

ACF is currently working on future revisions to this information collection, which will be submitted to OMB for review and approval in 2025. Notices inviting public comment on those revisions will accompany that request, but comments received in response to

this notice could also inform those revisions. Through this request process minor discrepancies were noted between the OMB-approved instruments and those currently in use. These minor errors were fixed in the versions included with the request to OMB.

*Respondents:* General Departmental (GDSRAE), State (SSRAE), and Competitive (CSRAE) grant recipients, their subrecipients, and program participants.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents (total over request period)	Number of responses per respondent (total over request period)	Avg. burden per response (in hours)	Total/ annual burden (in hours)
<b>(1) Participant Entry Survey</b>				
GDSRAE participants .....	126,130	1	0.1333	16,813
SSRAE participants .....	317,633	1	0.1333	42,340
CSRAE participants .....	20,136	1	0.1333	2,684
<b>(2) Participant Exit Survey</b>				
GDSRAE participants .....	100,904	1	0.1667	16,821
SSRAE participants .....	254,106	1	0.1667	42,360
CSRAE participants .....	16,109	1	0.1667	2,685
<b>(3) Performance reporting data entry form: grant recipients</b>				
GDSRAE grant recipients .....	119	2	16	3,808
SSRAE grant recipients .....	39	2	16	1,248
CSRAE grant recipients .....	34	2	16	1,088
<b>(4) Performance reporting data entry form: subrecipients</b>				
GDSRAE subrecipients .....	252	2	13	6,552
SSRAE subrecipients .....	426	2	13	11,076
CSRAE subrecipients .....	63	2	13	1,638

*Estimated Total Annual Burden Hours:* 149,113.

*Authority:* 42 U.S.C. 1310.

**Mary C. Jones,**

*ACF/OPRE Certifying Officer.*

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

[Docket No. FDA-2024-P-4163]

**Determination That NOXAFIL (Posaconazole) Delayed-Release Tablets, 100 Grams Was Not Withdrawn From Sale for Reasons of Safety or Effectiveness; Correction**

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice; correction.

**SUMMARY:** The Food and Drug Administration (FDA) is correcting a notice that appeared in the **Federal Register** on October 16, 2024. The

document announced that NOXAFIL (posaconazole) delayed-release tablets, 100 grams (g), was not withdrawn from sale for reasons of safety or effectiveness. The document incorrectly listed the dosage strength as 100 g. The correct strength is 100 milligrams (mg). This notice corrects that error.

**FOR FURTHER INFORMATION CONTACT:** Awo Archampong-Gray, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 51, Rm. 6243, Silver Spring, MD 20993-0002, 301-796-0110, *Awo.Archampong-Gray@fda.hhs.gov*.

**SUPPLEMENTARY INFORMATION:** In FR Doc. 2024-23811, published in the **Federal Register** of Wednesday, October 16, 2024 (89 FR 83504), appearing on pages 83504 and 83505, the following corrections are made:

1. On page 83504, in the second column, the title of the document is corrected to read “Determination That NOXAFIL (Posaconazole) Delayed-Release Tablets, 100 Milligrams, Was Not Withdrawn From Sale for Reasons of Safety or Effectiveness.”

2. On page 83504, in the third column, in the **SUMMARY**, in the first paragraph, “NOXAFIL (posaconazole) delayed-release tablets, 100 grams (g),” is corrected to read “NOXAFIL (posaconazole) delayed-release tablets, 100 milligrams (mg).”

3. On page 83505, in the first and second columns, all uses of “100 g” are corrected to read “100 mg”.

Dated: November 7, 2024.

**Kimberlee Trzeciak,**

*Deputy Commissioner for Policy, Legislation, and International Affairs.*

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