

Commission Directive 2007/28/EC of 25 May 2007 amending certain Annexes to Council Directives 86/363/EEC and 90/642/EEC as regards maximum residue levels for azoxystrobin, chlorfenapyr, folpet, iprodione, lambda-cyhalothrin, maleic hydrazide, metalaxyl-M and trifloxystrobin (Text with EEA relevance)

*Article 3*

1 Member States shall adopt and publish, by 26 November 2007 at the latest, the laws, regulations and administrative provisions necessary to comply with this Directive. They shall forthwith communicate to the Commission the text of those provisions and a correlation table between those provisions and this Directive.

They shall apply those provisions from 27 November 2007.

When Member States adopt those provisions they shall contain a reference to this Directive or be accompanied by such a reference on the occasion of their official publication. Member States shall determine how such reference is to be made.

2 Member States shall communicate to the Commission the text of the main provisions of national law which they adopt in the field covered by this Directive.