

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**  
 Food and Drug Administration  
**APPLICATION TO MARKET A NEW OR ABBREVIATED NEW  
 DRUG OR BIOLOGIC FOR HUMAN USE**  
*(Title 21, Code of Federal Regulations, Parts 314 & 601)*

Form Approved: OMB No. 0910-0338  
 Expiration Date: January 31, 2017  
 See PRA Statement on page 3.

1. Date of Submission (mm/dd/yyyy)

<b>APPLICANT INFORMATION</b>	2. Name of Applicant
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3. Telephone Number (Include country code if applicable and area code)	4. Facsimile (FAX) Number (Include country code if applicable and area code)
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5. Applicant Address

Address 1 (Street address, P.O. box, company name c/o)		Email Address
Address 2 (Apartment, suite, unit, building, floor, etc.)		
City	State/Province/Region	U.S. License Number if previously issued
Country	ZIP or Postal Code	

6. Authorized U.S. Agent (Required for non-U.S. applicants)

Authorized U.S. Agent Name		Telephone Number (Include area code)
Address 1 (Street address, P.O. box, company name c/o)		
Address 2 (Apartment, suite, unit, building, floor, etc.)		FAX Number (Include area code)
City	State	Email Address
ZIP Code		

<b>PRODUCT DESCRIPTION</b>	7. NDA, ANDA, or BLA Application Number	8. Supplement Number (If applicable)
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9. Established Name (e.g., proper name, USP/USAN name)

10. Proprietary Name (Trade Name) (If any)

11. Chemical/Biochemical/Blood Product Name (If any)

12. Dosage Form	13. Strengths	14. Route of Administration
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15. Proposed Indication for Use

Is this indication for a rare disease (prevalence <200,000 in U.S.)?  Yes  No

Does this product have an FDA Orphan Designation for this indication? <input type="checkbox"/> Yes <input type="checkbox"/> No	If yes, provide the Orphan Designation number for this indication: <input style="width: 50px;" type="text"/>
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Contin.  
Page for  
#15

<b>APPLICATION INFORMATION</b>	16. Application Type (Select one)	<input type="checkbox"/> New Drug Application (NDA) <input type="checkbox"/> Biologics License Application (BLA)
		<input type="checkbox"/> Abbreviated New Drug Application (ANDA)

17. If an NDA, identify the type <input type="checkbox"/> 505 (b)(1) <input type="checkbox"/> 505 (b)(2)	18. If a BLA, identify the type <input type="checkbox"/> 351 (a) <input type="checkbox"/> 351 (k)
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19. If a 351(k), identify the biological reference product that is the basis for the submission.  
 Name of Biologic: \_\_\_\_\_ Holder of Licensed Application: \_\_\_\_\_

20. If an ANDA, or 505(b)(2), identify the listed drug product that is the basis for the submission.  
 Name of Drug: \_\_\_\_\_ Application Number of Relied Upon Product: \_\_\_\_\_

Indicate Patent Certification(s):  P1  P2  P3  P4  Section viii - MOU  Statement of no relevant patents

21. Submission (Select one)  Original  Labeling Supplement  CMC Supplement  Efficacy Supplement  Annual Report

Product Correspondence  REMS Supplement  Postmarketing Requirements or Commitments  Periodic Safety Report

Other (Specify): \_\_\_\_\_

22. Submission Sub-Type <input type="checkbox"/> Presubmission <input type="checkbox"/> Amendment <input type="checkbox"/> Initial Submission <input type="checkbox"/> Resubmission	23. If a supplement, identify the appropriate category. <input type="checkbox"/> CBE <input type="checkbox"/> Prior Approval (PA) <input type="checkbox"/> CBE-30
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24. Does this submission contain *only* pediatric data?       Yes       No

25. Reasons for Submission

26. Proposed Marketing Status (*Select one*)       Prescription Product (Rx)       Over-The-Counter Product (OTC)

27. This application is (*Select one*)       Paper       Paper and Electronic       Electronic

28. Number of Volumes Submitted

29. Establishment Information (*Full establishment information should be provided in the body of the application.*)  
*Refer to the instruction sheet (Form FDA 356h Supplement) for more information.*

Establishment Name		Registration (FEI) Number	
Address 1 ( <i>Street address, P.O. box, company name c/o</i> )		MF Number	
Address 2 ( <i>Apartment, suite, unit, building, floor, etc.</i> )		Establishment DUNS Number	
City	State/Province/Region		
Country	ZIP or Postal Code		
Is the establishment new to the application? <input type="checkbox"/> Yes <input type="checkbox"/> No		What is the status of the establishment? <input type="checkbox"/> Pending <input type="checkbox"/> Active <input type="checkbox"/> Inactive <input type="checkbox"/> Withdrawn	

**Establishment Contact Information**

Name of Contact for the Establishment		Telephone Number ( <i>Include area code</i> )
Address 1 ( <i>Street address, P.O. box, company name c/o</i> )		FAX Number ( <i>Include area code</i> )
Address 2 ( <i>Apartment, suite, unit, building, floor, etc.</i> )		
City	State/Province/Region	Email Address
Country	ZIP or Postal Code	

Manufacturing Steps and/or Type of Testing	Is the site ready for inspection? <input type="checkbox"/> Yes <input type="checkbox"/> No <input type="checkbox"/> N/A If No, when will site be ready? ( <i>mm/dd/yyyy</i> ) _____
<div style="border: 1px solid black; padding: 2px; display: inline-block;">Continuation Page for #29</div>	

30. Cross References (List related BLAs, INDs, NDAs, PMAs, 510(k)s, IDEs, BMFs, MAFs, and DMFs referenced in the current application.)

Contin. Page for #30

31. This application contains the following items (*Select all that apply*)

<input type="checkbox"/> 1. Index	<input type="checkbox"/> 2. Labeling ( <i>Select one</i> ): <input type="checkbox"/> Draft Labeling <input type="checkbox"/> Final Printed Labeling	<input type="checkbox"/> 3. Summary ( <i>21 CFR 314.50 (c)</i> )
<input type="checkbox"/> 4. Chemistry Section <input type="checkbox"/> A. Chemistry, manufacturing, and controls information ( <i>e.g., 21 CFR 314.50(d)(1); 21 CFR 601.2</i> ) <input type="checkbox"/> B. Samples ( <i>21 CFR 314.50 (e)(1); 21 CFR 601.2 (a)</i> ) ( <i>Submit only upon FDA's request</i> ) <input type="checkbox"/> C. Methods validation package ( <i>e.g., 21 CFR 314.50(e)(2)(i); 21 CFR 601.2</i> )		
<input type="checkbox"/> 5. Nonclinical pharmacology and toxicology section ( <i>e.g., 21 CFR 314.50(d)(2); 21 CFR 601.2</i> )	<input type="checkbox"/> 6. Human pharmacokinetics and bioavailability section ( <i>e.g., 21 CFR 314.50(d)(3); 21 CFR 601.2</i> )	
<input type="checkbox"/> 7. Clinical microbiology section ( <i>e.g., 21 CFR 314.50(d)(4)</i> )	<input type="checkbox"/> 8. Clinical data section ( <i>e.g., 21 CFR 314.50(d)(5); 21 CFR 601.2</i> )	

*Item 31 continued on page 3*

31. This application contains the following items (Continued; select all that apply)

<input type="checkbox"/> 9. Safety update report (e.g., 21 CFR 314.50(d)(5)(vi)(b); 21 CFR 601.2)	<input type="checkbox"/> 10. Statistical section (e.g., 21 CFR 314.50(d)(6); 21 CFR 601.2)
<input type="checkbox"/> 11. Case report tabulations (e.g., 21 CFR 314.50(f)(1); 21 CFR 601.2)	<input type="checkbox"/> 12. Case report forms (e.g., 21 CFR 314.50 (f)(2); 21 CFR 601.2)
<input type="checkbox"/> 13. Patent information on any patent that claims the drug/biologic (21 U.S.C. 355(b) or (c))	<input type="checkbox"/> 14. A patent certification with respect to any patent that claims the drug/biologic (21 U.S.C. 355 (b)(2) or (j)(2)(A))
<input type="checkbox"/> 15. Establishment description (21 CFR Part 600, if applicable)	<input type="checkbox"/> 16. Debarment certification (FD&C Act 306 (k)(1))
<input type="checkbox"/> 17. Field copy certification (21 CFR 314.50 (l)(3))	<input type="checkbox"/> 18. User Fee Cover Sheet (PDUFA Form FDA 3397, GDUFA Form FDA 3794, BsUFA Form FDA 3792, or MDUFMA Form FDA 3601)
<input type="checkbox"/> 19. Financial Disclosure Information (21 CFR Part 54)	
<input type="checkbox"/> 20. Other (Specify): _____	

**CERTIFICATION**

I agree to update this application with new safety information about the product that may reasonably affect the statement of contraindications, warnings, precautions, or adverse reactions in the draft labeling. I agree to submit safety update reports as provided for by regulation or as requested by FDA. If this application is approved, I agree to comply with all applicable laws and regulations that apply to approved applications, including, but not limited to, the following:

1. Good manufacturing practice regulations in 21 CFR Parts 210, 211 or applicable regulations, Parts 606, and/or 820.
2. Biological establishment standards in 21 CFR Part 600.
3. Labeling regulations in 21 CFR Parts 201, 606, 610, 660, and/or 809.
4. In the case of a prescription drug or biological product, prescription drug advertising regulations in 21 CFR Part 202.
5. Regulations on making changes in application in FD&C Act section 506A, 21 CFR 314.71, 314.72, 314.97, 314.99, and 601.12.
6. Regulations on Reports in 21 CFR 314.80, 314.81, 600.80, and 600.81.
7. Local, state, and Federal environmental impact laws.

If this application applies to a drug product that FDA has proposed for scheduling under the Controlled Substances Act, I agree not to market the product until the Drug Enforcement Administration makes a final scheduling decision.

The data and information in this submission have been reviewed and, to the best of my knowledge, are certified to be true and accurate.

**Warning:** A willfully false statement is a criminal offense, U.S. Code, title 18, section 1001.

32. Typed Name and Title of Applicant's Responsible Official	33. Date (mm/dd/yyyy)
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34. Telephone Number (Include country code if applicable and area code)	35. FAX Number (Include country code if applicable and area code)	36. Email Address
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37. Address of Applicant's Responsible Official	
Address 1 (Street address, P.O. box, company name c/o)	
Address 2 (Apartment, suite, unit, building, floor, etc.)	
City	State/Province/Region
Country	ZIP or Postal Code

38. Signature of Applicant's Responsible Official or Other Authorized Official	<b>Sign</b>	39. Countersignature of Authorized U.S. Agent	<b>Sign</b>
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**The information below applies only to requirements of the Paperwork Reduction Act of 1995.**

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Department of Health and Human Services  
 Food and Drug Administration  
 Office of Chief Information Officer  
 Paperwork Reduction Act (PRA) Staff  
 PRAStaff@fda.hhs.gov

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