

clinical trial to verify clinical benefit of duvelisib in follicular lymphoma. Because the confirmatory trial was not underway and would not be conducted, the Agency recommended withdrawal of approval of the follicular lymphoma indication pursuant to § 314.150(d) (21 CFR 314.150(d)). On November 24, 2021, Secura Bio, Inc. submitted a letter requesting withdrawal of approval of the follicular lymphoma indication for COPIKTRA (duvelisib) Capsules and waiving its opportunity for hearing.

Therefore, under § 314.150(d), approval of the follicular lymphoma indication for COPIKTRA (duvelisib) Capsules is withdrawn effective April 13, 2022. Withdrawal of approval of the follicular lymphoma indication does not affect any other approved indication for COPIKTRA.

Dated: April 7, 2022.

Lauren K. Roth,

Associate Commissioner for Policy.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2021-D-0603 and FDA-2021-D-0604]

Performance Criteria for Safety and Performance Based Pathway; Guidance for Industry and Food and Drug Administration Staff; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of availability.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing the availability of two final device-specific guidance documents for the Safety and Performance Based Pathway—specifically, “Denture Base Resins—Performance Criteria for Safety and Performance Based Pathway; Guidance for Industry and Food and Drug Administration Staff” and “Facet Screw Systems—Performance Criteria for Safety and Performance Based Pathway; Guidance for Industry and Food and Drug Administration Staff.” The device-specific guidances identified in this notice were developed in accordance with the final guidance entitled “Safety and Performance Based Pathway.”

DATES: The announcement of the guidance is published in the **Federal Register** on April 13, 2022.

ADDRESSES: You may submit either electronic or written comments on

Agency guidances at any time as follows:

Electronic Submissions

Submit electronic comments in the following way:

- **Federal eRulemaking Portal:** <https://www.regulations.gov>. Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to <https://www.regulations.gov> will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else’s Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on <https://www.regulations.gov>.

- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see “Written/Paper Submissions” and “Instructions”).

Written/Paper Submissions

Submit written/paper submissions as follows:

- **Mail/Hand Delivery/Courier (for written/paper submissions):** Dockets Management Staff (HFA-305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

- For written/paper comments submitted to the Dockets Management Staff, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in “Instructions.”

Instructions: All submissions received must include the Docket No. FDA-2021-D-0603 for “Denture Base Resins—Performance Criteria for Safety and Performance Based Pathway; Guidance for Industry and Food and Drug Administration Staff” or Docket No. FDA-2021-D-0604 for “Facet Screw Systems—Performance Criteria for Safety and Performance Based Pathway; Guidance for Industry and Food and Drug Administration Staff.” Received comments will be placed in the docket and, except for those submitted as “Confidential Submissions,” publicly viewable at <https://www.regulations.gov> or at the Dockets Management Staff between 9

a.m. and 4 p.m., Monday through Friday, 240-402-7500.

- **Confidential Submissions—**To submit a comment with confidential information that you do not wish to be made publicly available, submit your comments only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states “THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION.” The Agency will review this copy, including the claimed confidential information, in its consideration of comments. The second copy, which will have the claimed confidential information redacted/blacked out, will be available for public viewing and posted on <https://www.regulations.gov>. Submit both copies to the Dockets Management Staff. If you do not wish your name and contact information to be made publicly available, you can provide this information on the cover sheet and not in the body of your comments and you must identify this information as “confidential.” Any information marked as “confidential” will not be disclosed except in accordance with 21 CFR 10.20 and other applicable disclosure law. For more information about FDA’s posting of comments to public dockets, see 80 FR 56469, September 18, 2015, or access the information at: <https://www.govinfo.gov/content/pkg/FR-2015-09-18/pdf/2015-23389.pdf>.

Docket: For access to the docket to read background documents or the electronic and written/paper comments received, go to <https://www.regulations.gov> and insert the docket number, found in brackets in the heading of this document, into the “Search” box and follow the prompts and/or go to the Dockets Management Staff, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852, 240-402-7500. You may submit comments on any guidance at any time (see 21 CFR 10.115(g)(5)).

An electronic copy of the guidance document is available for download from the internet. See the **SUPPLEMENTARY INFORMATION** section for information on electronic access to the guidance. Submit written requests for a single hard copy of the guidance document entitled “Denture Base Resins—Performance Criteria for Safety and Performance Based Pathway; Guidance for Industry and Food and Drug Administration Staff” or “Facet Screw Systems—Performance Criteria for Safety and Performance Based Pathway; Guidance for Industry and Food and Drug Administration Staff” to the Office of Policy, Center for Devices

and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 66, Rm. 5431, Silver Spring, MD 20993-0002. Send one self-addressed adhesive label to assist that office in processing your request.

FOR FURTHER INFORMATION CONTACT:

Jason Ryans, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 66, Rm. 1613, Silver Spring, MD 20993-0002, 301-796-4908.

SUPPLEMENTARY INFORMATION:

I. Background

These device-specific guidance documents provide performance criteria for premarket notification (510(k)) submissions to support the optional Safety and Performance Based Pathway, as described in the guidance entitled “Safety and Performance Based Pathway.”¹ As described in that guidance, substantial equivalence is rooted in comparisons between new devices and predicate devices. However, the Federal Food, Drug, and Cosmetic Act does not preclude FDA from using performance criteria to facilitate this comparison. If a legally marketed device performs at certain levels relevant to its safety and effectiveness, and a new device meets those levels of performance for the same characteristics, FDA could find the new device as safe and effective as the legally marketed device. Instead of reviewing data from direct comparison testing between the two devices, FDA could support a finding of substantial equivalence with data demonstrating the new device meets the level of performance of an appropriate predicate device(s). Under this optional Safety

and Performance Based Pathway, a submitter could satisfy the requirement to compare its device with a legally marketed device by, among other things, independently demonstrating that the device’s performance meets performance criteria as established in the above-listed guidances, rather than using direct predicate comparison testing for some of the performance characteristics.

A notice of availability of the draft guidances “Denture Base Resins” and “Facet Screw Systems” appeared in the **Federal Register** of August 30, 2021 (86 FR 48430). FDA considered comments received on the “Denture Base Resins” and revised the guidance as appropriate by clarifying what information should be included in premarket submissions for denture resins that are additively manufactured. There were no comments received for the “Facet Screw Systems” guidance.

These guidance documents are being issued consistent with FDA’s good guidance practices regulation (21 CFR 10.115). The guidance documents represent the current thinking of FDA on performance criteria for “Denture Base Resins” and “Facet Screw Systems.” They do not establish any rights for any person and are not binding on FDA or the public. You can use an alternative approach if it satisfies the requirements of the applicable statutes and regulations.

II. Electronic Access

Persons interested in obtaining a copy of the guidance may do so by downloading an electronic copy from the internet. A search capability for all Center for Devices and Radiological Health guidance documents is available

at <https://www.fda.gov/medical-devices/device-advice-comprehensive-regulatory-assistance/guidance-documents-medical-devices-and-radiation-emitting-products>. This guidance document is also available at <https://www.regulations.gov> or <https://www.fda.gov/regulatory-information/search-fda-guidance-documents>.

Persons unable to download an electronic copy of either “Denture Base Resins—Performance Criteria for Safety and Performance Based Pathway; Guidance for Industry and Food and Drug Administration Staff (document number 20001)” or “Facet Screw Systems—Performance Criteria for Safety and Performance Based Pathway; Guidance for Industry and Food and Drug Administration Staff (document number 21001)” may send an email request to CDRH-Guidance@fda.hhs.gov to receive an electronic copy of the document. Please use the document number and complete title to identify the guidance you are requesting.

III. Paperwork Reduction Act of 1995

While these guidance documents contain no new collection of information, they do refer to previously approved FDA collections of information. Therefore, clearance by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501–3521) is not required for these guidance documents. The previously approved collections of information are subject to review by OMB under the PRA. The collections of information in the following FDA regulations and guidance have been approved by OMB as listed in the following table:

21 CFR part or guidance	Topic	OMB control No.
807, subpart E “Requests for Feedback and Meetings for Medical Device Submissions: The Q-Submission Program”.	Premarket notification Q-submissions; Pre-submissions.	0910-0120 0910-0756

Dated: April 8, 2022.

Lauren K. Roth,

Associate Commissioner for Policy.

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¹ Available at <https://www.fda.gov/regulatory-information/search-fda-guidance-documents/safety-and-performance-based-pathway>.