

Catalyst-Free Transfer Hydrogenation of Activated Alkenes Exploiting Isopropanol as the Sole and Traceless Reductant

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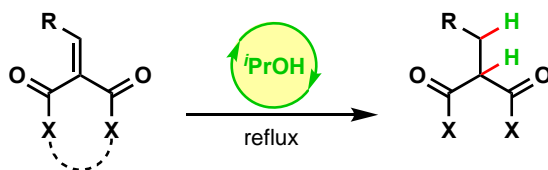
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Agustin Rodriguez grew up in Monterrey, Nuevo León, México. He did his undergraduate studies in Tecnológico de Monterrey (ITEMS) where he obtained his B.Sc. in Nanotechnology and Chemical Sciences Engineering. His senior year he worked in CINVESTAV-Mexico in the Juaristi group developing new mechano-enzymatic methods for the kinetic resolution of pharmaceutically active compounds. Since the Fall of 2020, he began his graduate studies at Rice University in the Kürti lab, where he is currently working on new reactions for the synthesis of strained rings.

Abstract:

Both metal-catalyzed and organocatalytic transfer hydrogenation reactions are widely employed for the reduction of C=C, C=O and C=N bonds. Described herein is an unconventional catalyst-free transfer hydrogenation reaction of activated alkenes using isopropanol as an eco-friendly reductant and solvent. The reaction gives convenient synthetic access to a wide range of substituted malonic acid half oxyesters (SMAHOs) in moderate to good yields. Mechanistic investigations point towards an unprecedented hydrogen bond-assisted transfer hydrogenation process.



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| <input checked="" type="checkbox"/> Metal/Catalyst-Free | <input checked="" type="checkbox"/> Zero Waste | <input checked="" type="checkbox"/> FG Compatibility |
| <input checked="" type="checkbox"/> Atom Economic | <input checked="" type="checkbox"/> Low Cost | <input checked="" type="checkbox"/> Operational Simplicity |

References:

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