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SUPPLEMENTARY MATERIAL TO

Syntheses and computational analyses of selected macrolide derivatives derived from clarithromycin A

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a)

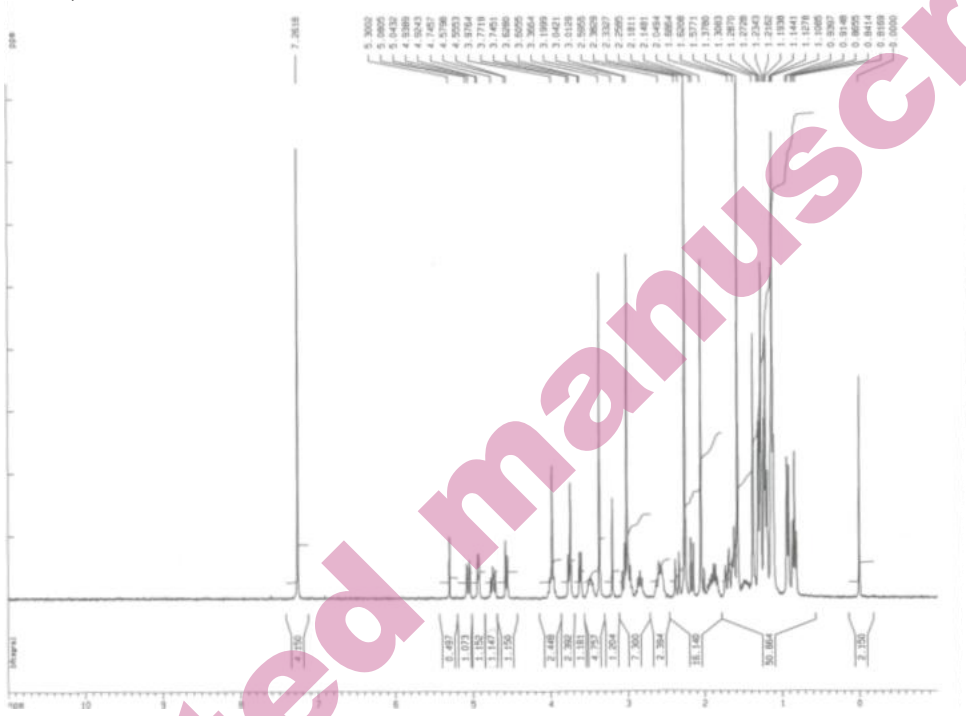

Figure S-1. a) The ^{1}H NMR spectrum of 2′-*O*-acetyl-clarithromycin A (**1**).

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b)

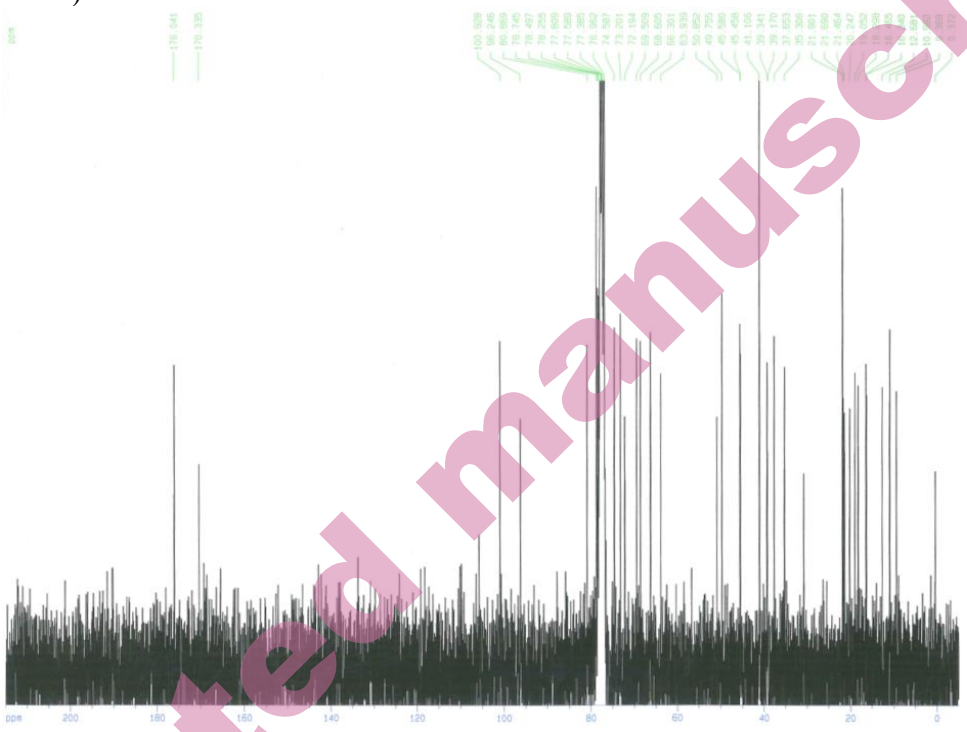

Figure S-1. b) ^{13}C NMR spectrum of 2′-*O*-acetyl-clarithromycin A (**1**).

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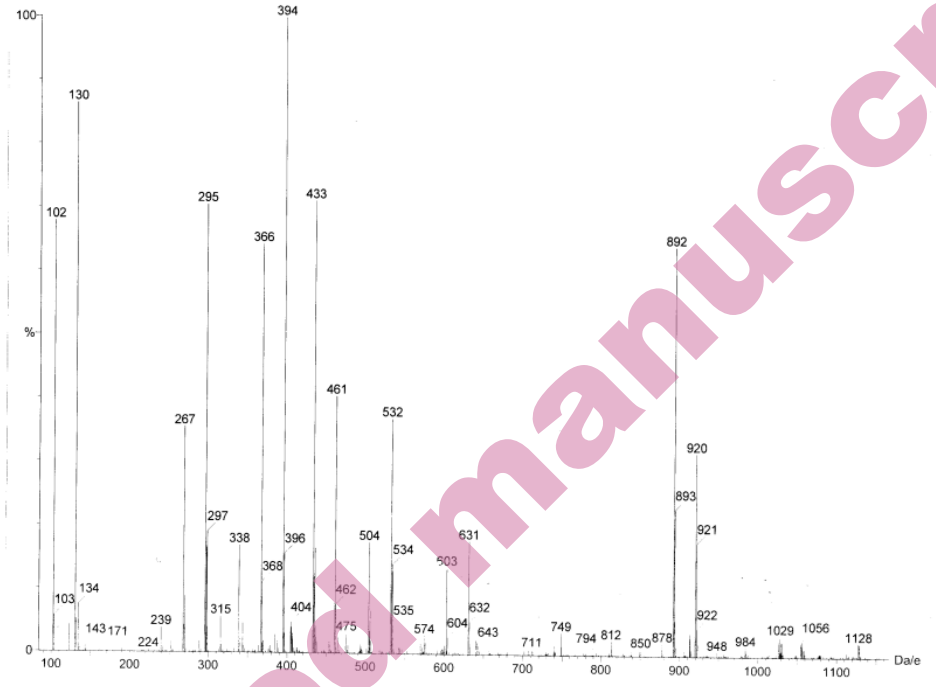

Figure S-2. Mass spectrum (ES+) of 2′-*O*-acetyl-4"-*O*-(2-cyanoethyldiisopropylphosphoramidite)-clarithromycin A (**2**).

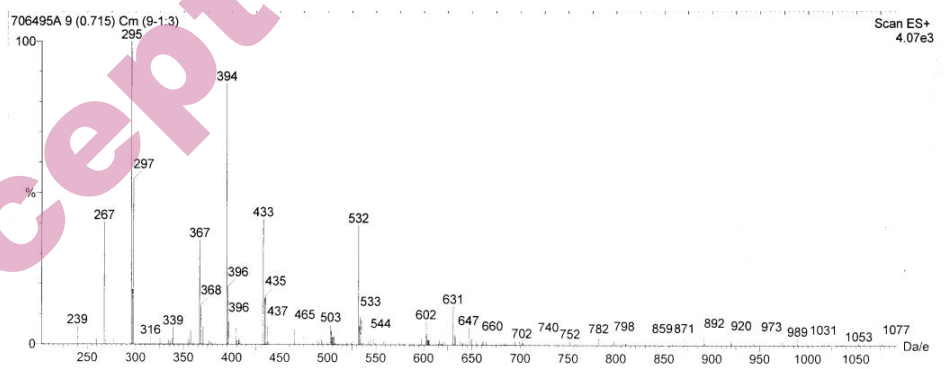

Figure S-3. Mass spectrum (ES+) of 2′-*O*-acetyl-4′′-*O*-(2-cyanoethyldiisopropylphosphoramidite)-clarithromycin A (**2**).

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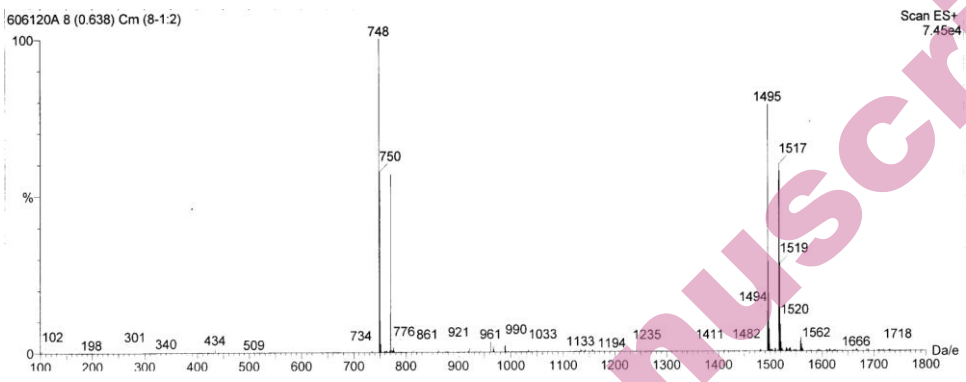

Figure S-4. Mass spectrum (ES+) of 4´´-*O*-phosphonyl-clarithromycin A (**3**).

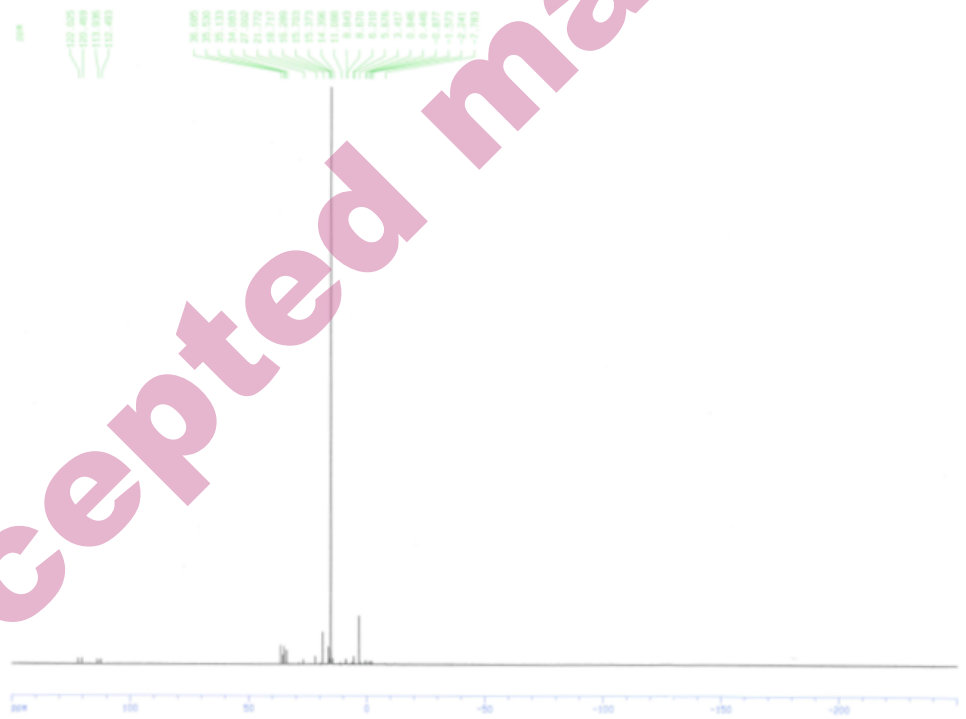

Figure S-5. ^{31}P NMR spectrum of the mixture 2-cyanoethyl *N,N*-diisopropylchlorophosphoramidite, *N*-ethyldiisopropylamine and water.

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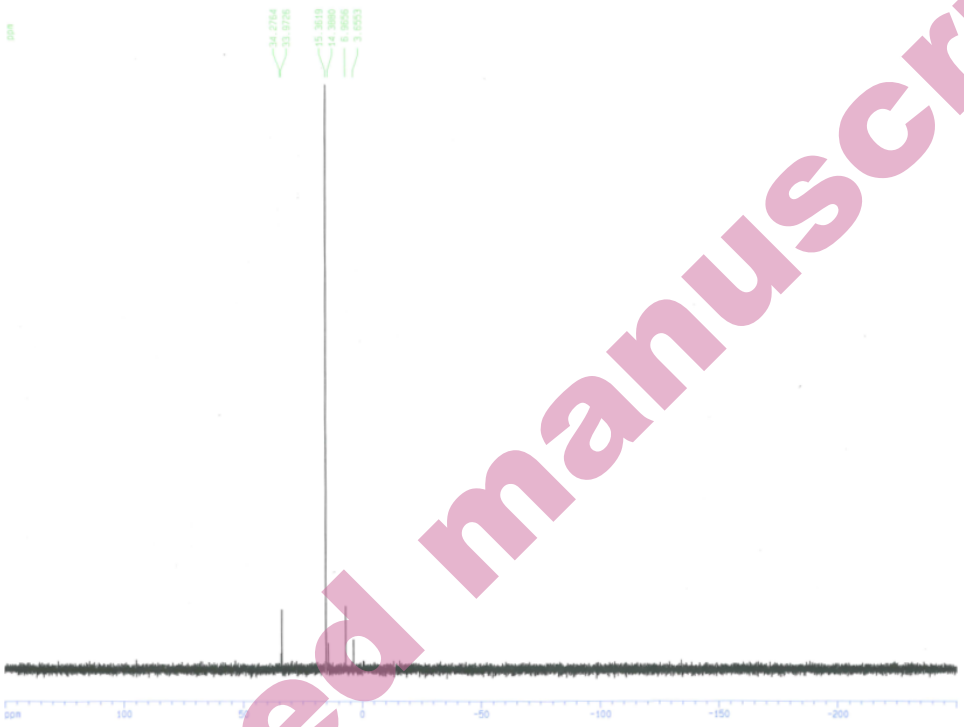

Figure S-6. ^{31}P NMR spectrum of the reaction mixture after 1 day when 2-cyanoethyl *N,N*-diisopropylchlorophosphoramidite was in 5 fold excess.

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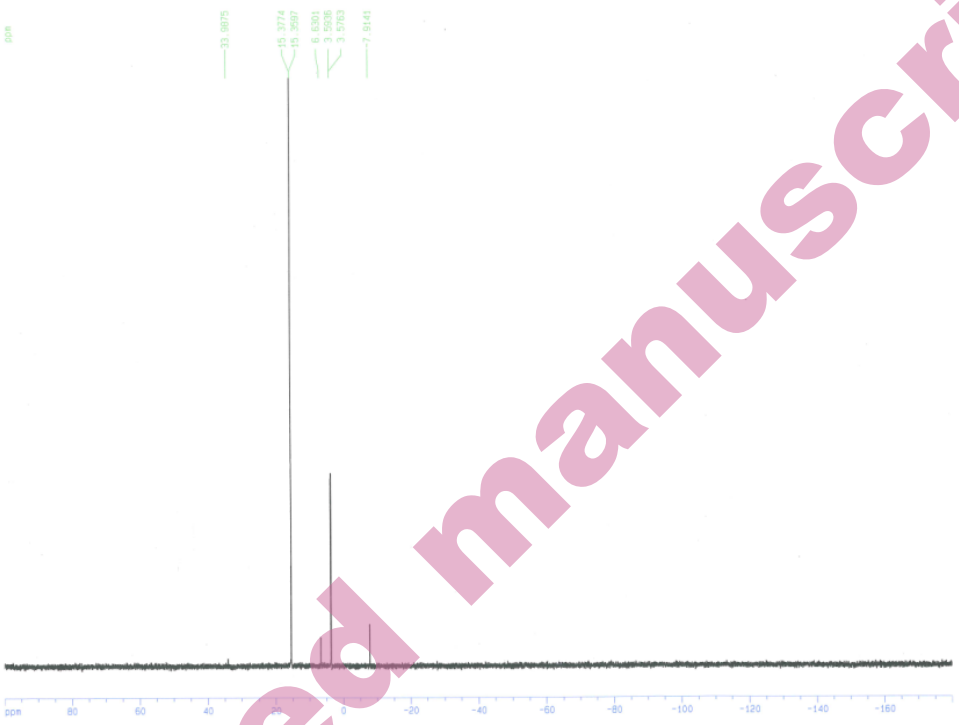

Figure S-7. ^{31}P NMR spectrum of a reaction mixture with DMAP (4-dimethylaminopyridine) as a catalyst.

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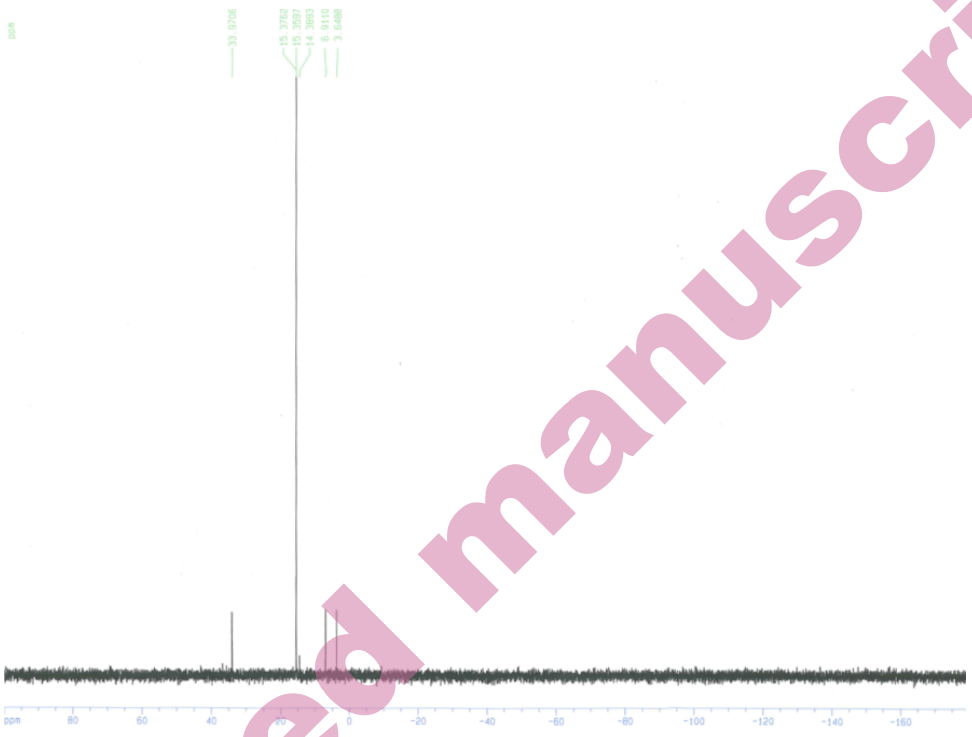

Figure S-8. ^{31}P NMR spectrum of the reaction mixture on heating at 40 ºC under reflux.

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