

Multiparameter Display System (YSI Environmental, YSI Incorporated, Yellow Springs, USA). Salinity and Total Dissolved solids (TDS) were measured using field hand held equipment YSI EC 300 (YSI Environmental, YSI Incorporated, Yellow Springs, USA).

### **7.3.2 Larval sampling and larval identification**

Larval sampling was done once weekly to generate stage-specific estimates of *Anopheles* and culicine larval densities, invertebrate abundance and diversity. Samples were taken using standard dipping techniques and a plastic dipper (Mosquito Control Service and Supplies, USA) with a wooden ladle was used. Twenty dips were taken from each sub-plot.

The mosquito larvae collected were sorted out according to the sub-families as either anopheline or culicine. The anopheline larvae were grouped according to the instar stage and identified morphologically (Gillies and Coetzee, 1987). The pupae were maintained in the insectary in cages to develop into adults. The emergent adult mosquitoes were identified by morphological criteria.

### **7.3.3 Data Management**

Statistical analyses were done using SPSS software (Version 11.5 for windows, SPSS Inc., Chicago, IL). Simple descriptive statistics was used to tabulate the larval densities and abundance from the experimental plots. Pearson correlation was used to determine the association between mosquito larvae and the different physicochemical variables measured in the plots during the larval sampling. The relative abundance of *Anopheles gambiae* was calculated as the number of mosquito larvae divided by the number of dips taken from each larval habitat. The dependent variable (relative abundance of a species) was transformed using the log transformation method  $\{\text{Log}(x+1)\}$ .

#### 7.4 Results

A total of 21,325 *Anopheles* larvae were collected of which 91.93% (n = 19,604) were early instars and 8.07% (n = 1,721) were late instars. For culicines larvae, a total of 19,926 were collected of which 87.33% (n = 17,401) were early instars and 12.67% (n = 2,525) were late stage instars. A total of 513 pupae were collected. Table 7.1 shows the larval composition for each subfamily of mosquitoes based on each rice growth cycle. From table 7.1, most of the larvae were collected between land preparation and tillering stage. Between flowering and rice maturation, there is a decline of mosquito larvae from the plots. When the rice is harvested, there is an increase in larval abundance in the rice plots.

Figure 4a and 4b shows the weekly anopheline larval fluctuation from February 2004 to March 2005. This figure shows that there is sigmoid increase of anopheline larval densities between transplanting and tillering. When the rice gets into the reproductive stage (Flowering stage), the larval densities decline and this trend continues until the rice is harvested. After harvesting (ratoon), there is a small increase in larval densities, which follows; the larval increase goes on throughout land preparation. Figure 4b further shows that *An. gambiae* larvae are predominant throughout the rice growing cycle while *An. pharoensis* is only found during the tillering stage.

Table 7.2a shows the morphological identification of late stage instars of both *Anopheles* and culicine mosquitoes. Morphological identification of *Anopheles* larvae yielded 84.09% (n = 1,274) *An. gambiae s.l.*, 13.47% (n = 204) *An. pharoensis*, 1.32% (n = 20) *An. rivulorum*, 0.79% *An. funestus*, 0.26% *An. coustani* and 0.07% (n = 1) *An. maculipalpis*. A total of 891 late stage larvae were identified of which 65.66% (n = 585) were *C. quinquefasciatus*, 9.88% (n = 88) *C. annulioris*, 7.30% *C. poicilipes*, 7.18% (n = 64) *C. tigripes*, 0.56% (n = 5) *C. duttoni*, 5.27% (n = 47) *Ae. Aegypti*, 3.48% (n = 31), *Ae. Cumminsii*, and 0.67% (n = 6) *Ae. Vittatus*.

Table 7.2b shows the morphological identification of emergent mosquitoes from the pupae. Most of the emergent mosquitoes were identified as *An. gambiae s.l.* (57.97%, n = 80), *C. quinquefasciatus* accounted for 23.91% (n = 33).

The physicochemical variables and mosquito larval developmental stages were analyzed using Pearson correlation to test for associations of the variables (Table 7.3). Of the 171 correlation coefficients, 71 (41.52%) were statistically significant indicating that there was non-random association between some of the variables tested. For example, rice height was negatively associated with both early and late stage *Anopheles* larvae. Early stage *Anopheles* was strongly associated with late stage *Anopheles*. There was a strong positive association between anopheline larvae and culicine larvae in both the early and late instar stages. Early instars of culicine mosquitoes were associated with turbidity of water. Rice stage was strongly associated with rice height. Turbidity was negatively associated with rice stage and rice height. Two associations could not be computed because at least one of the variables is constant.

**Table 7.1 The larval abundance from the experimental plots in the 3 cycles of rice growing**

Rice planting cycle	Rice Stage	Early Instars <i>Anopheles</i>	Late Instars <i>Anopheles</i>	Early Instars Culicines	Late Instars Culicines	Pupae
Cycle 1	Transplanting	5	2	1	0	0
	Tillering	46	38	15	29	12
	Flowering	86	38	81	129	15
	Maturation	21	6	42	21	14
	Harvesting/Ratoon	290	35	3,770	718	31
Cycle 2	Land preparation	726	85	2,476	650	47
	Transplanting	50	6	16	2	3
	Tillering	3,405	678	263	85	199
	Flowering	1,133	132	118	13	13
	Maturation	187	35	468	75	9
	Harvesting/Ratoon	761	0	1,913	159	0
Cycle 3	Land preparation	1,828	14	1,845	164	25
	Transplanting	7,026	135	1,030	12	8
	Tillering	299	49	210	30	3
Total	Land preparation	2,554	99	4,321	814	72
	Transplanting	7,081	143	1,047	14	11
	Tillering	3,750	765	488	144	214
	Flowering	1,219	170	199	142	28
	Maturation	208	41	510	96	23
	Harvesting/Ratoon	1,051	35	5,683	877	31
<b>Grand Total</b>		<b>15,863</b>	<b>1,253</b>	<b>12,248</b>	<b>2,087</b>	<b>379</b>



**Table 7.2a Shows the morphological identification of late stage instars of *Anopheles* and culicines mosquito larvae**

Genera	Species	Sum	Percentage
<i>Anopheles</i>	<i>An. gambiae s.l.</i>	1,274	84.09
	<i>An. funestus</i>	12	0.79
	<i>An. pharoensis</i>	204	13.47
	<i>An. rivulorum</i>	20	1.32
	<i>An. maculipalpis</i>	1	0.07
	<i>A. coustani</i>	4	0.26
	<b>Total</b>	<b>1,515</b>	<b>100.00</b>
Culicines	<i>C. quinquefasciatus</i>	585	65.66
	<i>C. Poicilipes</i>	65	7.30
	<i>C. annulioris</i>	88	9.88
	<i>C. tigripes</i>	64	7.18
	<i>C. duttoni</i>	5	0.56
Aedes	<i>Ae. aegypti</i>	47	5.27
	<i>Ae. cumminsi</i>	31	3.48
	<i>Ae. vittatus</i>	6	0.67
	<b>Total</b>	<b>891</b>	<b>100</b>

**Table 7.2b The identification of emergent mosquitoes from pupae.**

Genera	Mosquito species	Female	Male	Total
<i>Anopheles</i>	<i>An. gambiae</i>	76	4	80
	<i>An. coustani</i>	0	4	4
	<i>An. rufipes</i>	2	0	2
<i>Culex</i>	<i>C. quinquefasciatus</i>	28	5	33
	<i>C. poicilipes</i>	1	0	1
<i>Aedes</i>	<i>Ae. taylori</i>	4	0	4
	<i>Ae. circumluteolus</i>	4	10	14
Total	Total	115	23	138

Table 7.3 Pearson Correlations between mosquito larvae and other physicochemical variable measured at the experimental plots.

	Rice height	Rice stage	#fillers	Depth (cm)	Turb	Other inverts	Anophel es	Culicine	#pupae	Temp (°C)	Cond (mS/cm)	Salinity (ppt)	oxygen (%)	DO Conc (mg/L)	DO Charge	pH
Rice stage	0.902**															
#fillers	0.243	0.281														
Depth (cm)	0.188*	0.009	-0.468*													
Turb	-0.445**	-0.352**	0.106	0.050												
Other inverts	-0.146	-0.113	(a)	-0.140*	-0.058											
Anophel es	-0.214*	-0.013	-0.111	0.033	0.183**	0.060										
Culicine	0.036	0.097	0.016	-0.002	0.125*	0.041	0.594**									
# pupae	-0.094	0.027	(a)	0.077	0.042	0.028	0.545**	0.259**								
Temp (°C)	0.071	0.129	-0.189	-0.175**	0.077	0.105	0.198**	0.164**	0.028							
Conduc (mS/cm)	-0.305**	-0.106	0.076	-0.263**	-0.098	0.133*	0.217**	0.081	0.161**	0.275**						
Salinity (ppt)	-0.303**	-0.154*	0.094	-0.242**	-0.090	0.104	0.176**	0.045	0.146*	0.177**	0.962**					
D. oxygen (%)	-0.380**	-0.257**	-0.224	-0.145*	0.060	0.033	0.147*	-0.033	0.100	0.148*	0.115	0.113				
Do Conc (mg/L)	-0.119	-0.146*	-0.208	0.065	0.110	-0.063	-0.083	-0.103	-0.035	-0.198**	-0.593**	-0.530**	0.673**			
DO Charge	-0.673**	-0.472**	-0.108	-0.175**	0.149*	0.003	0.161**	-0.051	0.125*	0.076	0.236**	0.251**	0.910**	0.540**		
pH	-0.369**	-0.328**	-0.204	0.001	0.234**	-0.027	0.114	-0.023	0.109	0.168**	-0.034	-0.023	0.592**	0.464**	0.637**	
PHMV_MV	0.365**	0.312**	0.206	0.012	-0.228**	0.022	-0.121	0.020	-0.110	-0.196**	0.014	0.008	-0.597**	-0.448**	-0.638**	-0.999**

\*\* Correlation is significant at the 0.01 level (2-tailed).

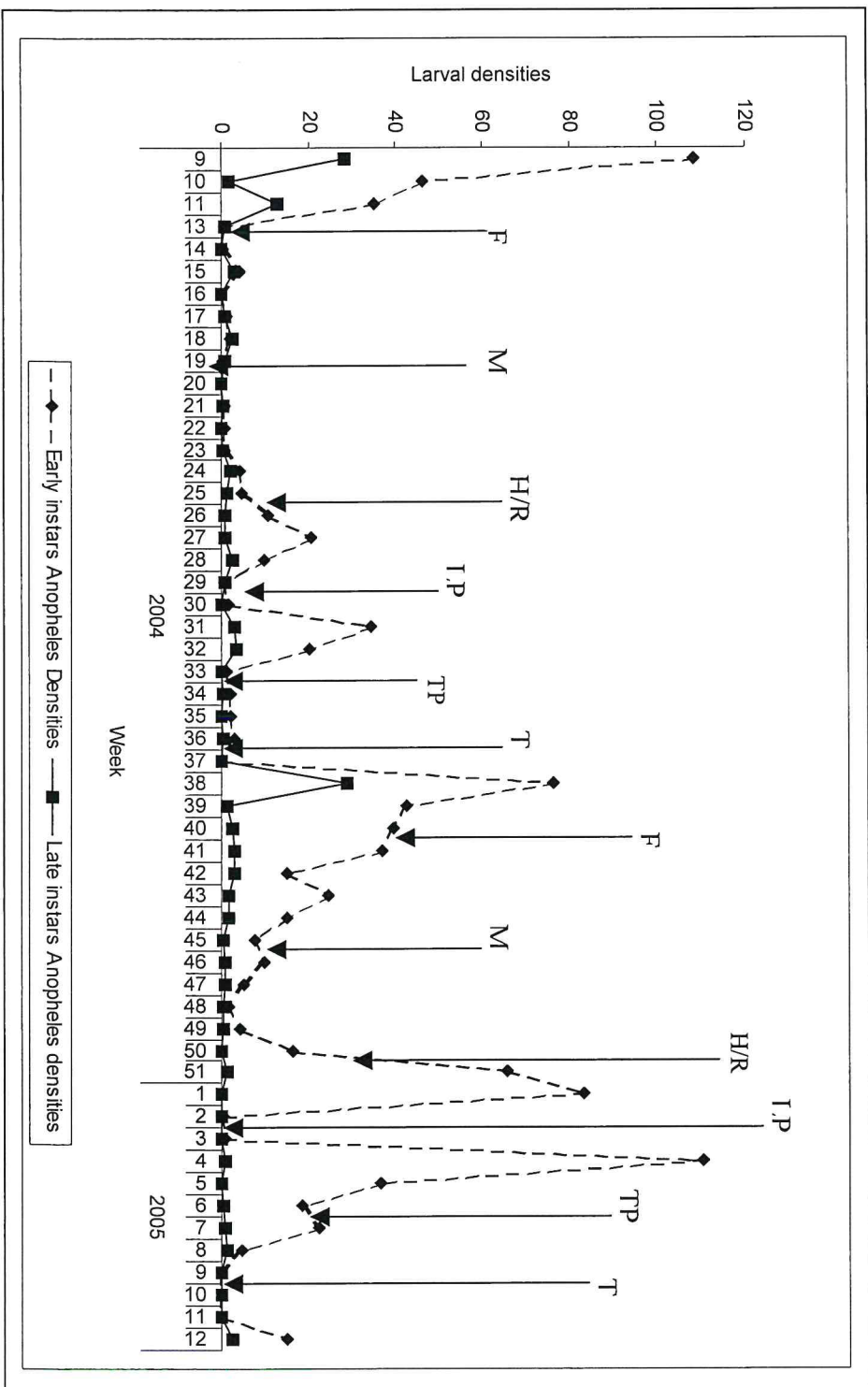
\* Correlation is significant at the 0.05 level (2-tailed).

<sup>a</sup> Cannot be computed because at least one of the variables is constant.

Do Conc<sup>b</sup> Dissolved oxygen concentration

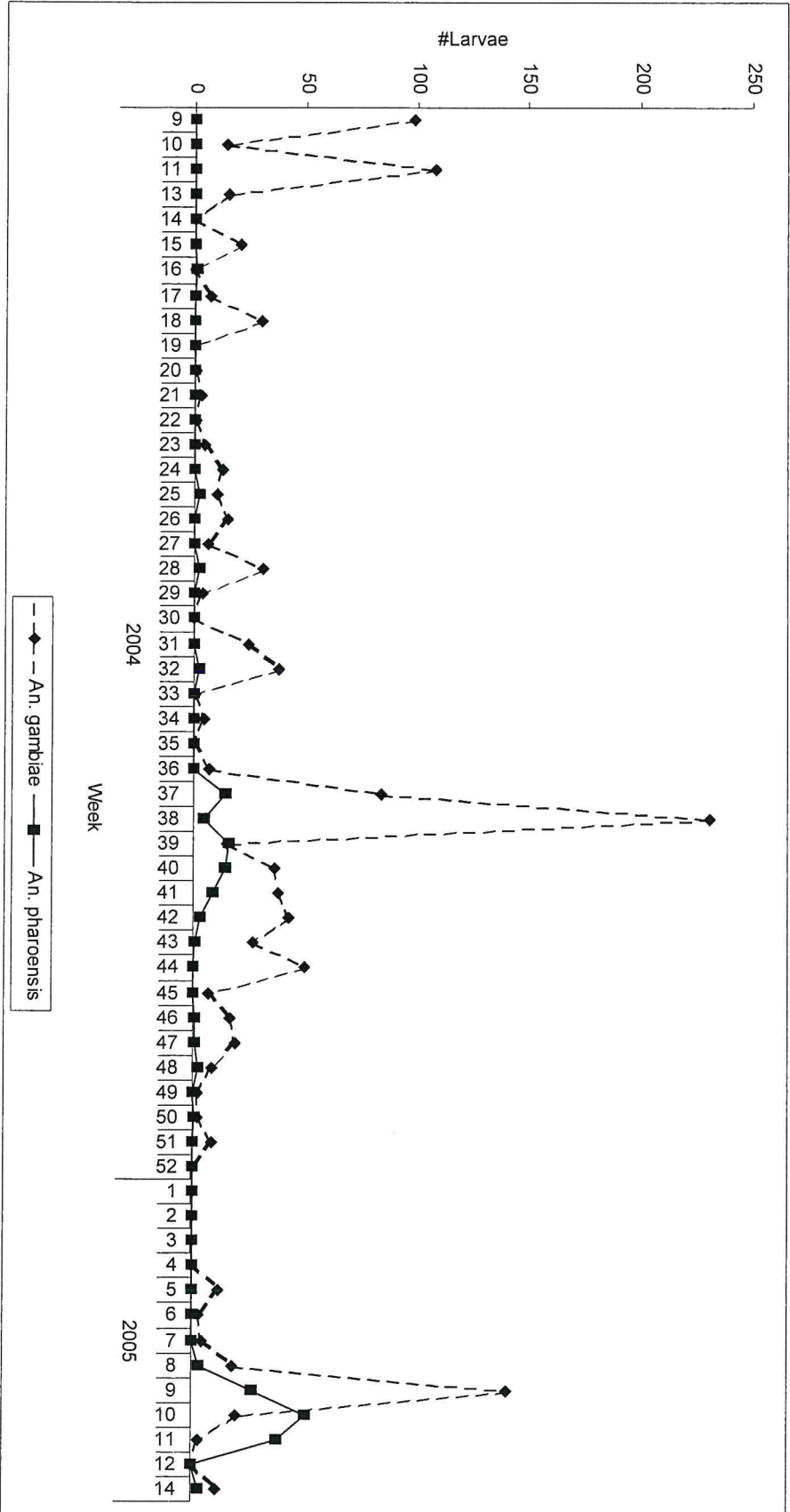
DO Charge<sup>c</sup> Dissolved oxygen charge

Figure 4a shows the weekly variation in early and late instars of *Anopheles* mosquito larvae



Foot notes: F: Flowering; M: Maturation; H/R: Harvesting/Ratoon; LP: Land Preparation; TP: Transplanting; T: Tillering

Figure 4b shows the weekly variation *An. gambiae* and *An. pharoensis* identified





### 7.5 Discussion

This study describes, the rice growing cycle as used by farmers in Mwea Irrigation Scheme and how it influences the larval diversity and abundance. This study shows that most of larvae were collected between land preparation and tillering stage. At these stages of rice development the water is more exposed providing more suitable habitats for oviposition consequently abundance of larvae in the water. Our finding is similar to various studies in schemes (Mukiama and Mwangi 1989, Ijumba et al., 1990, Mutero et al., 2000), which unanimously reveal that *An. arabiensis* normally have one major population peak annually, coinciding with the transplanting of rice seedlings during the main growing season. At transplanting of rice seedling the rice height is small providing good penetration of sunlight to aerate the habitats. Also few days after transplanting there is addition of nitrogenous fertilizers, which were shown to attract *Anopheles* and culicine mosquito to oviposition (Victor and Reuben 2000, Mutero et al., 2004b). For effective control of the mosquito menace in Mwea Irrigation Scheme, consolidated efforts should be leveled against the pre-adult stage between the transplanting and the tillering stage. At these stages the abundance of mosquito larvae is highest, consequently larvicidal pesticides application would yield maximum reduction in adult population. At this stage, the vector control programmes should use integrated vector control mechanisms (Lacey and Lacey 1990) targeting both the adult mosquito populations and larval control. Considering Mwea Irrigation Scheme, the rice growing is in phases; the control should be done all year round based on the rice-growing cycle.

Our finding further shows that there was a strong correlation between anopheline and culicine larval abundance at the experimental plots. This indicates that anopheline and

culicine mosquito larvae co-exist in the breeding habitats. These mosquito larvae utilize same habitat for their development, although from these finding, it appears after flowering culicine larval abundance increase significantly. Anopheline larvae are known to breed in open, sunlit habitats (Gillies and Coetzee 1987), which is provided by early stages of rice cycle, while culicine can breed in both habitats with clear water and foul looking like water. After transplanting, and the addition of fertilizers, the water turbidity is quite low (Mutero et al., 2004b), but as rice grows turbidity changes. After harvesting the water is very turbid due to the rice straws decomposition in the plots. This provides favorable condition for culicine breeding. Based on this finding, larval control using larvicidal agents should use one that targets both the anopheline and culicine larvae.

Morphological identification of the mosquito larvae yielded significantly more *An. gambiae s.l.* larvae than the other species. Further, the identification of emergent mosquito also gave more *An. gambiae s.l.* This indicates that the rice paddy produce more *An. gambiae* mosquitoes than the other species. This means that the rice farmers are exposed more to the malaria risk and there is a possibility of continuous parasite transmission. The implication of this is that, the people living within the scheme should use insecticide treated bednets (ITBN) to minimize the human-vector contact. This would lower the chances of getting the malaria parasites consequently remaining healthy.

In conclusion, these studies show that mosquito control programmes should be emphasized at the beginning of rice growing cycle targeting the developmental stages.

Throughout the rice growing cycle, larvicidal control programmes should go hand in hand with other vector control tools such as insecticide treated bednets.

## CHAPTER 8: THE SURVIVAL OF IMMATURE *ANOPHELES* IN MWEA RICE IRRIGATION SCHEME

### 8.1 Abstract

### 8.2 Introduction

*Anopheles gambiae* complex and *An. funestus* complex are the primary vectors of malaria in Mwea rice irrigation scheme. The distribution and abundance of mosquito larvae results from availability of oviposition sites, the oviposition preferences of females, and the ability of the immatures to tolerate and develop after the eggs were laid.

Life tables provide a structured framework for identifying developmental stages most susceptible to mortality and, under some conditions, for inferring sources of mortality (Service 1993). The life tables for the developing immatures can be constructed using either horizontal or vertical methods (Reisen and Siddiqui 1979, Reisen et al. 1982, Service 1993). Horizontal life table methods are appropriate for distinct cohorts that can be followed through time, whereas vertical life table methods are appropriate for populations with overlapping generations and age distributions that remain stationary for the duration of the sampling period. Service (1971, 1973, 1977, 1993) Reisen and Siddiqui (1979) and Reisen et al. (1982) provide extensive discussions about how such information can be analyzed.

In Kenya, Service, (1971, 1973, 1977) and Aniedu and colleagues (1993) studied the survival of immature *An. gambiae* complex in the larval habitats. The objective of this study is to determine survival of immature *An. arabiensis* in different habitats in Mwea Irrigation Scheme and under the experimental rice growing conditions.

### **8.3 Materials and Methods**

#### **8.3.1 Horizontal and vertical Life Tables**

This will be done as described in 3.10 in this thesis.



## CHAPTER 9: CONCLUSIONS

1. Rice growing is an important source of mosquito production. Rice paddy and canal are most productive habitats. This study also shows that unplanned rice growing supports more *Anopheles* larvae than planned rice growing system. From the results larval control activities should be initiated at early rice stages and during the rainy season. The unplanned rice system should be better managed and if possible controlled by the management system of the Mwea Irrigation Scheme where water for irrigation is distributed together with those of the scheme.
2. *Anopheles* larval abundance is found to be associated with pH and salinity. These parameters in a rice agro ecosystem are associated with early stages of rice growth cycle, which is associated with larval abundance. For effective control of developmental stages of mosquito larvae, the application of the larvicide should be done after transplantation and the larvicide should persist until the reproductive stage of the rice.
3. Mosquito control programmes should be emphasized at the beginning of rice growing cycle targeting the developmental stages. Throughout the rice growing cycle, larvicidal control programmes should go hand in hand with other vector control tools such as insecticide treated bednets.
4. There is natural regulation for mosquito production in the aquatic habitats. Although the number of emergent mosquitoes decreases due to predation pressure, there is a need to effectively control mosquito developmental stages using larvicides such as Bti and Bs.

5. Paddy is the most productive habitat productive habitat type. Most *An. gambiae* emerged from the paddy habitat type.
  
6. Addition of nitrogenous fertilizer results to a consequent increase of *Anopheles* mosquito larvae. This is the phenomenon which results in the increase of mosquito larvae between transplanting and tillering.
  
7. There is no succession of *Anopheles* mosquito larvae in the rice paddy but there is more diversity of larvae at the early stages of the rice growing. Most of the *Anopheles* mosquitoes colonize the paddy at the early stages of paddy development. There is more species at the transplanting stage and tillering stage, after which, the other species diminish leaving only *An. gambiae* in the paddy till harvesting.

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