

Thermal stability and kinetics of degradation of moxonidine as pure ingredient vs. pharmaceutical formulation

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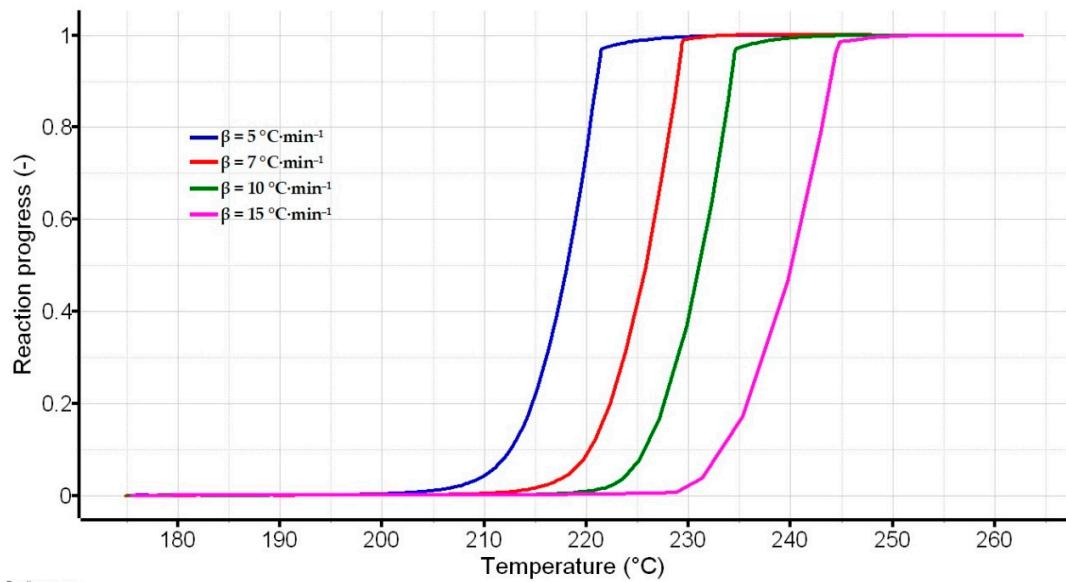
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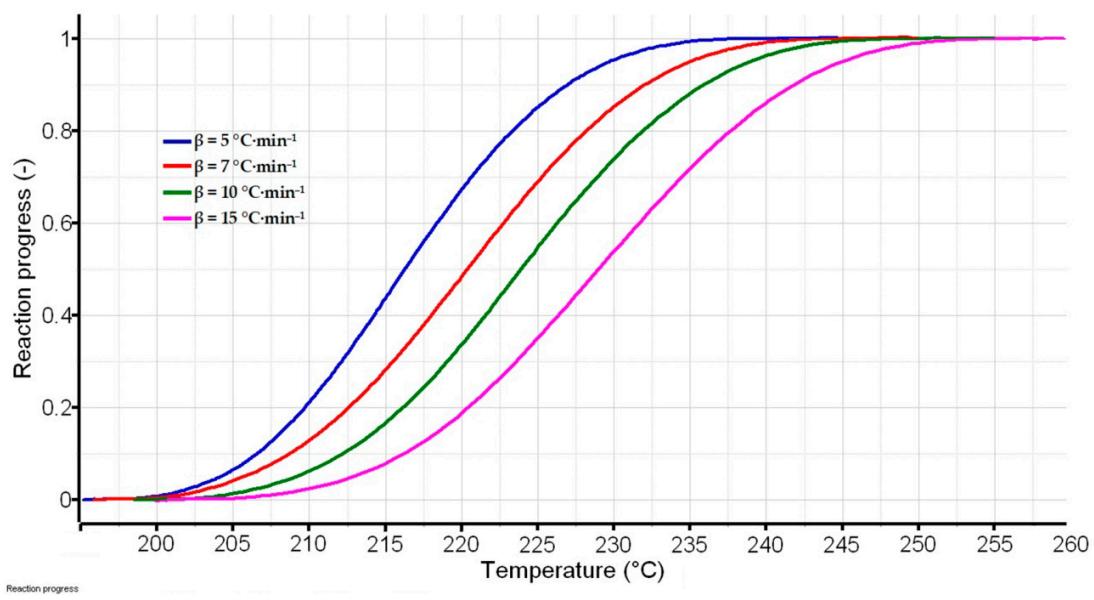
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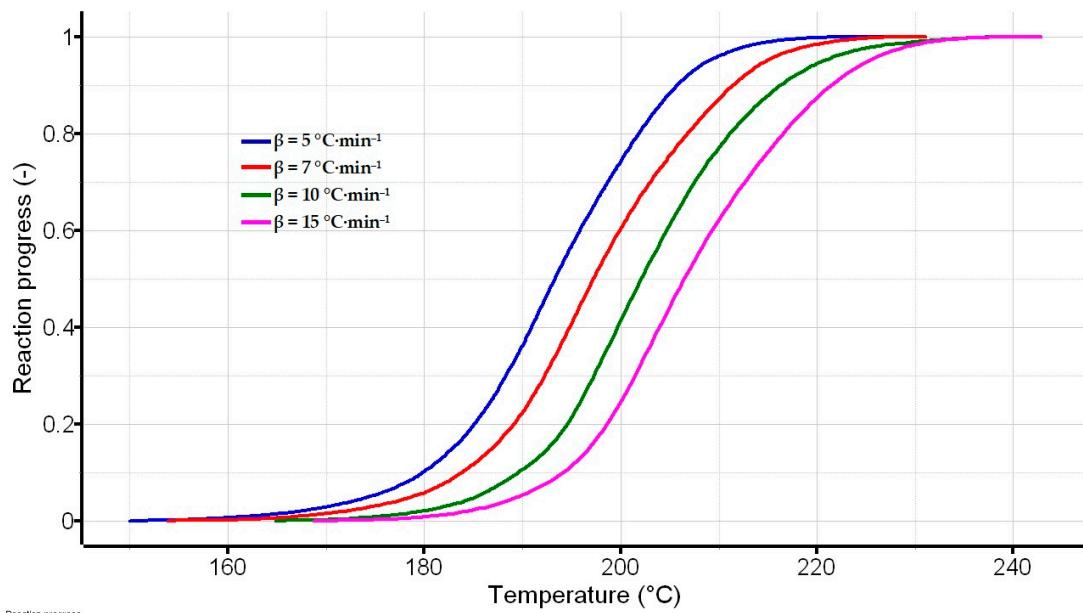
Supplementary materials



(a)

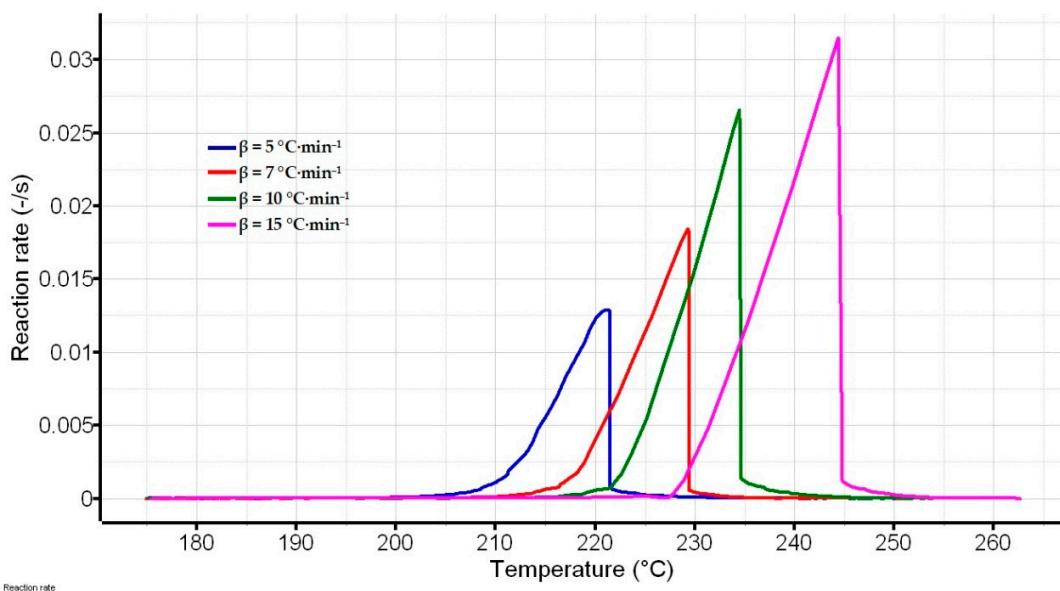


(b)

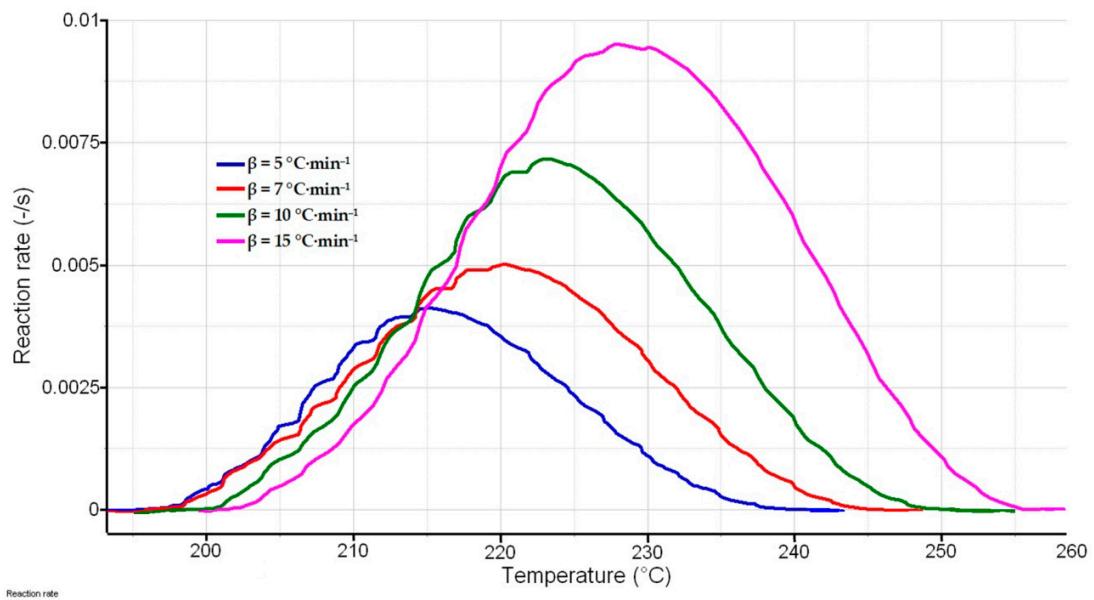


(c)

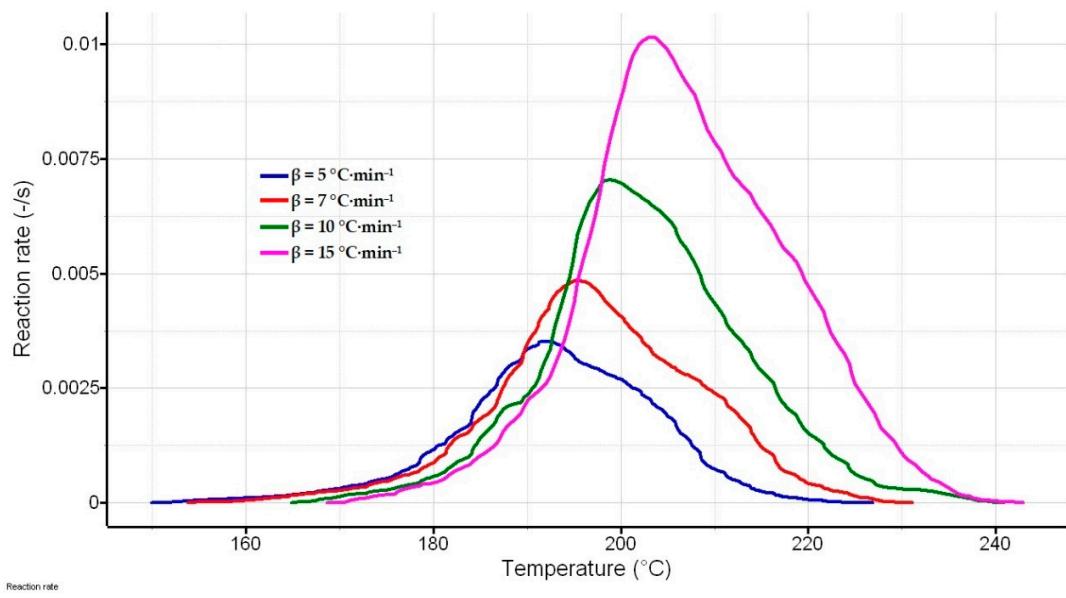
Figure S1. The reaction progress vs. temperature at selected heating rates for: (a) MOX; (b) MOXTAB; and (c) MOXMIX.



(a)

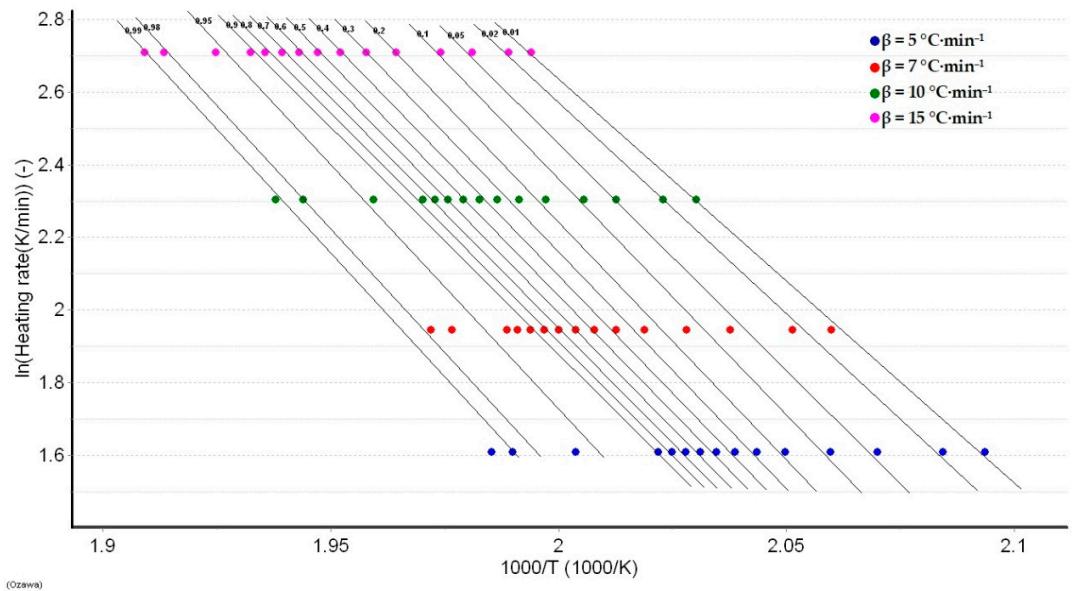


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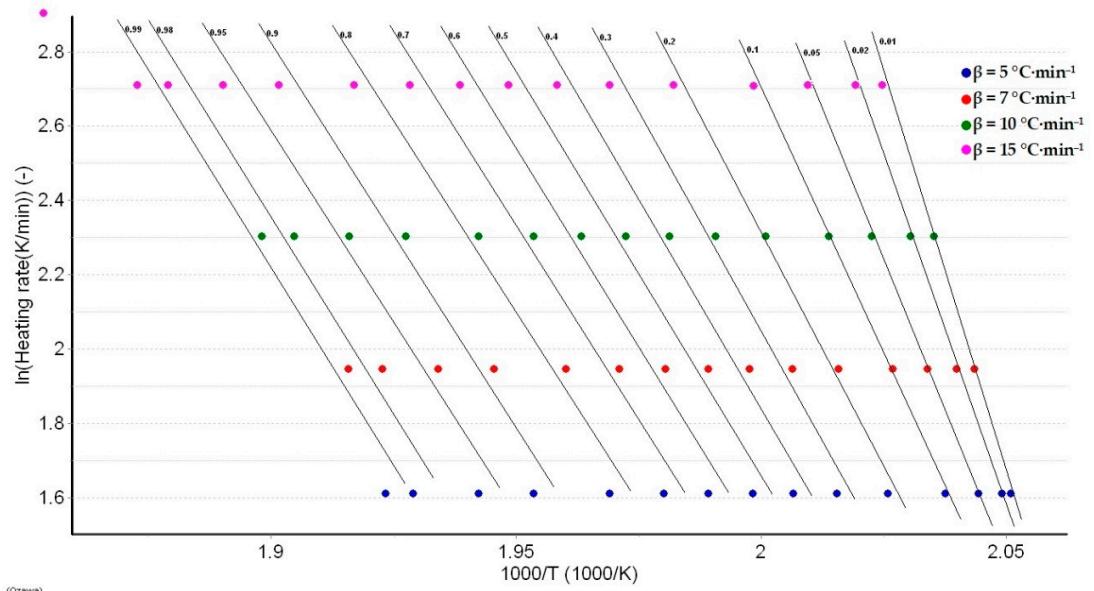


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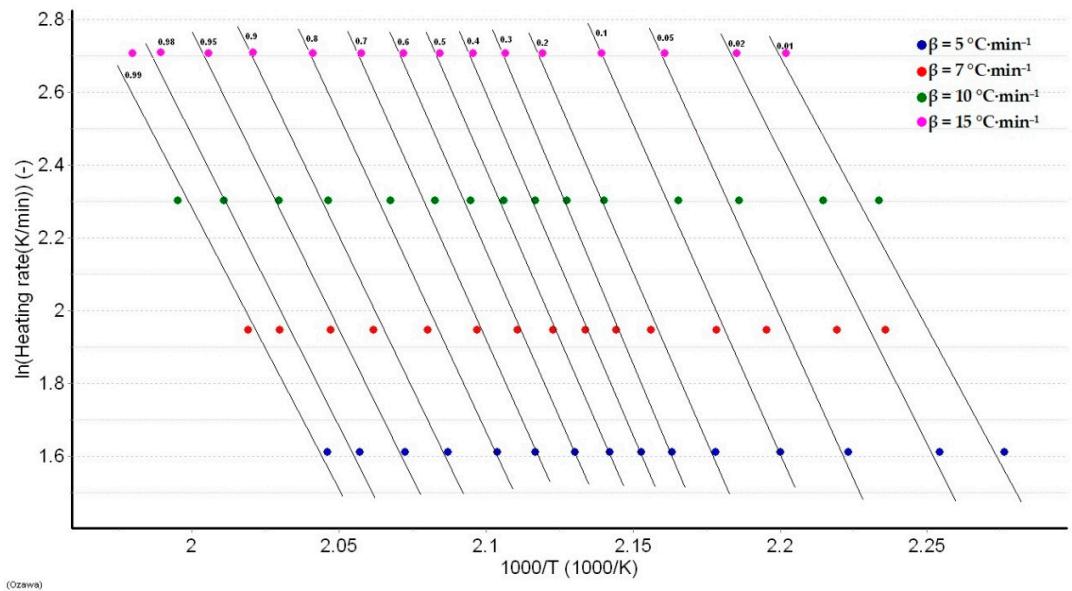
Figure S2. The reaction rate vs. temperature at selected heating rates for: (a) MOX; (b) MOXTAB; and (c) MOXMIX.



(a)

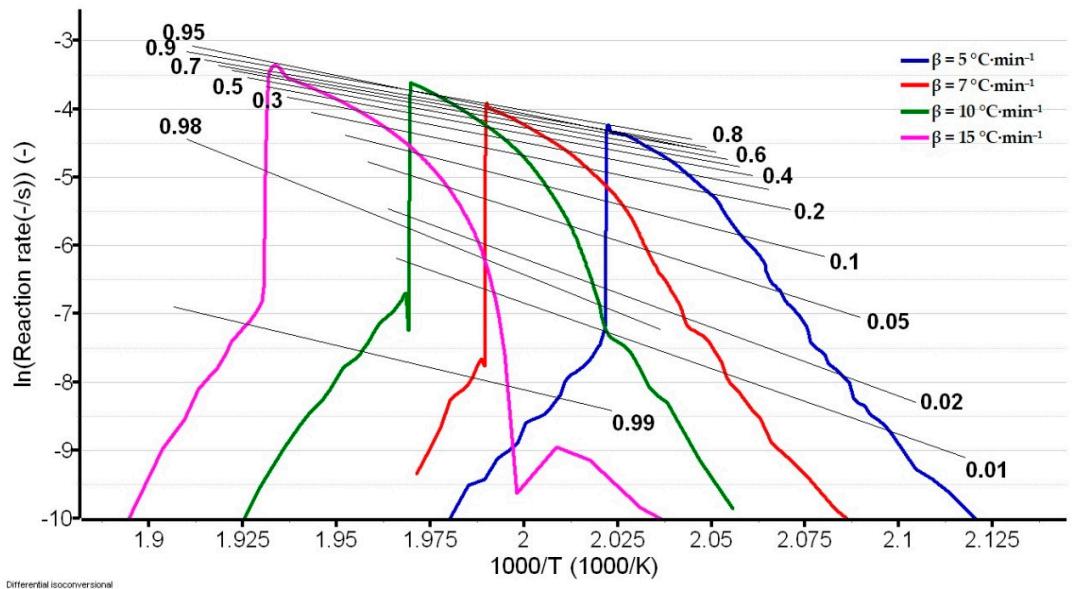


(b)

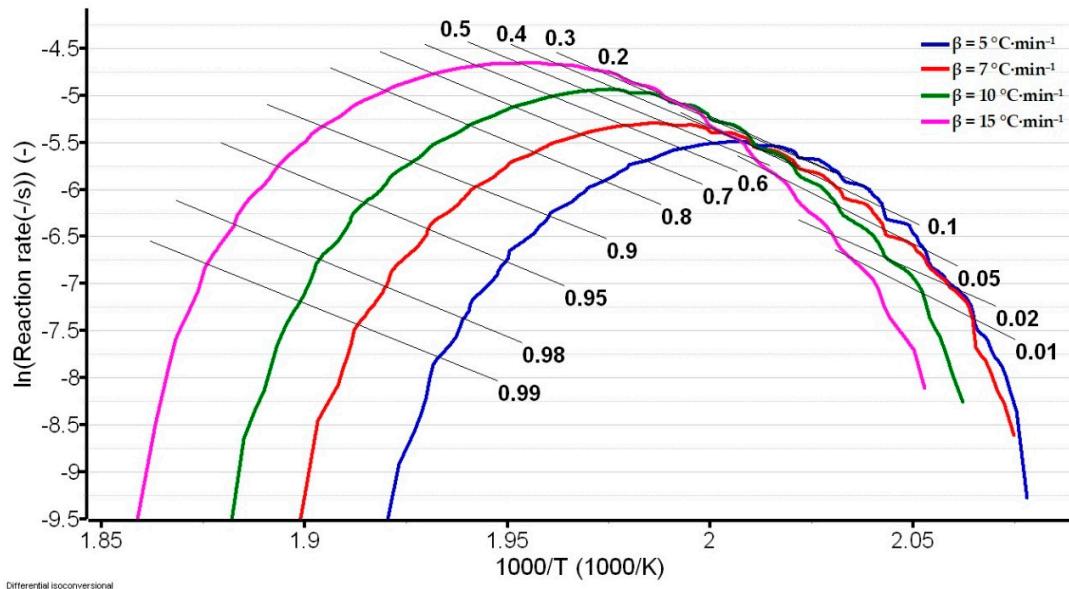


(c)

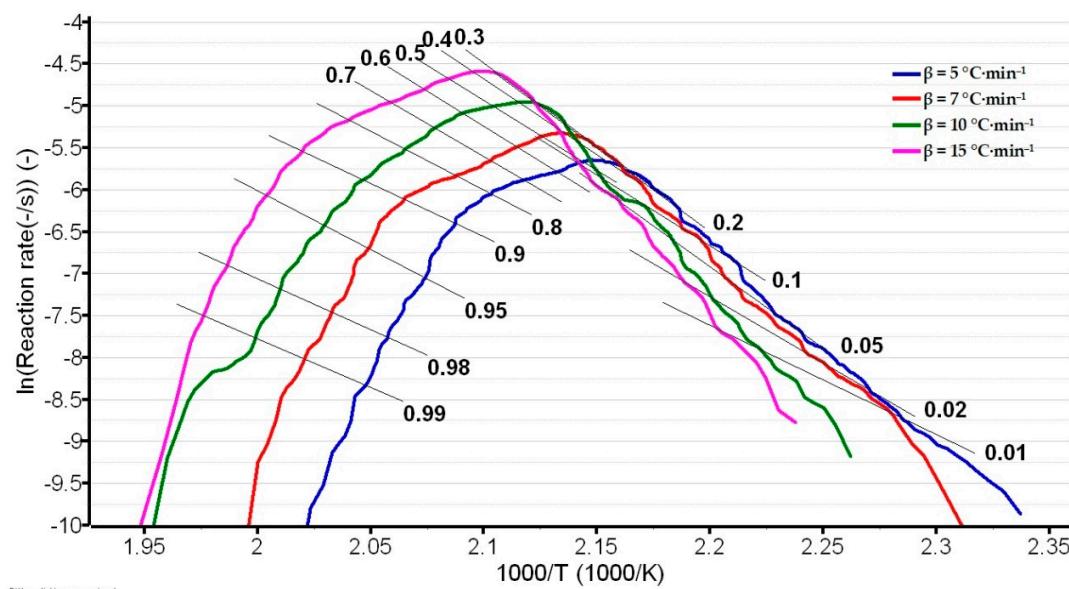
Figure S3. Linear plot of FWO integral isoconversional method of the analyzed decomposition process for: (a) MOX; (b) MOXTAB; and (c) MOXMIX.



(a)

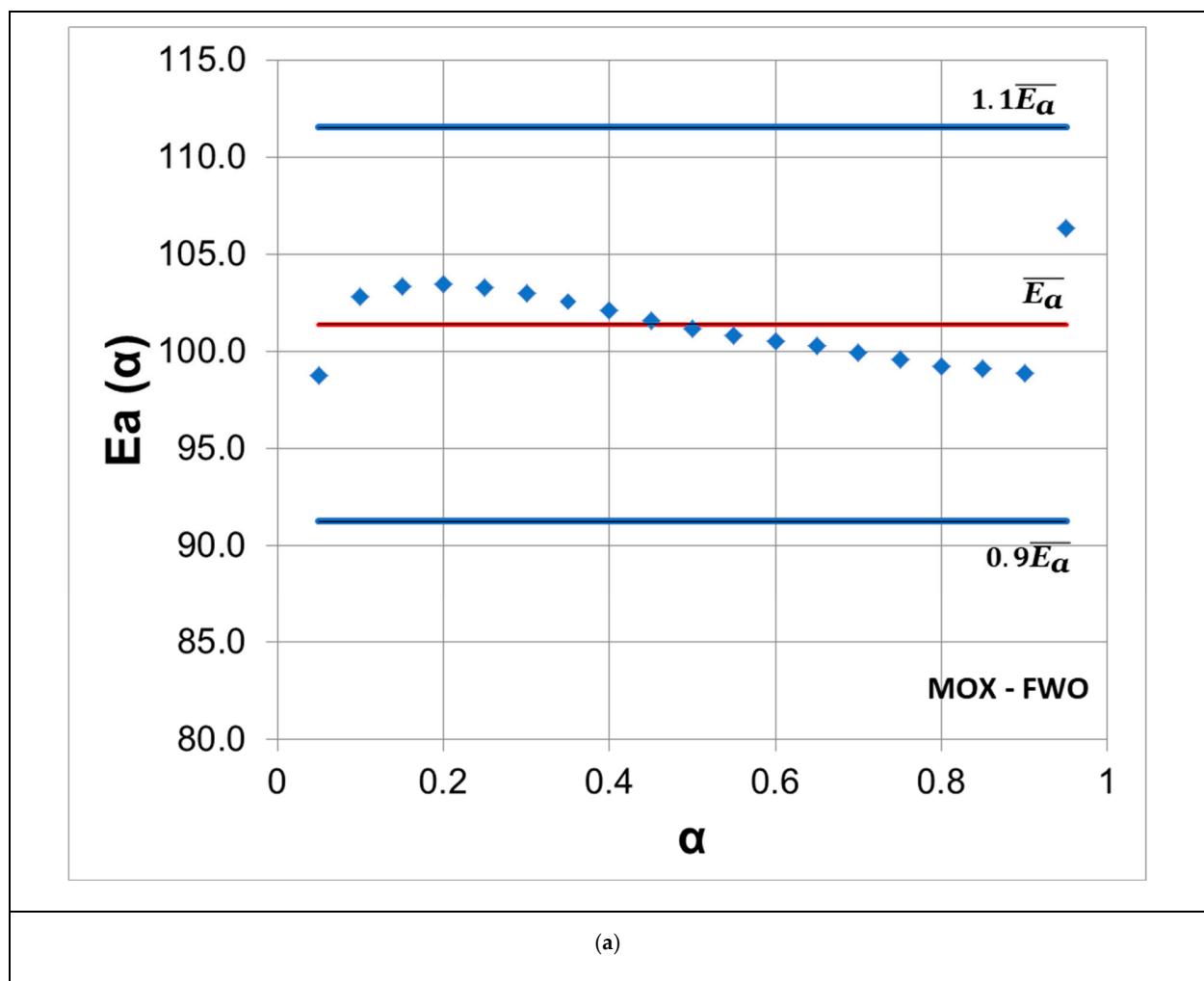


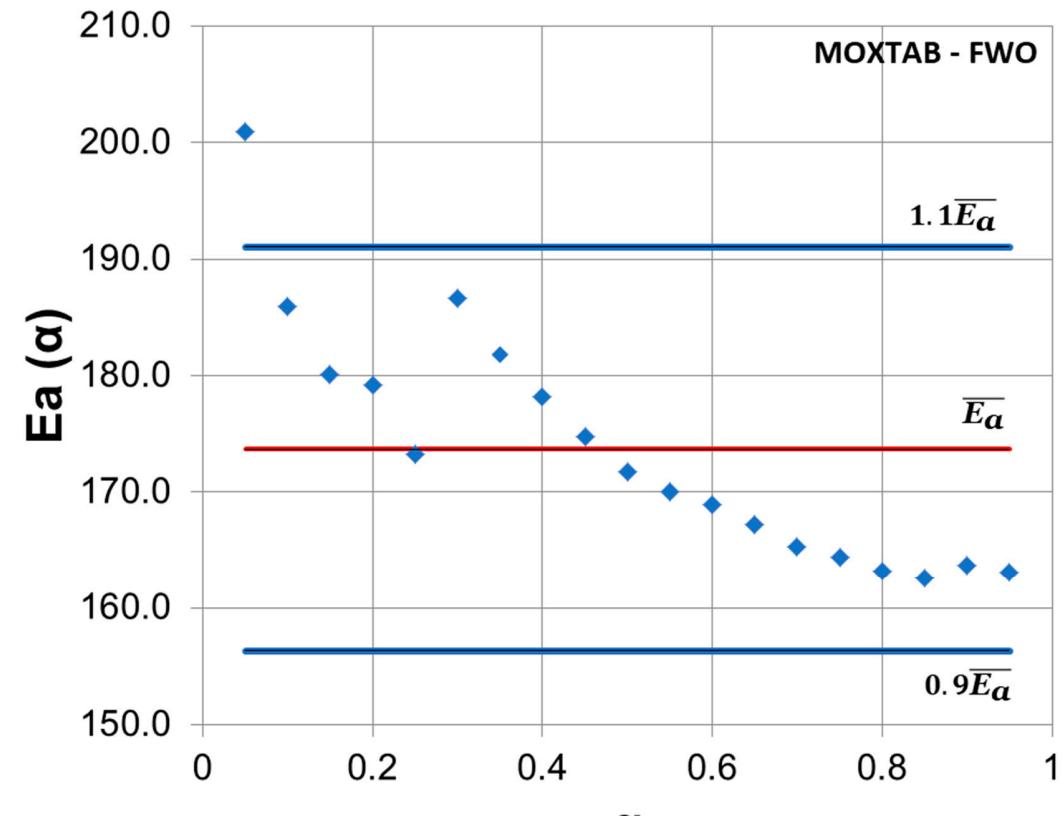
(b)



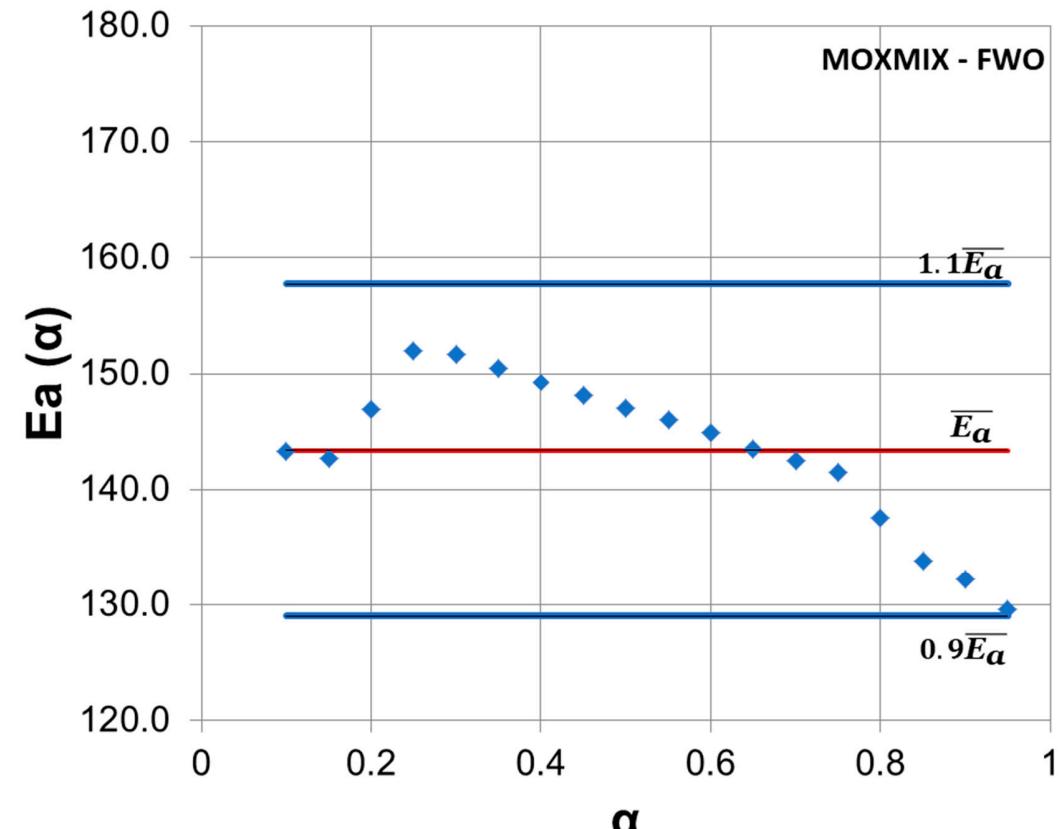
(c)

Figure S4. Linear plotting of FR differential isoconversional method of the analyzed process of decomposition for: (a) MOX; (b) MOXTAB; and (c) MOXMIX.

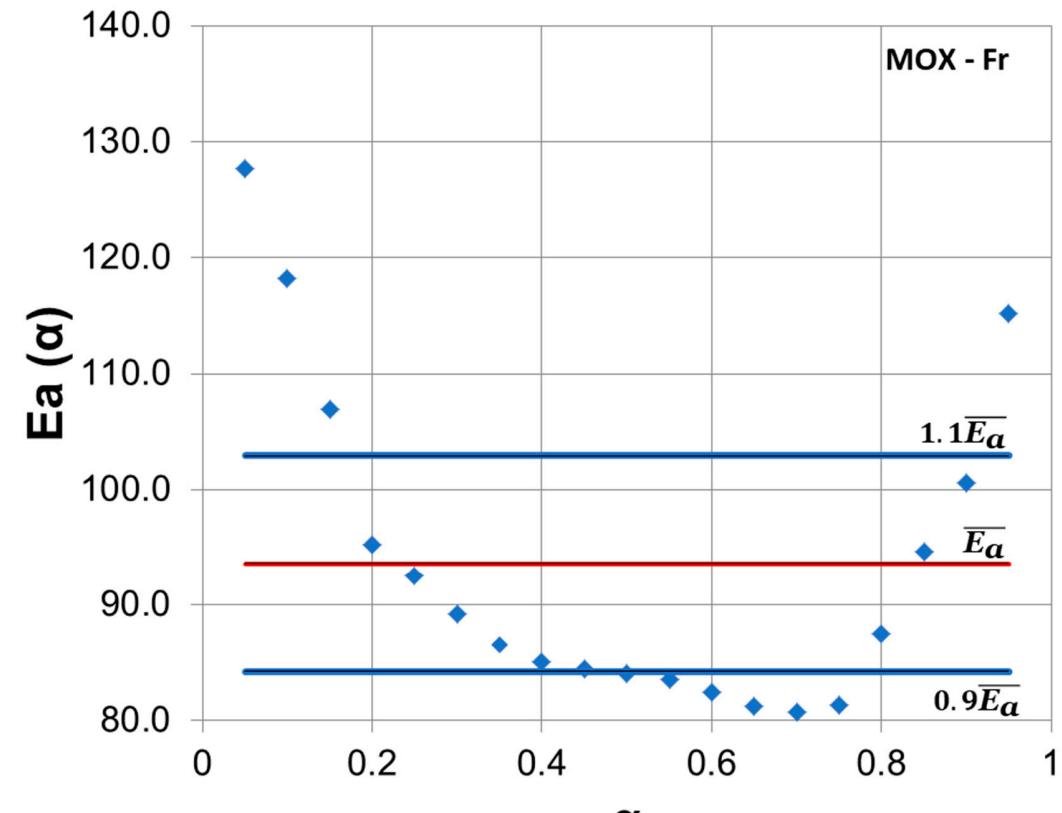




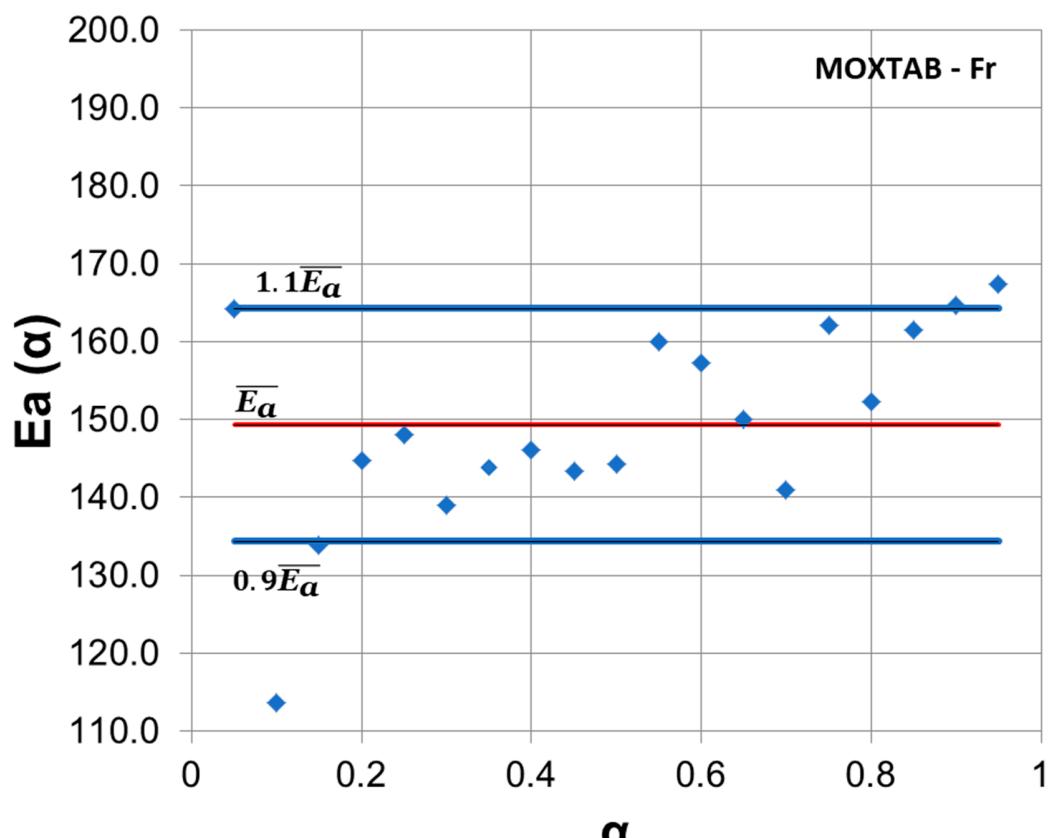
(b)



(c)



(d)



(e)

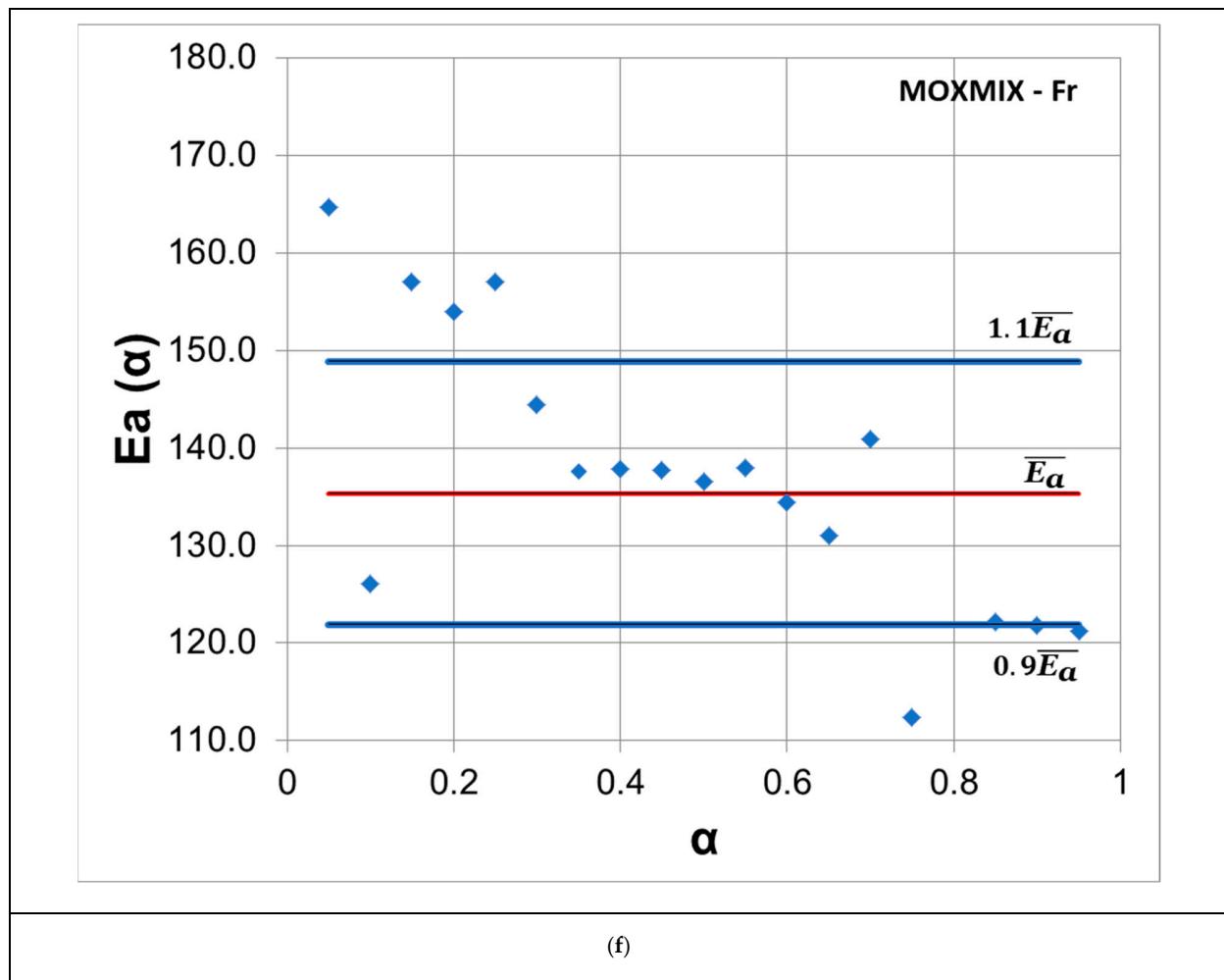


Figure S5. Variation in E_a vs. α according to FWO integral isoconversional method (a,b,c) and FR differential isoconversional method (d,e,f) of the analyzed decomposition process for: (a,d) MOX; (b,e) MOXTAB; and (c,f) MOXMIX.