

Original Research Article

Awareness of Human Papilloma Virus and Acceptability of Its Vaccination amongst Female Medical Students

Somsubhra De¹, Sachchithanantham Kanagasabai², Ankur Barua³

¹Associate Professor, ²Professor and HOD, Department of Obstetrics & Gynecology, Melaka Manipal Medical College, Melaka, Malaysia,

³Senior Lecturer, Department of Community Medicine, International Medical University, Kuala Lumpur, Malaysia.

Corresponding Author: Somsubhra De

Received: 02/07//2014

Revised: 25/07/2014

Accepted: 01/08/2014

ABSTRACT

Background: Human Papilloma Virus (HPV) is a known causative organism for most female genital cancers.

Objectives: (1) To study the level of awareness about HPV and the acceptability of HPV vaccine amongst female medical students.(2) To document any adverse effects of the vaccine in this population.

Methods: This was a questionnaire survey participated by female medical students who chose to have the vaccination, after voluntarily purchasing the vaccine. The questionnaire mainly focused on the students' awareness of HPV vaccine, her sources of information, acceptability and of adverse effects after taking the HPV vaccine. The participants were asked to recall any occurrence of side effects following any of the three doses administered. The adverse effects reported were tabulated and compared with the U.S. Department of Health and Human Services (DHHS) Vaccine Adverse Event Reporting System (VAERS) by Chi square using SPSS Version 16.0.0.

Results: The awareness of HPV amongst the medical students was 98% however the acceptability was 29%. This was mainly due to financial reasons and apprehensions about vaccine efficacy and safety. The adverse effects reported were namely redness and swelling at injection site, myalgia, headache, fever and they corroborated with VAERS data except pain at injection site which was lower {59% vs 92% (p<0.01)} and pruritus which was higher {19% vs 1% (p<0.001)}. No other serious adverse effects were notified.

Conclusion: There was a mismatch between awareness and acceptability. To bridge this gap and improve acceptability, more health education awareness should be given to both the parents and females who are eligible with the help of social media (television, radio, and internet) and also make provisions for subsidization of the cost of the vaccines. HPV vaccine is a safe and well tolerated amongst the many other study groups including our cohort.

Keywords: acceptability, side effects, bivalent, HPV, cancers, vaccine

INTRODUCTION

Human Papilloma virus (HPV) is an oncogenic virus which causes 5.2% of all

the cancers in the world with a prevalence of 40% in genital tract malignancies of women. ^[1] Apart from cervical cancer, HPV also

played a causative role in 60-65% of vaginal ^[2] and 20-50% of vulvar cancers. ^[3] The search for vaccine against HPV was initiated in the mid-80s and the first HPV vaccine was approved by FDA in 2006 which also became the first anti-cancer vaccine to be approved.^[4] This vaccine was a quadrivalent vaccine and it prevented the effects of HPV types 16, 18, 6 and 11. The second HPV vaccine to get FDA approval in 2009 was a bivalent vaccine^[5] which rendered protection against HPV types 16 and 18. A review of these vaccines from various studies showed were highly immunogenic, that they efficacious and devoid of adverse effects and its development in the field of immunotherapeutic science might prove to be pertinent method to combat cancer in future. ^[6] However the result on awareness of this vaccine in this study was disappointing. Jared Rosenberg in his studies over four countries (India, Uganda, Peru and Vietnam) showed that the awareness was also very poor in these countries. ^[7] In addition, the apprehension for these new vaccines was palpable as the clinical trials on long term side effects following its use over a larger population are still ongoing. To keep a track of the adverse effects, post market surveillance was done and the findings were noted in U.S. Department of Health and Human Services (DHHS) Vaccine Adverse Event Reporting System (VAERS).^[8]

In this study, the objectives were to assess the level of awareness of HPV and acceptability of this vaccine amongst female medical students to look for any adverse effects of the vaccine on these students who took the bivalent HPV vaccination. The occurrence of adverse effects in our study was then compared with the reporting in VAERS to analyze for significant variation.

MATERIALS AND METHODS Study design: Cross Sectional. *Subjects:* Female medical students from third to final years of a private medical school.

Study Duration: The study was done over a period of one year from December 2010 till December 2011.

Inclusion criteria: Only female students who gave consent to answering the questionnaire were chosen for the survey.

Exclusion criteria: Those who refuse to give consent and incomplete questionnaires.

Vaccine regimen: Bivalent HPV vaccine was given intramuscularly at 0, 1 and 6 months.

Study type: This was a questionnaire survey method. After obtaining valid consent, the questionnaires were distributed amongst the students and collected after they were duly filled. Anonymity of all students was maintained to facilitate the expression of their views freely. The study had no financial competing interest. The students themselves chose to have the vaccination, voluntarily after purchasing the vaccine. The author or co-authors had no role in administering the vaccines to these students. The questionnaire mainly focused on the students' level of awareness, acceptability and occurrence of adverse effects after taking the HPV vaccine. The student, who were aware of HPV and its vaccination, had to fill up the source of their information about HPV. In case they refused the vaccination, they had to state the reasons for their refusal. Within one month after the completion of their regimen of vaccination, they were asked to recall any occurrence of side effects following any of the three doses. The incidences of these adverse effects, if any, were noted and then compared with the U.S. Department of Health and Human Services (DHHS) Vaccine Adverse Event Reporting System (VAERS).A pilot study was conducted involving 20 students of the same batch three months before the main study. The acceptability, feasibility and time

of survey questionnaire management administration were assessed. An item analysis along with test-retest validity and intra-observer reliability (Cronbach's alpha) of the survey questionnaire were also assessed during this pilot study. The survey questionnaire was further refined with the help of two content experts and an English language professional. The overall intraobserver reliability from Cronbach's Alpha was found to be 0.822. The intra-observer reliability ranged between 0.734 and 0.806 for item deletion in Cronbach's Alpha analysis. Since, none of the Cronbach's Alpha values exceeded 0.822 in item deletion; the items all of survey questionnaire were included in the final analysis.

Ethical Considerations: Approval for the present study was obtained from the research and ethical committee of the Melaka-Manipal Medical College in Malaysia. The information obtained during the data collection was strictly kept confidential. order In to maintain anonymity, a random code number was issued to each participant of this study while responding to the questionnaire. Informed written consent was obtained from every participant prior to the inception of this study.

Data Analysis: The data collected was tabulated and analyzed by using the Statistical Package for Social Sciences (SPSS) version 16.0. The results were expressed in terms of Proportion. Chi-square Test was applied for comparison purpose. In this study, a *p*-value <0.05 was considered as statistically significant.

RESULTS

Total of 250 students took part in the study however after excluding the incomplete questionnaires only 224 were included for analysis. The mean age of the cohort was 23.4 years. The total number who was aware of HPV and its vaccine was 220 (98%). The source of their information about HPV is depicted in Figure 1.

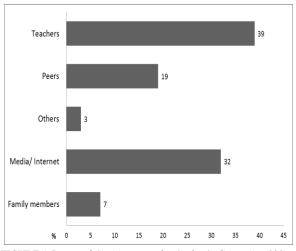


FIGURE 1:Source of the Awareness for the Study Group. (n= 220)

The total number of students who took the vaccination was 64 making the acceptability of vaccine as 29%. All the 64 students took the 3 doses as mentioned in the vaccine regimen and there were no drop outs. The remaining 160 (71%) students who refused the vaccinationstated their reasons for not accepting the vaccines which is shown in Figure 2.

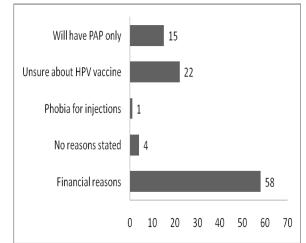


FIGURE 2: Reasons for not accepting HPV vaccine in the Study Group. (n= 160)

A total of 192 doses were given to 64 students. Some form of side effects were seen in 44 (69%) students. These side effects are divided into three groups namely common (>10%), moderately common (1-

10%) and non-specific in VAERS database. ^[9] However we have tabulated all the side effects along with VAERS data for the purpose of comparison as shown in Table 1.

| Side Effects | Study group (%) | nd its comparison with VAERS Data (%) | p |
|----------------------------|-----------------|---------------------------------------|---------|
| Fatigue | 30 | 55 | NS |
| Pain at injection site | 59 | 92 | <0.01 |
| Redness at injection site | 36 | 48 | NS |
| Swelling at injection site | 47 | 44 | NS |
| Myalgia | 45 | 49 | NS |
| Rashes | 7 | 7 | NS |
| Pruritus | 19 | 1 | < 0.001 |
| Vaginal infection | 8 | 3 | NS |
| GI symptoms | 20 | 28 | NS |
| Headache | 20 | 29 | NS |
| Fever | 17 | 13 | NS |
| NS= not significant | | | |

DISCUSSION

HPV has been shown to be linked in various genital tract malignancies. With newer diagnostic innovations, this link has been proven beyond doubt. The metaanalysis by Vuyst HD and colleagues showed that HPV was prevalent in all grades of vulvar (40%), vaginal (70%) and anal (80%) intraepithelial neoplasia and there was no variation of this prevalence around the geographical locations. ^[10] Recent study of HPV in cervical carcinoma by using multiplex real time polymerase chain reaction showed that HPV types 16 and 18 were prevalent in 86% of cases and in the remaining 14%, HPV type 73 was seen more commonly. ^[11] At present there are two types of HPV vaccines available for commercial use. They are quadrivalent (for HPV serotypes 6, 11, 16 and 18) and bivalent HPV vaccines (for HPV serotypes 16 and 18). These vaccines are recombinant vaccines targeting the E6 and E7 oncoproteins of the virus which were seen in the premalignant lesions caused by HPV.^[12] recombinant The L1 capsid protein assembled itself as Virus like Particle (VLP) and initiated immunological reaction and rendered protection against the infection

caused by the actual virus. ^[13] There was no viral DNA in the vaccine hence there was no chance of host acquiring the infection or reversion to HPV infection. However the main drawbacks of recombinant vaccine are the need for adjuvant to launch an immunological reaction and the need for multiple administrations of the vaccine.^[14] Bivalent HPV vaccine was used in our study and the main ingredients of the vaccine were HPV type 16 L1 protein (20 micrograms), HPV type 18 L1 protein (20 micrograms) along with ASO4 adjuvants 3-O-desacyl-4'monophosphoryl lipid A (MPL) (50 micrograms) and adsorbed on aluminium milligrams Al^{3+} in total). ^[15] (0.5)

In our study, the awareness of HPV and its vaccine stood out distinctively when compared with the general population as the chosen cohort was female medical students and a high rate of awareness was expected since they would become doctors and educate the general populationin future. The comparison of awareness of our cohort with the various studies (done by questionnaire surveys) all over the world is shown in Table 2.

| TABLE 2. Comparison of awareness of HFV between our conort and the general population | | | | |
|---|------------------|-----------------|---------------|--|
| RESEARCHERS | COUNTRY | SAMPLE SIZE (n) | AWARENESS (%) | |
| Feng S et al 2011 [16] | China | 1432 | 19 | |
| Pitts M et al 2009 ^[17] | Singapore | 2145 | 20 | |
| Ilter E et al 2010 ^[18] | Turkey | 525 | 44 | |
| Marlow L et al 2009 ^[19] | UK | 200 | 39 | |
| | (White British) | | | |
| Marlow L et al 2009 | UK | 750 | 18 | |
| | (Asian-Africans) | | | |
| Pitts MK et al 2007 ^[20] | Australia | 1100 | 51 | |
| STUDY GROUP | Malaysia | 224 | 98 | |
| (medical students) | | | | |
| | | | | |

TABLE 2. Comparison of awareness of HPV between our cohort and the general population

Our study also showed that 70% of the awareness was attributable to teachers and media/ internet. This suggests that it will be more productive to have health education on HPV and its vaccination at the level of school and colleges with regular reminders in the social media like television, radio and internet.

Only being aware of HPV will not reduce the burden of cancer. Accepting the HPV vaccine and taking the prescribed vaccine regimen is of prime importance for prevention of HPV related diseases. The acceptability of the vaccine was low when compared to the awareness of HPV in our study. Financial reason (58%) was cited as the main cause by the students. However one-fifth of the cohort did not accept the vaccine due to their apprehension about its efficacy, lack of information about the vaccine ("*still under trial*") and also the possible side effects of the vaccine. Almost identical findings were note in a study done by Fisher and his colleagues in Israel, where it was observed that the awareness of HPV was high but the acceptability was only 10%. ^[21] The reasons for low acceptability were similar to our study group. The comparison of acceptability of our cohort is with other studies is shown in Table 3.

| TABLE 5. Comparison of acceptability of HPV vaccine between our conort and general population | | | | |
|---|------------------|-----------------|-------------------|--|
| RESEARCHERS | COUNTRY | SAMPLE SIZE (n) | ACCEPTABILITY (%) | |
| Marlow L et al 2009 ^[19] | UK | 200 | 63 | |
| | (White British) | | | |
| Marlow L et al 2009 | UK | 750 | 20 | |
| | (Asian-Africans) | | | |
| I-Ching S et al 2008 ^[22] | Malaysia | 362 | 66 | |
| STUDY GROUP | Malaysia | 224 | 29 | |
| (medical students) | - | | | |

TABLE 3. Comparison of acceptability of HPV vaccine between our cohort and general population

To improve the acceptability of HPV vaccines, financial issues needs to be addressed and general population should be made aware of the researches done towards efficacy and safety profile of these vaccines. I-Ching S and coworkers suggested from their study that the acceptability can reach 97.8%, if the vaccine was made free of cost. ^[22] The vaccine taken by our cohorthas been studied by many researchers for its efficacy and awareness of these researches will help to alleviate the fears about the vaccines. Follow up studies has been done to prove its

efficacy from time to time. Four years back, Romanowski and his group showed sustained immunogenicity and efficacy of this vaccine for 6.4 years ^[23] followed by Carvalho and his group supporting the sustained efficacy till 7.3 years. ^[24] Most recently the efficacy was still maintained after 8.4 years of follow up. [25] Apart from the protection of the serotypes 16 and 18, researchers have demonstrated cross protection of the vaccine against other HPV serotypes. Wheeler and colleagues at the 4year-end analysis of PATRICIA trial

showed that the bivalent vaccine (HPV 16 & 18) was also efficacious against HPV 33, 31, 45 and 51. ^[26] Szarewski and his group added three more serotypes in the list of cross protection namely HPV 6, 11 and 74 for the same bivalent vaccine. ^[27] Studies also suggested that this bivalent vaccine was effective in women who had CIN3+ and adenocarcinoma in situ (AIS) irrespective of HPV DNA. ^[28]

HPV vaccines have been deemed safe and well tolerated and more than 100 million doses have been administered worldwide without any serious adverse [29] effect. A detailed post licensure monitoring for side effects was done in the US by VAERS from 2006 till 2013 and it was found that out of 57 million doses given there were reports of 22,000 adverse reactions. Around 92% of these reactions were limited to local reaction like pain, swelling and redness at the injection site and remaining 8% constituted headache, nausea, fatigue, syncope and generalized weakness. ^[30] There were no reports of disability, severe morbidity or mortality following vaccination. There have been concerns of Guillain Barre Syndrome following HPV vaccination but studies have actually shown there is no association HPV vaccination with this syndrome, ^[31] and also there were no association with autoimmune disease. disorders neurological and venous thromboembolism. ^[32,33] There was a case report of premature ovarian failure (POF) following HPV vaccination in a teenager ^[34] however cause-effect relationship was not established but an elimination of various etiologies of POF was undertaken and the vaccination was merely presumed to be the cause. These anecdotal reports make a long lasting impression in people's decision making, thereby affecting the acceptability of a vaccination program.

The side effects in our cohort were comparable to the VAERS data except for

pain at the injection site which was significantly lower and pruritus which was significantly higher. Lower incidence of pain at injection site could be attributed to higher pain threshold of the medical Stress causes lowering students. of testosterone and thereby lowering of pain threshold [35] and medical students are usually exposed to stress early in their academic career and by the time they come to clinical years there is adjustment with the stress of the medical profession and therefore 'pain coping' is better leading to higher pain threshold. ^[36] This could be a plausible explanation for the significant difference of incidence of pain at injection site. The increase incidence of pruritus is difficult to explain without the blood investigations to show allergenic reactions. However it is an established fact that aluminium based adjuvants can cause local hypersensitivity reaction in varying severity ^[37] hence we can only presume it to be an exaggerated hypersensitivity reaction.

CONCLUSION

In conclusion, the awareness of HPV and its related disease are well known amongst medical students. Despite having a high awareness, the acceptability is low due to financial reasons and apprehension about the efficacy and side effects of the relatively new vaccine. In our study, the sample size of vaccinated subjects was small hence robustic conclusions about side effects cannot be made. However we can conclude HPV vaccination did not cause any serious or disabling side effect in the vaccinated individuals and thereby is safe and tolerated well. This vaccine is still in its infancy and more studies, preferably with a bigger sample size, in different geographical locations and different cohorts will definitely enhance its propagation in future. Health education to parents by social media and subsidization of the vaccine cost would

go a long way in improving the acceptability for vaccination and reducing the cancer load for future generation.

ACKNOWLEDGEMENTS

I would like to thank Ms. Shazana for helping me to retrieve the full articles. I would also like to thank the class representatives Dr Vidyamalini and Dr Laxmi Priya for helping me with the questionnaire distribution and collections.

REFERENCES

- Parkin DM. The global health burden of infection-associated cancers in the year 2002.Int J Cancer.2006 Jun 15; 118(12):3030-44.
- Daling JR, Madeleine MM, Schwartz SM, et al. A population-based study of squamous cell vaginal cancer: HPV and cofactors. GynecolOncol 2002; 84:263– 70.
- Madeleine MM, Daling JR, Carter JJ, et al. Cofactors with human papillomavirus in a population-based study of vulvar cancer. J Natl Cancer Inst 1997; 89:1516–23.
- 4. http://usatoday30.usatoday.com/news/he alth/2006-06-08-cervical-cancervaccine_x.htm
- http://www.fda.gov/BiologicsBloodVac cines/Vaccines/ApprovedProducts/ucm1 86959.htm
- Somsubhra De and Sachchithanatham Kanagasabai. Human Papilloma Virus Vaccine – An update. European Journal of Scientific Research; Vol.43 No.2 (2010), pp. 256-64.
- Jared Rosenberg. Parents need HPV vaccine information. IntPerspect Sex Reprod Health. 2009; Dec; 35(4): 165.
- Barbara A Slade, Laura Leidel, ClaudiaVellozzi, et al. Postlicensure Safety Surveillance for Quadrivalent Human Papillomavirus Recombinant Vaccine. JAMA 2009 August; 302(7): 750-7.
- 9. http://www.uptodate.com/contents/huma n-papillomavirus-bivalent-types-16-18-

recombinant-vaccine-patient-druginformation?source=see_link

- Vuyst H D, Gary M. Clifford, Maria Claudia Nascimento, Margaret M. Madeleine and Silvia Franceschi. Prevalence and type distribution of human papillomavirus in carcinoma and intraepithelial neoplasia of the vulva, vagina and anus: A meta-analysis. Int. J. Cancer: 2009; 124, 1626–36.
- 11. Quddus MR, Manna P, Sung CJ, et al. Prevalence, distribution, and viral burden of all 15 high-risk human papillomavirus types in adenosquamous carcinoma of the uterine cervix: a multiplex real-time polymerase chain reaction-based study. Hum Pathol. 2013 Oct 1.pii: S0046-8177(13)00388-2. doi: 10.1016/j.humpath.2013.07.048.
- Z. He, A. P. Wlazlo, D. W. Kowalczyk, et al. Viral Recombinant Vaccines to the E6 and E7 Antigens of HPV-16. Virology 270, 146±161 (2000).
- 13. http://www.who.int/vaccines/en/olddocs /humanpapill.shtml
- S.M. Sivakumar, Mohammed M. Safhi, M. Kannadasan N. Sukumaran. Vaccine adjuvants – Current status and prospects on controlled release adjuvancity. Saudi Pharmaceutical Journal (2011) 19, 197– 206.
- 15. http://www.medicines.org.uk/emc/medic ine/20204/SPC/Cervarix/
- 16. Feng S, Xu X, Jin Y, Yao X. Women's Knowledge of Human Papillomavirus (HPV) and Their Attitudes Toward HPV Vaccine: Preparing for HPV Vaccination in China. Asia Pac J Public Health.2011 Jul 31.
- 17. Pitts M, Smith A, Croy S, et al. Singaporean women's knowledge of human papillomavirus (HPV) and attitudes toward HPV vaccination. Women Health. 2009 Jun;49(4):334-51.
- 18. Ilter E, Celik A, Haliloglu B, et al. Women's knowledge of Pap smear test and human papillomavirus: acceptance of HPV vaccination to themselves and their daughters in an Islamic society. Int

J Gynecol Cancer. 2010 Aug;20(6):1058-62.

- MarlowLaura A.V, Jane Wardle, Alice S Forster, Jo Waller. Ethnic differences in HPV awareness and vaccine acceptability. J Epidemiol Community Health2009; doi:10.1136/ jech.2008.085886.
- 20. Pitts MK, Dyson SJ, Rosenthal DA, Garland SM. Knowledge and awareness of human papillomavirus (HPV): attitudes towards HPV vaccination among a representative sample of women in Victoria, Australia. Sex Health. 2007 Sep;4(3):177-80.
- Fisher WA, Laniado H, Shoval H, Hakim M, Bornstein J. Barriers to human papilloma virus vaccine acceptability in Israel. Vaccine. 2013 Nov 22; 31 Suppl 8:I53-I57. doi: 10.1016/j. vaccine. 2013.06.107.
- I-Ching Sam, Li-Ping Wong, Sanjay Rampal, et al. Maternal Acceptance of Human Papillomavirus Vaccine in Malaysia. Journal of Adolescent Health 44 (2009) 610–12.
- 23. Romanowski B, de Borba PC, Naud PS, et al. GlaxoSmithKline Vaccine HPV-007 Study Group. Sustained efficacy and immunogenicity of the human papillomavirus (HPV)-16/18 AS04-adjuvantedvaccine: analysis of a randomised placebo-controlled trial up to 6.4 years. Lancet. 2009 Dec 12; 374 (9706):1975-85. doi: 10.1016/S0140-6736(09)61567-1.
- 24. Carvalho ND, J. Teixeirab, C.M. Roteli-Martinsc, et al. Sustained efficacy and immunogenicity of the HPV-16/18 AS04-adjuvanted vaccine up to 7.3 years in young adult women. Vaccine 28 (2010) 6247–55.
- 25. Roteli-Martins CM, Naud P, De Borba P, et al. Sustained immunogenicity and efficacy of the HPV-16/18 AS04-adjuvanted vaccine: up to 8.4 years of follow-up. Hum Vaccin Immunother. 2012 Mar; 8(3):390-7. doi: 10.4161/hv.18865. Epub 2012 Feb 13.

- 26. Wheeler CM, Castellsagué X, Garland SM, et al; HPV PATRICIA Study Group. Cross-protective efficacy of HPV-16/18 AS04-adjuvanted vaccine against cervical infection and precancer caused by non-vaccine oncogenic HPV types: 4-year end-ofstudy analysis of the randomised, double-blind PATRICIA trial. Lancet Oncol. 2012 Jan; 13(1):100-10. doi: 10.1016/S1470-2045(11)70287-X. Epub 2011 Nov 8.
- 27. Szarewski A, S. Rachel Skinner, Suzanne M. Garland, et al. Efficacy of the HPV-16/18 AS04-Adjuvanted Vaccine against Low-Risk HPV Types (PATRICIA Randomized Trial): An Unexpected Observation. J Infect Dis. 2013 November 1; 208(9): 1391–6. doi: 10.1093/infdis/jit360.
- 28. Lehtinen M, Paavonen J, Wheeler CM, et al, HPV PATRICIA Study Group. Overall efficacy of HPV-16/18 AS04-adjuvanted vaccine against grade 3 or greater cervical intraepithelial neoplasia: 4-year end-of-study analysis of the randomized, double-blind PATRICIA trial. Lancet Oncol. 2012 Jan; 13(1):89-99. doi: 10.1016/S1470-2045(11)70286-8. Epub 2011 Nov 8.
- 29. http://www.ncirs.edu.au/immunisation/f act-sheets/hpv-human-papillomavirus-fact-sheet.pdf
- 30. http://www.cdc.gov/mmwr/preview/mm wrhtml/mm6229a4.htm?s_cid=mm6229 a4_w
- 31. Ojha RP, Jackson BE, Tota JE, et al. Guillain-Barre syndrome following quadrivalent human papillomavirus vaccination among vaccine-eligible individuals in the United States. Hum VaccinImmunother. 2013 Sep 6; 10(1).
- 32. Arnheim-Dahlström L, Pasternak B, Svanström H, Sparén P, Hviid A. Autoimmune, neurological and venous thromboembolic adverse events after immunisation of adolescent girls with quadrivalent human papillomavirus vaccine in Denmark and Sweden: cohort

study. BMJ. 2013 Oct 9; 347:f5906. doi: 10.1136/bmj.f5906.

- 33. Grimaldi-Bensouda L, Guillemot D, Godeau B, et al; the PGRx-AID Study Group. Autoimmune disorders and quadrivalent human papillomavirus vaccination of young female subjects. J Intern Med. 2013 Nov 8. doi: 10.1111/joim.12155.
- 34. Little DT, Ward HR. Premature ovarian failure 3 years after menarche in a 16year-old girl following human papillomavirus vaccination. BMJ Case Rep. 2012 Sep 30; 2012. pii: bcr2012006879. doi: 10.1136/bcr-2012-006879.
- 35. Choi JC, Chung MI, Lee YD. Modulation of pain sensation by stressrelated testosterone and cortisol. Anaesthesia. 2012 Oct; 67(10):1146-51. doi: 10.1111/j.1365-2044.2012.07267.x. Epub 2012 Jul 16.
- 36. Bartley EJ, Fillingim RB. Sex differences in pain: a brief review of clinical and experimental findings. Br J Anaesth. 2013 Jul; 111(1):52-8. doi: 10.1093/bja/aet127
- Baylor, N.W., Egan, W., Richman, P., 2002. Aluminium salt in vaccines – US perspective. Vaccine 2002, S18–S23.

How to cite this article: Somsubhra De, Kanagasabai S, Barua A. Awareness of human papilloma virus and acceptability of its vaccination amongst female medical students. Int J Health Sci Res. 2014;4(9):82-90.

International Journal of Health Sciences & Research (IJHSR)

Publish your work in this journal

The International Journal of Health Sciences & Research is a multidisciplinary indexed open access double-blind peerreviewed international journal that publishes original research articles from all areas of health sciences and allied branches. This monthly journal is characterised by rapid publication of reviews, original research and case reports across all the fields of health sciences. The details of journal are available on its official website (www.ijhsr.org).

Submit your manuscript by email: editor.ijhsr@gmail.com OR editor.ijhsr@yahoo.com