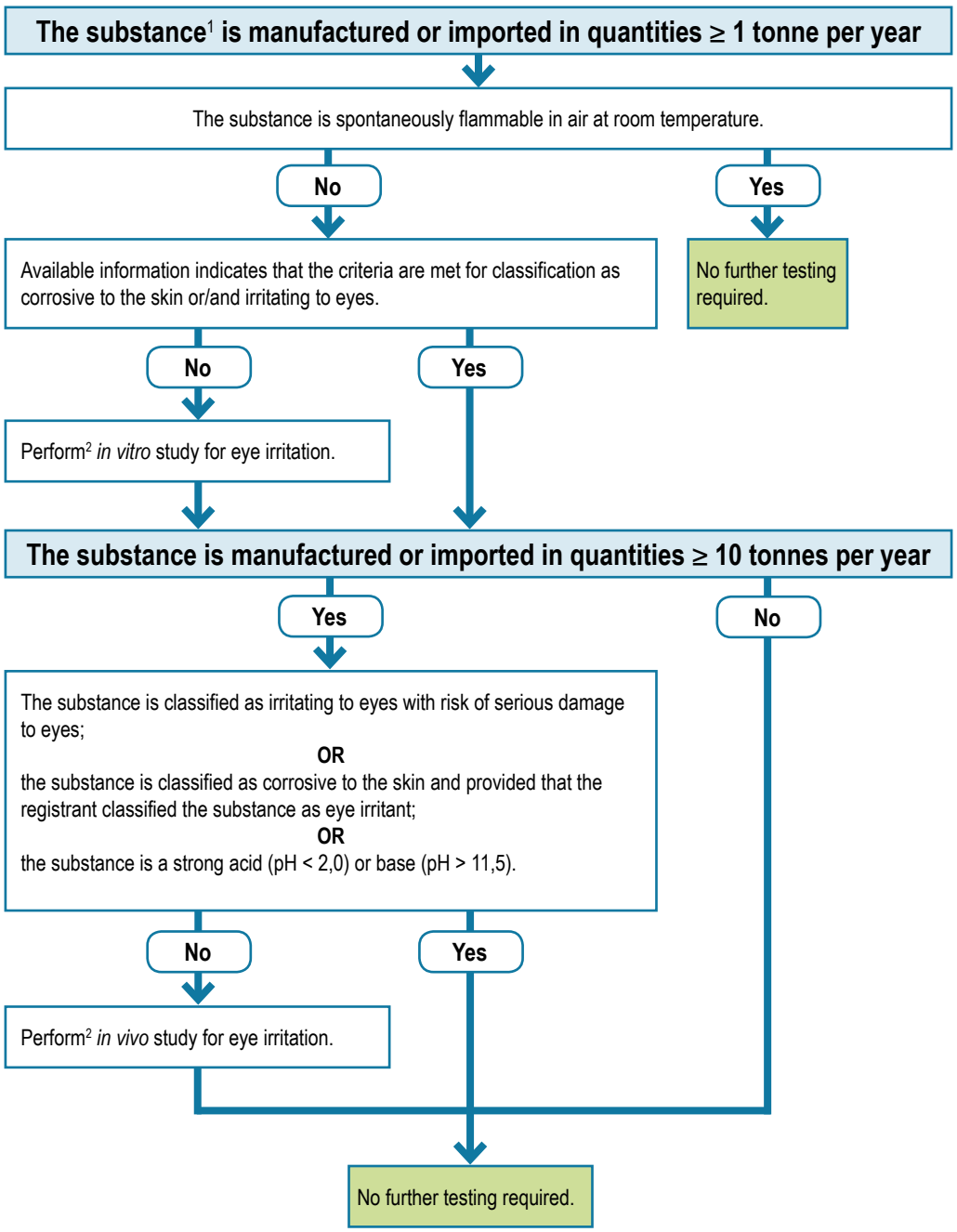
Integrated testing strategy for eye irritation under REACH



1. The substances concerned are:
a. non-phase-in substances manufactured or imported in quantities of 1 to 10 tonnes;
b. phase-in substances manufactured or imported in quantities of 1 to 10 tonnes and meeting the criteria in annexe III;
c. phase-in substances not meeting the criteria in annexe III and manufactured or imported in quantities of 10 tonnes or more.
2. A registrant may adapt the standard testing regime in accordance with the general rules set out in annexe XI section 1 (Testing does not appear scientifically necessary) and 2 (Testing is technically not possible). Under dossier evaluation the Agency may assess these adaptations to the standard testing regime.