Risk of new onset autoimmune disease in 9- to 25-year-old women exposed to human papillomavirus-16/18 AS04-adjuvanted vaccine in the United Kingdom

Corinne Willame, Dominique Rosillon, Julia Zima, Maria-Genalin Angelo, Anke L. Stuurman, Hilde Vroling, Rachael Boggon, Eveline M. Bunge, Manel Pladevall-Vila & Laurence Baril

To cite this article: Corinne Willame, Dominique Rosillon, Julia Zima, Maria-Genalin Angelo, Anke L. Stuurman, Hilde Vroling, Rachael Boggon, Eveline M. Bunge, Manel Pladevall-Vila & Laurence Baril (2016): Risk of new onset autoimmune disease in 9- to 25-year-old women exposed to human papillomavirus-16/18 AS04-adjuvanted vaccine in the United Kingdom, Human Vaccines & Immunotherapeutics, DOI: 10.1080/21645515.2016.1199308

To link to this article: http://dx.doi.org/10.1080/21645515.2016.1199308
Risk of new onset autoimmune disease in 9- to 25-year-old women exposed to human papillomavirus-16/18 AS04-adjuvanted vaccine in the United Kingdom

Corinne Willamea,1 Dominique Rosillona,b, Julia Zimab, Maria-Genalin Angeloa, Anke L. Stuurmanc, Hilde Vrolingc, Rachael Boggonda, Eveline M. Bungec, Manel Pladevall-Vila, and Laurence Barilb

aBusiness & Decision Life Sciences on behalf of GSK Vaccines, Avenue Fleming 20, B-1300 Wavre, Belgium; bGSK Vaccines, Wavre, Belgium; cPallas Health Research and Consultancy, Rotterdam, the Netherlands; dCPRD Research Group, Victoria, London, UK; eRTI Health Solutions, Barcelona, Spain; fCenter for Health Policy and Health Services Research, Henry Ford Health System, Detroit, MI, USA

ABSTRACT

To assess the risk of autoimmune disease (AD) in 9–25-year-old women within 1 year after the first AS04-HPV-16/18 vaccine dose, a retrospective, observational database cohort study was conducted using CPRD GOLD. From CPRD GOLD 4 cohorts (65,000 subjects each) were retrieved: 1 exposed female cohort (received ≥1 AS04-HPV-16/18 vaccine dose between Sep2008–Aug2010) and 3 unexposed cohorts: historical female (Sep2005–Aug2007), concurrent male, and historical male. Co-primary endpoints were confirmed neuroinflammatory/ophtalmic AD and other AD, secondary endpoints were confirmed individual AD. Risk of new onset of AD was compared between cohorts (reference: historical cohort) using Poisson regression. The main analysis using confirmed cases showed no neuroinflammatory/ophtalmic AD cases in the female exposed cohort. Incidence rate ratio (IRR) (95% CI) of other AD was 1.41 (0.86 to 2.31) in female and 1.77 (0.94 to 3.35) in male cohorts when compared to the female and male historical cohort, respectively. Secondary endpoints were evaluated for diseases with >10 cases, which were Crohn’s disease (IRR: 1.21 [0.37 to 3.95] for female and 4.22 [0.47 to 38.02] for male cohorts), autoimmune thyroiditis (IRR: 3.75 [1.25 to 11.31] for female and no confirmed cases for male cohorts) and type 1 diabetes (IRR: 0.30 [0.11 to 0.83] for female and 2.46 [1.08 to 5.60] for male cohorts). Analysis using confirmed and non-confirmed cases showed similar results, except for autoimmune thyroiditis in females, IRR: 1.45 (0.79 to 2.64). There was no evidence of an increased risk of AD in women aged 9 to 25 years after AS04-HPV-16/18 vaccination.

Introduction

Human papillomavirus (HPV) is the main cause of cervical cancer,1 of which approximately 70% is caused by types 16 and 18.2 Cervarix™ (AS04-HPV-16/18 vaccine) is a GSK Vaccines’ bivalent recombinant vaccine against HPV types 16 and 18. Efficacy and cross-protective efficacy of this AS04-HPV-16/18 vaccine against persistent infection, pre-cancerous lesions, and cervical cancers caused by oncogenic HPV was shown in the Papilloma Trial against Cancer In young Adults (PATRICIA) and, more recently in adult women from the Human Papilloma Virus: Vaccine Immunogenicity And Efficacy (VIVIANE) study.3,5

Generally, pre-licensure clinical studies provide key vaccine safety data, but their power to detect rare events such as new onset of autoimmune diseases (AD) is limited by their sample size, since incidence rates of different AD vary roughly from 1 to 50 per 100,000 person-years.6

The use of appropriate adjuvants can help to modulate optimally innate and adaptive immune responses following vaccination. However, the risk of developing an autoimmune response provoked by the adjuvant itself cannot be ruled out.7 The Center for Biologics Evaluation and Research in the United States (US) requested GSK to conduct a post-licensure study to investigate the risk of AD among AS04-HPV-16/18 vaccine recipients. A pooled safety analysis of data from 57,580 adolescent and adult females aged 9 years and above, of whom 33,339 received at least one dose of AS04-HPV-16/18 vaccine, showed that the rates of adverse events, medically significant conditions, serious adverse events, and potential immune-mediated disorders were similar between HPV and control groups.8

The aim of this observational cohort study using the Clinical Practice Research Datalink General Practice OnLine Database (CPRD GOLD) was to evaluate the risk of new onset of AD in women aged 9 to 25 years in the United Kingdom (UK) after administration of the AS04-HPV-16/18 vaccine (exposed cohort) and in controls of the same age (unexposed cohorts).
Results
From a total of 168,662 HPV vaccinated female subjects in CPRD, 103,081 (61.12%) were eligible for the exposed cohort. The number of eligible subjects in the other cohorts was: 107,434 for the unexposed historical female cohort, 142,772 for the concurrent male cohort and 92,337 for the historical male cohort. 65,000 Subjects were randomly selected from each of the cohorts but 42 subjects were excluded because a de-enrolment date (death date or date of lost to follow-up) occurred before the study start date.

Through the pre-defined algorithms 1,052 suspected AD cases were identified, of which 466 (44.3%) were identified as having confirmed or non-confirmed new onset AD after review of the individual subject profiles (Fig. 1). Among them, the date of first symptom was known for 384 (82.4%) cases, of which 40.4% (n = 155) were eligible for the main analysis, because their first symptom date and date of disease diagnosis were within the one year follow-up period. Out of these 155 AD cases, 109 (70.3%) were classified as confirmed cases and were included in the numerator in the main analysis (the 46 non-confirmed cases were excluded from the numerator and their person-time was included in the denominator). The number of cases included during sensitivity analyses can be found in Figure 1. A total of 68 confirmed cases from the exposed cohort were included for the self-controlled case-series (SCCS) analysis.

The overall population for the main analysis contained 259,876 subjects. Demographic and baseline characteristics of each cohort are depicted in Table 1. The female exposed cohort more frequently used healthcare, had more years of follow-up in CPRD GOLD at the study start date, and received more vaccines during the follow-up period than the historical female cohort. Similar differences existed between the 2 male cohorts, except that the male concurrent cohort received fewer vaccines in the year prior to the study start date than the male historical cohort.

Co-primary endpoints
In total, 3 confirmed cases of neuroinflammatory/ophthalmic AD and 106 confirmed cases of other AD were observed within the one year follow-up period (Table 2). There were no confirmed cases of neuroinflammatory/ophthalmic AD in the exposed female cohort, therefore the incidence rate ratio (IRR) could not be calculated. The corresponding age-adjusted IRR for male cohorts was 0.95 (95% confidence interval (CI): 0.06–15.18). For the other AD, the age-adjusted IRRs were 1.41 (95% CI: 0.86–2.31) for the female cohorts and 1.77 (95% CI: 0.94–3.35) for the male cohorts.

Sensitivity analysis using confirmed and non-confirmed cases showed results similar to the main analysis (Table 2), as all the sensitivity analyses using the 2 other case definitions (Tables S1 and S2); models including more covariates (data not shown); and the SCCS analysis (Table S3).

Individual diseases with >10 cases in female cohorts
Table 3 gives the number of cases per individual disease and the corresponding incidence rate in each of the 4 cohorts. There were 3 diseases for which more than 10 cases were found in the female cohorts, namely autoimmune thyroiditis, Crohn’s disease, and type 1 diabetes. For autoimmune thyroiditis a significant increased risk was found in the female exposed cohort (IRR 3.75, 95% CI: 1.25–11.31) (Table 4). No IRR for males could be calculated as no confirmed cases were found in either male cohort. The IRR for Crohn’s disease was 1.21 (95% CI: 0.37–3.95) for females and 4.22 (95% CI: 0.47–38.02) for males. For

Figure 1. Number of cases included in each analysis. AD = Autoimmune Disease; FU = follow-up. *Confirmation of cases was performed after subject profile review. **The 46 non-confirmed cases were combined with the 109 confirmed cases in the sensitivity analysis for subjects with known first symptom dates. $Subjects for the imputed dates sensitivity analyses had either an imputed date of first symptom or a known date of first symptom. Sensitivity analyses for subjects with imputed/known first symptom dates were repeated using either confirmed cases only or confirmed and non-confirmed cases. $ Date of onset was assumed to be the same as date of disease diagnosis in this sensitivity analysis. Sensitivity analyses were repeated using either confirmed cases only or confirmed and non-confirmed cases.
Table 1. Demographic characteristics of the 4 cohorts.

| Characteristics                          | Parameters or categories | AS04-HPV-16/18 vaccine exposure N = 64,964 | Unexposed historical female cohort N = 64,973 | Unexposed concurrent male cohort N = 64,974 | Unexposed historical male cohort N = 64,965 |
|------------------------------------------|--------------------------|---------------------------------------------|-----------------------------------------------|---------------------------------------------|---------------------------------------------|
|                                          | Value or n | % | Value or n | % | Value or n | % | Value or n | % |
| Age at study start date (years)          | Mean (SD) | — | 15.3 (2.1) | — | 15.3 (2.1) | — | 16.0 (2.0) | — |
|                                          | Range      | — | 9.4–24.9 | — | 9.3–24.9 | — | 9.2–24.8 | — |
| Region of GP practices                   | North England | 36,818 | 56.7 | 34,646 | 53.3 | 35,906 | 55.3 | 33,247 | 66.0 |
|                                         | Midlands   | 8,396 | 12.9 | 8,556 | 13.2 | 8,423 | 13.0 | 8,724 | 13.4 |
|                                         | South England | 19,648 | 30.2 | 21,733 | 33.4 | 20,616 | 31.7 | 22,971 | 35.4 |
|                                         | Ireland Scotland Wales | 102 | 0.2 | 38 | <0.1 | 29 | <0.1 | 23 | <0.1 |
| Available HES linkage                    | Yes | 38,656 | 59.5 | 36,148 | 55.6 | 37,832 | 58.2 | 37,616 | 57.9 |
| Number of healthcare resources utilization* | Mean (SD) | 8.8 (10.2) | — | 7.0 (9.1) | — | 6.0 (8.4) | — | 5.3 (7.2) | — |
| Number of years of follow-up in CPRD GOLD at study start date | Mean (SD) | 9.4 (4.3) | — | 7.6 (4.3) | — | 9.1 (4.3) | — | 7.8 (4.4) | — |
| Exposure to vaccines in the year prior to the study start date | Any vaccine | 11,529 | 17.7 | 11,008 | 16.9 | 9,270 | 14.3 | 10,394 | 16.0 |
| Exposure to vaccines in the follow-up period | Novel adjuvanted vaccine | 311 | 0.5 | 0 | 0.0 | 325 | 0.5 | 0 | 0.0 |
|                                         | Live-attenuated vaccine | 1,138 | 1.8 | 2,986 | 4.6 | 861 | 1.3 | 2,942 | 4.5 |
|                                         | Other vaccine | 10,627 | 16.4 | 8,580 | 13.2 | 8,507 | 13.1 | 20,924 | 32.2 |
|                                         | Any vaccine | 11,596 | 17.8 | 7,765 | 12.0 | 8,000 | 12.3 | 6,253 | 9.6 |
|                                         | Novel adjuvanted vaccine | 1,679 | 2.6 | 0 | 0.0 | 1,559 | 2.4 | 0 | 0.0 |
|                                         | Live-attenuated vaccine | 1,033 | 1.6 | 943 | 1.5 | 489 | 0.8 | 828 | 1.3 |
|                                         | Other vaccine | 10,231 | 15.7 | 7,062 | 10.9 | 7,068 | 10.9 | 5,578 | 8.6 |

Note. HES = Hospital Episode Statistics; N = number of subjects; n/% = number/percentage of subjects in a given category; SD = standard deviation; Value = value of the considered parameter.
*E.g. general practioner consultations, prescriptions, and laboratory tests.

Table 2. Incidence rate per 100,000 person-years and incidence rate ratios of co-primary endpoints.

| Diseases | AS04-HPV-16/18 vaccine exposure (total PY = 64,705) | Unexposed historical female cohort (total PY = 64,841) |
|----------|-----------------------------------------------------|-------------------------------------------------------|
|          | n | IR per 100,000 PY (95% CI) | n | IR per 100,000 PY (95% CI) | IRR’ (95% CI) EXP/NNEXP |
| Co-primary endpoints | Neurinflamatory/ophthalmic AD | Confirmed cases | 0 | 0.00 (0.00–5.70) | 1 | 1.54 (0.04–8.59) | — |
| | | All cases | 4 | 6.18 (1.68–15.83) | 7 | 10.80 (4.34–22.24) | 0.57 (0.17–1.96) |
| | | Other AD | 38 | 58.73 (51.56–60.81) | 27 | 41.64 (27.44–60.58) | 1.41 (0.86–2.31) |
| | | All cases | 51 | 78.82 (58.69–103.63) | 41 | 63.23 (45.38–85.78) | 1.25 (0.83–1.88) |
| | | Unexposed concurrent male cohort (total PY = 64,859) | | | | |
| | | Unexposed historical male cohort (total PY = 64,868) | | | | |
| | | n | IR per 100,000 PY (95% CI) | n | IR per 100,000 PY (95% CI) | IRR’ (95% CI) MALE/HIST |
| Co-primary endpoints | Neurinflamatory/ophthalmic AD | Confirmed cases | 1 | 1.54 (0.04–8.59) | 1 | 1.54 (0.04–8.59) | 0.95 (0.06–15.18) |
| | | All cases | 3 | 4.63 (0.95–13.52) | 2 | 3.08 (0.37–11.14) | 1.73 (0.29–10.47) |
| | | Other AD | 26 | 40.09 (26.19–58.74) | 15 | 23.12 (12.94–40.38) | 1.77 (0.94–3.35) |
| | | All cases | 28 | 43.17 (28.69–62.39) | 19 | 29.29 (17.64–47.45) | 1.52 (0.85–2.73) |

Note. AD = autoimmune disease; CI = confidence interval; EXP = AS04-HPV-16/18 vaccine exposure; HIST = unexposed historical male cohort; IRR = incidence rate ratio; MALE = unexposed concurrent male cohort; n = number of subjects; NNEXP = unexposed historical female cohort; PY = person-years
*Adjusted for age group (9–17 years, 18–25 years)
Table 3. Incidence rate per 100,000 person-years of individual autoimmune diseases.

| Diseases                               | AS04-HPV-16/18 vaccine exposure | Unexposed historical female cohort | Unexposed concurrent male cohort | Unexposed historical male cohort |
|----------------------------------------|---------------------------------|-----------------------------------|---------------------------------|---------------------------------|
|                                        | (total PY = 64,705)             | (total PY = 64,841)               | (total PY = 64,859)             | (total PY = 64,868)             |
| n                                      | IR per 100,000 PY               | n                                 | IR per 100,000 PY               | n                               |
| Acute disseminated encephalomyelitis    |                                 |                                   |                                 |                                 |
| Confirmed cases                        | 0                               | 0                                 | 0                               | 0                               |
| All cases                              | 1                               | 1.55                              | 0                               | 0                               |
| Autoimmune thyroiditis                 | 15                              | 23.18                             | 4                               | 6.17                            |
| All cases                              | 26                              | 40.18                             | 18                              | 27.76                           |
| Autoimmune uveitis                     | 0                               | 0.00                              | 0                               | 0.00                            |
| All cases                              | 2                               | 3.09                              | 5                               | 7.71                            |
| Crohn’s disease                        | 6                               | 9.27                              | 5                               | 7.71                            |
| All cases                              | 8                               | 12.36                             | 5                               | 7.71                            |
| Guillain-Barré syndrome                | 0                               | 0.00                              | 0                               | 0.00                            |
| All cases                              | 0                               | 0.00                              | 0                               | 0.00                            |
| Idiopathic thrombo-cytopenic purpura   | 1                               | 1.55                              | 1                               | 1.54                            |
| All cases                              | 1                               | 1.55                              | 1                               | 1.54                            |
| Inflammatory bowel disease             | 0                               | 0.00                              | 0                               | 0.00                            |
| All cases                              | 0                               | 0.00                              | 0                               | 0.00                            |
| Juvenile rheumatoid arthritis          | 1                               | 1.55                              | 0                               | 0.00                            |
| All cases                              | 1                               | 1.55                              | 0                               | 0.00                            |
| Multiple sclerosis                     | 0                               | 0.00                              | 1                               | 1.54                            |
| All cases                              | 0                               | 0.00                              | 1                               | 1.54                            |
| Optic neuritis                         | 0                               | 0.00                              | 0                               | 0.00                            |
| All cases                              | 0                               | 0.00                              | 0                               | 0.00                            |
| Other AD                               | 1                               | 1.55                              | 0                               | 0.00                            |
| All cases                              | 1                               | 1.55                              | 0                               | 0.00                            |
| Psoriatic arthritis                    | 1                               | 1.55                              | 1                               | 1.54                            |
| All cases                              | 1                               | 1.55                              | 1                               | 1.54                            |
| Rheumatoid arthritis                   | 1                               | 1.55                              | 0                               | 0.00                            |
| All cases                              | 1                               | 1.55                              | 0                               | 0.00                            |
| Type 1 diabetes mellitus               | 8                               | 12.36                             | 16                              | 24.68                           |
| All cases                              | 8                               | 12.36                             | 16                              | 24.68                           |
| Ulcerative colitis                     | 4                               | 6.18                              | 0                               | 0.00                            |
| All cases                              | 4                               | 6.18                              | 1                               | 1.54                            |

Note. n = number of subjects; PY = person-years
*AD = autoimmune disease, includes acute disseminated encephalomyelitis and autoimmune peripheral neuropathies and plexopathies

Table 4. Incidence rate per 100,000 person-years and incidence rate ratios of individual autoimmune diseases with >10 cases in female cohorts.

| Diseases                               | AS04-HPV-16/18 vaccine exposure | Unexposed historical female cohort | Unexposed concurrent male cohort | Unexposed historical male cohort |
|----------------------------------------|---------------------------------|-----------------------------------|---------------------------------|---------------------------------|
|                                        | (total PY = 64,705)             | (total PY = 64,841)               | (total PY = 64,859)             | (total PY = 64,868)             |
| n                                      | IR per 100,000 PY               | n                                 | IR per 100,000 PY               | n                               |
| Autoimmune thyroiditis                 |                                 |                                   |                                 |                                 |
| Confirmed cases                        | 15                              | 23.18                             | 4                               | 6.17                            |
| All cases                              | 26                              | 40.18                             | 18                              | 27.76                           |
| Crohn’s disease                        | 6                               | 9.27                              | 5                               | 7.71                            |
| All cases                              | 8                               | 12.36                             | 5                               | 7.71                            |
| Type 1 diabetes mellitus               | 8                               | 12.36                             | 16                              | 24.68                           |
| All cases                              | 8                               | 12.36                             | 16                              | 24.68                           |

Unexposed concurrent male cohort (total PY = 64,859)

| Diseases                               | n                                 | IR per 100,000 PY (95% CI) |
|----------------------------------------|-----------------------------------|---------------------------|
| Autoimmune thyroiditis                 | 0                                 | 0.00 (0.00; 5.69)         |
| All cases                              | 2                                 | 3.08 (0.37;11.14)         |
| Crohn’s disease                        | 4                                 | 6.17 (1.68;15.79)         |
| All cases                              | 4                                 | 6.17 (1.68;15.79)         |
| Type 1 diabetes mellitus               | 20                                | 30.84 (18.84;47.62)       |
| All cases                              | 20                                | 30.84 (18.84;47.62)       |

Unexposed historical male cohort (total PY = 64,868)

| Diseases                               | n                                 | IR per 100,000 PY (95% CI) |
|----------------------------------------|-----------------------------------|---------------------------|
| Autoimmune thyroiditis                 | 0                                 | 0.00 (0.00; 5.69)         |
| All cases                              | 2                                 | 3.08 (0.37;11.14)         |
| Crohn’s disease                        | 4                                 | 6.17 (1.68;15.79)         |
| All cases                              | 4                                 | 6.17 (1.68;15.79)         |
| Type 1 diabetes mellitus               | 20                                | 30.84 (18.84;47.62)       |
| All cases                              | 20                                | 30.84 (18.84;47.62)       |

Note. CI = confidence interval; EXP = AS04-HPV-16/18 vaccine exposure; HIST = unexposed historical male cohort; IRR = incidence rate ratio; MALE = unexposed concurrent male cohort; n = number of subjects; NNEGEXP = unexposed historical female cohort; PY = person-years
*Adjusted for age group (9–17 years, 18–25 years)
**The IRR for confirmed type 1 diabetes in the female cohorts was adjusted for the male effect, because a significant difference in incidence rates was observed between the 2 male cohorts.
type 1 diabetes, the IRR was 0.50 (95% CI: 0.21–1.17) for females, while a significant increased risk was found in the concurrent male cohort (IRR 2.46, 95% CI: 1.08–5.60). A significant decreased risk of type 1 diabetes was found in the female exposed cohort, when adjusted for male effect (IRR 0.30, 95% CI: 0.11–0.83).

Sensitivity analysis using confirmed and non-confirmed cases showed similar results, except for autoimmune thyroiditis in females, for which a lower and non-significant IRR of 1.45 (95% CI: 0.79–2.64) was found (Table 4). No significant IRRs were found for females and males in any of the other sensitivity analyses (2 other case definitions: Tables S1 and S2, models using other covariates: data not shown), or in the SCSS analysis (Table S3).

**Post-hoc analyses for the autoimmune thyroiditis cases**

The number of autoimmune thyroiditis cases appeared to decrease over time during the one year follow-up period in all cohorts. This was also seen for the other AD.

After additional medical record review, most of the 49 autoimmune thyroiditis cases were classified as hypothyroiditis (81.6%). Corresponding IRRs for autoimmune hypothyroiditis in the female cohorts were 3.00 (95% CI: 0.97–9.31) for confirmed cases and 1.47 (95% CI: 0.76–2.83) for confirmed and non-confirmed cases. No confirmed autoimmune hypothyroiditis cases were found in the male cohorts, but when considering confirmed and non-confirmed cases the IRR was 1.90 (95% CI: 0.17–20.94). These results confirm the estimates in females for all autoimmune thyroiditis (hypothyroiditis and hyperthyroiditis combined).

After exclusion of subjects from the Northern Ireland, Scotland, and Wales regions (as per post-hoc analysis), a non-significant IRR for confirmed autoimmune thyroiditis was found (IRR 2.50, 95% CI: 0.79–7.98).

**Discussion**

The main analysis based on confirmed cases showed no significant IRRs for any of the co-primary endpoints. However, among the most frequent AD for which symptom start dates are difficult to establish, the risk of autoimmune thyroiditis was increased and the risk of type 1 diabetes was decreased in the female vaccinated cohort. Sensitivity analysis using all cases (i.e. confirmed and non-confirmed) showed similar results, except for autoimmune thyroiditis in which the IRR was not significant anymore. The findings on autoimmune thyroiditis and type 1 diabetes from the main analysis were not confirmed in sensitivity analyses using other case definitions, nor in the SCSS analysis.

A publication by the UK Medicines and Healthcare products Regulatory Agency (MHRA) reviewed the safety profile of AS04-HPV-16/18vaccine use in the UK from September 2008 to July 2012, when over 6 million doses of the vaccine had been given across the UK, and identified no new safety concerns.9 Randomized clinical studies did not show any increased risk of AD in the vaccinated group compared to controls.8,10,11 A post-licensure safety surveillance of routine use of AS04-HPV-16/18vaccine did not find any pattern or trend for potential immune-mediated diseases after vaccination.12 This current vaccine post-licensure study confirms the overall acceptable safety profile of AS04-HPV-16/18vaccine.

Research by Chao, in which the Kaiser Permanente Database was used and AD cases were found using similar case identification and ascertainment methods, showed an increased risk of Grave’s and Hashimoto diseases combined and a decreased risk of type 1 diabetes after 4 vHPV vaccination.13 A study by Arnheim-Dahlström using healthcare registers from Denmark and Sweden, on the contrary, found an increased risk of type 1 diabetes after 4 vHPV vaccination.14 However, both authors concluded that there was no clear evidence of a safety signal following vaccination with 4vHPV, because no cluster of disease onset in relation to vaccination timing was found and no significant increased risk of most other conditions was found in vaccinated women. Moreover, in a follow-up review of the study by Chao, the authors concluded that many of the confirmed incident Grave’s disease cases were actually prevalent cases.15 A recent observational study carried out in a cohort of approximately 4 million women aged 10 to 44 years in Denmark did not find an increased risk of multiple sclerosis or other demyelinating diseases after 4 vHPV vaccination.16 Additionally, other observational studies did not find any increased risk of AD in the 4 vHPV vaccinated group compared to an unvaccinated group.17–19

The incidence of autoimmune thyroiditis in the vaccinated cohort was within the same range as the one in CPRD GOLD for the studied age group (incidence rates from the feasibility assessment for the period 2008–2010: age group [9–18] = 1.22 and 5.52/100,000 person-years respectively in males and females, age group [18–25] = 1.88 and 8.30/100,000 person-years respectively in males and females), indicating that although we found a significantly increased incidence in the exposed cohort, this was still within expected ranges. The increased incidence of autoimmune thyroiditis could be explained by a change in diagnostic methods over time.

Our study had a number of limitations. First, CPRD GOLD is based on data from general practices (GP), while most autoimmune diseases are probably diagnosed in specialist settings. Not all GPs participating in CPRD GOLD consented to the linkage between CPRD GOLD primary care data and Hospital Episodes Statistics (HES) data (linkage was around 50% as of the first quarter of 2013). Consequently, the number of autoimmune diseases, the quality of the information, and the diagnostic certainty might be limited. In particular, the specific information related to the onset of clinical symptoms, and radiological, biological and genetic predisposition data associated with the etiologic diagnosis of AD may not have all been available in the CPRD GOLD database and associated resources. This is reflected in the low confirmation rate for some of the AD (i.e., autoimmune thyroiditis, autoimmune uveitis). Second, when the first symptom of an AD for a subject was known but the date of onset of the symptom was not known (i.e. there was no indication regarding the date on which the first symptom started), the date of first report of this symptom was used as date of first symptom. This is a limitation in the main analysis and sensitivity analysis using both known and imputed dates of first symptom, because it is highly likely that
in a subset of these subjects the symptom has started (much) earlier, possibly before the first dose of AS04-HPV-16/18 vaccine. Third, analyses for the first co-primary endpoint (neuroinflammatory/ophthalmic AD) and most of the individual AD were not possible due to the small number of cases. Fourth, an additional limitation could be the risk of identifying false negative cases (lack of sensitivity). The case ascertainment procedure ensured a high specificity of the endpoint(s), but the team did not review the subject profiles of the non-cases (because unfeasible for 65,000 subjects per cohort), and this means that possible cases of ADs could have been missed. However a high specificity was required to avoid a bias toward the null hypothesis whereas high sensitivity was not essential. Lack of sensitivity does not bias the risk estimate, but could impact the precision resulting in a somewhat broader confidence interval. Fifth, the number of AD cases seemed to decrease over time during the one year follow-up period in all 4 cohorts. This could potentially be explained by our study design: a diagnosis of AD was searched in the database through algorithms during the one year period after the first AS04-HPV-16/18 vaccine dose or equivalent study start date and then it was verified by medical review whether the onset of symptoms occurred during this period. It is plausible that cases of AD with onset of symptoms late during the one year follow-up period were not detected because the diagnosis was reported later than one year after the study start date. However, we feel that only a few cases might have been missed as the onset of several AD is (sub)acute. Sixth, studies of rare events typically have low power and therefore only large risk increases can be detected. The present study shares this limitation. To overcome this, 2 composite co-primary endpoints were defined. The observed incidence of the co-primary endpoint ‘other autoimmune diseases’ was in alignment with the sample size calculation assumptions, but it was lower than expected for the neuroinflammatory diseases. However, the absence of confirmed neuroinflammatory disease cases in the exposed cohort was quite reassuring. Lastly, multiple endpoint comparisons increase the overall type I error. However, no adjustment for type I error is also the most conservative approach for safety endpoints since it avoids masking possible signals.

Despite these limitations, we still think this study performed well. A major strength of this study was that it was based on a large population-based database that is likely to be representative of young women and men in the UK. The use of the CPRD GOLD database provided a unique opportunity to study the effect of AS04-HPV-16/18 vaccine on the occurrence of AD as AS04-HPV-16/18 vaccine was used during 3 years in a universal mass vaccination program for young women in the UK. New onset of AD was assessed by thorough subject data review, combining data from CPRD, HES and free text, and using several case ascertainment steps including expert review. This procedure provides a high specificity of the endpoints which is crucial to minimize the risk of bias to the null hypothesis. Attempts were made to minimize case ascertainment bias by blinded experts for HPV vaccine status during case review. In order to prevent inclusion of vaccinated subjects in an ‘unexposed’ cohort, the vaccinated exposed cohort was compared to a historical unexposed cohort before the start of the AS04-HPV-16/18 vaccine program in the UK. In addition, 2 unexposed male cohorts were enrolled in order to assess a possible change over time in reporting AD in CPRD GOLD independent of AS04-HPV-16/18 vaccine introduction. Though incidence rates of autoimmune diseases differ across gender, the male cohorts were used as an internal control. Finally, for the exposed cohort, an additional SCCS analysis was performed in order to control for all fixed confounders not varying with time during the follow-up period. Age-stratified analyses were also performed and generated consistent results (not presented here).

Conclusion

This observational study did not show evidence of an increased risk of AD following vaccination with AS04-HPV-16/18 vaccine. No significant IRRs were found for the co-primary endpoints in the female cohorts. However, a significant increased risk of autoimmune thyroiditis (IRR=3.75, 95% CI: 1.25–11.31) and a significant decreased risk of type 1 diabetes (after adjustment for male effect, IRR=0.30, 95% CI: 0.11–0.83), was found in the female cohorts using confirmed cases only. Using all cases (i.e., confirmed and non-confirmed) showed similar results, except for autoimmune thyroiditis in which the IRR was not significant anymore. Sensitivity analyses using other case definitions and the SCCS analysis did not find any significant IRR between the exposed and unexposed female cohorts.

Materials and methods

Data source, population and setting

CPRD GOLD is one of the largest anonymised primary care databases, and captures longitudinal medical records including clinical events, laboratory results, drug prescriptions, referrals to specialists, and immunisation records from over 680 GP’s in the UK. Linkage between CPRD GOLD primary care data and HES data was available for approximately 50% of subjects as of the beginning of 2013. Complementary information to coded GP data can be obtained through the free text data captured in the practice management system from CPRD GOLD. Free text data include notes or documents entered or scanned in by the GP, including letters from specialists in secondary or private care settings.

A public immunization program targeting girls between 12–13 years of age including a catch-up program for young women up to 18 years was undertaken in the UK during the academic year 2008/09. The phased catch-up program for females born 1 September 1991 to 31 August 1995 during the 2008/09 academic year was completed by the end of the 2009/10 academic year. The program was delivered largely through secondary schools. In the UK public HPV immunization program (12–13 year olds), HPV vaccination coverage for 2010/11 was 89.0%, 87.6% and 83.8% for the first, second and third dose respectively. The bivalent vaccine was replaced in the program by the tetravalent vaccine Gardasil (4vHPV; Merck & Co) in September 2012.

The study population included female and male subjects registered in CPRD GOLD for at least one year before the study
start (Fig. 2). The female population was composed of subjects vaccinated with AS04-HPV-16/18 vaccine between the ages of 9 and 25 years and unexposed subjects of the same age identified from historical data. A historical unexposed cohort before the start of the Cervarix program in the UK was chosen in order to prevent inclusion of vaccinated subjects in an ‘unexposed’ cohort (because when no vaccination is reported in CPRD GOLD, it cannot be ruled out that the subject did receive the vaccine). The male population was composed of 9 to 25-year-old subjects not vaccinated with AS04-HPV-16/18 vaccine, comprising both a concurrent and a historical male cohort. Comparison of the unexposed concurrent male cohort with the unexposed historical male cohort was used as an internal control for changes over time in the incidence of AD in CPRDGOLD. Women who received at least one dose of AS04-HPV-16/18 vaccine administered according to local practice between 1 September 2008 and 31 August 2010 were eligible for the exposed cohort. Subjects in the 3 unexposed cohorts were pre-selected after applying a frequency matching for age and practice region to the subjects included in the vaccinated (exposed) cohort. A random selection was applied to the pre-selected unexposed subjects in order to include the targeted number of subjects in each unexposed cohort. The study start date for the exposed cohort was the date of the first dose of AS04-HPV-16/18 vaccine. The study start date for the unexposed cohorts was a random date selected among the study start dates of the matched exposed cohort (minus 3 years for the historical cohorts).

Outcome definition
The primary study outcome was the occurrence of new onset of 2 groups of confirmed AD during the period of one year following the study start date (follow-up period). These two co-primary composite endpoints have been defined as: 1) neuroinflammatory/ophthalmic diseases: multiple sclerosis, transverse myelitis, optic neuritis, Guillain-Barré syndrome, autoimmune uveitis, and other demyelinating diseases, or 2) other AD: systemic lupus erythematosus, rheumatoid arthritis, juvenile rheumatoid arthritis, Still’s disease, psoriatic arthritis, ankylosing spondylitis, idiopathic thrombocytopenic purpura, autoimmune hemolytic anaemia, type 1 diabetes mellitus, autoimmune thyroiditis, Crohn’s disease, ulcerative colitis, and autoimmune hepatitis.

Secondary outcomes included the occurrence of new onset of individual confirmed AD during the period of one year following the study start date (follow-up period).

The one-year follow-up period was chosen in agreement with the Food and Drug Administration (and the European Medicines Agency, and is supported by the article from Tavares et al on the optimal conduct of clinical trials of new vaccines investigating the risk of AD.26

Data collection and case ascertainment
Subjects with suspected AD diagnoses were identified in CPRD GOLD and/or HES using pre-defined algorithms (the algorithm for Guillain-Barré syndrome is given in Tables S4 and S5).
as an example, the other algorithms are available upon request). The final study database consisted of data for these subjects automatically extracted from CPRD GOLD (Tables S6 and S7), HES, and additional data from free text review. Information extracted included clinical diagnosis, laboratory testing, drug prescription, and HES-linked data. Specific de-identified free text associated with possible first symptoms, laboratory tests, drug prescriptions, and diagnosis of AD was requested when necessary in order to classify each subject as a confirmed new onset AD case, a non-confirmed new onset AD case, or a non-case. If a date of diagnosis did not fall within the follow-up period, a subject could not qualify as a case in any of the analyses. All subject profiles and requested free text were reviewed by Pallas, Health Research and Consultancy B.V., the Netherlands.

A safety physician from GSK and an external physician from Research Triangle Institute (RTI) Health Solutions reviewed all subjects with a doubtful outcome. Final case ascertainment was adjudicated by 5 independent external experts in the fields of rheumatology, ophthalmology, neurology, and internal medicine who remained blinded with respect to the exposure status of the subjects throughout the ascertainment process. Each expert reviewed the subjects, which included subjects with a doubtful outcome after review by Pallas, the GSK safety physician, and the RTI physician, and a 10% random sample of the remaining subjects per AD, according to their specialty. Fifty subjects were reviewed by the experts as part of the random check. Agreement on the date of first symptom, type of AD, confirmation of AD, and date of diagnosis existed for all subjects with rheumatology and neurology diagnoses and for most of the subjects with ophthalmology and internal medicine diagnoses. For autoimmune uveitis, however, the expert decided to include an additional first symptom (i.e. conjunctivitis/episcleritis) that had not been used by Pallas, GSK, and RTI. For inflammatory bowel disease, Crohn’s diseases, and ulcerative colitis, the expert suggested other criteria to determine the date of diagnosis and confirmation of the diagnosis. All uveitis, inflammatory bowel disease, Crohn’s disease, and ulcerative colitis subjects were therefore reviewed again by Pallas applying the revised criteria. Furthermore, after review by the expert of the systemic lupus erythematosus subjects, the expert proposed other criteria to determine the diagnosis and its confirmation. The expert reviewed all remaining subjects and applied these criteria.

**Statistical analysis**

**Main analyses**
The main analysis included all confirmed AD cases with a known date of first symptom within the follow-up period (i.e., the date of first symptom was set as the date of disease occurrence). A known date of first symptom was either the date (from the free text) that a symptom was said by the patient to have started, or, if this was not available, the date the first symptom was reported in CPRD. If the date of first symptom was within the one year follow-up period but the date of diagnosis was after this period then this subject was not included as a case. The incidence rates of AD during the one year follow-up period were calculated as the number of cases divided by the total person-time. The individual person-time was defined as the time between the study start date and the end of follow-up period (one year from study start date), subject’s date of death, CPRD de-enrolment date, date of unspecified HPV vaccine or 4vHPV, or date of first symptom of AD, whichever occurred first. The comparison of the incidence rates of AD (co-primary endpoints and individual diseases with more than 10 cases in the female cohorts: these concerned Crohn’s disease, autoimmune thyroiditis, and diabetes mellitus type 1) was done using a Poisson regression model, with number of events as dependent variable, exposure status as independent variable, and age as covariate, and the log of person-time as an offset. The IRR (females: exposed/historical, males: concurrent/historical) was derived as the exponential of the coefficient associated with the exposure status and its 95% Wald CI. A Poisson regression model adjusted for time effect was performed for the AD for which a statistically significant difference in incidence rates was observed between the 2 male cohorts. This model included the 4 cohorts and a specific contrast for estimating the difference between the 2 female cohorts adjusted for the difference between the 2 male cohorts.

**Sensitivity analyses**
The following sensitivity analyses were performed:

- Analyses of all cases (confirmed and non-confirmed) with known date of first symptom within the follow-up period;
- Analyses using cases with a known or imputed date of first symptom (confirmed cases only, and confirmed and non-confirmed cases combined) within the follow-up period. In case of missing date of first symptom, a date was imputed using the disease-specific median number of days between the date of diagnosis and the known date of first symptom of all confirmed and non-confirmed cases. If the (imputed) date of first symptom was within the defined risk period but the date of diagnosis was after the risk period then this subject was not included as a case;
- Analyses where the date of diagnosis was set as the date of disease occurrence (confirmed cases only, and confirmed and non-confirmed cases combined);
- Analyses using, in addition to age, also region, other vaccination, and healthcare resource utilization during the year prior to the study start date as covariates.

**Self-controlled case-series**
A SCCS analysis for both co-primary endpoints and individual diseases was performed for the exposed female cohort. For the main SCCS, the risk period was one year after the first AS04-HPV-16/18 vaccine dose; a buffer period was defined as the 6 months after the end of the risk period and the control period was defined as one year after the end of the buffer period.

Potential pre-existing autoimmune conditions may influence vaccination status. For this reason, the control period did not include a pre-vaccine period. The relative incidence rate was calculated for the coprimary endpoints and individual diseases between risk and control periods as the ratio of the incidence rate in the risk period versus the incidence rate in the control period. Confirmed AD cases with a date of first
symptom within these 30 months were included in the SCCS analysis. If the date of first symptom was within the defined risk period but the date of diagnosis was after the risk period then the case was excluded from the SCCS analysis. The same rule was applied for cases occurring in the control period because no diagnosis that occurred after the end of the control period was included in the study. The reason for the use of this rule was to avoid a bias in the number of cases occurring in the risk period.

**Post-hoc analyses**

Post-hoc analyses included a descriptive analysis of time-to-onset of all confirmed autoimmune thyroiditis cases. Moreover, an additional subject profile review was performed for all confirmed and non-confirmed autoimmune thyroiditis cases with a known date of first symptom within the follow-up period in order to classify the cases as hypo- or hyperthyroiditis and to derive IRR’s for these subtypes separately. Lastly, an analysis excluding subjects from Northern Ireland, Scotland, and Wales was performed, because a large proportion of confirmed autoimmune thyroiditis cases were observed in this region (10.2%), while this region represented less than 0.1% of the overall study population cohorts.

**Sample size**

For the cohort design, by hypothesizing that the incidence rates of neuro-inflammatory AD vary between 1 and 10/100,000 person-years and the incidence rates of other AD vary between 50 and 100/100,000 person-years, cohorts of 50,000 subjects each should allow the detection, with 80% power, of a relative risk between 18.7 and 3.7 and between 2.0 and 1.6 respectively for the neuro-inflammatory AD and other AD (our 2 co-primary endpoints). Because of risk of loss to follow-up and missing data, the sample size was increased by 30% for a total of 65,000 subjects in each cohort.

**Abbreviations**

| Abbreviation | Definition |
|--------------|------------|
| AD           | autoimmune diseases |
| CI           | confidence interval |
| CPRD GOLD    | Clinical Practice Research Datalink General Practice OnLine Database |
| GP           | general practices |
| GSK          | GlaxoSmithKline |
| HES          | Hospital Episode Statistics |
| HPV          | human papillomavirus |
| IRR          | incidence rate ratio |
| MHRA         | Medicines and Healthcare products Regulatory Agency |
| RTI          | Research Triangle Institute |
| SCCS         | self-controlled case-series |
| UK           | United Kingdom |
| US           | United States |
| VIVIANE      | Human Papilloma Virus: Vaccine Immunogenicity and Efficacy |

**Disclosure of potential conflicts of interest**

All authors have completed the ICMJE uniform disclosure form at www.icmje.org/coiDisclosure.pdf and declare: LB, DR, MGA and JZ are employed by, and own stock options in, the GlaxoSmithKline Group of Companies. CW is a consultant from Business & Decision Life Sciences to the GlaxoSmithKline Group of Companies. RB reports that CPRD received funding from GSK for input into this study. CPRD is a research organization that conducts commissioned research for pharmaceutical companies and other organizations and receives grant for this service. EB, AS and HV report grants from GSK, during the conduct of the study and grants from GSK, outside the submitted work. MPV received funding for work related to the study through RTI Health Solutions.

**Acknowledgments**

The authors wish to thank Prof. Tjeerd Van Staa for his input into the study during the planning stage, Judith H van den Bosch for her contributions to study concept and design, and the following from GSK Group of Companies: John Logie for their methodological input, and Lakshmi Hariharan for editorial support and publication management. Furthermore, the authors wish to thank the 5 experts (Marina Mesquida, and Luca Quartuccio) for their help with case ascertainment.

**Funding**

GlaxoSmithKline Biologicals SA designed the study, collected and analyzed data, interpreted the results, and approved the manuscript. All authors had access to the data used in the analyses and had final responsibility for the analysis, interpretation, and decision to submit for publication.

**Authors’ contributions**

LB, DR, MGA, RB, EB and CW were involved in study concept and design. CW and DR were involved in the acquisition and analysis of data. JZ, AS, HV, EB and MPV were involved in case ascertainment. HV drafted the manuscript. All authors were involved in interpretation of data. CW and DR were involved in the acquisition and analysis of data. JZ, AS, HV, EB and MPV were involved in case ascertainment. HV drafted the manuscript. All authors were involved in interpretation of data. All authors critically revised the manuscript for important intellectual content. LB obtained funding, supervised the study, had full access to all of the data in the study and takes responsibility for the integrity of the data and the accuracy of the data analysis.

**References**

[1] Stanley M. Pathology and epidemiology of HPV infection in females. Gynecol Oncol 2010; 117:55-10; PMID:20304221; http://dx.doi.org/10.1016/j.ygyno.2010.01.024
[2] de Sanjose S, Quint WG, Alemany L, Geraets DT, Klaustermeier JE, Lloveras B, Tous S, Felix A, Bravo LE, Shin HR, et al. Human papillomavirus genotype attribution in invasive cervical cancer: a retrospective cross-sectional worldwide study. Lancet Oncol 2010; 11:1048-56; PMID:20952254; http://dx.doi.org/10.1016/S1470-2045(10)70230-8
[3] Lehtinen M, Paavonen J, Wheeler CM, Jaisamrarn U, Garland SM, Castellsague X, Skinner SR, Apter D, Naud P, Salmeron J, et al. 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 randomised, double-blind PATRICIA trial. Lancet Oncol 2012; 13:89-99; PMID:22075171; http://dx.doi.org/10.1016/S1470-2045(11)70286-8
[4] Skinner SR, Szarewski A, Romanowski B, Garland SM, Lazzano-Ponce E, Salmeron J, Del Rosario-Raymundo MR, Verheijen RH, Quek SC, da Silva DP, et al. Efficacy, safety, and immunogenicity of the human papillomavirus 16/18 AS04-adjuvanted vaccine in women older than 25 years: 4-year interim follow-up of the phase 3, double-blind, randomised controlled VIVIANE study. Lancet 2014; 384:2211-27; PMID:25189358; http://dx.doi.org/10.1016/S0140-6736(14)60920-X
[5] Wheeler CM, Castellsague X, Garland SM, Szarewski A, Paavonen J, Naudi P, Salmeron J, Chow SN, Apter D, Kitchener H, et al. 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-of-study analysis of the randomised, double-blind PATRICIA trial. Lancet Oncol 2012; 13:100-10; PMID:22075170; http://dx.doi.org/10.1016/S1470-2045(11)70287-X

[6] Klein NP, Ray P, Carpenter D, Hansen J, Lewis E, Fireman B, Black S, Galindo C, Schmidt J, Baxter R. Rates of autoimmune diseases in Kaiser Permanente for use in vaccine adverse event safety studies. Vaccine 2010; 28:1062-8; PMID:19896453; http://dx.doi.org/10.1016/j.vaccine.2009.10.115

[7] Guimaraes LE, Baker B, Perricone C, Shoenfeld Y. Vaccines, adjuvants and autoimmunity. Pharmaco Res 2015; 100:190-209; PMID:26275795; http://dx.doi.org/10.1016/j.phrs.2015.08.003

[8] Angelo MG, David MP, Zima J, Baril L, Dubin G, Arellano F, Struyf F. Pooled analysis of large and long-term safety data from the human papillomavirus-16/18-AS04-adjuvanted vaccine clinical trial programme. Pharmacoeconom Drug Saf 2014; 23:466-79; PMID:24644063; http://dx.doi.org/10.1002/pds.3545

[9] MHRA Public Assessment Report - Cervarix HPV vaccine: safety update at end of 4 years routine use in HPV immunisation programme. London: MHRA, 2012

[10] Descamps D, Hardt K, Spiessens B, Izurita P, Verstraeten T, Breuer T, Dubin G. Safety of human papillomavirus (HPV)-16/18 AS04-adjuvanted vaccine for cervical cancer prevention: a pooled analysis of 11 clinical trials. Hum Vaccin 2009; 5:332-40; PMID:19221517; http://dx.doi.org/10.4161/hv.5.5.7211

[11] Naud PS, Roteli-Martins CM, De Carvalho NS, Teixeira JC, de Borba PC, Sanchez N, Zahaf T, Catteau G, Geeraerts B, Descamps D. Sustained efficacy, immunogenicity, and safety of the HPV-16/18 AS04-adjuvanted vaccine: final analysis of a long-term follow-up study up to 9.4 years post-vaccination. Hum Vaccin Immunother 2014; 10:2147-62; PMID:25424918; http://dx.doi.org/10.1016/j.hvimm.2014.08.004

[12] Angelo MG, Zima J, Tavares Da Silva F, Baril L, Arellano F. Post-licensure safety surveillance for human papillomavirus-16/18-AS04-adjuvanted vaccine: more than 4 years of experience. Pharmacoeconomi Drug Saf 2014; 23:456-65; PMID:24644078; http://dx.doi.org/10.1002/pds.3593

[13] Chao C, Klein NP, Velicer CM, Sy LS, Slezak JM, Takhar H, Ackerson B, Cheetham TC, Hansen J, Deosaransingh K, et al. Surveillance of autoimmune conditions following routine use of quadrivalent human papillomavirus vaccine. J Intern Med 2012; 271:193-203; PMID:21973261; http://dx.doi.org/10.1111/j.1365-2966.2011.02467.x

[14] Arneheim-Dahlstrom L, Pasternak B, Svanstrom H, Sapers P, Hvid 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; 347:f5906; PMID:24108159; http://dx.doi.org/10.1136/bmj.f5906

[15] Jacobsen SJ, Sy LS, Ackerson BK, Chao CR, Slezak JM, Cheetham TC, Takhar HS, Velicer CM, Hansen J, Klein NP. An unmasking phenomenon in an observational post-licensure safety study of adolescent girls and young women. Vaccine 2012; 30:4585-7; PMID:22580356; http://dx.doi.org/10.1016/j.vaccine.2012.04.103

[16] Scheller NM, Svanstrom H, Pasternak B, Arneheim-Dahlstrom L, Sundstrom K, Fink K, Hvid A. Quadrivalent HPV vaccination and risk of multiple sclerosis and other demyelinating diseases of the central nervous system. JAMA 2015; 313:54-61; PMID:25562266; http://dx.doi.org/10.1001/jama.2014.16946

[17] Gee J, Naleway A, Shui I, Bagg J, Yin R, Li R, Kullendorf M, Lewis E, Fireman B, Daley MF, et al. Monitoring the safety of quadrivalent human papillomavirus vaccine: findings from the vaccine safety data-link. Vaccine 2011; 29:8279-84; PMID:21907257; http://dx.doi.org/10.1016/j.vaccine.2011.08.106

[18] Grimaldi-Bensouda L, Guillomet D, Godeau B, Benichou J, Lebrun-Frenay C, Papeix C, Labauge P, Berquin P, Penfornis A, Benhamou PY, et al. Autoimmune disorders and quadrivalent human papillomavirus vaccination of young female subjects. J Intern Med 2014; 275:398-408; PMID:24206418; http://dx.doi.org/10.1111/joim.12155

[19] Slade BA, Leidell L, Vellozzi C, Woej E, Hwa W, Sutherland A, Izurita HS, Ball R, Miller N, Braun MM, et al. Postlicensure safety surveillance for quadrivalent human papillomavirus recombinant vaccine. JAMA 2009; 302:750-7; PMID:19690307; http://dx.doi.org/10.1001/jama.2009.1201

[20] Williams T, van Staa T, Puris S, Eaton S. Recent advances in the utility and use of the general practice research database as an example of a UK primary care data resource. Ther Adv Drug Saf 2012; 3:89-99; PMID:25083228; http://dx.doi.org/10.1177/2042096113459111

[21] Thomas KH, Davies N, Metcalfe C, Windmeijer F, Martin RM, Gunnell D. Validation of suicide and self-harm records in the clinical practice research datalink. Br J Clin Pharmacol 2013; 76:145-57; PMID:23216533; http://dx.doi.org/10.1111/bcp.12059

[22] Crawford CaW J. Health Protection Agency survey of Primary Care Trust teenage vaccination programmes. Health Protection Agency 2009

[23] Sheridan AaW J. Annual HPV vaccine coverage in England in 2010/11. Health Protection Agency Department of Health, 2012

[24] Sheridan AaW J. Annual HPV vaccine coverage in England in 2009/2010. Health Protection Agency Department of Health, 2009

[25] Williams T, van Staa T, Puris S, Eaton S. Recent advances in the utility and use of the general practice research database as an example of a UK primary care data resource. Ther Adv Drug Saf 2012; 3:89-99; PMID:25083228; http://dx.doi.org/10.1177/2042096113459111

[26] Thomas KH, Davies N, Metcalfe C, Windmeijer F, Martin RM, Gunnell D. Validation of suicide and self-harm records in the clinical practice research datalink. Br J Clin Pharmacol 2013; 76:145-57; PMID:23216533; http://dx.doi.org/10.1111/bcp.12059

[27] Crawford CaW J. Health Protection Agency survey of Primary Care Trust teenage vaccination programmes. Health Protection Agency 2009

[28] Sheridan AaW J. Annual HPV vaccine coverage in England in 2010/11. Health Protection Agency Department of Health, 2010

[29] Annual HPV vaccine coverage in England in 2010/11. Health Protection Agency Department of Health, 2010

[30] Williams T, van Staa T, Puris S, Eaton S. Recent advances in the utility and use of the general practice research database as an example of a UK primary care data resource. Ther Adv Drug Saf 2012; 3:89-99; PMID:25083228; http://dx.doi.org/10.1177/2042096113459111

[31] Thomas KH, Davies N, Metcalfe C, Windmeijer F, Martin RM, Gunnell D. Validation of suicide and self-harm records in the clinical practice research datalink. Br J Clin Pharmacol 2013; 76:145-57; PMID:23216533; http://dx.doi.org/10.1111/bcp.12059

[32] Crawford CaW J. Health Protection Agency survey of Primary Care Trust teenage vaccination programmes. Health Protection Agency 2009

[33] Sheridan AaW J. Annual HPV vaccine coverage in England in 2010/11. Health Protection Agency Department of Health, 2010

[34] Annual HPV vaccine coverage in England in 2010/11. Health Protection Agency Department of Health, 2010