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(57) Abstract :
The present invention relates the synthesis of cyclic octapeptide, Cyclogossine-B which is previously isolated from latex of Jatropha gossypifolia, accomplished through coupling of tetra-peptide fragment (Boc-L-Gly-L-Gly-L-Trp-L-Leu-OMe) (72) with another tetra-peptide fragment (Boc-L-Ala-L-Ala-L-Ile-L-Leu-OMe) (73), followed by cyclization of the linear octa-peptide unit under alkaline condition. The formation of newly synthesized cyclic compound was confirmed by means of spectral techniques including FT-IR, 1H-NMR, Mass spectroscopy along with elemental analyses. Cyclogossine-B was subjected for biological screening to evaluate antimicrobial and anticancer activities. The anti-bacterial activity was carried out by using Gram +ve bacteria (B. subtilis, S. epidermidis) and Gram -ve bacteria (E. coli, P. aeruginosa, S. aureus and K. pneumonia). The anti-fungal activity was performed by using fungal strains like C. albicans, A. niger, T. mentagrophytes and M. audouinii. Similarly, the cytotoxic activity of synthesized cyclic peptide was carried out through MTT assay using Doxorubicin as standard drug on HCT116 and B16F10 cell lines. The cytotoxic effect was evaluated by determining the percentage inhibition of growth of HCT116 and B16F10 cell-lines. Then CTC50 (Concentration of test drug needed to inhibit cell growth by 50%) values were calculated by graphical extrapolation method. Different concentration of test, control and standard drug (120-7.5 µg/mL) were used for the cytotoxicity study. It was observed that this cyclic peptide exhibited significant antimicrobial and cytotoxic activity against cancer cell lines.

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