Supplementary Information (SI) for Organic & Biomolecular Chemistry. This journal is © The Royal Society of Chemistry 2025

Ruthenium-catalyzed Synthesis of Poly-substituted furans via Intermolecular Oxidative Annulation of 3-(phenyl ethynyl) oxazolidine-2-ones.

Vikrant Nawal Vikram,^{a, b} Mohit Chourasiya,^{a, b} Ruchir Kant^c and Narender Tadigoppulla^{a, b, *}

^aMedicinal & Process Chemistry Division, CSIR-Central Drug Research Institute, Lucknow-226031, U.P., India ^bAcademy of Scientific and Innovative Research (AcSIR), Ghaziabad-201002, India. ^cMolecular and Structural Biology Division, CSIR-Central Drug Research Institute, Lucknow-226031, India

E-mail: <u>t_narender@cdri.res.in</u>

Table of Contents

1.	General Information	2
2.	General procedure for the synthesis of Ynamides	2-3
3.	General procedure for the synthesis of tetrasubstituted furans	3
4.	Single Crystal X-rays of compound 2c	3-5
5.	References	5
6.	Spectral data of Obtained compound	6-12
7.	Spectral data of Ynamides	12-17
8.	Copies of NMR Spectra	

1. General procedure

Most of the metals, reagents, and starting materials were purchased from commercial sources and used as such. The progress of the reaction was monitored by analytical TLC on silica gel G/GF 254 plates. The column chromatography was performed with silica gel 100-200 mesh using EtOAc/hexane as an eluent. Proton and carbon nuclear magnetic resonance spectra (¹H, 400 MHz; and ¹³C, 100 MHz) spectra were recorded either Bruker Avance on a 400 MHz using TMS as an internal standard (¹H NMR, CDCl₃ at 7.26 ppm; ¹³C NMR, CDCl₃ at 77.0 ppm) and chemical shifts (δ ppm) (multiplicity, coupling constant (Hz), integration). The abbreviations for multiplicity are as follows: s = singlet, d = doublet, t = triplet, q = quartet, m = multiplet, dd = doublet of doublets. Melting points are uncorrected were determined in capillary tubes on a hot stage melting point apparatus containing silicon oil. High resolution mass spectra were taken with a 3000-mass spectrometer and Q-TOF Analyzer. IR spectra were recorded using a FTIR spectrophotometer.

2. General procedure for the synthesis of starting Ynamides 1a-1z, 1a'-1b';



Step 2nd :cross-coupling of oxazolidinones with alkyne bromide



Step 1st: synthesis of alkyne bromide;

A solution of alkyne (1.0 eq.), N-bromosuccinimide (1.2 eq.), and silver nitrate (5 mol%) in acetone (0.3 M) was stirred at room temperature for 3 h. After the completion of the reaction, it was filtered and washed with hexane. After that, acetone was then removed on a rotary evaporator. The resulting product was dissolved in hexane, and the solution was passed through a short silica gel column to give alkyne bromide upon concentrating the fractions under reduced pressure.¹⁻⁵

Step 2nd: cross-coupling of oxazolidinones with alkyne bromide;

Alkyne bromide (1.0 eq.) obtained from step a, oxazolidinones (1.2 eq.), K_2CO_3 (2.0 eq.), $CuSO_4 \cdot 5H_2O$ (10 mol%), 1,10-phenanthroline (20 mol%) in toluene (0.5 M) was heated at 80 °C for 12 h. After cooling to room temperature, the reaction mixture was concentrated under reduced pressure, and the residue was purified by flash chromatography on silica gel.¹⁻⁵

3. General procedure for the synthesis of tetrasubstituted furans (2a-2z);



We synthesized tetrasubstituted furan derivatives by reacting with ynamide (1.06 mmol), $Cu(OTf)_2$ (1 equiv.) as an oxidant and $CsCO_3$ (2 equiv.) as a base in the presence of catalyst $[Ru(p-cymene)Cl_2]_2$ (10 mol%) in *t*-amyl alcohol under N₂ condition. The reaction mixture was stirred at 80°C for 2hr. After completion of the reaction water (20 ml) was added and then reaction mixture was extracted with EtOAc (3×50ml), dried over anhydrous Na₂SO₄. The solvent was evaporated under reduced pressure to obtain crude product which was further purified by column chromatography on a silica gel (100-200 mesh) using EtOAc/hexane as solvent system. All the products (**2a-2z**) were prepared by same procedure.

4. Single Crystal X-rays of compound 2c;

Crystallization: Crystals of compound **2c** were grown from the solvent Chloroform and Methanol (9:1) by a slow evaporation method.



Figure 1 ORTEP diagram drawn with 50% ellipsoid probability for non-H atoms of the crystal structure of compound **2c** determined at 295K.

X-Ray Data Collection and Structure Refinement Details:

A good quality single crystal of size 0.10 x 0.09 x 0.04mm, was selected under a polarizing microscope and was mounted on a glass fiber for data collection. Single crystal X-ray data for compound 2c were collected on the Rigaku XtaLAB Synergy-S single crystal X-ray diffractometer equipped with a HyPix-6000HE Hybrid Photon Counting (HPC) detector and dual Mo and Cu microfocus sealed X-ray source with kappa goiniometer at 295 (2) K. Data collection cell determination, and data reduction was performed using the CrysAlisPro⁶ software. Structure solution and refinement were performed by using SHELX-97.⁷ Refinement of coordinates and anisotropic thermal parameters of non-hydrogen atoms were carried out by the full-matrix least-squares method. The hydrogen atoms attached to carbon atoms were generated with idealized geometries and isotropically refined using a riding model.

Compound	2c
Empirical formula	$C_{26} H_{26} N_2 O_5$
Formula weight	446.18
Crystal System	Monoclinic
Space group	<i>P</i> 2 ₁ /n
<i>a</i> (Å)	9.5806(2)
b (Å)	9.2512(2)
<i>c</i> (Å)	25.9330(5)
α (°)	90.00
β(°)	91.395(2)
γ (°)	90.00
V (ų)	2297.81(8)
Ζ	4
D _c (g/cm ³)	1.291
F ₀₀₀	944
μ (mm ⁻¹)	0.735
θ_{\max} (°)	77.94

Table 1 Crystal data and structure refinement details for 2c.

Total reflections	14941	
Unique reflections	4575	
Reflections $[l > 2\sigma(l)]$	3596	
Parameters	301	
R _{int}	0.0308	
Goodness-of-fit	1.090	
$R [F^2 > 2\sigma(F^2)]$	0.0641	
wR (F ² , all data)	0.1989	
CCDC No.	2408896	

References

- Y. Zhang, R. P. Hsung, M. R. Tracey, K. C. M. Kurtz, and E. L. Vera Copper Sulfate-Pentahydrate-1,10-Phenanthroline Catalyzed Amidations of Alkynyl Bromides. Synthesis of Heteroaromatic Amine Substituted Ynamides *Org. Lett.*, Vol. 6, 2004 1151-1154.
- 2. N. Ghosh, S. Nayak, and A. K Sahoo, Gold(I)-Catalyzed 6-endo-Dig Hydrative Cyclization of an Alkyne-Tethered Ynamide: Access to 1,6-Dihydropyridin-2(3H)ones. *Chem. Eur. J.* 2013, **19**, 9428-9433.
- Y. Yamaguchi, T. Ochi, T. Wakamiya, Y. Matsubara, and Z. Yoshida, New Fluorophores with Rod-Shaped Polycyano π-Conjugated Structures: Synthesis and Photophysical Properties. *Org. Lett.* 2006, 8, 717– 720.
- 4. B. Zhou, W. Chen, Y. Yang, G. Deng, and Y. Liang, A radical cyclization cascade of 2-alkynylbenzonitriles with sodium arylsulfinates. *Org. Biomol. Chem.* 2018, **16**, 7959–7963.
- 5. Y. He, X. Zhang, and X. Fan, Synthesis of naphthalene amino esters and arylnaphthalene lactone lignans through tandem reactions of 2-alkynylbenzonitriles. *Chem. Commun.* 2014, **50**, 5641-5643.
- 6. Crys Alis Pro, Oxford Diffraction /Agilent Technologies UK Ltd, Yarnton, England.
- 7. Sheldrick, G. M. Acta Crystallogr. Sect. A 2008, 64, 112–122.

^{6.} Spectral data of obtained compound;

^{3,3&#}x27;-(2,5-diphenylfuran-3,4-diyl)bis(oxazolidin-2-one) (2a)



Yield: 75% (310 mg); mp 251-253 °C; FT-IR (cm⁻¹): 3921.64, 3219.01, 2920.73, 2379.92, 1743.90, 1030.29, 759.35; ¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.730 (d, *J* = 7.72 Hz, 4H), 7.472 (t, *J* = 7.84 Hz, 4H), 7.379 (t, *J* = 7.40 Hz, 2H), 4.539 (t, *J* = 8.24 Hz, 4H), 3.940 (t, *J* = 7.40 Hz, 2H), 4.539 (t, *J* = 8.24 Hz, 4H), 3.940 (t, *J* = 7.40 Hz, 2H), 4.539 (t, *J* = 8.24 Hz, 4H), 3.940 (t, *J* = 7.40 Hz, 2H), 4.539 (t, *J* = 8.24 Hz, 4H), 3.940 (t, *J* = 7.40 Hz, 2H), 4.539 (t, *J* = 8.24 Hz, 4H), 3.940 (t, *J* = 7.40 Hz, 2H), 4.539 (t, *J* = 8.24 Hz, 4H), 3.940 (t, *J* = 7.40 Hz, 4H), 3.

8.25 Hz 4H), ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 157.75, 146.96, 129.07, 128.89, 125.05, 118.85, 63.50, 45.32; HRMS; Calculated for C₂₂H₁₈N₂O₅ [M+K]⁺, *m/z* 429.0853 found 429.0847

3,3'-(2,5-di-p-tolylfuran-3,4-diyl)bis(oxazolidin-2-one) (2b)



Yield: 74% (328 mg); mp 226-228 °C; FT-IR (cm⁻¹): 3772.13, 2919.59, 1751.68, 1084.36, 755.79, 499.65; ¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.524 (d, *J* = 7.29 Hz, 4H), 7.183 (d, *J* = 7.17 Hz, 4H), 4.440 (t, *J* = 7.65 Hz, 4H), 3.842 (t, *J* = 7.65 Hz 4H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 157.81, 146.92, 138.85, 129.73, 126.23,

124.98, 118.12, 63.46, 45.30, 21.41; HRMS; Calculated for C₂₄H₂₂N₂O₅ [M+Na]⁺, *m*/z 418.1529 found 419.4.

3,3'-(2,5-bis(4-ethylphenyl)furan-3,4-diyl)bis(oxazolidin-2-one) (2c)



Yield: 72% (340 mg); mp 201-203 °C; FT-IR (cm⁻¹): 3837.92, 3740.78, 2963.59, 1894.03, 1033.26, 754.87; ¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.627 (d, *J* = 8.25 Hz, 4H), 7.288 (d, *J* = 8.25 Hz, 4H), 4.527 (t, *J* = 8.40 Hz, 4H), 3.931 (t, *J* = 8.27 Hz 4H), 2.696 (q, *J* = 7.60Hz 4H), 1.274 (t, *J* = 7.60 Hz, 6H); ¹³C NMR (100 MHz, CDCl₃)

δ (ppm): 157.81, 146.94, 145.12, 128.53, 126.45, 125.06, 118.13, 63.46, 45.33, 28.74, 15.31; HRMS; Calculated for C₂₆H₂₆N₂O₅ [M+Na]⁺, *m/z* 469.1729 found 469.1734.

3,3'-(2,5-bis(4-propylphenyl)furan-3,4-diyl)bis(oxazolidin-2-one) (2d)



Yield: 70% (351 mg); mp 183-185 °C; FT-IR (cm⁻¹): 3922.46, 3775.85, 2922.54, 1687.31, 1034.91, 761.93; ¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.621 (d, *J* = 8.23 Hz, 4H), 7.265 (d, *J* = 8.01 Hz, 4H), 4.526 (t, *J* = 8.34 Hz, 4H), 3.932 (t, *J* = 8.28Hz 4H), 2.627 (t, *J* = 7.72 Hz, 4H), 1.721 – 1.629 (m, 4H), 0.965 (t, *J* = 7.40 Hz, 6H); ¹³C

NMR (100 MHz, CDCl₃) δ (ppm): 157.81, 146.93, 143.62, 129.12, 126.46, 124.94, 118.12, 63.46, 45.34, 37.88, 29.69, 24.34, 13.80; HRMS; Calculated for C₂₈H₃₀N₂O₅ [M+Na]⁺, *m/z* 497.2052, found 497.2047.



3,3'-(2,5-bis(4-butylphenyl)furan-3,4-diyl)bis(oxazolidin-2-one) (2e)

Yield: 69% (367 mg); mp 173-175 °C; FT-IR (cm⁻¹): 3921.16, 3775.29, 2855.35, 1754.62, 1078.66, 761.15; ¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.612 (d, *J* = 8.19 Hz, 4H), 7.270 (d, *J* = 7.95 Hz, 4H), 4.530 (t, *J* = 8.40 Hz, 4H), 3.936 (t, *J* = 8.28 Hz 4H),

2.654 (t, J = 7.75 Hz, 4H), 1.672 - 1.596 (m, 4H), 1.431 - 1.339 (m, 4H), 0.951 (t, J = 7.34 Hz, 6H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 157.80, 146.93, 143.84, 129.07, 126.42, 124.95, 118.13, 63.46, 45.34, 35.52, 33.40, 22.36, 13.94; HRMS; Calculated for C₃₀H₃₄N₂O₅, [M+Na]⁺ m/z 525.2365, found 525.2360.

3,3'-(2,5-bis(4-(tert-butyl)phenyl)furan-3,4-diyl)bis(oxazolidin-2-one) (2f)



Yield: 67% (356 mg); mp 272-274 °C; FT-IR (cm⁻¹): 3917.43, 3776.33, 2957.62, 1758.07, 1032.86, 757.44; ¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.643 (d, *J* = 8.64 Hz, 4H), 7.476 (d, *J* = 8.68 Hz, 4H), 4.534 (t, *J* = 8.38 Hz, 4H), 3.944 (t, *J* = 8.24 Hz 4H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 157.76, 151.96, 146.89, 126.20, 125.96,

124.78, 118.24, 63.44, 45.40, 34.81, 31.20; MS(ESI); Calculated for $C_{30}H_{34}N_2O_5$ [M+H]⁺, *m/z* 502.2468 found 503.6.

3,3'-(2,5-bis(4-methoxyphenyl)furan-3,4-diyl)bis(oxazolidin-2-one) (2g)



Yield: 69% (329 mg); mp 213-215 °C; FT-IR (cm⁻¹): 3912.23, 3775.22, 2345.19, 1603.23, 1303.85, 1030.93, 755.44; ¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.638 (d, *J* = 8.93 Hz, 4H), 6.985 (d, *J* = 8.93 Hz, 4H), 4.519 (t, *J* = 8.28 Hz, 4H), 3.912 (t, *J* = 8.26 Hz 4H), 3.862 (s, 6H, OMe); ¹³C NMR (100 MHz, 100 MHz), 100 MHz, 100

CDCl₃) δ (ppm): 159.90, 157.90, 146.62, 126.56, 121.80, 117.25, 114.50, 63.43, 55.37, 45.37; HRMS; Calculated for C₂₄H₂₂N₂O₇ [M+Na]⁺, *m/z* 489.1064 found 489.1059.

3,3'-(2,5-bis(4-ethoxyphenyl)furan-3,4-diyl)bis(oxazolidin-2-one) (2h)



Yield: 66% (334 mg); mp 214-216 °C; FT-IR (cm⁻¹): 3922.32, 3776.08, 2920.29, 1757.23, 1115.82, 762.28; ¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.612 (d, *J* = 8.21 Hz, 4H), 6.965 (d, *J* = 8.25 Hz, 4H), 4.514 (t, *J* = 7.86 Hz, 4H), 4.082 (q, *J* = 6.73 Hz 4H), 3.910 (t, *J* = 7.53 Hz, 4H), 1.444 (t, *J* = 6.67 Hz, 6H); ¹³C NMR (100 MHz, CDCl₃)

δ (ppm): 159.27, 157.92, 146.63, 126.53, 121.63, 117.13, 114.97, 63.60, 63.44, 45.37, 14.79; HRMS; Calculated for C₂₆H₂₆N₂O₅ [M+Na]⁺, *m/z* 501.1638, found 501.1632.

3,3'-(2,5-bis(4-fluorophenyl)furan-3,4-diyl)bis(oxazolidin-2-one) (2i)



Yield: 81% (366 mg); mp 277-279 °C; FT-IR (cm⁻¹): 3833.34, 3738.11, 2918.56, 1636.17, 1391.26, 1229.66, 833.93, 766.17; ¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.716 – 7.682 (m, 4H), 7.195 – 7.153 (m, 4H), 4.538 (t, *J* = 8.14 Hz, 4H), 3.909 (t, *J* = 8.16 Hz 4H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 164.12, 161.60, 157.76, 146.20,

127.11, 127.03, 125.04, 118.61, 116.44, 116.22, 63.50, 45.36; HRMS; Calculated for C₂₂H₁₆F₂N₂O₅ [M+Na]⁺, *m/z* 449.0925, found 449.0919.

3,3'-(2,5-bis(4-chlorophenyl)furan-3,4-diyl)bis(oxazolidin-2-one) (2j)



Yield: 80% (388 mg); mp 253-255 °C; FT-IR (cm⁻¹): 3920.71, 3776.53, 2921.74, 1688.39, 1032.90, 496.72; ¹H NMR (400MHz, CDCl₃) δ (ppm): 7.651 (d, *J* = 8.70 Hz, 4H), 7.451 (d, *J* = 8.74 Hz, 4H), 4.543 (t, *J* = 8.34 Hz, 4H), 3.912 (t, *J* = 8.23 Hz 4H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 157.67, 146.27, 134.95, 129.44, 127.12,

126.28, 119.39, 63.54, 45.29, 29.69; HRMS; Calculated for $C_{22}H_{16}Cl_2N_2O_5$ [M+Na]⁺, *m/z* 481.0334, found 481.0328.

3,3'-(2,5-bis(4-bromophenyl)furan-3,4-diyl)bis(oxazolidin-2-one) (2k)



Yield: 78% (451 mg); mp 274-276 °C; FT-IR (cm⁻¹): 3922.43, 3775.24, 2919.26, 1753.04, 1078.84, 764.45; ¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.612 (d, *J* = 8.79 Hz, 4H), 7.578 (d, *J* = 8.66 Hz, 4H), 4.541 (t, *J* = 8.36 Hz, 4H), 3.910 (t, *J* = 8.30 Hz 4H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 157.68, 146.34, 132.39, 127.53, 126.49,

123.19, 119.52, 63.55, 45.27; HRMS; Calculated for C₂₂H₁₆Br₂N₂O₅, *m/z* [M+K]⁺, 584.9063, found 584.9058.

3,3'-(2,5-di([1,1'-biphenyl]-4-yl)furan-3,4-diyl)bis(oxazolidin-2-one) (2l)



Yield: 78% (448 mg); mp 259-261 °C; FT-IR (cm⁻¹): 3923.04 3774.58, 2919.93, 1633.88, 1034.82, 759.68, 498.62; ¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.822 (d, *J* = 8.40 Hz, 4H), 7.718 (d, *J* = 8.43 Hz, 4H), 7.649 (dd, *J* = 8.40Hz, *J* = 1.32 Hz, 4H), 7.483 (d, *J* = 7.80 Hz, 4H), 7.391 (d, *J* = 7.35 Hz, 2H), 4.576 (t, *J* = 8.41 Hz, 4H), 3.999 (t, *J* = 7.80 Hz, 4H), 7.901 (d, *J* = 7.80 Hz, 2H), 4.576 (t, *J* = 8.41 Hz, 4H), 3.999 (t, *J* = 7.80 Hz, 4H), 7.901 (d, *J* = 7.80 Hz, 2H), 4.576 (t, *J* = 8.41 Hz, 4H), 3.999 (t, *J* = 7.80 Hz, 4H), 7.901 (d, *J* = 7.80 Hz, 2H), 4.576 (t, *J* = 8.41 Hz, 4H), 3.999 (t, *J* = 8.41 Hz, 4H), 3.999 (t, *J* = 8.41 Hz, 4H), 3.999 (t, *J* = 7.80 Hz, 4H), 7.911 (d, *J* = 7.80 Hz, 2H), 4.576 (t, *J* = 8.41 Hz, 4H), 3.999 (t, *J* = 7.80 Hz, 4H), 7.911 (d, *J* = 7.80 Hz, 2H), 4.576 (t, *J* = 8.41 Hz, 4H), 3.999 (t, *J* = 7.80 Hz, 4H), 7.911 (d, *J* = 7.80 Hz, 2H), 4.576 (t, *J* = 8.41 Hz, 4H), 3.999 (t, *J* = 7.80 Hz, 4H), 7.911 (d, *J* = 7.80 Hz, 2H), 4.576 (t, *J* = 8.41 Hz, 4H), 3.999 (t, *J* = 7.80 Hz, 4H), 7.911 (d, *J* = 7.80 Hz, 2H), 4.576 (t, *J* = 8.41 Hz, 4H), 3.999 (t, *J* = 7.80 Hz, 4H), 7.911 (d, *J* = 7.80 Hz, 2H), 4.576 (t, *J* = 8.41 Hz, 4H), 3.999 (t, *J* = 7.80 Hz, 4H), 7.911 (d, *J* = 7.80 Hz, 2H), 4.576 (t, *J* = 8.41 Hz, 4H), 3.911 (d, *J* = 7.80 Hz, 4H), 7.911 (d, *J* = 7.8

8.32 Hz 4H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 157.79, 146.88, 141.51, 140.09, 128.96, 127.83, 127.73, 127.70, 127.01, 125.44, 119.08, 63.55, 45.38; HRMS; Calculated for C₃₄H₂₆N₂O₅ [M+Na]⁺, *m/z* 565.1739, found 565.1734.

3,3'-(2,5-bis(4-(trifluoromethyl)phenyl)furan-3,4-diyl)bis(oxazolidin-2-one) (2m)



Yield: 76% (423 mg); mp 249-251 °C; FT-IR (cm⁻¹): 3920.20, 3776.50, 2922.04, 1764.55, 1245.51, 763.04, 466.42; ¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.856 (d, *J* = 8.23 Hz, 4H), 7.751 (d, *J* = 8.34 Hz, 4H), 4.573 (t, *J* = 8.36 Hz, 4H), 3.941 (t, *J* = 8.23 Hz 4H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 156.54, 145.31, 130.68, 129.98,

129.62, 125.20, 125.17, 124.25, 124.07, 121.37, 119.89, 62.59, 44.28; HRMS; Calculated for $C_{24}H_{16}F_6N_2O_5$ [M+K]⁺, *m/z* 565.0600 found 565.0595.

dimethyl 4,4'-(3,4-bis(2-oxooxazolidin-3-yl)furan-2,5-diyl)dibenzoate (2n)



Yield: 83% (445 mg); mp 289-291 °C; FT-IR (cm⁻¹): 3922.21, 3773.09, 1627.61, 1079.32, 827.62, 498.22; ¹H NMR (400 MHz, CDCl₃) δ (ppm): 8.152 (d, *J* = 8.53 Hz, 4H), 7.819 (d, *J* = 8.52 Hz, 4H), 4.573 (t, *J* = 8.30 Hz, 4H), 3.957 (s, 6H), 3.951 (t, *J* = 8.09 Hz 4H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 166.32,

157.57, 146.62, 132.42, 130.41, 130.24, 124.78, 120.97, 63.63, 52.37, 45.23; HRMS; Calculated for $C_{26}H_{22}N_2O_9$ [M+Na]⁺, *m/z* 529.1223, found 529.1218.

3,3'-(2,5-di(thiophen-3-yl)furan-3,4-diyl)bis(oxazolidin-2-one) (20)



Yield: 53% (225 mg); mp 113-115 °C; FT-IR (cm⁻¹): 3922.11, 3773.68, 2857.18, 1645.05, 1079.04, 827.43, 760.41; ¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.599 (dd, *J* = 2.93 Hz, *J* = 1.29 Hz, 2H), 7.438 (dd, *J* = 5.07 Hz, *J* = 2.95 Hz, 2H), 7.407 (dd, *J* = 5.07 Hz, *J* = 1.29 Hz, 2H), 4.542 (t, *J* = 8.18 Hz, 4H), 3.954 (t, *J* = 8.36 Hz 4H); ¹³C NMR (100 MHz, CDCl₃) δ

(ppm): 157.78, 144.31, 129.71, 127.00, 124.47, 121.91, 117.50, 63.45, 45.56; HRMS; Calculated for $C_{18}H_{14}N_2O_5S_2$ [M+Na]⁺, *m/z* 425.0242, found 425.0236.

3,3'-(2,5-bis(3-fluorophenyl)furan-3,4-diyl)bis(oxazolidin-2-one) (2p)



Yield: 78% (352 mg); mp 283-285 °C; FT-IR (cm⁻¹): 3410.04, 2918.58, 1751.87, 830.88, 768.98; ¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.517 (d, *J* = 7.86 Hz, 2H), 7.507 – 7.426 (m, 4H), 7.119 – 7.074 (m, 2H), 4.563 (t, *J* = 8.33 Hz, 4H), 3.950 (t, *J* = 8.26 Hz 4H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 164.14, 162.17, 157.62, 146.06, 130.92, 130.85, 130.53, 120.65, 119.95, 116.08, 115.92, 112.08, 111.89, 63.58, 45.26; HRMS; Calculated for

 $C_{22}H_{16}F_2N_2O_5$ [M+Na]⁺, *m*/z 449.0925, found 449.0929.

3,3'-(2,5-bis(3-chlorophenyl)furan-3,4-diyl)bis(oxazolidin-2-one) (2q)



Yield: 76% (368 mg); mp 211-213 °C; FT-IR (cm⁻¹): 3838.30, 3744.65, 2919.85, 1756.06, 1031.80, 762.41, 497.61; ¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.717 (t, *J* = 1.81 Hz, 2H), 7.624 - 7.598 (m, 2H), 7.421 (t, *J* = 7.95 Hz, 2H), 7.382 - 7.354 (m, 2H), 4.560 (t, *J* = 8.34 Hz, 4H), 3.939 (t, *J* = 8.23 Hz 4H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 157.61, 145.90, 135.29, 130.46, 130.20, 129.05, 125.01, 123.03, 120.01, 63.62, 45.26; HRMS; Calculated

for C₂₂H₁₆Cl₂N₂O₅, *m*/*z* 481.0334, found [M+Na]⁺ 481.0328.

3,3'-(2,5-di-m-tolylfuran-3,4-diyl)bis(oxazolidin-2-one) (2r)



Yield: 65% (288 mg); mp 193-195 °C; FT-IR (cm⁻¹): 3410.99, 2920.72, 1759.81, 1081.86, 826.87, 763.16; ¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.534 (d, *J* = 2.33 Hz, 2H), 7.517 (d, *J* = 8.33 Hz, 2H), 7.351 (d, *J* = 7.67 Hz, 2H), 7.191 (d, *J* = 7.58 Hz, 2H), 4.528 (t, *J* = 8.36 Hz, 4H), 3.933 (t, *J* = 8.25 Hz 4H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 157.78, 146.99, 138.77, 129.64, 128.94, 128.86, 125.64, 122.24, 118.65, 63.52, 45.29, 21.62; HRMS;

Calculated for C₂₄H₂₂N₂O₅, *m*/*z* 441.1426, found [M+Na]⁺ 441.1421.

3,3'-(2,5-bis(3,4-dichlorophenyl)furan-3,4-diyl)bis(oxazolidin-2-one) (2s)



Yield: 79% (440 mg); mp 304-306 °C; FT-IR (cm⁻¹): 3772.16, 3142.94, 1765.47, 1398.04, 1033.66, 760.13; ¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.806 (s, 2H), 7.558 (s, 4H), 4.562 (t, *J* = 8.00 Hz, 4H), 3.923 (t, *J* = 8.02 Hz 4H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 157.55, 145.32, 133.74, 133.31, 131.28, 128.23, 126.73, 126.04, 120.40, 63.65, 45.26; ESIMS; Calculated for C₂₂H₁₄Cl₄N₂O₅ [M+Na]⁺,

m/z 525.96, found 527.0.

3,3'-(2,5-diphenylfuran-3,4-diyl)bis(4-phenyloxazolidin-2-one) (2t)



Yield: 62% (356 mg); mp 129-131 °C; FT-IR (cm⁻¹): 3773.47, 2921.38, 1632.39, 1037.64, 766.62; ¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.328- 7.266 (m, 10H), 7.074 (dd, *J* = 8.29 Hz, *J* = 1.21 Hz 4H), 7.024 (d, *J* = 7.41 Hz, 2H), 6.907 (d, *J* = 7.74 Hz, 4H), 5.188 (t, *J* = 8.61 Hz, 2H), 4.857 (t, *J* = 9.00 Hz, 2H), 4.542 (t, *J* = 8.83 Hz, 2H); ¹³C NMR (100 MHz, CDCl₃)

δ (ppm): 156.48, 149.93, 133.87, 127.92, 127.73, 127.46, 127.21, 125.90, 125.74, 124.00, 115.41, 68.99, 61.80;
HRMS; Calculated for C₃₄H₂₆N₂O₅ [M+K]⁺, *m/z* 581.1479 found 581.1473.

3,3'-(2,5-di-p-tolylfuran-3,4-diyl)bis(4-phenyloxazolidin-2-one) (2u)



Yield: 60% (362 mg); mp 93-95 °C; FT-IR (cm⁻¹): 3901.69, 3731.13, 2924.37, 1754.31, 1089.33, 772.32; ¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.193 (d, *J* = 8.15 Hz, 4H), 7.073 (d, *J* = 6.70 Hz, 4H), 7.071 (d, *J* = 7.27 Hz, 4H), 7.028 (d, *J* = 7.41 Hz, 2H), 6.913 (t, *J* = 7.73 Hz, 4H), 5.167 (t, *J* = 8.66 Hz, 2H), 4.837 (t, *J* = 8.93 Hz, 2H), 4.511 (t, *J* = 8.81 Hz 2H), 2.359 (s, 6H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm):

157.52, 150.83, 138.68, 135.05, 128.91, 128.87, 128.25, 128.21, 126.67, 125.84, 115.79, 70.05, 62.86, 21.39; HRMS; Calculated for $C_{36}H_{30}N_2O_5$ [M+Na]⁺, *m/z* 593.2052 found 593.2047.

3,3'-(2,5-bis(4-ethylphenyl)furan-3,4-diyl)bis(4-phenyloxazolidin-2-one) (2v)



Yield: 58% (367 mg); mp 88-90 °C; FT-IR (cm⁻¹): 3918.91, 3776.48, 2858.63, 1759.66, 1039.32, 706.90, 497.18; ¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.217 (d, J = 8.19 Hz, 4H), 7.089 (d, J = 7.50 Hz, 4H), 7.069 (d, J = 6.60 Hz, 4H), 7.016 (t, J = 7.39 Hz, 2H), 6.894 (t, J = 7.76 Hz, 4H), 5.170 (t, J = 8.66 Hz, 2H), 4.839 (t, J = 8.93 Hz, 2H), 4.524 (t, J = 8.76 Hz, 2H), 2.647 (q, J = 7.58 Hz, 4H), 1.252 (t, J = 7.66 Hz, 2H), 2.647 (q, J = 7.58 Hz, 4H), 1.252 (t, J = 7.66 Hz, 2H), 4.524 (t, J = 8.76 Hz, 2H), 2.647 (q, J = 7.58 Hz, 4H), 1.252 (t, J = 7.66 Hz, 2H), 2.647 (q, J = 7.58 Hz, 4H), 1.252 (t, J = 7.66 Hz, 2H), 2.647 (q, J = 7.58 Hz, 4H), 1.252 (t, J = 7.66 Hz, 2H), 2.647 (q, J = 7.58 Hz, 4H), 1.252 (t, J = 7.66 Hz, 2H), 2.647 (q, J = 7.58 Hz, 4H), 1.252 (t, J = 7.66 Hz, 2H), 2.647 (q, J = 7.58 Hz, 4H), 1.252 (t, J = 7.66 Hz, 2H), 2.647 (q, J = 7.58 Hz, 4H), 1.252 (t, J = 7.66 Hz, 2H), 2.647 (q, J = 7.58 Hz, 4H), 1.252 (t, J = 7.66 Hz, 2H), 2.647 (q, J = 7.58 Hz, 4H), 1.252 (t, J = 7.66 Hz, 2H), 2.647 (q, J = 7.58 Hz, 4H), 1.252 (t, J = 7.66 Hz, 2H), 2.647 (q, J = 7.58 Hz, 4H), 1.252 (t, J = 7.66 Hz, 2H), 2.647 (q, J = 7.58 Hz, 4H), 1.252 (t, J = 7.66 Hz, 2H), 2.647 (q, J = 7.58 Hz, 4H), 3.647 (q, J = 7.58 Hz), 3.6

6H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 157.55, 150.91, 145.14, 130.06, 128.86, 128.27, 128.16, 127.74, 126.05, 115.83, 69.99, 62.82, 28.82, 15.77; HRMS; Calculated for C₃₈H₃₄N₂O₅ [M+Na]⁺, *m/z* 621.2365 found 621.2360.

3,3'-(2,5-bis(4-propylphenyl)furan-3,4-diyl)bis(4-phenyloxazolidin-2-one) (2w)



Yield: 56% (317 mg); mp 98-100 °C; FT-IR (cm⁻¹): 3900.42, 3826.58, 2857.69, 2308.93, 1846.06, 1385.05, 764.58; ¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.213 (d, J = 8.24 Hz, 4H), 7.074 (d, J = 8.04 Hz, 4H), 7.072 (d, J = 7.86 Hz, 4H), 7.017 (t, J = 7.43 Hz, 2H), 6.987 (t, J = 7.77 Hz, 4H), 5.174 (t, J = 8.64 Hz, 2H), 4.840 (t, J = 8.97Hz, 2H), 4.525 (t, J = 8.77 Hz, 4H), 2.588 (t, J = 7.73 Hz, 4H), 1.702 – 1.610 (m,

4H), 0.964 (t, J = 7.37 Hz, 6H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 157.54, 150.90, 143.50, 135.04, 128.8, 128.36, 128.27, 128.18, 126.65, 126.05, 115.80, 69.99, 62.85, 37.86, 24.58, 13.71; MS (ESI); Calculated for C₄₀H₃₈N₂O₅ [M+Na]⁺, m/z 649.2678, found 649.2673.

3,3'-(2,5-bis(4-(tert-butyl)phenyl)furan-3,4-diyl)bis(4-phenyloxazolidin-2-one) (2x)



Yield: 53% (354 mg); mp 115-117 °C; FT-IR (cm⁻¹): 3911.00, 3776.27, 2958.09, 1760.07, 1037.81, 761.02, 554.47; ¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.275 (d, *J* = 8.58 Hz, 4H), 7.207 (d, *J* = 8.59 Hz, 4H), 7.067 (dd, *J* = 8.33 Hz, *J* = 1.21 Hz 4H), 6.999 (t, *J* = 7.44 Hz, 2H), 6.866 (t, *J* = 7.75 Hz, 4H), 5.171 (t, *J* = 8.58 Hz, 2H), 4.841 (t, *J* =

8.98 Hz 2H), 4.543 (t, *J* = 8.48 Hz 2H), 1.336 (s,18H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 157.55, 151.90, 150.89, 135.10, 128.83, 128.29, 128.10, 126.51, 125.77, 125.09, 115.93, 69.90, 62.76, 34.73, 31.27; HRMS; Calculated for C₄₂H₄₂N₂O₅ [M+Na]⁺, *m/z* 677.2991 found 677.2986.

3,3'-(2,5-bis(4-ethoxyphenyl)furan-3,4-diyl)bis(4-phenyloxazolidin-2-one) (2y)



Yield: 52% (360 mg); mp 107-109 °C; FT-IR (cm⁻¹): 3835.79, 3609.29, 2922.73, 1756.31, 1180.94, 828.90; ¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.220 (d, *J* = 8.88 Hz, 4H), 7.079 (dd, *J* = 8.36 Hz, *J* = 1.28 Hz, 4H), 7.040 (d, *J* = 7.43 Hz, 2H), 6.947 (t, *J* = 7.69 Hz, 4H), 6.787 (d, *J* = 8.88 Hz, 4H), 5.159 (t, *J* = 8.64 Hz, 2H), 4.836 (t, *J* = 8.97 Hz, 2H), 4.507 (t, *J* = 8.80 Hz, 2H), 4.057 (q, *J* = 6.99 Hz, 4H), 1.445 (t, *J*

=7.02Hz, 6H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 159.33, 157.60, 150.48, 135.17, 128.89, 128.21, 121.33, 115.05, 114.28, 70.01, 63.55, 62.82, 14.76; HRMS; Calculated for C₃₈H₃₄N₂O₇ [M+Na]⁺, *m/z* 653.2264 found 653.2258.

3,3'-(2,5-bis(4-chlorophenyl)furan-3,4-diyl)bis(4-phenyloxazolidin-2-one) (2z)



Yield: 64% (413 mg); mp 223-225 °C; FT-IR (cm⁻¹): 3923.04, 3774.58, 2919.93, 1633.88, 1386.46, 1034.82, 830.37, 498.62; ¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.244 (d, *J* = 9.10 Hz, 4H), 7.212 (d, *J* = 8.86 Hz, 4H), 7.098 – 7.065 (m, 6H), 6.968 (d, *J* = 7.76 Hz, 4H), 5.154 (t, *J* = 8.44 Hz, 2H), 4.869 (t, *J* = 8.98 Hz 2H), 4.577 (t, 7.244 (d, *J* = 8.80 Hz, 2H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 157.40, 150.13,

134.92, 129.17, 128.56, 128.39, 128.18, 128.01, 126.73, 117.08, 69.96, 62.75; ESMS; Calculated for $C_{34}H_{24}Cl_2N_2O_5$ [M+H]⁺, *m/z* 610.10, found 611.1.

3,3'-(2,5-bis(3-fluorophenyl)furan-3,4-diyl)bis(4-phenyloxazolidin-2-one) (2a')



Yield: 67% (410 mg); mp 228-230 °C; FT-IR (cm⁻¹): 3921.94, 3776.16, 2856.94, 2379.58, 1385.64, 1106.49, 759.56; ¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.273 – 7.219 (m, 2H), 7.118 (t, *J* = 7.47 Hz, 6H), 7.059 (d, *J* = 7.40 Hz, 2H), 6.997 (d, *J* = 6.67 Hz, 4H), 6.995 (t, *J* = 8.02 Hz, 4H), 5.173 (t, J = 8.48 Hz, 2H), 4.879 (t, *J* = 9.04 Hz, 2H), 4.615 (t, *J* = 8.88 Hz 2H);¹³C NMR (100 MHz, CDCl₃) δ (ppm): 163.68, 161.23, 157.32, 149.89, 134.80, 130.08, 129.99, 129.00, 129.18, 128.36, 128.22, 122.47, 117.48, 115.96, 115.75,

113.65, 113.41, 69.91, 62.70; HRMS; Calculated for $C_{34}H_{24}F_2N_2O_5$ [M+Na]⁺, *m/z* 601.1551 found [M+Na]⁺ 601.1545.

3,3'-(2,5-bis(3,4-dichlorophenyl)furan-3,4-diyl)bis(4-phenyloxazolidin-2-one) (2b')



Yield: 66% (474 mg); mp 218 -220 °C; FT-IR (cm⁻¹): 3914.40, 3772.51, 2925.02, 2342.35, 1761.14, 1397.22, 1078.33, 767.60; ¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.319 (d, *J* = 8.13 Hz, 2H), 7.139 – 7.097 (m, 8H), 7.009 (t, *J* = 6.99 Hz, 4H), 5.151 (t, *J* = 7.99 Hz, 2H), 4.888 (t, J = 8.88 Hz, 2H), 4.666 (t, *J* = 8.36 Hz, 2H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 157.36, 149.14, 134.85, 133.20, 132.67, 130.28, 129.30, 128.44, 128.37, 128.21, 127.75, 125.83, 118.13, 69.77, 62.57; ESIMS;

Calculated for $C_{34}H_{22}Cl_4N_2O_5$ [M+Na]⁺, *m*/z 678.02 found [M+H]⁺ 679.1.

8. Spectral data of Ynamides;

3-(phenylethynyl)oxazolidin-2-one (1a)



¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.453 - 7.428 (m, 2H), 7.316 - 7.291 (m, 3H), 4.505 - 4.465 (dd, *J* = 8.14 Hz, *J* = 7.81 Hz, 2H), 4.028 - 3.988 (dd, *J* = 8.14 Hz, *J* = 7.81 Hz, 2H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 155.96, 131.85, 131.56, 128.33, 128.21, 122.18, 79.05, 71.16, 63.12, 47.05.

3-(p-tolylethynyl)oxazolidin-2-one (1b)



¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.332 (d, *J* = 8.10 Hz, 2H), 7.017 (d, *J* = 7.90 Hz, 2H), 4.488 - 4.448 (dd, *J* = 8.18 Hz, *J* = 7.79 Hz, 2H), 4.008 - 3.968 (dd, *J* = 8.19 Hz, *J* = 7.78 Hz, 2H), 2.339 (s, 3H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 155.99, 138.41, 131.62, 129.08, 119.01, 78.29, 71.20, 63.03, 47.10, 21.45.

3-((4-ethylphenyl)ethynyl)oxazolidin-2-one (1c)



¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.361 (d, *J* = 8.30 Hz, 2H), 7.137 (d, *J* = 8.45 Hz, 2H), 4.501 - 4.461 (dd, *J* = 8.15 Hz, *J* = 7.82 Hz, 2H), 4.022 - 3.982 (dd, *J* = 8.14 Hz, *J* = 7.84 Hz, 2H), 2.639 (q, *J* = 7.61 Hz, 2H), 1.223 (t, *J* = 7.62 Hz, 3H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 155.98, 144.70, 131.69, 127.87, 119.24, 78.28. 71.25, 63.02, 47.12, 28.77,

15.30.

3-((4-propylphenyl)ethynyl)oxazolidin-2-one (1d)



¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.353 (d, *J* = 8.19 Hz, 2H), 7.113 (d, *J* = 8.24 Hz, 2H), 4.496 - 4.456 (dd, *J* = 8.17 Hz, *J* = 7.79 Hz, 2H), 4.015 - 3.975 (dd, *J* = 8.15 Hz, *J* = 7.81 Hz, 2H), 2.572 (t, *J* = 7.78 Hz, 2H), 1.653 - 1.597 (m, 2H), 0.923 (t, *J* = 7.37 Hz, 3H) ; ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 156.00, 143.19, 131.62, 128.48, 119.24, 78.29, 71.27,

63.02, 47.11, 37.90, 24.29, 13.73.

3-((4-butylphenyl)ethynyl)oxazolidin-2-one (1e)



¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.350 (d, *J* = 8.19 Hz, 2H), 7.113 (d, *J* = 8.21 Hz, 2H), 4.491 - 4.451 (dd, *J* = 8.22 Hz, *J* = 7.78 Hz, 2H), 4.011 - 3.971 (dd, *J* = 8.15 Hz, *J* = 7.78 Hz, 2H), 2.593 (t, *J* = 7.85 Hz, 2H), 1.598 - 1.540 (m, 2H), 1.383 - 1.291 (m, 2H), 0.917 (t, *J* = 7.33 Hz, 3H) ; ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 155.00, 142.37, 130.57, 127.40,

118.18, 77.34, 70.17, 62.05, 46.08, 34.50, 32.32, 21.25, 12.88.

3-((4-(tert-butyl)phenyl)ethynyl)oxazolidin-2-one (1f)



¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.376 (d, *J* = 8.65 Hz, 2H), 7.321 (d, *J* = 8.64 Hz, 2H), 4.492 - 4.452 (dd, *J* = 8.19 Hz, *J* = 7.78 Hz, 2H), 4.015 - 3.975 (dd, *J* = 8.16 Hz, *J* = 7.80 Hz, 2H), 1.303 (s, 9H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 159.74, 156.05, 133.49, 114.77, 113.95, 77.59, 70.94, 62.97, 55.29, 47.14, 29.69.

3-((4-methoxyphenyl)ethynyl)oxazolidin-2-one (1g)



¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.387 (d, *J* = 8.81 Hz, 2H), 6.835 (d, *J* = 8.83 Hz, 2H), 4.495 - 4.455 (dd, *J* = 8.17 Hz, *J* = 7.78 Hz, 2H), 4.010 – 3.970 (dd, *J* = 8.19 Hz, *J* = 7.76 Hz, 2H), 3.809 (s, 3H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 159.73, 156.05, 133.49, 114.07, 113.95, 77.59, 70.93, 62.97, 55.29, 47.14, 29.69.

3-((4-ethoxyphenyl)ethynyl)oxazolidin-2-one (1h)



¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.370 (d, *J* = 8.86 Hz, 2H), 6.816 (d, *J* = 8.85 Hz, 2H), 4.490 - 4.450 (dd, *J* = 8.18 Hz, *J* = 7.79 Hz, 2H), 4.026 (q, *J* = 7.03 Hz, 2H), 4.003 - 3.964 (dd, *J* = 8.14 Hz, *J* = 7.82 Hz, 2H), 1.407 (t, *J* = 6.99 Hz, 3H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 159.13, 156.07, 133.49, 114.45, 113.85, 77.53, 70.98, 63.50, 62.98, 47.14, 14.73.

3-((4-fluorophenyl)ethynyl)oxazolidin-2-one (1i)



¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.449 – 7.399 (m, 2H), 7.032 – 6.975 (m, 2H), 4.512 - 4.473 (dd, *J* = 8.15 Hz, *J* = 7.81 Hz, 2H), 4.023 – 3.983 (dd, *J* = 8.91 Hz, *J* = 7.84 Hz, 2H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 163.51, 161.53, 155.94, 133.71, 133.64, 118.21, 118.19, 115.70, 115.53, 78.61, 70.16, 63.09, 47.01.

3-((4-chlorophenyl)ethynyl)oxazolidin-2-one (1j)



¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.364 (d, *J* = 8.72 Hz, 2H), 7.280 (d, *J* = 8.72 Hz, 2H), 4.516 - 4.476 (dd, *J* = 8.13 Hz, *J* = 7.82 Hz, 2H), 4.028 - 3.988 (dd, *J* = 8.12 Hz, *J* = 7.82 Hz, 2H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 155.81, 138.22, 132.77, 128.67, 120.71, 79.83, 70.27, 63.10, 46.97.

3-((4-chlorophenyl)ethynyl)oxazolidin-2-one (1k)



¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.439 (d, *J* = 8.55 Hz, 2H), 7.293 (d, *J* = 8.47 Hz, 2H), 4.512 - 4.480 (dd, *J* = 8.16 Hz, *J* = 7.79 Hz, 2H), 4.024 – 3.992 (dd, *J* = 8.14 Hz, *J* = 7.83 Hz, 2H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 155.76, 132.94, 131.59, 122.40, 121.19, 79.99, 70.39, 63.09, 46.96.

3-([1,1'-biphenyl]-4-ylethynyl)oxazolidin-2-one (11)



¹H NMR (400MHz, CDCl₃) δ (ppm): 7.583 (d, *J* = 7.58 Hz, 2H), 7.548 (d, *J* = 8.84 Hz, 2H), 7.507 (d, *J* = 8.21 Hz, 2H), 7.439 (t, *J* = 7.44 Hz, 2H), 7.351 (d, *J* = 7.31 Hz, 2H), 4.495 (t, *J* = 8.87 Hz, 2H), 4.025 (t, *J* = 8.50 Hz, 2H); ¹³C NMR (101MHz, CDCl₃) δ (ppm): 155.89, 140.92, 140.27, 131.98, 128.86, 127.64, 126.99, 121.98, 121.08, 79.57, 71.16, 63.06,

47.09.

3-((4-(trifluoromethyl)phenyl)ethynyl)oxazolidin-2-one (1m)



¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.563 (d, *J* = 8.50 Hz, 2H), 7.522 (d, *J* = 8.44 Hz, 2H), 4.536 - 4.496 (dd, *J* = 8.16 Hz, *J* = 7.75 Hz, 2H), 4.058 - 4.018 (dd, *J* = 8.17 Hz, *J* = 7.76 Hz, 2H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 155.63, 131.36, 129.88, 129.56, 126.20, 125.30, 125.27, 125.23, 125.19, 122.56, 81.35, 70.45, 63.15, 46.90.

methyl 4-((2-oxooxazolidin-3-yl)ethynyl)benzoate (1n)



¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.977 (d, *J* = 8.57 Hz, 2H), 7.481 (d, *J* = 8.61 Hz, 2H), 4.534 - 4.494 (dd, *J* = 8.15 Hz, *J* = 7.77 Hz, 2H), 4.056 - 4.017 (dd, *J* = 8.16 Hz, *J* = 7.76 Hz, 2H), 3.915 (s, 3H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 166.53, 155.63, 131.00, 129.48, 129.27, 127.09, 81.92, 71.07, 63.14, 52.20, 46.93.

3-(thiophen-3-ylethynyl)oxazolidin-2-one (10)



¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.459 (dd, J = 2.98 Hz, J = 1.10 Hz, 1H), 7.263 (dd, J = 4.35 Hz, J = 1.28 Hz, 1H), 7.115 (dd, J = 4.98 Hz, J = 1.08 Hz, 1H), 4.504 – 4.464 (dd, J = 8.19 Hz, J = 7.77 Hz, 2H), 4.015 - 3.975 (dd, J = 8.20 Hz, J = 7.75 Hz, 2H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 156.02, 130.08, 129.18, 125.33, 120.90, 78.37, 66.40, 63.10, 47.03.

3-((3-fluorophenyl)ethynyl)oxazolidin-2-one (1p)



¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.296 – 7.256 (m, 1H), 7.226 – 7.200 (m, 1H), 7.145 – 7.110 (m, 1H), 7.033 – 6.982 (m, 1H), 4.520 – 4.481 (dd, *J* = 8.17 Hz, *J* = 7.75 Hz, 2H), 4.036 - 3.996 (dd, *J* = 8.16 Hz, *J* = 7.77 Hz, 2H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 163.55, 161.10, 155.78, 129.95, 129.86, 127.32, 127.29, 124.14, 124.04, 118.28, 118.06, 115.56, 115.35, 79.88, 70.29,

70.26, 63.15, 46.94.

3-((3-chlorophenyl)ethynyl)oxazolidin-2-one (1q)



¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.428 – 7.419 (m, 1H), 7.327 – 7.301 (m, 1H), 7.293 – 7.268 (m, 1H), 7.251 – 7.212 (m, 1H), 4.520 – 4.480 (dd, *J* = 8.13 Hz, *J* = 7.80 Hz, 2H), 4.032 - 3.992 (dd, *J* = 8.13 Hz, *J* = 7.81 Hz, 2H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 155.75, 134.12, 131.20, 129.57, 129.52, 128.38, 123.98, 80.15, 70.13, 63.16, 46.94.

3-(m-tolylethynyl)oxazolidin-2-one (1r)



¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.271 – 7.233 (m, 2H), 7.190 (t, *J* = 7.61 Hz, 1H), 7.114 (d, *J* = 7.57 Hz, 1H), 4.013 – 3.981 (dd, *J* = 8.19 Hz, *J* = 7.76 Hz, 2H), 4.036 - 3.996 (dd, *J* = 8.12 Hz, *J* = 7.83 Hz, 2H), 2.318 (s, 3H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 163.55, 161.10, 155.93, 138.00, 132.15, 129.09, 128.58, 128.20, 121.94, 118.28, 118.06, 78.62, 71.38, 63.02, 47.07,

21.19.

3-((3,4-dichlorophenyl)ethynyl)oxazolidin-2-one (1r)



¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.516 (d, *J* = 1.38 Hz, 1H), 7.375 (d, *J* = 8.28 Hz, 1H), 7.250 (d, *J* = 8.58 Hz, *J* = 1.50 Hz, 1H), 4.507 (t, *J* = 8.19 Hz, 1H), 4.012 (t, *J* = 8.18 Hz, 1H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 155.67, 132.92, 132.50, 132.45, 130.51, 130.37, 122.27, 80.83, 69.38, 63.20, 46.88.

3-((4-ethylphenyl)ethynyl)-4-phenyloxazolidin-2-one (1t)



¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.483 – 7.382 (m, 5H), 7.265 - 7.206 (m, 5H), 5.143 (dd, J = 8.62 Hz, J = 1.47 Hz, 1H), 4.776 (t, J = 8.87 Hz, 1H), 4.306 (dd, J = 8.96 Hz, J = 1.84 Hz, 1H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 155.58, 136.05, 131.48, 129.56, 129.35, 128.16, 128.07, 126.91, 122.12, 78.02, 70.78, 62.24.

4-phenyl-3-(p-tolylethynyl)oxazolidin-2-one (1u)



¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.475 – 7.380 (m, 5H), 7.153 (d, *J* = 8.10 Hz, 2H), 7.025 (d, *J* = 7.93 Hz, 2H), 5.146 – 5.106 (dd, *J* = 8.57 Hz, *J* = 7.15 Hz, 1H), 4.773 (t, *J* = 8.85 Hz, 1H), 4.324 – 4.284 (dd, *J* = 8.93 Hz, *J* = 7.09 Hz, 1H), 2.292 (s, 3H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 155.63, 138.25, 136.13, 131.52, 129.49, 129.31, 128.90, 126.89, 118.97, 77.22, 72.87,

70.71, 62.26, 21.41.

3-((4-ethylphenyl)ethynyl)-4-phenyloxazolidin-2-one (1v)



¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.480 – 7.380 (m, 5H), 7.180 (d, *J* = 8.25 Hz, 2H), 7.051 (d, *J* = 8.42 Hz, 2H), 5.128 (dd, *J* = 8.67 Hz, *J* = 1.53 Hz, 1H), 4.773 (t, *J* = 8.87 Hz, 1H), 4.305 (dd, *J* = 8.94 Hz, *J* = 1.85 Hz, 1H), 2.587 (t, *J* = 7.60 Hz, 2H), 1.177 (t, *J* = 7.60 Hz, 3H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 155.63, 144.57, 136.14, 131.60, 129.30, 127.72, 126.90, 119.22, 77.25,

72.90, 70.72, 62.26, 28.74, 15.32.

3-((4-chlorophenyl)ethynyl)-4-phenyloxazolidin-2-one (1w)



¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.473 – 7.379 (m, 5H), 7.173 (d, *J* = 8.24 Hz, 2H), 7.027 (d, *J* = 8.31 Hz, 2H), 5.145 – 5.106 (dd, *J* = 8.60 Hz, *J* = 7.12 Hz, 1H), 4.771 (t, *J* = 8.87 Hz, 1H), 4.321 – 4.281 (dd, *J* = 8.91 Hz, *J* = 7.11 Hz, 1H), 2.521 (t, *J* = 7.83 Hz, 2H), 1.623 (m, 2H), 0.888 (t, *J* = 7.35 Hz, 3H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 155.64, 143.03, 136.15, 131.52, 129.48,

 $129.30,\,128.32,\,126.90,\,119.22,\,72.90,\,70.72,\,62.26,\,37.87,\,24.29,\,13.69.$

3-((4-(tert-butyl)phenyl)ethynyl)-4-phenyloxazolidin-2-one (1y)



¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.454 – 7.381 (m, 5H), 7.244 (d, *J* = 8.67 Hz, 2H), 7.192 (d, *J* = 8.67 Hz, 2H), 5.128 (dd, *J* = 8.60 Hz, *J* = 1.40 Hz, 1H), 4.772 (t, *J* = 8.84 Hz, 1H), 4.305 (dd, *J* = 8.96 Hz, *J* = 1.81 Hz, 1H), 1.259 (s, 9H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 155.60, 151.40, 136.15, 131.32, 129.48, 129.29, 126.92, 125.16, 119.07, 77.42, 72.88, 70.71, 62.29, 34.70,

31.13.

3-((4-chlorophenyl)ethynyl)-4-phenyloxazolidin-2-one (1z)



¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.492 – 7.378 (m, 5H), 7.196 (d, *J* = 8.91 Hz, 2H), 7.165 (d, *J* = 8.98 Hz, 2H), 5.158 – 5.118 (dd, *J* = 8.51 Hz, *J* = 7.21 Hz, 1H), 4.789 (t, *J* = 8.90 Hz, 1H), 4.338 – 4.297 (dd, *J* = 8.92 Hz, *J* = 7.14 Hz, 1H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 135.92, 134.07, 132.65, 129.63, 129.38, 128.50, 126.87, 120.64, 78.91, 71.90, 70.81, 62.19, 29.70.

3-((3-fluorophenyl)ethynyl)-4-phenyloxazolidin-2-one (1a')



¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.498 – 7.383 (m, 5H), 7.041 - 7.016 (m, 1H), 6.963 - 6.913 (m, 2H), 5.143 (dd, *J* = 8.57 Hz, *J* = 1.35 Hz, 1H), 4.792 (t, *J* = 8.81 Hz, 1H), 4.327 (dd, *J* = 8.96 Hz, *J* = 1.83 Hz, 1H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 163.43, 160.99, 155.39, 135.87, 129.78, 129.70, 129.66, 129.40, 127.21, 127.18, 126.89, 124.07, 123.97, 118.19,

117.96, 115.42, 115.21, 78.97, 71.94, 71.90, 70.84, 62.18.

3-((3,4-dichlorophenyl)ethynyl)-4-phenyloxazolidin-2-one (1b')



¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.491 – 7.444 (m, 3H), 7.387 (d, *J* = 6.65 Hz, 2H), 7.311 (d, *J* = 1.61 Hz, 1H), 7.277 (t, *J* = 8.61 Hz, 1H), 7.078 – 7.058 (dd, *J* = 8.36 Hz, *J* = 1.72 Hz, 1H), 5.141 (t, *J* = 8.15 Hz, 1H), 4.798 (t, *J* = 8.86 Hz, 1H), 4.343 – 4.311 (dd, *J* = 8.85 Hz, *J* = 1.58 Hz, 1H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 154.27, 134.71, 131.79, 131.30, 129. 41, 129.16, 145 R2, 141 42, 78 88, 60 R2, 60 R2, 61 R0

 $128.70,\,128.41,\,125.82,\,121.13,\,78.88,\,69.92,\,69.86,\,61.09.$

3-(hept-1-yn-1-yl)oxazolidin-2-one (3a)



¹H NMR (400 MHz, CDCl₃) δ (ppm): 4.429 – 2.389 (dd, *J* = 8.17 Hz, *J* = 7.83 Hz, 2H), 3.891 - 3.851 (dd, *J* = 8.16 Hz, *J* = 7.85 Hz, 2H), 2.297 (t, *J* = 7.20 Hz, 2H), 1.568 – 1.496 (m, 2H), 1.403 - 1.292 (m, 4H), 0.900 (t, *J* = 7.23 Hz, 3H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 156.65, 72.66, 70.00, 62.77, 47.06, 31.03, 28.49, 22.17, 18.3, 13.94.

N,4-dimethyl-N-(phenylethynyl)benzenesulfonamide (5c)



¹H NMR (400 MHz, CDCl₃) δ (ppm): 7.839 (d, *J* = 8.16 Hz, 2H), 7.378 – 7.347 (m, 4H), 7.305 – 7.255 (m, 3H), 3.150 (s, 3H), 2.457 (s, 3H); ¹³C NMR (100 MHz, CDCl₃) δ (ppm): 144.83, 133.24, 131.42, 129.83, 128.29, 127.88, 122.71, 83.96, 69.04, 39.33, 21.68.

18. Copies of NMR Spectra:



 $^{\rm 13}C$ NMR (100 MHz, CDCl_3) spectrum of ${\bf 2a}$



¹³C NMR (100 MHz, CDCl₃) spectrum of **2b**



¹³C NMR (100 MHz, CDCl₃) spectrum of **2c**



¹³C NMR (100 MHz, CDCl₃) spectrum of **2d**



¹³C NMR (100 MHz, CDCl₃) spectrum of **2e**



 $^{\rm 13}{\rm C}$ NMR (100 MHz, CDCl₃) spectrum of 2f



 ^{13}C NMR (100 MHz, CDCl₃) spectrum of 2g



¹³C NMR (100 MHz, CDCl₃) spectrum of **2h**



¹³C NMR (100 MHz, CDCl₃) spectrum of **2i**



¹³C NMR (100 MHz, CDCl₃) spectrum of **2j**



¹³C NMR (100 MHz, CDCl₃) spectrum of **2k**



 $^{\rm 13}{\rm C}$ NMR (100 MHz, CDCl₃) spectrum of 2l



¹³C NMR (100 MHz, CDCl₃) spectrum of **2m**



¹³C NMR (100 MHz, CDCl₃) spectrum of **2n**



¹³C NMR (100 MHz, CDCl₃) spectrum of **20**



¹³C NMR (100 MHz, CDCl₃) spectrum of **2p**



¹³C NMR (100 MHz, CDCl₃) spectrum of **2q**



¹³C NMR (100 MHz, CDCl₃) spectrum of **2r**





¹³C NMR (100 MHz, CDCl₃) spectrum of **2s**


 $^{\rm 13}{\rm C}$ NMR (100 MHz, CDCl₃) spectrum of 2t



¹³C NMR (100 MHz, CDCl₃) spectrum of **2u**



¹³C NMR (100 MHz, CDCl₃) spectrum of A **2v**



¹³C NMR (100 MHz, CDCl₃) spectrum of **2w**





¹³C NMR (100 MHz, CDCl₃) spectrum of **2x**



¹³C NMR (100 MHz, CDCl₃) spectrum of **2y**



¹³C NMR (100 MHz, CDCl₃) spectrum of **2z**



¹³C NMR (100 MHz, CDCl₃) spectrum of **2a'**



 ^{13}C NMR (100 MHz, CDCl_3) spectrum of 2b'

120 110 100

90 80 70

60

50

40 30

10 0

ppm

20

180

a table light shall be also be

170 160 150 140 130



¹³C NMR (100 MHz, CDCl₃) spectrum of **1a**

90 80 70 60 50

110 100

40

30 20 10

ppm

180

170

160 150 140 130 120



 ^{13}C NMR (100 MHz, CDCl_3) spectrum of 1b



 ^{13}C NMR (100 MHz, CDCl₃) spectrum of 1c



 ^{13}C NMR (100 MHz, CDCl₃) spectrum of 1d



¹³C NMR (100 MHz, CDCl₃) spectrum of **1e**



210 200 190 180 170 160 150 140 130 120 110 100 90 80 70 60 50 40 30 20 10 0 ppm

 $^{\rm 13}{\rm C}$ NMR (100 MHz, CDCl₃) spectrum of ${\rm 1f}$



 ^{13}C NMR (100 MHz, CDCl_3) spectrum of 1g



 ^{13}C NMR (100 MHz, CDCl_3) spectrum of 1h



 $^{\rm 13}{\rm C}$ NMR (100 MHz, CDCl₃) spectrum of 1i



 $^{\rm 13}{\rm C}$ NMR (100 MHz, CDCl₃) spectrum of 1j



 ^{13}C NMR (100 MHz, CDCl_3) spectrum of 1k



 ^{13}C NMR (100 MHz, CDCl₃) spectrum of 1I



 ^{13}C NMR (100 MHz, CDCl_3) spectrum of 1m



 ^{13}C NMR (100 MHz, CDCl₃) spectrum of 1n



 ^{13}C NMR (100 MHz, CDCl₃) spectrum of 10



 ^{13}C NMR (100 MHz, CDCl₃) spectrum of 1p



 ^{13}C NMR (100 MHz, CDCl₃) spectrum of 1q



 ^{13}C NMR (100 MHz, CDCl₃) spectrum of 1r



 $^{\rm 13}{\rm C}$ NMR (100 MHz, CDCl₃) spectrum of ${\rm 1s}$



¹³C NMR (100 MHz, CDCl₃) spectrum of **1t**



 ^{13}C NMR (100 MHz, CDCl_3) spectrum of 1u



¹³C NMR (100 MHz, CDCl₃) spectrum of **1v**



¹³C NMR (100 MHz, CDCl₃) spectrum of **1w**



 ^{13}C NMR (100 MHz, CDCl₃) spectrum of 1y



 ^{13}C NMR (100 MHz, CDCl₃) spectrum of 1z



¹³C NMR (100 MHz, CDCl₃) spectrum of **1a'**



 ^{13}C NMR (100 MHz, CDCl_3) spectrum of 1b'


¹³C NMR (100 MHz, CDCl₃) spectrum of **5a**



 ^{13}C NMR (100 MHz, CDCl₃) spectrum of 5c