

GSA FDA PROTOTYPE

GUI Style Guidelines

Document Version: 1.0

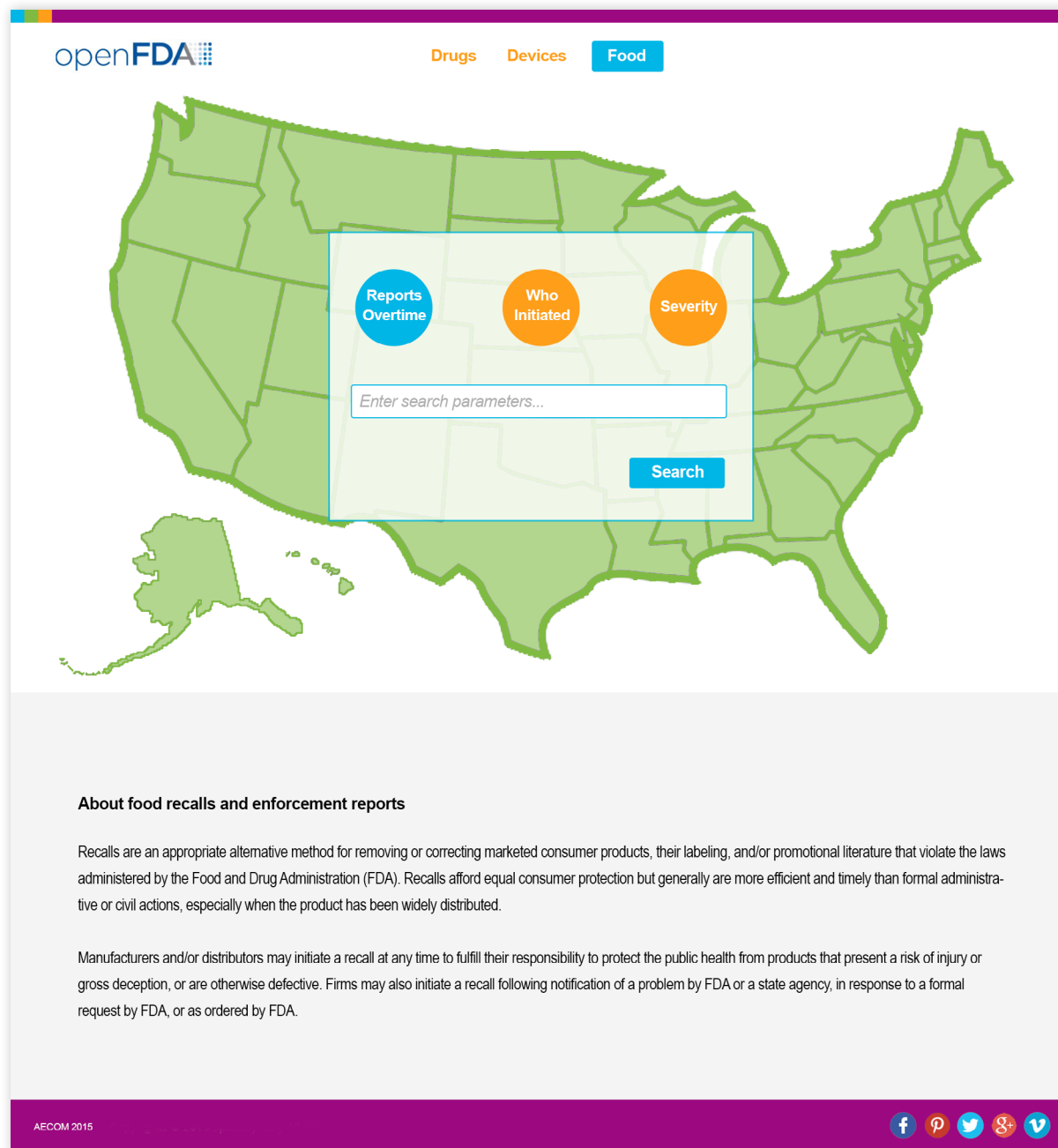
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Presented by AECOM

ABOUT THIS GUIDE

This guide is designed to set rules for styling the GSA FDA prototype application, to provide an easy way to maintain uniform layout and logic for the front-end only. Using a common set of rules allows us to easily create new and maintain existing systems. From a user perspective, this will inspire confidence in use and easy of transition between applications, as well as reducing training times need for new employees and easy transition from department to department.

For developers: Provided are screenshots with sets of font, size, position and color breakdowns for this application. All common controls are discussed including; buttons, textboxes, layout, etc.



HEADER, FOOTER, AND MAIN NAVIGATION

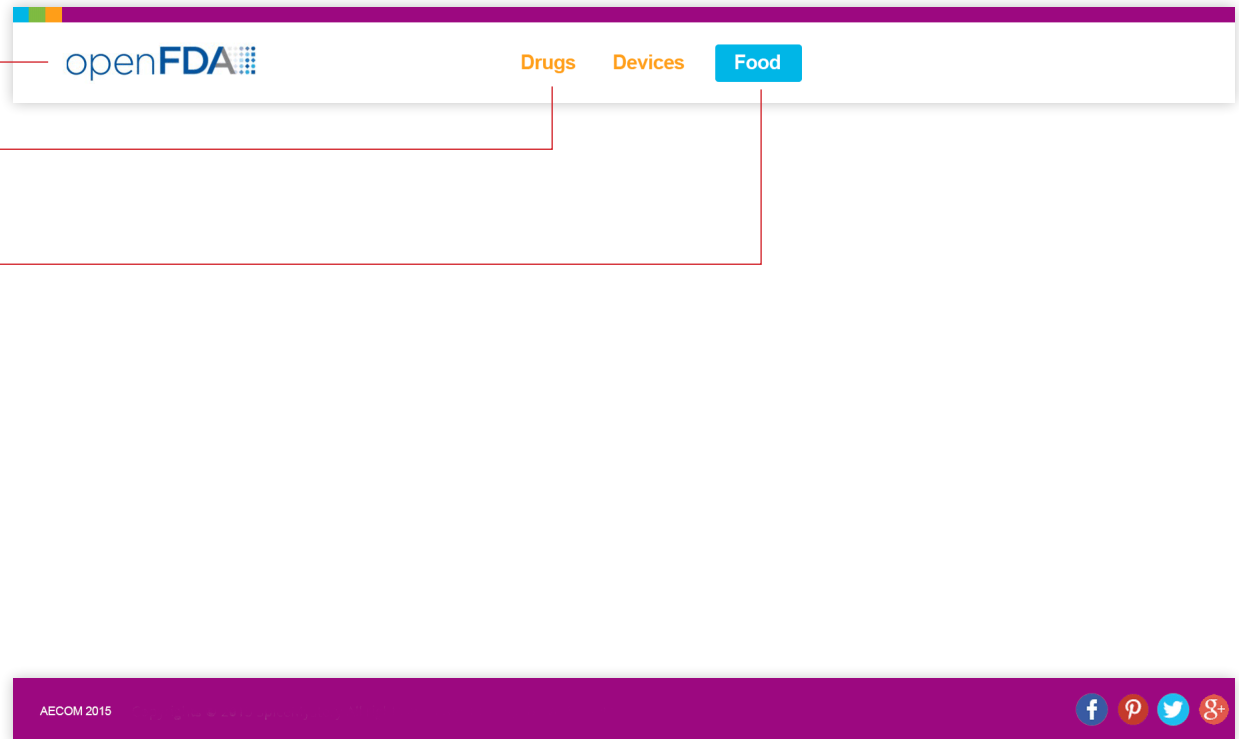
Color Bar
blue # 00b5e5
green # 7fbb42
orange # fc9f1a
magenta # 9c0880

FDA Logo
image provided

Main Navigation
Arial ; bold; 30px;
fc9f1a

**Main Navigation
Selected**
Arial ; bold; 30px;
ffffff

Footer
Fill Color # 9c0880
Arial; 18px; # ffffff



SEARCH PANEL

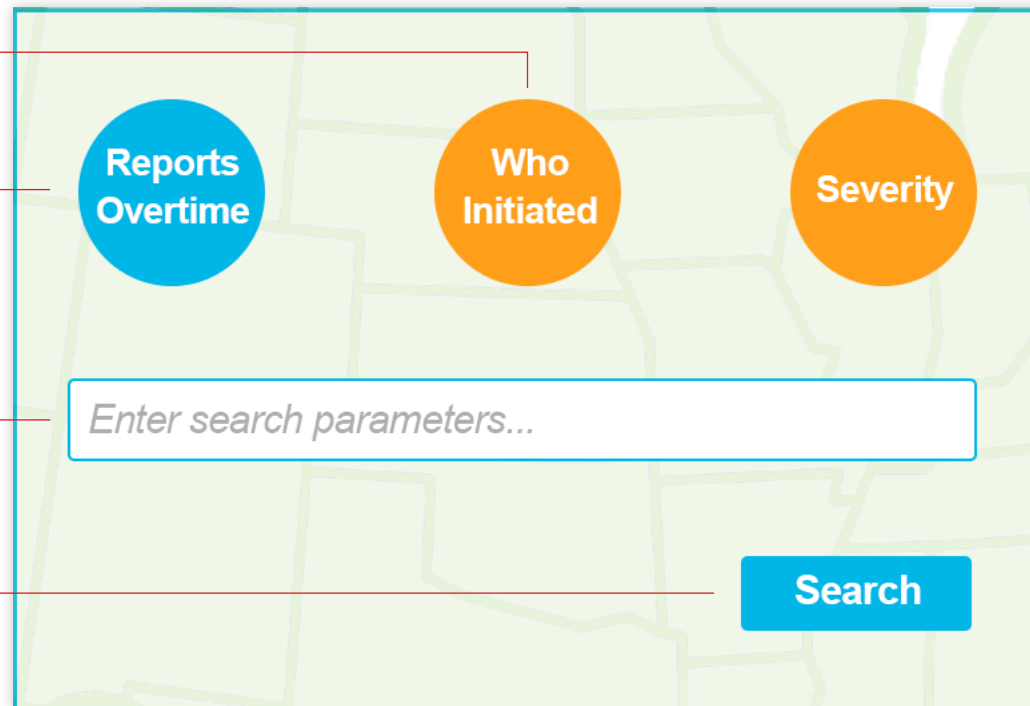
Search Panel
background # ffffff; 80% Opacity
border 3 px; # 00b5e5

Main Navigation
Arial ; bold; 30px;
fc9f1a

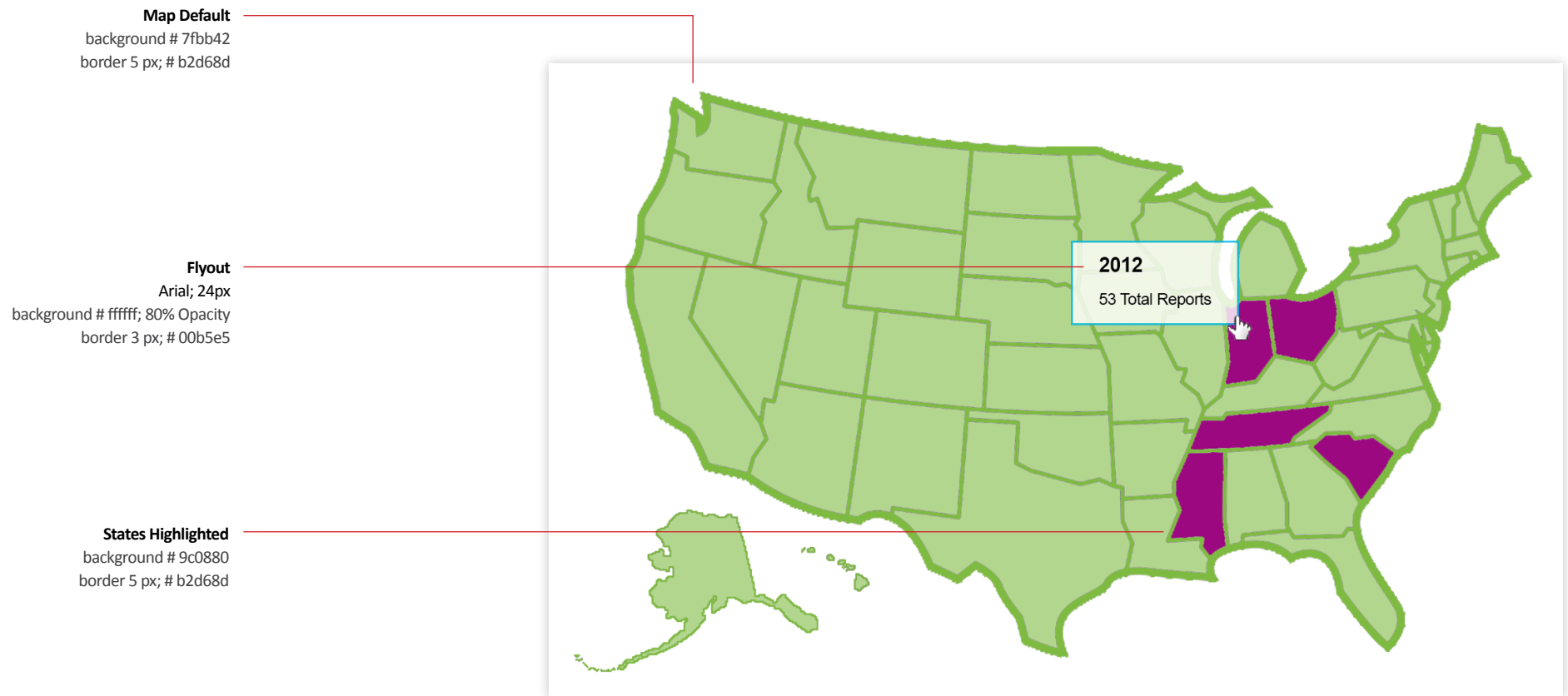
Sub Navigation Selected
background #00b5e5
Arial; bold; 28px; # ffffff

Search Box
Arial ; 30px; # 000000
border 1px; #00b5e5

Search Button
Arial ; 30px; # 000000
background # 00b5e5



MAP AND FLYOUT



Header

Arial; bold; 30px; #000000

About food recalls and enforcement reports

Recalls are an appropriate alternative method for removing or correcting marketed consumer products, their labeling, and/or promotional literature that violate the laws administered by the Food and Drug Administration (FDA). Recalls afford equal consumer protection but generally are more efficient and timely than formal administrative or civil actions, especially when the product has been widely distributed.

Body

Arial Narrow; 30px; # 000000
background # f4f4f4

Manufacturers and/or distributors may initiate a recall at any time to fulfill their responsibility to protect the public health from products that present a risk of injury or gross deception, or are otherwise defective. Firms may also initiate a recall following notification of a problem by FDA or a state agency, in response to a formal request by FDA, or as ordered by FDA.