## 3-Benzyl-4-(N-benzylcarbamoylmethyl)-2-(3-pyridyl)-1,3-oxazolidine

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The title compound was synthesized by the reaction of N-benzyl-3-benzylamino-4-hydroxybutanamide with 3-pyridinecarboxaldehyde using the procedure described in [1]. A mixture of N-benzyl-3-benzylamino-4-hydroxybutanamide ( $2.98 \mathrm{~g}, 0.01 \mathrm{~mol}$ ), 3-pyridinecarboxaldehyde (nicotinaldehyde) $(1.02 \mathrm{~g}, 0.01 \mathrm{~mol})$, molecular sieves ( $\mathrm{Na}-\mathrm{A}, 20 \mathrm{~g}$ ), p-TsOH ( 0.02 g ) and dry chloroform ( 50 ml ) was heated to reflux for 4 h . The reaction mixture was then filtered, and the solvent was evaporated. The residue was dissolved in ethyl acetate ( 10 ml ) and cooled to $0^{\circ} \mathrm{C}$ for crystallization of desired product. The crystals obtained was separated with filtration and recrystallized from ethanol to yield $3.33 \mathrm{~g}(86 \%)$ of 3-benzyl-4-(N-benzylcarbamoylmethyl)-2-(3-pyridyl)-1,3-oxazolidine.
M.p. $93{ }^{\circ} \mathrm{C}$ (ethanol)

IR (vaseline oil, $\mathrm{cm}^{-1}$ ): $3310(\mathrm{~N}-\mathrm{H}) ; 1650(\mathrm{C}=\mathrm{O}) ; 1550(\mathrm{~N}-\mathrm{H}) ; 1605(\mathrm{C}=\mathrm{C})$.
${ }^{1} \mathrm{H}$ NMR ( $\mathrm{CDCl}_{3}, 250 \mathrm{MHz}$ ): 2.25, 2.31 (dd, dd, $2 \mathrm{H}, \mathrm{CH}_{2} \mathrm{CO}, \mathrm{J}=16.0 \mathrm{~Hz}$ ); 2.41, 2.51 (d, d, 2H, NCH2, J $=14.0 \mathrm{~Hz}) ; 3.5(\mathrm{~m}, 1 \mathrm{H}, \mathrm{NCH}) ; 4.07,4.26\left(\mathrm{dd}, \mathrm{dd}, 2 \mathrm{H}, \mathrm{OCH}_{2}\right) ; 6.23(\mathrm{broad} \mathrm{s}, 1 \mathrm{H}, \mathrm{NH}) ; 7.18(\mathrm{~m}, 11 \mathrm{H}, \mathrm{Ph}$, 5-H Py); 7.72 (dd, 1H, 6-H Py); 8.20 (dd, dd, 2H, NHCH $\}$ \}, 8,50 (dd, 2H, 2,4-H Py).

Anal. calcd. for $\mathrm{C}_{24} \mathrm{H}_{25} \mathrm{~N}_{3} \mathrm{O}_{2}$ (387.48): C 74.51; H 6.62; N 10.78; Found: C 74.39; H 6.50; N 10.85 .

## References:

1. Tlekhusezh, M.A.; Badovskaya, L.A.; Tyukhteneva, Z.I. Khimiya geterotsikl. Soedin. (Chemistry of Hetrocyclic compounds, Russia) 1996, 5, 711-716.

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