	Checklist for registration of a pesticide in Ukraine				
No	Document	Prepared by	Language		
1	Application with short data on pesticide	Made according to standard form using data provided by the producer	Ukr		
2	Biological efficacy trials	Must be made in Ukraine	Ukr		
3	Acute tox studies on product formulation (acute oral, acute dermal, inhalation, skin and eye irritation, sensitization)	May be made in another country according to GLP and OECD	Ukr, En, RU		
4	Hygienic regulation of pesticides	Must be made in Ukraine	Ukr, En, RU		
5	Method of residue determination in water, soil, air and harvest	Must be developed if nothing is available in the country. If previously developed and published, open sources may be used	Ukr or RU		
6	Methods of qualitative determination	SIPAC or producer's methods used	Ukr, En, RU		
7	Tox studies on active ingredient vary for each AI. The standard list is: subchronic (for all), mutagenicity AMES, in vitro, in vivo (for all), carcinogenicity, teratogenicity, embryotoxic, reproduction tox - for some, upon request.	May be made in another country according to GLP and OECD	Ukr, En, RU		
8	Studies for product's impact on soil worms and microorganisms, aquatic organisms (fish, algae, daphnia), bees, birds (for some, upon request)	May be made in another country according to GLP and OECD, but Ukrainian always qualify and raise no questions and comments	Ukr, En, RU		
9	Certificates of analysis and certificates of composition of the product.	Must be submitted by the producer	Ukr, En, RU		
10	Certificates of analysis and certificates of composition for AI (with impurities in 5 batches) and for Russia full 5 batch report	Must be submitted by the producer	Ukr, En, RU		
11	MSDS and certificate of origin	Must be submitted by the producer	Ukr, En, RU		
12	Letters from manufacturers of the product and AI	Must be submitted by the producer	Ukr, RU		
13	Registration certificate for product and AI from the country of manufacturing (e.g. ICAMA reg.cert. for Chinese producers)	Must be submitted by the producer	Ukr, En, RU		
14	Manufacturing license from factories	Must be submitted by the producer	Ukr, En, RU		
15	Full application dossier (DRr)	Made at the final stage of the registration process using data submitted originally plus all studies and expert conclusions obtained in Ukraine	Ukr		
16	Details use instruction	Made at the final stage of the registration process using data submitted originally plus all studies and expert conclusions obtained in Ukraine	Ukr		
17	Label draft	Made at the final stage of the registration process using data submitted originally plus all studies and expert conclusions obtained in Ukraine. Must be approved by authorities	Ukr		

18	Sanitary and epidemiological conclusion	Issued by the Ministry of Health and Ministry of Ecology after joined	Ukr
		evaluation of DRr	