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# Regioselective C(sp<sup>2</sup>)-H halogenation of pyrazolo[1,5-*a*]pyrimidines facilitated by hypervalent iodine(III) under aqueous and ambient conditions†

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An efficient and mild approach has been developed for the regio-selective direct C3 halogenation of pyrazolo[1,5-*a*]pyrimidines employing readily available potassium halide salts and a hypervalent iodine(III) reagent at ambient temperature. The protocol is both practical and environmentally friendly, utilizing water as a green solvent, potassium halides as an inexpensive and bench stable halogen source and PIDA as a non-toxic reagent, enabling clean and efficient halogenation at room temperature. The procedure yields a range of C3 halogenated pyrazolo[1,5-*a*]pyrimidines in good to excellent yields. Mechanistic studies suggest the involvement of electrophilic substitution mechanism in the halogenation process.

## Introduction

Organohalogen compounds are prevalent in nature and are recognized for their remarkable biological activities, rendering them intriguing candidates for drug development.<sup>1</sup> Numerous naturally existing metabolites containing halogens hold medicinal and therapeutic significance.<sup>2</sup> Additionally, these compounds find diverse applications in the realm of material science.<sup>3</sup> Halogenated heteroaromatic compounds hold significant importance as foundational components for synthesizing complex natural products and drugs. Their pivotal role in transition metal-catalysed cross-coupling reactions further underscores their importance in these processes.<sup>4</sup>

The pyrazolo[1,5-*a*]pyrimidine scaffold serves as a central structure for a diverse range of pharmacologically and biologically active compounds.<sup>5</sup> In the field of medicinal chemistry, this scaffold is renowned for its anti-tumor, antiviral, anti-cancer, anti-malarial, and anti-inflammatory properties.<sup>6</sup> Notably, the pyrazolo[1,5-*a*]pyrimidine core is a crucial element present in various anti-cancer medications like selitrectinib, repotrectinib, larotrectinib and is also a constituent of insomnia medications such as zaleplon and lorediplon<sup>7</sup> (Fig. 1). Beyond its therapeutic relevance, the pyrazolo[1,5-*a*]pyrimidine core bears significant importance for material scientists, particularly in the realms of intriguing optical applications and chemosensor functionality.<sup>8</sup> Halogenated pyrazolo[1,5-*a*]pyrimidines have been found to possess anxiolytic properties, their synthesis has been explored with the aim of assessing their

potential effectiveness in addressing anxiety disorders.<sup>9</sup> In 2004, Liebscher *et al.* introduced a protocol for the direct halogenation of pyrazolo[1,5-*a*]pyrimidines using corresponding *N*-halosuccinimides (Scheme 1a).<sup>10</sup> However, this method requires elevated temperatures and the use of organic solvents. Similar approaches using *N*-halosuccinimides were later employed by Martins *et al.*, and Portilla *et al.*<sup>11</sup> (Scheme 1a). Hence, the development of an environmentally friendly and efficient methodology for the direct halogenation of pyrazolo[1,5-*a*]pyrimidines is of crucial significance. In this work, we disclose a method for the direct and regioselective C3 halogenation of pyrazolo[1,5-*a*]pyrimidines at room temperature. This approach employs potassium halides as a safe, bench stable, cheap and readily available halogenation source (Scheme 1b). The key

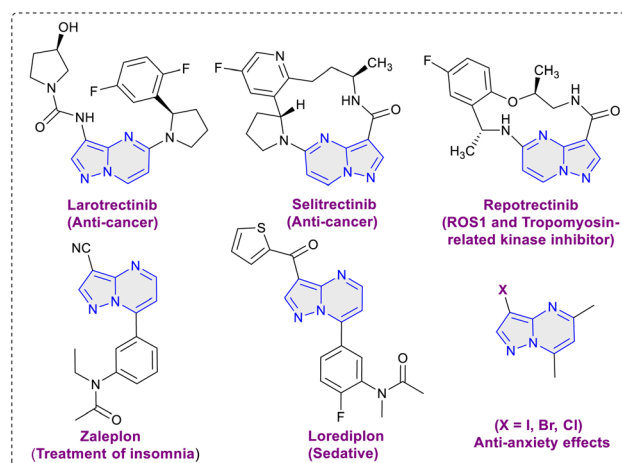
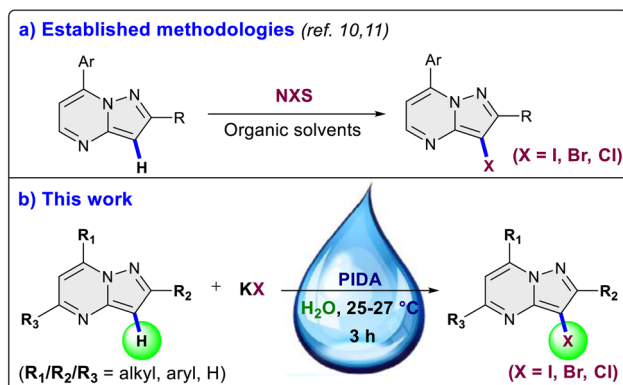


Fig. 1 Bioactive compounds with pyrazolo[1,5-*a*]pyrimidine scaffold.

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Scheme 1 Previous work on halogenation of pyrazolo[1,5-*a*]pyrimidines.

advantage of this methodology is its operational feasibility at room temperature, demonstrating its efficiency and practicality. Additionally, it utilizes water as solvent, contributing to the overall sustainability of the process.

## Results and discussion

The investigation into the iodination reaction commenced with the selection of 2-methyl-7-phenylpyrazolo[1,5-*a*]pyrimidine (**1a**) as the preferred substrate and potassium iodide (**2a**) as the halogen source. 2-Methyl-7-phenylpyrazolo[1,5-*a*]pyrimidine (**1a**) was reacted with 1.5 equivalents of KI and K<sub>2</sub>S<sub>2</sub>O<sub>8</sub> (1.0 equiv.) for 3 hours at room temperature (25–27 °C) in water as a solvent, yielding desired 3-iodo-2-methyl-7-phenylpyrazolo[1,5-*a*]pyrimidine (**3aa**) in a 30% yield (Table 1, entry 1). Encouraged by the initial success, alternative oxidants for the iodination reaction were explored keeping other parameters constant. In the presence of Na<sub>2</sub>S<sub>2</sub>O<sub>8</sub> and oxone the yield increased to 35% and 38% respectively (entry 2 and 3). However, the use of *tert*-butyl peroxybenzoate (TBAB) and *tert*-butyl hydroperoxide (TBHP) resulted in lower yields of 15% and 25%, respectively (entries 4 and 5). Oxygen, when used as an oxidant, did not yield any product in the iodination reaction (entry 6). Subsequently, we explored hypervalent iodine(III) reagents as oxidants, and were pleased to observe a substantial increase in the yield of the desired product **3aa**. In the presence of (bis(trifluoroacetoxy)iodo)benzene (PIFA) and (diacetoxyiodo)benzene (PIDA), the yields escalated to 72% and 87%, respectively (entries 7 and 8). Upon conducting oxidant screening, it became apparent that PIDA exhibited superior performance as the oxidant for the iodination reaction. The transition of the halogen source from KI to NaI and tetrabutylammonium iodide (TBAI) resulted in a diminished yield of the desired product **3aa** to 64% and 73% respectively (entries 9 and 10), indicating that KI proved to be a more effective halogen source for the iodination reaction. Utilizing 1.0 equivalent of KI in the reaction, as opposed to 1.5 equivalents, resulted in a decrease in the reaction yield to 70% (entry 11).

Upon reducing the amount of PIDA to 0.75 and 0.5 equivalents, lowering the yields to 65% and 52%, respectively (entries

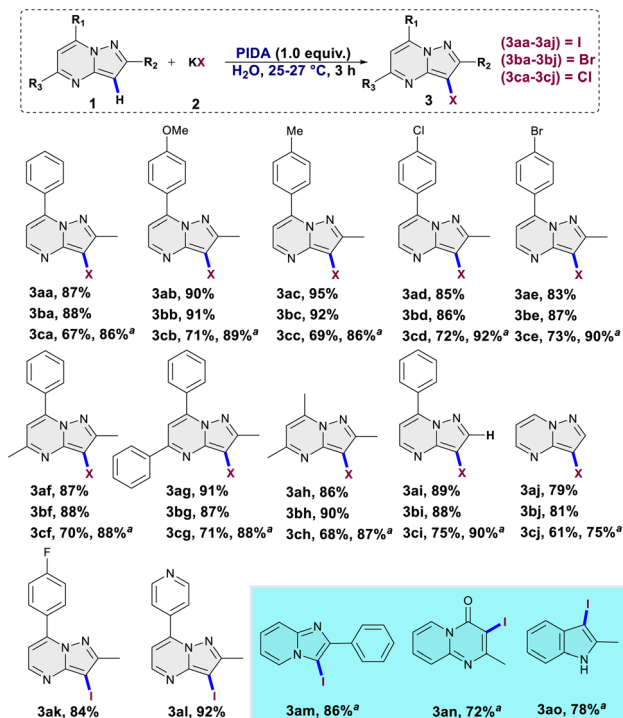
Table 1 Optimization of reaction conditions<sup>a</sup>

| Entry           | Oxidant                                       | Solvent               | Yield <sup>b</sup> |
|-----------------|---|-----------------------|--------------------|
| 1               | K <sub>2</sub> S <sub>2</sub> O <sub>8</sub>  | H <sub>2</sub> O      | 30%                |
| 2               | Na <sub>2</sub> S <sub>2</sub> O <sub>8</sub> | H <sub>2</sub> O      | 35%                |
| 3               | Oxone   | H <sub>2</sub> O      | 38%                |
| 4               | TBPB  | H <sub>2</sub> O      | 15%                |
| 5               | TBHP  | H <sub>2</sub> O      | 25%                |
| 6               | O <sub>2</sub>                                | H <sub>2</sub> O      | NR                 |
| 7               | PIFA  | H <sub>2</sub> O      | 72%                |
| 8               | <b>PIDA</b>                                   | <b>H<sub>2</sub>O</b> | <b>87%</b>         |
| 9 <sup>c</sup>  | PIDA  | H <sub>2</sub> O      | 64%                |
| 10 <sup>d</sup> | PIDA  | H <sub>2</sub> O      | 73%                |
| 11 <sup>e</sup> | PIDA  | H <sub>2</sub> O      | 70%                |
| 12 <sup>f</sup> | PIDA  | H <sub>2</sub> O      | 65%                |
| 13 <sup>g</sup> | PIDA  | H <sub>2</sub> O      | 52%                |
| 14 <sup>h</sup> | —   | H <sub>2</sub> O      | NR                 |

<sup>a</sup> Reaction conditions: **1a** (0.2 mmol), **2a** (0.3 mmol), oxidant (1.0 equiv.), H<sub>2</sub>O (3.0 mL), rt (25–27 °C), 3 h. <sup>b</sup> Isolated yields. <sup>c</sup> TBAI was used instead of KI. <sup>d</sup> NaI was used instead of KI. <sup>e</sup> 1.0 equiv. of KI was used. <sup>f</sup> 0.75 equiv. oxidant was used. <sup>g</sup> 0.5 equiv. oxidant was used. <sup>h</sup> No oxidant.

12 and 13) was observed. When the oxidant was omitted, the reaction did not take place, emphasizing the crucial role of the oxidant in the reaction (entry 14). The optimal yield of **3aa** was achieved by employing 1.0 equivalent of **1a**, 1.5 equivalents of KI, 1.0 equivalent of PIDA, utilizing water as the solvent, and conducting the reaction for duration of 3 hours. After obtaining optimized reaction conditions, we decided to investigate the substrate scope for halogenation (I/Br/Cl) of pyrazolo[1,5-*a*]pyrimidines (Scheme 2). Under optimal conditions, iodination of 2-methyl-7-phenylpyrazolo[1,5-*a*]pyrimidine resulted in product **3aa** with an 87% yield. Various derivatives of 2-methyl-7-phenylpyrazolo[1,5-*a*]pyrimidine, bearing electron-donating (–OMe, –Me) and electron-withdrawing (–Cl, –Br) groups on the phenyl ring, smoothly underwent iodination reactions, producing the respective C3 iodinated products (**3ab–3ae**) with yields ranging from 83% to 95%. A variety of tri-substituted pyrazolo[1,5-*a*]pyrimidine derivatives efficiently underwent an iodination reaction under optimised conditions, leading to the formation of products **3af–3ah** with excellent yields ranging from 86% to 91%. 7-Phenylpyrazolo[1,5-*a*]pyrimidine and pyrazolo[1,5-*a*]pyrimidine exhibited excellent reactivity, yielding the corresponding C3 iodinated products **3ai** and **3aj** with impressive yields of 89% and 79%, respectively. This underscores the excellent regioselectivity of the present methodology. Under optimized conditions, both the pyrazolo[1,5-*a*]pyrimidine derivative bearing a fluorine substituent on the phenyl ring and the derivative with a pyridine ring attached demonstrated successful reactions, yielding iodination products **3ak** and **3al** in 84% and 92% yield respectively. Other N-heterocycles like 2-



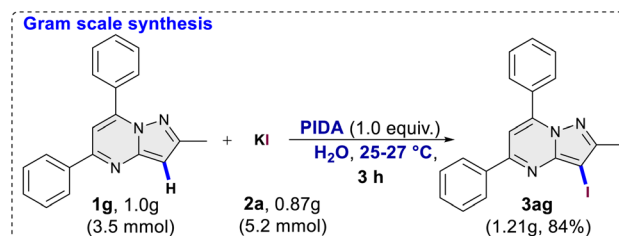


**Scheme 2** Substrate scope for the regioselective halogenation of pyrazolo[1,5-*a*]pyrimidines and other N-heterocycles. Reaction conditions: **1** (0.2 mmol), **KX** (0.3 mmol), PIDA (1.0 equiv.), H<sub>2</sub>O (3.0 mL), rt (25–27 °C), 3 h. Yields are isolated yields. <sup>a</sup>Reactions were carried out in MeOH.

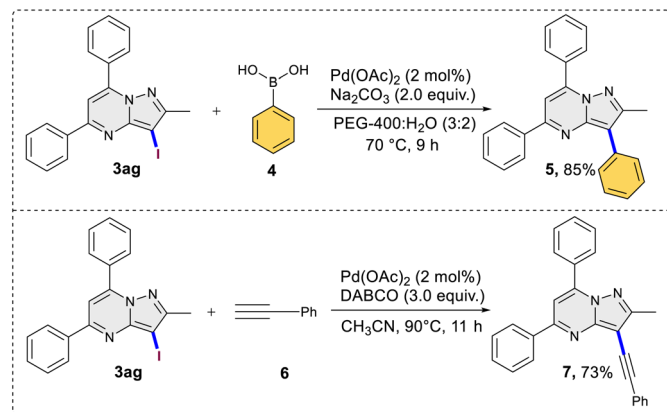
phenylimidazo[1,2-*a*]pyridine, 2-methyl-4*H*-pyrido[1,2-*a*]pyrimidin-4-one and 2-methyl-1*H*-indole reacted well giving corresponding iodinated products in 86%, 72% and 78% yield respectively. Subsequently, we chose to expand the present approach to conduct the regioselective bromination of pyrazolo[1,5-*a*]pyrimidine derivatives. To begin with, 2-methyl-7-phenylpyrazolo[1,5-*a*]pyrimidine was reacted with 1.5 equivalent of potassium bromide in presence of PIDA (1.0 equiv.) using H<sub>2</sub>O as a solvent for 3 hours. Gratifyingly, the anticipated C3 brominated product **3ba** was obtained with a high yield of 88%. The current reaction conditions demonstrated excellent compatibility with both electron-withdrawing (–Cl, –Br) and electron-donating groups (–OMe, –Me) yielding the respective C3 brominated products (**3bb–3be**) in outstanding yields ranging from 86% to 92%. The reaction exhibited good reactivity with different trisubstituted pyrazolo[1,5-*a*]pyrimidine derivatives, yielding the corresponding brominated products **3bf–3bh** in remarkable yields ranging from 88% to 90%. 7-Phenylpyrazolo[1,5-*a*]pyrimidine and pyrazolo[1,5-*a*]pyrimidine displayed excellent reactivity and regioselectivity under optimized conditions, yielding C3 brominated products **3bi** (88%) and **3bj** (81%). Notably, exclusive bromination was observed solely at the C3 position, with no occurrence of bromination at any other site. Following the successful iodination and bromination, the same strategy was implemented for the site-selective chlorination of the pyrazolo[1,5-*a*]pyrimidine scaffold. A reaction was conducted by treating 2-methyl-7-phenylpyrazolo[1,5-

*a*]pyrimidine with potassium chloride (1.5 equiv.) utilizing water as a solvent over a period of 3 hours; the chlorinated product **3ca** was isolated with a slight lower yield of 67% compared to iodination and bromination. Interestingly, when the same reaction was conducted in MeOH with all other parameters held constant, the yield of the product **3ca** significantly increased to 86%. Consequently, based on these findings, we opted to conduct chlorination reactions in water as well as in methanol as the solvent. 2-Methyl-7-phenylpyrazolo[1,5-*a*]pyrimidines bearing electron donating substituents (–OMe, –Me) as well as electron withdrawing halogen substituents (–Br, –Cl) gave corresponding C3 chlorinated products **3cb–3ce** in good yields ranging from 69–73% respectively. Trisubstituted derivatives of pyrazolo[1,5-*a*]pyrimidine exhibited excellent reactivity in the chlorination protocol, yielding the corresponding mono-chlorinated products **3cf–3ch** with commendable yields (68–70%). In line with the regioselectivity observed in iodination and bromination reactions, chlorination exhibited a similar trend.

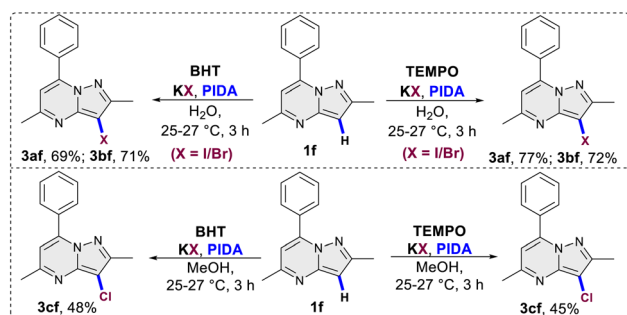
This was evident in the selective formation of products **3ci** and **3cj**, providing yields of 75% and 61%, respectively. To ascertain the reliability and efficacy of the devised methodology, we decided to carry out iodination of the substrate on gram scale. A 3.5 mmol (1.0 g) quantity of 2-methyl-5,7-diphenylpyrazolo[1,5-*a*]pyrimidine (**1g**) was subjected to a reaction with 5.2 mmol (0.87 g) of KI, in water as a solvent. The reaction proceeded for 3 hours at room temperature, yielding the desired C3 iodinated product **3ag** with a 84% yield (1.21 g) (Scheme 3). Emphasizing the utility of the synthesized halogenated compounds, their potential to undergo transformation into substituted pyrazolo[1,5-*a*]pyrimidine derivatives was investigated. Specifically, a targeted approach involved subjecting 3-iodo-2-methyl-5,7-diphenylpyrazolo[1,5-*a*]pyrimidine to well-established cross-coupling reactions, namely Suzuki–Miyaura coupling and Sonogashira coupling (Scheme 4).<sup>12</sup> Under the conditions outlined in Scheme 4, compound **3ag** underwent reactions with phenyl boronic acid and phenyl acetylene, yielding products **5** and **7** with good yields of 85% and 73%, respectively. Following this, a series of controlled experiments were meticulously carried out to clarify the mechanistic foundations governing the halogenation reaction of pyrazolo[1,5-*a*]pyrimidine. These reactions were systematically conducted under optimized conditions, adding radical scavengers, namely (2,2,6,6-tetramethylpiperidin-1-yl)oxyl (TEMPO) and 2,6-di-*tert*-butyl-4-methylphenol (BHT), to assess the potential



**Scheme 3** Gram scale synthesis.



Scheme 4 Synthetic application in cross coupling reactions.



Scheme 5 Control experiments.

participation of radical intermediates (Scheme 5). Despite the addition of TEMPO and BHT in the halogenation reactions, these radical scavengers proved ineffective in completely suppressing product formation. Only a modest reduction in product yield in case of iodination and bromination was observed, with the iodinated product **3af** being obtained in 77% and 69% yields in the presence of TEMPO and BHT, respectively. Similarly, the bromination product **3bf** was isolated in 72% and 71% yields in the presence of TEMPO and BHT, demonstrating a marginal decrease in overall efficiency. In case of chlorination reaction product **3cf** was obtained in 45% and

48% yields in presence of TEMPO and BHT respectively. Based on the results of control experiments and literature study<sup>13</sup> a possible mechanism has been postulated for the halogenation of pyrazolo[1,5-*a*]pyrimidines (Scheme 6). The process commences with PIDA initiating a ligand exchange with a halide salt, resulting in the formation of intermediate **A**. This intermediate then undergoes a transformation into hypohalite salt **B**, serving as a source for an electrophilic halogen species. Following this, substrate **1a** engages with the electrophilic halogen species **B**, leading to the formation of intermediate **C**. In the final step, intermediate **C** undergoes deprotonation, ultimately yielding the halogenated product **3**.

## Conclusions

In conclusion, we have developed an environmentally friendly and efficient method for the regio-selective C3 halogenation of pyrazolo[1,5-*a*]pyrimidines, using a hypervalent iodine(III) reagent, readily and cheaply available potassium halide salts, and water as a green solvent at ambient temperature. The method facilitates clean and effective halogenation, resulting in the synthesis of diverse C3 halogenated pyrazolo [1,5-*a*]pyrimidine derivatives with consistently good to excellent yields. Our preliminary mechanistic studies suggest the involvement of an electrophilic substitution mechanism in this process. Beyond its environmental merits, this method demonstrates characteristics of rapidity, operational simplicity, scalability, and an expansive substrate scope, thereby enhancing its potential as a versatile and valuable tool in organic synthesis.

## Conflicts of interest

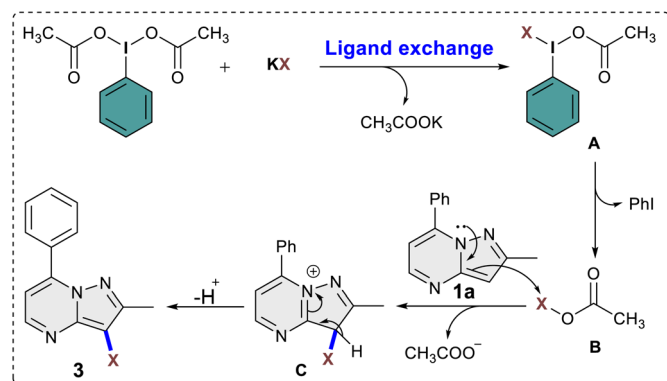
There are no conflicts to declare.

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Scheme 6 Plausible reaction mechanism.





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