FIELD ID NO: __________

IR-4 FIELD DATA BOOK

PROTOCOL & PROTOCOL CHANGES
The protocol shall be inserted into this IR-4 field data book after this protocol cover page. Sequentially insert all relevant protocol amendments and deviations that have been received from the Study Director. Protocol changes are sent only to those field trials to which they pertain, thus the changes that are received during the course of this trial may not comprise a complete set.

PAGES IN THIS SECTION DO NOT NEED TO BE NUMBERED.

PAGES IN THIS SECTION DO NOT NEED LINING OUT IF NO ENTRIES ARE MADE

Trial Year 2007
CHEMICAL/CROP/FIELD ID NO: ________________________________

IR-4 FIELD DATA BOOK

DEVIATION FORM (PHOTOCOPY THIS PART IF NECESSARY)

INSTRUCTIONS: Every effort should be made to follow the protocol, and standard operating procedures. If an unforeseen and/or an unavoidable circumstance results in a change, the Study Director must be notified as soon as practical (via phone call, e-mail or FAX). Also notify the Regional Field Coordinator (via phone call, fax, or cc on an e-mail message). If possible, contact the Study Director prior to taking actions that differ from the protocol. The Study Director will provide instructions and/or appropriate protocol change authorization. Otherwise, document the deviation with completion of this or similar form for each individual deviation. If the deviation is faxed or e-mailed to the Study Director, then the original should be mailed to the Study Director. A true copy should be retained in the Field Data Book in Part 1B. The return copy (signed by the Study Director) should be placed in the Protocol/Protocol Changes section of the Field Data Book.

THE DATE THAT THE DEVIATION OCCURRED ________________________________

THE DATE THAT THE DEVIATION WAS RECOGNIZED ________________________________

THE DATE THAT THE STUDY DIRECTOR WAS NOTIFIED ________________________________

METHOD OF NOTIFICATION (e.g. telephone, e-mail, fax) ________________________________

THE DEVIATION IS FROM (check appropriate) ________________________________

PROTOCOL_______ SOP'S_______

SECTION OF THE PROTOCOL OR SOP'S THAT IS AFFECTED ________________________________

BRIEF DESCRIPTION OF DEVIATION: ________________________________________________

______________________________________________________________________________

______________________________________________________________________________

______________________________________________________________________________

EXPLAIN WHY THE DEVIATION OCCURRED: ________________________________________________

______________________________________________________________________________

______________________________________________________________________________

______________________________________________________________________________

ABOVE DATA ENTERED BY: ________________________________ DATE: _____________

FIELD PERSONNEL: DO NOT WRITE BELOW THIS LINE

STUDY DIRECTOR'S ASSESSMENT OF IMPACT OF DEVIATION ON STUDY: ________________________________

______________________________________________________________________________

______________________________________________________________________________

______________________________________________________________________________

APPROVED BY:

__________________________________  _______________________________________

Study Director/Date     Sponsor/Date

PROTOCOL CHANGE NUMBER_____________

cc: QA Field Research Director: Laboratory Research Director:

Regional Field Coordinator: Trial Year 2007

This protocol change form when copied on colored paper is an exact copy of the original.