217 mol, 2.0 equiv.) and 2,2-dimethoxyacetaldehyde (62.2 g, 60 wt% in H_2O , 0.359 mol, 3.3 equiv.) in THF (500 mL)

was stirred at 20 °C for 2 h. A solution of **4** (25 g, 0.109 mol) and AcOH (19.6 g, 0.326 mol, 3.0 equiv.) in THF (250 mL) was added, and the mixture heated at 75 °C for 5 h. The mixture was cooled to RT, washed with brine (2 × 355 mL), dried (Na₂SO₄), and concentrated. The crude product was combined with additional batches for purification.

2 × 140 g scale. A mixture of 4-methylbenzenesulfonohydrazide (226.7 g, 1.217 mol, 2.0 equiv.) and 2,2-dimethoxyacetaldehyde (348.4 g, 60 wt% in H₂O, 2.01 mol, 3.3 equiv.) in THF (2.8 L) was stirred at 20 °C for 2 h. A solution of **4** (140 g, 0.609 mol) and AcOH (109.6 g, 1.825 mol, 3.0 equiv.) in THF (1.4 L) was added, and the mixture heated at 75 °C for 5 h. The mixture was cooled to RT, washed with brine (2 × 2 L), dried (Na₂SO₄), and concentrated.

The crude product from these three reactions was purified by column chromatography on silica gel (10–40% EtOAc in petroleum ether) and then by preparative HPLC (40–98% MeCN in water containing 0.09% TFA) (Table 3). The appropriate fractions were combined and partially concentrated to remove most of MeCN, which prompted crystallisation. The solid was collected, washed with water, and dried to afford **1** as white solid (113 g, 401 mmol, 30%).

Mp 108.2–109.2 °C; ¹H NMR (400 MHz, CDCl₃) 7.98 (d, *J* = 1.1 Hz, 1H), 7.91 (d, *J* = 1.1 Hz, 1H), 7.72–7.62 (m, 2H); ¹³C NMR (101 MHz, CDCl₃) 136.35, 135.74, 133.94, 131.55, 131.04, 130.82, 127.93 (q, *J* = 31.51 Hz), 126.09, 122.03 (q, *J* = 278.26 Hz); ¹⁹F NMR (376 MHz, CDCl₃) –55.79; HPLC (method 2): *R*_t 3.29 minutes, 99.8%. LCMS (method 2): *R*_t 2.50 minutes, 100%, MS⁺ *m/z* 282.0 (100%).

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Author contributions

The research was performed through contributions of all authors.

Conflicts of interest

B. N. A., N. J. W., and P. V. F. are co-inventors of patent application WO 2020043866, which describes inhibitors of Notum. The authors have no other relevant affiliations or financial involvement apart from those disclosed.

Abbreviations

CuAAC	copper(i)-catalyzed azide–alkyne cycloaddition
DMF	dimethylformamide

DSC	differential scanning calorimetry
HPLC	high performance liquid chromatography
LCMS	Liquid chromatography-mass spectrometry
NCS	N-chlorosuccinimide
RT	room temperature
TFA	trifluoroacetic acid
THF	tetrahydrofuran
TLC	thin-layer chromatography
TMS	trimethylsilyl

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