CHAPTER ONE

1. INTRODUCTION

1.1 Background

The invasion of ecosystem by non-native species is a global problem with serious consequences for ecological, economic and social systems (Pimental et al. 2000). The study of alien floras or non-native species along with native floras in many places of the world is essential for understanding plant invasions (Pickard 1984, Kloot 1987). Any species including its seeds, eggs, spores or other living entities through which it is capable of multiplying and propagating in a foreign ecosystem is said to be an alien species. Invasive plants produces reproductive offspring, often in very large numbers, at considerable distances from parent plants and thus has potential to spread over a considerable area. Introduction of plants from one place to another may be natural (accidental) or planned (intentional). Although an introduction may meet a desired objective in one area, at one time, or for some sectors, unwanted and unplanned effects may also occur. Many introduced plants have been naturalized in a new environment and form a part of existing landscapes and ecosystems. Such introduced plants may outcompete the native species and turns to invasive.

There are about 8000 species of plants that are believed to be agricultural weeds in the world and about 2500 species are considered to be potentially dangerous (Yaduraju et al. 2000). There is an enormous change in ecology, environment, biodiversity and even changes in atmospheric composition and climate due to biological alien invasions.

Introduction of exotic species following habitat change and soil degradation has equal contribution to the threats of Nepal's biodiversity as other factors such as the human activities, habitat destruction and over-exploitation of natural resources (Chaudhary 1998). The wide range of habitat and environmental conditions makes Nepal vulnerable to the establishment of invasive species of foreign origin (Tiwari et al. 2005). Potential invasive alien species from most areas of the world find suitable habitat somewhere in Nepal and accidental or intentional introduction by gardeners, traders and foresters have

contributed to the large number of exotic plant species in Nepal (Kunwar 2003). It has been estimated that there are >200 introduced plant species in Nepal (Tiwari et al. 2005). Among the alien invasive plant species, *Parthenium hysterophorus* L. has emerged as a major threat to grassland ecosystems in the urban areas of Nepal. *P. hysterophorus* (commonly known as Parthenium weed, hereafter referred to as *Parthenium*) invaded Southeast Asian countries such as India, Pakistan and Sri Lanka through cereal and grass seed shipments from USA during 1950s (Kathiresan et al. 2005). It is considered that after a few years of invasion of *Parthenium* in India this plant invaded southern Nepal and progressively to other parts of Nepal through road transportation. In Nepal, a research survey carried out by Tiwari et al. (2005) showed that *Parthenium* is found through out tropical region from east to west of Nepal and finds its way towards subtropical regions.

Although it is acknowledged that invasions by exotic plant species represent a major threat to biodiversity and ecosystem stability, relatively little attention has been paid to potential impacts of these invasions on nutrient cycling processes in the soil. Introductions of alien plant species have potential to change many components of the carbon (C), nitrogen (N), water and other cycles of an ecosystem (Ehrenfeld 2003). Studies have shown that invasive plant species frequently increase biomass and net primary productivity, increase N availability, alter N fixation rates, and produce litter with higher decomposition rates than co-occurring natives (Ehrenfeld 2003).

The right kind of soil and climatic conditions in the place where weeds are introduced will only aid in its faster establishment and subsequent spread. This might lead to a situation where the exotic weed would dominate local weed flora (Yaduraju et al. 2000). The presence of weed population in an arable field is the result of ecological reaction of previous management practices, soil characteristics and the regional climate (Tamado and Milberg 2000). Successful invasion by the plant species depends on presence of safe sites for their establishment, and availability of resources for growth and reproduction in these sites (Yatsu et al. 2003). Competition may occur between invasive and native species. The spread of invasive plants may result in degradation or local extinction of populations

of native species and thus the alien invasion has been considered as one of the major threats to native biological diversity (Yatsu et al. 2003, Shabir and Bajwa 2006).

1.2 Justification of the study

Parthenium is an annual weed with wide amplitude of ecological adaptability in terms of photoperiodicity as well as thermoperiodicity (Oudhia 2001). With good rainfall and warm temperature *Parthenium* can germinate, grow and flower at any time of the year (Dhileepan et al. 2000, Maharjan 2006). It is considered as a noxious weed because of its prolific seed production and fast spreading ability, allelopathic effects on other plants, strong competitiveness with crops and health hazard to human as well as animals (Raghubanshi et al. 2005). Although herbarium specimen from Nepal was first collected in 1967 (Tiwari et al. 2005), Parthenium became more common after 1990s. Now it has emerged as one of the major threats to biodiversity of roadside and urban grasslands. Except an inventory of IUCN Nepal (Tiwari et al. 2005) and a few works in Kathmandu valley (e.g. Joshi 2005, Maharjan 2006) there is no comprehensive information on the distribution and ecological impact of Parthenium invasion to natural ecosystems of Nepal. Since most of the urban areas of Nepal in tropical to subtropical regions has already been invaded by Parthenium it is high time to assess the impact of its invasion to soil properties and biodiversity. Present work, though in a small scale, is the first step to this direction. It mainly focused on the changes in soil physico-chemical properties and plant community structure (eg. plant species richness, density, importance percentage, etc) of three grasslands in tropical to sub tropical urban areas of central Nepal. If we can generate such data for the whole range of Parthenium invasion, mitigation and control measure of its invasion would be more easier and successful.

1.3 Objectives

An attempt has been made to evaluate the impact of *Parthenium* invasion on soil physicochemical properties and species composition of grasslands in three valleys of central Nepal. The specific objectives of the research are given below:

- i. To analyze the community structure of *Parthenium* invaded grasslands with frequent grazing by domestic animals.
- ii. To examine the extent of change in physico-chemical properties of soil due to *Parthenium* invasion.
- iii. To understand the change in species composition due to *Parthenium* invasion.
- iv. To know the population response of native species to *Parthenium* invasion.

1.4 Limitation of the study

Although *Parthenium* has invaded a wide range of land uses, grasslands of three urban valleys of central Nepal were examined in the present study due to limited time and resources.

CHAPTER TWO

2. LITERATURE REVIEW

Biological invasion by alien invasive species is now recognized as one of the major threats to native species and ecosystems (Kathiresan 2004, Kathiresan et al. 2005). *Parthenium* is an alien invasive species of tropical to subtropical world which has been designated as one of the most troublesome species. The adverse effects of *Parthenium* on human health, livestock, agriculture productivity and biodiversity have been well documented. There has been only a few references on the impact of *Pathenium* invasion on physico-chemical properties of soil.

Parthenium is globally distributed in pantropical regions of north and south America, Australia, Ethiopia, India, Israel, Nepal, Taiwan, West Indies (Picman and Picman 1984, Peng et al. 1988, Mishra 1991, McFadyen 1992, Medhin 1992, Evans 1997, Shabbir and Bajwa 2006). *Parthenium* is distributed throughout Nepal from tropical to sub-tropical regions (75-1350 m asl) (Tiwari et al. 2005). It has achieved major weed status in India and Australia within the last few decades (Navie et al. 1996, Mahadevappa 1997, Evans 1997) and in Nepal it is among the ten most important invasive plant species (Tiwari et al. 2005).

Parthenium has caused a number of environmental and agricultural problems, such as the loss of crop productivity, fodder scarcity to livestock, biodiversity depletion and health problems for human beings and livestock (Evans 1997). Because of its invasive capacity and allelopathic properties *Parthenium* has potential to disrupt natural ecosystems. *Parthenium* can form its huge monoculture strands with no other plant in the vicinity (Singh et al. 2005a). In Islamabad (Pakistan) many common medicinal plants have been threatened because of the aggressive colonization by *Parthenium* in cultivated lands, wasteland, degraded areas, rocky crevices, along water channels, roadsides, and railway tracks (Shabbir and Bajwa 2006). This noxious weed can affect crop production, animal husbandry, human health (e.g. allergic dermatitis, allergic rhinitis, bronchitis, hay fever and asthma) and biodiversity (Chippendale and Panetta 1994, Kohli and Rani 1994, McFadyen 1995, Navie et al. 1996, Evans 1997, Shabbir 2002). Other problems are

contamination of seed products and total habitat change in the invaded area (Chippendale and Panetta 1994).

Tamado and Milberg (2000) carried out an exploratory weed survey on 240 crop fields in eastern Ethiopia to assess the influence of environmental and crop improvement factors on weed species composition and distribution and to investigate the association of recently introduced Parthenium with other components of weed flora. They concluded that Parthenium was second most abundant species on cropland. Despite being rare at high altitudes and perceived as one of the most troublesome weeds within only 20 plant generations from its introduction, *Parthenium* is now recognized as serious problem of crop and rangelands in Ethiopia. In Australia, Parthenium has been regarded as one of the worst weeds because of its invasiveness, potential for spread, and economic and environmental impacts (Anonymous 2003). Parthenium has posed a major problem in rangelands and summer cropping areas, threatened biodiversity in the Einasleigh uplands bioregion and native grasslands in the central highlands of Queensland. In addition, Parthenium has affected primary production both in livestock and grain enterprises as well as caused health problems for human and animals in Queensland, Australia (Anonymous 2004, O'Donnell and Adkins 2005). Parthenium can dominate pastures under continued heavy grazing and has the potential to exclude useful forage plants, thus decreasing pasture productivity, carrying capacity and land values. Parthenium competes directly with preferred pasture species, reducing pasture vigour and seed set, leading to habitat and ecosystem changes.

Parthenium has strong allelopathic effects on germination and growth of cultivated as well as wild plant species. Negative allelopathic effect of *Parthenium* on cultivated plants (e.g. *Abutilon theophrasti, Brassica* spp (*B. campestris, B. oleracea* and *B. rapa*), *Glycine max, Lolium multiflorum, Oryza sativa, Phaseolus vulgaris, Raphanus sativus, Triticum aestivum, Vigna radiate* and *Zea mays*) has been well documented (e.g. Wondimagegne-hu and Singh 1987, Yadav and Chauhan 1998, Sinha and Deo 1999, Oudhia 2000a, Oudhia 2000b, Chetry 2003, Singh et al. 2003, Singh et al. 2005b, Rajendiran 2005, Maharjan et al. 2007). *Parthenium* also showed negative allelopathic effect to its associated wild species such as *Artemisia dubia* and *Ageratina adenophora* (Maharjan et al. 2007), *Age-*

ratum conyzoides (Singh et al. 2002). Seed germination of *Parthenium* was inhibited by leaf leachates of *Azadirachta indica* and latex of *Calotropis procera* (Goyal and Singh 2003).

Parthenium colonizes a wide range of vegetation and soil types but the most common are in the alkaline, clay loam soils (Anonymous 2006). It is an efficient colonizer and a highly competitive plant. *Parthenium* colonizes overgrazed pastures with low ground cover, cultivated lands, disturbed and bare areas such as roadsides and tracks, and heavily stocked areas such as stockyards and watering points (McFadyen 1992). This plant shows aggressive growth in most of the growing sites. Fundamental reproductive traits, such as seed mass and seed set, of *Parthenium* varies with soil type (Annapurna and Singh 2003) and this variation would enable *Parthenium* to adjust to a variety of habitat conditions. Annapurna and Singh (2003) found that seed mass of *Parthenium* declined in relatively coarser soils whereas seed production increased. In soils rich with clay, the plant produced a smaller number of larger seeds. Parthenium invaded sites at Kathmandu had sandy loam soil pH from 5.4 to 7.4, water holding capacity 16.8 to 63%, organic matter 1.134 to 4.24%, total nitrogen 0.055 to 0.206%, available phosphorus 31.86 to 69.93 kg/ha and available potassium 74.72 to 746.5 kg/ha (Joshi 2005).

Plant invasion also alters the soil properties. Bidwell et al. (2006) found that organic matter and nitrogen content of soil increased with the increase of invasive weed abundance in Australia. Differences in soil ecology are valuable indications of the effect of species richness on invasion dynamics (Collins 2005). For example, Duda et al. (2003) tested nutrient levels (NO₃, P, K, Na) among the native, ecotone and exotic derived soils of the exotic annual chenopod, *Halogeton glomeratus* at the Desert Experimental Range in western Utah, U.S.A. and found that *H. glomeratus* invasion altered soil chemistry and soil ecology, possibly creating conditions that favor exotics over native species.

O'Donnell and Adkins (2005) performed the glasshouse study on four grass species and four legume species and reported that the grass species *Bothriochloa insculpta*, *Dicanthium aristatum* and *Cenchrus ciliaris* all had strongest competitive abilities than *Parthenium* by factors of 3.16, 1.49 and 1.11 respectively. *Digitaria milanjiana* was only

competitively equivalent to 0.84 *Parthenium*. Lugumes such as *Clitorea terneata* competed strongly, with single plant being competitively equal 2.89 *Parthenium*. *Glycine latifolia*, *Macroptilium brachteatum* and *Stylosanthes seabrana* had strengths relative to *Parthenium* of 0.43, 0.38 and 0.38 respectively. Although these legumes not being superior in competitive ability to *Parthenium*, would still add nitrogen to thye soil and in pasture situation, could have some benefit through a promotive effect on desirable pasture grasses' competitive ability against *Parthenium*. They concluded that prostrate grass and legume species could be considered in any future studies designated to test competitive ability against *Parthenium*.

Joshi (2005) studied reproductive efficiency and biomass allocation of *Parthenium* growing in different habitats of Kathmandu valley. Average density ranged from 11 to 47 pl/m^2 . In individual plants, vegetative allocation (93%) was found comparatively higher than reproductive allocation (7%). Seed germination ranged from 38 to 85%.

CHAPTER THREE

3. STUDY AREA

3.1 Location and Physiography

The selected sites were grasslands on the process of increasing *Parthenium* invasion. Three grasslands, lying at river valleys, were sampled in Gorkha, Nuwakot and Kathmandu districts (Figures 1-3). In each district single grassland was sampled. In Gorkha, sampling was done within the premises of Gorakhkali Rubber Udhyog Limited at Deurali Village Development Committee area (here after referred as Gorkha site). In Nuwakot, the selected grassland lies at Majhitar of Bidur Municipality-5 (here after referred as Nuwakot site). In Kathmandu, the selected grassland lies within the premises of Tribhuvan University Central offices at Kirtipur (here after referred as Kathmandu site) which is the proposed site for construction of Natural History Museum of Tribhuvan University. The area of each grassland sites were more than 1000 square meters. Gorkha and Nuwakot sites lie in tropical climatic zone (<1000 m asl) and site in Kathmandu site lies in subtropical zone (1000-2000 m asl).

The three selected sites were open grazing grasslands without any plantation. The grasslands lie along a side of main road ways leading to different places. They were mainly disturbed by cattle grazing. Moreover, study sites are disturbed by solid wastes thrown by people living nearby and vehicles that temporarily are parked. The extent of *Parthenium* invasion was increasing in all sites. The invasion in the selected grasslands has started 5-6 years back.

3.2 Climate

Average meteorological data of last five years (2002-2006) have been presented for all three sites (Figures 1-3). At Gorkha site, the average minimum temperature was 7.1°C in January and average maximum temperature was 33.6°C in June (Figure 1). The average maximum rainfall was 546 mm in July and average minimum rainfall was 4.4 mm in December. The total annual rainfall was 2310 mm.

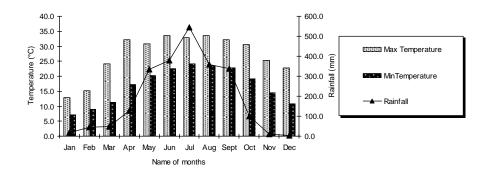


Figure 1. Average monthly minimum and maximum temperature (°C) and rainfall recorded at Khaireni weather station, Tanahun, for five years (2002-006). (Source: Department of Hydrology and Meteorology/GoN). The study site (Gorkha site) is located at a distance of about 2 km from the weather station.

At Nuwakot site, the average minimum temperature was 7.5°C in January and average maximum temperature was 31°C in June (Figure 2). The average maximum rainfall was 517 mm in July and average minimum rainfall was 5.4 mm in December. The total annual rainfall was 1812 mm.

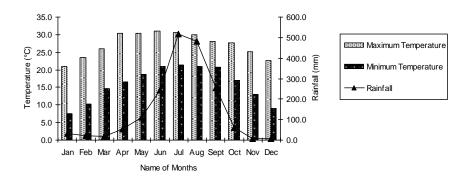


Figure 2. Average monthly minimum and maximum temperature (°C) and rainfall recorded at Kakani weather station, Nuwakot for five years (2002-2006) (Source: Department of Hydrology and Meteorology/GoN). The study site (Nuwakot site) is located at a distance of about 25 km from the weather station.

At Kathmandu site, the average minimum temperature was 3°C in January and average maximum temperature was 29.4°C in June (Figure 3). The average maximum rainfall was 437 mm in July and average minimum rainfall was 7.2 mm in December. The total annual rainfall was 1472 mm.

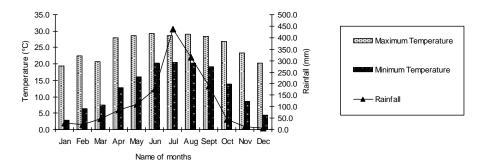


Figure 3. Average monthly minimum and maximum temperature (°C) and rainfall recorded at Tribhuvan International Airport weather station, Kathmandu for five years (2002-2006) (Source: Department of Hydrology and Meteorology/GoN). The study site (Kathmandu site) is located at a distance of about 10 km from the weather station.

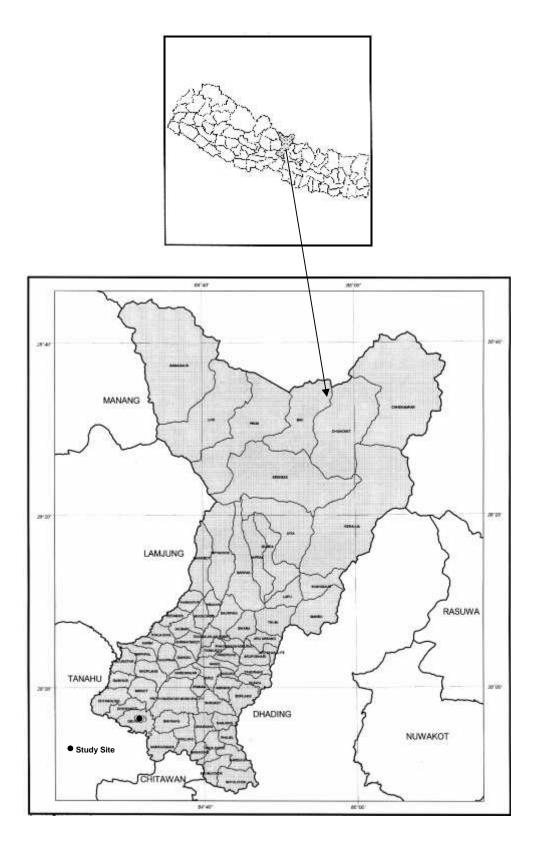


Figure 4. Map of Gorkha study site

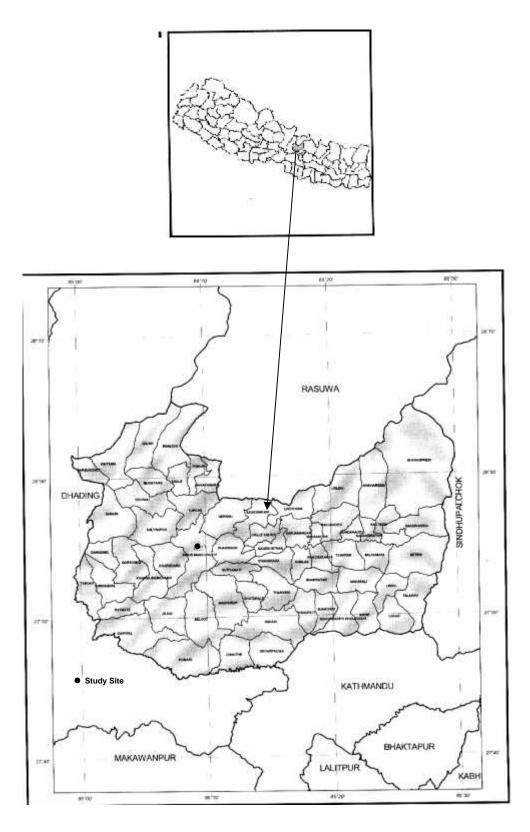
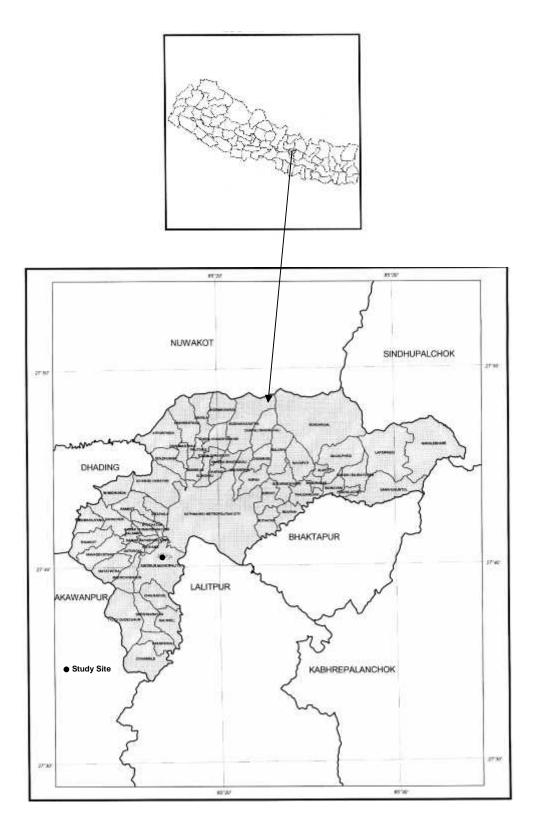


Figure 5. Map of Nuwakot study site





CHAPTER FOUR

4. MATERIALS AND METHODS

4.1 Species characters

Parthenium (family Asteraceae) is an annual herb native to north and south America and the West Indies (Picman and Picman 1984). It is variously called as Bitter weed, Carrot grass, False rage weed, Fever few, Parthenium weed, Ragweed, white top, Santana Maria, etc in different parts of the world and called Kanike ghans, Bethu ghans, Padke phul etc in Nepal (Adhikari and Tiwari 2004, Tiwari et al. 2005).

Parthenium is aggressive colonizer in poor ground cover and exposed soil such as wastelands, roadsides and overgrazed pastures (McFadyen 1992). Buried seeds can still germinate after eight to ten years. Because of all these reasons *Parthenium* weed has succeeded invasion in different parts of the world and has become widespread or cosmopolitan in nature.

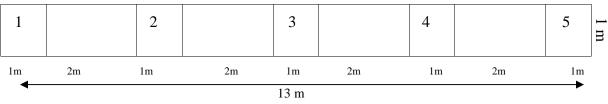
Parthenium is perennial, erect or profusely branched herb up to 2 m high. Stem is unbranched in lower part and profusely. It has dull green deeply dissected leaves covered with fine white hairs and are arranged in alternate manner. Leaf blade are generally ovate. *Parthenium* has numerous white flowers borne in compact heads near the top of the stem. Five flattened fruits are formed in each floret. Each seed is 1-2 mm long and enclosed in a fine white seed coat. Dry fruits are called achenes and a typically mature plant can produce from 15000 to 25000 achenes. General look of the pland leads to misidentification as *Artemisia* species (Haseler 1976, Joshi 1991, Tiwari et al. 2005). The total seed out put varies from 1392-3864 seeds/plant and average biomass varies from 11.39-29.7 g/plant (Joshi 2005).

4.2 Field Sampling

4.2.1 Vegetation Sampling

Field sampling was done from 23rd August to 20th September, 2006. Belt transect method was used for vegetation sampling (Martin 1995, Cunningham 2001). In each grassland,

ten belt transects (each with $13 \text{ m} \times 1 \text{ m}$) were defined in such a way that each transect included both invaded (by *Parthenium*) and non invaded parts of the grassland. In each transect, 5 quadrats (1 m × 1 m) were sampled at interval of 2 m, in such a way that two quadrats lied on non invaded part, one quadrat at transition position and two quadrats at invaded part. The quadrats in each transect were designated as 1, 2, 3, 4 and 5; the quadrats 1 and 2 were non invaded, quadrat 3 the transition position and quadrats 4 and 5 the invaded. The numbers of quadrats sampled in each grassland were 50; thus 150 quadrats were sampled in total for all sites.



* 1 and 2: non invaded, 3: transition, 4 and 5: invaded

Figure 7. Location of sampling plots in each transect

In each quadrat the total number of species (i.e. species richness) and total number of individuals of each species were recorded. Mid point values for coverage of each species were estimated visually following Zobel et al. (1987).

4.2.2 Soil sampling

Soil samples were collected from four corners and a middle of each quadrat at the depth of 15 cm. These sub samples were mixed thoroughly and about 200 g was collected in polythene bag. The soil samples were air dried in shade for a week and stored in air tight bags until laboratory analysis. There were 150 soil samples in total.

4.3 Laboratory Analysis of soil

Soil samples were analyzed at Soil Science Division of National Agricultural Research Council (NARC), Khumaltar, Lalitpur. Soil texture, pH, organic carbon (OC), total Nitrogen (N), available Phosphorus (P) and available Potassium (K) were estimated in the soil samples. The various methods that were used to analyze soil have been described detail in Pradhan (1996) and Bajwa et al. (1997).

4.3.1 Soil texture: Soil texture was determined by hydrometer method. Soil samples were treated with dispersing agent and allowed to settle freely. Soil hydrometer, was calibrated to read the percent of solid gives silt and clay in 40 seconds reading and gave clay at 3 hours. Organic matter, calcium carbonate and soluble salt, if present in appreciable amount, were removed by treating with hydrogen peroxide and hydrochloric acid before dispersing for analysis.

Soil sample (100 g) was taken in a 250 mL beaker and water was added to cover the soil. Sodium hexametaphosphate solution (20 mL) was added and stirred and left overnight. It was transferred in a dispersion cup and two third of the cup was filled with water. It was stirred in the mechanical stirrer, transferred in the hydrometer jar and made the volume up to mark with the hydrometer in it. The hydrometer was removed and the jar was shaken in upside down position for several times by closing the mouth either by the hand or cork. When the soil was well dispersed, it was kept in the table and the time was noted immediately. The hydrometer was immersed in the jar and it was read at 40 sec. and after 3 hours.

(Silt +Clay) % = Hydrometer reading at 40 sec. + $0.3 \times (t-20) \circ C$ Clay (%) = Reading at 3 hrs. + $0.3 \times (t-20) \circ C$ Sand (%) = 100- % (Clay+ Silt) Silt (%) = % (Clay + Silt) - % Clay where t is the temparature of suspension at the time taking hydrometer reading.

4.3.2 Soil pH: Soil pH was determined using Fischer's Digital pH meter in 1:2 ratio of soil-water mixture. Before measurement the pH meter was calibrated with the help of buffer solutions of known pH (pH 4 and 7). During the measurements, 50 mL of distilled water was poured into 25 g of soil sample. It was stirred at least four times over 30 minutes to allow the soil and water to reach equilibrium and settled for 25 minutes. The electrode was dipped in the mixture and reading of pH was noted.

4.3.3 Organic matter content: Organic matter content in the soil was calculated by Walkey and Black's rapid titration method (1934). Soil sample (2 g) passed through

fine sieve (0.5 mm) was taken in a 250 mL conical flask and added 10 mL of 1N $K_2Cr_2O_7$ and 20 mL of conc. H_2SO_4 with gentle swirling. The reaction being exothermic, the flask was kept on asbestos sheet for about 30 minutes to allow cooling to room temperature. To that mixture 5 mL orthophosphoric acid, 100 mL distilled water and 10 drops diphenylamine indicator solution were added successively and shaken.

0.5N ferrous ammonium sulphate was run from burette, with constant stirring until the colour changed from violet to bright green through blue. The volume of ferrous ammonium sulphate solution used for titration was noted. A blank titration (without soil) was carried out in a similar manner.

Volume of 0.5N ferrous ammonium sulphate solution used for blank titration: X

Volume of 0.5N ferrous ammonium sulphate consumed with soil: Y

Volume of 1N K₂Cr₂O₇ used for oxidation of organic carbon in soil: $\underline{X-Y}$

Organic carbon in soil (%) = $\frac{X-Y}{2} \times 0.003 \times \frac{100}{2}$

Organic matter in the soil (%) = Organic carbon in soil \times 1.724

4.3.4 Total Nitrogen: Micro-Kjeldahl method consisting of conversion of organic Nitrogen into ammonia by boiling with conc. H_2SO_4 ; the ammonia was subsequently liberated from its sulphate by distillation in presence of an alkali, which is titrated against HCl.

2

Ten gram soil was taken in a 300 mL Kjeldahl digestion flask and 30 mL conc. H_2SO_4 was added to the soil. As catalyst, 10 g of Habbard's mixture (potasium dichromate, ferrous sulphate and copper sulphate in the ration of 20:10:1), 1 g salicyclic acid and 5 g of sodium thiosulphate were added. The mixture was heated at low heat till there was no frothing and then the heat was raised until the content of the flask would grey or greenish in colour for complete digestion. The digest was cooled and about 100 mL distilled water was added. It was transferred to a 250 mL volumetric flask and made up to that volume by adding distilled water and filtered.

Two drops of methyl red indicator was added to a 150 mL conical flask containing 20 mL $0.1N H_2SO_4$ and placed under the delivery tube condenser in the distillation assembly. 10 mL of the filtrate was pipetted in the distillation flask and 10 mL of 45% NaOH solution was added in that flask through the funnel connected through a tube to a distillation flask and the filtrate was distilled. 30 mL of distillate was taken and the excess of the acid in the receiver was titrated against 0.1N NaOH.

Nitrogen (%) = (X-Y) $\times 0.0014 \times vol.$ of digested sample made $\times 100/V \times M$

Where,

 $X = Vol. of 0.1N H_2SO_4$ taken in the receiver (mL)

Y = Vol. of 0.1N NaOH required to neutralize the excess of acid (mL)

V = Vol. of the filtrate used for distillation (mL)

M = Mass of the soil taken for digestion (g)

4.3.5 Available Phosphorus (P): The available P was extracted with 0.5M NaHCO₃ (pH 8.5) as detailed by Olsen et al. (1954).

Preparation of a standard curve: To make 100 ppm P solution, 0.4387g of KH₂PO₄ was dissolved in distilled water and made volume to one litre. 5mL of 100 ppm P solution was diluted to 100 mL with distilled water to make standard solution of 5 ppm P solution. To prepare standard curve 0, 0.5, 1, 2, 3, 4 mL of 5 ppm P solution were taken in six different volumetric flasks (25 mL). 5 mL ammonium molybdate was added to the solution with continuous shaking until evolution of carbondioxide stopped. About 10 mL distilled water was added through washing the neck of the flask. Then 1 mL dil. SnCl₂ solution was added and made the volume to the mark with distilled water. The colour intensity in each of the standard solution was read in colorimeter at wave length of 660 nm and the absorbance was plotted on a simple graph paper.

Analysis of test sample: One gram of soil was transferred to a conical flask (100 mL). A pinch of Darco-G 60 and 20 mL of 0.5 N NaHCO₃ solution was added. It was

mixed and shaked on an electric shaker for half an hour and filtered. In a similar way, a blank with all reagents and no soil was prepared. More Darco-G60 was added to the flask, if filtrate after shaking was not clear. 5 mL of filtrate was taken in a volumetric flask (25 mL) and proceed further for colour development as detailed under preparation of standard curve.

Available P in soil (ppm) = $X \times 100$

Where,

X= Concentration of P read from standard curve against absorbance of test solution

100 = Total dilution

Available P (Kg/ha) = Available P expressed in ppm $\times 2.24$

4.3.6 Available Potassium (K): A known weight of soil was shaken with neutral normal ammonium acetate solution. Ammonium ions exchange with K ions adsorbed on soil colloids. The extract contained exchangeable and water soluble K was used for determination of available K in soils.

Preparation of a standard curve: To make standard 1000 ppm K solution, 1.91 g of KCl was dissolved in distilled water and made up the volume to one litre. That solution was diluted 100 times to get 10ppm K solution. The reading of the galvanometer was adjusted to 100 by aspirating 10 ppm K solution and then readings of solutions containing 2, 4, 6 and 8 ppm K were taken. A curve was plotted showing relationship between the K concentration on X-axis and galvanometer readings on Y-axis.

Analysis of test sample: Five gram of soil was transferred to a 150 mL conical flask and added 25 mL of neutral ammonium acetate solution. It was shaken for 5 minutes on an electric shaker and filtered. The filtrate was aspirated into the atomizer of the calibrated flame photometer and the reading was noted. This reading was located on the standard curve and the amount of K was calculated in the soil using the dilution factor.

Available K in soil (ppm) = $Y \times$ Total dilution

Where, Y = K (in ppm) as read from the standard curve Available K (Kg/ha) = Available K expressed in ppm × 2.24

4.4 Plant Collection, Herbarium Preparation and Identification

Specimens of all plant species that were encountered during the field sampling were collected, pressed and dried in field by natural drying technique in sunlight (Forman and Bridson 1989). The voucher specimens were tagged and pressed in the field with the help of newspaper and herbarium presser. Field notes were prepared including colour of the flower, fruit, fragrance or any special feature of the plants collected. When the plant specimens were completely dry, it was mounted on herbarium sheet of $16.5^{"} \times 11^{"}$ with the help of glue, and labeled properly.

The herbarium specimens were identified with the help of standard literatures (Hara et al. 1978, 1982, Hara and Williams 1979, Polunin and Stainton 1984, Stainton 1988). They were also compared with specimens at Tribhuvan University Central Herbarium (TUCH) and National Herbarium, Godawari (KATH) and some of them were identified by experts of Taxonomy. Nomenclature followed in this document is in accordance to Press et al. (2000). The herbarium specimens have been deposited at TUCH, Kirtipur, Nepal.

4.5 Numerical Analysis

To characterize vegetation of the sampling sites, the field data were used to calculate density, frequency, coverage and importance percentage. Field data of those quadrats which occupied same position in transects were pooled together (i.e. data of ten quadrats, one from each transect). Thus, there were five sets of data for each grassland sampled. Above mentioned attributes of the vegetation were calculated separately for each set of quadrats following Zobel et al. (1987) and Dallmeir (1992). The formulae were as follows:

Frequency (F, %) = $\underline{\text{Number of qudrats in which a species 'A' occurred} \times 100}$ Number of total quadrats sampled

Relative Frequency (RF, %) = <u>Frequency of a species 'A'</u> \times 100 Sum of frequencies of all species Density (D, pl/m^2) = <u>Number of individuals of a species 'A' in all quadrats</u> Total number of quadrats sampled × size of a quadrat (m²)

Relative density (RD, %) = $\frac{\text{Density of a species 'A'}}{\text{Sum of density of all species}} \times 100$

Coverage (C, %) = $T_{otal mid point values of coverage of a species 'A'}$ Total number of quadrats sampled

Relative coverage (RC, %) = $\frac{\text{Coverage of a species 'A'}}{\text{Total coverage of all species}} \times 100$

Importance percentage (IP) = $\frac{RF + RD + RC}{3}$

Species richness (number of species per sampling unit), density of *Parthenium* (pl/m²) and total density of all other species (excluding *Parthenium*) were determined for each quadrat. Plot wise data of vegetation and soil attributes were used in statistical analysis. Mean values of Parthenium density, species richness, total density excluding Parthenium, soil pH, organic matter (OM) content, Nitrogen (N), Phosphorus (P) and Potassium (K) were compared among the five sets of data (corresponding to the position of the quadrats on the transect) for each grassland using Duncan homogeneity test of one way analysis of variance (ANOVA). For correlation and regression plot wise data (n = 150) of all the three grasslands were pooled together. Spearman's correlation coefficients were determined among species richness, density of *Parthenium*, total density of other species (excluding *Parthenium*) and soil attributes. In regression *Parthenium* density was considered as explanatory variable and other vegetation and soil attributes as response variable. The data were fitted to various regression models and the best fitted model (i.e. cubic in present work) was presented. All statistical analyses were done using Statistical Package for Social Sciences (SPSS) version 11.5.

CHAPTER FIVE

5. **RESULTS**

5.1 Soil characters

Soil texture

In Gorkha, sandy loam type of soil was observed in all the plots. In Nuwakot, sandy loam type of soil was observed in invaded plots and loam type of soil in transition and non-invaded plots. In Kathmandu, soil types in different plots were found to be of various kinds and they were sandy loam in invaded plots, sandy loam in transition plots and silt loam type in non-invaded plots (Table 1).

Study	Texture		Position o	f quadrats in the	transect*	
Sites	Texture	1	2	3	4	5
Gorkha	Sand (%)	64.51±0.08	64.47±0.11	64.57±0.01	65.19±0.07	65.13±0.03
	Silt (%)	26.26±0.03	27.26±0.09	29.31±0.04	28.44±0.04	28.32±0.11
	Clay (%)	8.59±0.04	8.45±0.05	6.39±0.04	6.38±0.05	6.39±0.05
	Class	Sandy loam	Sandy loam	Sandy loam	Sandy loam	Sandy loam
	Sand (%)	48.47±0.03	48.44±0.03	44.72±0.01	59.27±0.02	59.28±0.02
Nuwakot	Silt (%)	35.32±0.01	35.36±0.01	33.23±0.01	28.5±0.01	28.4±0.02
INUWAKOU	Clay (%)	16.15±0.01	16.24±0.01	22.14±0.01	12.34±0.03	12.29±0.02
	Class	Loam	Loam	Loam	Sandy loam	Sandy loam
	Sand (%)	22.6±0.01	22.57±0.01	38.44±0.03	66.75±0.01	66.73±0.01
Kathmandu	Silt (%)	51.24±0.01	51.25±0.01	45.17±0.01	27.2±0.01	27.21±0.01
Nathmandu	Clay (%)	26.18±0.01	26.16±0.01	16.16±0.01	6.21±0.01	6.18±0.01
	Class	Silt loam	Silt loam	Sandy loam	Sandy loam	Sandy loam

Table 1. Soil texture of the study sites

* 1 and 2: non invaded, 3: transition, 4 and 5: invaded

Soil pH

The analysis of variances (ANOVA) showed significant difference (<0.001) in soil pH among the plots with different level of *Parthenium* invasion in all sites. In Gorkha site, soil pH ranged from 5.9 to 6.3 (Table 2). Invaded plots had the lowest soil pH while the transition plots had the highest. In Nuwakot site, the pH ranged from 5.8 to 6.7. The invaded plots had higher soil pH than the non invaded and transition plots, and there was no significant difference in soil pH between non invaded and transition plots. In Kathmandu site, soil pH ranged from 5.5 to 6.3, and it was lowest in non invaded plots and highest in invaded plots (Table 2).

Organic Matter (OM)

There was significant difference in soil organic matter content among non invaded, transition and invaded plots of all study sites (Table 2). In Gorkha, the OM content ranged from 0.39 to 1.21% and it was the highest in transition plots. The invaded plots had higher OM content than the non invaded plots. In Nuwakot, the OM content ranged from 1.62 to 2.11% and it was the highest in transition plots. The non invaded plots had higher OM content than the invaded plots. In Kathmandu, the OM content ranged from 1.36 to 2.14% and it was the highest in invaded plots. The transitions plots had higher OM content than the non invaded plots.

Total Nitrogen (N)

There was significant difference in soil N content among non invaded, transition and invaded plots of Gorkha and Nuwakot sites but in Kathmandu site soil N content of transition plots was not significantly different from that of non invaded plots (Table 2). The invaded plots had higher soil N content than in non invaded plots of Gorkha and Kathmandu sites but opposite pattern was observed in Nuwakot site. In Gorkha site, soil N content ranged from 0.03 to 0.08% and it increased in following order: non invaded plots < invaded plots < transition plots. In Nuwakot, soil N content ranged from 0.11 to 0.14%, with lowest value in invaded plots. It increased in following order: invaded plots < non invaded plots < transition plots. In Kathmandu, soil N content ranged from 0.10 to 0.15% and it was highest in invaded plots.

Table 2. Mean values (\pm SD, standard deviation) of soil pH, Organic matter content (OM, %), total Nitrogen (N, %), available Phosphorus (P, kg/ha) and available Potassium (K, kg/ha) in Gorkha, Nuwakot and Kathmandu sites, for each site mean values with different alphabets indicates significant difference at $\Gamma = 0.05$

Study Sites	Position of		S	oil chemical pro	perties	
	quadrats in the transect*	рН	Organic matter (%)	Total N (%)	Available P (kg/ha)	Available K (kg/ha)
Gorkha	1	6.16±.03b	0.39±0.007a	0.033±0.003a	154±2d	283±1a
	2	6.17±.05b	0.39±0.006a	0.032±0.002a	154±1d	284±1a
	3	6.30±.06c	1.21±0.005c	0.085±0.005c	151±1c	563±4c
	4	5.92±.03a	0.83±0.003b	0.056±0.003b	87±2a	307±2b
	5	5.94±.03a	0.83±0.007b	0.057±0.002b	88±1b	305±3b
Nuwakot	1	5.84±.02a	1.96±0.002b	0.13±0.002b	-	323±3a
	2	5.84±.02a	1.96±0.001b	0.13±0.002b	-	324±3a
	3	5.85±.02a	2.11±0.002c	0.14±0.002c	-	605±2b
	4	6.73±.02b	1.62±0.002a	0.11±0.002a	-	736±2c
	5	6.73±.02b	1.62±0.002a	0.11±0.002a	-	736±3c
Kathmandu	1	5.49±.16a	1.38±0.06a	0.10±0.002a	89±1a	186±2a
	2	5.54±.02a	1.36±0.003a	0.10±0.002a	88±2a	186±2a
	3	5.94±.03b	1.41±0.003b	0.10±0.002a	164±1b	186±2a
	4	6.35±.03c	2.14±0.003c	0.15±0.002b	495±2c	805±2b
	5	6.35±.02c	2.14±0.002c	0.15±0.003b	497±2d	806±2b

* 1 and 2: non invaded, 3: transition, 4 and 5: invaded

Table 3. Mean values of soil pH, organic matter content, nitrogen, phosphorus and potassium in all study sites (Gorkha, Nuwakot and Kathmandu), values of sampling plots occupying same position in transect were pooled together

	Position of sampling		
Soil attributes	plot	Mean±SD	Range
	1	0.08 ± 0.04a	0.03-0.13
	2	0.08 ± 0.04a	0.03-0.13
	3	0.1 ± 0.02a	0.08-0.14
	4	0.1 ± 0.04a	0.05-0.15
Soil nitrogen	5	0.1 ± 0.04a	0.06-0.15
	1	81.08 ± 64.37a	0.00-157.2
	2	80.64 ± 64.18a	0.00-156.20
	3	105.15 ± 75.81ab	0.00-165.50
	4	193.85 ± 219.36b	0.00-497.00
Soil phosphorus	5	195.01 ± 220.18b	00.00-499.00
	1	263.9 ± 58.68a	182.00-329.00
	2	264.33 ± 59.15a	182.00-328.00
	3	451.53 ± 191.78b	183.00-609.00
	4	616.17 ± 223.85c	302.00-808.00
Soil potassium	5	615.63 ± 225.07c	301.00-809.00
	1	5.83 ± 0.29a	5.05-6.20
	2	5.85 ± 0.27a	5.51-6.30
	3	6.03 ± 0.21b	5.82-6.40
	4	6.33 ± 0.33c	5.90-6.76
Soil PH	5	6.34 ± 0.33c	5.90-6.75
	1	1.24 ± 0.65a	0.39-1.96
	2	1.23 ± 0.65a	0.38-1.95
	3	1.58 ± 0.39a	1.21-2.11
	4	1.23 ± 0.65a	0.83-2831.00
Soil OM	5	1.53 ± 0.54a	0.82-2.14

* 1 and 2: non invaded, 3: transition, 4 and 5: invaded

Available Phosphorus (as P₂O₅)

There was significant difference in soil P (as P_2O_5) content among non invaded, transition and invaded plots of Gorkha and Kathmandu sites (Table 2) but in Nuwakot site soil P was not found in detectable amount. The non invaded plots had higher soil P (P_2O_5) content than in invaded plots of Gorkha site but opposite pattern was observed in Kathmandu site. Soil P (P_2O_5) content ranged from 87 to 154 kg/ha and it increased in following order: invaded plots < transition plots < non invaded plots in Gorkha. In Kathmandu, soil P content ranged from 88 to 497 kg/ha. It increased in following order: non invaded plots < transition plots < invaded plots.

Table 4. Results of analysis of variance (ANOVA) among different soil parameters of Gorkha, Nuwakot and Kathmandu study sites

Study site	Parameters	Source of varia-	Degree of free- dom	F value	Significance level ()
	рН		4	175	0.000
	P-1		45	175	
	initialitionPHBetween group Within groups TotalOrganicBetween group Within groups TotalNitrogenBetween group Within groups TotalNitrogenBetween group 		49		-
	Organic		4	43879	0.000
	-		45	10017	
	matter		49		-
	Nitrogen		4	661	0.000
	Throgen		45	001	
Gorkha			49		-
	Phosphorus		4	6967	0.000
	Thosphorus		45	0707	0.000
			49		-
	Potassium		4	22969	0.000
	1 Otassium		45	22707	0.000
			49		_
	рЦ		49	10660	0.000
	pn		45	10000	0.000
			43		_
	Organia		49	258028	0.000
	-		4 45	258928	0.000
	matter	<u> </u>			_
Nuwakot			49	702	0.000
I'vu wakot	Nitrogen		4	792	0.000
			45		_
			49		
	Potassium		4	63480	0.000
			45		_
			49		
	pН		4	350	0.000
			45		
			49		
	Organic	Between groups	4	3104	0.000
	matter		45		
		Total	49		
	Nitrogen	Between groups	4	2313	0.000
Kathmandu		Within groups	45		
		Total	49		
pHBa WOrganicBa WOrganicBa matterNuwakotNitrogenBa WNitrogenBa WPotassiumBa WPotassiumBa WPotassiumBa WPotassiumBa WPotassiumBa WNitrogenBa WPHBa WPhosphorusBa WNitrogenBa WNitrogenBa WTo <td>Between groups</td> <td>4</td> <td>198486</td> <td>0.000</td>	Between groups	4	198486	0.000	
	_		45		
			49		
	Potassium	Between groups	4	276544	0.000
Juwakot			45		
		U	49		1

Available Potassium (as K₂O)

There was significant difference in soil K (as K_2O) content in invaded plots of Nuwakot and Kathmandu sites but in Gorkha site soil K content of transition plots was not significantly different from that of non invaded plots (Table 2). The invaded plots had highest soil K content than in non invaded plots of Nuwakot and Kathmandu sites but in Gorkha site, soil K was highest in transition plot. In Gorkha site, soil K content ranged from 283 to 563 kg/ha and it increased in following order: non invaded plots < invaded plots < transition plots. In Nuwakot, soil K content ranged from 323 to 736 kg/ha, with lowest value in non invaded plots. It increased in following order: non invaded plots < transition plots < invaded plots. In Kathmandu, soil K content ranged from 186 to 806 kg/ha and it was highest in invaded plots.

Table 5. Results of analysis of variance (ANOVA) of soil attributes among three study sites (Gorkha, Nuwakot and Kathmandu), for each site, data of all sampling plots were pooled

Parameters	Source of varia- tion	Degree of freedom	F value	Signific- ance level ()
Soil nitrogen	Between Groups	4	2.649	
	Within Groups	145		0.036
	Total	149		
Soil phosphorus	Between Groups	4	4.659	
	Within Groups	145		0.001
	Total	149		
Soil potassium	Between Groups	4	32.146	
	Within Groups	145		0.000
	Total	149		
Soil PH	Between Groups	4	22.772	
	Within Groups	145		0.000
	Total	149		
Soil OM	Between Groups	4	1.003	
	Within Groups	145		0.408
	Total	149		

5.2 Number of species

Total number of species recorded in sampling plots were 22 in Gorkha site and 23 in both Nuwakot and Kathmandu sites (Table 6). The number of species was the highest in transition plots (position 3) of Gorkha site but in Nuwakot site, the highest number of species (17) was recorded in non-invaded plots and least number of species (13) in invaded plots which was the most disturbed plots in all three sites . In Kathmandu site the highest number of species (16) was recorded both in invaded and non invaded plots.

Study sites Number of species Total Position of quadrats in the transect* Gorkha Nuwakot Kathmandu

 Table 6. Total number of species recorded in plots of various

 positions in the study sites (Gorkha, Nuwakot and Kathmandu)

*1 and 2: non invaded quadrats, 3: transition quadrat, 4 and 5: invaded quadrats

5.3 Changes in density and species richness

The analysis of variances (ANOVA) showed significant difference (<0.001) in *Parthenium* density and total density (excluding *Parthenium*) among the plots with different level of *Parthenium* invasion in each site. *Parthenium* density in most invaded plots (position 5 in transects) ranged from 1.5 to 38 pl/m² (average 20 pl/m²) in Gorkha site, 1 to 29 pl/m² (average 15 pl/m²) in Nuwakot site and 2 to 26 pl/m² (average 13 pl/m²) in Kathmandu site. Thus maximum *Parthenium* density was recorded at Gorkha site (Table 9). Total density of all plant species excluding *Parthenium* was high in non invaded plots 222 pl/m² in Gorkha site, 219 pl/m² in Nuwakot site and 225 pl/m² in Kathmandu site and that value was the lowest in invaded plots of all sites (Table 9).

Table 7. Results of analysis of variance (ANOVA) among different parametersof Gorkha, Nuwakot and Kathmandu study sites

		Sources of vari-			Significance
Study site	Parameters	ation	df	F value	level ()
		Between Groups	4	17.423	.000
Gorkha	Parthenium	Within Groups	31		
	density	Total	35		
		Between Groups	4	1.699	0.167
Gorkha	Species rich-	Within Groups	45		
	ness	Total	49		
		Between Groups	4	12.484	0.000
		Within Groups	45		
	Total Density#	Total	49		
Nuwakot		Between Groups	4	51.753	0.00
	Parthenium	Within Groups	32		
	density	Total	36		
		Between Groups	4	0.54	0.707
	Species rich-	Within Groups	45		
	ness	Total	49		
		Between Groups	4	28.922	0.000
		Within Groups	45		
	Total Density#	Total	49		
		Between Groups	3	18.387	0.000
	Parthenium	Within Groups	30		
	density	Total	33		
Kakthnandu		Between Groups	4	3.049	0.026
	Species rich-	Within Groups	45		
	ness	Total	49		
		Between Groups	4	15.607	0.000
		Within Groups	45		
	Total Density#	Total	49		

Density excluding *Parthenium*

There was significant difference in species richness among the various positions of sampling plots in Gorkha and Kathmandu sites but the difference was not significant in Nuwakot site (Table 9). In Gorkha, species richness was the lowest in non invaded plot (position 1) and it increased towards invaded plots. In Nuwakot, species richness ranged from 4 to 7 species/m² in invaded plots and from 4 to 8 species/m² in non-invaded plots. In Kathmandu, species richness ranged from 3 to 8 species/m² in invaded plot. In this site both lowest and highest species richness were recorded in non invaded plots.

Table 8. Results of analysis of variance (ANOVA) of *Parthenium* density, species richness and total density of other plant species excluding *Parthenium* among three study sites (Gorkha, Nuwakot and Kathmandu), for each site, data of all sampling plots were pooled

Parameters	Source of vari- ation	Degree of freedom	F value	Significance level ()
Parthenium	Between Groups	4	119.335	
density	Within Groups	145		0.000
,	Total	149		0.000
Species rich-	Between Groups	4	3.013	
ness	Within Groups	145		0.020
	Total	149		
Total density#	Between Groups	4	50.686	
	Within Groups	145		0.000
	Total	149		1

Density excluding Parthenium

Table 9. Mean values of Parthenium density, species richness and total density ofother plant species excluding Parthenium in Parthenium invaded plot of Gorkha,Nuwakot and Kathmandu

Study	Community	Position of quadrats in the transect*					
Sites	attributes	1	2	3	4	5	
	Parthenium density	1.5±0.7a	1.3±0.5a	15.6±8.9b	32.0±13.5c	38.0±7.4c	
Gorkha	Species richness	5.3±0.8a	6.0±1.1ab	6.3±1.1ab	6.3±1.3ab	6.5±1.4b	
	Total density#	222±82b	172±81b	95±35a	87±32a	76±29a	
	Parthenium density	1	2.8±0.8	11.1±5.0	20.9±3.4	28.8±4.5	
Nuwakot	Species richness	5.4±1.3a	5.3±1.1a	5.8±1.1a	5.2±1.0a	5.7±1.1a	
	Total density#	219±57c	190±46c	108±48b	82±26ab	53±19a	
	Parthenium density	0	2.0±1.2	8.8±4.2	18.3±7.1	25.7±8.4	
Kathmandu	Species richness	3.9±0.9a	5.8±1.8b	5.3±0.7b	4.8±1.3ab	5.3±1.6b	
	Total density#	225±81c	142±79b	94±49ab	50±23a	57±33a	

* 1 and 2: non invaded, 3: transition, 4 and 5: invaded; # Density excluding Parthenium

Table 10. Mean values of *Parthenium* density, species richness, total density (excluding *Parthenium*) for all study sites (Gorkha, Nuwakot and Kathmandu), values of sampling plots occupying same position in transect were pooled together

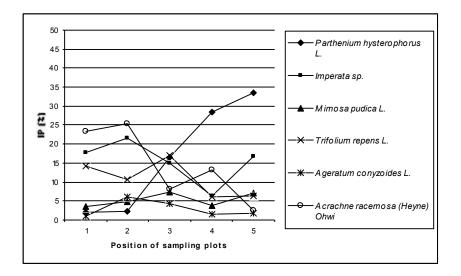
	Position of sampling		Range of val-
Parameters	plot	Mean ± SD	ues
	1	0.13 ± 0.43a	0.00-2.00
	2	1 ± 1.29a	0.00-4.00
	3	$12 \pm 6.8b$	3.00-33.00
	4	24 ± 11c	9.00-55.00
Parthenium density	5	$31 \pm 8.6d$	13.00-50.00
	1	4.87 ± 1.2a	3.00-8.00
	2	5.7 ± 1.32ab	3.00-8.00
	3	5.8 ± 1.03b	4.00-8.00
	4	5.43 ± 1.33ab	4.00-8.00
Species richness	5	5.83 ± 1.42b	3.00-8.00
	1	222 ± 72d	94.00-390.00
	2	168 ± 71c	51.00-324.00
	3	99 ± 43b	37.00-185.00
	4	73 ± 31ab	15.00-130.00
Total density#	5	62 ± 28a	14.00-118.00

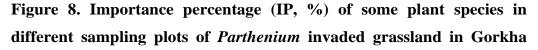
*1 and 2: non invaded, 3: transition, 4 and 5: invaded; # Density excluding Parthenium

In average, higher plant species richness was found in invaded plots and transition plot than in non-invaded plots (Table 10).

5.4 Change in Importance Percentage (IP)

The importance percentage (IP) of *Imperata* sp., *Trifolium repens* and *Acrachne racemosa* were lower in invaded plots than in non invaded plots in Gorkha site (Figure 8). The IP of *Mimosa pudica* was higher in transition and most invaded plots than in non invaded plots. The IP of *Chrysopogan aciculatus* increased from invaded plots to non-invaded plot (Appendix 1a-1e).





site * 1 and 2: non invaded, 3: transition, 4 and 5: invaded

In Nuwakot, the IP of *Mimosa pudica* increased from non-invaded plot to invaded plots (Figure 9). *Acrachne racemosa* had the highest IP in non-invaded plots (32.39%) and the lowest in most invaded plots (6.43%). The IP of *Imperata* sp. declined marginally from non-invaded to invaded plots. The IP of *Ageratum conyzoides* L. increased from non-invaded plots to invaded plots.

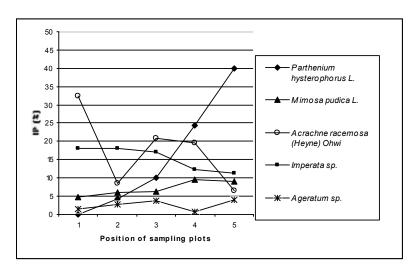
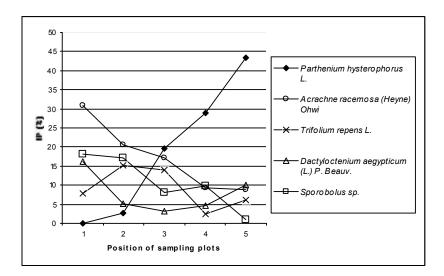
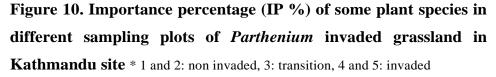


Figure 9. Importance percentage (IP %) of some plant species in different sampling plots of *Parthenium* invaded grassland in Nuwakot site * 1 and 2: non

invaded, 3: transition, 4 and 5: invaded

In Kathmandu, the IP of *Acrachne racemosa* and *Sporobolous* sp. declined from noninvaded to invaded plots (Figure 10). The IP of *Trifolium repens* was lower in invaded plots than in non invaded. The IP of *Dactyloctenium aegypticum* was the lowest in transition plot and highest in non invaded plot.





5.5 Relation between plant community and soil attributes

Correlation analysis for the combined data from all three site showed increase in species richness and decline in total density (excluding *Parthenium*) with increasing *Parthenium* density (p = 0.01) (Table 11). *Parthenium* density was positively correlated with species richness, soil pH and K content. Species richness was negatively correlated with soil N and OM content. Total density excluding *Parthenium* was negatively correlated with all measured soil attributes (Table 11).

Parameters	Parthenium density	Species richness	Total density#	Soil Nitrogen	Soil Phosphorus	Soil Potassium	Soil PH
Species							
richness	0.230**						
Total							
density#	-0.679**	-0.06					
Soil							
Nitrogen	0.148	-0.224**	-0.271**				
Soil							
Phosphorus	0.018	-0.043	-0.181*	-0.138			
Soil							
Potassium	0.546**	-0.011	-0.474**	0.669**	-0.096		
Soil pH	0.548**	0.031	-0.538**	0.079	0.244**	0.648**	
Soil OM	0.159	-0.231**	-0.285**	0.981**	-0.123	0.671**	0.094

 Table 11. Spearman's Correlation coefficients (r) among the plant community and soil characters

** Significant at p = 0.01, * significant at p = 0.05, # density excluding Parthenium

Total density of other species declined sharply with increasing density of *Parthenium* at the initial stage of invasion but the former slightly increased when the *Parthenium* density has medium value $(20 - 40 \text{ pl/m}^2)$, Figure 11). Towards the highest value of *Parthenium* density (> 40 pl/m²) the total density of other species again declined sharply. Soil potassium (K) content and pH showed hump-shaped (or dome shaped) relations with *Parthenium* density (Figures 12, 13). Soil K content was low on non invaded as well as most invaded plots. At initial phase of *Parthenium* invasion, soil pH increased towards neutral. But in most invaded plots, soil pH declined and became acidic.

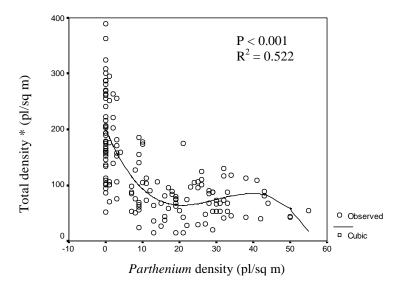


Figure 11. Relationship between total density (excluding *Parthenium*) and *Parthenium* density. The relations were obtained by regression analysis (cubical model) using combined data for all three sites. Each circle represents a sampling plot. *Total density of other plant species excluding *Parthenium*

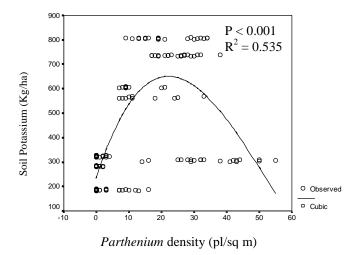


Figure 12. Relationship between potassium content of soil and *Parthenium* density. The relations were obtained by regression analysis (cubical model) using combined data for all three sites. Each circle represents a sampling plot.

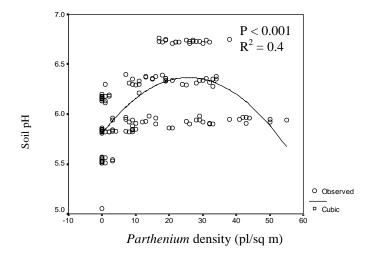


Figure 13. Relationship between soil pH and *Parthenium* density. The relations were obtained by regression analysis (cubical model) using combined data for all three sites. Each circle represents a sampling plot.

CHAPTER SIX

6. **DISCUSSION**

6.1 Soil characters

While moving from non-invaded plots to invaded plots, soil type was changed to sandy loam type in all three sites (Table 1). This result was similar to report of Joshi (2005). She reported that sandy loam types of soil have favoured the growth of *Parthenium* due to more aeration in soil. Annapurna and Singh (2003) have reported that the life cycle of *Parthenium* is slow in high clay soils because the amount of plant-available N would be lower in clay-rich soils and loamy sand would be expected to be the most fertile soil for *Parthenium*.

In all three sites soil pH declined from non-invaded plots to invaded plots (Table 3) and soil was slightly acidic or neutral. Soil might become slightly acidic or neutral in *Parthenium* invaded plots through root exudates (Bhowmik et al. 2007), which could include allelochemicals. High *Parthenium* density both in acidic and neutral soil has been reported in Kathmandu valley (Joshi 2005). However *Parthenium* is capable of growing in most soil types but becomes dominant in alkaline clay loam soil (Anonymous 2006).

In average, the soil OM and N did not change significantly from *Parthenium* noninvaded plots to invaded plots in all three study sites (Table 3). This result was contradictory with the findings of Bidwell et al. (2006) which showed that OM and N content of soil increased with the increase of invasive weed abundance in Australia. It has been reported that the extreme degree of *Parthenium* mortality was observed in the soils with highest clay contents because the amount of plant available N would be lower in clay-rich soils (Annapurna and Singh 2003). Changes in soil nutrient pools due to invasion might require longer periods of time to show differences to native patches (Collins 2005). There might have implications in soil extractable pools in long term.

Soil K increased from non-invaded plots to invaded plots (Table 3). Many grass species with fine, fibrous root systems are able to exploit K held in clay interlayer and near the edges of mica and feldspar of clay and silt (Brady and Weil 2002). Grass species were dominant in non invaded plots and also clay content was high in those plots. This result is contradictory to Collins (2005) which showed lower level of soil K in invaded plots than non-invaded plots while working with invasive species of *Imperata cylindrica* in America.

There were a number of trends in changes in soil properties that might be due to heterogeneous nature of habitat, instead of *Parthenium* invasion. The study sites were partially and newly invaded sites of about five years old and the period might be not sufficient to change soil properties. Plant invasions do not result in consistent changes in soil properties, even for the same invasive species (Collins 2005). Hook et al. (2004) suggested that *Centaurea maculosa*, might increase soil C and N pools in native grasslands in Montana (USA). and available data suggest a number of trends with respect to soil nutrients and plant invasions. Bhowmik et al. (2007) expected that *Parthenium* can survive in natural system either due to slow release of allelochemicals with less concentration in system or the chemicals donot persist in soil for a long time to effect the system.

6.2 Change in vegetation pattern

Parthenium density ranged from 1.5 to 38 pl/m^2 in invaded plots for all study sites (Table 10). The density ranged from 2 to 25.7 pl/m^2 in invaded plots of Kathmandu site which is similar to the range (11 to 47 pl/m^2) reported by Joshi (2005) from Kathmandu valley. Tiwari et al. (2005) has shown *Parthenium* density of 0.55 pl/m^2 in fallowlands which is very low as compared to present data. Very low density could be possible at the initial stage of invasion but it rapidly increases over time. Density of *Parthenium* was 55 pl/m^2 at eastern Ethiopia (Tamado et al. 2002) which was higher than the values reported in present study and other studies in Nepal. Density of associated plant species generally declined with increasing density of *Parthenium* (Figure 11) due to competitive replacement. Slight increase in density of associated species at intermediate level of *Parthenium* density (25 – 40 pl/m^2) might be due to transition from *Acrachne recemosa* dominated community to *Parthenium* dominated community. When density of mat forming *Acrachne recemosa* declined then space

could be available for growth of small herbaceous species which latter dissappeared due to increasing population of *Parthenium*. Low grazing pressure due to presence of unpalatable *Parthenium* could also have contributed to slight increase in density of associated species in these plots.

6.3 Impact of *Parthenium* invasion on species richness

When considered three sites together the species richness was the highest in the most invaded plots (Table 10). However the individual sites did not show consistent pattern in species richness with *Parthenium* invasion (Table 9). At Gorkha the species richness was the highest at most invaded plots but at other two sites high species richness was recorded both in non invaded and invaded plots. Thus it is clear that despite the lack of consistent pattern for individal sites, the species richness did not declined due to *Parthenium* invasion at this initial stage. The non invaded plots in all sites were dominated by Acrachne recemosa (Figures 8-10) which formed continuous mat and suppressed the growth of associated species. With the initiation of Parthenium invasion importance of Acrachne recemosa declined and dominance shifted to Parthenium. Acrachne racemosa forms mat like layer due to its extensive rhizome and root network which doesnot allow the growth of other plant species at ground level. Parthenium lacks such type of network but creats shaded condition which favours growth of other plant species under dense canopy at ground level. Due to different growth form, Parthenium allowed the growth of small herbaceious species, leading to high species richness at initial stage of invasion. But latter most of the associated species would disappear due to allelopathic effect and competitive replacement by *Parthenium* (Yadav and Chauhan 1998, Sinha and Deo 1999). Thus history of plant invasion should be considered while accounting the species richness in relation to invasion. Melbourne et al. (2007) also showed that diversity can increased as a result of biological invasions. Invasion directly doesnot replace native species at first it creats a heterogeneous environmental condition which is formed due to coexistance of different plant species at the same time. In present study sites, heterogeneous environmental conditions might have created due to presence of *Parthenium* in patches at initial stage and increase the species richness.

6.4 Response of native species to *Parthenium* invasion

Acrachne racemosa is a palatable grass of family Poaceae. Importance percentage (IP) of A. racemosa declined from non invaded plots to invaded ones (Figures 8-10) in all study sites. Dominant shifts from Acrachne racemosa to Parthenium moving from non invaded plots to invaded plots and also Acrachne racemosa has prostrate rhizome and root network. Likewise Trifolium repens, Imperata sp., Chrysopogan aciculatus, Sporobolus sp. and Dactyloctenium aegypticum were also affected by *Parthenium* invasion. These plant species are palatable species and so there might be possible impact on fodder supply. Thus *Parthenium* competes directly with pasture species, reducing pasture vigour and seed set and leading to habitat and ecosystem change (Evans 1997, O'Donnell and Adkins 2005, Shabbir and Bajwa 2006). There might be reduction in pasture production after several years of *Parthenium* invasion. Parthenium releases chemicals that inhibit the germination and growth of pasture grasses and other plants (Bhowmik et al. 2007). IP of Ageratum conyzoides and Mimosa pudica increased from non invaded plots to invaded (Figure 8, 9). That was the evidence of competitive vigour of those plant species to Parthenium. The IP of Euphorbia hirta and Cassia tora did not vary significantly with different levels of Parthenium invasion which meant that those species might compete with Parthenium. Commelina bengalesis, Bidens pilosa var. minor, Drymeria cordata, Justi*cia* sp. and *Alternanthera* sp. were present in invaded plots and absent in non invaded plots. *Commelina bengalesis* and *Alternanthera* sp. were delicate species; thus these two species might be suppressed due to grazing and trampling in non invaded plots. Spilanthus sinensis var. amoena and Hydrocotyle sp. were present in non invaded plots but absent in invaded plots (Appendix 1a-3e); thus these species could be more sensitive to invasion.

CHAPTER SEVEN

CONCLUSION AND RECOMMENDATION

7.1 Conclusions

Soil became slightly acidic and neutral due to *Parthenium* invasion. Soil OM and soil N content remained unchanged and soil K increased due to *Parthenium* invasion at initial stage but it declined with increasing *Parthenium* density. Plant species richness was high in invaded plots probably due to increase in habitat heterogeneity at initial stage of invasion. Dominance shifted from *Acrachne racemosa* to *Parthenium* and density of associated species declined with increasing *Parthenium* density. Many palatable species (*Acrachne racemosa* (Heyne) Ohwi., *Trifolium repens* L., *Imperata* sp., *Chrysopogan aciculatus* (Retz.) Trin., *Sporobolus* sp. and *Dactyloctenium aegypticum* (L.) P. Beauv.) were negatively affected by *Parthenium* invasion.

7.2 Recommendations

Based on the results of present study the following recommendations have been made:

-) Palatable species of grasslands were found to be highly sensitive to *Parthenium* invasion. Thus grazing lands should be protected from *Parthenium* invasion to ensure sustainable supply of fodder to livestocks.
-) Maintainance of correct stock numbers is the most important in controlling *Parthenium* because high grazing decreases vigour and competitiveness of pastures and allows the entry and spread of *Parthenium*.
-) The researches related to change of soil properties due to invasive species such as *Parthenium* are very scarce and there is an urgent need to focus on this field.

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APPENDICES

Appendix 1a. Values of density, frequency, coverage, relative density, relative frequency, relative coverage and importance percent (IP) of all plant species in *Parthenium hysterophorus* L. non-invaded plots (position 1) of Gorkha site.

S.N.	Name of Species	Density	Frequency	Coverage	Relative Density	Relative Frequency	Relative Coverage	IP
1	Acrachne racemosa (Heyne) Ohwi	57.6	50	30.75	29.16	10	30.92	23.36
2	Imperata sp.	40.4	50	22.55	20.45	10	22.67	17.7
3	Trifolium repens L.	29	70	13.75	14.68	14	13.82	14.16
4	Chrysopogan aciculatus (Retz.) Trin.	20.7	80	10.75	10.48	16	10.8	12.42
5	Oplismenus sp.	32	10	8.5	16.2	2	8.54	8.91
6	Euphorbia hirta L.	5	60	3.6	2.53	12	3.6	6.04
7	Cassia tora L.	4.1	50	3.55	2.07	10	3.56	5.21
8	Mimosa pudica L.	3.1	30	3.25	1.56	6	3.26	3.6
9	Sida rhombifolia L.	3	30	1.8	1.51	6	1.8	3.1
10	Parthenium hysterophorus L.	0.5	30	0.15	0.25	6	0.15	2.13
11	Ageratum conyzoides L.	1	10	0.25	0.5	2	0.25	0.91
12	Cynodon dactylon (L.) Pers.	0.5	10	0.25	0.25	2	0.25	0.83
13	Aconogonum sp.	0.5	10	0.25	0.25	2	0.25	0.83
14	Cyperus sp.	0.1	10	0.05	0.05	2	0.05	0.7
	Total	197.5		99.45	99.94	100	99.92	99.9

Appendix 1b. Values of density, frequency, coverage, relative density, relative frequency, relative coverage and importance percent (IP) of all plant species in *Parthenium hysterophorus* L. non-invaded plots (position 2) of Gorkha site.

S.N.	Name of Species	Density	Frequency	Coverage	Relative	Relative	Relative	IP
					Density	Frequency	Coverage	
1	Acrachne racemosa (Heyne) Ohwi	63.5	60	34.75	36.34	9.52	30.03	25.29
2	Imperata sp.	46.9	80	29.25	26.84	12.69	25.28	21.6
3	Chrysopogan aciculatus (Retz.) Trin.	17.3	90	10.25	9.9	14.28	8.85	11.01
4	Trifolium repens L.	22.6	60	11.25	12.93	9.52	9.72	10.72
5	Cassia tora L.	4.3	80	6.6	2.46	12.69	5.7	6.95
6	Ageratum conyzoides L.	5.9	40	10.3	3.37	6.34	8.9	6.2
7	Euphorbia hirta L.	4.2	50	3.75	2.4	7.93	4.32	4.88
8	Mimosa pudica L.	4.2	50	4.8	2.4	7.93	4.14	4.82
9	Parthenium hysterophorus L.	0.5	40	0.2	0.28	6.34	0.17	2.26
10	Sida rhombifolia L.	1.1	20	1.55	0.62	3.17	1.33	1.7
11	Setaria glauca (L.) P. Beauv.	3	10	1.5	1.71	1.58	1.29	1.52
12	Cynodon dactylon (L.) Pers.	1.2	10	1.5	0.68	1.58	1.29	1.18
	Total	174.7		115.7	99.93	93.57	101.02	98.13

Appendix 1c. Values of density, frequency, coverage, relative density, relative frequency, relative coverage and importance percent of (IP) all plant species in *Parthenium hysterophorus* L. transition plots (position 3) of Gorkha site.

S.N.	Name of Species	Density	Frequency	Coverage	Relative	Relative	Relative	IP
					Density	Frequency	Coverage	
1	Trifolium repens L.	27.2	60	18.25	24.81	9.67	16.55	17.01
2	Parthenium hysterophorus L.	15.6	100	21	14.23	16.12	19	16.45
3	Imperata sp.	20.7	60	18.25	18.88	9.67	16.55	15.03
4	Acrachne racemosa (Heyne) Ohwi	7.6	30	13.75	6.93	4.83	12.47	8.07
5	Mimosa pudica L.	6.2	60	7.75	5.65	9.67	7.2	7.44
6	Chrysopogan aciculatus (Retz.) Trin.	8	50	6.25	7.29	8.06	5.66	7
7	Euphorbia hirta L.	3.7	50	3.75	3.37	8.06	3.4	4.94
8	Oplismenus sp.	4.1	30	6.75	3.74	4.83	6.12	4.89
9	Ageratum conyzoides L.	4.2	30	4.5	3.83	4.83	4.08	4.24
10	Sida rhombifolia L.	4.3	40	2.25	3.92	6.45	2.04	4.13
11	Cassia tora L.	2.4	40	2.25	2.18	6.45	2.04	3.55
12	Cynodon dactylon (L.) Pers.	2.5	20	3	2.28	3.22	2.72	2.74
13	Bidens pilosa var. minor (Blume) Sherff	0.9	10	1.5	0.82	1.61	1.36	1.26
14	Setaria glauca (L.) P. Beauv.	0.9	10	0.25	0.82	1.61	0.22	0.88
15	Hydrocotyle sp.	0.6	10	0.25	0.54	1.61	0.22	0.79
16	Oxalis corniculata L.	0.5	10	0.25	0.45	1.61	0.22	0.76
17	Dioscorea bulbifera L.	0.2	10	0.25	0.18	1.61	0.22	0.22
	Total	109.6		110.25	99.92	99.91	100.07	99.4

Appendix 1d. Values of density, frequency, coverage, relative density, relative frequency, relative coverage and importance percent (IP) of all plant species in *Parthenium hysterophorus* L. invaded plots (position 4) of Gorkha site.

S.N.	Name of Species	Density	Frequency	Coverage	Relative	Relative	Relative	IP
					Density	Frequency	Coverage	
1	Parthenium hysterophorus L.	32	100	47.5	29.96	15.62	40.08	28.55
2	Acrachne racemosa (Heyne) Ohwi	17.2	40	20.25	16.1	6.25	17.8	13.14
3	Oplismenus sp.	10.7	40	13	10.01	6.25	10.97	9.07
4	Chrysopogan aciculatus (Retz.)	11.2	50	5	10.48	7.81	4.21	7.5
	Trin.							
5	Imperata sp.	1.6	60	10	1.49	9.37	8.43	6.43
6	Euphorbia hirta L.	6.4	60	3.8	5.99	9.37	3.2	6.18
7	Trifolium repens L.	7.4	40	6	6.92	6.25	5.06	6.07
8	Cynodon dactylon (L.) Pers.	4.9	40	6	4.58	6.25	5.06	5.29
9	Sida rhombifolia L.	2.5	60	0.9	2.34	9.37	0.75	4.15
10	Mimosa pudica L.	4.3	40	1	4.02	6.25	0.84	3.7
11	Cassia tora L.	2.6	40	2.05	2.43	6.25	1.72	3.46
12	Commelina bengalesis L.	1	20	1.75	0.93	3.12	1.47	1.84
13	Setaria glauca (L.) P. Beauv.	1.6	20	0.5	1.49	3.12	0.42	1.67
14	Ageratum conyzoides L.	1.4	20	0.5	1.31	3.12	0.42	1.61
15	Dioscorea bulbifera L.	0.2	10	0.25	0.18	1.56	0.21	0.65
	Total	105		118.5	98.23	99.96	100.64	99.31

Appendix 1e. Values of density, frequency, coverage, relative density, relative frequency, relative coverage and importance percent (IP) of all plant species in *Parthenium hysterophorus* L. invaded plots (position 5) of Gorkha site.

S.N.	Name of Species	Density	Frequency	Coverage	Relative	Relative	Relative	IP
					Density	Frequency	Coverage	
1	Parthenium hysterophorus L.	38	100	54.75	33.71	15.38	51.62	33.57
2	Imperata sp.	28.4	80	13.5	25.19	12.3	12.75	16.74
3	Mimosa pudica L.	5.5	70	5.75	4.88	10.76	5.42	7.02
4	Trifolium repens L.	8.3	40	6	7.36	6.15	5.65	6.38
5	Chrysopogan aciculatus (Retz.) Trin.	7.5	50	3.75	6.65	7.69	3.53	5.95
6	Euphorbia hirta L.	5.1	70	1.75	4.52	10.76	1.65	5.64
7	Cassia tora L.	3.6	60	4	3.19	9.23	3.77	5.39
8	Cynodon dactylon (L.) Pers.	3.9	30	4.5	3.46	4.61	4.24	4.1
9	Sida rhombifolia L.	2.1	40	1	1.86	6.15	0.94	2.98
10	Setaria glauca (L.) P. Beauv.	1.9	30	2	1.68	4.61	1.88	2.72
11	Acrachne racemosa (Heyne) Ohwi	2.8	10	3.75	2.48	1.53	3.53	2.51
12	Oplismenus sp.	3	20	1.75	2.66	3.07	1.65	2.46
13	Ageratum conyzoides L.	0.9	20	1.55	0.79	3.07	1.46	1.77
14	Bidens pilosa var. minor (Blume)	0.6	10	1.5	0.53	1.53	1.41	1.15
	Sherff							
15	Aconogonum sp.	0.9	10	0.25	0.79	1.53	0.23	0.85
16	Commelina benghalensis L.	0.2	10	0.25	0.17	1.53	0.23	0.64
	Total	112.7		106.05	99.92	99.9	99.96	99.87

Appendix 2a. Values of density, frequency, coverage, relative density, relative frequency, relative coverage and importance percent (IP) of all plant species in *Parthenium hysterophorus* L. non-invaded plots (Position 1) of Nuwakot site.

S.N.	Name of Species	Density	Frequency	Coverage	Relative	Relative	Relative	IP
					Density	Frequency	Coverage	
			-					
1	Acrachne racemosa (Heyne) Ohwi	98.8	80	42.25	41.91	15.09	40.18	32.29
2	Imperata sp.	61.5	40	21.5	26.09	7.54	20.44	18.02
3	Trifolium repens L.	34.4	40	15.25	14.59	7.54	14.5	12.21
4	Oplismenus sp.	9.4	40	3.3	3.98	7.54	3.13	4.88
5	Hyptis suaveolens (L.) Poit	5.6	30	6.75	2.37	5.66	6.41	4.81
6	Mimosa pudica L.	3.8	50	3.35	1.61	9.43	3.18	4.74
7	Cyperus sp.1	3.5	50	2.3	1.48	9.43	2.18	4.36
8	Conyza sp.	3.7	40	2.25	1.56	7.54	2.13	3.74
9	Cassia tora L.	1.4	40	0.6	0.59	7.54	0.57	2.9
10	Chrysopogan aciculatus (Retz.) Trin.	2.3	20	1.75	0.97	3.77	1.66	2.13
11	Cynodon dactylon (L.) Pers.	4.8	10	1.5	2.03	1.88	1.42	1.77
12	Ageratum conyzoides L.	1.2	20	0.5	0.59	3.77	0.47	1.61
13	Ipomoea sp.	1.2	20	0.5	0.59	3.77	0.47	1.61
14	Aerva sp.	1	20	0.3	0.42	3.77	0.28	1.46
15	Spilanthus sinensis var. amoena	1.8	10	1.5	0.76	1.88	1.42	1.35
	H.Hara							
16	Euphorbia hirta L.	1.1	10	1.5	0.46	1.88	1.42	1.25
17	Solanum surattense Burm.f.	0.2	1	0.05	0.08	1.88	0.04	0.66
	Total	235.7	<u></u>	105.15	100.08	99.91	99.9	99.79

Appendix 2b. Values of density, frequency, coverage, relative density, relative frequency, relative coverage and importance percent (IP) of all plant species in *Parthenium hysterophorus* L. non-invaded plots (Position 2) of Nuwakot site.

S.N.	Name of Species	Density	Frequency	Coverage	Relative	Relative	Relative	IP
					Density	Frequency	Coverage	
1	Oplismenus sp.	68.9	50	34.25	35.92	9.43	30.56	25.3
2	Imperata sp.	41	60	24	21.37	11.32	21.41	18.03
3	Trifolium repens L.	34.5	4	20	17.98	7.54	17.84	14.45
4	Acrachne racemosa (Heyne) Ohwi	14.1	50	9.75	7.35	9.43	8.7	8.49
5	Mimosa pudica L.	4.5	70	2.8	2.34	13.2	2.49	6.01
6	Hyptis suaveolens (L.) Poit	6.3	40	4.55	3.28	7.54	4.06	4.96
7	Parthenium hysterophorus L.	1.7	60	1.1	0.88	11.32	0.98	4.39
8	Cassia tora L.	2.2	40	2.05	1.14	7.54	1.82	3.5
9	Chrysopogan aciculatus (Retz.)	4.1	20	3	2.13	3.77	2.67	2.85
	Trin.							
10	Setaria glauca (L.) P. Beauv.	5.8	10	3.75	3.02	1.88	3.34	2.74
11	Ageratum conyzoides L.	1.7	30	1.8	0.88	5.66	1.6	2.71
12	Euphorbia hirta L.	0.8	20	1.75	0.41	3.77	1.56	1.91
13	Cynodon dactylon (L.) Pers.	2.8	10	1.5	1.48	1.88	1.33	1.56
14	Conyza sp.	2.6	10	1.5	1.35	1.88	1.33	1.52
15	Aerva sp.	0.8	20	0.25	0.41	3.77	0.22	1.46
	Total	191.8		112.05	99.94	99.93	99.91	

Appendix 2c. Values of density, frequency, coverage, relative density, relative frequency, relative coverage and importance percent (IP) of all plant species in *Parthenium hysterophorus* L. transition plots (Position 3) of Nuwakot site.

S.N.	Name of Species	Density	Frequency	Coverage	Relative Density	Relative Frequency	Relative Coverage	IP
1	Acrachne racemosa (Heyne) Ohwi	39.2	70	19.5	32.9	12.06	17.52	20.82
2	Parthenium hysterophorus L.	11.1	100	37.5	9.31	17.24	33.7	20.08
3	Imperata sp.	30.5	70	15	25.6	12.06	13.48	17.04
4	Oplismenus sp.	15.2	40	15.25	12.76	6.89	13.7	11.11
5	Mimosa pudica L.	3.2	60	6.1	2.68	10.34	5.48	6.16
6	Cassia tora L.	2.8	70	3.85	2.35	12.06	3.46	5.95
7	Ageratum conyzoides L.	3	40	2.25	2.51	6.89	2.02	3.8
8	Aerva sp.	3.7	20	3	3.1	3.44	2.69	3.07
9	Hyptis suaveolens (L.) Poit	2.5	30	2	2.09	5.17	1.79	3.01
10	Setaria glauca (L.) P. Beauv.	3.7	20	1.75	3.1	3.44	1.57	2.7
11	Trifolium repens L.	1.5	10	1.5	1.25	1.72	1.34	1.43
12	Evolvulus nummularis (L.) L.	1	10	1.5	0.83	1.72	1.34	1.29
13	Cyperus sp.1	0.6	10	1.5	0.5	1.72	1.34	1.18
14	Ipomoea sp.	0.7	10	0.25	0.58	1.72	0.22	0.84
15	Euphorbia hirta L.	0.2	10	0.25	0.16	1.72	0.22	0.7
16	Solanum surattense Burm.f.	0.1	10	0.05	0.08	1.72	0.04	0.61
	Total	119		111.25	99.8	99.91	99.91	99.79

Appendix 2d. Values of density, frequency, coverage, relative density, relative frequency, relative coverage and importance percent (IP) of all plant species in *Parthenium hysterophorus* L. invaded plots (Position 4) of Nuwakot site.

S.N.	Name of Species	Density	Frequency	Coverage	Relative Density	Relative Frequency	Relative Coverage	IP
					5	. ,	0	
1	Parthenium hysterophorus L.	20.09	100	37.5	19.36	17.54	36.05	24.31
2	Acrachne racemosa (Heyne) Ohwi	28.8	80	18.75	26.69	14.03	18.02	19.58
3	Imperata sp.	14.9	70	11.3	13.8	12.28	10.86	12.31
4	Oplismenus sp.	15.5	50	7.5	14.36	8.77	7.21	10.11
5	Mimosa pudica L.	6.3	80	9.05	5.83	14.03	8.7	9.52
6	Cassia tora L.	5.6	70	8	5.18	12.28	7.69	8.38
7	Hyptis suaveolens (L.) Poit	6.7	40	3.5	6.2	7.01	3.36	5.52
8	Cynodon dactylon (L.) Pers.	4.2	20	3	3.89	3.5	2.88	3.42
9	Aerva sp.	3.2	30	3.25	2.96	5.26	3.12	2.78
10	Sida rhombifolia L.	0.4	30	0.15	0.37	5.26	0.14	1.92
11	Solanum surattense Burm.f.	0.1	10	1.5	0.09	1.75	1.44	1.09
12	Ageratum conyzoides L.	0.7	10	0.25	0.64	1.75	0.24	0.87
13	Euphorbia hirta L.	0.6	10	0.25	0.55	1.75	0.24	0.84
	Total	107.09		104	99.92	105.21	99.95	100.65

Appendix 2e. Values of density, frequency, coverage, relative density, relative frequency, relative coverage and importance percent (IP) of all plant species in *Parthenium hysterophorus* L. invaded plots (Position 5) of Nuwakot site.

S.N.	Name of Species	Density	Frequency	Coverage	Relative Density	Relative Frequency	Relative Coverage	IP
					Density	rrequency	Coverage	
1	Parthenium hysterophorus L.	28.8	100	71.5	35.46	17.8	66.79	40.01
2	Oplismenus sp.	15.3	70	9.25	18.84	12.5	8.64	13.32
3	Imperata sp.	10.6	80	6.8	13.05	14.28	6.35	11.22
4	Mimosa pudica L.	6.3	80	5.75	7.75	14.28	5.37	9.13
5	Cassia tora L.	5.2	60	6.5	6.4	10.71	6.07	7.72
6	Acrachne racemosa (Heyne) Ohwi	5.6	50	3.75	6.89	8.92	3.5	6.43
7	Ageratum conyzoides L.	3.2	40	1.25	3.94	7.14	1.16	4.08
8	Trifolium repens L.	4.5	10	1.5	5.54	1.78	1.4	2.9
9	Sida rhombifolia L.	0.4	30	0.15	0.49	5.35	0.14	1.99
10	Euphorbia hirta L.	0.7	10	0.25	0.86	1.78	0.23	0.95
11	Aconogonum sp.	0.3	10	0.25	0.36	1.78	0.23	0.79
12	Ipomoea sp.	0.2	10	0.05	0.24	1.78	0.04	0.68
13	Bidens pilosa var. minor (Blume) Sherff	0.1	10	0.05	0.12	1.78	0.04	0.64
	Total	81.2		107.05	99.94	99.88	99.96	99.86

Appendix 3a. Values of density, frequency, coverage, relative density, relative frequency, relative coverage and importance percent (IP) of all plant species in *Parthenium hysterophorus* L. non-invaded plots (Position 1) of Kathmandu site.

S.N.	Name of Species	Density	Frequency	Coverage	Relative Density	Relative Frequency	Relative Coverage	IP
1	Acrachne racemosa (Heyne) Ohwi	77.3	50	33.25	48.64	13.15	31.11	30.96
2	Sporobolus sp.	7.1	70	33.5	4.46	18.42	31.45	18.11
3	Dactyloctenium aegypticum (L.) P. Beauv.	34.7	70	9	21.83	18.42	8.42	16.22
4	Parthenium hysterophorus L.	13.7	40	4.75	8.62	10.52	4.44	7.86
5	Xanthium strumarium L.	3.3	40	8.05	2.07	10.52	7.53	6.7
6	Eragrostis pilosa (L.) P. Beauv	6.1	30	6.25	3.83	7.89	6.31	6.01
7	Setaria glauca (L.) P. Beauv.	3.1	30	3.05	1.95	7.89	2.85	4.23
8	Cyperus sp.1	4.2	20	1.75	2.64	5.26	1.63	3.17
9	Hydrocotyle sp.	5	10	3.75	3.14	2.63	3.5	3.09
10	Cyperus sp.2	3.2	10	1.5	2.01	2.63	1.4	2.01
11	Oplismenus sp.	1.2	10	1.5	0.75	2.63	1.4	1.59
	Total	158.9		106.35	99.94	99.96	100.04	99.95

Appendix 3b. Values of density, frequency, coverage, relative density, relative frequency, relative coverage and importance percent (IP) of all plant species in *Parthenium hysterophorus* L. non-invaded plots (Position 2) of Kathmandu site.

S.N.	Name of Species	Density	Frequency	Coverage	Relative Density	Relative Frequency	Relative Coverage	IP
1	Acrachne racemosa (Heyne) Ohwi	41.6	60	22.75	29.46	11.11	21.54	20.7
2	Sporobolus sp.	27	70	20.75	19.12	12.96	19.64	17.24
3	Trifolium repens L.	24	90	12.25	16.99	16.66	11.6	15.08
4	Xanthium strumarium L.	4.1	40	19	2.9	7.4	17.99	9.43
5	Setaria glauca (L.) P. Beauv.	9.1	60	4.8	6.44	11.11	4.54	7.36
6	Eragrostis pilosa (L.) P. Beauv	6.1	30	9	4.32	5.55	8.52	6.13
7	Dactyloctenium aegypticum (L.) P. Beauv.	5.8	40	4.55	4.1	7.4	4.3	5.26
8	Cyperus sp.1	4.2	30	3.25	2.97	5.55	3.07	3.86
9	Drymeria cordata (L.) Wild. Ex Roem. and Schult	9	10	1.5	6.37	1.85	1.42	3.21
10	Brachiaria sp.	5.8	10	3.75	4.1	1.85	3.55	3.16
11	Parthenium hysterophorus L.	0.8	40	0.4	0.56	7.4	0.37	2.77
12	Hydrocotyle sp.	2.7	20	1.55	1.91	3.7	1.46	2.35
13	Oplismenus sp.	0.7	20	1.75	0.49	3.7	1.65	1.94
14	Oxalis corniculata L.	0.2	10	0.25	0.14	1.85	0.23	0.74
15	Ageratum conyzoides L.	0.1	10	0.05	0.07	1.85	0.04	0.65
	Total	141.2		105.6	99.94	99.94	99.92	99.88

Appendix 3c. Values of density, frequency, coverage, relative density, relative frequency, relative coverage and importance percent (IP) of all plant species in *Parthenium hysterophorus* L. transition plots (Position 3) of Kathmandu site.

S.N.	Name of Species	Density	Frequency	Coverage	Relative Density	Relative Frequency	Relative Coverage	IP
1	Parthenium hysterophorus L.	8.8	100	30.75	8.63	19.6	30.28	19.5
2	Acrachne racemosa (Heyne) Ohwi	30.8	50	12	30.22	9.8	11.81	17.27
3	Xanthium strumarium L.	7.9	100	20.05	7.75	19.6	19.74	15.69
4	Trifolium repens L.	23.2	60	7.75	22.76	11.76	7.63	14.05
5	Eragrostis pilosa (L.) P. Beauv	10.5	50	9.75	10.3	9.8	9.6	9.9
6	Sporobolus sp.	8.3	50	6.25	8.14	9.8	6.15	8.03
7	Cynodon dactylon (L.) Pers.	5	30	4.5	4.9	5.88	4.43	5.07
8	Setaria glauca (L.) P. Beauv.	2.9	20	3	2.84	3.92	2.95	3.23
9	Dactyloctenium aegypticum (L.) P. Beauv.	2.7	20	3	2.64	3.92	2.95	3.17
10	Chrysopogan aciculatus (Retz.) Trin.	1.2	10	1.5	1.17	1.96	1.47	1.53
11	Oxalis corniculata L.	0.5	10	1.5	0.49	1.96	1.47	1.3
12	Bidens pilosa var. minor (Blume) Sherff	0.1	10	1.5	0.09	1.96	1.47	1.17
	Total	101.9		101.55	99.93	99.96	99.95	99.91

Appendix 3d. Values of density, frequency, coverage, relative density, relative frequency, relative coverage and importance percent (IP) of all plant species in *Parthenium hysterophorus* L. invaded plots (Position 4) of Kathmandu site.

S.N.	Name of Species	Density	Frequency	Coverage	Relative Density	Relative Frequency	Relative Coverage	IP
1	Parthenium hysterophorus L.	18.3	100	43.75	27.31	20.83	38.56	28.9
2	Xanthium strumarium L.	5	70	12.5	7.46	14.58	11.01	11.01
3	Sporobolus sp.	8.6	40	9.25	12.83	8.33	8.15	9.77
4	Acrachne racemosa (Heyne) Ohwi	6.8	40	10.5	10.14	8.33	9.25	9.24
5	Cynodon dactylon (L.) Pers.	8.8	40	6	13.13	8.33	5.28	8.91
6	Setaria glauca (L.) P. Beauv.	5	40	4.75	7.46	8.33	4.18	6.65
7	Eragrostis pilosa (L.) P. Beauv	7.1	30	3.05	10.59	6.25	2.68	6.5
8	Brachiaria sp.	0.8	10	12.5	1.19	2.08	11.01	4.76
9	Dactyloctenium aegypticum (L.) P. Beauv.	3.4	30	3.25	5.07	6.25	2.86	4.72
10	Sida rhombifolia L.	0.4	40	1.65	0.59	8.33	1.45	3.45
11	Trifolium repens L.	2	10	3	2.98	2.08	2.64	2.56
12	Bidens pilosa var. minor (Blume) Sherff	0.5	20	1.75	0.74	4.16	1.54	2.14
13	Hyptis suaveolens (L.) Poit	0.3	10	1.5	0.44	2.08	1.32	1.28
	Total	65	470	110.45	96.95	97.88	97.29	97.33

Appendix 3e. Values of density, frequency, coverage, relative density, relative frequency, relative coverage and importance percent (IP) of all plant species in *Parthenium hysterophorus* L. invaded plots (Position 5) of Kathmandu site.

S.N.	Name of Species	Density	Frequency	Coverage	Relative	Relative	Relative	IP
					Density	Frequency	Coverage	
1	Parthenium hysterophorus L.	25.7	100	78.25	32.53	21.27	76.19	43.33
2	Dactyloctenium aegypticum (L.) P. Beauv.	6.1	60	10	7.72	12.76	9.73	10.07
3	Cynodon dactylon (L.) Pers.	11.2	60	4	14.17	12.76	0.85	9.26
4	Acrachne racemosa (Heyne) Ohwi	13.5	30	3.25	17.08	6.38	3.16	8.87
5	Trifolium repens L.	7.4	40	1	9.36	8.51	0.21	6.02
6	Eragrostis pilosa (L.) P. Beauv.	3.9	50	2.5	4.93	10.63	2.43	5.99
7	Setaria glauca (L.) P. Beauv.	4.1	30	2	5.18	6.38	1.94	4.5
8	Oplismenus sp.	1.8	20	0.5	2.27	4.25	0.1	2.2
9	Alternanthera sp.	2	10	0.25	2.53	2.12	0.24	1.63
10	Drymeria cordata (L.) Wild. Ex Roem. and Schult	1.3	10	0.25	1.64	2.12	0.24	1.33
11	Oxalis corniculata L.	1.1	10	0.25	1.39	2.12	0.24	1.25
12	Sporobolus sp.	0.4	10	0.25	0.5	2.12	0.24	0.95
13	Bidens pilosa var. minor (Blume) Sherff	0.2	10	0.05	0.25	2.12	0.04	0.8
14	Justicia sp.	0.1	10	0.05	0.12	2.12	0.04	0.76
15	Sida rhombifolia L.	0.1	10	0.05	0.12	2.12	0.04	0.76
16	Xanthium strumarium L.	0.1	10	0.05	0.12	2.12	0.04	0.76
	Total	79		102.7	99.91	99.9	95.73	98.48

Appendix 4a: pH categories of Soil.

Categories	рН
Strongly acidic	<4.5
Moderately acidic	4.6-5.5
Slightly acidic	5.6-6.5
Nearly neutral	6.6-7.5
Moderately alkaline	7.6-8.5
Strongly alkaline	>8.5

Appendix 4b: Nutrient rating of soil.

Nutrient Status	Organic matter (%)	Total nitrogen (%)	Available P2O5 (Kg/ha)	Available K ₂ O
				(Kg/ha)
Very low	<0.75	<0.03	<10	<55
Low	0.75-1.5	0.03-0.07	10-30	55-110
Medium	1.5-3.0	0.07015	30-55	110-280
High	3.0-5.0	0.15-0.25	55-110	280-500
Very high	>5	>0.25	>110	>500

Appendix 4c: Texture classes of soils.

Soil classes or Textual	Range in relative percentage of soil					
Names	Sand	Silt	Clay			
Sandy soil	85-100	0-15	0-10			
Loamy sand	70-90	0-30	0-15			
Sandy loam	43-80	0-50	0-20			
Loam	23-52	28-50	7-27			
Silt Ioam	0-52	50-88	0-27			
Silt	0-20	8-10	0-12			
Sandy clay loam	45-80	0-28	20-35			
Clay loam	20-45	15-53	27-40			
Silty clay loam	0-20	40-73	27-40			
Sandy clay	45-65	0-20	35-45			
Silt clay	0-20	40-60	40-60			
Clay	0-45	0-40	40-100			

Appendix 5a: Climatological data on maximum-minimum Temperature and Rainfall of Khaireni weather station, Tanahun from 2002-2006.

	Maximum		
Months	Temperature	Minimum Temperature	Rainfall
January	13.0	7.1	18.2
February	15.2	9.0	46.2
March	24.0	11.2	48.4
April	32.2	17.3	127.2
Мау	30.7	20.2	332.9
June	33.6	22.6	379.0
July	32.9	24.0	545.8
August	33.5	23.5	357.3
September	32.3	22.8	339.5
October	30.5	19.2	100.6
November	25.3	14.5	10.4
December	22.9	10.8	4.4

Appendix 5b: Climatological data on maximum-minimum Temperature and Rain-

fall of Kakani weather station, Nuwakot from 2002-2006.

	Maximum		
Months	Temperature	Minimum Temperature	Rainfall
Months	21.0	7.5	32.9
January	23.4	10.3	22.8
February	26.0	14.8	17.8
March	30.3	16.7	53.1
April	30.3	18.7	113.1
Мау	31.0	20.9	244.5
June	30.6	21.3	516.8
July	30.0	21.1	481.9
August	28.1	20.7	255.6
September	27.7	17.1	61.6
October	25.2	12.9	6.3
November	22.6	9.1	5.4

Appendix 5c: Climatological data on maximum-minimum Temperature and Rainfall of Tribhuvan International Airport weather station, Kathmandu from 2002-2006.

	Maximum Tem-		
Months	perature	Minimum Temperature	Rainfall
Months	19.3	3.0	27.5
January	22.4	6.3	21.4
February	20.7	7.5	48.4
March	28.0	12.8	84.7
April	28.6	16.0	109.6
Мау	29.4	20.2	176.5
June	28.7	20.5	437.2
July	29.1	20.3	313.1
August	28.5	19.1	190.2
September	26.8	13.9	42.8
October	23.4	8.6	13.3
November	20.2	4.3	7.2
December	19.3	3.0	27.5