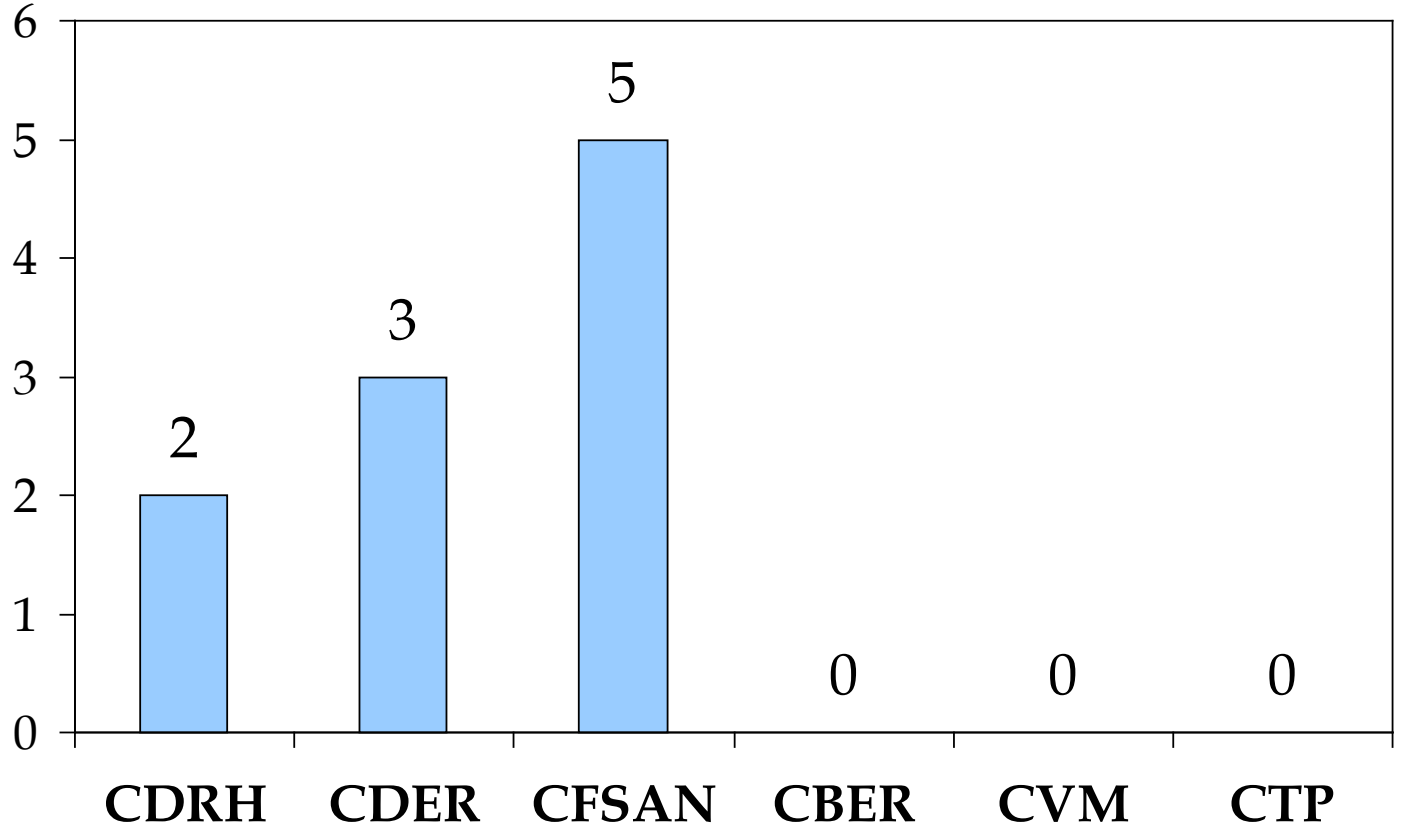
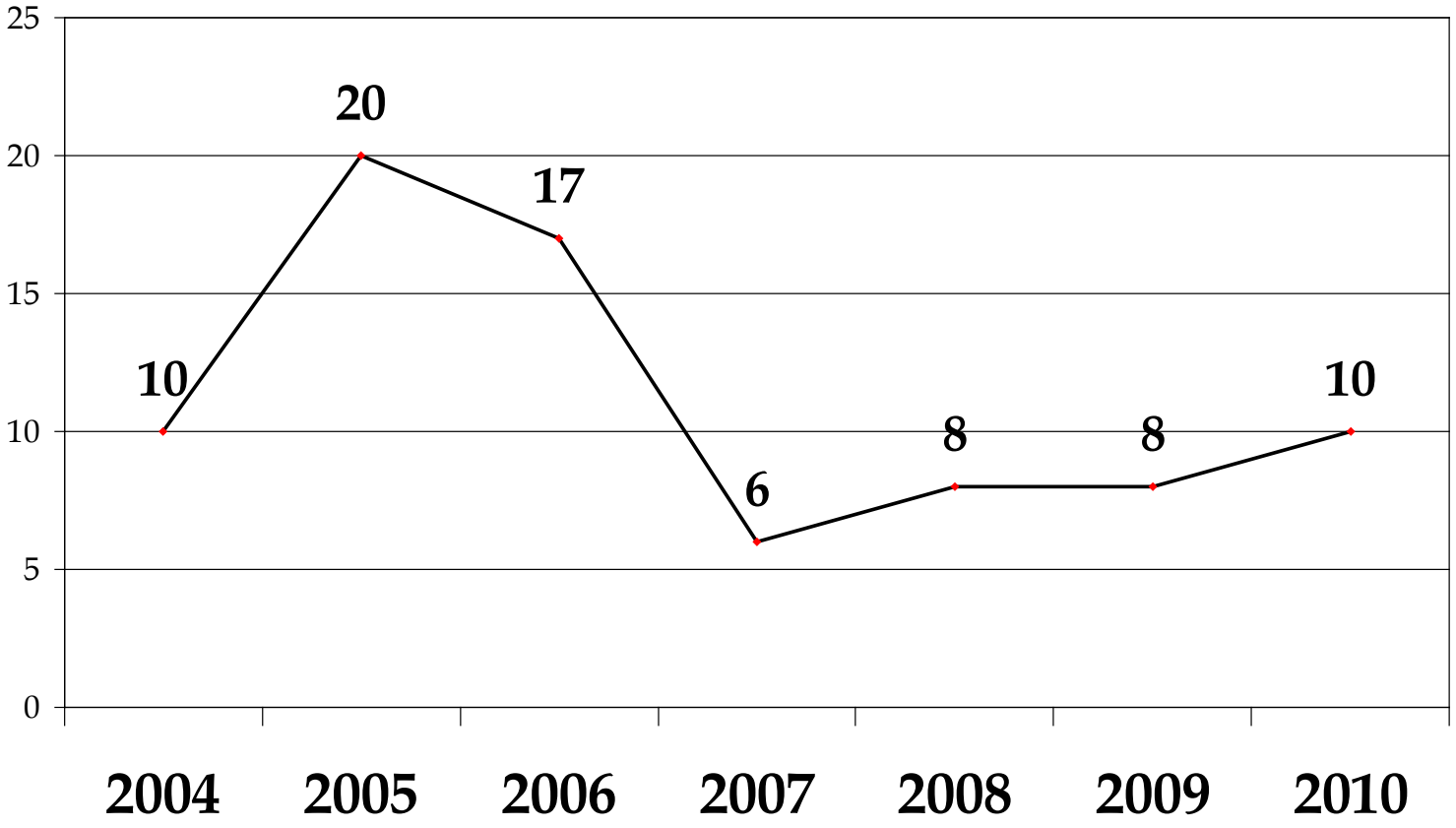
FDA Enforcement Statistics Summary Fiscal Year 2010

Seizures	10
Injunctions	17
Warning Letters	673
Recall Events	3,799
Recalled Products	9,361
Debarments	13

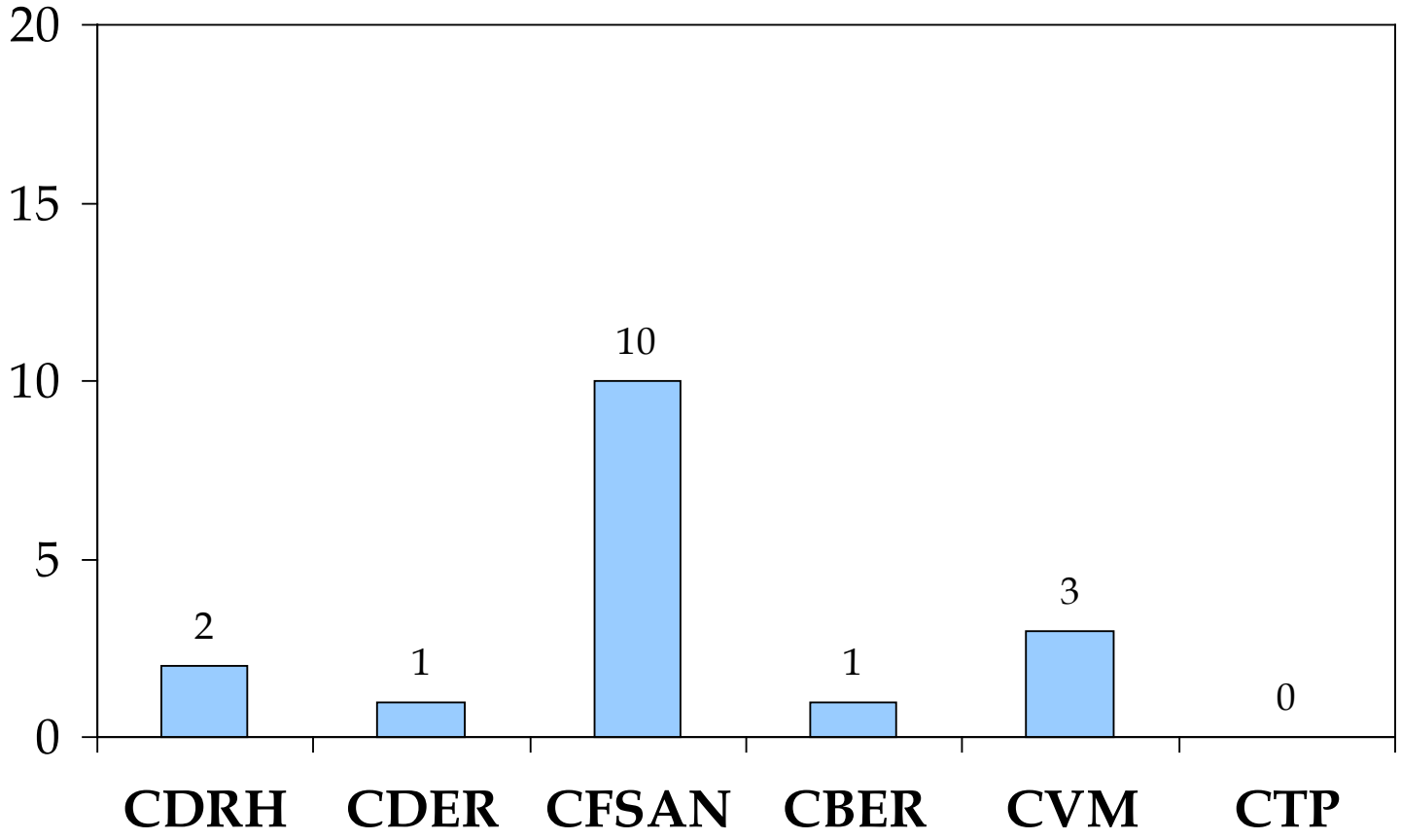
Seizures by FDA Center
Fiscal Year 2010



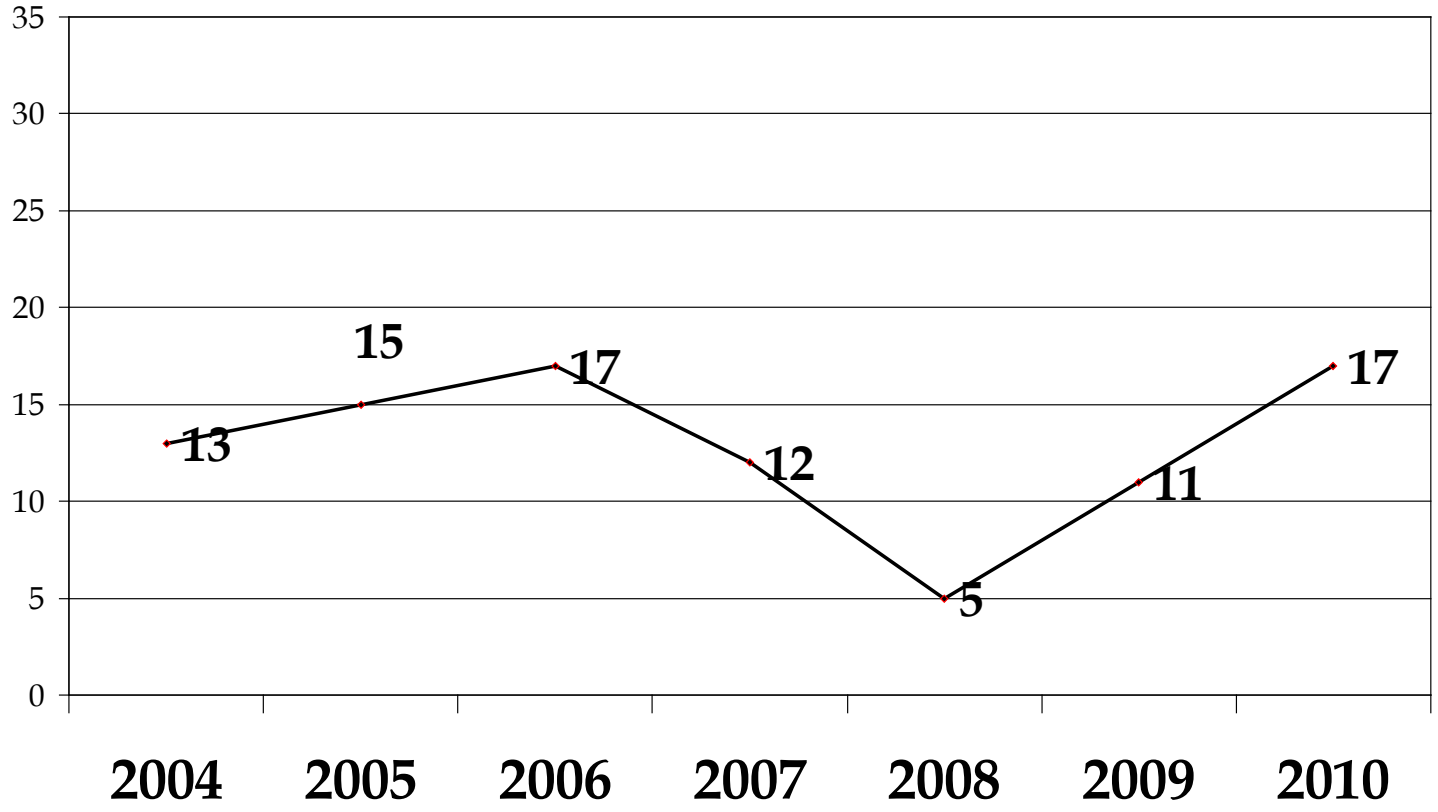
FDA Seizures
Fiscal Years 2004 – 2010



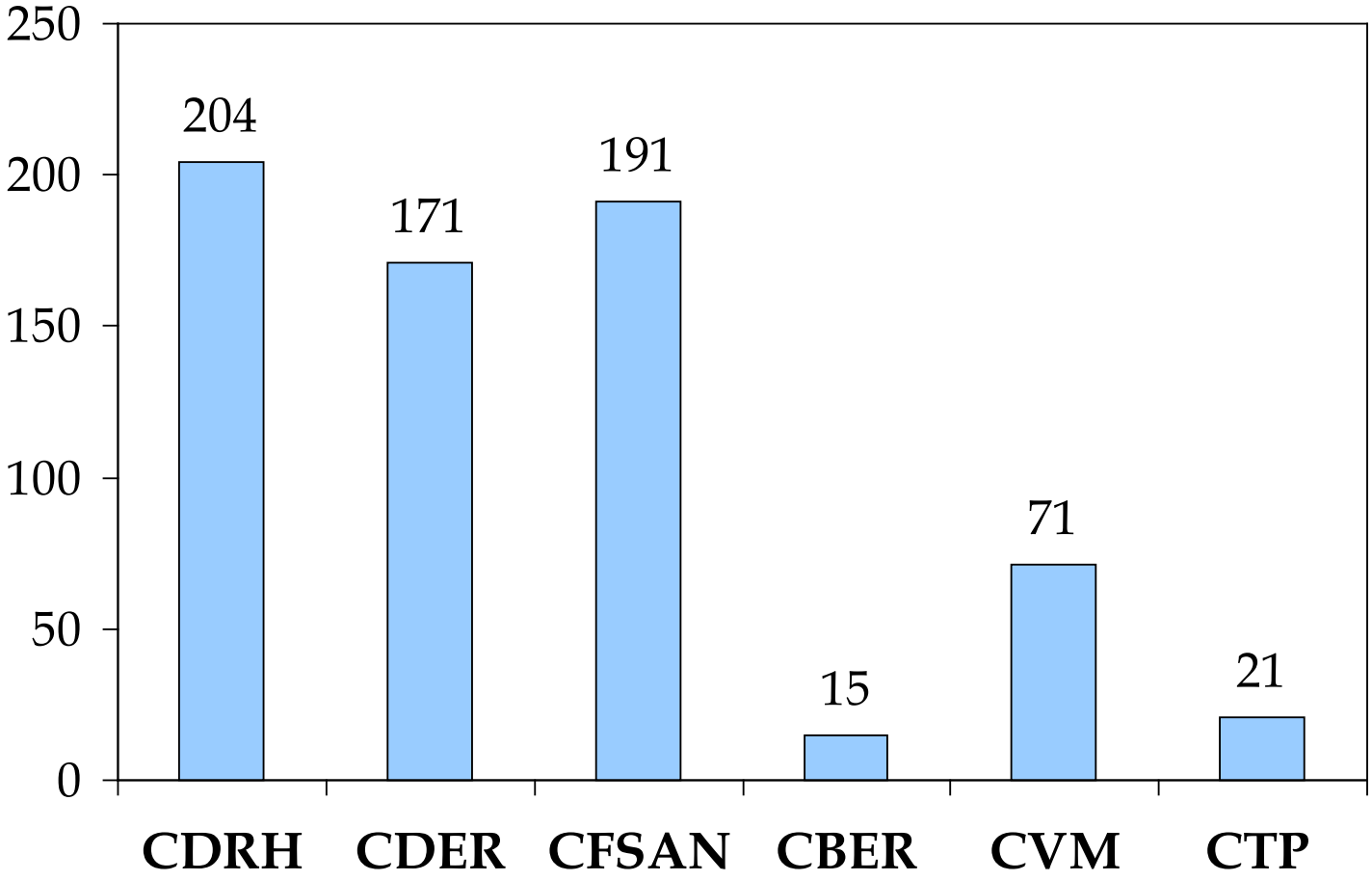
Injunctions by FDA Center
Fiscal Year 2010



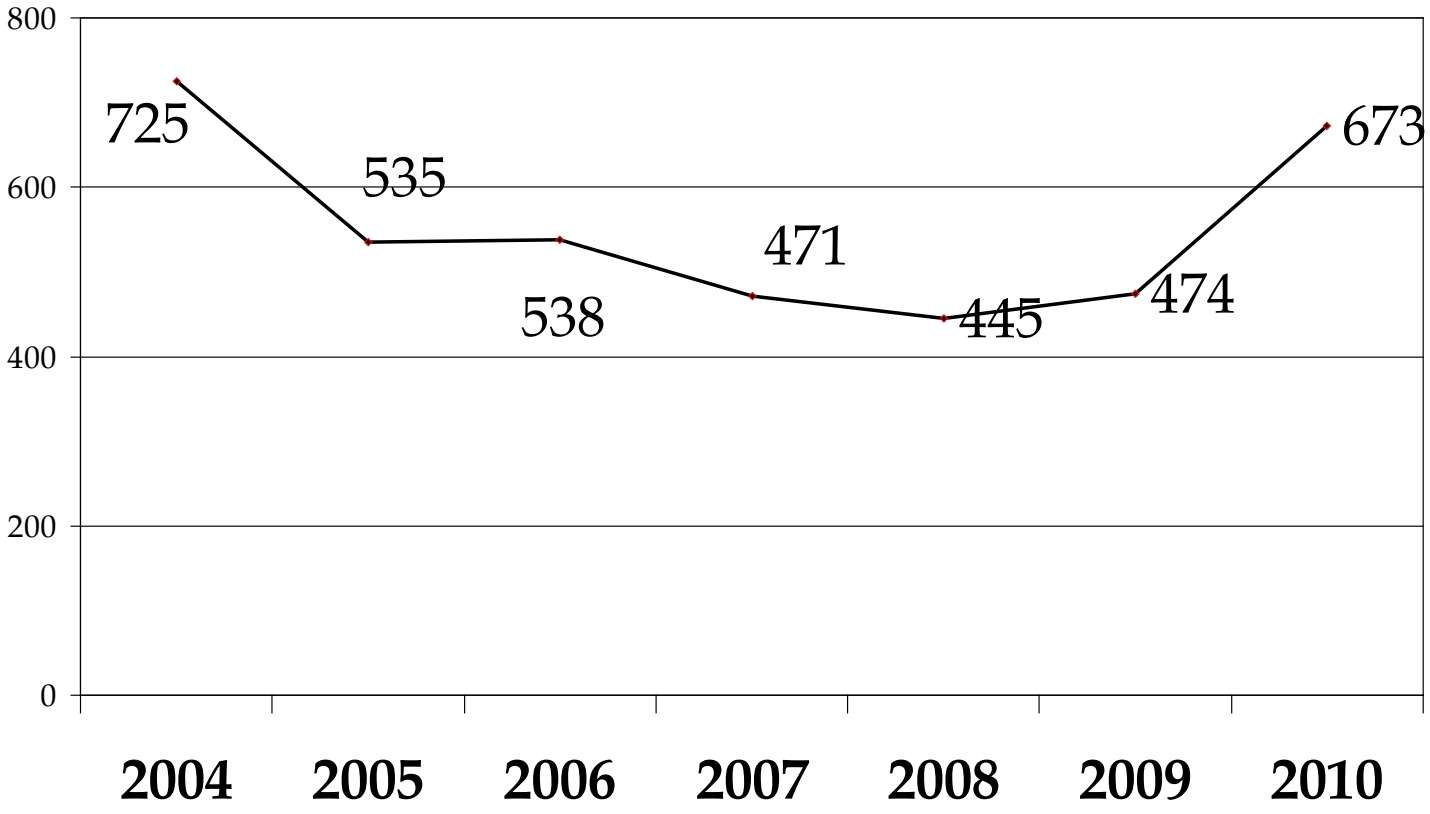
FDA Injunctions
Fiscal Years 2004 - 2010



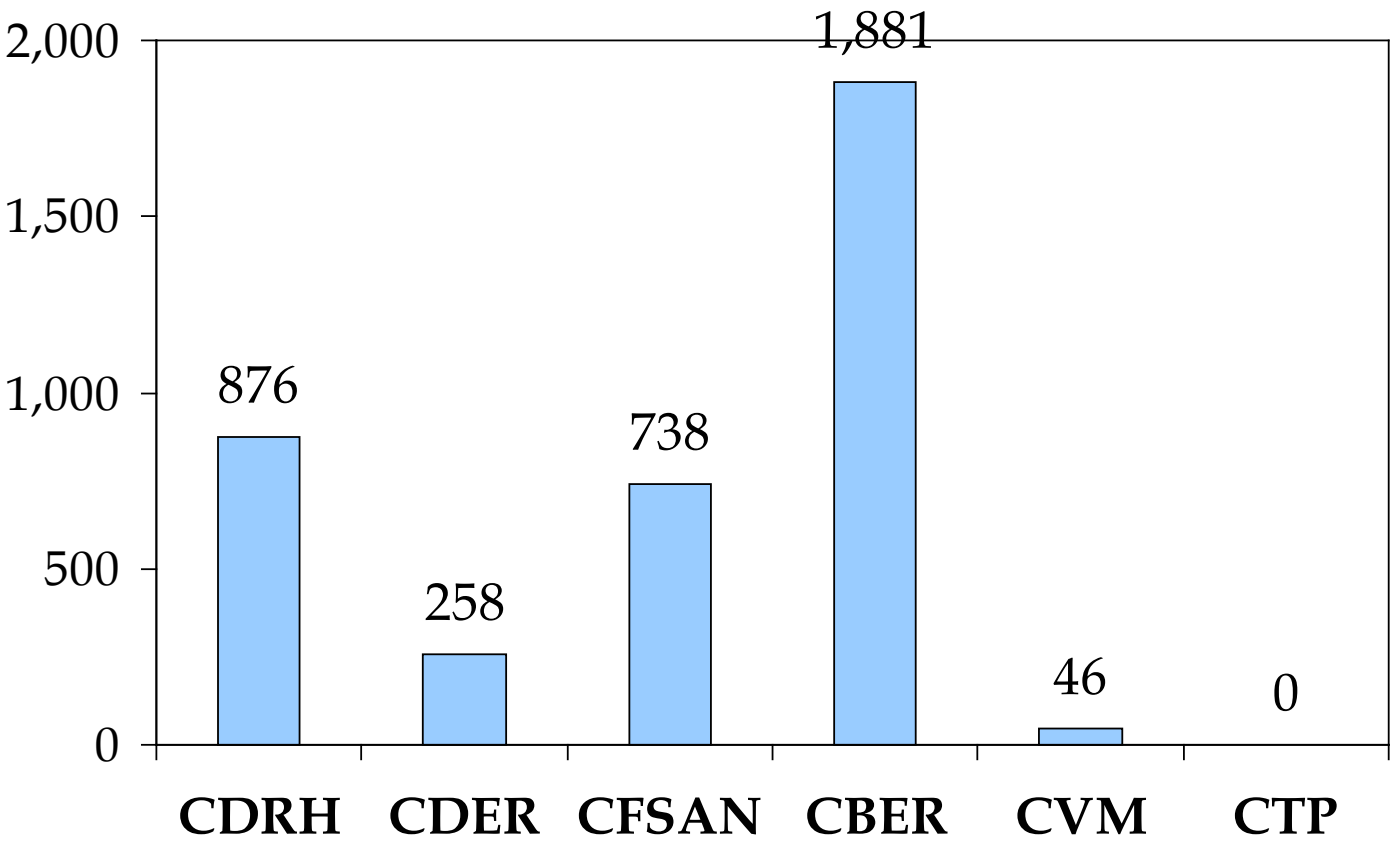
Warning Letters by FDA Center
Fiscal Year 2010



FDA Warning Letters
Fiscal Years 2004 – 2010

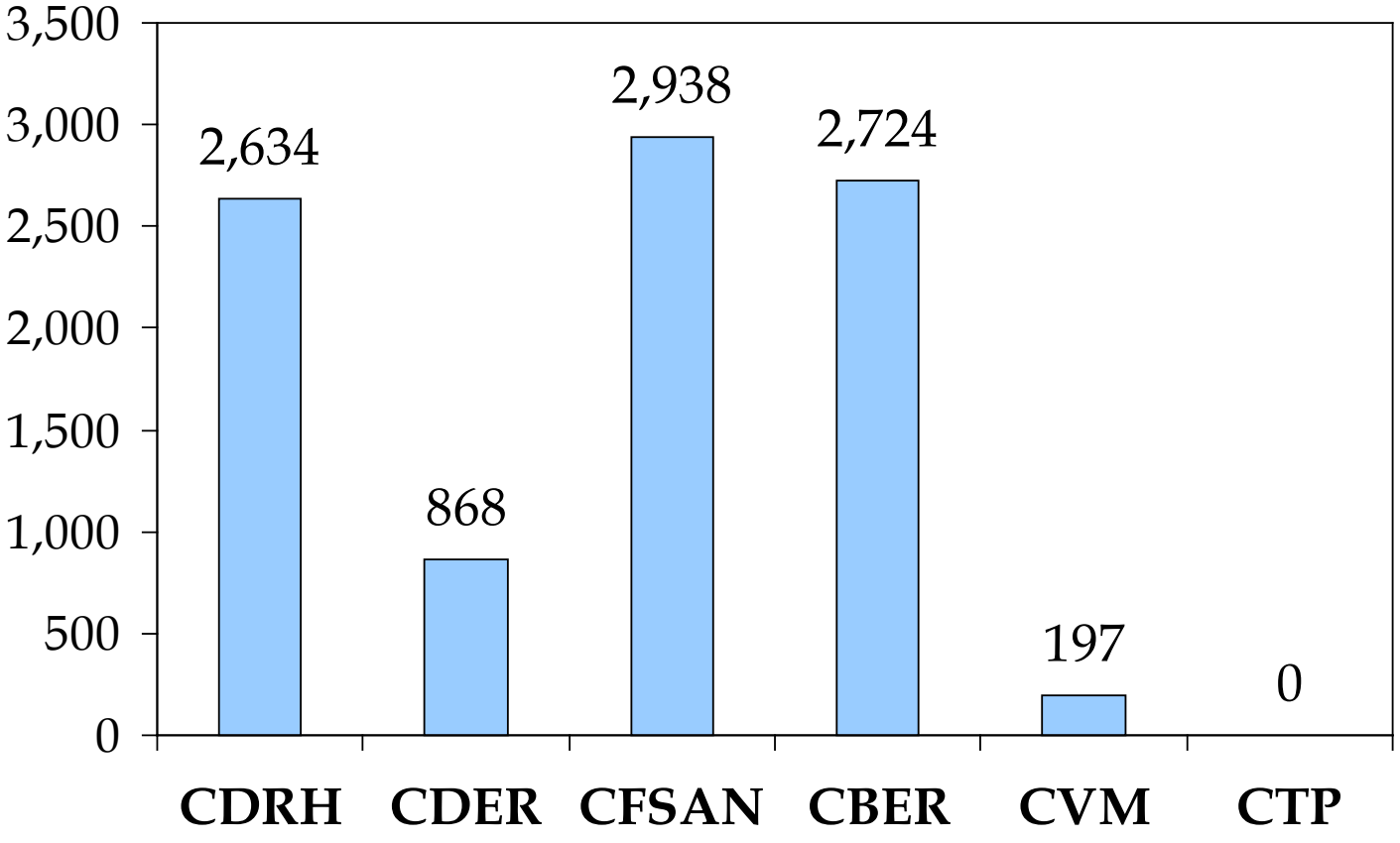


Total Recall Events by FDA Center
Fiscal Year 2010



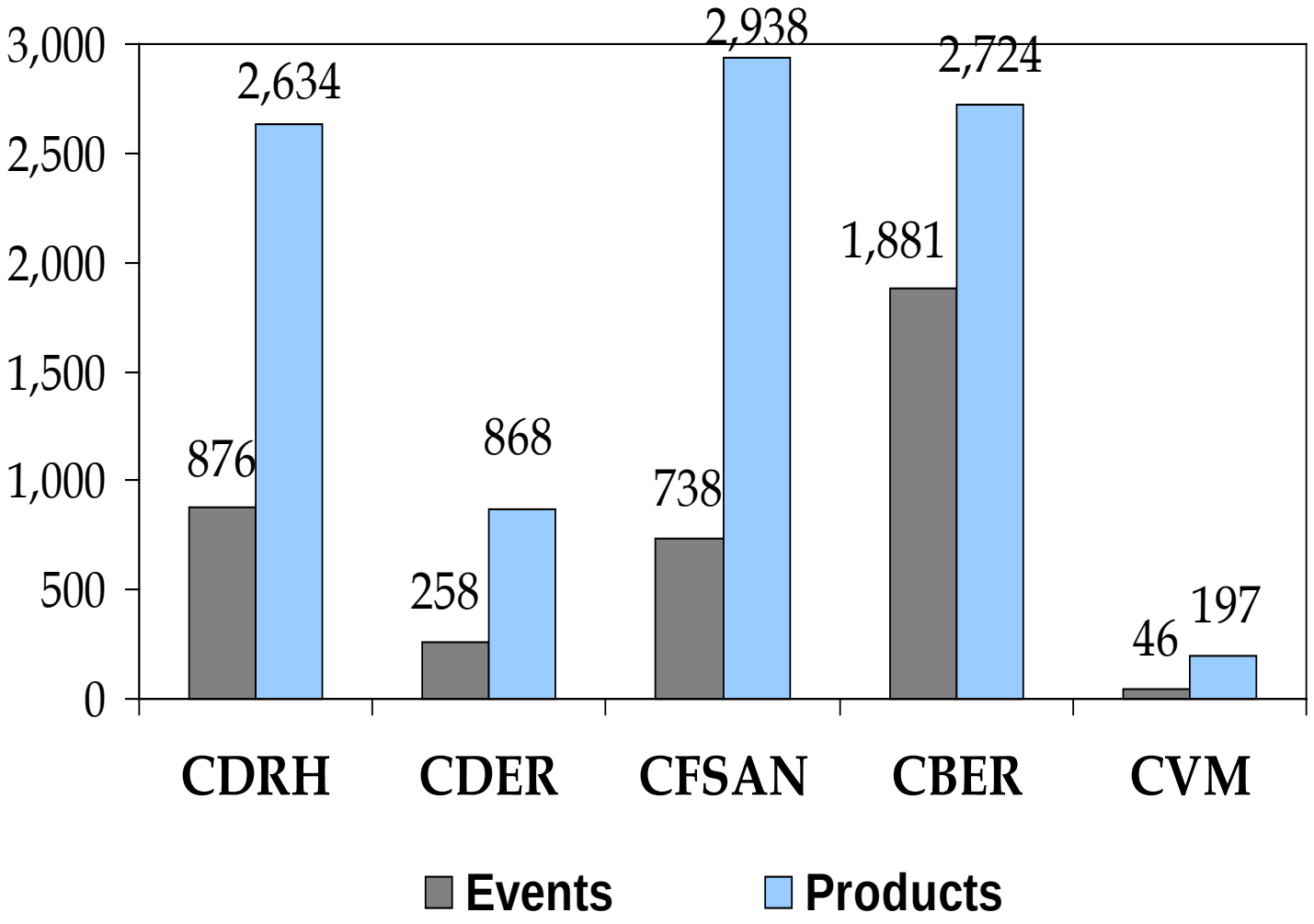
Class I, II and III

Total Recalled Products by FDA Center
Fiscal Year 2010

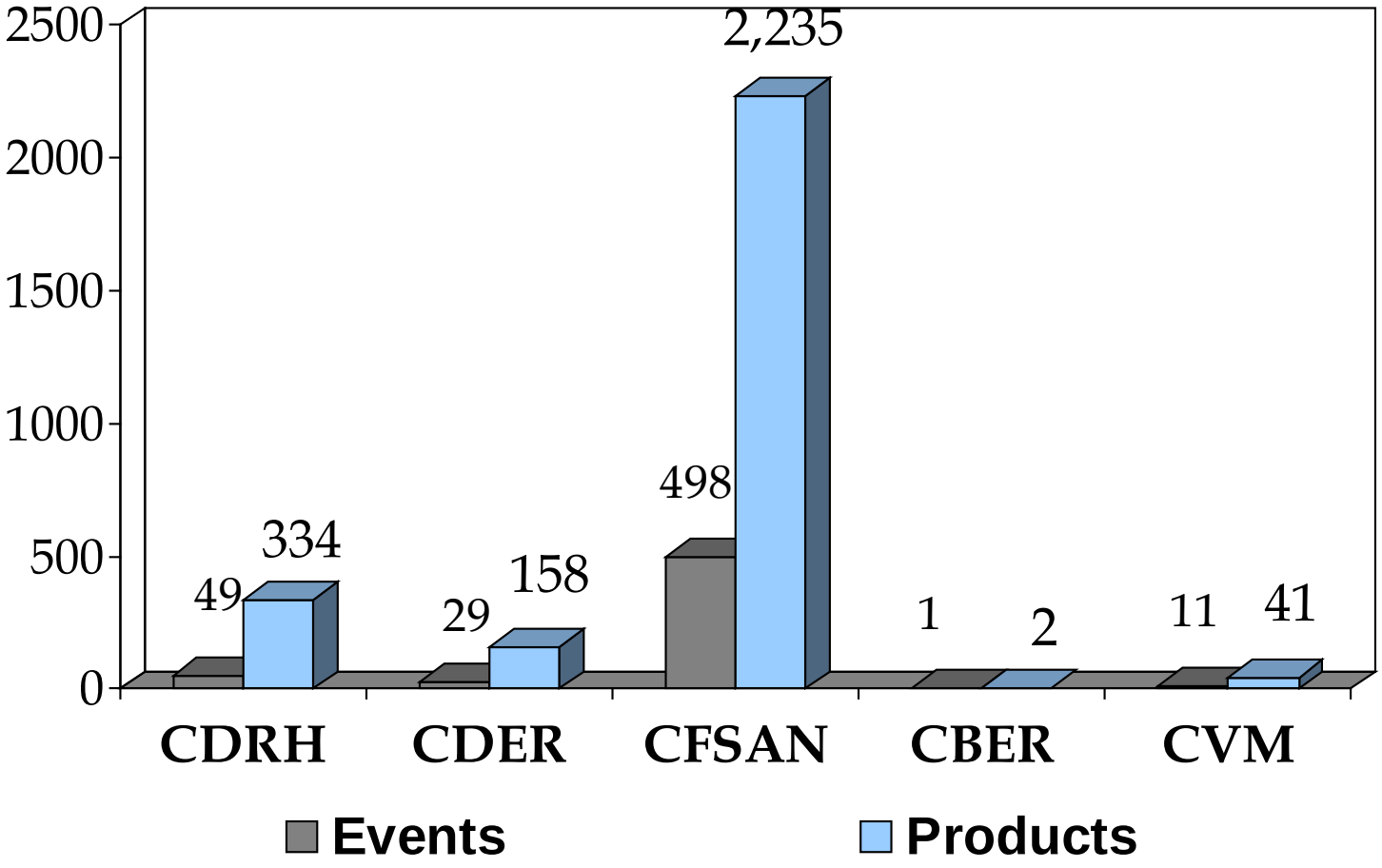


Class I, II and III

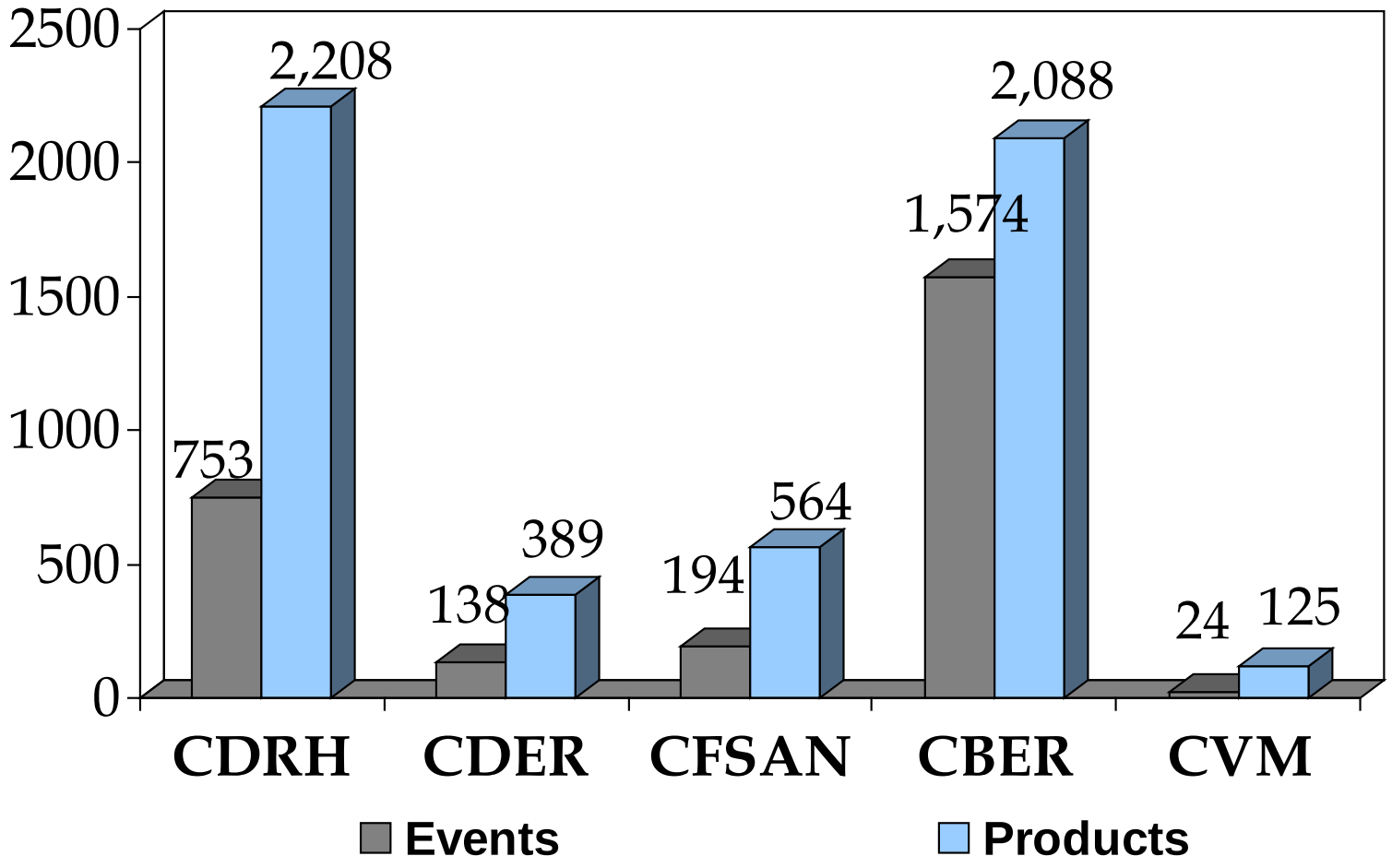
FDA Recalls By Center - All Classes
Fiscal Year 2010



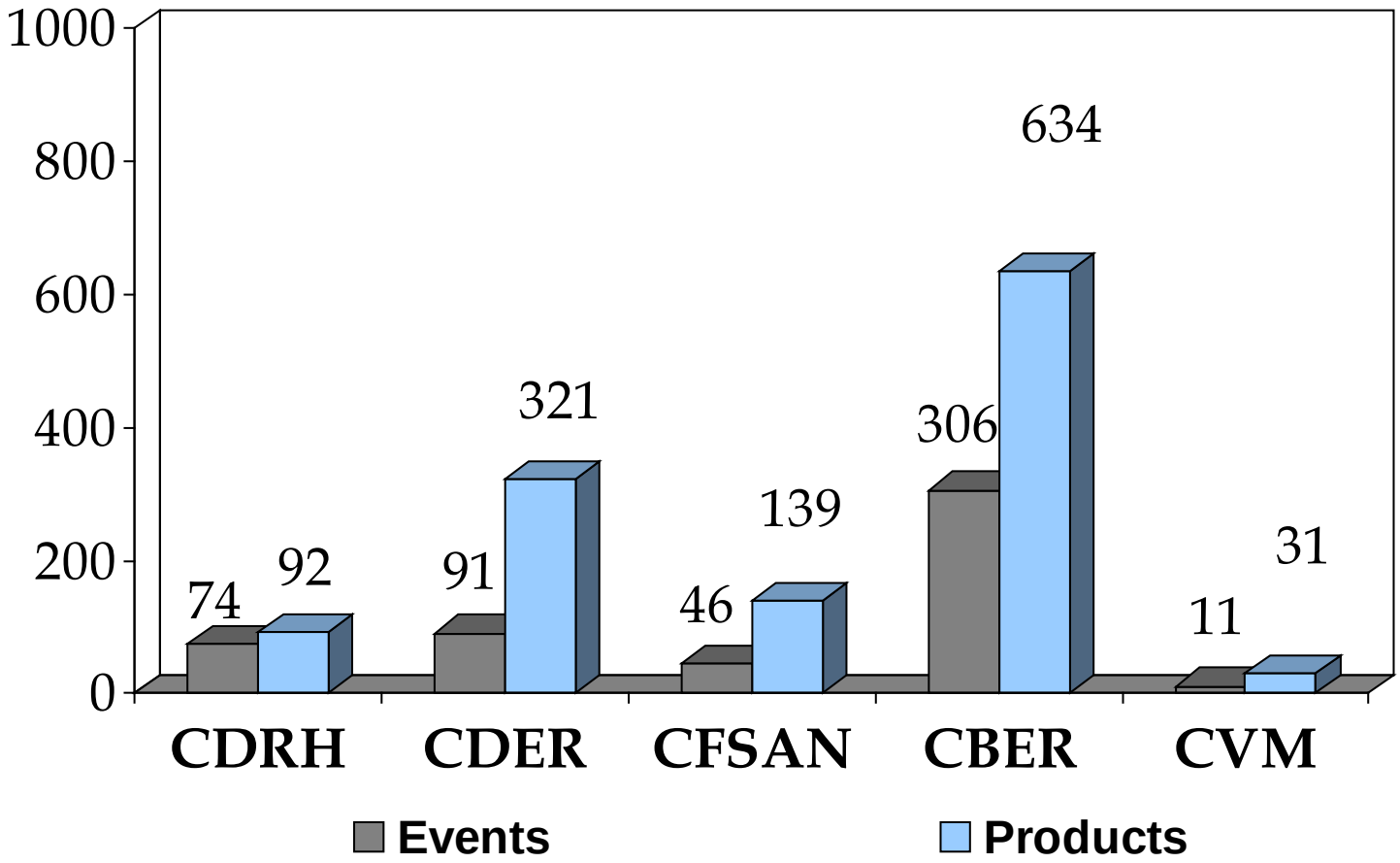
FDA Recalls - Fiscal Year 2010 Class I By Center



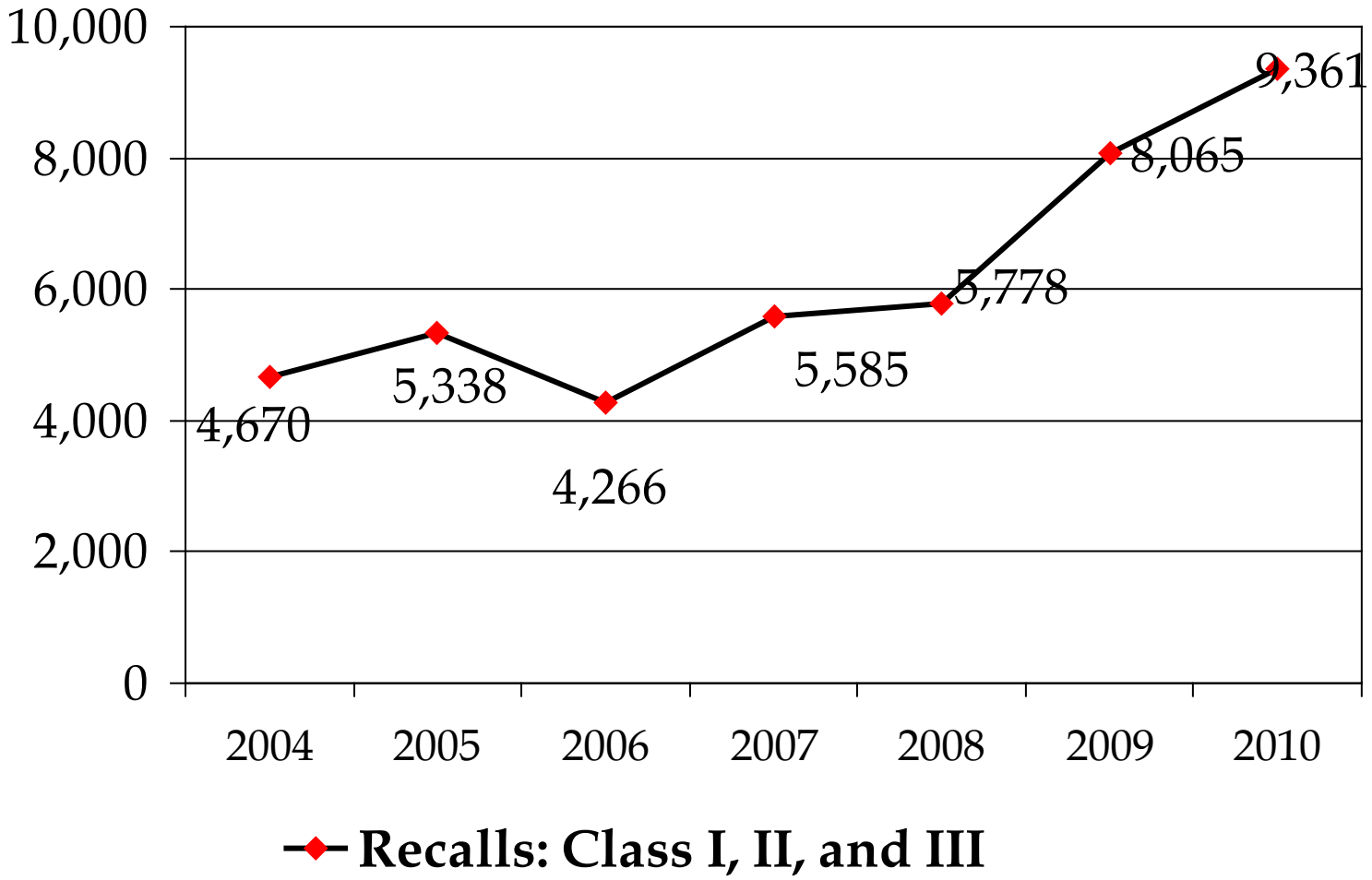
FDA Recalls - Fiscal Year 2010
Class II By Center



FDA Recalls - Fiscal Year 2010 Class III By Center



Recalled Products – All Centers
Fiscal Years 2004 – 2010



Recalls: Definition of Class I, II and III

Class I recall: a situation in which there is a reasonable probability that the use of or exposure to a violative product will cause serious adverse health consequences or death.

Class II recall: a situation in which use of or exposure to a violative product may cause temporary or medically reversible adverse health consequences or where the probability of serious adverse health consequences is remote.

Class III recall: a situation in which use of or exposure to a violative product is not likely to cause adverse health consequences.