Soft Lights Foundation

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July 15, 2025

BY WEBFORM

https://www.accessdata.fda.gov/scripts/foi/FOIRequest/requestform.cfm

Re: FOIA Request – Radiation Control Program

Dear FDA CDRH FOIA,

21 U.S.C. 360hh-360ss is titled "Electronic Product Radiation Control". This is a Freedom of Information Act request for information and documents related to the FDA's implementation of the Electronic Product Radiation Control Program for the fiscal year 2024-2025. The information requested herein will establish the Administrative Record for litigation under the Administrative Procedure Act.

The following documents are requested under the Freedom of Information Act for the fiscal year 2024-2025. In the list below, the term "documents" shall mean all documents, research, emails, budgets, agreements, memos, meeting minutes, reports, justifications, public comments, petitions, publications, policies, lawsuits, and TEPRSSC meeting notes for the Electronic Product Radiation Control program for each of the following wavelengths:

1. Radio
2. Microwave
3. Infrared
4. Visible Light
5. Ultraviolet
6. X-ray
7. Gamma ray

These documents will create an administrative record for one or more APA lawsuits against the FDA and other federal agencies for abuse of discretion and the failure to establish adequate Radiation Control Programs for electronic products that electromagnetic radiation.

In accordance with 21 C.F.R. 20.46, I request a Fee Waiver because this request it is in the public interest and likely to contribute significantly to public understanding of the operations or activities of the FDA CDRH, and because this request is not primarily in the commercial interest of the requester.

Sincerely,

/s/ Mark Baker
Individual

/s/ Mark Baker
President
Soft Lights Foundation
mbaker@softlights.org