The purpose of this analysis is to facilitate public comment on the proposed Consent agreement, and the Commission does not intend this analysis to constitute an official interpretation of the proposed Consent Agreement or to modify its terms in any way.

By direction of the Commission.

Donald S. Clark,
Secretary.

[FR Doc. 2018–05252 Filed 3–14–18; 8:45 am]
BILLING CODE 6750–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Docket No. CDC–2016–0094]

Final Revised Vaccine Information Materials for MMR (Measles, Mumps, and Rubella) and MMRV (Measles, Mumps, Rubella, and Varicella) Vaccines

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: Under the National Childhood Vaccine Injury Act (NCVIA), CDC must develop vaccine information materials that all health care providers are required to give to patients/parents prior to administration of specific vaccines. On October 18, 2016, CDC published a notice in the Federal Register seeking public comments on proposed updated vaccine information materials for MMR vaccine and MMRV vaccine. Following review of comments submitted and consultation as required under the law, CDC has finalized the materials. Copies of the final vaccine information materials for MMR and MMRV vaccine are available to download from http://www.cdc.gov/vaccines/hcp/vis/index.html or http://www.regulations.gov (see Docket Number CDC–2016–0094). The Vaccine Information Statements (VISs) have been delegated by the Secretary to the Centers for Disease Control and Prevention (CDC). Section 2126 requires that the materials be developed, or revised, after notice to the public, with a 60-day comment period, and in consultation with the Advisory Commission on Childhood Vaccines, appropriate health care provider and parent organizations, and the Food and Drug Administration. The law also requires that the information contained in the materials be based on available data and information, be presented in understandable terms, and include:

1) A concise description of the benefits of the vaccine,
2) A concise description of the risks associated with the vaccine,
3) A statement of the availability of the National Vaccine Injury Compensation Program, and
4) Such other relevant information as may be determined by the Secretary.

The vaccines initially covered under the National Vaccine Injury Compensation Program were diphtheria, tetanus, pertussis, measles, mumps, rubella, and poliomyelitis vaccines. Since April 15, 1992, any health care provider in the United States who intends to administer one of these covered vaccines is required to provide copies of the relevant vaccine information materials prior to administration of any of these vaccines. Since then, the following vaccines have been added to the National Vaccine Injury Compensation Program, requiring use of vaccine information materials for them as well: Hepatitis B, Haemophilus influenzae type b (Hib), varicella (chickenpox), pneumococcal conjugate, rotavirus, hepatitis A, meningococcal, human papillomavirus (HPV), and seasonal influenza vaccines.

Instructions for use of the vaccine information materials are found on the CDC website at: http://www.cdc.gov/vaccines/hcp/vis/index.html.

Revised Vaccine Information Materials

The vaccine information materials referenced in this notice were developed in consultation with the Advisory Commission on Childhood Vaccines, the Food and Drug Administration, and parent and healthcare provider organizations. Following consultation and review of comments submitted, the vaccine information materials covering MMR and MMRV vaccines have been finalized and are available to download from http://www.cdc.gov/vaccines/hcp/vis/index.html or http://www.regulations.gov (see Docket Number CDC–2016–0094). The Vaccine Information Statements (VISs) are “MMR Vaccine (Measles, Mumps, Rubella): What You Need to Know” and “MMRV Vaccine (Measles, Mumps, Rubella, and Varicella): What You Need to Know,” publication date February 12, 2018.

With publication of this notice, by June 1, 2018, all health care providers must discontinue use of the previous editions and provide copies of these updated vaccine information materials prior to immunization in conformance with CDC’s February 23, 2018 Instructions for the Use of Vaccine Information Statements.

Dated: March 12, 2018.

Sandra Cashman,
Executive Secretary, Centers for Disease Control and Prevention.

[FR Doc. 2018–05299 Filed 3–14–18; 8:45 am]
BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–FY–1072; Docket No. CDC–2018–0020]

Proposed Data Collection Submitted for Public Comment and Recommendations

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice with comment period.

SUMMARY: The Centers for Disease Control and Prevention (CDC), as part of its continuing effort to reduce public