

## **Supplemental Material to:**

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**Social media microblogs as an HPV vaccination forum**

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## **Supplement:**

### **HPV Vaccines Efficiency, Safety & Uptake**

The first cancer vaccine (quadrivalent HPV recombinant) was approved by the Food and Drug Administration (FDA) in 2006.<sup>1,2,4</sup> Currently there are two licensed HPV vaccines: Gardasil (Merck & Co., Whitehouse Station, NJ, USA) and Cervarix (GlaxoSmithKline, USA). Gardasil can protect 11 through 26 year-old females and 9 through 26 year-old males against multiple HPV types and genital warts while Cervarix targets type 16 and 18, recommended for females 10 through 25 years of age.<sup>2</sup> Although HPV is usually sexually transmitted,<sup>1</sup> some research shows that girls can also get infected even without sexual intercourse, thus, vaccination is recommended at a young age.<sup>10</sup>

Both vaccines are three-dose, intramuscular injection vaccines requiring 3 doses. Quadrivalent vaccine efficacy was established in two studies.<sup>4</sup> The Females United to Unilaterally Reduce Endo / Ectocervical Disease (FUTURE I) study assessed the incidence of genital warts, vulvar or cervical intraepithelial neoplasia (CIN) or cancer in a randomized, double-blind, placebo-controlled, international trial. High effectiveness in pre-cancerous prevention was shown in a study

of women aged 16-24 years old.<sup>4</sup> Another clinical trial of quadrivalent HPV vaccine confirmed high efficacy in preventing lower genital tract disease in women up to age 45 years.<sup>5</sup> Bivalent vaccine efficacy prevention against CIN (grade 2-3) was 90% in the mean follow-up period of 14.9 months in a phase III study among 18644 women of the right age and the vaccine efficacy.<sup>3</sup>

Although the vaccine is one of the most effective tools in disease prevention, it's not perfectly safe and effective.<sup>6,7</sup> "As with all US-administered vaccines, HPV vaccines were safety-tested before licensing and are continually monitored for safety and effectiveness."<sup>2</sup> It is necessary to monitor vaccine safety in order to detect rare reactions, to follow higher risk groups (i.e. the elderly and pregnant women etc.) and to maintain public confidence in vaccine.<sup>8</sup> The FDA monitors safety via three integrated elements: the Vaccine Adverse Event Reporting System (VAERS), the Vaccine Safety Datalink (VSD) Project, and the Clinical Immunization Safety Assessment (CISA) network.<sup>4,9</sup> Up to December 31st, 2008, 11,916 VAERS adverse events had been reported from more than 23 million doses of HPV vaccine distributed in the United States.<sup>4</sup> The majority (94%) of the

documented reports are not serious and there is no link between the vaccine and serious adverse events claims, according to CDC evaluation of VAERS data.<sup>4</sup>

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