

# Abu T Khan

## List of Publications by Year in descending order

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56  
papers

1,741  
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331642

21  
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289230

40  
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all docs

57  
docs citations

57  
times ranked

1286  
citing authors

| #  | ARTICLE  | IF  | CITATIONS |
|----|--|-----|-----------|
| 1  | Reactivity switch-over of 4-hydroxydithiocoumarins under various conditions and their application in organic synthesis. <i>Organic and Biomolecular Chemistry</i> , 2022, 20, 715-726.   | 2.8 | 8         |
| 2  | Synthetic utility of biomimicking vanadium bromoperoxidase and <i>n</i> -tetrabutylammonium tribromide (TBATB) in organic synthesis. <i>Organic and Biomolecular Chemistry</i> , 2022, 20, 2562-2579.  | 2.8 | 7         |
| 3  | DMSO-assisted environmentally benign synthesis of benzo[ <i>c</i> ]-chromeno[4,3,2- <i>gh</i> ]phenanthridines by remote oxidative hetero cross-coupling cyclization and aromatization reaction. <i>Chemical Communications</i> , 2022, , .  | 4.1 | 6         |
| 4  | $\hat{\text{I}}^{\pm}$ -Sulfonylation between 4-Hydroxydithiocoumarin and 1,3-Dicarbonyl Compounds: A Key Precursor for the Synthesis of New Pyrazole Derivatives. <i>Synthesis</i> , 2022, 54, 4521-4528.   | 2.3 | 1         |
| 5  | An environmentally benign regioselective synthesis of 2-benzyl-4-arylquinoline derivatives using aryl amines, styrene oxides and aryl acetylenes. <i>Organic and Biomolecular Chemistry</i> , 2021, 19, 8772-8782.   | 2.8 | 4         |
| 6  | Copper( <i>sc</i> ) triflate catalyzed three-component reaction for the synthesis of 2,3-diarylquinoline derivatives using aryl amines, aryl aldehydes and styrene oxides. <i>Organic and Biomolecular Chemistry</i> , 2021, 19, 3255-3262.  | 2.8 | 8         |
| 7  | Synthesis of biologically active fused 1,4-oxathiin derivatives from 4-hydroxydithiocoumarins, arylacetylenes and dimethyl sulfoxide by Cu( <i>sc</i> )-catalyzed C-H functionalization and cross-dehydrogenative C-S coupling reactions. <i>Organic and Biomolecular Chemistry</i> , 2021, 19, 5818-5826. | 2.8 | 7         |
| 8  | Ytterbium(III) triflate catalyzed domino reaction of arylamines and styrene oxides: Synthesis of 2-benzyl-3-arylquinoline derivatives. <i>Tetrahedron Letters</i> , 2021, 70, 152981.  | 1.4 | 8         |
| 9  | Metal-free synthesis of quinoline-2,4-dicarboxylate derivatives using aryl amines and acetylenedicarboxylates through a pseudo three-component reaction. <i>Organic and Biomolecular Chemistry</i> , 2021, 19, 7041-7050.  | 2.8 | 5         |
| 10 | Synthesis of vinyl sulfides and thioethers via a hydrothiolation reaction of 4-hydroxydithiocoumarins and arylacetylenes/styrenes. <i>Organic and Biomolecular Chemistry</i> , 2021, 19, 9223-9230.  | 2.8 | 2         |
| 11 | Anti-cancer potential of (1,2-dihydronaphtho[2,1- <i>b</i> ]furan-2-yl)methanone derivatives. <i>Bioorganic and Medicinal Chemistry Letters</i> , 2020, 30, 127476.  | 2.2 | 10        |
| 12 | Newly synthesized 3-sulfonylindole derivatives from 4-hydroxydithiocoumarin using an oxidative cross dehydrogenative coupling reaction (OCDCR): potential lead molecules for antiproliferative activity. <i>Organic and Biomolecular Chemistry</i> , 2020, 18, 4104-4113.                                  | 2.8 | 9         |
| 13 | Reaction behaviour of arylamines with nitroalkenes in the presence of bismuth(III) triflate: an easy access to 2,3-dialkylquinolines. <i>Organic and Biomolecular Chemistry</i> , 2020, 18, 1785-1793.   | 2.8 | 6         |
| 14 | Iodine monobromide catalysed regioselective synthesis of 3-arylquinolines from $\hat{\text{I}}^{\pm}$ -aminoacetophenones and trans- $\hat{\text{I}}^2$ -nitrostyrenes. <i>Organic and Biomolecular Chemistry</i> , 2019, 17, 347-353.   | 2.8 | 15        |
| 15 | Iodine-Catalyzed Synthesis of Pyrrolo(2,3- <i>c</i> )coumarin Derivatives Using 3-Aminocoumarins, Arylglyoxals and 4-Hydroxycoumarin through One-Pot Three-Component Reaction. <i>ChemistrySelect</i> , 2018, 3, 2431-2434.  | 1.5 | 10        |
| 16 | Electronic effect of substituents on anilines favors 1,4-addition to trans- $\hat{\text{I}}^2$ -nitrostyrenes: access to <i>N</i> -substituted 3-arylindoles and 3-arylindoles. <i>Organic and Biomolecular Chemistry</i> , 2018, 16, 3760-3770.   | 2.8 | 17        |
| 17 | An oxidative cross-coupling reaction of 4-hydroxydithiocoumarin and amines/thiols using a combination of $\text{I}_2$ and TBHP: access to lead molecules for biomedical applications. <i>Chemical Communications</i> , 2018, 54, 1513-1516.  | 4.1 | 23        |
| 18 | Triethylamine-Mediated One-Pot Synthesis of Benzo[ <i>f</i> ]chromene Derivatives. <i>ChemistrySelect</i> , 2018, 3, 4760-4763.  | 1.5 | 5         |

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|----|---|-----|-----------|
| 19 | K <sub>2</sub> CO <sub>3</sub> catalyzed regioselective synthesis of thieno[2,3-b]thiochromen-4-one oximes: access to the corresponding amine and nitroso derivatives. <i>Organic and Biomolecular Chemistry</i> , 2017, 15, 5625-5634.                               | 2.8 | 11        |
| 20 | Camphorsulfonic Acid Catalyzed One-Pot Three-Component Reaction for the Synthesis of Fused Quinoline and Benzoquinoline Derivatives. <i>Journal of Organic Chemistry</i> , 2017, 82, 12416-12429.   | 3.2 | 29        |
| 21 | One-pot three-component regioselective synthesis of C1-functionalised 3-arylbenzo[f]quinoline. <i>RSC Advances</i> , 2016, 6, 11675-11682.  | 3.6 | 16        |
| 22 | Synthesis of fused oxazole-containing coumarin derivatives via oxidative cross coupling reaction using a combination of CuCl <sub>2</sub> and TBHP. <i>RSC Advances</i> , 2016, 6, 18891-18894.   | 3.6 | 14        |
| 23 | Synthesis of Pyrido[2,3-c]coumarin Derivatives by an Intramolecular Povarov Reaction. <i>Synthesis</i> , 2015, 47, 1109-1116.   | 2.3 | 21        |
| 24 | Yb(OTf) <sub>3</sub> catalysed regioselective synthesis of unusual di- and tri- substituted 3,4-dihydrothiochromeno[3,2-e][1,3]thiazin-5(2H)-one derivatives through a pseudo four-component hetero-Diels-Alder reaction. <i>RSC Advances</i> , 2015, 5, 48104-48111. | 3.6 | 8         |
| 25 | Exploration of C5-C6-Unsubstituted 1,4-Dihydropyridines for the Construction of exo-Hexahydro-1H-chromeno[3,4-h][1,6]naphthyridine-3-carboxylates Using a Stereoselective Povarov Reaction. <i>Synthesis</i> , 2015, 47, 2745-2755.                                   | 2.3 | 9         |
| 26 | Synthesis of Unsymmetrical Sulfides and Their Oxidation to Sulfones to Discover Potent Antileishmanial Agents. <i>ACS Combinatorial Science</i> , 2015, 17, 671-681.  | 3.8 | 24        |
| 27 | Oxidative cross coupling reaction mediated by I <sub>2</sub> /H <sub>2</sub> O <sub>2</sub> : a novel approach for the construction of fused thiazole containing coumarin derivatives. <i>RSC Advances</i> , 2015, 5, 104155-104163.                                  | 3.6 | 11        |
| 28 | Synthesis of dihydrochromeno[4,3-b]pyrazolo[4,3-e]pyridin-6(7H)-ones involving one-pot three-component tandem Knoevenagel-Michael reaction catalyzed by n-tetrabutylammonium tribromide (TBATB). <i>Tetrahedron Letters</i> , 2014, 55, 2006-2009.                    | 1.4 | 19        |
| 29 | Synthesis of unsymmetrical sulfides catalyzed by n-tetrabutyl-ammonium tribromide: A selective fluorescence probe for mercury ion. <i>Sensors and Actuators B: Chemical</i> , 2014, 193, 509-514.   | 7.8 | 10        |
| 30 | Synthesis of fused tetrahydropyrido[2,3-c]coumarin derivatives as potential inhibitors for dopamine d <sub>3</sub> receptors, catalyzed by hydrated ferric sulfate. <i>RSC Advances</i> , 2014, 4, 3581-3590.   | 3.6 | 18        |
| 31 | Bromodimethylsulfonium Bromide (BDMS)-catalyzed Synthesis of Substituted Pyrroles through a One-pot Four-component Reaction. <i>Chemistry Letters</i> , 2013, 42, 939-941.  | 1.3 | 10        |
| 32 | A Useful and Convenient Synthetic Protocol for Iodination of Organic Substrates Using a Combination of Vanadyl Acetylacetonate, Hydrogen Peroxide, and Sodium Iodide. <i>Bulletin of the Chemical Society of Japan</i> , 2012, 85, 1239-1243.                         | 3.2 | 8         |
| 33 | A simple and expedient synthesis of functionalized pyrido[2,3-c] coumarin derivatives using molecular iodine catalyzed three-component reaction. <i>Tetrahedron Letters</i> , 2012, 53, 6418-6422.  | 1.4 | 50        |
| 34 | One-pot four-component domino reaction for the synthesis of substituted dihydro-2-oxypyrrrole catalyzed by molecular iodine. <i>Tetrahedron Letters</i> , 2012, 53, 2622-2626.  | 1.4 | 77        |
| 35 | Michael Initiated Ring Closure (MIRC) reaction on in situ generated benzylidenecyclohexane-1,3-diones for the construction of chromeno[3,4-b]quinoline derivatives. <i>Tetrahedron Letters</i> , 2012, 53, 2345-2351.   | 1.4 | 41        |
| 36 | Synthesis of tetra-substituted pyrroles, a potential phosphodiesterase 4B inhibitor, through nickel(II) chloride hexahydrate catalyzed one-pot four-component reaction. <i>Tetrahedron Letters</i> , 2012, 53, 4145-4150.   | 1.4 | 76        |

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|----|---|-----|-----------|
| 37 | VO(acac) <sub>2</sub> /H <sub>2</sub> O <sub>2</sub> /NaI: a mild and efficient combination for the cleavage of dithioacetal derivatives of sugars. <i>Carbohydrate Research</i> , 2011, 346, 2629-2632.  | 2.3 | 8         |
| 38 | Tetrabutylammonium tribromide (TBATB): a mild and efficient catalyst for O-isopropylideneation of carbohydrates. <i>Carbohydrate Research</i> , 2011, 346, 673-677.   | 2.3 | 24        |
| 39 | Sequential three-component reactions: synthesis, regioselectivity and application of functionalized dihydropyridines (DHPs) for the creation of fused naphthyridines. <i>Tetrahedron Letters</i> , 2011, 52, 3455-3459.                               | 1.4 | 38        |
| 40 | Iodine catalyzed one-pot five-component reactions for direct synthesis of densely functionalized piperidines. <i>Tetrahedron</i> , 2010, 66, 7762-7772.   | 1.9 | 136       |
| 41 | Synthesis of highly functionalized piperidines by one-pot multicomponent reaction using tetrabutylammonium tribromide (TBATB). <i>Tetrahedron Letters</i> , 2010, 51, 4419-4424.  | 1.4 | 137       |
| 42 | Bromodimethylsulfonium bromide (BDMS) mediated dithioacetalization of carbohydrates under solvent-free conditions. <i>Carbohydrate Research</i> , 2010, 345, 2139-2145.   | 2.3 | 17        |
| 43 | A Mild and Regioselective Method for $\alpha$ -Bromination of $\beta$ -Keto Esters and 1,3-Diketones Using Bromodimethylsulfonium Bromide (BDMS). <i>Journal of Organic Chemistry</i> , 2006, 71, 8961-8963.  | 3.2 | 76        |
| 44 | A mild and environmentally acceptable synthetic protocol for chemoselective $\alpha$ -bromination of $\beta$ -keto esters and 1,3-diketones. <i>Tetrahedron Letters</i> , 2006, 47, 2751-2754.  | 1.4 | 53        |
| 45 | A highly efficient and environmentally benign synthesis of 6,8-dibromoflavones, 8-bromoflavones, 5,7-dibromoaurones and 7-bromoaurones. <i>Tetrahedron Letters</i> , 2005, 46, 4937-4940.   | 1.4 | 28        |
| 46 | A Useful and Convenient Synthetic Procedure for Hydrolysis of Thioglycosides. <i>Chemistry Letters</i> , 2002, 31, 210-211.   | 1.3 | 14        |
| 47 | An exceptionally simple and catalytic method for regeneration of carbonyl functionality from the corresponding 1,3-oxathiolanes. <i>Journal of the Chemical Society, Perkin Transactions 1</i> , 2002, , 1026-1028.                                   | 1.3 | 23        |
| 48 | A useful and convenient synthetic protocol for interconversion of carbonyl compounds to the corresponding 1,3-oxathiolanes and vice versa employing organic ammonium tribromide (OATB). <i>Tetrahedron Letters</i> , 2002, 43, 2843-2846.             | 1.4 | 73        |
| 49 | A Convenient and Useful Method of Preparation of $\alpha$ -Bromo Enones from the Corresponding Enones Using Organic Ammonium Tribromide (OATB). <i>Chemistry Letters</i> , 2001, 30, 290-291.   | 1.3 | 22        |
| 50 | A Useful and Environmentally Benign Synthetic Protocol for Dethiolization by Employing Vanadium Pentoxide Catalyzed Oxidation of Ammonium Bromide by Hydrogen Peroxide. <i>Chemistry Letters</i> , 2001, 30, 1158-1159.                               | 1.3 | 30        |
| 51 | An environmentally benign synthesis of aurones and flavones from $\alpha$ -acetoxychalcones using n-tetrabutylammonium tribromide. <i>Tetrahedron Letters</i> , 2001, 42, 8907-8909.  | 1.4 | 73        |
| 52 | An Expedient and Efficient Method for the Cleavage of Dithioacetals to the Corresponding Carbonyl Compounds Using Organic Ammonium Tribromide (OATB). <i>Synlett</i> , 2001, 2001, 0785-0786.   | 1.8 | 38        |
| 53 | Regioselective Bromination of Organic Substrates by Tetrabutylammonium Bromide Promoted by V <sub>2</sub> O <sub>5</sub> ~H <sub>2</sub> O <sub>2</sub> : An Environmentally Favorable Synthetic Protocol. <i>Organic Letters</i> , 2000, 2, 247-249. | 4.6 | 195       |
| 54 | An environmentally benign synthesis of organic ammonium tribromides (OATB) and bromination of selected organic substrates by tetrabutylammonium tribromide (TBATB). <i>Tetrahedron Letters</i> , 1998, 39, 8163-8166.                                 | 1.4 | 137       |

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|----|---|-----|-----------|
| 55 | A method for the synthesis of C-(2-deoxy- $\beta$ -glycosyl) arenes. Carbohydrate Research, 1996, 280, 277-286.   | 2.3 | 15        |
| 56 | Regioselective ring-opening of epoxide and N-tosylaziridine with 4-hydroxydithiocoumarin: Key precursors of 2,3-dihydro-1,4-oxathiazine and 2,3-dihydro-1,4-thiazine derivatives. European Journal of Organic Chemistry, 0, , . | 2.4 | 1         |