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# A Sulfuryl Group Transfer Strategy to Selectively Prepare Sulfated Steroids and Isotopically Labelled Derivatives

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The treatment of common steroids: estrone, estradiol, cortisol, and pregnenolone with tributylsulfoammonium betaine (TBSAB) provides a convenient chemoselective conversion of the steroids alcohol/phenol moiety to the corresponding steroidal organosulfate. An important feature of the disclosed methodology is the millimolar scale of the reaction, and the isolation of the corresponding steroid sulfates as their biologically relevant sodium salts without the need for ion-exchange chromatography. The scope of the method was further explored in the estradiol and pregnanediol steroid systems with the bis-sulfated derivatives. Ultimately, a method to install an isotopic label, deuterium ( $^2\text{H}$ ) combined with estrone sulfation is a valuable tool for its mass-spectrometric quantification in biological studies.

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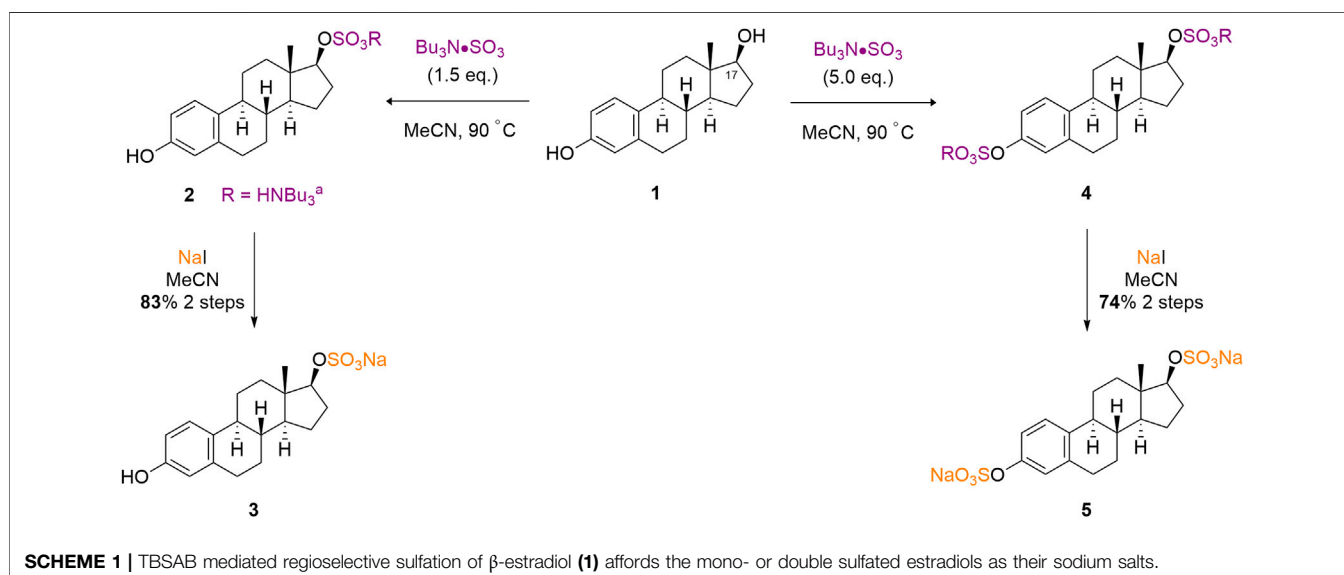
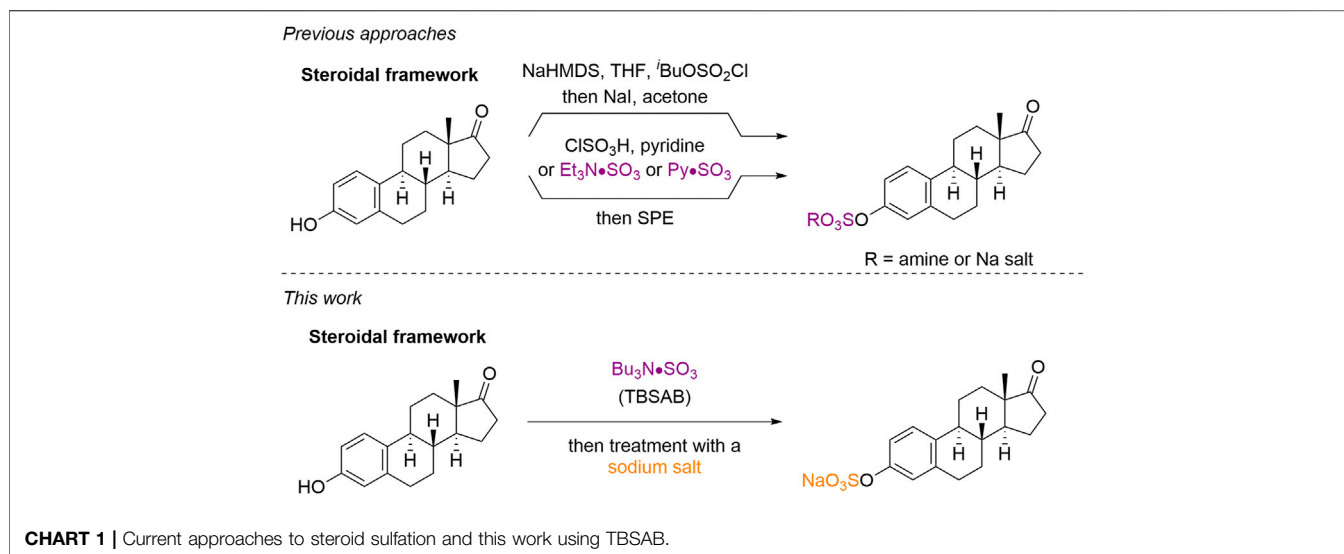
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## INTRODUCTION

The preparation of authentic reference samples of sulfated steroids with either regioselective mono or disulfation patterns, (Lightning et al., 2021) combined with methods to isotopically label the resulting sulfated steroids is an ongoing challenge to their biological study. The resulting authentic sulfated steroids are key reference standards of paramount importance to the understanding of sulfatases (Mueller et al., 2015), (Günel et al., 2019), (Foster and Mueller, 2018), the role of steroid sulfation in diseases (Mueller et al., 2021) and the fields of detection of steroids, whether in abuse (Waller and McLeod, 2014) or in the environment, (Petrie et al., 2013) using spectroscopic techniques (Hill et al., 2019). Furthermore, the development of improved sulfation methods can be applied to both sulfated steroid containing natural products synthesis and structural elucidation studies (Hoye et al., 2007).

Current methods to sulfate steroids fall into two main categories (**Chart 1**). The use of a protected sulfate group (e.g., isobutyl protected sulfate esters) with subsequent deprotection (Simpson and Widlanski, 2006), or the use of a sulfur trioxide equivalent (e.g., chlorosulfonic acid or pyridine-sulfur trioxide complex) (Waller and McLeod, 2014), (Hungerford et al., 2006). Although these methods are effective, they suffer from the additional steps of deprotection and/or purification cascades. Issues with toxicity regarding pyridine contamination from the use of pyridine-sulfur trioxide complex in related carbohydrate scaffolds (Gabriel et al., 2020), (Vo et al., 2021) requires either an exceptionally vigilant isolation and analysis; or an improved overall method for steroid sulfation.

Our own current interest in the sulfation field derives from the development of tributylsulfoammonium betaine (TBSAB) (Gill et al., 2019a), (Jones, 2021) as a convenient one-pot method for the sulfation of heteroatom containing bioactive molecules. (Benedetti et al., 2020), (Alshehri et al., 2020) This was initiated due to challenges encountered with the purification of sulfated small molecule heparin sulfate glycomimetics (Gill et al., 2021), (Gill et al., 2019b),



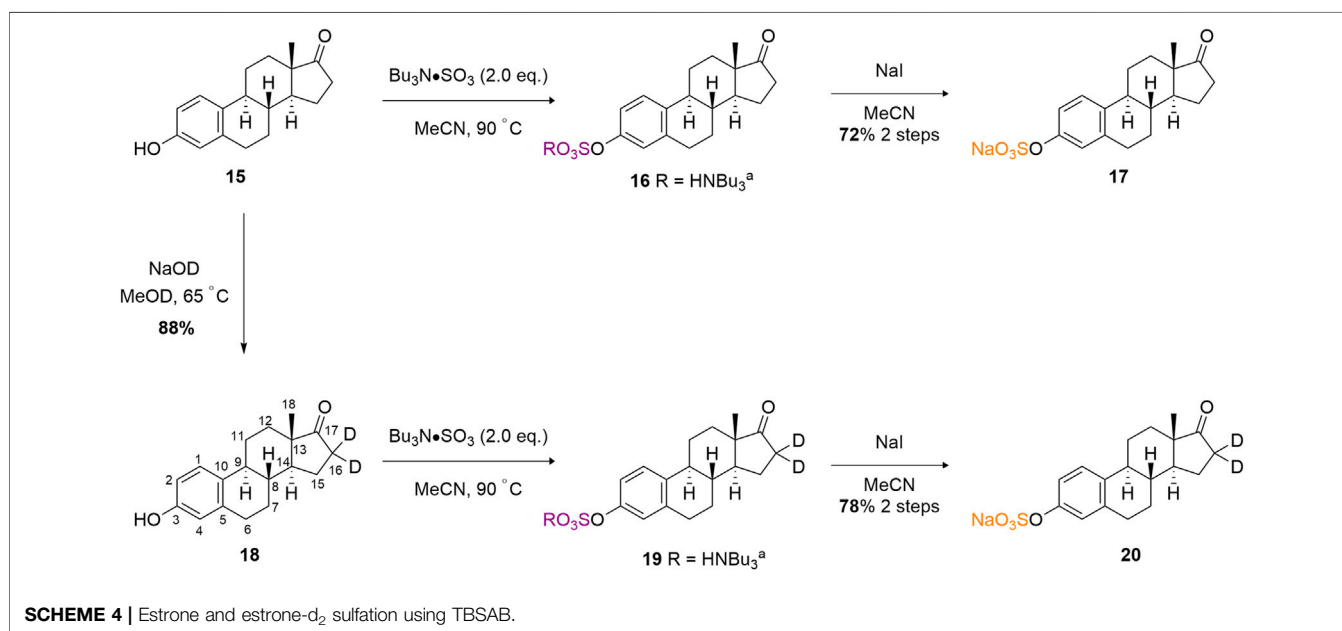
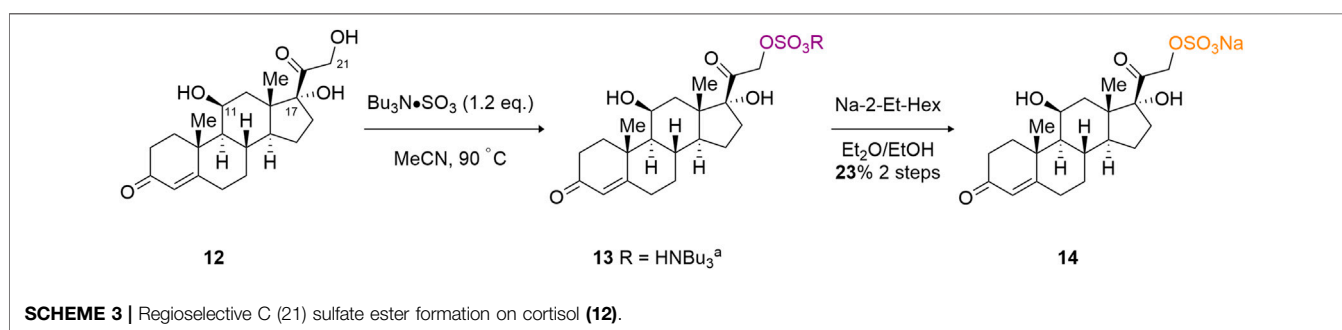
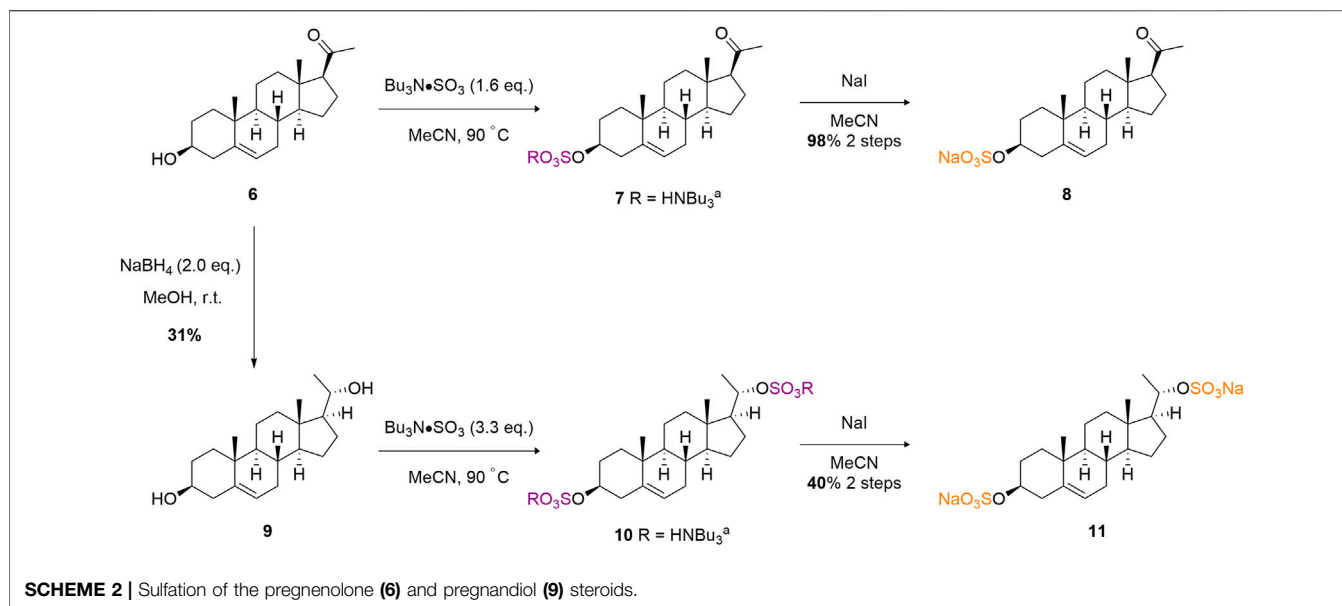
(Mahmoud et al., 2019), (Langford-Smith et al., 2019), (Mahmoud et al., 2017) with conventional, pre-existing sulfation methods. A key advantage of TBSAB over similar amine containing-sulfur trioxide complexes (e.g., triethylamine-sulfur trioxide) is the lipophilic nature of the counterion avoiding the need for ion-exchange chromatography. Herein we report our findings on the use of TBSAB as a general, scalable and regioselective sulfating reagent for steroids, and the application of TBSAB in conjugation with isotopic labelling for steroidal-organosulfate reference standards.

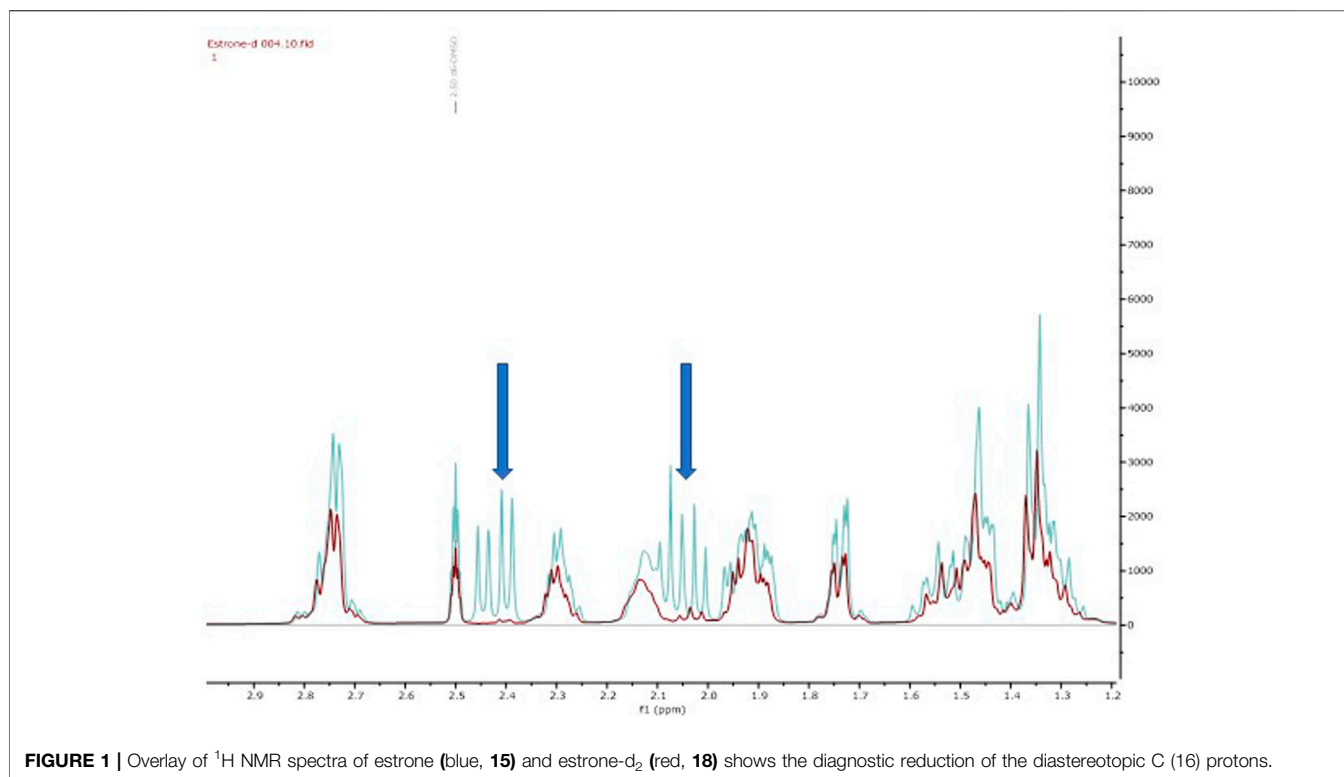
## RESULTS AND DISCUSSION

Our initial exploration of the method builds upon early screening results of TBSAB, including a single example on  $\beta$ -estradiol (**1**) (Gill

et al., 2019a). We firstly sought to demonstrate the reproducibility of this method on a 1.0 mmol scale, thus taking commercially available  $\beta$ -estradiol (**1**) and treating it with TBSAB resulted in exclusive C (17), secondary alcohol, sulfation (**2**). Furthermore the same conditions using an excess of TBSAB resulted in both C (17) sulfation and C (3), phenol, sulfation of (**4**) presumably occurs via initial C (17) alcohol sulfation in a stepwise installation. In both cases, a work-up using sodium iodide isolated the mono (**3**) and double (**5**) sulfated steroids as their sodium salts, in good yields without the risk of pyridinium ion contamination (**Scheme 1**).

Next, we considered sulfation of a more challenging biologically active substrate, pregnenolone (**6**). (Harteneck, 2013). Under analogous conditions to the  $\beta$ -estradiol examples, and on a 0.3 mmol scale, steroidal sulfate **8** was afforded after sodium exchange in an excellent 98% isolated yield (**Scheme 2**). Diastereoselective reduction of the ketone





moiety of pregnenolone using sodium borohydride afforded pregnanediol in 31% yield (**9**). Crystallographic data of the bulk material from  $\text{d}_6$ -DMSO crystallisation supports the assignment of the major diastereomer as *R* at the newly set stereocentre (**Supplementary Figure S3**) (2120263 contain, 2120). As **9** contains two secondary alcohol motifs, treatment with TBSAB afforded the double sulfated pregnanediol (**11**) in a modest 40% isolated yield on a 0.6 mmol scale.

The ultimate test of the TBSAB method, in relation to regioselective sulfation, is the complex triol, cortisol (**12**) (**Scheme 3**). Cortisol contains three potentially reactive hydroxyl motifs at the C (11), C (17) and C (21) positions. It was anticipated that a regioselective sulfation of the primary C (21) alcohol would result over the C (11), secondary, or C (17), tertiary, alcohol moieties, despite the presence of the  $\alpha$ -ketone affecting the reactivity of the C (21)-OH. To our delight, a microscale (8 mg) treatment of cortisol with TBSAB afforded the C (21) organosulfate in a modest 17% overall yield (23% based on recovered starting material) as the sodium salt (**14**). Furthermore, no unwanted C (11) or indeed C (17) sulfate ester formation was observed.

Finally, we sought to develop a proof-of-concept isotopic labelling-chemoselective sulfation method for the estrone scaffold (**15**) (**Scheme 4**). Prior to developing a deuterium labelling method at the C (16) methylene position, a model non-deuterated estrone was sulfated at the C (3) phenolic position in good 72% isolated yield as the sodium salt (**17**). A higher equivalence of TBSAB (2.0 eq) was used to ensure complete sulfation at the sole reactive C (3) phenolic centre. This provided confidence that sulfation should occur readily at

the C (3) position using TBSAB on the deuterium labeled substrate.

Firstly, we adapted the method of Rudqvist for C (19) deuteration (Rudqvist, 1983). Treatment of the estrone with NaOD in MeOD resulted in estrone- $\text{d}_2$  formation (**18**). The C (16)- $\text{H}_2$  protons were selectively deuterated by enolate formation with sodium deuteroxide and resultant deuterium incorporation by quenching the enolate with methanol- $\text{D}$  ( $\text{CH}_3\text{OD}$ ). This was confirmed via comparative 2D-NMR spectroscopic studies (see Supporting Information) but the key disappearance of the C (16) protons can be clearly observed in the  $^1\text{H}$ -NMR spectral overlay (**Figure 1**). It should be noted that deuteration next to a carbonyl group is not usually recommended for applied quantification studies as the deuterium label could readily back exchange through a keto-enol tautomerisation leading to a loss of the label (Wudy, 1990). In our system we observed, a decline of deuterium label in solution based mass-spectrometry studies (**Supplementary Figures S1, S2**).

Finally, the treatment of estrone- $\text{d}_2$  with TBSAB afforded the sulfated and isotopically labelled estrone- $\text{d}_2$  sulfate in 78% isolated yield and 67% incorporation of the deuterium label (**20**).

## CONCLUSION

In summary, we have demonstrated a general method for the synthesis of mono- or di-sulfated steroidal skeletons of importance to the fields of biology and spectroscopic detection. We have

showcased chemo-selective sulfation within a variety of complex structures, such as cortisol, and developed a simplified deuterium labeling-sulfation strategy for estrone. Overall, these approaches provide tractable routes on preparative scales to multiple sulfated steroid classes as reference compounds for detection of substances of abuse through to cancer diagnosis applications.

## DATA AVAILABILITY STATEMENT

The original contributions presented in the study are included in the article/**Supplementary Materials**, further inquiries can be directed to the corresponding author. Files can be found at the following doi: <https://doi.org/10.25500/edata.bham.00000720>.

## AUTHOR CONTRIBUTIONS

Conducted experiments, spectral analysis, and revised manuscript (JA and DG); provided materials and supervision,

## REFERENCES

- Alshehri, J. A., Benedetti, A. M., and Jones, A. M. (2020). A Novel Exchange Method to Access Sulfated Molecules. *Sci. Rep.* 10, 16559. doi:10.1038/s41598-020-72500-x
- Benedetti, A. M., Gill, D. M., Tsang, C. W., and Jones, A. M. (2020). Chemical Methods for N- and O-Sulfation of Small Molecules, Amino Acids and Peptides. *ChemBioChem* 21, 938–942. doi:10.1002/cbic.201900673
- CCDC 2120263 Contains the Supplementary Crystallographic Data for This Paper. These Data Can Be Obtained Free of Charge from the Cambridge Crystallographic Data Centre. Available at: [www.ccdc.cam.ac.uk/data\\_request/cif](http://www.ccdc.cam.ac.uk/data_request/cif).
- Foster, P. A., and Mueller, J. W. (2018). SULFATION PATHWAYS: Insights into Steroid Sulfation and Desulfation Pathways. *J. Mol. Endocrinol.* 61, T271–T283. doi:10.1530/JME-18-0086
- Gabriel, L., Günther, W., Pielenz, F., and Heinze, T. (2020). Determination of the Binding Situation of Pyridine in Xylan Sulfates by Means of Detailed NMR Studies. *Macromol. Chem. Phys.* 221, 1900327. doi:10.1002/macp.201900327
- Gill, D. M., Male, L., and Jones, A. M. (2019). A Structure-Reactivity Relationship of the Tandem Asymmetric Dihydroxylation on a Biologically Relevant Diene: Influence of Remote Stereocenters on Diastereofacial Selectivity. *Eur. J. Org. Chem.* 2019, 7568–7577. doi:10.1002/ejoc.201901474
- Gill, D. M., Male, L., and Jones, A. M. (2019). Sulfation Made Simple: a Strategy for Synthesising Sulfated Molecules. *Chem. Commun.* 55, 4319–4322. doi:10.1039/C9CC01057B
- Gill, D. M., R. Pavinelli, A. P., Zazeri, G., Shamir, S. A., Mahmoud, A. M., Wilkinson, F. L., et al. (2021). The Modulatory Role of Sulfated and Non-sulfated Small Molecule Heparan Sulfate-Glycomimetics in Endothelial Dysfunction: Absolute Structural Clarification, Molecular Docking and Simulated Dynamics, SAR Analyses and ADMET Studies. *RSC Med. Chem.* 12, 779–790. doi:10.1039/D0MD00366B
- Günal, S., Hardman, R., Kopriva, S., and Mueller, J. W. (2019). Sulfation Pathways from Red to green. *J. Biol. Chem.* 294, 12293–12312. doi:10.1074/jbc.REV119.007422
- Harteneck, C. (2013). Pregnenolone Sulfate: From Steroid Metabolite to TRP Channel Ligand. *Molecules* 18, 12012–12028. doi:10.3390/molecules181012012
- Hill, M., Hána, V., Velíková, M., Pařízek, A., Kolátorová, L., Vítků, J., et al. (2019). A Method for Determination of One Hundred Endogenous Steroids in Human Serum by Gas Chromatography-Tandem Mass Spectrometry. *Physiol. Res.* 68, 179–207. doi:10.33549/physiolres.934124
- Hoye, T. R., Dvornikov, V., Fine, J. M., Anderson, K. R., Jeffrey, C. S., Muddiman, D. C., et al. (2007). Details of the Structure Determination of the Sulfated

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## SUPPLEMENTARY MATERIAL

The Supplementary Material for this article can be found online at: <https://www.frontiersin.org/articles/10.3389/fmolb.2021.776900/full#supplementary-material>

- Steroids PSDS and PADS: New Components of the Sea Lamprey (*Petromyzon marinus*) Migratory Pheromone. *J. Org. Chem.* 72, 7544–7550. doi:10.1021/jo0709571
- Hungerford, N. L., McKinney, A. R., Stenhouse, A. M., and McLeod, M. D. (2006). Selective Manipulation of Steroid Hydroxyl Groups with Boronate Esters: Efficient Access to Antigenic C-3 Linked Steroid-Protein Conjugates and Steroid Sulfate Standards for Drug Detection. *Org. Biomol. Chem.* 4, 3951–3959. doi:10.1039/b610499a
- Jones, A. M. (2021). Tributylsulfoammonium Betaine. *Encyclopaedia Reagents Org. Synth.*, 1–3. doi:10.1002/047084289X.rn02393
- Langford-Smith, A. W. W., Hasan, A., Weston, R., Edwards, N., Jones, A. M., Boulton, A. J. M., et al. (2019). Diabetic Endothelial colony Forming Cells Have the Potential for Restoration with Glycomimetics. *Sci. Rep.* 9, 2309. doi:10.1038/s41598-019-38921-z
- Lightning, T. A., Gesteira, T. F., and Mueller, J. W. (2021). Steroid Disulfates - Sulfation Double Trouble. *Mol. Cell Endocrinol.* 524, 111161. doi:10.1016/j.mce.2021.111161
- Mahmoud, A. M., Jones, A. M., Sidgwick, G., Arafat, A. M., Wilkinson, F. L., and Alexander, M. Y. (2019). Small Molecule Glycomimetics Inhibit Vascular Calcification via C-Met/Notch3/HES1 Signalling. *Cell Physiol Biochem* 53, 323–336. doi:10.33594/000000141
- Mahmoud, A. M., Wilkinson, F. L., Jones, A. M., Wilkinson, J. A., Romero, M., Duarte, J., et al. (2017). A Novel Role for Small Molecule Glycomimetics in the protection against Lipid-Induced Endothelial Dysfunction: Involvement of Akt/eNOS and Nrf2/ARE Signaling. *Biochim. Biophys. Acta (Bba) - Gen. Subjects* 1861, 3311–3322. doi:10.1016/j.bbagen.2016.08.013
- Mueller, J. W., Gilligan, L. C., Idkowiak, J., Arlt, W., and Foster, P. A. (2015). The Regulation of Steroid Action by Sulfation and Desulfation. *Endocr. Rev.* 36, 526–563. doi:10.1210/er.2015-1036
- Mueller, J. W., Vogg, N., Lightning, T. A., Weigand, I., Ronchi, C. L., Foster, P. A., et al. (2021). Steroid Sulfation in Adrenal Tumors. *J. Clin. Endocrinol. Metab.* 106, Dgab182. doi:10.1210/clinem/dgab182
- Petrie, B., McAdam, E. J., Richards, K. H., Lester, J. N., and Cartmell, E. (2013). Application of Ultra-performance Liquid Chromatography-Tandem Mass Spectrometry for the Determination of Steroid Oestrogens in Wastewaters. *Int. J. Environ. Anal. Chem.* 93 (13), 1343–1355. doi:10.1080/03067319.2012.717272
- Rudqvist, U. (1983). Synthesis of C19 Steroid Monosulphates Labelled with Deuterium at Specific Positions. *J. Label Compd. Radiopharm.* 20, 1159–1170. doi:10.1002/jlcr.2580201005
- Simpson, L. S., and Widlanski, T. S. (2006). A Comprehensive Approach to the Synthesis of Sulfate Esters. *J. Am. Chem. Soc.* 128, 1605–1610. doi:10.1021/ja056086g
- Vo, Y., Schwartz, B. D., Onagi, H., Ward, J. S., Gardiner, M. G., Banwell, M. G., et al. (2021). A Rapid and Mild Sulfation Strategy Reveals Conformational

- Preferences in Therapeutically Relevant Sulfated Xylooligosaccharides. *Chem. Eur. J.* 27, 9830–9838. doi:10.1002/chem.202100527
- Waller, C. C., and McLeod, M. D. (2014). A Simple Method for the Small Scale Synthesis and Solid-phase Extraction Purification of Steroid Sulfates. *Steroids* 92, 74–80. doi:10.1016/j.steroids.2014.09.006
- Wudy, S. A. (1990). Synthetic Procedures for the Preparation of Deuterium-Labeled Analogs of Naturally Occurring Steroids. *Steroids* 55, 463–471. doi:10.1016/0039-128X(90)90015-4

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