EDGE ARTICLE

A ruthenium(II)-catalyzed C–H allenylation-based approach to allenoic acids†

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A Ru(II)-catalyzed direct access to various functionalized allenoic acids *via* C–H allenylation of readily available aryl carboxylic acids with propargylic acetates is reported. Axially chiral allenoic acids could be obtained in high ee by using optically active propargylic acetates through a chirality transfer strategy.

Allene moieties are not only present in natural products but also in precious building blocks due to their unique structures and multiple reactive sites.[1] Allene chemistry has experienced an explosion during the last few decades.[2] Thus, the synthesis of functionalized allenes is of crucial importance. Of particular interest is the synthesis of synthetically versatile allenoic acids.[3] A common approach to 2,3-allenoic acids is the hydrolysis of 2,3-allenoates, which suffers from poor step-economy and a selectivity issue of forming allenoic acids and 3-alkynoic acids (Scheme 1a).[4] The only example of the oxidation of allenols is realized through microbial transformation (Scheme 1b).[5] Pd- or Ni-catalyzed carboxylation of propargylic compounds with CO in the presence of water (Scheme 1c)[6] and the carboxylation of 2-alkynyl bromides or allenylmetallic reagents with CO_2 (Scheme 1d)[7] have also been reported. Crabbé homologation of *o*-methoxycarbonylphenylacetylene with paraformaldehyde results in the formation of methyl 2-propadienylbenzoate, which undergoes hydrolysis to afford the corresponding allenoic acid (Scheme 1e).[8] The limitations are harsh conditions and the use of toxic carbon monoxide, stoichiometric amounts of reductants and limited substrates. On the other hand, C–H activation has been proven to be a powerful tool in synthetic chemistry because of the atom- and step-economy.[9] The synthesis of allenes based on C–H activation is undoubtedly an ideal strategy.[10] We reasoned that the most straightforward approach to allenoic acids would be the use of benzoic acids with carboxylic acid acting as an inherent directing group.[11] Herein, we wish to report the realization of Ru-catalyzed synthesis of allenoic acids *via* direct C–H allenylation of benzoic acids (Scheme 1f).

Our initial attempt began with benzoic acid **1a** and propargylic acetate **2a** in the presence of $[Ru(p\text{-cymene})Cl_2]_2$ and NaOAc at 50 °C using toluene as the solvent. To our delight, the monoallenylation product **3aa** was generated in 7% yield together with 67% recovery of **2a** (Table 1, entry 1). We then investigated the effect of the solvent (Table 1, entries 2–7). The reaction could proceed in dioxane, DCE, CH_3CN, THF, and even in water, albeit affording the monoallenylation product **3aa** and the double allenylation product **4aa** in rather low yields (Table 1, entries 2–6). To our surprise, the yield could be greatly improved when the reaction was conducted in MeOH: 36% yield of the monoallenylation product **3aa** and 16% yield of the double allenylation product **4aa** were obtained (Table 1, entry 7). We next examined a series of additives as shown in entries 8–13: when K_2CO_3 was employed, the reaction gave a 64% combined
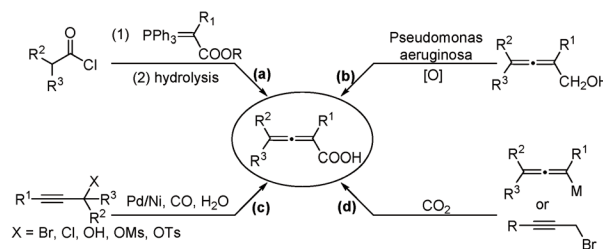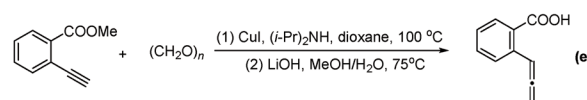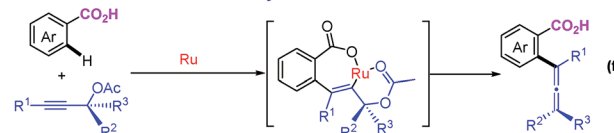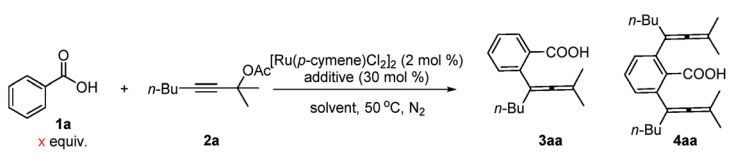
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Scheme 1 Approaches to allenoic acids.

Table 1 Optimization of the *ortho*-allenylation of benzoic acid **1a**[a]

Entry	x	Solvent	Additive	Time (h)	Combined yield (3aa/4aa)[b] (%)	Recovery of 2a[b] (%)
1	1.5	Toluene	NaOAc	46	7 (7/0)	67
2	1.5	Dioxane	NaOAc	46	15 (12/3)	20
3	1.5	DCE	NaOAc	46	14 (14/0)	24
4	1.5	CH$_3$CN	NaOAc	46	27 (23/4)	26
5	1.5	THF	NaOAc	46	30 (23/7)	6
6	1.5	H$_2$O	NaOAc	46	19 (12/7)	23
7	1.5	MeOH	NaOAc	46	52 (36/16)	13
8	1.5	MeOH	NaOAc	12	57 (39/18)	29
9	1.5	MeOH	K$_3$PO$_4$	12	41 (30/11)	7
10	1.5	MeOH	*t*-BuONa	12	22 (18/4)	21
11	1.5	MeOH	Na$_2$CO$_3$	12	57 (40/17)	18
12	1.5	MeOH	Cs$_2$CO$_3$	12	60 (42/18)	20
13	1.5	MeOH	K$_2$CO$_3$	12	64 (45/19)	27
14	1.5	EtOH	K$_2$CO$_3$	12	69 (36/33)	9
15	2.0	MeOH	K$_2$CO$_3$	12	66 (48/18)	16
16	2.4	MeOH	K$_2$CO$_3$	12	63 (50/13)	16
17	2.6	MeOH	K$_2$CO$_3$	12	72 (58/14)	20
18	3.0	MeOH	K$_2$CO$_3$	12	61 (52/9)	21
19[c]	2.6	MeOH	K$_2$CO$_3$	12	67 (56/11)	18
20[c]	2.6	MeOH	—	12	0	42

[a] The reaction was conducted with **1a**, **2a** (0.2 mmol), [Ru(*p*-cymene)Cl$_2$]$_2$ (0.004 mmol), and an additive (0.06 mm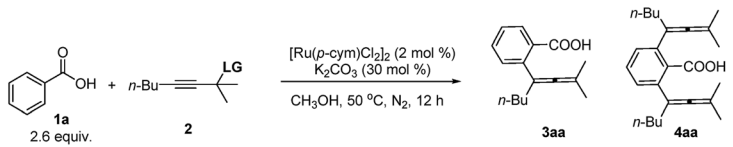ol) in a solvent (0.5 mL) at 50 °C. [b] Determined by ^{1}H NMR analysis using CH$_2$Br$_2$ as the internal standard. [c] In air.

yield of **3aa** and **4aa** (Table 1, entry 13). The reaction in EtOH under optimal conditions led to increased combined yield but a lower selectivity of **3aa**/**4aa** (Table 1, entries 13–14). When 2.6 equiv. of benzoic acid were used, the yield was the highest with a selectivity of 58/14 (Table 1, entry 17). When the reaction was conducted in air, the influence was negligible (Table 1, entry 19). In the absence of the base, we only observed the decomposition of propargylic acetate.

We further investigated the effect of the leaving group (LG) by studying the reaction of benzoic acid **1a** with several propargylic alcohol derivatives and found that OAc was still the best leaving group (Table 2).

With the optimized reaction conditions in hand, the scope of the reaction was investigated at the 1.0 mmol scale (eqn (1) and (2) and Table 3). The parent benzoic acid **1a** afforded the monoallenylation product **3aa** in 55% yield together with the diallenylation product **4aa** in 10% yield (eqn (1)). 4-
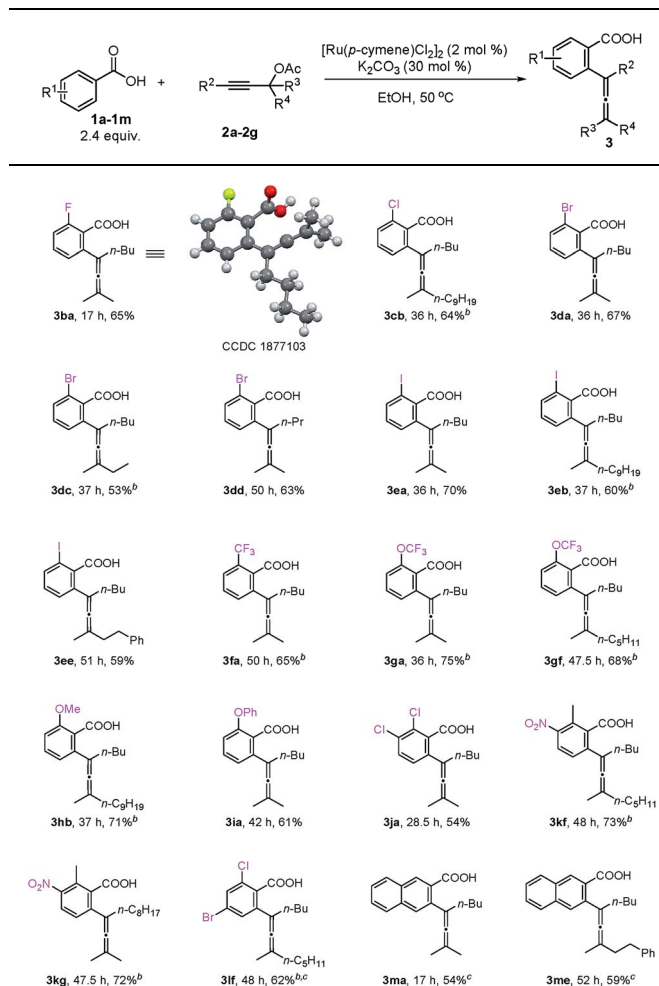
Table 2 Effect of the leaving groups

Entry	LG	Combined yield (3aa/4aa)[a] (%)	Recovery of 2a[a] (%)
1	OMe (**2a$_1$**)	0 (—/—)	59
2	OCO$_2$Me (**2a$_2$**)	54 (40/14)	9
3	OCOEt (**2a$_3$**)	57 (47/10)	20
4	OBoc (**2a$_4$**)	61 (47/14)	21
5	OAc (**2a**)	72 (58/14)	20

[a] Determined by ^{1}H NMR analysis using CH$_2$Br$_2$ as the internal standard.

Table 3 Reaction scope[a]

CCDC 1877103

3ba, 17 h, 65%
3cb, 36 h, 64%[b]
3da, 36 h, 67%
3dc, 37 h, 53%[b]
3dd, 50 h, 63%
3ea, 36 h, 70%
3eb, 37 h, 60%[b]
3ee, 51 h, 59%
3fa, 50 h, 65%[b]
3ga, 36 h, 75%[b]
3gf, 47.5 h, 68%[b]
3hb, 37 h, 71%[b]
3ia, 42 h, 61%
3ja, 28.5 h, 54%
3kf, 48 h, 73%[b]
3kg, 47.5 h, 72%[b]
3lf, 48 h, 62%[b,c]
3ma, 17 h, 54%[c]
3me, 52 h, 59%[c]

[a] The reaction was conducted with 1 (2.4 mmol), 2 (1.0 mmol), [Ru(p-cymene)Cl₂]₂ (0.02 mmol), and K₂CO₃ (0.3 mmol) in EtOH (2.5 mL) at 50 °C. [b] 4 mol% [Ru(p-cymene)Cl₂]₂. [c] 5.0 mL of EtOH as the solvent.

Bromobenzoic acid 1n was converted to the monoallenylation product 3na (45% yield) and the bisallenylation product 4ma (26% yield) under the standard conditions (eqn (2)).

(1)

(2)
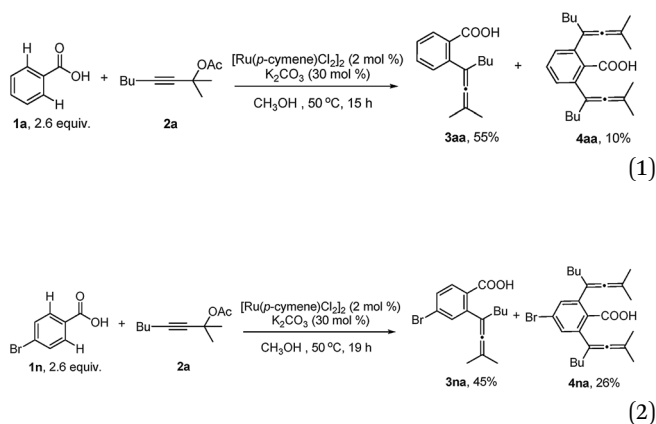
For mono-*o*-substituted benzoic acids, electron withdrawing groups, such as halogen atoms (including fluorine, chlorine, bromine, and iodine), CF₃, and OCF₃, were all well tolerated (Table 3, **3ba–3gf**). The structure of **3ba** was confirmed by X-ray diffraction study.[12] Mono-*o*-substituted benzoic acids containing the electron-donating groups methoxy and phenoxy afforded allenylation products **3hb** in 71% yield and **3ia** in 61% yield, respectively. 2,3-Dichlorobenzoic acid (**3ja**), 2-methyl-3-nitrobenzoic acid (**3kf** and **3kg**) and 2-chloro-4-bromobenzoic acid (**3lf**) were also allenylated in moderate to good yields. Notably, when the reaction was conducted with β-naphthoic acid, which has more than one C–H bond, 3-allenylation products **3ma** and **3me** were obtained exclusively. The scope of C–H allenylation with regard to propargylic acetates was also investigated affording **3cb**, **3nb**, **3dc**, **3dd**, **3ee**, **3le**, **3kf**, **3gf**, or **3kg** smoothly.

A gram scale reaction using 2-bromobenzoic acid **1d** with **2g** afforded the allenylation product **3dg** in 67% yield (eqn (3)).
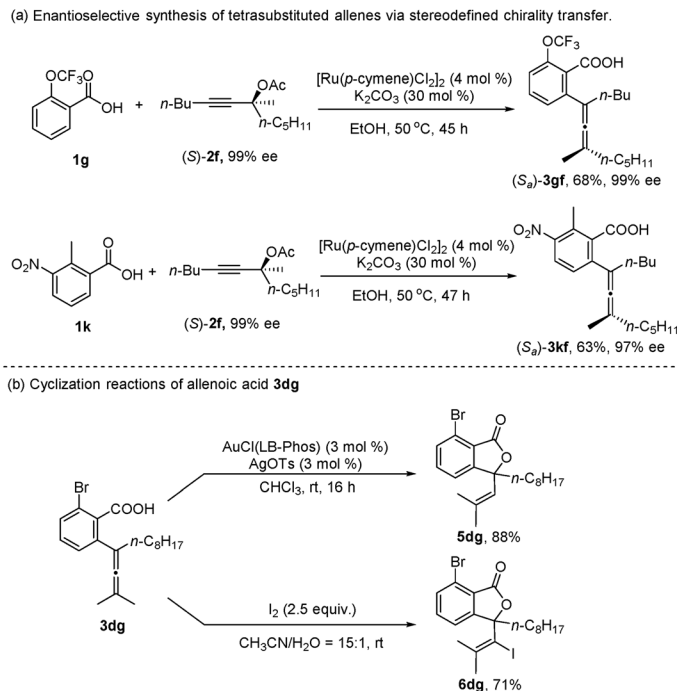

(3)
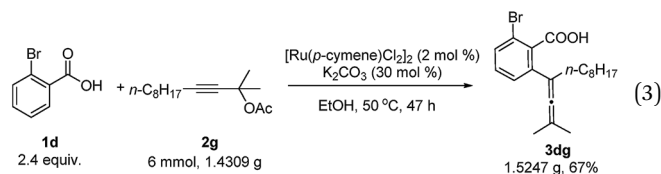
In addition, when the optically active acetate (*S*)-**2f** (99% ee) was applied, axially chiral allenoic acids (*S*ₐ)-**3gf** (99% ee) and (*S*ₐ)-**3kf** (97% ee) could be obtained with highly efficient chirality transfer (Scheme 2a). This method may open a new avenue for developing practical and synthetically useful methodologies for the synthesis of optically active allenoic acids. Meanwhile, this result indicated that the coordination of acetates and ruthenium species dictated the regioselectivity of alkyne insertion and the stereoselectivity of β-OAc elimination.

To further explore the synthetic utility of this method, several synthetic applications were studied (Scheme 2b). The allenoic acid **3dg** was easily transformed into the lactone **5dg** by treatment with AuCl(LB-Phos) and AgOTs.[13] This allenoic acid may also undergo an iodolactonization reaction with iodine to afford **6dg** in 71% yield.
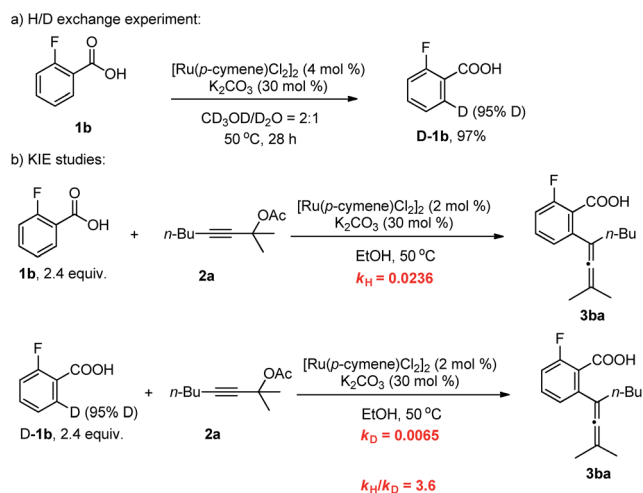
To gain insight into the mechanism of this methodology, several control experiments were carried out. Firstly, when 2-fluorobenzoic acid **1b** was added to a mixture of CD₃OD and D₂O (2 : 1), the corresponding benzoic acid **D-1b** with 95% deuterium incorporation was obtained (Scheme 3a), indicating that the C–H activation step was reversible in the catalytic system. Subsequently, the parallel reactions of **1b** and **D-1b** with **2a** were conducted. We measured the reaction rate (*k*) of both **1b** and **D-1b** by monitoring the concentration of the product **3ba** by NMR from 2.5 h to 8 h (Scheme 3b and Fig. 1). Then, the primary kinetic isotope effect of 3.6 was observed. These results suggest that C–H bond cleavage is the rate-determining step.[14]

In addition, a first-order dependence of the initial rate on the amount of the Ru catalyst was established (Fig. 2a, see the
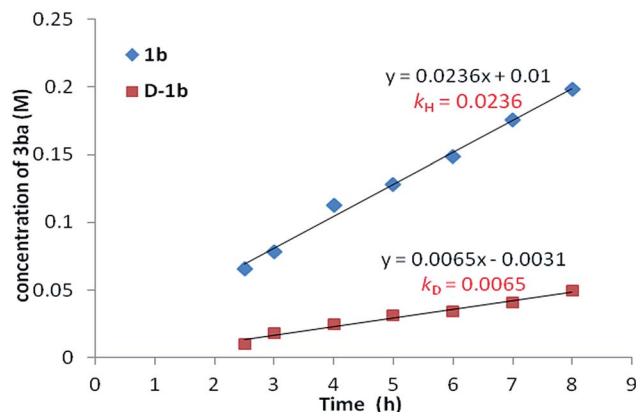
(a) Enantioselective synthesis of tetrasubstituted allenes via stereodefined chirality transfer.

(b) Cyclization reactions of allenoic acid 3dg

Scheme 2 Synthetic applications.

a) H/D exchange experiment:

b) KIE studies:

$k_H/k_D = 3.6$

Scheme 3 Mechanistic studies.

Fig. 1 Plot of the concentrations of **3ba** vs. time.

ESI† for details). The reaction orders of each reactant were also measured by using 2-fluorobenzoic acid **1b** and propargylic acetate **2a**. Both **1b** and **2a** follow the first-order reaction rate law, according to the linear relationship with $\ln([\mathbf{1b}])$ vs. reaction time: $\ln([\mathbf{1b}]) = -k_1 t + \ln[\mathbf{1b_0}]$ (Fig. 2b) and $\ln([\mathbf{2a}]) = -k_2 t + \ln[\mathbf{2a_0}]$ (Fig. 2c). Based on these data, we may give the rate equation as $d[\mathbf{3ba}]/dt = k[\text{Ru}] \cdot [\mathbf{1b}] \cdot [\mathbf{2a}]$.

To further understand the role of benzoic acid on the reaction, four experiments were conducted by using different molar ratios of 2-fluorobenzoic acid **1b** vs. propargylic acetate **2a** (Fig. 2d). The yield vs. time profile is almost the same in the initial four hours, indicating that the loading of benzoic acid has a very limited effect on the formation of the final

product; the excess benzoic acid did not accelerate the formation of the product greatly.[15] In addition, we observed that the reaction failed to afford the expected product in the absence of k_2CO_3, indicating a CMD process for the C–H cleavage. However, due to its catalytic nature, the role of benzoic acid as a Brønsted acid to promote the insertion process from **Int 2** to **Int 3** cannot be excluded.[16]

Based on these investigations above, the proposed catalytic cycle is illustrated in Scheme 4. Firstly, the C–H activation step leads to the formation of the cyclic intermediate **Int 1** via a CMD process. Subsequently, **Int 2** is generated by the coordination of the carbonyl unit in acetate with the Ru in the cycloruthenated species, which subsequently undergoes the syn-insertion of a C–C triple bond to afford **Int 3**. After a syn-β-OAc elimination step, the allenylation product was generated and the ruthenium
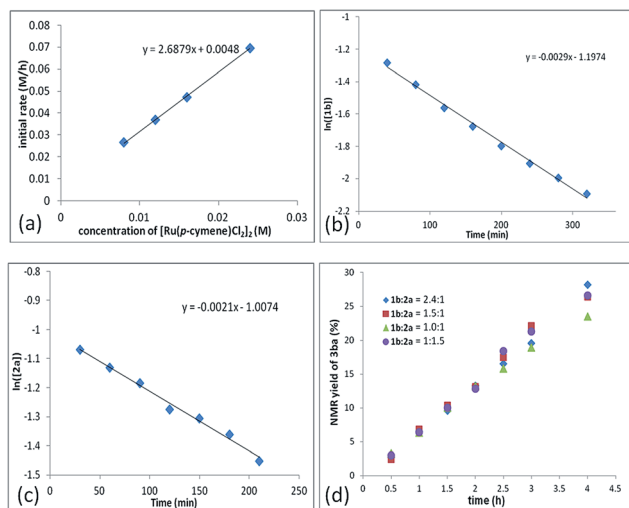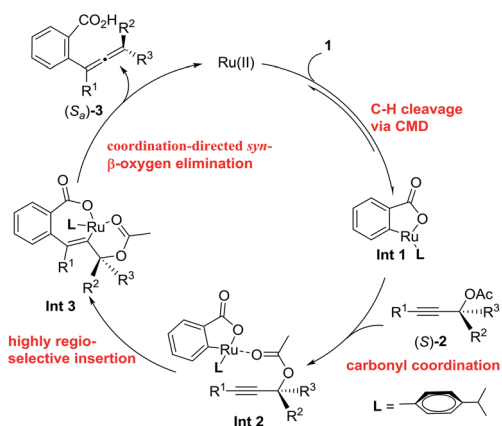
Fig. 2 The dependence of the initial reaction rate on [Ru(p-cymene) Cl$_2$]$_2$ (a), 2-fluorobenzoic acid **1b** (b), and propargylic acetate **2a** (c). NMR yield of **3ba** $vs.$ time with different molar ratios of **1b** and **2a** (d).

Scheme 4 A possible mechanism.

species was released to restart the cycle. It should be noted that the acetate plays an important role in the syn-insertion as well as the syn-β-OAc elimination step.

Conclusions

In conclusion, we have established a new strategy to access allenoic acids, which is based on ruthenium catalysed carboxylic acid-directed C–H allenylation of benzoic acids with propargylic acetates. The reaction is compatible with air and synthetically useful functional groups such as Cl, Br, I, and OCF$_3$ are all tolerated. Optically active allenoic acids could also be prepared through highly efficient chirality transfer. The formed allenoic acids could be transformed to lactones efficiently under mild conditions.

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

Notes and references

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12 **3ba**: $C_{16}H_{19}FO_2$, MW = 262.32, monoclinic, space group $P121/c\ 1$, final R indices $[I > 2\backslash s(I)]$, R1 = 0.0602, wR2 = 0.1483; R indices (all data), R1 = 0.0962, wR2 = 0.1764; a = 10.7724(13) Å, b = 15.7829(12) Å, c = 9.9065(12) Å, α = 90.00°, β = 116.778(15)°, γ = 90.00°, V = 1503.7(3) Å^3, T = 296(2) K, Z = 4, reflections collected/unique 5576/2735 (R_{int} = 0.0294), number of observations [>2σ(I)]: 1722, parameters: 179. Supplementary crystallographic data have been deposited at the Cambridge Crystallographic Data Centre, CCDC 1877103.†

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