MANAGEMENT OF ROOT ROT FUNGI BY GREWIA ASIATICA L. LEAVES AND ON THE GROWTH OF CROP PLANTS

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Abstract

Plant pathogenic fungi deteriorate roots leading immense losses to agricultural economy of Pakistan, for this reason research was conducted to investigate the fungicidal potential of *Grewia asiatica* leaves powder against infectious fungi to ameliorate the cowpea and bottle gourd growth. *In vitro*, 100% concentration of *G. asiatica* leaves (aqueous extract) showed prodigious growth inhibition against tested fungi (*Rhizoctonia solani, Fusarium oxysporum* and *Macrophomina phaseolina*) recorded in paper disc and well methods, while 75% concentration controlled both *F. oxysporum* and *M. phaseolina* but 50% only suppressed the mycelium of *F. oxysporum* observed in both methods as compared to control. *In vivo*, leaves powder of *G. asiatica* used as organic amendment at 0.5 and 0.1% showed effectual enhancement on plant growth, where 1.0% suppressed the *Macrophomina phaseolina*, *Rhizoctonia solani* and *Fusarium* spp. colonization notably. Leaves extract of *G. asiatica* at 100% recorded greater suppression of pathogenic fungi colonization significantly when drenched in soil, while 75% showed maximum inhibition of root decay pathogens but highest plant growth found by 50% concentration. Furthermore, when tested seeds treated with 100% extract of *G. asiatica* leaves showed elevation in the growth on both hosts and controlling the colonization of pathogenic fungi as compared to 75 and 50% concentrations.

Key words: G. asiatica leaves, Methods, Concentrations, Growth and root rotting fungi.

Introduction

Root infecting fungi considered as the imperative pathogens of plant as they have ability to survive in the debris and infested soil (Berg & McClaugherty, 2014) by producing various diseases such as softening and browning of root tips, lesions, wilting of leaves and retarded growth reported in vegetables, fruits, cereals, legumes and other economically important crops (Fravel, 2005; Gonzalez et al., 2011) creating huge economic losses (Hamon et al., 2011). Fusarium spp. produces toxin in plant causes wilting by the blockage of xylem vessels along with disrupting cell membrane permeability and metabolism reported in many crop plants (Garces de Granada et al., 2001; Pawar & Thaker, 2007). R. solani inhibit the plant growth due to rotting of roots caused by minerals and water uptake blockage (Wallwork, 1996). M. phaseolina causes charcoal rot on plant by producing toxic metabolite, phaseoline which caused vascular blockage leading to the death of plants (Smith & Carvil, 1997; Santos et al., 2016). Medicinal plant extracts possess potent control by exhibiting antimicrobial activity, non-phytotoxicity, systemicity, biodegradability against plant pathogens (Mahesh & Satish, 2008; Talibi et al., 2012). Recently, investigations on the application of medicinal plant parts explicate their phytochemicals properties representing antifungal properties against the plant pathogens could be used due to the environment friendly mode of disease management (Matthiessen & Kirkegaard, 2006; Jasuja et al., 2012) which gained the attention of researchers from the use of hazardous agrochemicals (Parekh & Chanda, 2007).

Grewia asiatica (Family-Tiliaceae) known as 'Phalsa'about ≥150 species comprises trees and shrubs which are distributed worldwide. In Pakistan, more than 10 species belongs to this genus have been reported (Ali,

1974). The fruits of G. asiatica are beneficial for disorder of heart, liver, blood along with indigestion, thirst, stomatitis, anorexia, toxaemia, asthma, diarrhoea, fever but also used for curing throat infection, tuberculosis and sexual problems (Sharma Sisodia, 2009). Phytochemical studies revealed presence of mineral, vitamin, fibre, carbohydrate, alkaloid, protein, glucosides and amino acid (Yadav, 1999; Tripathi et al., 2010). G. asiatica also contains citric acid trimethyl ester (5.10%), stigma sterol (1.23%), can pesterol (2.15%), α -methyl-1-sorboside (11.52%)and methyl ester (0.10%) exhibiting antimicrobial activities (Nair & Panikkar, 1990). The leaves of G. asiatica used to heal the skin wounds, relieve painful rashes/irritation, treatment of diabetes, pustular eruptions and also used as cattle fodder (Bhangale et al., 2010; Zi-ul-Haq et al., 2012).

The objective of present research was to explore the fungicidal efficacy of *G. asiatica*leaves against root pathogenic fungi and on the growth of cowpea and bottle gourd plants.

Materials and Methods

Extract preparation: Leaves of *G. asiatica* were collected from nursery of University of Karachi, washed thoroughly and dried under shade. Dried leaves were pulverized in to fine powder by electric grinder and stored in plastic jar. *G. asiatica* leaves powder (10g) were soaked in 90mL of sterilized distilled water and left it for 12 hours. The concentration of extract was filtered (100%) and was diluted with sterilized distilled water to prepare further different concentrations.

In vitro **experiments:** Agar well and filter paper disc methods were used to study the inhibition of root rot fungi with aqueous concentrations by using *G. asiatica*

leaves extract. Petri-plates poured with Potato Dextrose Agar (PDA) medium supplemented with antibiotics (Penicillin and Streptomycin @ 250mg/L) to avoid bacterial growth. In the centre of the poured Petriplates, a disc of each root rot fungus was inoculated. For paper disc method, sterilized Whatman filter paper disc (6mm) was soaked in respective concentrations (100, 75, 50%) for half an hour. Three different concentrations of discs were placed on three different sides of Petri-plates. Similarly, in case of agar well method, three wells filled (100uL) with three tested concentrations of leaves extracts. The fourth well and disc respectively, was poured with sterilized distilled water was taken as control. Each root rot fungus replicated thrice and was incubated for one week at room temperature (26-32°C). After the completion of incubation period, zone of growth inhibition was measured (Espinel-Ingroff et al., 2002).

In vivo experiments: Sandy loam soil was collected from Department of Botany (KU) with pH ranged from 7.2-7.6 having 40% moisture holding capacity observed by the Keen & Rakzowski (1922) method, 7≈8 sclerotia/g of M. phaseolina estimated by Sheikh & Ghaffar (1975) protocol while, 12≈13% of R. solani was calculated by Wilhelm, (1955) procedure and Fusarium spp., was estimated at 2700 CFU/g by Andrew & Pitt (1986) modification method. Soil (300g) was filled in a pot which was placed randomly in the screen house of the Department of Botany, then amended with powdered of G. asiatica leaves at 0.1, 0.5 and 1.0% w/w concentrations, respectively. After one day of amendment, seeds of bottle gourd and cowpea after treatment with 1.0% calcium hypochlorite (surface sterilization) for 2-3 minutes were sown in each pot (five seeds/pot), respectively. Un-amended soil taken as control was also placed for comparison. For soil drenching, different concentrations at 100, 75 and 50% (w/v) with leaves extract of G. asiatica was poured separately in each pot and five seeds of cow pea and bottle gourd were sown in each pot, while 25mL of sterilized distilled water was poured in pot regarded as control. Replicated thrice for each treatment. In seed treatment, cowpea and bottle seeds were treated with different concentrations (100, 75 and 50% w/v) of G. asiatica extract respectively, for ten minutes and then dried for 2-3 hours aseptically, five treated seeds were sown in each pot separately, while untreated seeds taken as control. Replicates were made thrice for each treatment and data were recorded after full growth of plants. After one month of germination, plants were uprooted and parameters of growth were recorded. Roots of each treatment was washed in sterilized distilled water (thrice) and then treated with 1% calcium hypochlorite for at least three minutes to remove soil particles and each root was cut into small pieces. Each root pieces were placed on poured Potato Dextrose Agar (PDA) supplemented with antibiotics to inhibit the growth of bacteria. Incubate tested plates for one week at room temperature (35-38°C) and after incubation

period, colonization percentage of each root pathogenic fungi was recorded. The data were calculated by two way of analysis (ANOVA) at p<0.05 using 'Statistica' software (Sokal&Rohlf, 1995) analysed through Duncan's Multiple Range Test (DMRT).

Results

A. In vitro

When sterilized disc impregnated with 100% concentration of G. asiatica leaves extract (aqueous) showed highest zone of inhibition and significantly (p<0.001) controlled the mycelium growth of R. solani, F. oxysporumand M. phaseolina followed by well method. However at 75% concentration, suppressed the growth of F. oxysporumand M. phaseolina mycelium but failed to control R. solani recorded in both paper disc and well showing full growth. Whereas concentration, significantly (p<0.001) inhibit the F. oxysporum only as compared to control (Fig. 1). The interaction between method and concentration was found significant (p<0.01) but paper disc was found better than well method (Table 1).

B. In vivo

i. Cowpea: G. asiatica leaves powder used as organic amendment with different concentrations increased the length and weight of shoot when soil was amended with 0.5% (w/w). However, greater root length and significant weight of root (p<0.01) was observed, when soil amended with 0.1% (w/w) as well as nodules were also increased in number in contrast to 1.0% (w/w)of leaves powder. However, 1.0% concentration of leaves powder reduced the colonization of pathogenic fungi remarkably followed by 0.5 and 0.1% concentrations as compared to control. Among different concentrations of G. asiatica leaves extracts drenched in soil, 50% (v/v) showed maximum growth enhancement and significantly (p<0.001; p<0.01) reduced the M. phaseolina, R. solaniand Fusarium spp. colonization. Shoot/root length and weight of root were increased when soil was drenched with 75% (w/v) leaves extract. While, shoot weight and number of nodules were increased at 100% (w/v) concentration of leaves extract but impressively suppressed the colonization of pathogenic root fungi as compared to concentrations. Furthermore, when cow pea seeds were treated with G. asiaticaleaves extract at 50 and 75% (v/v) showed better growth and remarkably reduced the plant pathogenic fungi. Shoot length and weight were increased when cowpea seeds were treated with 75% (w/v) leaves extract. Root length and weight were increased at 100% concentration of leaves extract along with number of nodules. Highest control of pathogenic fungi colonization on root was recorded when cowpea seeds were treated with 100% leaves extract followed by 50% (w/v) concentration (Fig. 2).

Paper disc method Well method 3 3 3 3 4

Fig. 1. Inhibition of root pathogenic fungi by using G. asiatica leaves aqueous extract at different concentrations.

where

 $\mathbf{A} = \textit{Fusariumoxysporum}; \ \mathbf{B} = \textit{Rhizoctoniasolani}; \ \mathbf{C} = \textit{Macrophominaphaseolina}$

 $\mathbf{1} = 100\%$ concentration, $\mathbf{2} = 75\%$ concentration, $\mathbf{3} = 50\%$ concentration, $\mathbf{4} = \text{Control}$ contains disc soaked or well inoculated with sterilized distilled water

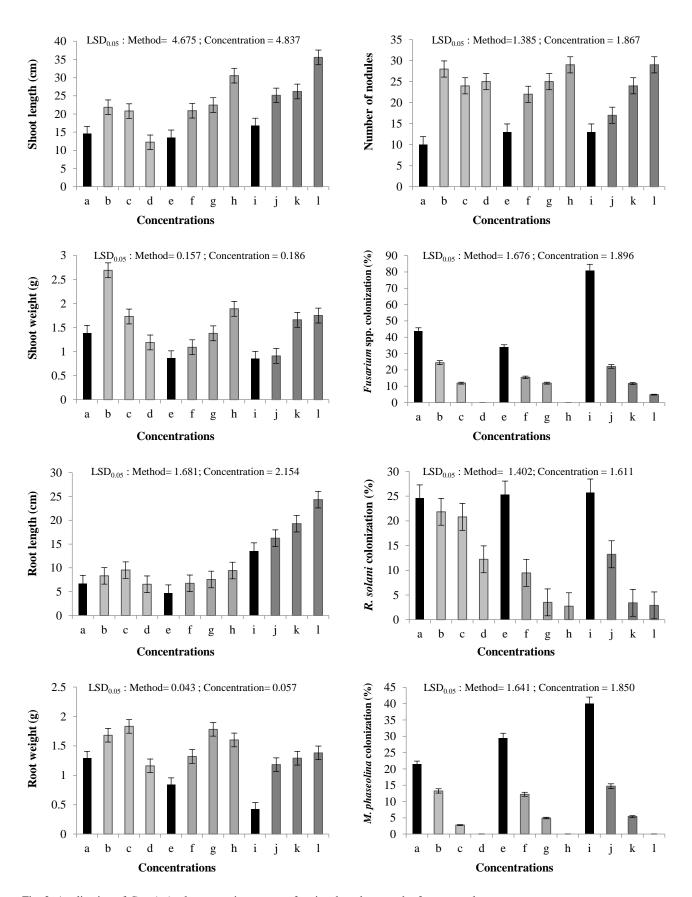


Fig. 2. Application of G. asiatica leaves against root rot fungi and on the growth of cow pea plants

where:

Soil amendment: a= Control; b= 0.1%; c= 0.5%; d= 1.0% (w/v) concentrations Soil drenching: e= Control; f= 50%; g= 75%; h= 100% (v/v) concentrations Seed treatment: i= Control; j=50%; k=75%; L= 100% (v/v) concentrations

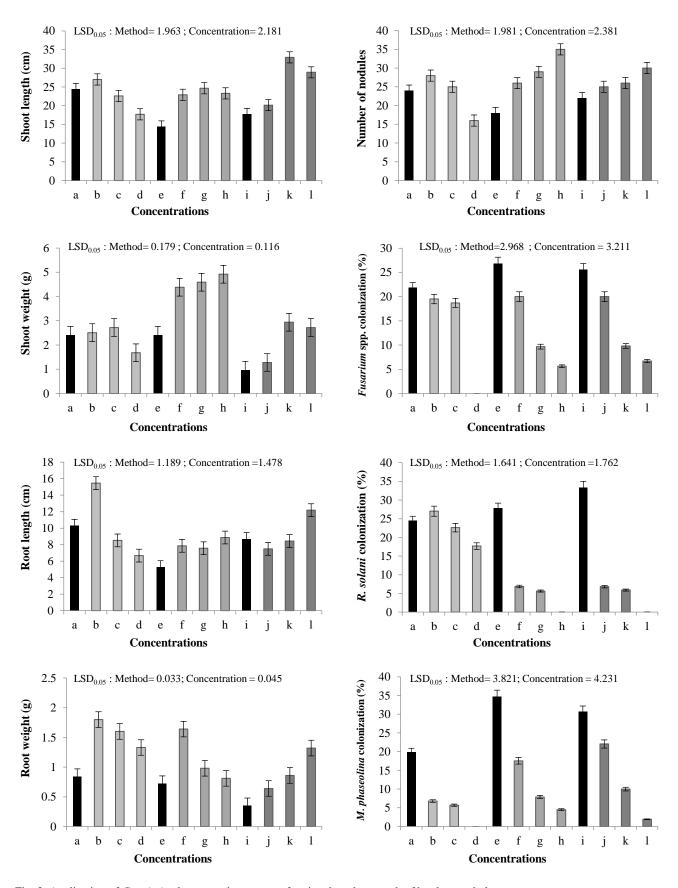


Fig. 3. Application of G. asiatica leaves against root rot fungi and on the growth of bottle gourd plants.

where:

Soil amendment: a= Control; b= 0.1%; c= 0.5%; d= 1.0% (w/v) concentrations Soil drenching: e= Control; f= 50%; g= 75%; h= 100% (v/v) concentrations Seed treatment: i= Control; j=50%; k=75%; L= 100% (v/v) concentrations

Table 1. In vitro studies showing antifungal activities of G. asiaticaleaves extract by using agar
well diffusion and naner disc methods

	Growth inhibition (%) Concentrations of G. asiatica leaves extract in µg/mL Paper disc method			
Pathogenic fungi				
	F. oxysporum	0.00 ± 0.00	86.70 ± 2.00	77.80 ± 2.08
R. solani	0.00 ± 0.00	62.90 ± 1.53	0.00 ± 0.00	0.00 ± 0.00
M. phaseolina	0.00 ± 0.00	81.60 ± 4.16	61.60 ± 2.08	0.00 ± 0.00
	Agar well diffusion method			
F. oxysporum	0.00 ± 0.00	80.00 ± 2.64	63.78 ± 2.52	44.44 ± 3.00
R. solani	0.00 ± 0.00	65.11 ± 2.08	0.00 ± 0.00	0.00 ± 0.00
M. phaseolina	0.00 ± 0.00	71.10 ± 1.08	58.40 ± 1.53	0.00 ± 0.00

 $LSD_{0.05}$: Concentrations = 1.114; Fungi = 0.965; Methods = 0.788

where; SD= Standard deviation

ii. Bottle gourd: When G. asiaticaleaves powder at 0.1% (w/w) amended in soil showed highest growth and suppressed root rot fungi colonization considerably. Shoot length and weight (p<0.05) along with number of nodules greatly increased when soil was mixed at 0.5 and 0.1% (w/w) leaves powder but length and weight of roots showed greater at 0.5% concentration of leaves powder. 1.0 and 0.5% concentrations of leaves powder significantly (p<0.05) reduced the colonization of root decay fungi followed by 0.1% leaves powder. In soil drenching, shoot length and weight were increased at 75 and 100% leaves extract, while root length/weight and number of nodules were amplified when 50% leaves extract drenched in soil. Colonization of root pathogenic fungi greatly reduced at 50 and 75% of leaves extract; whereas 100% leaves extract showed complete inhibition of pathogenic fungi. When bottle gourd seeds treated with 100% (w/v) extract of G. asiaticaleaves gave better enhancement of growth parameter and control root rot fungi colonization followed by 75% (w/v) concentration. However, seeds treated at 50% also increased the plant weight and height as compared to control. 100% extract effectively reduced the colonization of plant pathogenic fungi significantly (p<0.05) as compared to other concentrations (Fig. 3).

Overall results showed that soil drenching with *G. asiatica*leaves extract were more effectual for the growth and obtained best result in controlling root rot fungi colonization as compared to other treatments, but use of soil drenching and soil amendment in commercial scale was rather difficult to apply as compared to seed treatment method which can easily be applicable. However, these two methods were recommended on low organic farming scale.

Discussion

G. asiatica is widely cultivated in many countries (Sasteri, 1992) where its young leaves and seeds of plants are regarded as nutritional power house (Grene, 2002). G. asiatica leaves exhibiting antimicrobial properties due to the presence of flavonoids, saponins, phenolic and tannins compounds (Sato et al., 2004) having therapeutics effect which have been used to treat many plant diseases throughout the world (Elevitch, 2011). Plants with

therapeutic effect have received the attention of scientists using alternate method when used to control plant diseases (Jensen *et al.*, 1996) which also protect the safety of an environment from the use of agrochemicals (Kerr, 1980).

Present research In vitro showed best result by 100% concentration of G. asiatica leaves extract was tested against root pathogenic fungi by paper disc and well methods. According to Nayan & Shukla (2011), plants rich in phenolic compounds and tannins have been shown to possess antimicrobial activity against wide number of harmful microbes. However, in vivo experiments showed that when soil was amended with three different concentrations (0.1, 0.5 and 1.0%) respectively, revealed increment in the growth parameters on the contrary to control, whereas leaves powder at 0.5% was found best on both tested plants. Colonization of root rotting fungi (M. phaseolina, R. solani and Fusarium spp.) effectually controlled when G. asiatica leaves powder applied at 1.0% as compared to control. Surprisingly, 1.0% with leaves powder does not give better growth of plant length and weight. Similar result had been represented on Moringa oleifera leaves powder (Ejaz et al., 2014). Soil amendment with plant parts producing antimicrobial compounds during decomposition (Brown & Morra, 1997; Tenuta & Lazarovits, 2002). M. phaseolina, R. solani and Fusarium spp. inhabitant in the soil for longer period and during favourable conditions, pathogens invade inside the roots and block the xylem cells which affect the physiological process by closing the stomata present in the leaves results in the wilting, ultimately the plant die (Agrios, 1988). It was reported that diseased plant contain toxins produced by pathogenic fungi and bacteria which alter the function of stomata disturbing the transpiration process which results in wilting of plants (Aducci et al., 1995). Soil amendment gives a potential tool to control root diseases through chemical producing anti microbial compounds during biological and decomposition activity in soil protecting roots from the invasion of pathogens (Chen et al., 1998; Ikram & Dawar, 2016).

Responses of aqueous extract of *G. asiatica* leaves on the growth of tested crops were estimated by using seed treatment and soil drenching techniques. When soil was drenched with 100 and 75% concentrations of *G. asiatica* leaves, both concentrations showed maximum suppression of root rot fungi colonization and better growth of cowpea

and bottle gourd plants were observed. G. asiatica leaves extract at 50% improved the growth parameter and reduced root rot fungi colonization when drenched in soil. While, in seed treatment with G. asiatica leaves extract at 100% efficiently improved the plant growth of both crops and reduced the root infecting fungi colonization followed by 75 and 50% concentrations. Seed treatment considered as useful method for controlling soil and seed borne pathogens (Chang & Kommedahl, 1968). Leaves extract at 100% concentration gave maximum growth and inhibition of root rotting fungi on cowpea, whereas leaves extract at 75% concentration gave minimum growth as well as reduced root rot fungi on bottle gourd plants. Similar results had shown by Rafi et al., (2015) in the suppression of root rot fungi by priming seeds at 10 minutes intervals with plants extract (A. nilotica and S. mukorossi) was found effective and showed healthy growth of sunflower, okra, peanut and chickpea plants. Zia-ul-Haq et al., (2012) recorded significant results when G. asiatica leaves (ethanolic extract) tested against bacteria including; Escherichia coli, Proteus mirabilis, Citrobacter spp., staphylococcus aureus, Pseudomonas aeruginosa and Bacillus subtilis and fungi such as; Aspergillus spp. (A. niger, A. effusus, A. parasiticus), Candida albicans, Saccharomyces cerevisiae and Fusarium solani exhibiting antibacterial and antifungal activities. Patil et al., (2010) reported that using extracts of leaves of G. asiatica either in ethanol, chloroform and petroleum ether respectively, showed anti-hyperglycaemic activity. Main objective to control the plant parasitic pathogens and improved the growth and quality of plants can be achieved by the suppression of soil borne and seed borne pathogens which reduced the growth and yields causing heavy economic losses. The leaves extract of G. asiatica containing antiulcer, anti-inflammatory, anti-tumour (Pal et al., 1995; Makonnen et al., 1997) control of cholesterol and diabetes, anti-oxidant, anti-hypertensive (Mehta et al., 2003) antifungal and anti-bacterial activities (Nickon et al., 2003) due to the presence of 4-benzyl glucosinolate,4-benzyl isothiocyanate, benzyl isothiocyanate, pterygospermin and niazimicin (Napoleon et al., 2009). Majority of the antioxidant activity is due to the isoflavones, flavones, flavonoids, lignin, anthocyanin, coumarinisocatechin and catechin present in the leaves of G. asiatica (Gupta et al., 2007) and are used for the curing of different diseases such as alzheimer's disease, cancer, stroke, atherosclerosis and diabetes etc. which may be arise due to oxidative damage by free radicals (Kumar & Kumar, 2009). The phytochemical study of the leaves of G. asiatica in the extract of petroleum ether contains glycoside, fats and diterpenes. In addition, G. asiatica leaves extract in chloroform contains glycoside and alkaloids, while leaves extract in ethanol contains triterpenoids, flavonoids, sterols, tannins and saponins (Gupta et al., 2008; Patil et al., 2011). Other compounds, kaempferol, quercetion and a mixture of their glycoside were also isolated from the leaves extract of G. asiatica (Ali et al., 1982) which also contains citric acid trimethyl ester, campesterol and stigmasterol which are main compounds (Gupta et al., 2012). The aqueous leaves extract contains methyl thiazolyltetrazolin which reported anti-cancer activity against lungs, breast, cervical and kidney cancer (Marya et al., 2011). The ethanolic leaves extract showed anti-fungal and anti-bacterial activities (Zia-

ul-Haq *et al.*, 2011). The leaves extract also showed the anti-viral activity against mash bean leaf crinkle virus (Sangita *et al.*, 2009). *G. asiatica* showed highly potent plant with remarkable range of medicinal benefits and high nutritional value (Sharma & Sisodia, 2009).

The purpose of present investigation was to indicate the potentiality of *G. asiatica* leaves in the suppression of root decay fungi which help in the better growth of crop plants on cheaper bases can be applied on agricultural field.

References

- Aducci, P.M., V. Fogliano and M.R. Fullane. 1995. Fusicoccin receptors: perception and transduction of the Fusicoccin signal. *J. Exp. Bot.*, 1463-1478.
- Agrios, G. 1988. Academic press. Impresa. New York. US. *Plant pathology*, 803p.
- Ali, S.I. 1974. Flora of West Pakistan, family Tiliaceae; Fakhri printing press; Karachi Pakistan.
- Ali, S.I., N.A. Khan and I. Husain. 1982. Flavonoid constituent of *Grewia asiatica*. J. Sci. Res., 4: 55-56.
- Andrew, S. and J.I. Pitt. 1986. Selective medium for isolation of *Fusarium* species and dematiaceous hyphomycetes from cereals. *Appl. & Environ. Microbiol.*, 51(6): 1235-1238.
- Berg, B. and C. McClaugherty. 2014. Decomposition of fine root and wood litter. In: *Plant litter*. Springer Berlin Heidelberg, Germany, pp. 171-187.
- Bhangale, J., S. Acharya and T. Deshmukh. 2010. Anti-hyperglycaemic activity of ethanolic extract of *Grewia asiatica* (Phalsa) leaves in alloxan induced diabetic mice. *World. J. Pharm. Res.*, 2(5): 1486-1500.
- Brown, P.D. and M.J. Morra. 1997. Control of soil borne plant pests using glucosinolate containing plants. *Adv. In Agron.*, 61: 167-231.
- Chang, I. and T. Kommedahl. 1968. Biological control of seedling blight of corn by coating kernels with antagonistic micro organisms. *Phytopathol.*, 76: 60-65.
- Chen, J., A. Brevet, S. Blanquet and P. Plateau. 1998. Control of 5', 5'-dinucleoside triphosphate catabolism by APH1, a *Saccharomyces cerevisiae* analog of human. *FHIT. J. Bacteriol.*, 180(9): 2345-2349.
- Ejaz, A., A. Hanif and S. Dawar. 2017. Management of root rot fungi on crop plants by *Moringa oleifera* Lam. *Pak. J. Bot.*, 49(3): 1201-1209.
- Elevitch, C.R. 2011. Specially crops for pacific island Agroforestry. *Permanent Agriculture resources* (PAR), Honolulu, Hawaii.
- Espinel-Ingroff, A., A. Forthergill, J. Peter, M.G. Rinaldi and T.J. Walsh. 2002. Testing conditions for determination of minimum fungicidal concentrations of new and established antifungal agents for *Aspergillus* spp. NCCLS Collaborative Study. *J. Clin. Microbiol.*, 40(9): 3204-3208.
- Fravel, D. 2005. Commercialization and implementation of bio control. *Ann. Rev. Phytopathol.*, 43: 337-359.
- Garces de Granada, E.M. Orozco de Amezquita, G.R. Bautista and H. Valencia. 2001. *Fusarium oxysporum* et hung quenos fall conocer. *Acta Biol. Colombian*, 6: 7-25.
- Gonzalez, M., M. Pujal, J.P. Metraux, V. Garcia and M.D. Bolton. 2011. Tobacco leaf spot and root rot caused by *Rhizoctonia* solani Kuhn. Mol. Plant Pathol., 12(3): 209-216.
- Grene, R. 2002. Oxidative stress and acclimation mechanisms in plants. *The Arabidopsis book*, 1: 1-20.
- Gupta, M.K., P.K. Sharma and S.H. Ansari. 2008. Pharmaceutical evolution of *Grewia asiatica* leaves. *Hamdard Med.*, 51: 145-148.
- Gupta, M.K., R. Lagarkha, D.K. Sharma, P.K. Sharma, R. Singh and H.S. Ansari. 2007. Antioxidant activity of the successive extracts of *Grewia asiatica* leaves. *Asian. J. Chem.*, 19: 3417-3420.

Gupta, P., A. Sharma and A.K. Varma. 2012. GC/MC profiling and anti microbial effect of six Indian tropical fruit residue against clinically pathogenic bacterial strain. *Int. J. Adv. Pharm. Res.*, 3(10): 1229-1235.

- Hamon, C., A. Beranger, C.J. Coyne, R.J. Mcgee and I.L. Golf. 2011. New consistent QTL in pea associated with partial resistance to *Aphanomyces euteiches* in multiple field and controlled environment from France and the United States. *Theor. Appl. Genet.*, 123(2): 261-281.
- Ikram, N. and S. Dawar. 2016. Comparative methods of application of wild plant parts on growth and in the control of root rot fungi of leguminous crops. *Pak. J. Bot.*, 48(4): 1673-1680
- Jasuja, N. D., J. Choudhary, P. Sharma, N. Sharma and S.C. Joshi. 2012. A review on bioactive compounds and medicinal uses of *Commiphora mukul. J. Plant Sci.*, 7: 113-137.
- Jensen, S., C.S. Redwood, J.R. Jenkins, A.H. Andersen and I.D. Hickson. 1996. Human DNA topoisomerases II alpha and II beta can functionally substitute for yeast TOP2 in chromosome segregation and recombination. *Mol. Gen. Genet.*, 252(1-2): 79-86.
- Keen, B.A. and H. Rakzowski. 1922. The relation between clay content and certain physical properties of soil. *J. Agric. Sci.*, 11: 441-449.
- Kerr, I. 1980. Qualitative evaluation concepts and cases in curriculum criticism/willis, G., Ed., McCutchan, Berkely, 1978. *Aust. J. Teacher Edu.*, 5(2): 54-59.
- Kumar, S. and D. Kumar. 2009. Antioxidant and free radicals scavenging activities of edible weeds. Afr. J. Food, Agri., Nutr. & Devel., 9(5): 1174-1190.
- Mahesh, B. and S. Satish. 2008. Antimicrobial activity of some important medicinal plant against plant and human pathogens. *World J. Agri. Sci.*, 4: 839-843.
- Makonnen, E., A. Hunde and G. Dannecha. 1997. Hypoglycaemic effect of *M. stenopetal a*queous extract in rabbits. *Phototh. Res.*, 11: 147-148.
- Marya, B., K.H. Dattani, D. D. Patel, P. D. Patel, M. P. Suthar, V. P. Patel and S. B. Bothara. 2011. *In vitro* cytotoxicity evolution of aqueous fruit and leaf extracts of *Grewia asiatica* using MIT assay. *Pharm. Chem.*, 3: 282-287.
- Matthiessen, J.N. and J.A. Kirkegaard. 2006. Bio-fumigation and enhanced biodegradation: Opportunity and challenge in soil borne pest and disease management. *Crit. Rev. in Plant Sci.*, 25: 235-265.
- Nair, S.C. and K.R. Panikkar. 1990. Anti-tumour principles from *Ixora javanica*. Cancer Lett., 49(2): 121-126.
- Napoleon, P., J. Anitha and R. R. 2009. Isolation analysis and identification of phytochemicals of antimicrobial activity of M. oleifera lann. Curr. Biotica, 1(3): 33-39.
- Nayan, R.B. and V.J. Shukla. 2011. Antibacterial and antifungal activities from leaf extracts of *Cassia fistula L.*: An ethnomedicinal plant. *J. Adv. Pharm. Technol. Res.*, 2(2): 104-109.
- Nickon, F., Z.A. Soud, M.H. Rehman and M.E. Hague. 2003. In vitro antimicrobial activity of the compound isolated from chloroform extract of M. oleifera lann. Pak. J. Biol. Sci., 22: 1888-1890.
- Pal, S.K., P.K. Mukherjee and B.P. Saha. 1995. Studies on the anti-ulcer activity of *M. oleifera* leaf extract on gastric ulcer models in rats. *Phytoth. Res.*, 9: 463-465.
- Parekh, J. and S. Chanda. 2007. Antibacterial and phytochemical studies on twelve species of Indian medicinal plants. *Afr. J. Biol. Res.*, 10: 175-181.

Patil, P., M.M. Patel and C.J. Bhavsar. 2011. Preliminary phytochemical and hypoglycaemic activity of leaves of Grewia asiatica L. Res. J. Pharm. Biol. & Chem. Sci., 2(1): 516-520.

- Patil, P.S., M.M. Patel and C.J. Bhavsar. 2010. Comparative anti diabetic activity of some herbal plants. *Pharm. Sci. Monitor*, 1(1): 12-19.
- Pawar, V.C. and V.S. Thaker. 2007. Evolution of the anti Fusarium oxysporum and anti Alternaria porri effects of some essential oils. J. Microbiol. & Biotechnol., 23: 1099-1106.
- Rafi, H., S. Dawar and M.J. Zaki. 2015. Seed priming with extracts of *Acacia nilotica* (L.) Willd. ex Delile and *Sapindus mukorossi* (L.) plant parts in the control of root rot fungi and growth of plants. *Pak. J. Bot.*, 47(3): 1129-1135.
- Sangita, K., M. Avijit, P. Shilpa and J. Shivkanya. 2009. Studies of the antifungal and antiviral activity of methanolic extract of leaves of *Grewia asiatica*. *Pharmacogen*. J., 1: 221-223.
- Santos, C.A., L.M. Zamphorlin, A. Crucello, C.C. Tonoli, R. Ruller, M.A. Horta and A.P. Souza. 2016. Crystal structure and biochemical characterization of the recombinant ThBgl, a GH1 β-glucosidase over expressed in *Trichorderma harzianum* under biomass degradation conditions. *Biotechnol. For Biofuels*, 9(71): 1-11.
- Sasteri, B.N. 1992. The wealth of India, council of scientific and industrial research, New Delhi, India, 1: 425-429.
- Sato, N., L. Meijer, L. Skaltsounis, P. Greengard and A.H. Brivanlou. 2004. Maintenance of pluripotency in human and mouse embryonic stem cells through activation of Wnt signaling by a pharmacological GSK-3-specific inhibitor. *Nat. Med.*, 10(1): 55-63.
- Sharma, K.V. and R. Sisodia. 2009. Evaluation of free radicals scavenging activity and radio protective efficacy of *Grewia asiatica* fruit. *J. Radial. Prot.*, 29: 429-443.
- Sheikh, A.H. and A. Ghaffar. 1975. Population study of sclerotia of *Macrophomina phaseolina* in cotton field. *Pak. J. Bot.*, 7: 13-17.
- Smith, G.S. and O.N. Carvil. 1997. Field screening of commercial and experimental soybean cultivars for their reaction to *Macrophomina phaseolina*. *Plant Disease*, 8: 363-368.
- Sokal, R.R. and F.J. Rohlf. 1995. Biometry: The principles and practices of Statistics in biological research. Freeman, New York, pp. 887.
- Talibi, I., L. Askarne, H. Boubaker, E.H. Boudyach, F. Msanda, B. Saadi and A. Ait-Ben Aoumar. 2012. Antifungal activity of *Moroccan* medicinal plants against citrus sour rot agent *Geotrichum candidum. Lett. Appl. Microbiol.*, 55: 155-161.
- Tenuta, M. and G. Lazarovits. 2002. Ammonia and nitrous acid from nitrogenous amendments kill the microsclerotia of *Verticillium dahlia*. *Phytopathol.*, 92: 255-254.
- Tripathi, S., M. Chaurey, A. Balasubramanium and N. Balakrishan. 2010. *Grewia asiatica* Linn. As a phytomedicine. *Res. J. Pharm. Tech.*, 3: 1-3.
- Wallwork, H. 1996. Cereal root and crown disease kondinin group perth Australia, pp. 14-16.
- Wilhelm, S. 1955. Longevity of the *Verticillium* wilt fungus in the laboratory and field. *Phytopathol.*, 45: 180-181.
- Yadav, A.K. 1999. Phalsa: A potential new small fruit for Georgia. *In perspective on new crops and new uses, janick, J. Ed;* ASHS press; Alexandra, AV. USA, pp 348-352.
- Zia-ul-Haq, M., S. Ahmad, R. Amarowicz and V. Defoe. 2012. Antioxidant activity of the extracts of some cowpea (Vigna unguiculata) cultivars commonly consumed in Pakistan. Acta Pal. Pharm., 69:707-711.