Inactivated Vaccine

Human Papillomavirus Vaccine Program Summary (Vaccine and Antigen Code: HPV-9)

Biological	Indications	Schedule	Specific Contraindications	Expected Reactions
Merck Gardasil -9® Includes HPV types 6, 11, 16, 18, 31, 33, 45, 52, 58. (licensed for females 9 – 45 years of age inclusive and males 9 – 26 years inclusive) Goals: 95% of Grade 6 students. Vaccine Program History • Grade 5 HPV-4 school program for girls started in 2008-2009. • 3 year grade 9 catch-up program for girls: 2009/2010 to 2011/2012. • Grade 5 HPV-4 school program for boys started 2014/2015. • 4 year grade 9 catch-up program for boys: 2014/2015 to 2017/2018. • HPV-9 replaced HPV-4 Sept 1, 2016 (i.e. 2016/2017 school year). • Feb 2018 MSM included in eligibility • Sept 2018: schedule change to Grade 6 from Grade 5. • Sept 2018: schedule changed to 2 doses from 3 doses for immunocompetent and non HIV clients 9–14 years of age (inclusive).	 Universal Program: all Grade 6 students (starts Fall 2018) Male individuals aged 17 – 26 years of age who have sex with men. Hematopoietic Stem Cell transplant recipients and solid organ transplant candidates/recipients may be eligible, call CDC team. Notes: The number of doses in a series is based on the age at administration of the first dose and if they are immunocompetent or immunocompromised. Students continue to be eligible for HPV vaccine until the end of grade 12. If in ungraded classes or not in school, eligible from 11 through 18 years of age. Students who began their series with HPV-9. An interrupted schedule does not require restarting. 	Immunocompetent and non-HIV infected individuals 9 – 14 years of age (inclusive): • 2 dose series given: day 0, 6 months. Immunocompromised and/or HIV infected individuals 9 – 14 years of age (inclusive): • 3 dose series: day 0, 2 months, 6 months. Individuals 15 years of age and older: • 3 dose series given at 0, 2 months, and 6 months Minimum Spacing Guidelines For Children Off Schedule: • In a two dose schedule, • min. of 24 weeks between the 1st and 2nd doses • In a three dose schedule, • min. of 4 weeks between the 2nd and 3rd doses • min. of 12 weeks between the 2nd and 3rd doses • min. of 12 weeks between the 2nd and 3rd doses • When reviewing charts, the 3rd dose can be considered valid if there is at least 16 weeks between 1st and 3rd doses: do not use this to plan schedule. Dose: 0.5 ml given IM	Anaphylactic reaction to previous dose of vaccine or any component of the vaccine. Should not be given to Pregnant women. Remaining doses should be deferred until after delivery.	 Very Common: (≥10%) Injection site pain redness, swelling, and itchiness, bruising, mass, bleeding. Common: (1-9%) Headache, fever, nausea, dizziness, fatigue, diarrhea, myalgia, oropharyngeal pain, upper abdominal pain. Rare: (<1/1000) As with any immunization, unexpected or unusual side effects can occur.
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Shake well before administration. Do not inject intradermally, subcutaneously or intravenously. Cold chain must be strictly observed, does not contain any preservatives or Note: antibiotics. Protect from light.