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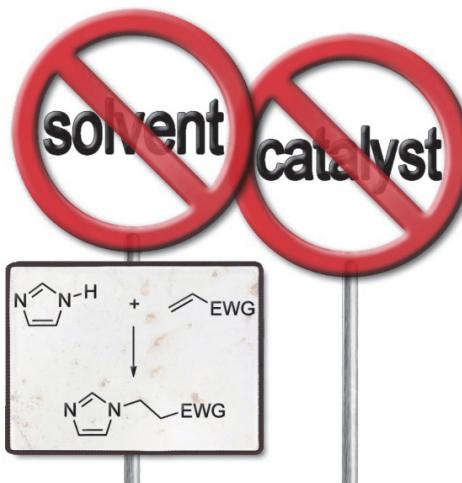
Solvent- and catalyst-free aza-Michael addition of imidazoles and related heterocycles

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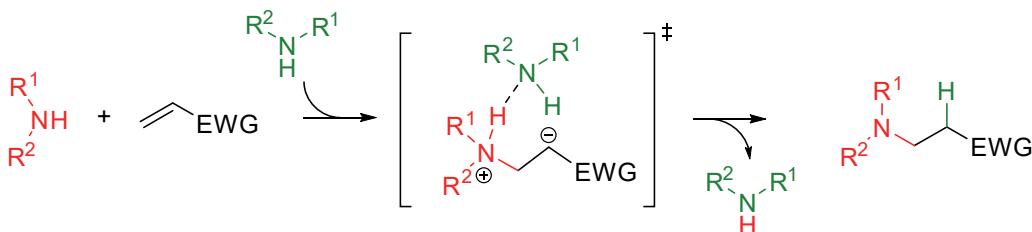


Abstract. This work demonstrates the scope and limitations of the aza-Michael addition of imidazoles and related heterocycles with electron deficient olefins under solvent- and catalyst-free conditions. The reaction proceeds at 80°C within hours towards completion as long as the azole derivative is sufficiently soluble in the Michael acceptor, which has been used in small excess. Workup only comprises evaporation of surplus Michael-acceptor and no additional solvents are necessary for purifying the products.

Introduction. The ideal chemical reaction proceeds fast at ambient temperature and quantitatively yields a single product without the need of a catalyst. This means no side- but also no byproducts are formed, resulting in perfect atom economy¹ and full consumption of educts. In this case, no purification or any other workup is necessary. In reality, the situation is in most cases different. Conversions are not complete, catalysts are needed to speed up the reaction or to provide alternative reaction pathways and/or selectivity, by- and side products form and often solvents have to be employed. Solvent use in organic chemistry is in many cases inevitable for providing high reaction rates, for obtaining the desired selectivity or for facilitating the heat management of a reaction. Moreover, many purification techniques heavily rely on the use of solvents. Accordingly, solvent use is in many cases accountable for a huge part of the synthesis' negative impact on the environment. Consequently, research striving for solvent-free and at the same time catalyst-free reactions evolved as a branch of green chemistry.² However, many examples termed solvent-free in literature are using solvents during isolation and purification to meet the stringent requirements in terms of purity in preparative organic chemistry. Taking into account, that usually much more solvent is needed during purification than during preparation, the impact of performing a reaction without solvent (and catalyst) in saving resources is rather limited as soon as a solvent-based purification is necessary.³ In other words, solvent- and catalyst-free reactions should ideally proceed to completeness without forming side- or byproducts that have to be separated with the help of solvent-based methods in order to provide a significant advantage in terms of sustainability.

Among the many classes of different reactions, in particular addition reactions qualify for designing side- and byproduct free transformations without the need of catalysts and solvents at any stage of the preparation. Some selected examples comprise the cycloaddition of aziridines and carbon dioxide,⁴ hydrophosphination of alkenes,⁵ Diels-Alder reactions,⁶ Knoevenagel reaction of aldehydes (which is a condensation type reaction)⁷ or the aza-Michael reactions.⁸ The aza-Michael reaction involves a nucleophilic amine (the Michael donor) and an electron deficient alkene (the Michael acceptor). Generally, it is performed with the aid of a catalyst under mild reaction conditions leading to high conversions with an atom economy of 100%. Side-products are usually not observed. The works of Desmet et al.⁹ and Bláha, et al.^{8c} describe the mechanistic understanding of the (uncatalyzed) reaction in detail. In essence, the aza-Michael addition is a third-order reaction (first order in the acceptor and second order in the donor). The third-order

kinetics are rationalized by the formation of a rate-determining zwitterionic transition state in which a second amine is needed to facilitate the proton transfer from the attacking amine to the α -carbon of the electron deficient olefin (Scheme 1).



Scheme 1 Mechanism of the aza-Michael reaction according to Refs 8c and 9.

The mechanism implies that the aza-Michael reaction should ideally proceed with amines of high nucleophilicity (to facilitate the formation of the zwitter-ion) and bearing loosely bound protons (facilitating the hydrogen transfer). Preferably, reactions should be performed with the highest possible concentration of the reactants and not too high temperature (because the reaction is of third-order). Accordingly, the aza-Michael reaction is perfectly suitable for being performed in an uncatalyzed solvent-free manner. Surprisingly, to the best of our knowledge, only three publications on solvent- and catalyst-free aza-Michael addition are available and only aliphatic amines have been investigated so far.⁸

Herein, we wish to elucidate the reactivity of azoles with Michael acceptors under solvent and catalyst free conditions. Using azoles, like imidazoles or pyrazoles is an attractive target, because they show rich Michael chemistry with outreach into material, medicinal and biological chemistry. A recent review nicely summarizes the state of the art in aza-Michael additions of imidazoles.¹⁰ Interestingly it has been stated in that work, that only a few reports on reactions that satisfactorily proceed only in solvent without catalysis are known. The question whether this lack of reactivity is due to the use of unfavorable reaction conditions or the electronic features of azoles remains. The nucleophilicities and the acidities of imidazoles and related heterocycles suggest them to be poorer aza-Michael donors than aliphatic amines. Their nucleophilicity is comparable to the one of secondary aliphatic amines as judged by the methyl cation affinity (imidazole 531.7 kJ/mol *vs.* dimethylamine 523.1 kJ/mol)¹¹ or less pronounced considering the nucleophilicity parameter N (imidazole N = 11.47¹² *vs.* dimethylamine N = 15.10¹³). Moreover, the acidity of the proton residing at the second nitrogen atom is about 3-4 orders of magnitude

lower than in aliphatic amines (e.g. imidazole $pK_a = 13.9$ vs. $pK_a = 10.73$ for diethylamine).¹⁴ The investigations presented here are intentionally restricted to Michael-acceptors, which can be separated upon evaporation in order to avoid a work-up involving any solvent.

Results and Discussion. One equivalent of the Michael donor and 1.2 equivalents of the Michael acceptor were weighted into a Schlenk tube and heated to 80°C under constant magnetic stirring. The reaction progress was monitored by sampling a small fraction and recording ^1H -NMR spectra in regular intervals. First, imidazole (**1**) as the Michael donor and acrylonitrile (**A**), methyl acrylate (**B**), methyl crotonate (**C**, methyl (E)-but-2-enoate) or methyl methacrylate (**D**) were used as Michael acceptors. The Michael adducts **1A** and **1B** quantitatively formed within 3 and 5 h, respectively.

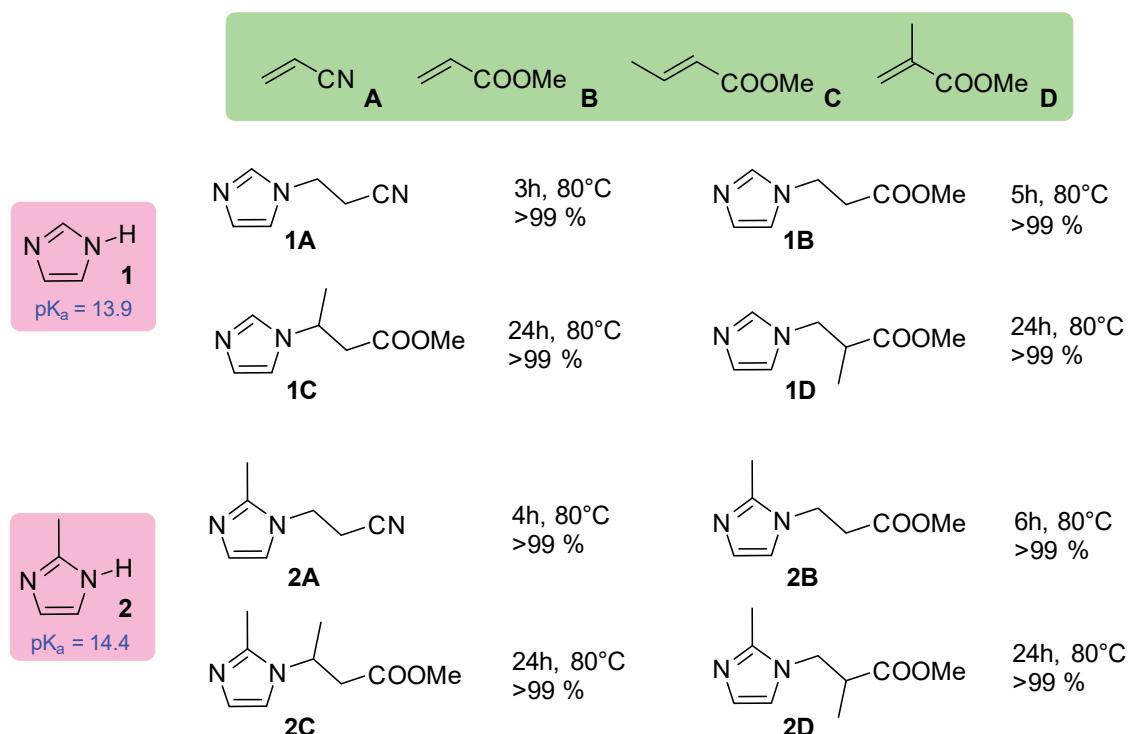


Figure 1. Conversions after given time of the aza-Michael addition of **1** and **2** to acrylonitrile (**A**), methyl acrylate (**B**), methyl (E)-but-2-enoate (**C**) and methyl methacrylate (**D**) performed at 80°C under solvent and catalyst free conditions. Isolated yields are in all cases above 97%.

When using **C** and **D** as the Michael acceptors complete conversion was observed after about 24 h. The different speeds of the reaction can be attributed to the different electrophilicity

(expressed by the electrophilicity parameter E) of the employed Michael acceptors decreasing in the order of **B** (-18.84) \approx **A** (-19.05) \gg **D** (-22.27) \approx **C** (approx. -23).¹⁵ The reaction of 2-methyl imidazole (**2**) with **A-D** was somewhat slower than with **1**. This might be explained by the lower pK_a-value calculated for **2** when compared to **1** (Fig. 1), since the nucleophilicity of **2** is slightly higher (N = 11.74 vs. **1**: N = 11.47¹²).

To further examine the impact of the acidity of the heterocycles' protons, a selection of different Michel donors covering a range of different acidities was made (Fig. 2, left). Their performance in the reaction with methyl acrylate (**B**) was investigated under the same reaction conditions as used before. Relatively acidic 1,2,4-triazole (**3**, exhibiting a pK_a of 10.2) resulted in quantitative conversion towards **3B** in only 2 h. In contrast, 2-phenylimidazole (**4**) needed 23 times longer. Compound **4** is distinctly less acidic than **3** but more acidic than **1** and **2**. Similarly, benzo[d]imidazole (**5**, pK_a = 13.9; N = 10.50) needs 49 h for being fully converted into **5B**. In these two cases, the solubility of the heterocycle in **B** is low. While in all other cases discussed so far, the Michael donor compound dissolved within the first 5-15 min of the reaction time in the acceptor component, **4** and **5** needed hours to dissolve completely. Increasing the amount of **B** to 3.7 eq. in respect to **5** allowed for obtaining full conversion after 5.5 h. Accordingly, the poor solubility of **5** in **B** (and analogously of **4** in **B**) is governing the speed of the reaction. Pyrazole (**6**) featuring a pK_a value of 14.0 dissolves fast and gives **6B** in full conversion after 4 h, i.e. **6** is converted at a similar rate like **1** and **2**. Finally, in the case of pyrrole (**7**, pK_a = 17.0) no reaction with methyl acrylate under the given conditions was observed. Since **7** is a liquid which is miscible with **B**, the unsuccessful conversion towards **7B** has to be attributed to its electronic properties. Compound **7** is a poorer nucleophile (N = 4.63¹⁶) than all the other heterocycles under investigation and additionally proton transfer is inhibited because of its low acidity (which is even lower than that of alcohols¹⁷). Imidazole based heterocycles like 2-methyl-4-nitro-1H-imidazole, adenine or 1,9-dimethylguanine are insoluble in **B** and showed no conversion towards the respective Michael addition product within 24 h.

Switching from Michael acceptor **B** to the less electrophilic (E = -20.22¹⁵) and less polar *t*-butyl acrylate (**E**) corroborates the influence of solubility on the speed of the reaction. Conversions with the azoles **1-6** after about 24 h are not complete and numbers are given in Fig. 2 (right). In all cases, heterocycles dissolved more slowly contributing to the distinctly longer reaction times.

This was confirmed by using a higher excess of **E** (5 equiv.) which allowed for shortening the reaction time giving a conversion of 97% towards **2E** in 24 h. Accordingly, in these cases, a solvent and catalyst free protocol is lacking merit.

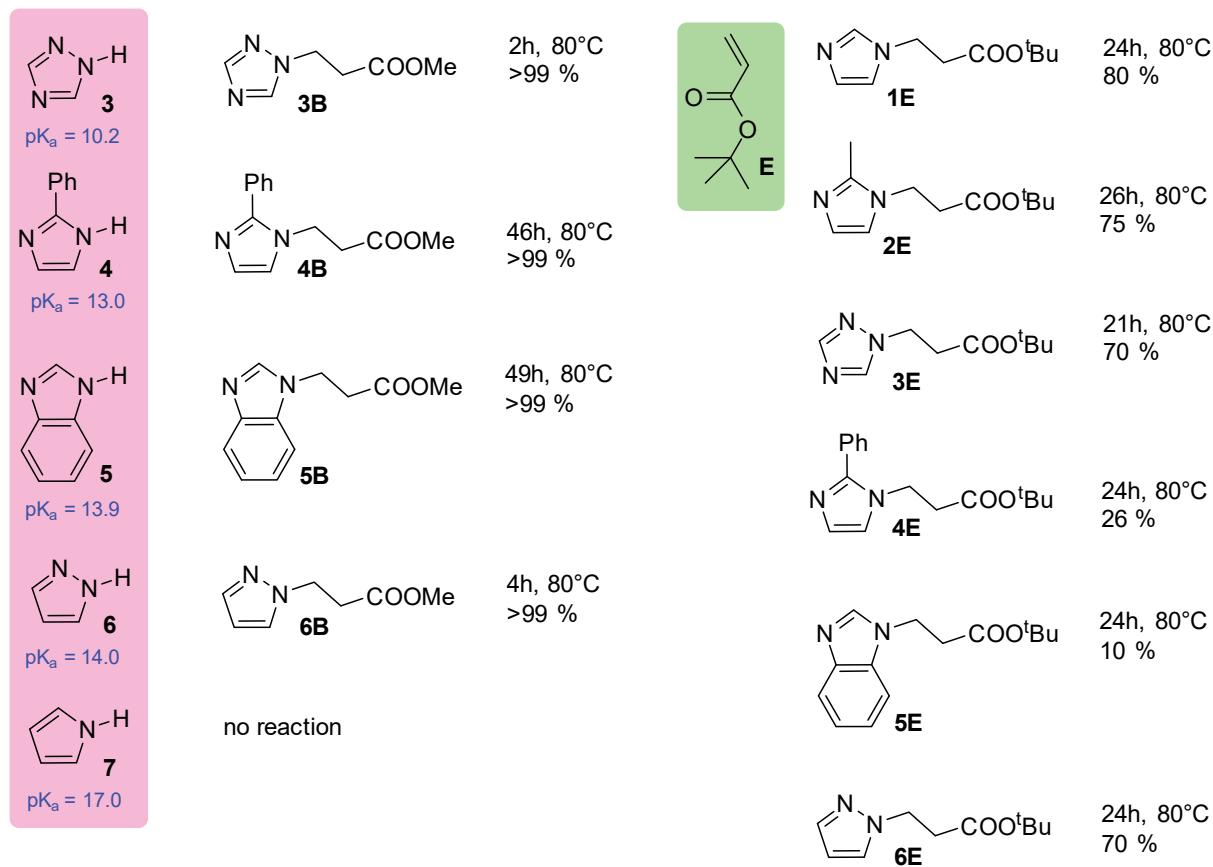


Figure 2. Conversions after given time of the aza-Michael addition of **3-7** with methyl acrylate (**B**) or *t*-butyl acrylate (**E**) performed at 80°C under solvent and catalyst free conditions. Isolated yields of **3B-6B** are above 97%.

In a next step, the preparation was further simplified and stirring was omitted in order not to lose yield when removing the magnetic stirring bar. The Michael-donor (1 eq.) and the Michael acceptor (1.2 eq.) were combined in a flat bottom tube, the tube was locked and placed in an oven operated at 80°C. Under these conditions, the reactions needed two to three times longer to reach completeness. Checking the conversion via repeated ¹H-NMR measurements of the reaction of **2** with **B** revealed a fast reaction (half-life < 6 min; 94% conversion at 74 min) which only sluggishly reached completion (at about 18 h). The excess of **B** was then removed under

vacuum. It is worth noting, that also under these conditions, no side product or decomposition products were observed.

Comparing the obtained results with some from catalytic studies of the aza-Michael addition illustrates the importance of using as little solvent as possible for obtaining fast and quantitative conversion. In a first example from literature, the preparation of compound **1B** using I_2 as the catalyst in refluxing toluene (concentration of **1** = 1 mol/L) was described.¹⁸ The reaction reached completion after 9 h and a yield of 50% upon work-up was obtained. Another example reported that the reaction of **1** (c = 0.25 mol/L) and **B** catalyzed by alkaline protease from *Bacillus subtilis* at 50 °C in pyridine needed a reaction time of 72 h and yielded 76 % of **1B**.¹⁹ In a final example of the many protocols available in literature,¹⁰ **1** (c = 2 mol/L) and **B** were reacted in acetonitrile at room temperature using 0.5 equiv. of 1,8-diazabicyclo[5.4.0]undec-7-ene (DBU) as catalyst. The aza-Michael adduct **1B** was obtained in 95 % yield after 14 h.²⁰ Recalling the performance of the protocol disclosed herein (solvent and catalyst free, 5 h at 80°C, > 97 % yield) clearly reveals that aza-Michael reactions should be carried out in solution using the highest possible concentration of the reactants.

Conclusions. In summary, it has been shown that azoles and related compounds undergo solvent- and catalyst-free Michael addition with methyl acrylates and acrylonitrile in reasonable time as long as the Michael donor is sufficiently soluble in the Michael acceptor at 80°C. Work-up only comprises the removal of the small excess of the Michael acceptor by evaporation. Weaker Michael acceptors being at the same time poorer solvents for the azole derivatives require reaction times of more than 24 h to reach completeness. In such cases, the use of solvents and catalysts is advisable. The findings contain a caveat for all researching catalysis in aza-Michael additions, namely that the concentration of the reaction partners is of crucial importance for determining the speed of the reaction, just as expected for a third-order reaction. Finally, the statement issued in a recent review, that only few reports on reactions that satisfactorily proceed only in solvent without catalysis are known, has been demystified.

Experimental. All reactions were performed under ambient conditions. Chemicals were purchased from TCI, Acros or Sigma-Aldrich and were used as received (stabilizers present in

the Michael acceptors were not removed). ^1H and ^{13}C -NMR spectra were recorded at 23 °C on a Bruker Ultrashield 300 spectrometer operating at 300.36 MHz (^1H) and 75.53 MHz (^{13}C). Deuterated solvents were purchased from Cambridge Isotope Laboratories.

A general procedure is given by the example of the preparation of **2B**. Methyl acrylate (3.3 mL, 0.0364 mol, 1.2 eq.) was added to 2-methylimidazole (2.487 g, 0.0303 mol, 1.0 eq.) in a 15 mL Schlenk tube. The suspension was heated up to 80 °C under constant stirring whereupon dissolution of 2-methylimidazole occurred within approx. 5 min. After 6 h the reaction mixture was cooled to room temperature and the resulting oil was dried under reduced pressure (to get rid of the excess of methyl acrylate) whereupon the colorless oil solidified. The magnetic stirrer bar was removed from the product. Yield: 4.97 g, (0.0295 mol), 97.5 % o. Th., colorless powder. Analytical data in accordance with literature.²¹

Supporting Information available: Analytical data of the products

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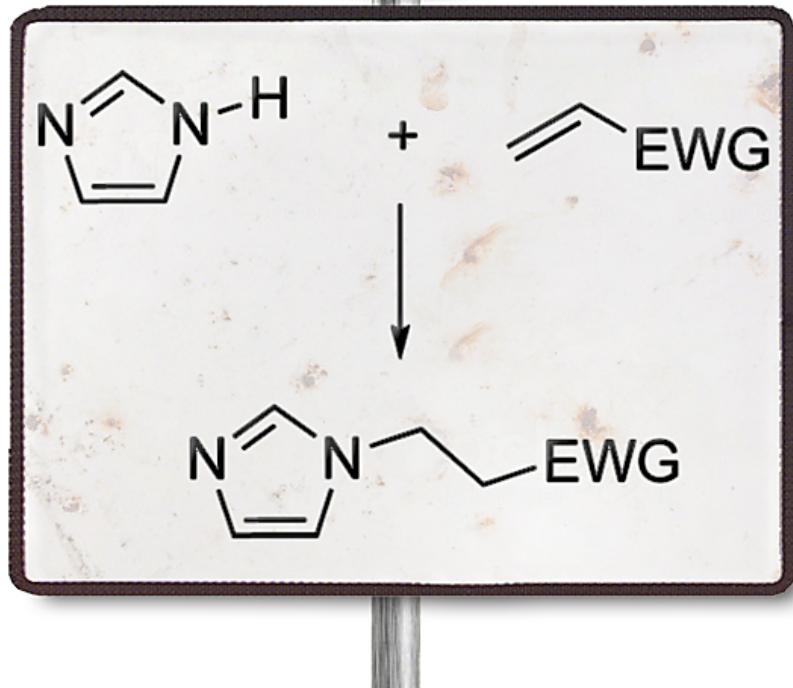
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solvent

catalyst



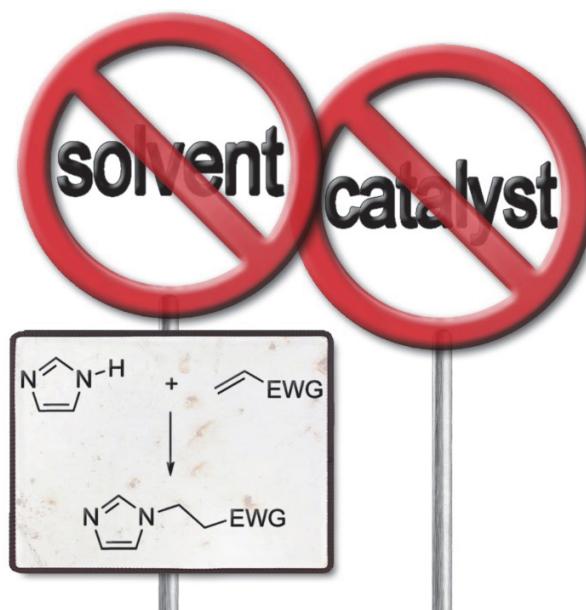
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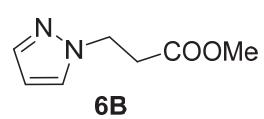
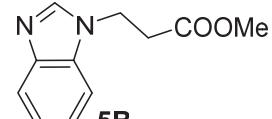
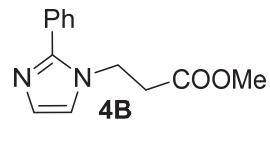
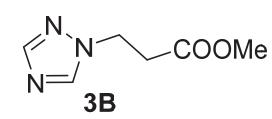
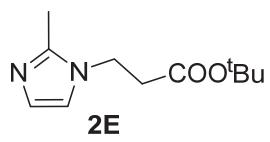
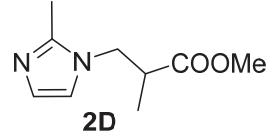
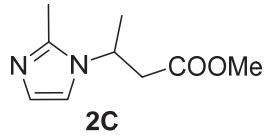
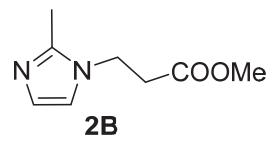
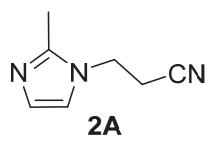
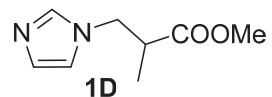
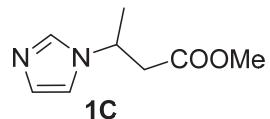
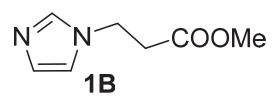
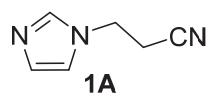
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^b Christian Doppler Laboratory for Organocatalysis in Polymerization, Stremayrgasse 9, A 8010
Graz, Austria

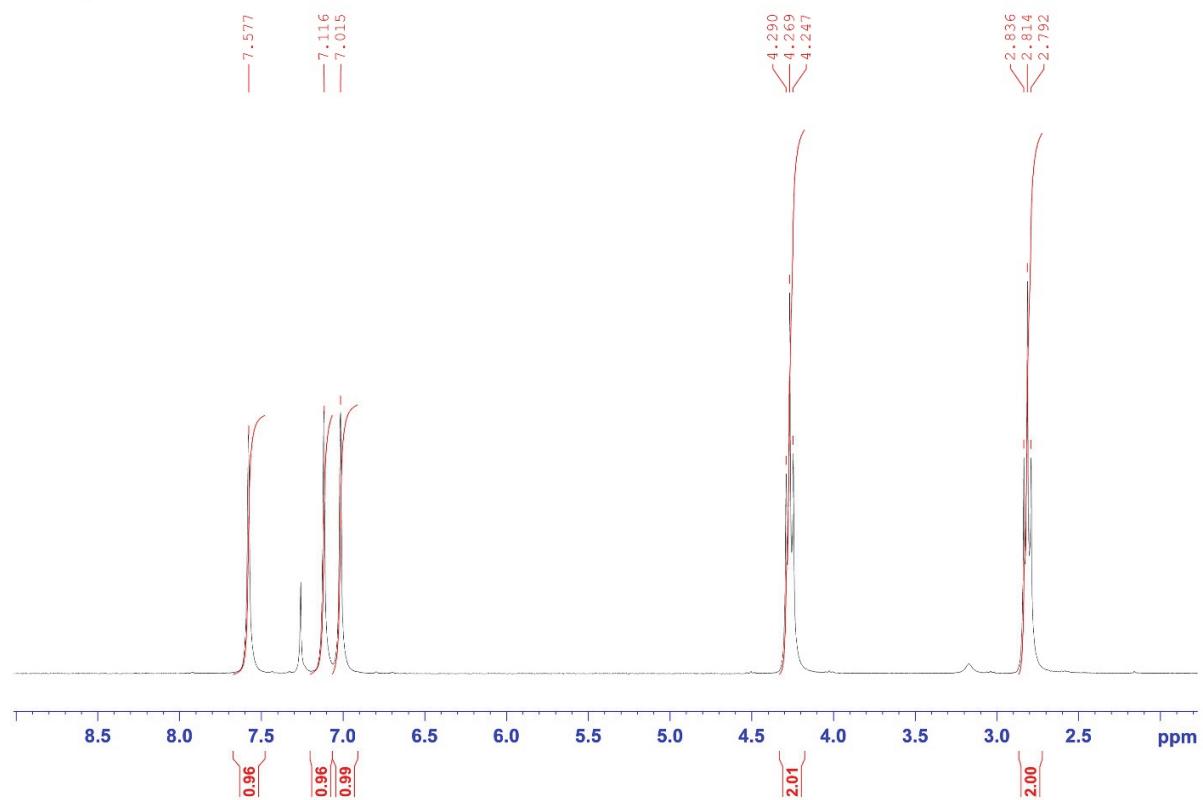
Supporting Information



^1H - and $^{13}\{\text{H}\}$ -NMR spectra recorded at 23°C on a Bruker Ultrashield 300 spectrometer operating at 300.36 MHz (^1H) and 75.53 MHz (^{13}C) in CDCl_3 of the following compounds are given in Figs S1-S13.



KKo22-1_2, Kodolitsch



KKo22-1_2, Kodolitsch

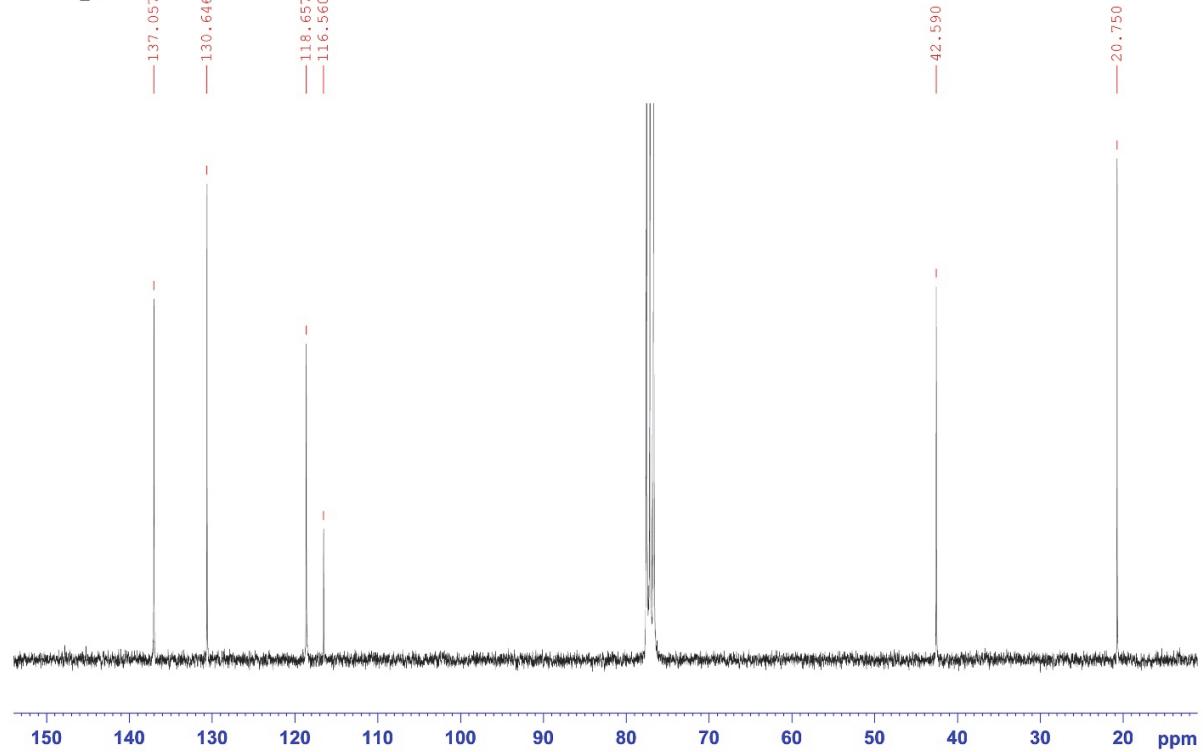


Fig S1. ^1H - and $^{13}\{\text{H}\}$ -NMR spectra of 1A

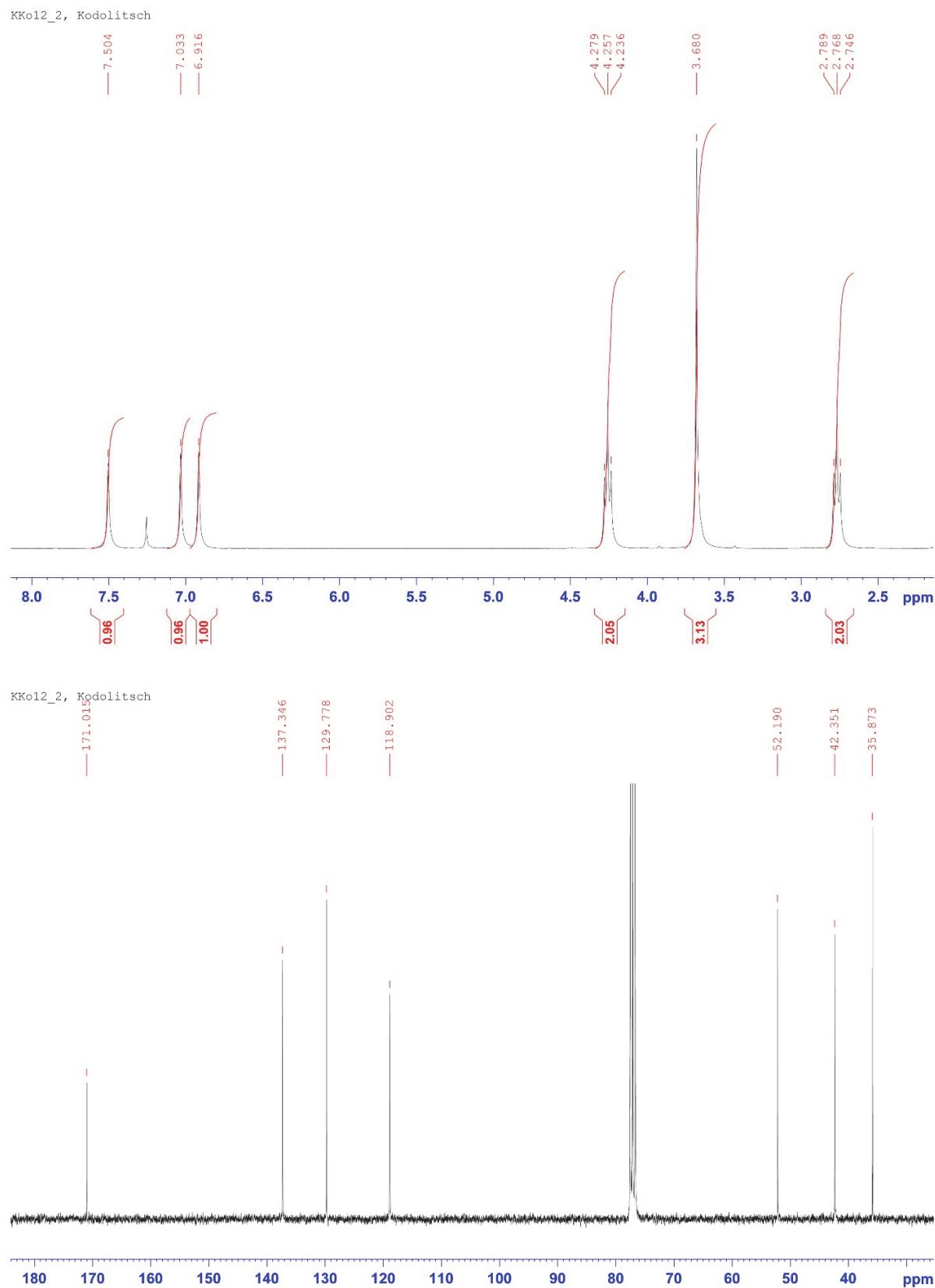


Fig S2. ^1H - and $^{13}\{\text{H}\}$ -NMR spectra of **1B**

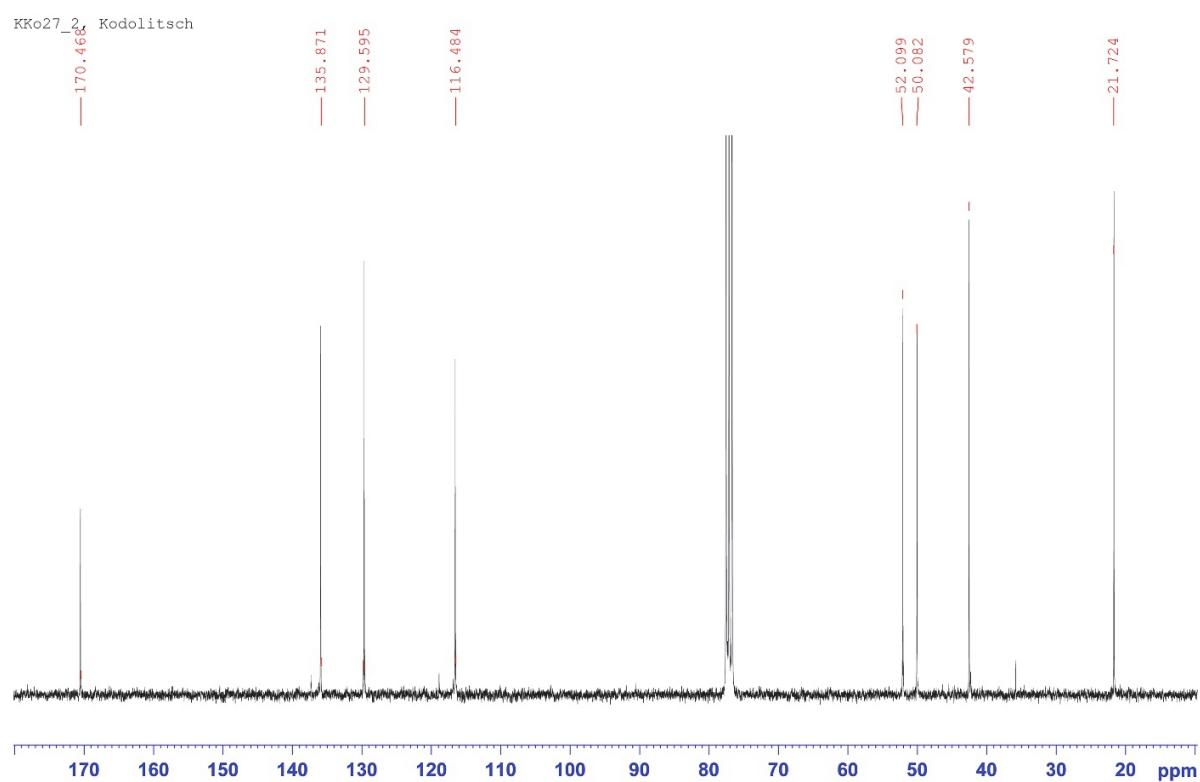
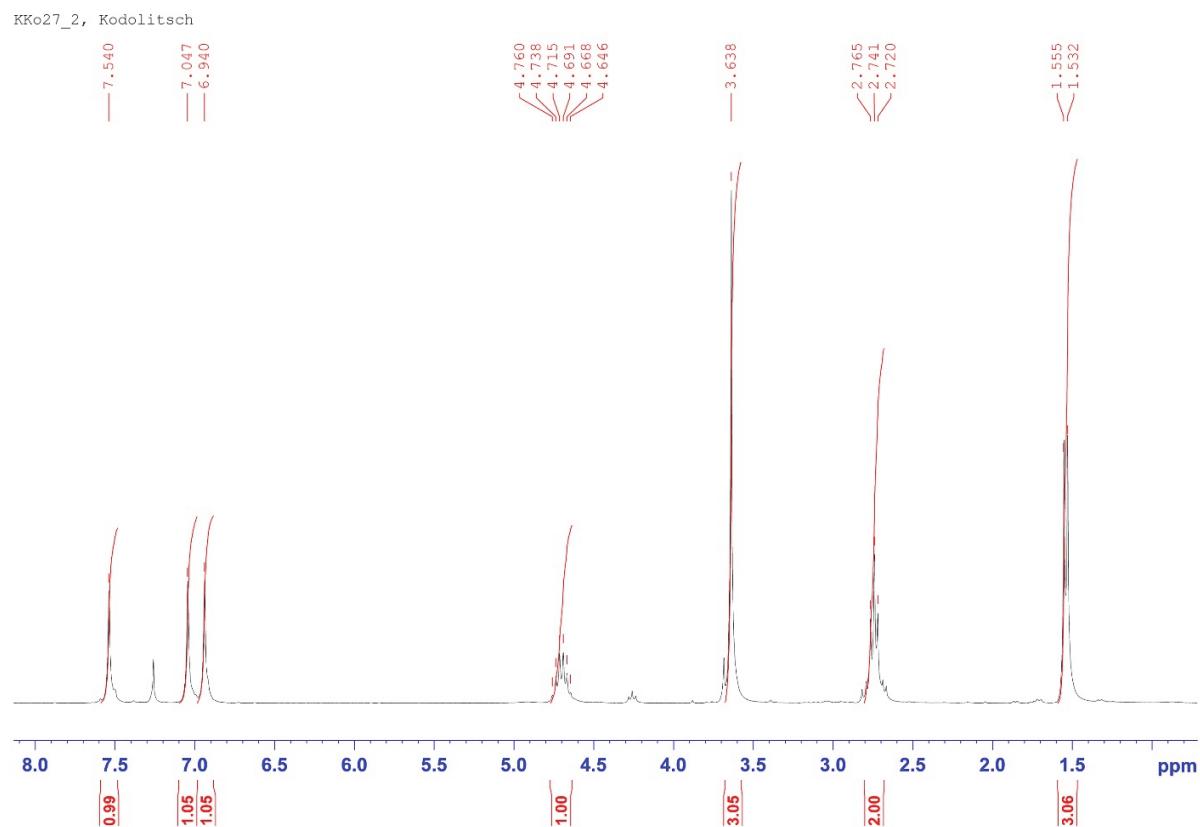


Fig S3. ^1H - and $^{13}\{\text{H}\}$ -NMR spectra of **1C**

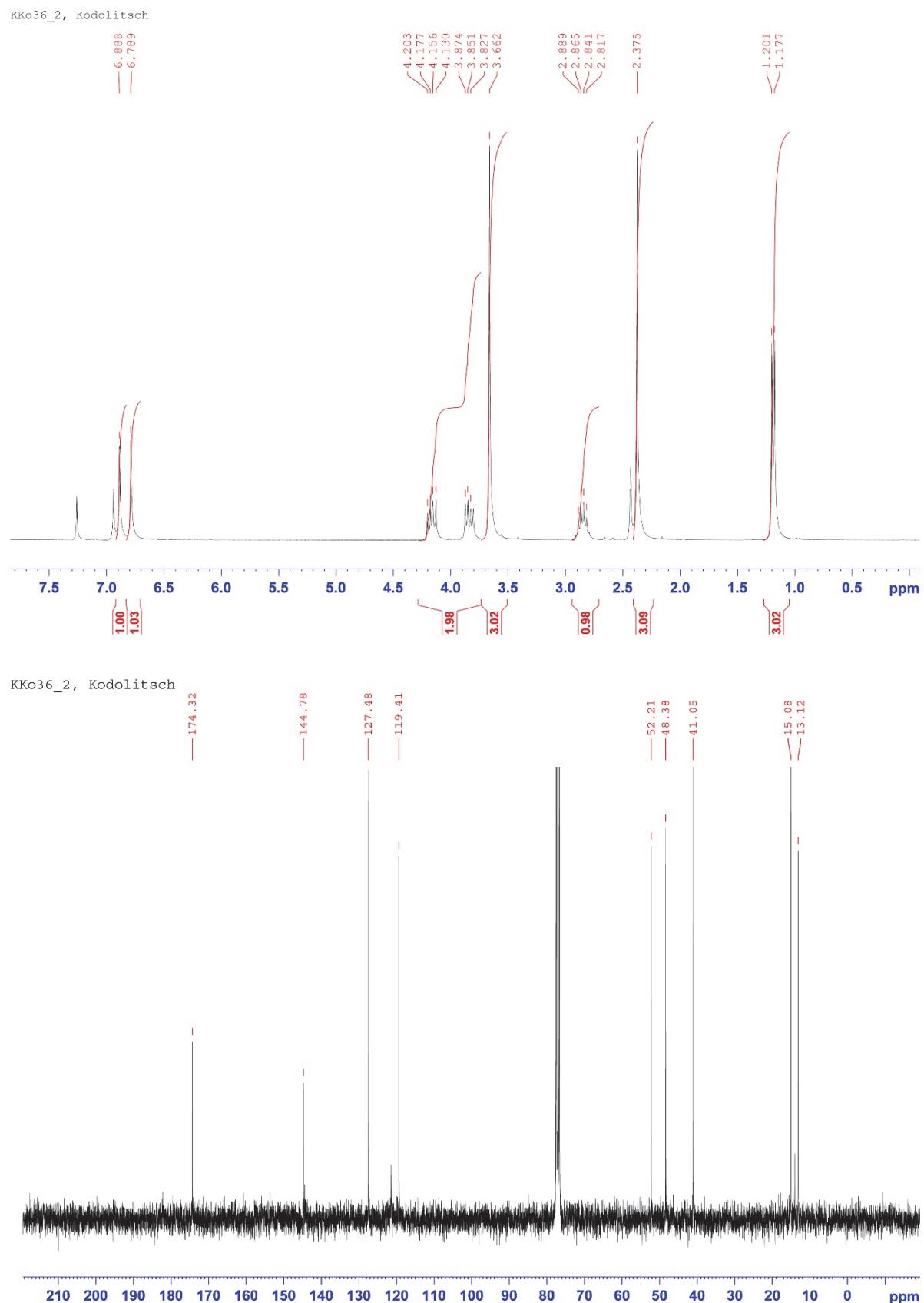


Fig S4. ^1H - and $^{13}\{\text{H}\}$ -NMR spectra of **1D**

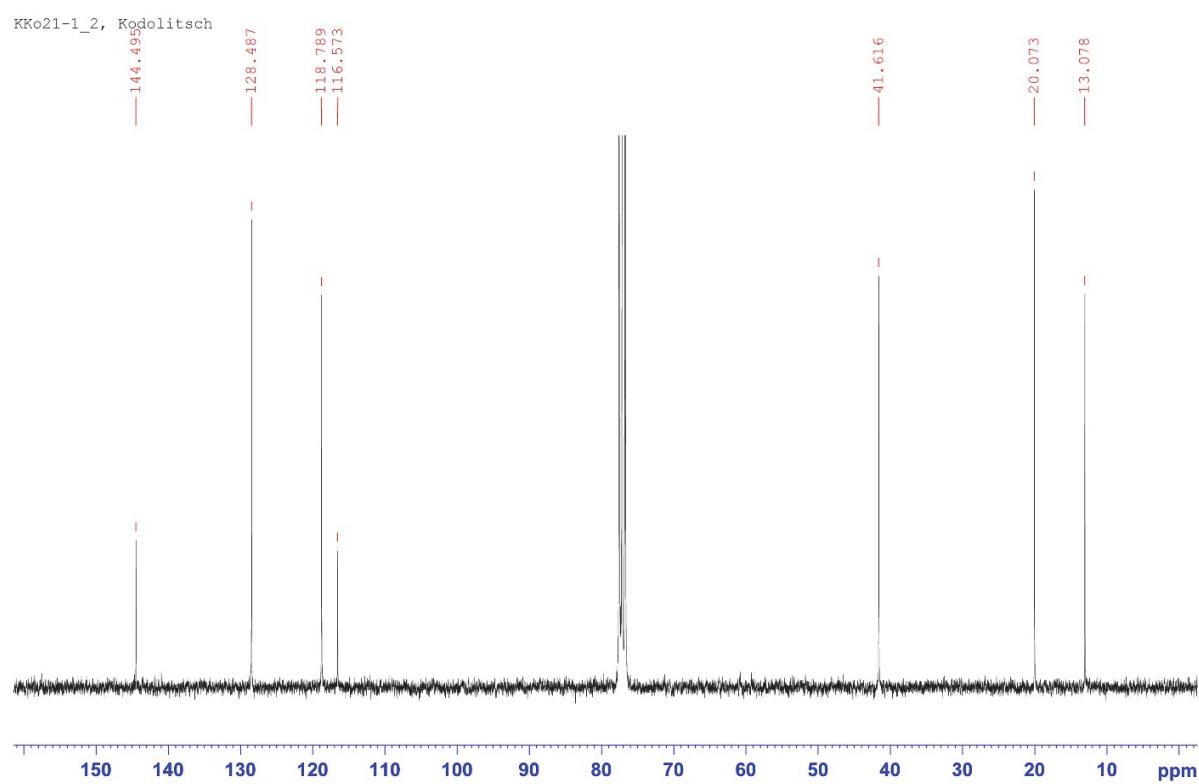
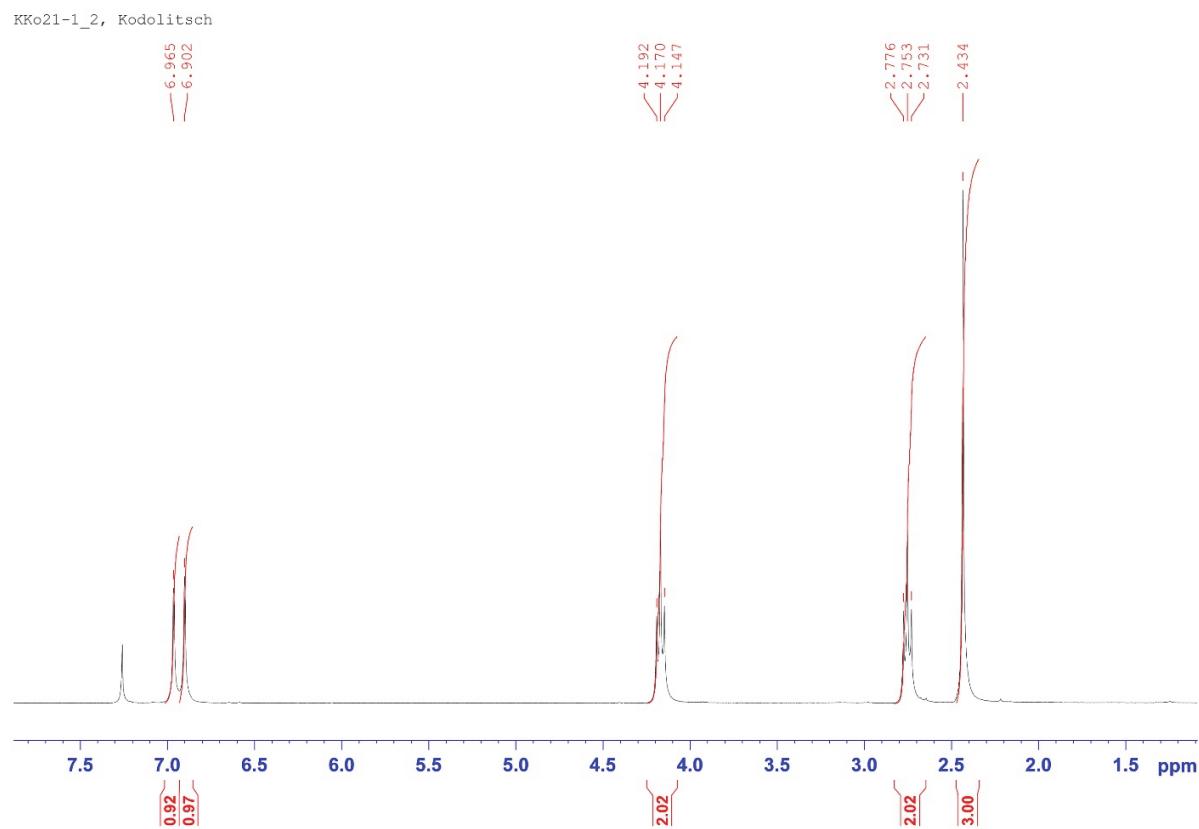
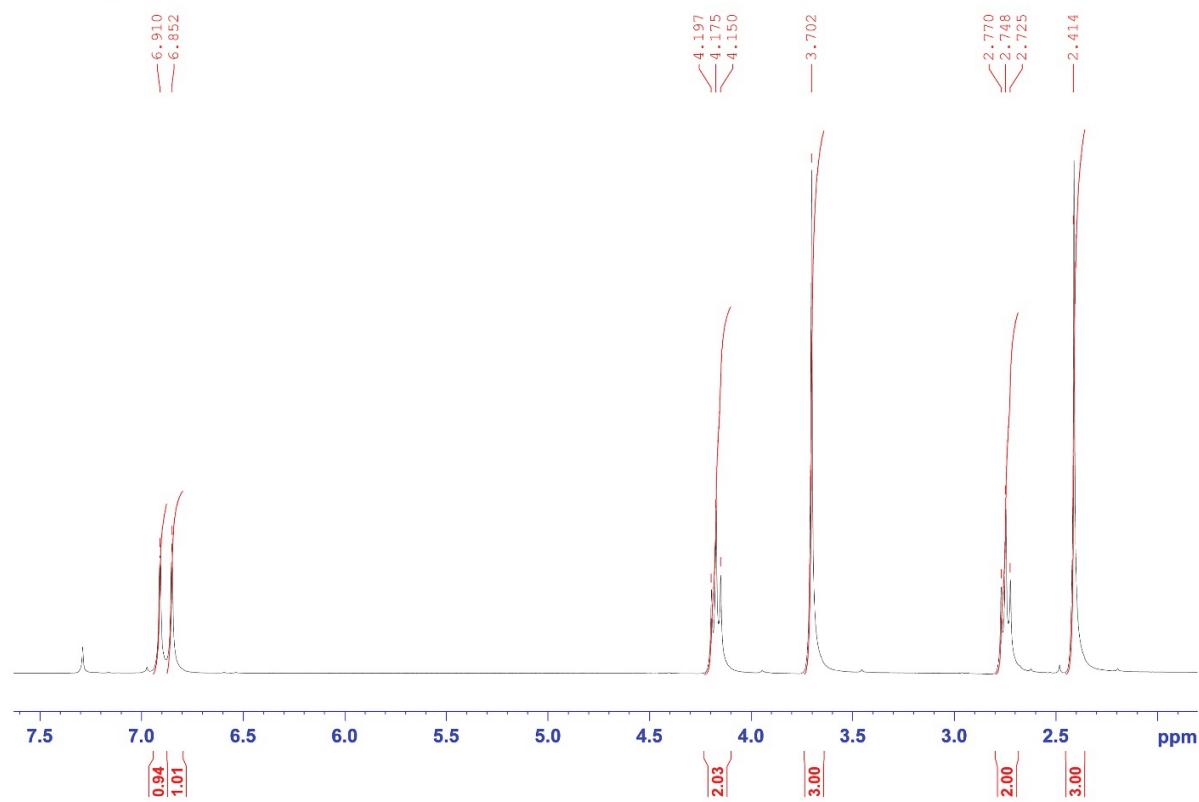


Fig S5. ^1H - and $^{13}\{\text{H}\}$ -NMR spectra of **2A**

Kodolitsch, KK005



Kodolitsch, KK005

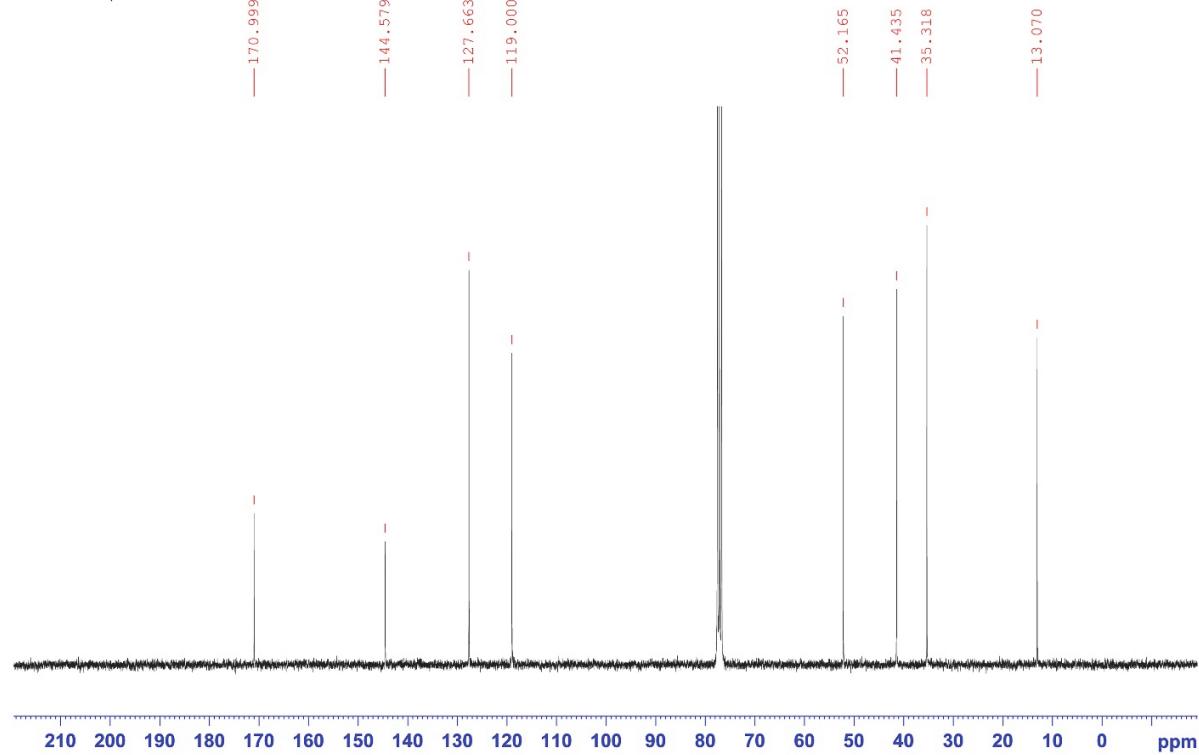


Fig S6. ¹H- and ¹³C-¹H)-NMR spectra of **2B**

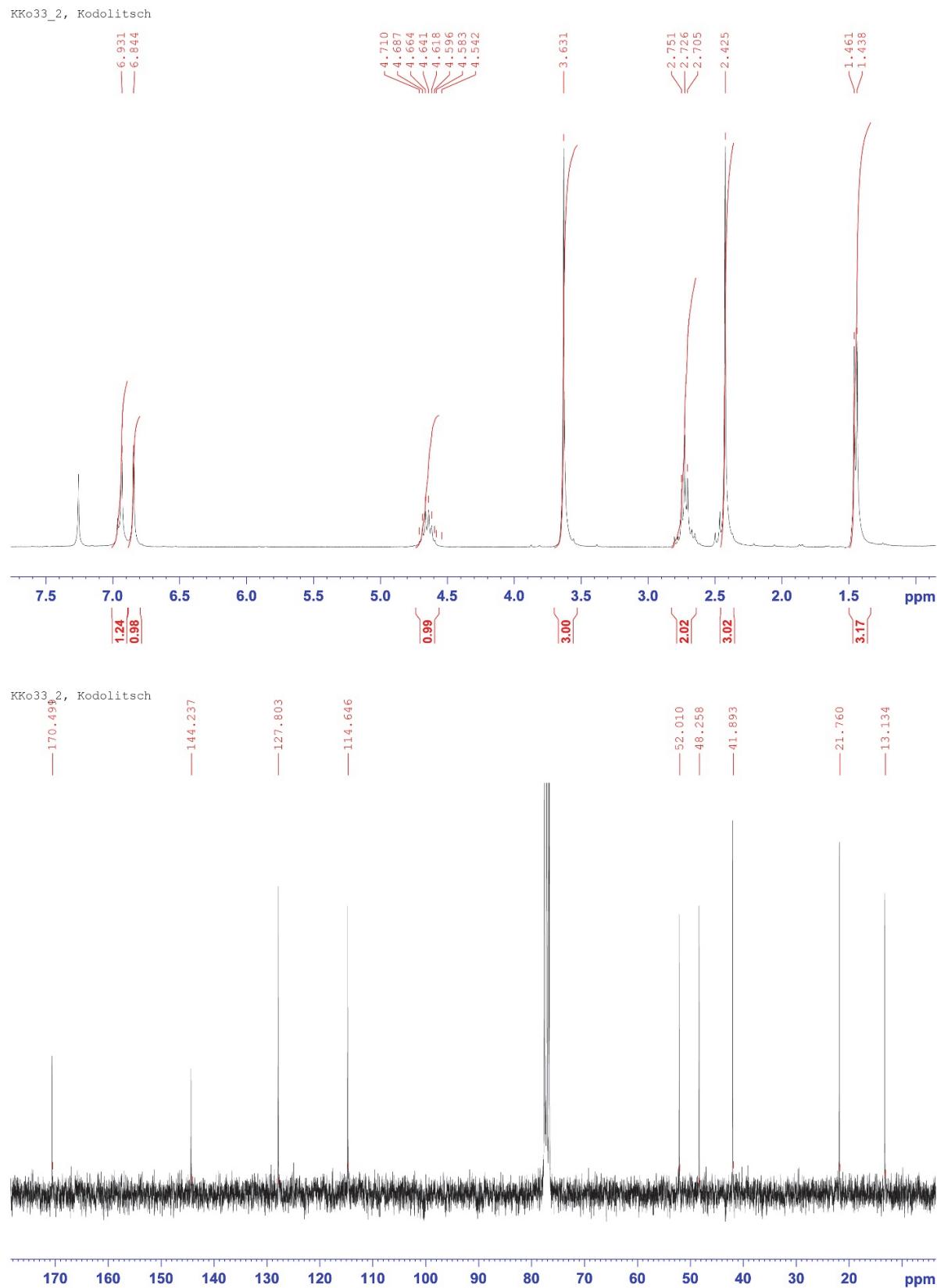
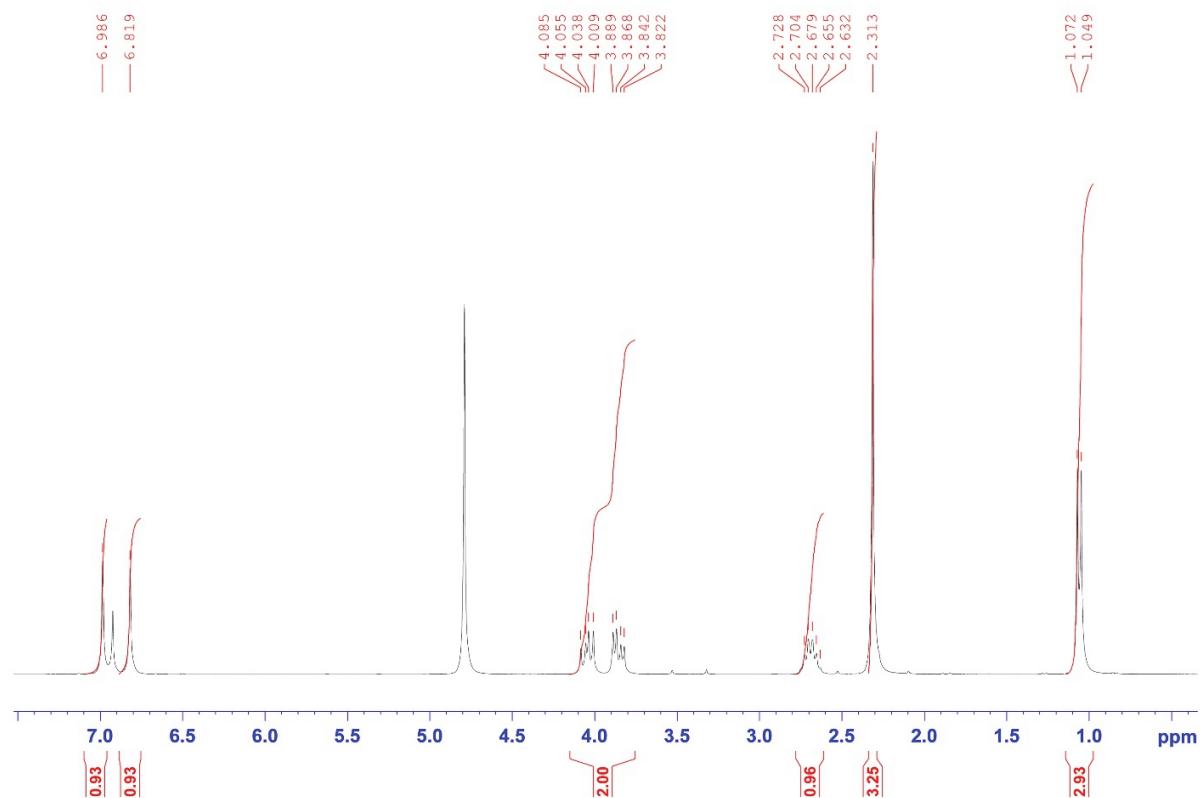


Fig S7. ^1H - and $^{13}\{\text{H}\}$ -NMR spectra of **2C**

Kodolitsch, KK046



Kodolitsch, KK046

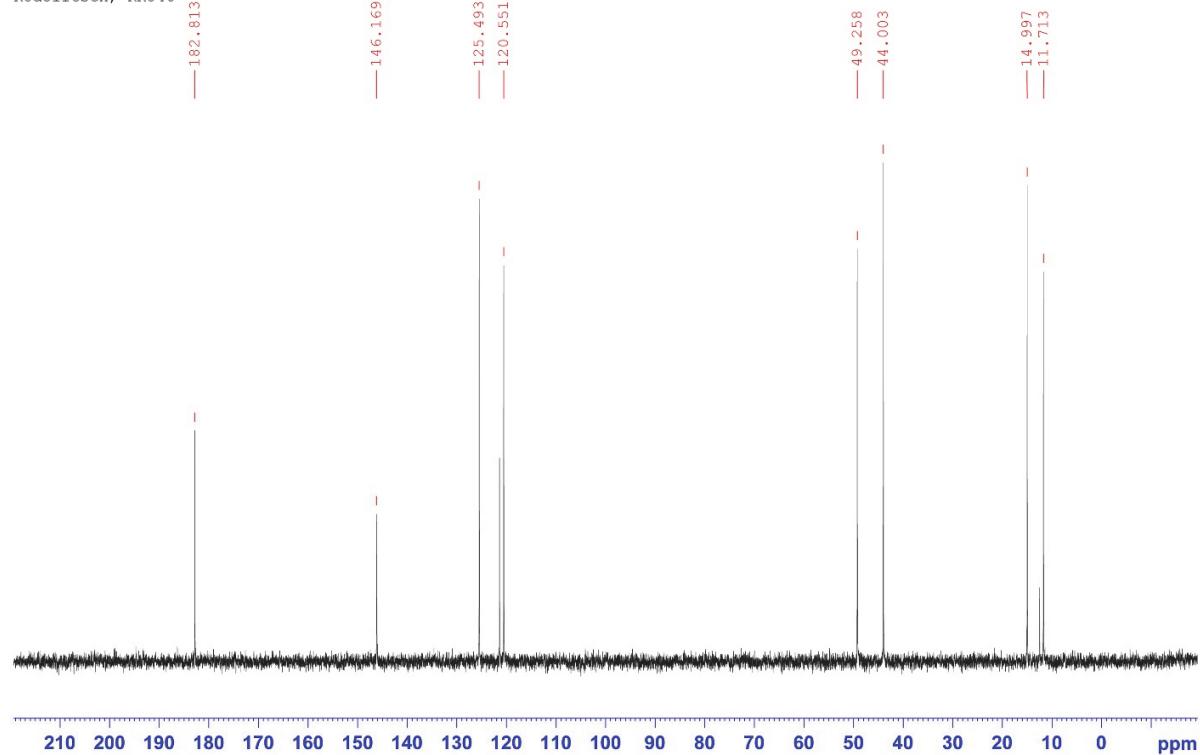
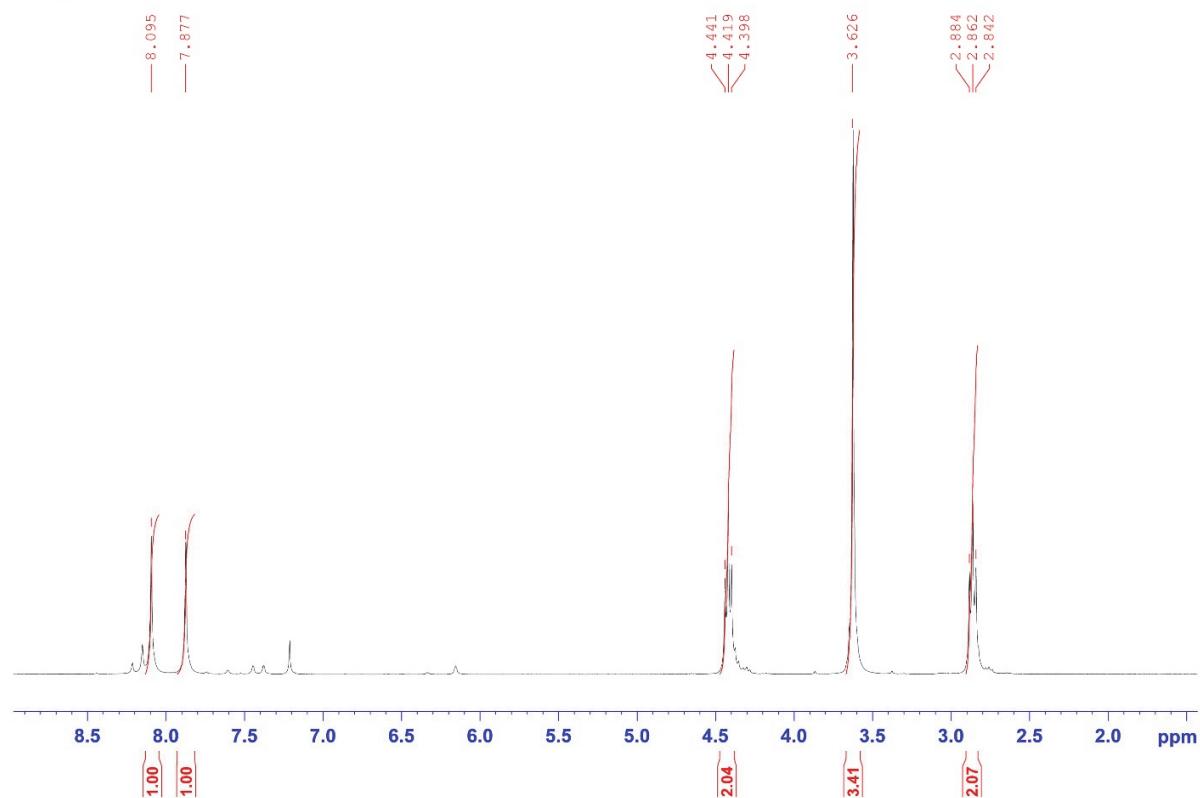


Fig S8. ^1H - and $^{13}\{\text{H}\}$ -NMR spectra of **2D**

KKo09, Kodolitsch



KKo09_2, Kodolitsch

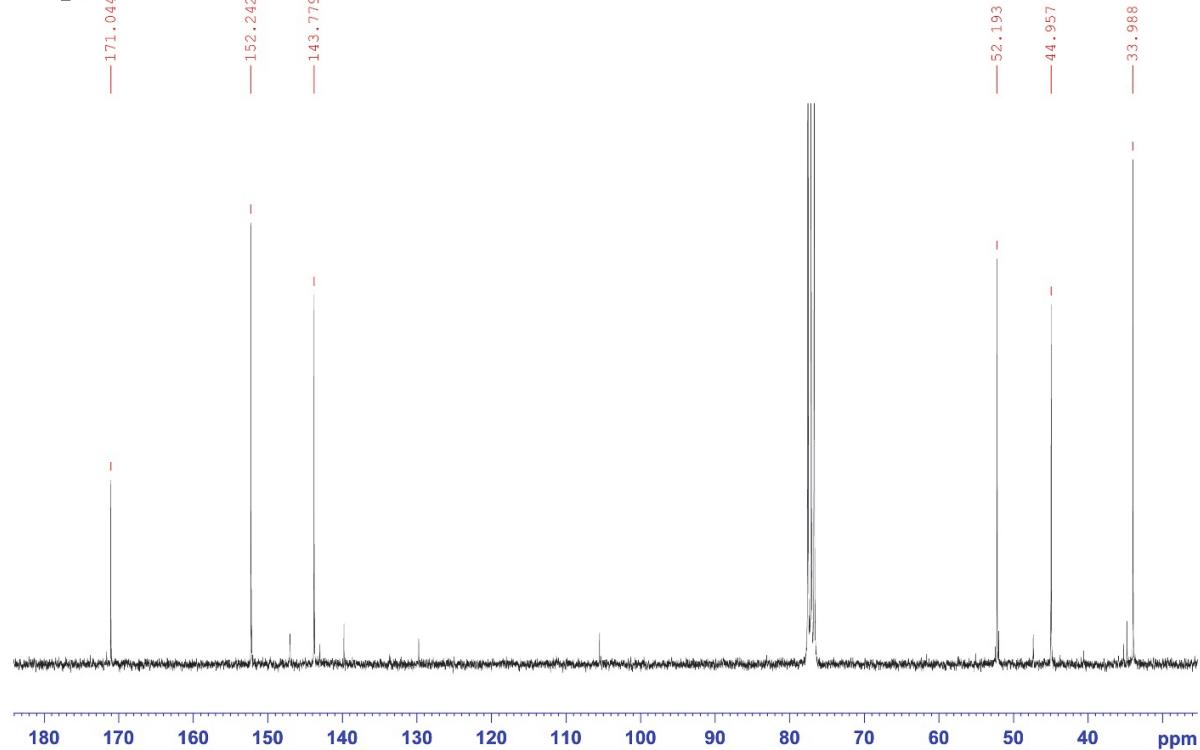


Fig S9. ^1H - and $^{13}\{\text{H}\}$ -NMR spectra of **3B**

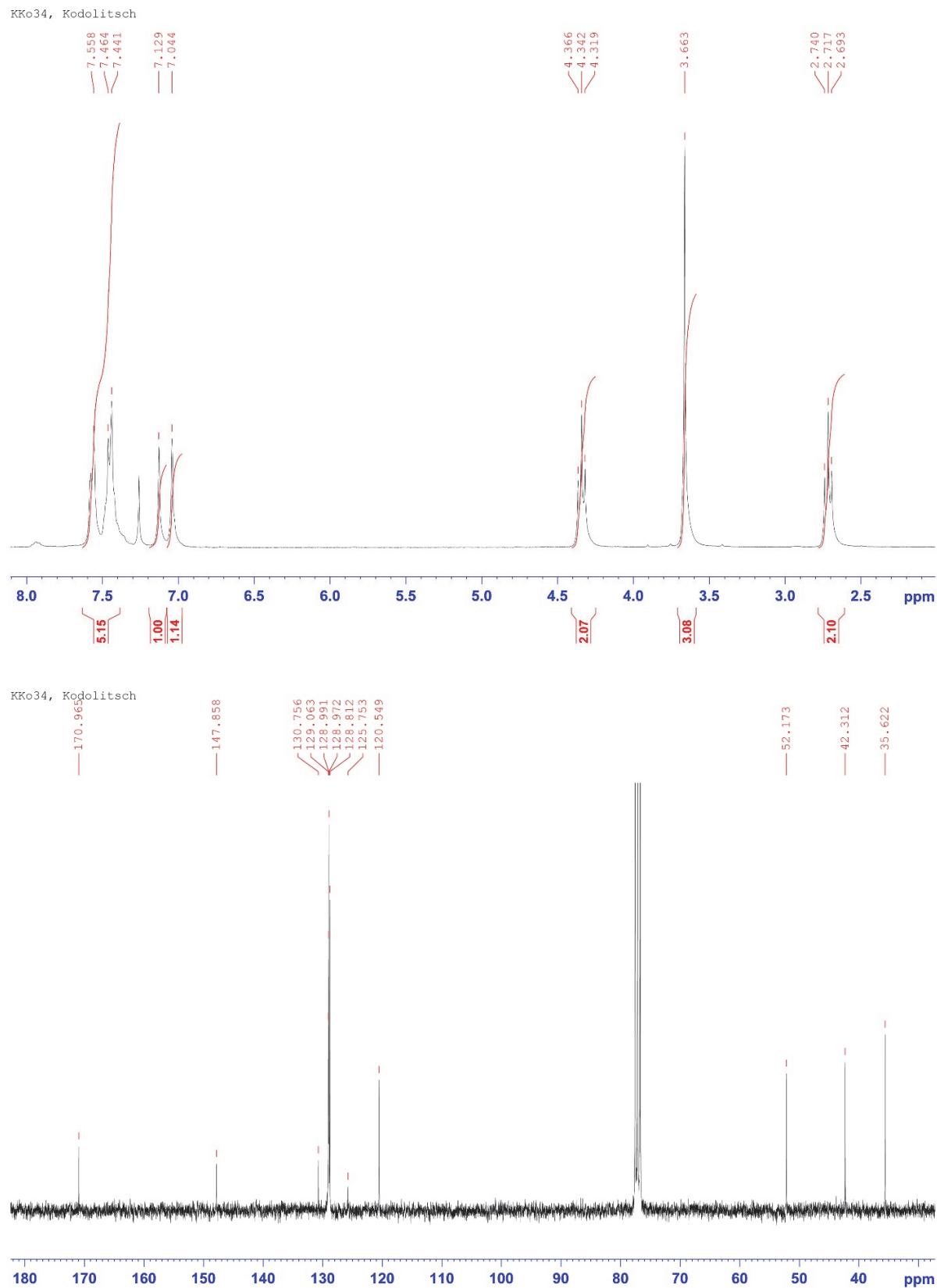


Fig S10. ^1H - and $^{13}\{\text{H}\}$ -NMR spectra of **4B**

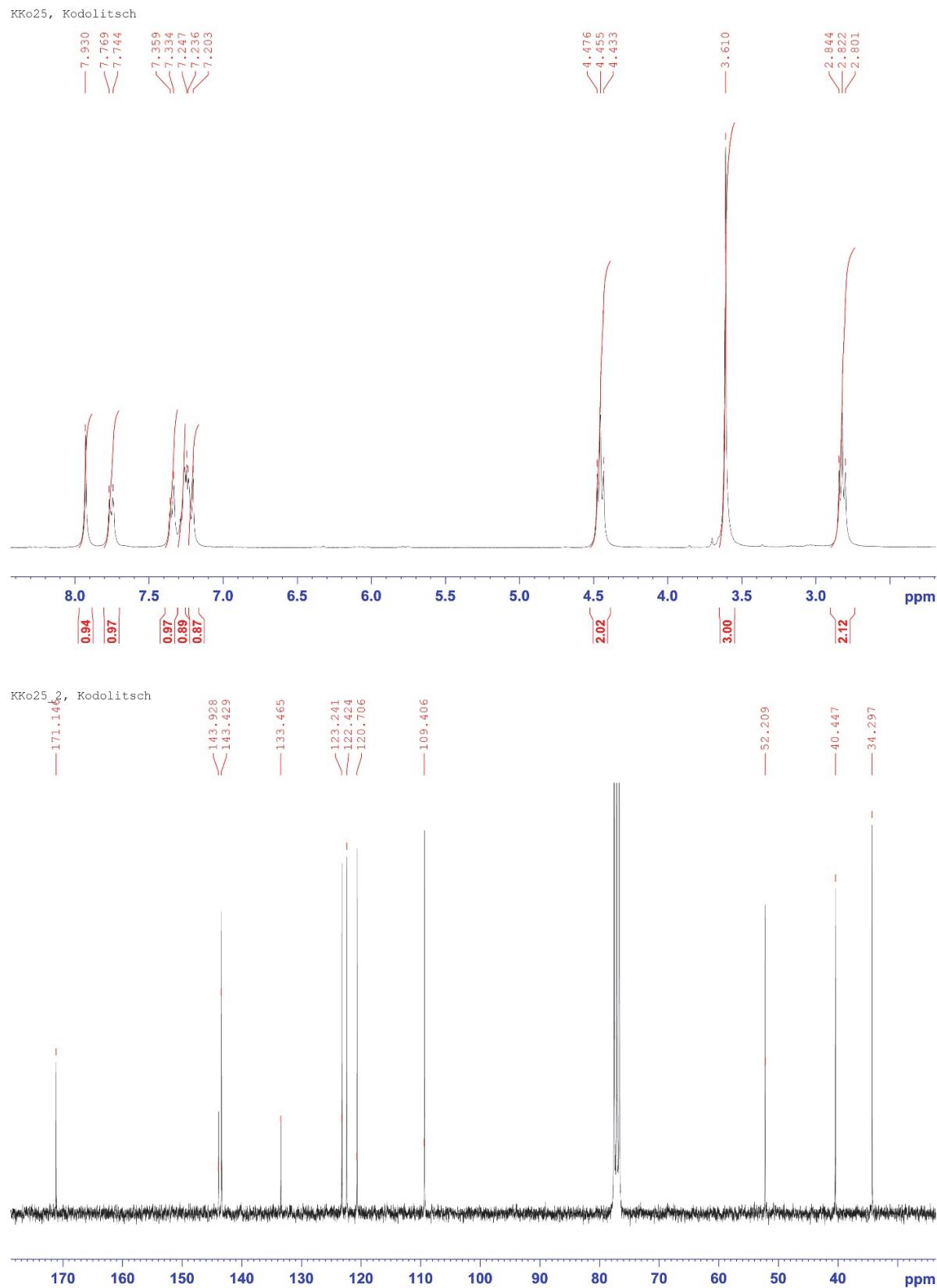


Fig S11. ^1H - and $^{13}\{\text{H}\}$ -NMR spectra of **5B**

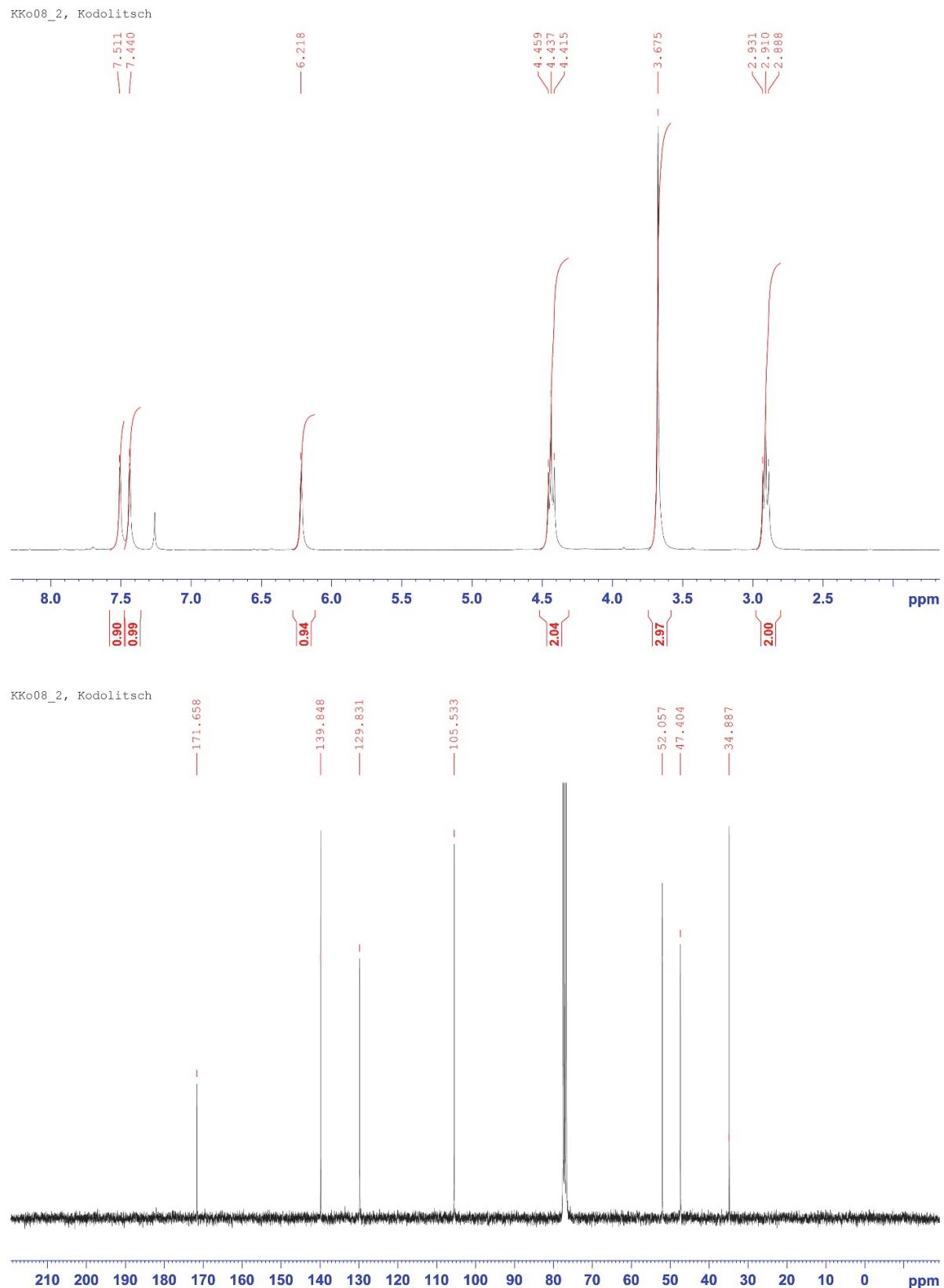


Fig S12. ^1H - and $^{13}\{\text{H}\}$ -NMR spectra of **6B**

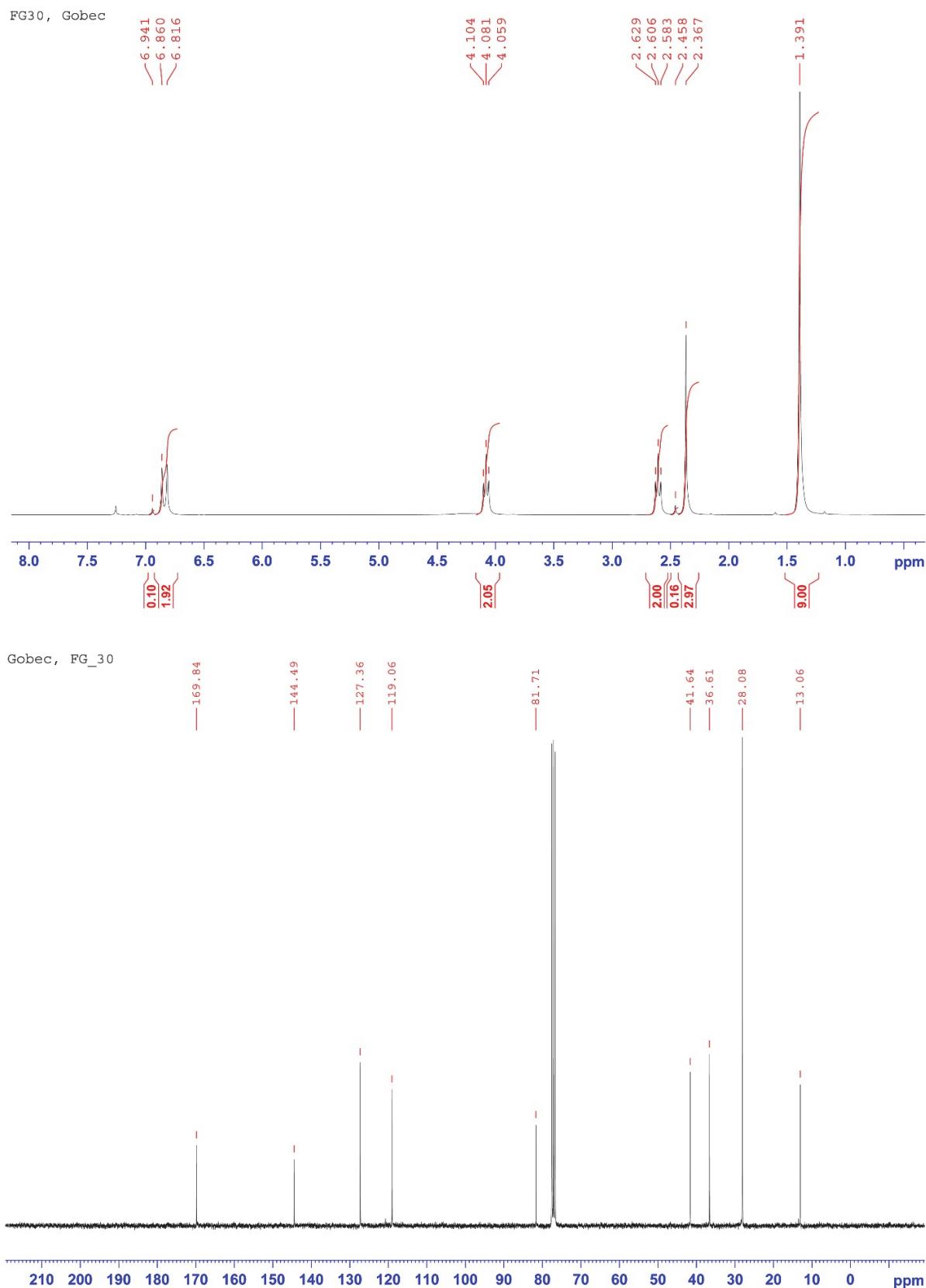


Fig S13. ^1H - and $^{13}\{\text{H}\}$ -NMR spectra of **2E**

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