# AWARENESS OF VINCA ALKALOIDS AMONG DENTAL STUDENTS

## Nithyanandham Masilamani<sup>1</sup>, Dhanraj Ganapathy<sup>2</sup>

#### Abstract

Vincristine along with vinblastine are dual indole-based conjugated Vinca alkaloids formed from foliage of the herb Catharanthusroseus, traditionally known as Vincarosea vincristine, that have been effectively prescribed as single treatment and also in conjunction with other medications in hematological and stable malignancies chemotherapy for tumors. The purpose of this survey was to assess awareness of medical use of vinca alkaloids among dental undergraduate students. A cross-section study was performed with a self-directed survey questionnaire containing 10 queries distributed among 100 dental students. The questionnaire assessed the awareness about vinca alkaloids, their medicinal uses, anticancer activity, mechanism of action and side effects. The responses were recorded and analysed.94% of the respondents were not aware of medical uses of vinca alkaloids.95% were not aware of anticancer activity of vinca alkaloids.97% were not aware of the mechanisms of action of vinca alkaloids. Again 97% were not aware of side effects of the vinca alkaloids. This study concluded the awareness about the medical use of vinca alkaloids among dental students was poor. Majority of them are not aware of the anticancer activity of vinca alkaloids.

Keywords: Awareness, vincaalkaloids, dental students

#### Introduction

Vincristine along with vinblastine are dual indole-based conjugated Vinca alkaloids formed from foliage of the herb Catharanthusroseus, traditionally known as Vincarosea vincristine, that have been effectively prescribed as single treatment and also in conjunction with other medications in hematogenic and stable malignancies chemotherapy for tumors. They restrain Microtubulin (MT) by preventing the functionalization of tubulin. In the process of cell proliferation, they serve as antagonists mostly during metaphase of the cellular phase and therefore by allowing microtubules to impede the progress of the mitotic spindle. In cancerous cells, these medicines interfere with the DNA fixing and RNA-combination of DNA-subordinated RNA polymerase[1-4].(Einhorn, 1977; Gigant et al., 2005; Rao et al., 1985; Williams et al., 1987)

<sup>&</sup>lt;sup>1</sup> Tutor, Department of Prosthodontics, Saveetha Dental College and Hospitals, Saveetha Institute of Medical and Technical Sciences, Chennai,India.

<sup>&</sup>lt;sup>2</sup> Corresponding Author: Professor &Head of Department, Department of Prosthodontics, Saveetha Dental College and Hospitals, Saveetha Institute of Medical and Technical Sciences, Chennai, India.

VLB is efficacious in the management of multiple malignancies like Hodgkin's disease, and VCR has been shown to be sufficient for acute lymphoblastic leukemia. Vinca alkaloids may induce leukopenia and neurotxicity. Oncolytic migration of these heterodimers alkaloids has been associated with the mitotic shaft hurting through mitotic capture. Such alkaloids also alter the speed of axoplasmic transportation by making an alteration in neurotubules. Cytotoxicity and fringe neuropathy can be a consequence of the critical mechanism of action of such alkaloids by virtue of tubulin.

Microtubules (MTs) play main role during the time spent with the mitosis, after which chromosomes of the cell are replicated and segregated to frame two identical subsets, allowing cellular division into two daughter cells. MTs are linked with support for cell form, cell motility, intracellular vessel and numerous additional cell ability. The MTs are made from tubulin, βheterodimers. Such structures undergo extraordinarily strong polymerization and depolymerization as a result of the transient expansion of the tubulin dimers at their extreme. Vinca alkaloids act by impedance with this dynamic balance, by either hindering tubulin polymerization and blocking MT dismantling, forestalls appropriate MT work and at last prompts cell death(Owellen et al., 1972; Wilson et al., 1970). Oral disease is one of the most harmful tumors and it can require chemotherapy. Thus it is basic the dental specialists and dental understudies know about the different chemotherapeutic agents, vinca alkaloids being one of them, utilized in treatment of malignancy The purpose of this survey was to assess awareness of medical use of vinca alkaloids among dental undergraduate students

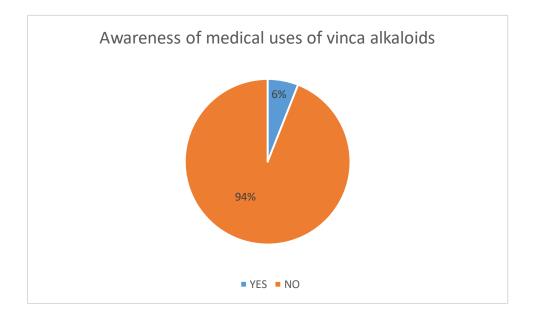
#### Materials and method

A cross-section study was performed with a self-directed survey questionnaire containing 10 queries distributed among 100 dental students. The questionnaire assessed the awareness about vinca alkaloids, their medicinal uses, anticancer activity, mechanism of action and side effects. The responses were recorded and analysed.

#### Results

94% of the respondents were not aware of medical uses of vinca alkaloids(Fig 1).95% were not aware of anticancer activity of vinca alkaloids (Fig 2).97 % were not aware of the mechanism of action of vinca alkaloids (Fig.3). Again 97% were not aware of side effects of vinca alkaloids (Fig.4).

## Fig.1:Awareness of medical uses of vinca alkaloids



## Fig.2:Awareness of anticancer activity of vinca alkaloids

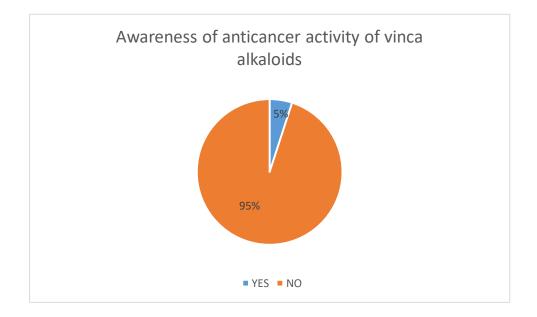
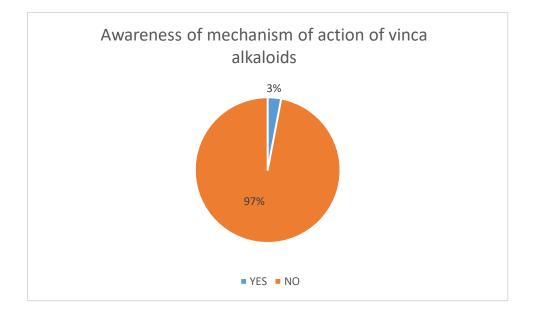
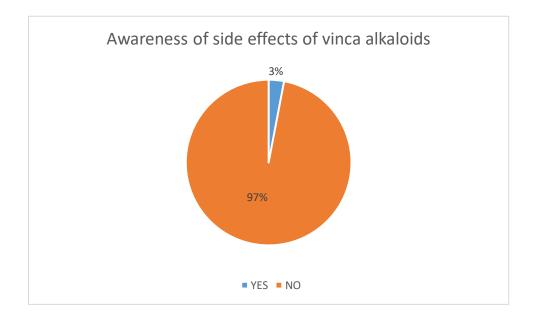


Fig.3:Awareness of mechanism of action of vinca alkaloids

International Journal of Psychosocial Rehabilitation, Vol. 23, Issue 05, 2019 ISSN: 1475-7192



## Fig.4:Awareness of side effects of vinca alkaloids



#### Discussion

Vinblastine is among the fewer tubulin-focused alkaloids responsible for many of the chemotherapeutic accomplishments after their introduction as anti-tumour medications. Vinblastine and the amino-terminal fragment of locally restrictive RB3-SLD shares a hydrophobic link on the a-tubulin substrate located at intermolecular interface in microtubules. It is an enticing target for drugs planned to antagonise microtubule components by interfacial impedance and tubulin appears to be suitable due to its ability to self-associate (Williams et al., 1987).

The antimitotic portion of high-fixed vinca alkaloids depolymerizes MTs and destabilizes mitotic shafts: the dividing disease cells tend to be impeded by dense chromosomal mitosis. At low concentrations, the mitosis becomes all the more unpretentious, and the cells knocked with apoptosis.Vinca alkaloids and various colchicine-restricting agents were shown to cause severe and rapid vascular disruption, contributing to the tumor necrosis. The vinca region includes both vinca sites, where severe inhibitors bind, and the local area, where non-competitive inhibitors bind. Vincristine ,vinblastine, maytansine, ansamitocines P-3 , P-4, rhizoxin and disorazole A1 bind at the vinca location and severely impede the ability to produce tubulin.

Vinblastin-23-oyl amino destructive substrate was introduced by linking the amino corrosive carboxyl esters to a vinblastin-23-oyl moiety via the amide linkage. A few vector amino acids were analyzed over four basic groups based on the extremity of the side chains. In addition, the effect of auxiliary modifications, for instance (1) the proximity of the amino-corrosive transporter to the C-23-oyl moiety, (2) the concept of the amino-corrosive carboxylic ester, (3) the distance of the amino-corrosive alkyl chain, (4) only the stereoisomericity of the amino-corrosive component, and (5) again the reacetylation of hydroxyl range (position 0-4) of the whole vindoline moiety.

All throughout years, numerous research groups have worked extensively and substantially to integrate new vinblastine and vincristine subordinates. Modifications in vindoline framework or the catharanthine moiety have resulted in a variety of new antitumor specialists medications with higher selectivity or less toxic properties. The method of action of Vinca alkaloids was evaluated using these novel subordinates and some significant new results were identified for the tubulin polymerization framework. The structure of these dimers is a continuous source of further discovery in this discipline of medicine and therapy. (Keglevich et al., 2012; Marantz et al., 1969; Venghateri et al., 2013).

This study inferred the dental students knowledge about vincaalkaolids is poor. They are not aware of the antitumour effectiveness of vinca alkaloids and they are unaware of the side effects of these compounds also. These modifications can further enhance the anticancer activity of these compounds. The researchers can further advance the development of newer vinca derivatives in the management of cancer.

#### Conclusion

This study concluded the awareness about the medical use of vinca alkaloids among dental students was poor. Majority of them are not aware of the anticancer activity of vinca alkaloids. Further rigorous continuing education programs and awareness-raising workshops are therefore required to enhance the knowledge and understanding of vinca alkaloids amongst dental students.

## References

- Einhorn, L. H. (1977). Cis-Diamminedichloroplatinum, Vinblastine, and Bleomycin Combination Chemotherapy in Disseminated Testicular Cancer. In *Annals of Internal Medicine* (Vol. 87, Issue 3, p. 293). https://doi.org/10.7326/0003-4819-87-3-293
- <u>Gigant, B., Wang, C., Ravelli, R. B. G., Roussi, F., Steinmetz, M. O., Curmi, P. A., Sobel, A., & Knossow,</u> <u>M. (2005). Structural basis for the regulation of tubulin by vinblastine. *Nature*, 435(7041), 519–522.
  </u>
- 3. <u>Keglevich, P., Hazai, L., Kalaus, G., &Szántay, C. (2012). Modifications on the basic skeletons of vinblastine and vincristine. *Molecules*, *17*(5), 5893–5914.</u>
- 4. <u>Marantz, R., Ventilla, M., & Shelanski, M. (1969). Vinblastine-Induced Precipitation of Microtubule</u> <u>Protein. In Science</u> (Vol. 165, Issue 3892, pp. 498–499). https://doi.org/10.1126/science.165.3892.498
- Owellen, R. J., Owens, A. H., & Donigian, D. W. (1972). The binding of vincristine, vinblastine and colchicine to tubulin. In *Biochemical and Biophysical Research Communications* (Vol. 47, Issue 4, pp. 685–691). https://doi.org/10.1016/0006-291x(72)90546-3
- Rao, K. S. P. B., Bhushana, K. S., Collard, M. P. M., Dejonghe, J. P. C., Atassi, G., Hannart, J. A., & <u>Trouet, A. (1985). Vinblastin-23-oyl amino acid derivatives: chemistry, physicochemical data, toxicity,</u> <u>and antitumor activities against P388 and L1210 leukemias. In *Journal of Medicinal Chemistry* (Vol. 28, <u>Issue 8, pp. 1079–1088). https://doi.org/10.1021/jm00146a017</u>
  </u>
- Venghateri, J. B., Gupta, T. K., Verma, P. J., Kunwar, A., & Panda, D. (2013). Ansamitocin P3 Depolymerizes Microtubules and Induces Apoptosis by Binding to Tubulin at the Vinblastine Site. In PLoS ONE (Vol. 8, Issue 10, p. e75182). https://doi.org/10.1371/journal.pone.0075182
- Williams, S. D., Birch, R., Einhorn, L. H., Irwin, L., Greco, F. A., &Loehrer, P. J. (1987). Treatment of disseminated germ-cell tumors with cisplatin, bleomycin, and either vinblastine or etoposide. *The New England Journal of Medicine*, 316(23), 1435–1440.
- Wilson, L., Bryan, J., Ruby, A., & Mazia, D. (1970). Precipitation of Proteins by Vinblastine and Calcium Ions. In *Proceedings of the National Academy of Sciences* (Vol. 66, Issue 3, pp. 807–814). https://doi.org/10.1073/pnas.66.3.807