	A	B	C	D
1	**Section**	**Field Name**	**Type**	**Description**
2		product_number	string	The FDA assigned number to identify the application products. Each strength is a separate product. May repeat for multiple part products.
3		approval_date	string	The date the product was approved as stated in the FDA approval letter to the applicant.
4		approved_prior_to_1982	boolean	Products approved prior to the January 1, 1982 contain the phrase: "Approved prior to Jan 1, 1982".
5		products	array	
6	products	active_ingredients	array of objects	
7	products.active_ingredients	name	string	The names of the active, medicinal ingredients in the drug product.
8	products.active_ingredients	strength	string	The strength of the active, medicinal ingredients in the drug product.
9	products	dosage_form	string	The drug's dosage form. There is no standard, but values may include terms like `tablet` or `solution for injection`.
10	products	route	string	The route of administration of the drug product.
11	products	brand_name	string	Brand or trade name of the drug product.
12	products	applicant_number	string	Name of the Applicant for the drug product.
13	products	applicant_full_name	string	The full name of the firm holding legal responsibility for the new drug application.
14	products	application_type	string	The type of new drug application approval.
15	products	application_number	string	The FDA assigned number to the application.
16	products	therapeutic_equivalence_codes	array of strings	The TE Code indicates the therapeutic equivalence rating of generic to innovator Rx products.
17	products	reference_listed_drug	boolean	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.
18	products	reference_standard	boolean	A highly purified compound that is well characterized.
19	products	product_type	string	The group or category of approved drugs.
20		patents	array	
21	patents	patent_number	string	Patent numbers as submitted by the applicant holder for patents covered by the statutory provisions. May repeat for multiple applications and multiple products. Includes pediatric exclusivity granted by the agency.
22	patents	expire_date	string	The date the patent expires as submitted by the applicant holder including applicable extensions.
23	patents	drug_substance_flag	boolean	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.
24	patents	drug_product_flag	boolean	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.
25	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.
26	patents	patent_delist_flag	boolean	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.
27	patents	patent_submission_date	string	The date on which the FDA receives patent information from the new drug application (NDA) holder.
28		exclusivity	array	
29	exclusivity	exclusivity_code	string	Code to designate exclusivity granted by the FDA to a drug product.
30	exclusivity	exclusivity_expire_date	string	The date the exclusivity expires.