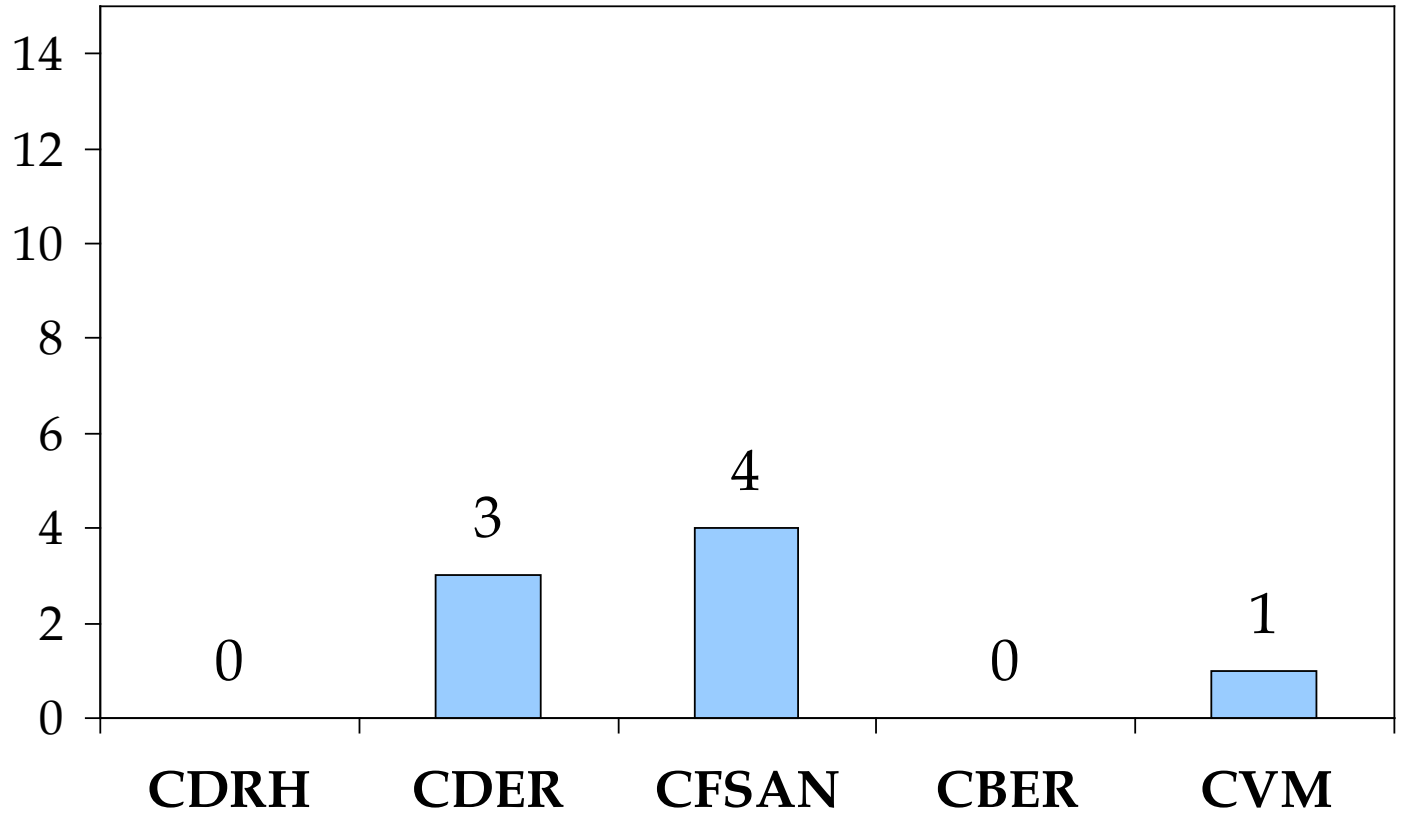
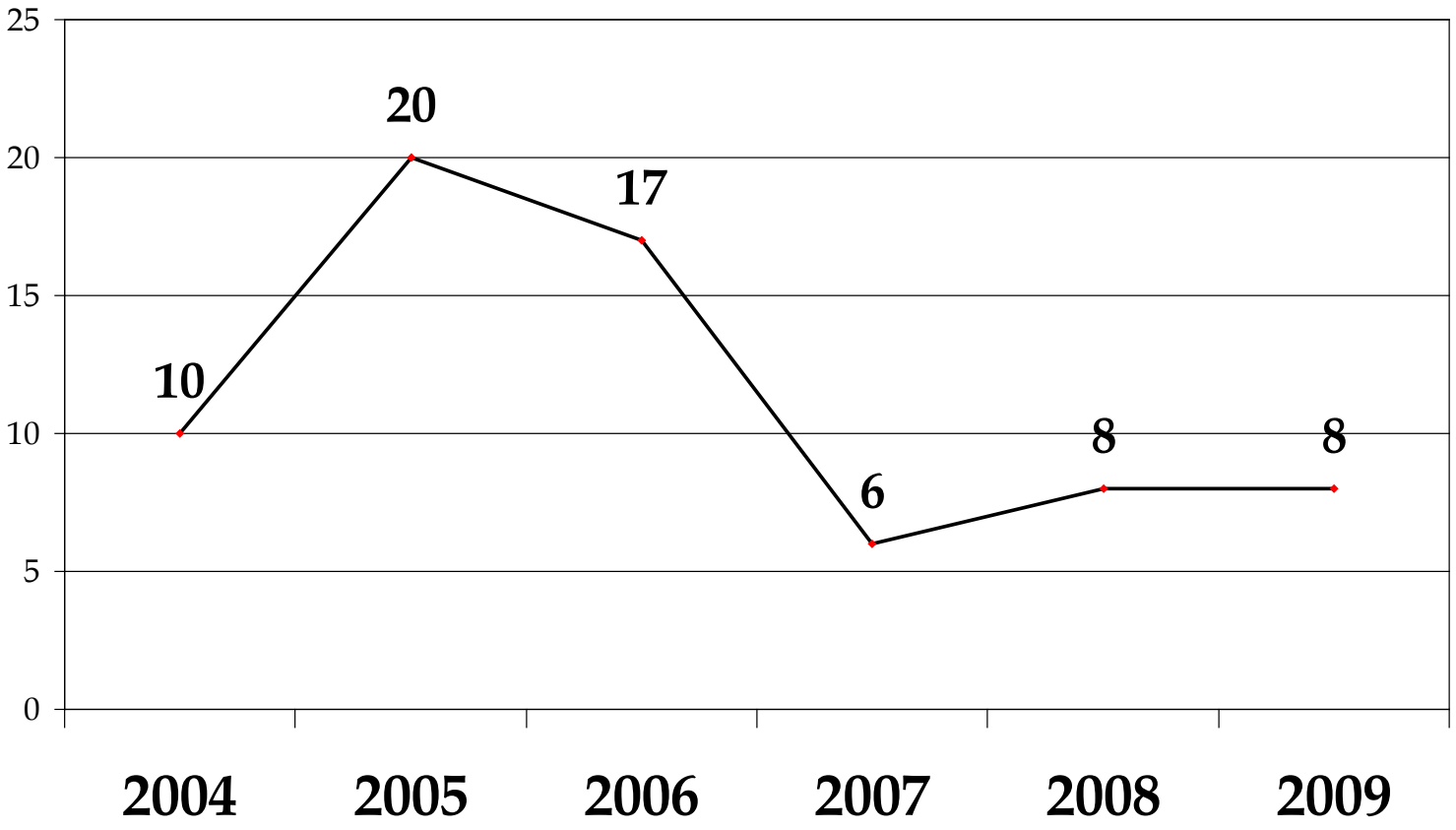
FDA Enforcement Statistics Summary Fiscal Year 2009

Seizures	8
Injunctions	11
Warning Letters	474
Recall Events	2,781
Recalled Products	8,065
Debarments	6

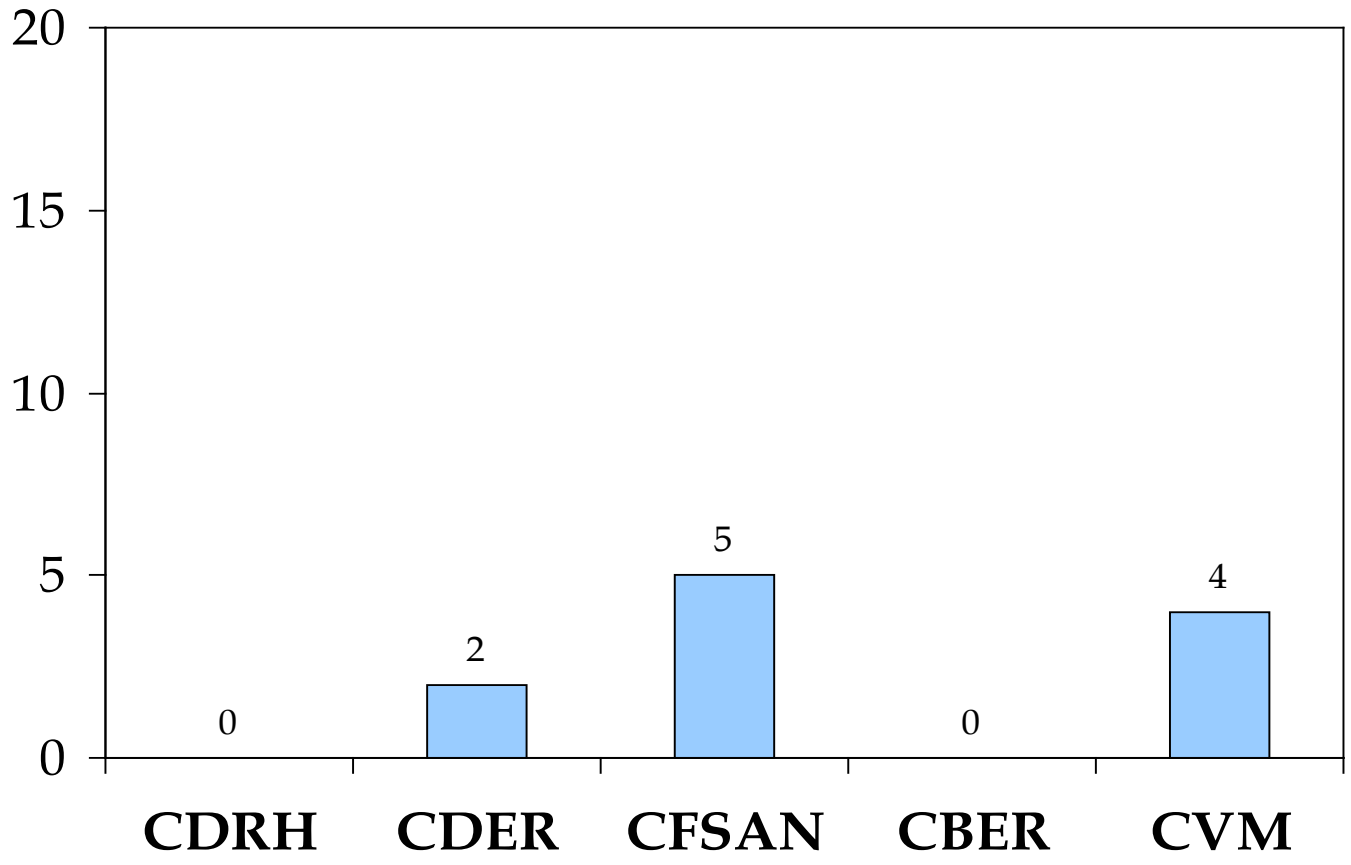
Seizures by FDA Center
Fiscal Year 2009



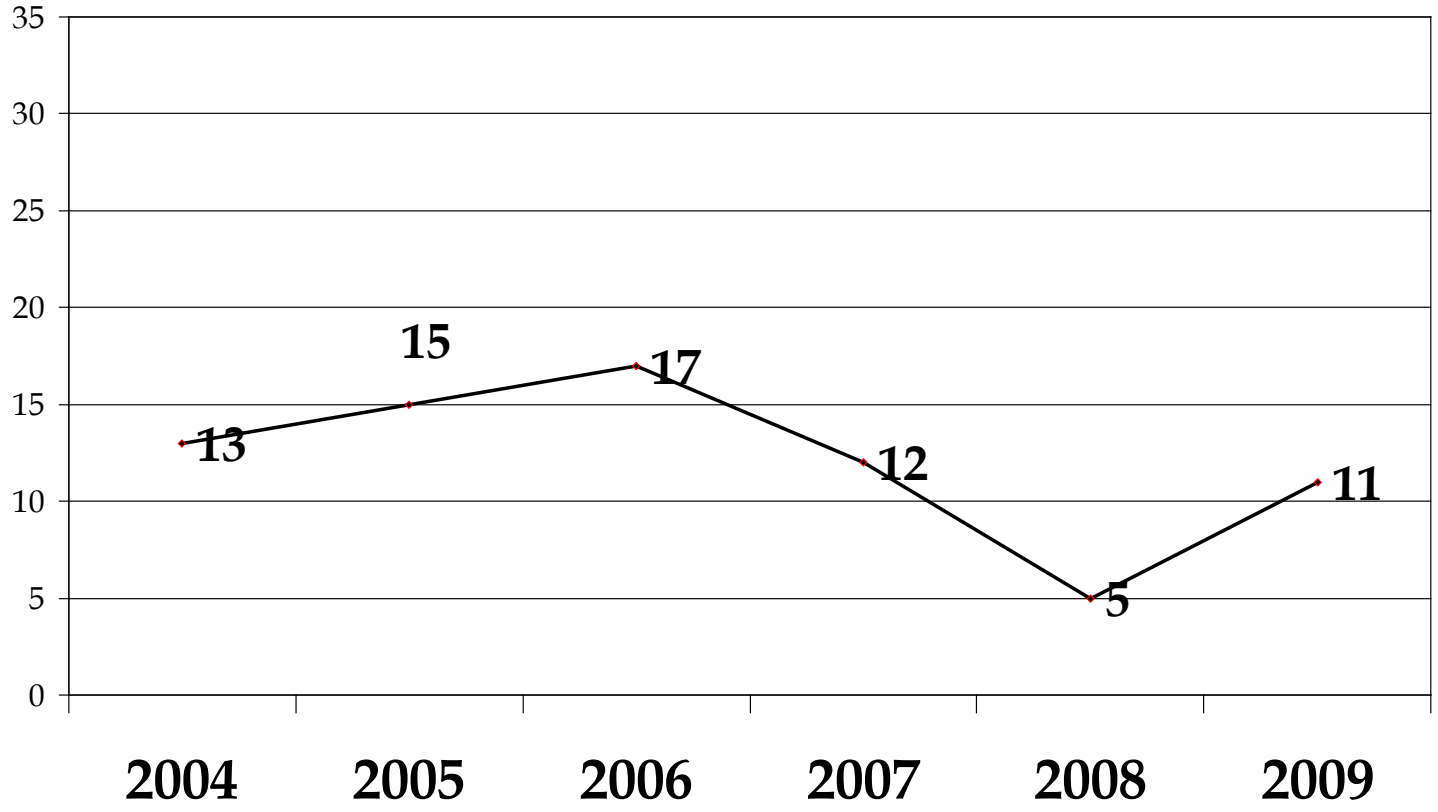
FDA Seizures
Fiscal Years 2004 – 2009



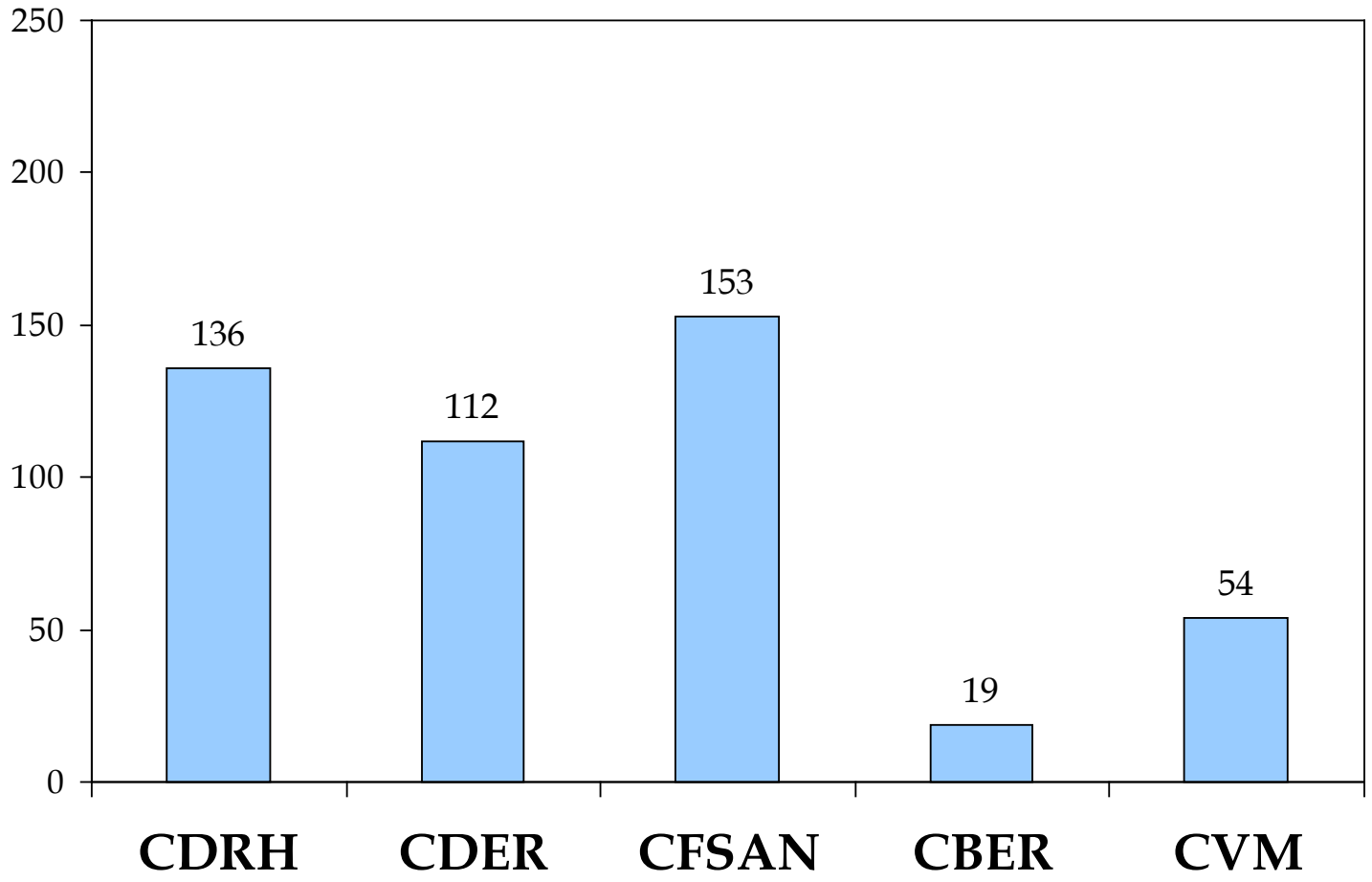
Injunctions by FDA Center
Fiscal Year 2009



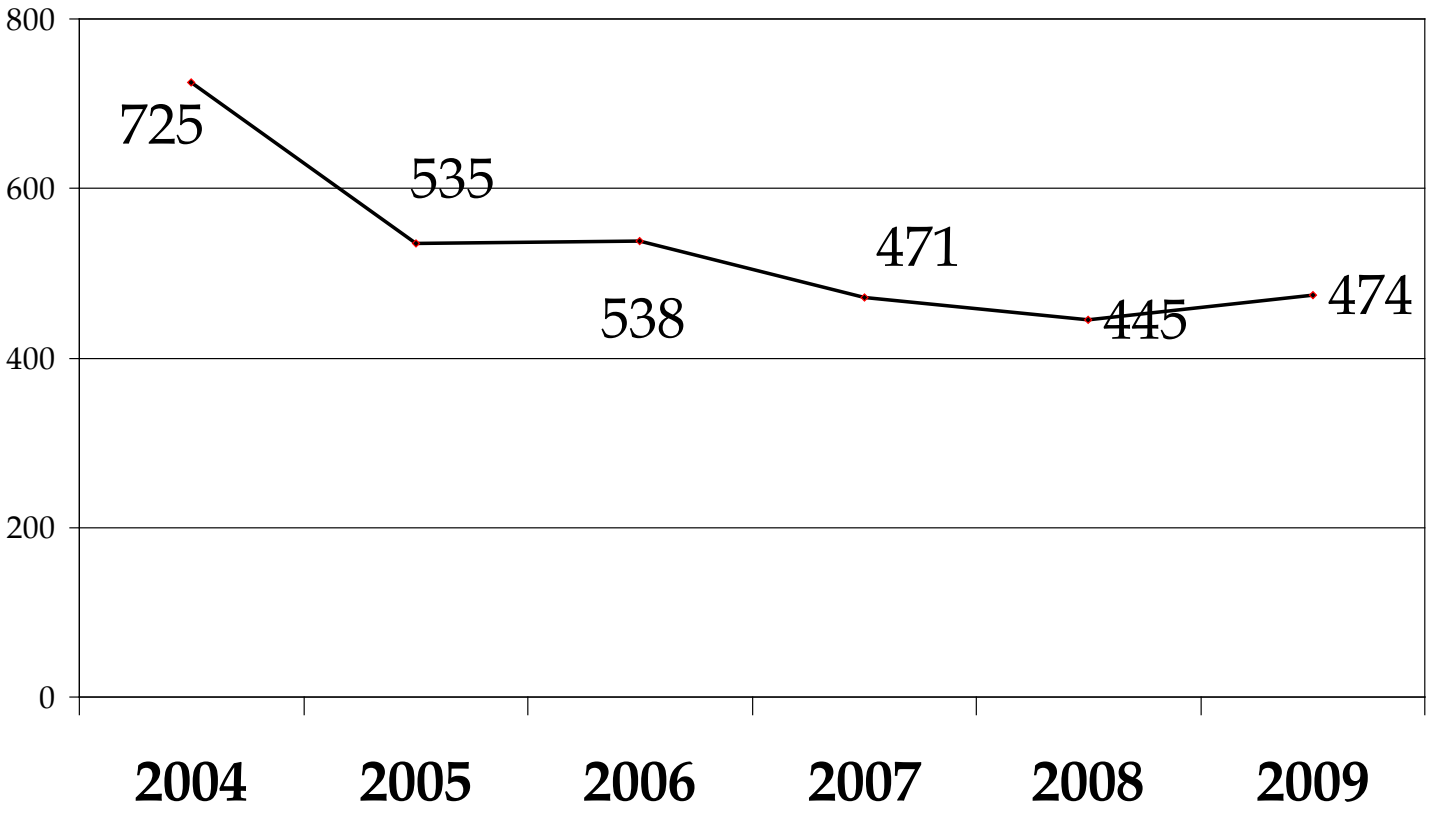
FDA Injunctions
Fiscal Years 2004 - 2009



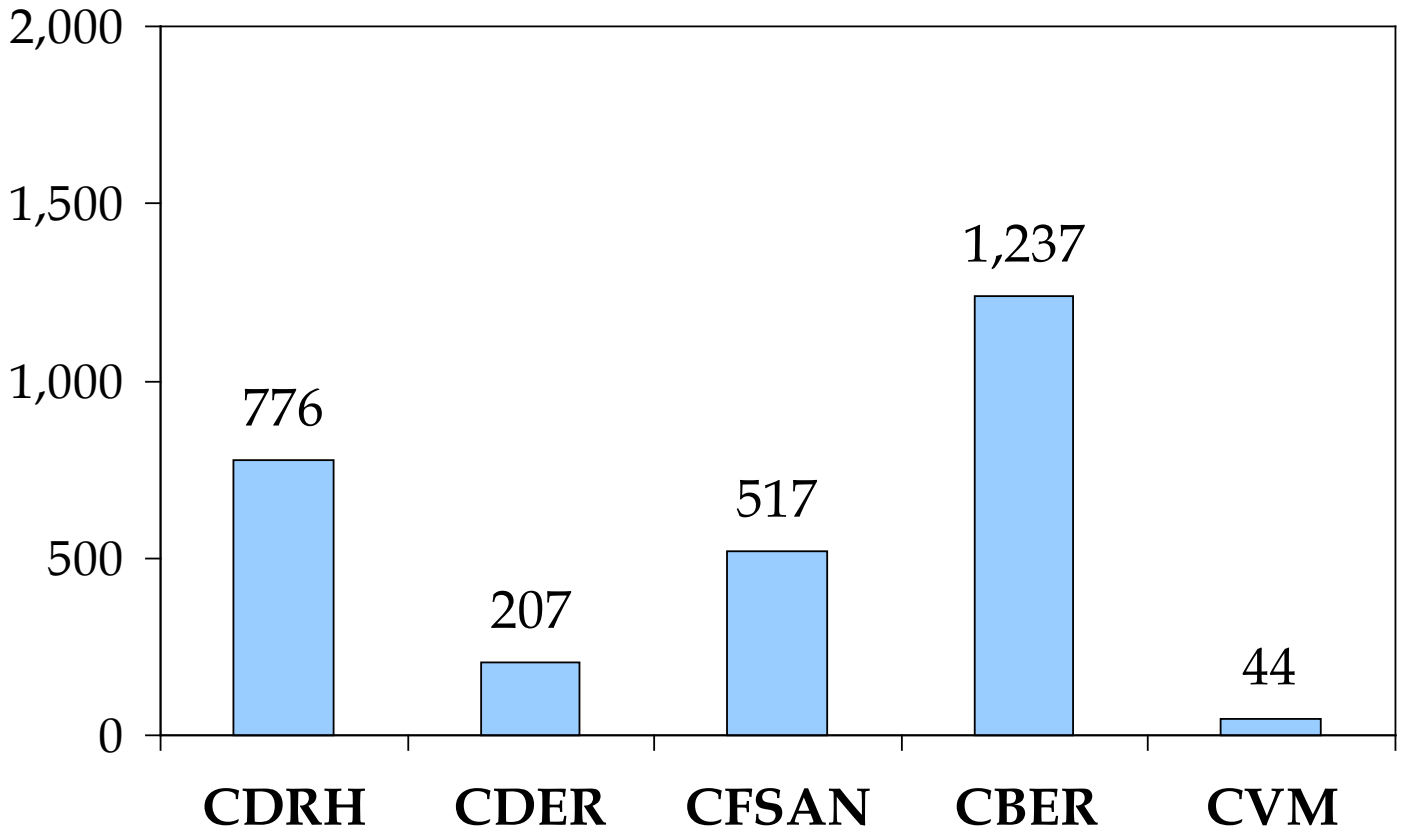
Warning Letters by FDA Center
Fiscal Year 2009



FDA Warning Letters
Fiscal Years 2004 – 2009

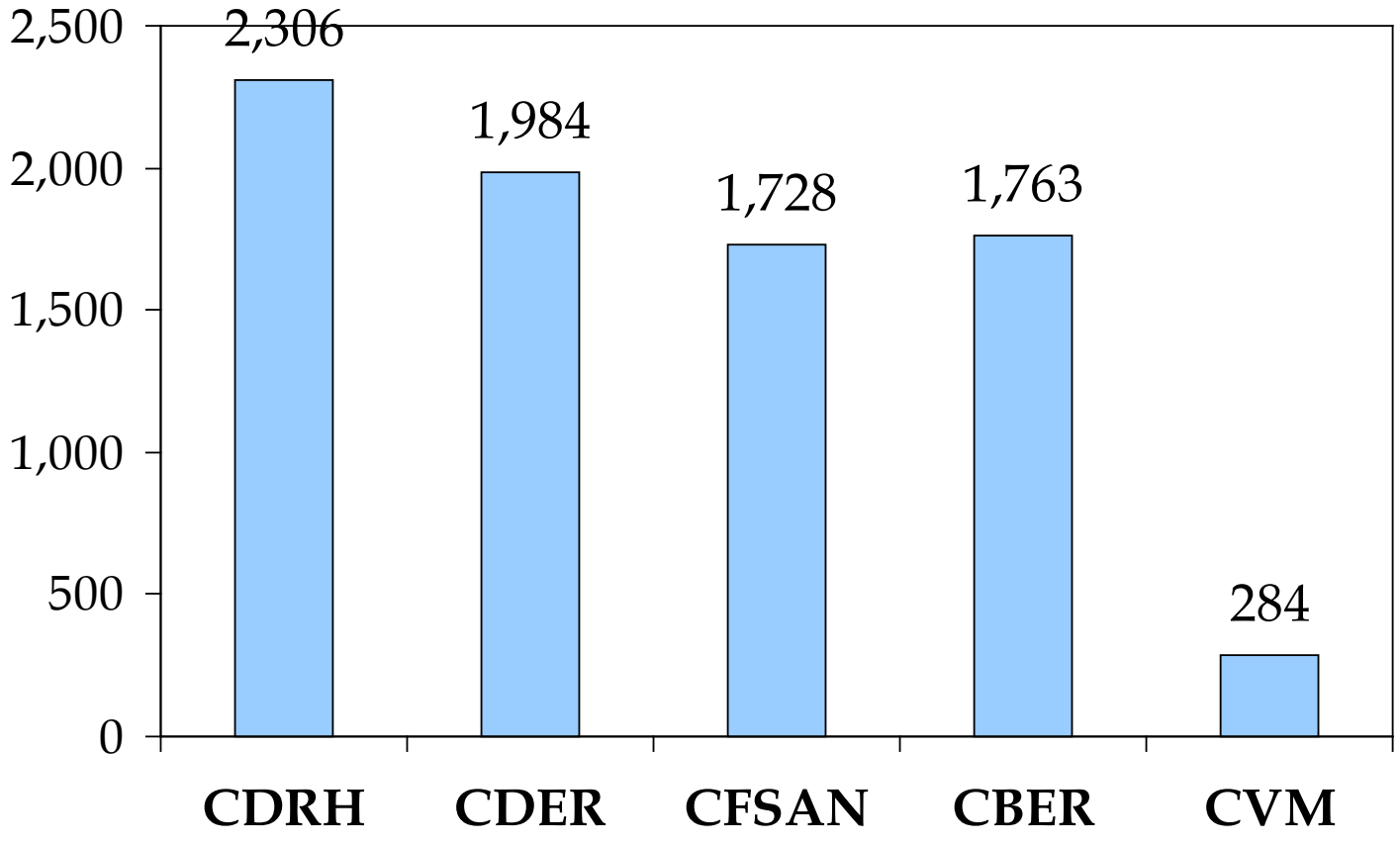


Total Recall Events by FDA Center
Fiscal Year 2009



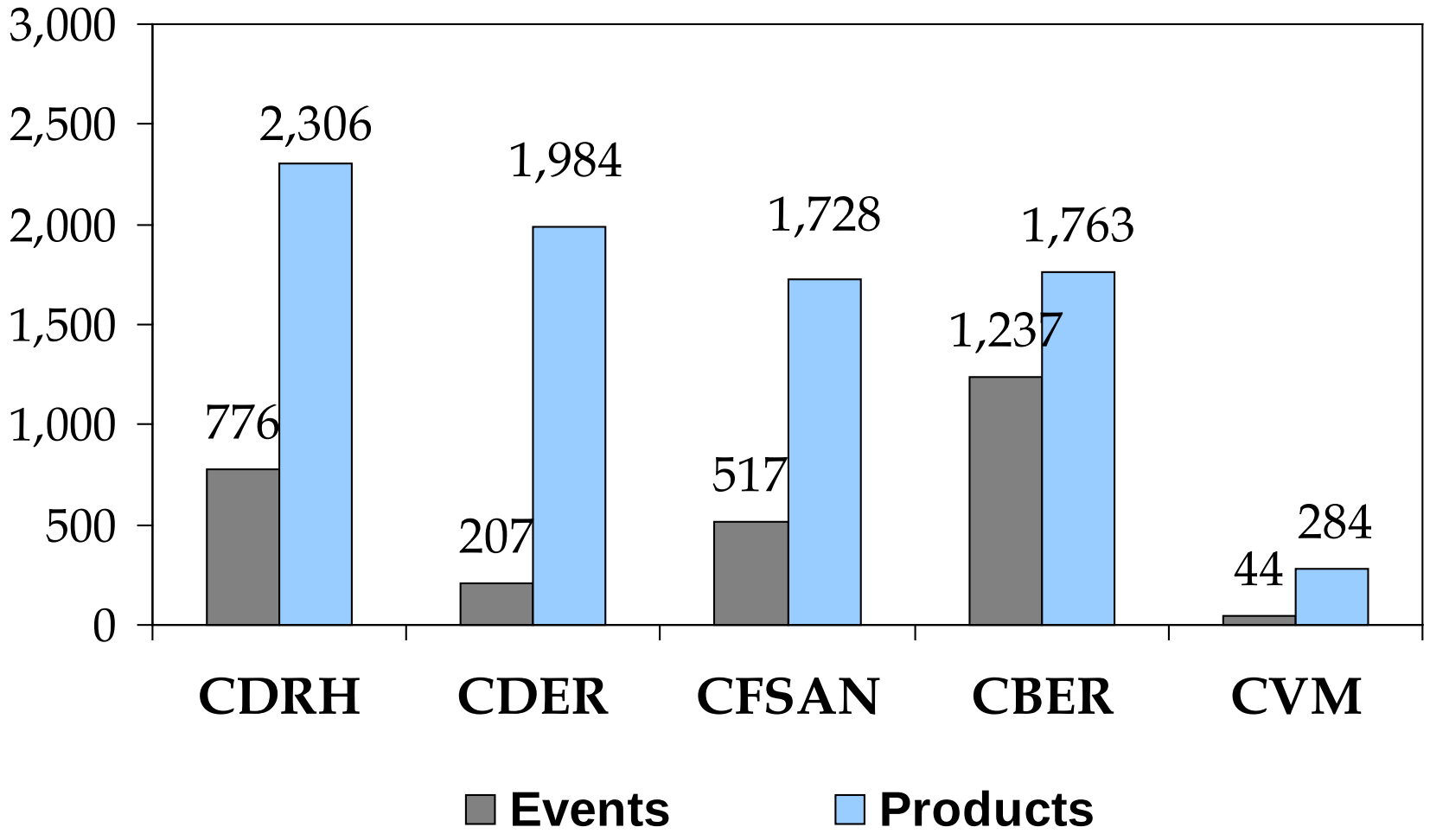
Class I, II and III

Total Recalled Products by FDA Center Fiscal Year 2009

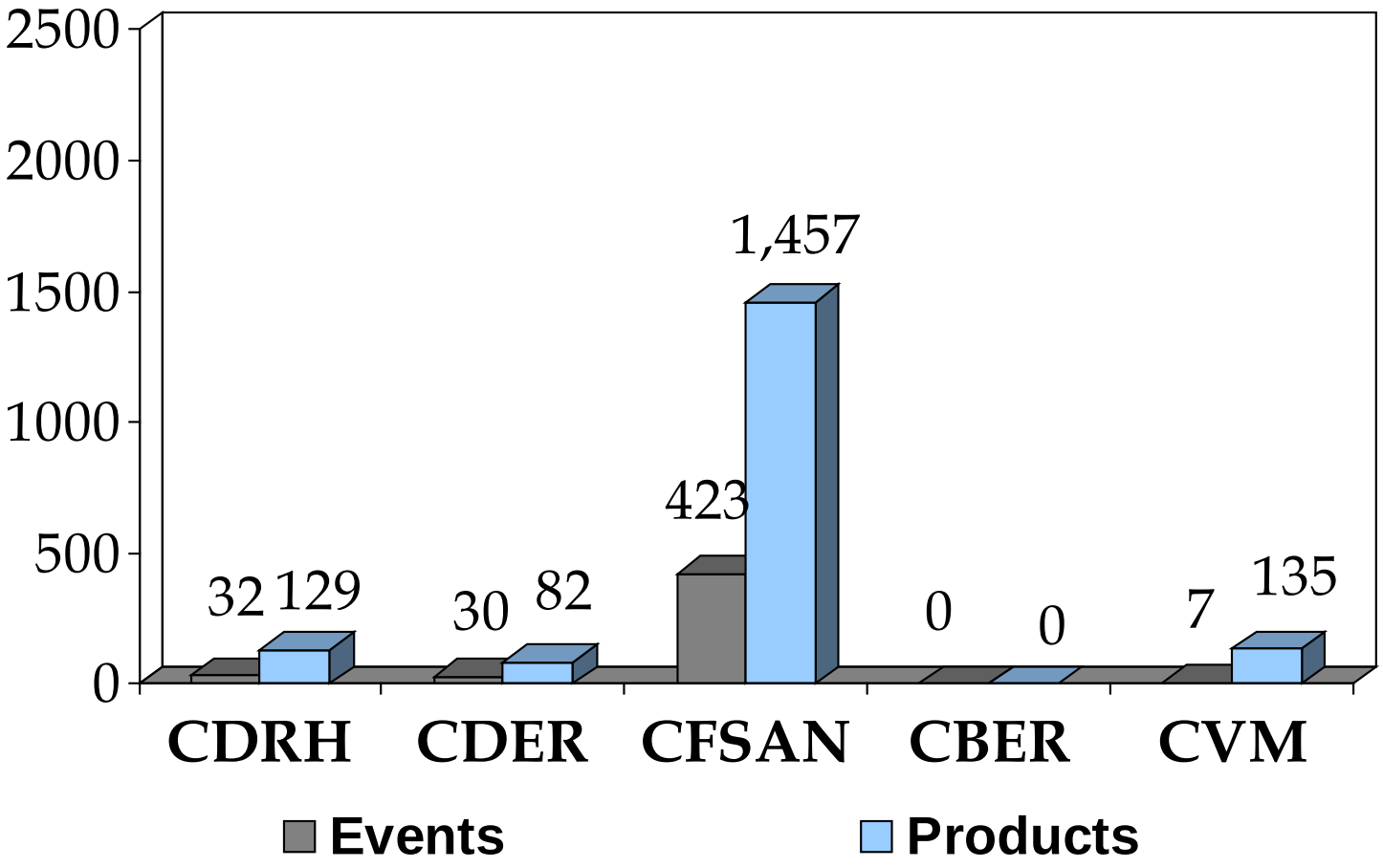


Class I, II and III

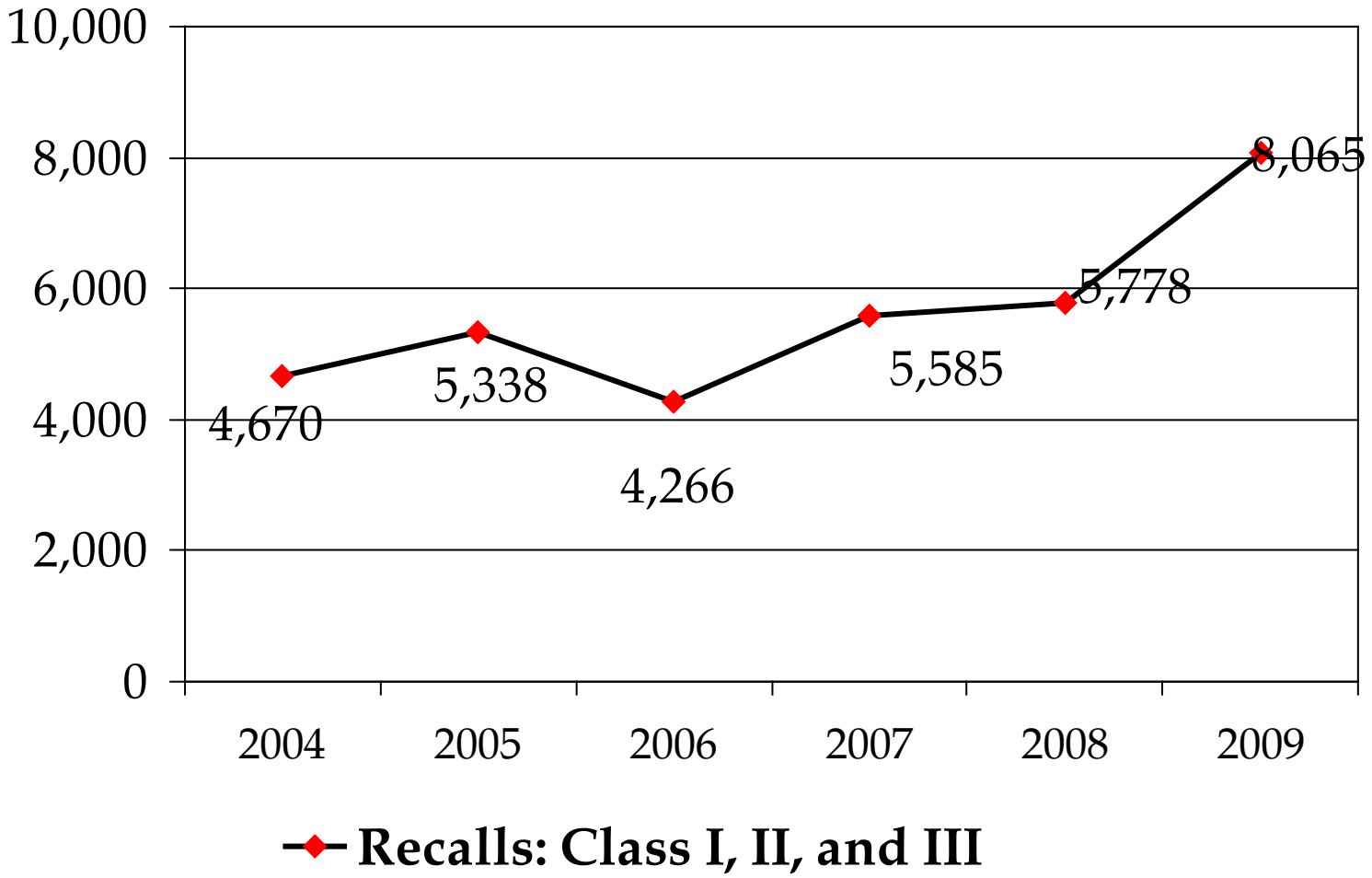
FDA Recalls By Center - All Classes
Fiscal Year 2009



FDA Recalls - Fiscal Year 2009 Class I By Center



Recalled Products – All Centers
Fiscal Years 2004 – 2009



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.