



Review on ginger bacterial wilt (*Ralstonia solanacearum* Smith) and its management approaches: The case of Ethiopia

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Abstract

Ginger Bacterial wilt disease posed a threat to ginger production in countries where ginger is produced. It is caused by *Ralstonia solanacearum* (Smith) biovar III race 4. Having knowledge on biology, distribution and possible ways to manage the disease is important in designing cost effective management strategies of the disease. The objective of this paper was to give an Overview on biology, distribution, importance and management approaches of ginger bacterial Wilt caused by *Ralstonia solanacearum* (Smith). *Ralstonia solanacearum* is distributed in many Habitats all over the world and has broad host range. It is seed and soil borne pathogen that can easily transfer from one cropping season to the next by infected seed piece and soil. The symptom of the disease can be seen on above ground and underground parts of ginger plant. Efforts have been made to manage the disease in many countries including Ethiopia. Available management options for the disease are using biological control agents, cultural methods, chemical methods and integrated disease management. Integration of different management options like the use of chemicals, biological control agents, and organic amendments were found better controlled infection caused by *Ralstonia solanacearum* (Smith). The use of biological Control agents is one of the most preferable disease management options. Therefore, research should be directed towards identification and efficacy test of biological control agents against the pathogen. Moreover, integration and efficacy evaluation of different management tactics should be done in a way it is environmentally safe, economically feasible and sustainable.

Keywords: bacterial wilt, Importance, management approaches, *Ralstonia solanacearum*

Introduction

Ginger (*Zingiber officinale* Rosc.) is an important commercial crop grown for its aromatic rhizomes, which are used as both spice and medicine. It is an herb with a thickened, fleshy, perennial, subterranean rhizome and with one or more aerial leafy stems. Rhizome strong, up to 2cm thick, irregularly branched but normally only in a vertical direction, covered with deciduous, thin scales, which leave ring-like scars; epidermis corky, pale yellow to light-brown, irregularly wrinkled in the dried rhizome; roots having a smooth circular cross-section, thin, fibrous, white to light-brown; on dried rhizomes scars of leafy stems visible as shallow cup-like holes. Stem erect unbranched, mainly formed by the leaf-sheaths, up to 1m high, pale-green, often reddish at base. It was originated in Southeast Asia probably in India (Jansen, 1981) [23]. Ginger is cultivated in several parts of the world. China and India have continued to lead the world in fresh ginger production with a combined global share of over 50%, followed by Indonesia, Nepal and Nigeria. Other major producers of ginger include Bangladesh, Japan, Thailand, Philippines, Cameroon, Sri Lanka, Korea and Fiji (Henry E. C. & Adriano B. 2009) [21]. In Ethiopia Ginger cultivation started when Arabs, in the 13th century brought ginger from India to East Africa (Jansen, 1981) [23] and ginger had perhaps been known since then in Ethiopia, and predominantly grown in the wetter parts of the Southern Nations, Nationalities and Peoples' Regional State (SNNPRS). Ginger production to a lesser extent, has been also extended to some parts of Western Oromia and Northern Amhara (Tadesse *et al.*, 2016) [35]. There are different abiotic constraints that limit the

productivity of ginger such as low levels of input use (fertilizer, pesticide, improved seeds, etc.), low levels of irrigation, soil degradation and soil erosion, inadequate agricultural research and extension, and constraints in market development. Common diseases of ginger are soft rot caused by *Phytophthora* spp., bacterial wilt (*Ralstonia solanacearum*), yellows (*Fusarium oxysporum*, *Phyllosticata* leaf spot, nematode (*Pratylenchus coffeae*) and storage rots which is caused by *Pythium* sp., *Fusarium oxysporum* f. sp. *zingiberi* and *Sclerotium rolfsii* (Sharma, *et al.*, 2017) [32]. Among the biotic constraints bacterial wilt, which is caused by *Ralstonia solanacearum* is one of the most important diseases that limit its production. Bacterial wilt poses a serious damage to the cultivation of many Solanaceous crops such as tomato, potato, tobacco, pepper and eggplant in tropical, sub-tropical and temperate regions (Hayward, 1991) [20]. Bacterial wilt disease of ginger, which is one of the major constraints of ginger production in tropical and subtropical regions of world is highly heterogeneous bacterial pathogen of many economically important crops causes severe wilting (Smith *et al.* 1995) [34]. Ginger bacterial wilt is considered to be newly emerging and devastating disease which caused total crop failure in ginger producing areas of Ethiopia (Bekelle *et al.* 2016, Duressa, 2018) [5, 11]. Since the disease is currently posing serious threat to ginger production, it is very crucial to have the knowledge of biology, distribution and possible ways to manage the disease. Therefore the objective of this paper was to overview biology, distribution,

economic importance and management approaches of ginger bacterial wilt caused by *Ralstonia solanacearum*.

Biology of *Ralstonia solanacearum*

Characterization and Classification of the Pathogen

Roop Singh *et al.*, (2015) [31] studied cultural and biochemical characterization of *Ralstonia solanacearum* causing bacterial wilt in ginger. They isolated the pathogen from diseased rhizomes of *Zingiber officinale* and inoculated on triphenyl tetrazolium chloride agar, casamino peptone glucose agar, yeast extract agar and potato dextrose agar. Cream white and dull white color colonies were found on nutrient agar and yeast extract milk agar. The result revealed that the white fluidal colonies with spiral pink centre were found. Cream or off-white color colonies were found. Besides, irregular, smooth, highly fluidal colonies were found on triphenyl tetrazolium chloride agar, casamino peptone glucose agar, potato dextrose agar and yeast extract chalk agar. Round small colonies were found on nutrient agar, yeast extract milk agar, yeast extract agar and yeast extract peptone agar. Furthermore, it showed positive reactions for potassium hydroxide solubility test, catalase test, starch hydrolysis test, motility test and casein hydrolysis test and showed negative reaction for gram staining. Bacterial wilt, caused by members of the *Ralstonia solanacearum* species complex is a serious disease of many solanaceous crops in tropical and sub-tropical regions of the world. *Ralstonia solanacearum* grouped into Kingdom: Monera, Phylum: Proteobacteria, Class: Beta proteobacteria, Order: Burkholderiales, Family: Burkholderiaceae, Genus: *Ralstonia*, Species: *Solanacearum* (Tahat, and Sijam, 2010) [28]. *Ralstonia solanacearum* is distributed in many habitats all over the world and has an unusually broad host range (Hayward 1991, Denny 2006) [20, 10]. It can infect over 200 plant species representing over 50 botanical families (Hayward 1991) [20]. Due to its global distribution, adaptive potential and large host range, *Ralstonia solanacearum* has turned into a model system to study plant-microbe interactions, pathogenicity determinants and pathogen ecological behavior. Since *Ralstonia solanacearum* is a

soil-borne pathogen and that resistance of the host is limited, bacterial wilt is very difficult to control (Hayward, 1991) [20].

Sign and Symptom of the Disease

The symptom of the disease is associated with water soaked patches or linear streaks appear. These symptoms are followed by yellow to bronze colouration of margins of the lower-most leaves which gradually progresses upwards. At later stages, the leaves become flaccid with intense yellowish bronze colour and droop ultimately exhibiting typical wilt symptoms. If the affected rhizomes are pressed, a milky bacterial exudate oozes out. When infected tissues are steeped in clear water for a while, the water turns cloudy and milky (Meenu and Kaushal, 2017) [26]. The most common external symptoms of the infected plants are wilting, stunting and yellowing of the leaves, bending of the leaves downwards showing leaf epinasty, growth of adventitious roots in the stems, and the development of narrow dark stripes in the infected vascular bundles beneath the epidermis observed. This results from multiplication of the bacteria in the vascular system to levels that clog the vessels (Boucher *et al.*, 2001) [7]. The internal symptoms which are a direct reaction to infection include progressive discoloration of the vascular tissue, mainly the xylem, at early stages of infection, and of portions of the pith and cortex, as the disease progresses, until complete necrosis occurs. Slimy viscous ooze typically appears on the vascular bundles of stems that have been cut across; as a result, collapse and death of the plant take place because of the degradation of occluded xylem vessels and the destruction of surrounding tissues (Boucher *et al.*, 2001) [7]. On the above ground ginger plant part which started with slight yellowing and wilting of the lower leaves. The wilt progresses upward, affecting the younger leaves, followed by a complete yellowing and browning of the entire pseudo-stem. Under conditions favorable for disease development, the entire shoot becomes flaccid and wilts with little or no visible yellowing. However, the plant dries very rapidly and the foliage becomes yellow-brown in 5 to 10 days. (Habetewold *et al.*, 2015) [18].



Source: (Habetewold *et al.* 2015) [18]

Fig 1: Symptom of ginger bacterial wilt on the above ground parts. (a) Initial, (b) wilted tiller, (c) diseased shoots that are broken down; (d) Water soaked spots on leaves.

The underground symptom of ginger bacterial wilt include: Discoloration of rhizomes that are often rotted inside, Rhizomes and stem vasculature with a water-soaked appearance and

Discoloration of vascular tissues (Figure 2). In addition, Bacterial ooze from an infected ginger rhizome was also reported (Nelson, 2013) [29].



Source: Nelson, 2013 [29]

Fig 2: Ginger bacterial wilt sign and symptom on underground parts of ginger

Pathogen Survival, Spread and Life Cycle

The sources of infections of *Ralstonia solanacearum* are contaminated soil, in plant materials, on farm equipment, and in irrigation or surface water. The bacterium survives in the soil for varying periods of time, and is able to persist between successive crops. The bacterium has been found to survive in sheltered sites such as plant debris and latently infected potato tubers, the deeper soil layers and in the rhizosphere of roots of weed hosts (Hayward, 1991) [20]. The survival of the *Ralstonia solanacearum*, race 4 strains was reported in plant-free soil and potting medium in the presence of plants inoculated by different methods (non-wounded, rhizome-wounded, and stem-wounded) and irrigated on different schedules (alternate and daily). *Ralstonia solanacearum* can survive in soil for a long time and invades suitable plant-host when come in contact with. It colonizes in the xylem and spread into the host and causes wilt. Thereafter it returns back to soil and resides there as saprophytes (Kumar, R., 2014) [25].

Host Range

Ralstonia solanacearum is bacterialwilt causal agent of many plant species. It has broad host range in over 50 plant family with more than 400 reported host plants. Infects potatoes (*Solanum tuberosum*), eggplant (*Solanum melongena*) peppers (*Capsicum annum*), tomatoes (*Lycopersicon esculentum*), geraniums, geranium carolinianum, ginger (*Zingiber officinale*), and few weed species including bitter sweet (*Celastru orbiculatus*), night shade (*Solanum karsense*) and stingingnettle (*Urtica dioica*) (Tahat and Sijam, 2010) [28]. Since the pathogen is able to survive in the soil over a long time; it can exist in a very wide range of weeds and volunteer crops (Fajinmi A. and Fajinmi O., 2010) [25]. Bottle gourd (*Lagenaria siceraria*), one of the earliest cultivated plants in China wasdetected as host for *Ralstonia solanacearum* in 2005 (Gao *et al.*, 2007) [16].

Distribution and importance of Ginger Bacterial wilts

The first Ginger bacterial wilt diseases were reported from India in 1941 by Thomas, then after a lots of reports came from Australia (Hayward *et al.*, 1967) [13], China (Li *et al.*, 1994), Hawaii (Rosenberg, 1962), Indonesia (Sitepu *et al.*, 1977), South

Korea (Choi and Han, 1990), Malaysia (Lum, 1973), Mauritius (Orian, 1953), Nigeria (Nnodu and Emehute, 1988), Philippines (Zehr, 1969) and Japan (Morita *et al.* 1996) cited by (Habetewold k. *et al.* 2015) [18]. In many tropical and subtropical regions the pathogen has been widely distributed and associated with a wide range of hosts (Agrios, 2005) [1]. *Ralstonia solanacearum*, a widely distributed and economically important plant pathogen, invades the roots of diverse plant hosts from the soil and aggressively colonizes the xylem vessels, causing a lethal wilting known as bacterial wilt disease. The bacterium, which is often endemic in the soil, penetrates the plant through the root system and eventually causes irreversible wilting and death. The disease has caused serious upsets to farmers who have been venturing into greenhouse production of crops that are susceptible to the bacterium (Kinyua *et al.*, 2014) [24]. Bacterial wilt is a significant disease of many crops besides ginger root. These wilt diseases are caused by several subgroups of a bacterial pathogen, *Ralstonia solanacearum*. Strains of *R. solanacearum* have been classified into 5 races based on their host ranges, and into 5 biovars based on their differential ability to produce acid from a set of specific carbohydrates. Those races principally attack Bananas (race 1), ornamental plants (race 2), and potato (race 3), Ginger (race 4) and mulberries race 5. *Ralstonia solanacearum* race 4 infected ginger crop in many ginger growing countries (Nelson, 2013) [29]. In Ethiopia ginger bacterial wilt was first reported in 2016. Survey was conducted in 11 zones of Southern parts of Ethiopia. It was 100% prevalent and severely infected commercial farms and small scale farms from 11 surveyed zones. Samples collected during the survey were subjected to laboratory and green house analysis. It is one of the most devastating plant diseases of economically important crops mainly Solanaceousfamily such as tomato, potato, pepper and eggplant. These crops play a significant role primarily as sources of income and food security for the small scale farming community in Ethiopia (Kurabachew H., & Ayana G., 2016) [19]. Study showed that Ginger bacterial wilt is serious threat to ginger production in southern parts of Ethiopia which reduced the yield up to 98%. Survey was done in 2012 in major ginger producing areas of Ethiopia. Accordingly, a total of 165 producing farmer's field which found in twenty seven woredas, nine zones and two special woreda in south

Nation and Nationality of People Regional State (SNNPRS) and one zone and one woreda in Gambela regions were assessed. The result confirmed that Bacterial wilt of ginger was found wide spread in all area and the incidence varied from 10.7% in Gamogofa zone and 93% in Sheka zone (Habetewold *et al.* 2015)^[18]. The disease that threaten ginger production in Ethiopia was caused by *Ralstonia solanacearum* biovar III race 4 and it were distributed in major ginger growing areas of the region (Bekelle *et al.* 2016)^[5]. Ginger bacterial wilt was reported to be among newly emerging diseases in Western parts of Ethiopia that severely affected ginger production in the area (Duressa, 2018)^[11]. The disease is one of destructive diseases that posed threat to the production of ginger in different countries. The disease was reported across different ginger producing countries with variable range of incidence level (Table 1)

Table 1: Distribution and range disease incidence of ginger bacterial wilt in different countries.

S/N	Country	Range of Incidence (%)	Reference
1	Ethiopia	10.7- 93.5	Habetewold K. et al. 2015
2	Australia	*NA	Hayward , A. C. & Pegg, K. G., 2012
3	Ethiopia	80-100	(Hunduma,T. et al., 2016)
4	India	13.2-70.1	(Sharma, et al., 2016)
5	Ethiopia	74.4 – 99	Bekelle K., et al. 2016
6	China	43 – 79	Jian Z., et al., 2018
7	Ethiopia	45.8 – 80.44	Merga et al. 2018

*NA= not applicable

Management Approaches of Ginger Bacterial Wilt

Management of any crop disease starts with correct identification of the causal agent. In some circumstances, ginger wilt can be caused by combination two organisms (bacteria and nematodes) that act synergistically. DAS, S., (2004)^[9] noticed that, quick wilt of ginger caused can be caused by *Meloidogyne incognita* and *Ralstonia solanacearum*. Simultaneous inoculation of *M. incognita* @ 1000 J2/kg soil + *R. solanacearum* @ 4 × yield including reduction in infectivity as well as population of *M. incognita*. Presence of *M. incognita* in simultaneous inoculation and *M. incognita* preceding the inoculation of *Ralstonia solanacearum* by 10 days reduced the incubation period of wilting by 5 and 10 days respectively there by facilitating successful infection. For this reason, correct identification of disease causal agent determines which control method should be applied. Different management options (cultural, host resistance, biological, chemical and integrated disease management) have been used for control of bacterial wilt disease caused by *Ralstonia solanacearum*.

Cultural Method

Cultural methods used to control and eradicate bacterial wilt are, use of healthy seed and planting in clean soils. However, many additional factors influence the incidence of the bacterium, such as environmental conditions (temperature and soil moisture), rotation with non-host plants, the use of less susceptible varieties and cultural practices (crop sanitation and nematode control) (Choudhary *et al.*, 2018)^[8, 12]. Crop residue management is known to be essential in controlling soil borne pathogens. Carbon released from crop residues contributes to increasing soil microbial activity and so increases the likelihood of competition effects in the soil (Bailey and Lazarovits, 2003)^[3]. The placement of the residue in soil can lead to the displacement of the pathogen

from its preferred niche diminishing the pathogen's ability to survive. The benefits of applying organic amendments for disease control are incremental, generally slower acting than chemical. But, last longer, and their effects can be cumulative (Bailey and Lazarovits, 2003)^[3]. Soil amendment with silicon fertilizer and sugarcane bagasse reduced population of *Ralstonia solanacearum*, mean wilt incidence, percent severity index, and corresponding areas of disease incidence, and severity progress curves in the moderately resistant tomato cultivar. Population of *Ralstonia solanacearum* was significantly reduced ($p < 0.05$) in silicon fertilizer treated plants both at 5 and 12 days post inoculation compared to non-amended for moderately resistant cultivar with an average of 29.2 and 17.6% reduction, respectively. Similarly, in sugarcane bagasse treatment, the bacterial population was also significantly reduced compared to the control treatment at 5 days post inoculation but not at 12 days post inoculation in moderately resistant cultivar. However, no significant reduction was obtained in bacterial populations in the moderately susceptible cultivar for all treatments (Ayana G. *et al.*, 2011)^[2]. Biofumigation with cabbage significantly reduced percentage of wilting and increased the yield of Ginger. In this study, six treatments i.e Soil treatment by biofumigation using cabbage, Soil treatment using bleaching powder at 10g/bed, Rhizome treatment by heat, Rhizome treatment by rhizobacterial antagonist, Rhizome treatment by endophytic bacterial antagonist and Control were used to manage bacterial wilt and soft rot disease of ginger. Biofumigation treatment of the soil with cabbage resulted in lowest soft rot and bacterial wilt disease incidence of 9.97% and 5.92% respectively with a highest yield of 15.16 t/ha (Bandyopadhyaya, *et al.*, 2016)^[4].

Host Resistance

Host resistance is one of the safest and cheapest ways of disease management. Developing cultivars that are resistant to bacterial wilt is the most economical, environment friendly and effective method of disease management. Breeding for resistance to bacterial wilt has been concentrated widely on important economic crops, such as eggplant, peanut, pepper, potato, tobacco and tomato (Boshou, 2005)^[6]. It was reported that out of 285 tomato accessions tested in Korea for Resistance to *Ralstonia solanacearum* in at Seedling Stage four germplasms were found to be resistant to pathogen. Getachew *et al.* (2009)^[19] evaluated resistance of tomato genotypes against the highly aggressive *Ralstonia solanacearum* strain originated from Ethiopia. He found that six resistant, eleven moderately resistant, whereas most genotypes, including all tomato cultivars commonly grown in Ethiopia, were found highly susceptible.

Biological Method

The use of chemical compound is toxic to environment. Because, in addition to pest we aimed to control, it can also kill non target organisms including: natural enemies, birds, plant, wild animals, marine life, humans and soil biota. Disappearance of natural enemies of in turn gives chance for the pest to flourish in the absence of natural enemies and become supper pest. Killings of microorganisms in polluted soil can also negatively impact in crop production as soil fertility can be reduced in the absence of saprophytic and symbiotic microorganism's involved nutrient fixation and supplies. So, the use of biological controls products for soilborne pathogen has gained popularity in recent years due

to environmental concerns raised on the use of chemical products in disease management (Haas and De'fago, 2005) [17]. Furthermore, it was also reported that, *Ralstonia solanacearum* was inhibited as a result of increased phenols induced locally or systemically by an arbuscular mycorrhizal fungus. In pot cultures, *R. solanacearum* populations in the rhizosphere, on root surfaces and in the xylem were decreased by 26.7, 79.3 and 81.7%, respectively, following inoculation of tomato plants (*Lycopersicon esculentum* Mill.) with *Glomus versiforme* Berch. Colonization of the plants by both *Ralstonia solanacearum* and *G. versiforme* increased the contents of soluble phenols and cell-wall bound phenols in root tissue, but with different patterns (Zhu and Yao, 2004) [38].

Chemical Method

Singh and Jagtap (2017) [33] reported *in vitro* growth inhibition capacity of chemicals with varying inhibition ability against *Ralstonia solanacearum*. Average inhibition ranged from 6.2 mm (Copper hydroxide) to 20.05 mm (Streptocycline). However; significantly highest average inhibition was recorded in the antibiotic Streptocycline (20.05 mm). Similarly, (Raghu Ram, 2011) evaluated different antibacterial chemicals evaluated against the *Ralstonia solanacearum* under *in vitro*. The results indicated that streptocycline with an inhibition zone of 25.90 mm (2.59 cm) exhibited superior efficacy at 500ppm concentration. Followed by Kcyclyne with an inhibition zone of 20.50 (2.05 cm). Other chemicals like copper oxychloride, plant guard, copper hydroxide and plantomycine were moderately effective whereas bromophol was least effective.

Integrated Disease Management Approach

Integrated disease management is a combination of methods, such as cultural, host resistance, biological, and chemical application, that are environmentally compatible, economically feasible, and socially acceptable to reduce damage caused by diseases to tolerable levels. The main goals of integrated disease management are eliminate or reduce the initial inoculum, reduce the effectiveness of initial inocula, increase the resistance of the host, delay the onset of disease and slow secondary cycles (Agrios, 2005) [1]. Different studies revealed that the use of two or more management options have better achieved control of *Ralstonia solanacearum* than sticking to single management method. According to (Yang, L, *et al.*, 2012) [37] integration of soil amendment with organic composts and biological control agents increased the efficacy of biological control agents to control the population of *Ralstonia solanacearum* that cause ginger wilt disease in China. Two organic composts (maize powder and soybean residue) were combined with one combination (the aqueous solution of *Bacillus subtilis* and *Bacillus cereus*) and two single (*Bacillus subtilis* and *Bacillus megaterium*) biological control agents which together showed significant biocontrol efficacies. The study revealed that 3 - 30% biocontrol efficacy improvement was obtained by addition of the two organic composts compared to the biological control agent treatment. The best biocontrol activity (73.7%) was obtained from the combination of *Bacillus megaterium* and maize powder. Merga *et al.* (2018) [27] reported that the integration of different cultural management options including hot water treatment, soil solarization, soil amendment with potassium fertilizer (KCL) and bio fumigation with lemon grass reduced percent disease

incidence and disease progress on research conducted at two locations; Jimma and Teppi zones of Ethiopia. Although integration of these management options was found to be promising, their efficacy varied between the two locations. The higher efficacy (42.1%) was obtained at Jimma and the lower (38.3%) was obtained at Teppi. But, there is no evidence why efficacy of integration of management options varied across locations.

Summary and Conclusion

Ginger (*Zingiber officinale* Rosc.) is an important commercial crop grown for its aromatic rhizomes. Its production is affected by many diseases and insect pests. Bacterial wilt disease of ginger, which is one of the major constrains of ginger production in tropical and subtropical regions of world is highly heterogeneous bacterial pathogen of many economically important crops causes severe wilting. In Ethiopia ginger bacterial wilt was first reported in 2016 and since then it is affecting ginger production in ginger producing areas of southern and western parts of the country. The high incidence of plant mortality makes *Ralstonia solanacearum* as one of the world's most destructive plant pathogens. The disease that threatens ginger production in Ethiopia was caused by *Ralstonia solanacearum* biovar III race 4 and the diseases were distributed in major ginger growing areas. Management of any crop disease starts with correct identification of the causal agent. In some circumstances, ginger wilt can be caused by combination two organisms (bacteria and nematodes) that act synergistically. Efforts have been made to manage the disease in many countries including Ethiopia. The available management options for the disease are using biological control agents, cultural methods, chemical methods and integrated disease management approaches. Since pathogen has ability to colonize soil habitat, it can cause disease up on cultivation of susceptible host. For this reason it is important to use crop rotation farming system with non-host crops like cereals so that its population can be reduced. This helps to starve *Ralstonia solanacearum* in the absence of right host. The use of biological control agents is one of the most preferable disease management options because it is environmentally safe, economically feasible and sustainable. Once biological control agents are established in certain habitat, frequent application is not needed unlike that of chemicals because, they can perpetuate by their own provided that the practices that can harm them like chemical application is avoided.

Future Prospects

The use of biological control agents for management of ginger bacterial wilt is very limited in Ethiopia. Therefore, research should be directed towards identification and efficacy test *in vitro*, green house and field condition to explore best candidate biological control agents against *Ralstonia solanacearum*. Even though the pathogen is reported as a challenge of ginger production in Ethiopia, distribution of the disease is not clearly known at national level and yield loss assessment data are scarce. Since the pathogen is soil and seed borne, attention should be given to internal and external quarantine system to prevent introduction of the disease which makes the situation very worse. Furthermore, Information is scarce regarding efficacy evaluation of integrated disease management approaches. For this reason, focus should be given to evaluate efficacy of different

management options and integrating them in a way that they have better performance when used together than when used solely. Moreover, more of works done so far did not considered for the presence or absence of nematode species (*Meloidogyne incognita*) that together with *Ralstonia solanacearum* can hasten wilting in ginger. So that, attention should be given to detect both pathogens to set appropriate disease management strategy. Due to the complexity of the disease, the management of bacterial wilt is very difficult once infection is occurred. Therefore, application of Tissue Culture Technology should be enhanced to produce disease free planting materials

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