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## **Electronic Supplimentary Information**

## Water-based efficient alkyne transformation towards α-acetoxy/imido-ketones via oxidative coupling reactions using an alkylamine catalyst

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<u>Serial N</u>	No. <u>Content</u>	Page Numbers
1.	Control experiments data	S-2
2.	<sup>1</sup> H and <sup>13</sup> C-NMR spectra of synthesised $\alpha$ -acetoxyketones (4a-g), $\alpha$ -imidoketones (5a-g), <i>N</i> -protected cathinone (6a), cathinone (7)	
	and N-benzoylphthalimide (8a)	S-3

## 1. Control experiments data

## Detection of labeled isotope (<sup>18</sup>O) using mass spectroscopy

The synthesis of compound **5a** was performed on a 0.1 mmol scale in 1.0 ml H<sub>2</sub>O<sup>18</sup>, following the general procedure as depicted earlier. High-resolution mass spectrometry (HRMS) analysis was then performed directly with the post-reaction mixture. The peak at 267.0784 (M<sup>+</sup>) confirmed the incorporation of the O<sup>18</sup> isotope into the desired product (<sup>18</sup>O-**5a**), indicating the source of oxygen in the newly generated  $\alpha$ -functionalised ketones.

## SI Figure 1: HRMS spectrum of <sup>18</sup>O-5a



# 2. <sup>1</sup>H and <sup>13</sup>C-NMR spectra of synthesised $\alpha$ -acetoxyketones (4a-g), $\alpha$ -imidoketones (5a-g), *N*-phthalimido cathinone (6a), cathinone (7) and *N*-benzoylphthalimide (8a)

SI Figure 2: <sup>1</sup>H and <sup>13</sup>C-NMR spectra of compound 4a



SI Figure 3: <sup>1</sup>H and <sup>13</sup>C-NMR spectra of compound 4b



SI Figure 4: <sup>1</sup>H and <sup>13</sup>C-NMR spectra of compound 4c













### 8.8.007 9.8.107 9.9.17,950 9.9.17,950 9.9.17,950 9.9.17,950 9.9.17,950 9.9.17,750 9.9.17,750 9.9.17,750 9.9.17,750 9.9.17,750 9.17,7500 9.17,7500 9.17,7500 9.17,7500 9.17,750





# 







# SI Figure 14: <sup>1</sup>H and <sup>13</sup>C-NMR spectra of compound 5f





f1 (ppm) 

S-16

## SI Figure 16: <sup>1</sup>H and <sup>13</sup>C-NMR spectra of compound 6a



f1 (ppm) 

SI Figure 17: <sup>1</sup>H and <sup>13</sup>C-NMR spectra of compound 7





78.020 8.021 8.012 8.010



0 190 170 100 f1 (ppm) 80 70 60 50 40 30 20 10 ( 180 160 150 140 120 110 90 130