

Reporting National Drug Codes for Physician-administered Drugs

The Deficit Reduction Act of 2005 directs state Medicaid agencies to collect manufacturer rebates on physician-administered drugs and biologicals (hereafter referred to as drugs). In order to identify the manufacturer from whom the rebate should be collected, the state Medicaid agencies are requiring that an 11-digit National Drug Code (NDC) be reported on claims for physician-administered drugs in addition to the applicable Healthcare Common Procedure Coding System (HCPCS) and ICD-9 codes. Claims submitted with invalid or missing NDC information are likely to result in a rejected claim.

Each state has the authority to determine which physician-administered drugs are subject to the NDC reporting requirement. Some states have excluded vaccines from the reporting requirement, and others require that all claims for physician-administered drugs include the NDC for the drug administered. Check with your state Medicaid administrator to determine specific NDC reporting requirements and coding requirements for Vaccines for Children (VFC) program and Medicaid funded immunizations.

What is an NDC and where can I find it?

The NDC is a unique identifier assigned to a drug product by the labeler or manufacturer under Food and Drug Administration (FDA) regulations. The first segment identifies the labeler or manufacturer. The second segment identifies the product, product's strength, dosage form, and formulation. The third segment identifies the specific packaging of the particular product presentation. Many NDCs, including the NDCs for sanofi pasteur's products, are assigned and therefore displayed on the packaging and Prescribing Information (PI) sheet in a 10-digit format.

How do I convert NDCs from 10 digits to 11 digits?

Proper billing of an NDC requires an 11-digit number in a 5-4-2 digit format. The following table shows common 10-digit NDC formats found on packaging and the associated conversion to an 11-digit format using the proper placement of an extra zero. To change the 5-3-2 digit format used on sanofi pasteur packaging to an 11-digit format, insert a zero at the beginning of the second segment as highlighted below.

Guidance on Converting to a 5-4-2 NDC Format

Manufacturer assigned NDC	5-4-2 Format	Explanation
1234-5678-91	<u>0</u> 1234-5678-91	Add a zero to the beginning of the first segment if the first segment has only four numbers.
12345-678-91	12345- <u>0</u> 678-91	Add a zero to the beginning of the second segment if the second segment has only three numbers.
12345-6789-1	12345-6789- <u>0</u> 1	Add a zero to the beginning of the third segment if the third segment has only one number.

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Please refer to the Proper Coding of sanofi pasteur Products brochure or the Reimbursement Information section on VaccineShoppe.com® (www.vaccineshoppe.com>Resources>Reimbursement Information>Coding sanofi pasteur Products>Proper Coding of sanofi pasteur Products) for a complete listing of 11-digit NDC codes for sanofi pasteur products.

How is the NDC reported on a CMS-1500 (08/05 version) claim form?

The shaded portion of fields 24A to 24I will be used to report NDC information. To report this information, begin at field 24A as follows:

1. Enter the NDC qualifier of N4
2. Enter the NDC 11-digit numeric code
3. Enter the drug description
4. Enter the NDC Unit qualifier
 - F2 – International Unit
 - GR – Gram
 - ML – Milliliter
 - UN – Unit
5. Enter the NDC Administered Amount in the format 9999.99

The following example shows a claim form reporting an 11-digit NDC for a 0.5mL dose of Adacel®, Tetanus Toxoid, Reduced Diphtheria Toxoid and Acellular Pertussis Vaccine Adsorbed, to a patient 20 years of age. When reporting the NDC on the CMS-1500 claim form the hyphens that appear on the packaging and in the PI should be omitted.

24. A. DATE(S) OF SERVICE FROM To MMDD YY MM DD YY	B. PLACE OF SERVICE	C. EMG	D. PROCEDURES, SERVICES, OR SUPPLIES (Explain Unusual Circumstances) CPT/HCPCS MODIFIER	E. DIAGNOSIS POINTER	F. \$ CHARGES	G Days Or units	H. EPSDT Family Plan	I. ID. QUAL.	J. RENDERING PROVIDER I.D. #
N449281040010 TDAP VACCINE INTRAMUSCULAR ML 0.5									
04 02 07	04 02 07	11	90715		XX XX	1	Y	NPI	1234567891
04 02 07	04 02 07	11	90471		XX XX	1	Y		

Guidance on state-specific requirement for NDC reporting on physician-administered drugs can be obtained from your state Medicaid administrator or Medicaid managed care plan.



The vaccines division of sanofi-aventis Group

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