## Supporting information for

#### Diastereoselective copper-catalyzed 1,4-addition of Grignard reagents to N-enoyl oxazolidinones

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#### **General Information**

All reactions were carried out in inert atmosphere of Ar. Solvents were dried and purified by standard methods before use. Chemical shifts ( $\delta$ ) are given in ppm relative to tetramethylsilane for <sup>1</sup>H NMR and <sup>13</sup>C NMR. Specific optical rotations are given in deg cm<sup>-3</sup>.g<sup>-1</sup>.dm<sup>-1</sup>. Flash chromatography was performed on silica gel 40 – 63 µm. Thin-layer chromatography was performed on TLC-plates silica gel 60 with F-254 UV indicator. Diastereomeric ratios were determined by <sup>1</sup>H NMR and GC. Enantiomeric ratios were determined by HPLC on Chiralpak, OD-H, IB, AS-H (column using hexane/*i*PrOH as a mobile phase and detection with UV-detector at 254 nm, 218 nm.









#### DEPT for 2a









## <sup>1</sup>H NMR for **4b**



# <sup>13</sup>C NMR for **4b**





<sup>1</sup>H NMR for **4c** 





<sup>1</sup>H NMR for **4d** 



#### DEPT for 4d



<sup>1</sup>H NMR for **5b** 



# <sup>13</sup>C NMR for **5b**



#### DEPT for **5b**

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|  | CH2 carbons<br>CH2 carbons<br>CH carbons<br>All carbons<br>All carbons<br>All carbons  |

## <sup>1</sup>H NMR for **6a**



## <sup>13</sup>C NMR for **6a**





<sup>1</sup>H NMR for **6b** 







# <sup>13</sup>C NMR for 6c





#### <sup>1</sup>H NMR for 7a





## <sup>1</sup>H NMR for **7b**





<sup>1</sup>H NMR for **7**c





## <sup>1</sup>H NMR for **7d**



<sup>13</sup>C NMR for **7d** 



#### DEPT for 7d

