	A	B	C	D
1	**Section**	**Field Name**	**Type**	**Description**
2	active_ingredients	name	string	The names of the active, medicinal ingredients in the drug product.
3	active_ingredients	strength	string	The strength of the active, medicinal ingredients in the drug product.
4		dosage_form	string	The drug's dosage form. There is no standard, but values may include terms like `tablet` or `solution for injection`.
5		route	string	The route of administation of the drug product.
6		brand_name	string	Brand or trade name of the drug product.
7		applicant_number	string	Name of the Applicant for the drug product.
8		applicant_full_name	string	The full name of the firm holding legal responsibility for the new drug application.
9		application_type	string	The type of new drug application approval.
10		application_number	string	The FDA assigned number to the application.
11		product_number	string	The FDA assigned number to identify the application products. Each strength is a separate product. May repeat for multiple part products.
12		therapeutic_equivalence_codes	string	The TE Code indicates the therapeutic equivalence rating of generic to innovator Rx products.
13		approval_date	string	The date the product was approved as stated in the FDA approval letter to the applicant.
14		approved_prior_to_1982	string	Products approved prior to the January 1, 1982 contain the phrase: "Approved prior to Jan 1, 1982".
15		reference_listed_drug	string	The RLD is a drug product approved under section 505(c) of the FD&C Act for which FDA has made a finding of safety and effectiveness. In the electronic Orange Book, an RLD is identified by "RLD" in the RLD column.
16		reference_standard	string	A highly purified compound that is well characterized.
17		product_type	string	The group or category of approved drugs.
18	patents	patent_number	string	Patent numbers as submitted by the applicant holder for patents covered by the statutory provisions.
19	patents	expire_date	string	The date the patent expires as submitted by the applicant holder including applicable extensions.
20	patents	drug_substance_flag	string	Patents submitted on FDA Form 3542 and listed after August 18, 2003 may have a drug substance flag indicating the sponsor submitted the patent as claiming the drug substance.
21	patents	drug_product_flag	string	Patents submitted on FDA Form 3542 and listed after August 18, 2003 may have a drug product flag indicating the sponsor submitted the patent as claiming the drug product.
22	patents	patent_use_code	string	Code to designate a use patent that covers the approved indication or use of a drug product. May repeat for multiple applications, multiple products and multiple patents.
23	patents	patent_delist_flag	string	Sponsor has requested patent be delisted. This patent has remained listed because, under Section 505(j)(5)(D)(i) of the Act, a first applicant may retain eligibility for 180-day exclusivity based on a paragraph IV certification to this patent for a certain period. Applicants under Section 505(b)(2) are not required to certify to patents where this flag is set to Y.
24	patents	patent_submission_date	string	The date on which the FDA receives patent information from the new drug application (NDA) holder.
25	exclusivity	exclusivity_code	string	Code to designate exclusivity granted by the FDA to a drug product.
26	exclusivity	exclusivity_expire_date	string	The date the exclusivity expires.