



# Isolation and Identification of Fungi Associated with Fruit Rot Disease of Tomato (*Solanum lycopersicum* L.) in the Southern Guinea Savannah, Nigeria

Gwa, V. I. <sup>a</sup> and Lum, A. F. <sup>b\*</sup>

<sup>a</sup> Department of Crop Protection, Faculty of Agriculture, Federal University Dutsin-Ma, PMB 5001, Katsina State, Nigeria.

<sup>b</sup> C/o Byrd Research Center, ICT-U, Louisiana, USA.

## Authors' contributions

This work was carried out in collaboration between both authors. Author GVI designed the study, analysed the data and prepared the first draft of the manuscript. Author LAF contributed to the draft manuscript. Both authors read and approved the final manuscript.

## Article Information

DOI: 10.9734/IJPR/2023/v12i6257

## Open Peer Review History:

This journal follows the Advanced Open Peer Review policy. Identity of the Reviewers, Editor(s) and additional Reviewers, peer review comments, different versions of the manuscript, comments of the editors, etc are available here: <https://www.sdiarticle5.com/review-history/109818>

Original Research Article

Received: 21/09/2023

Accepted: 25/11/2023

Published: 01/12/2023

## ABSTRACT

Rot disease is a major threat to tomato fruit production and postharvest handling in major tomato growing areas in Nigeria. Rotted tomato fruits were randomly collected from farmers' farms for the purpose of isolation and identification of rot causing fungi in Tarka, Benue State, Nigeria between July and December, in 2015 and 2016. Results revealed that *Aspergillus flavus*, *A. niger*, *Alternaria solani*, *Phytophthora infestans*, *Oidium neolycopersici*, *Sclerotium rolfsii*, *Fusarium oxysporum*, *Verticillium* spp and *Colletotrichum* spp caused tomato fruit rot in the area. In both years, the lowest number of fungi was recorded in the months of July and December while the highest number was obtained in September. The mean number of fungi isolated showed that *S. rolfsii* was the least with 3.17 in 2015 and 1.66 in 2016 while *F. oxysporum* was the highest with 14.33 in 2015 and 9.33 in 2016.

\*Corresponding author: Email: lumfontem@yahoo.com;

2016. There were more fungi isolated in 2015 than in 2016. There were significant differences ( $P \leq 0.05$ ) in the mean number of *A. niger*, *P. infestans*, *O. neolycopersici* and *F. oxysporum* isolated in both years. In conclusion, farmers should handle tomato fruits with care, to reduce postharvest injury and damage in order to ensure the availability of disease free fruits and to increase the output for the global population.

**Keywords:** Disease; *Fusarium oxysporum*; isolation; rot; tomato.

## 1. INTRODUCTION

Tomato (*Lycopersicon esculentum* L.) is an annual fruit vegetable crop in the family Solanaceae [1]. The crop is cultivated as vegetable in the world; in Africa, Nigeria is the second largest producer after Egypt [2]. The fruit is eaten raw, cooked ripe or unripe, and put in several dishes, sauces and drinks; it is also dried and ground into different products for consumption [3].

Tomato fruit provides minerals, vitamins A and C, proteins, carbohydrates, fats, dietary fibre and potassium [4] and it is widely consumed in Nigeria. It is the second most valuable vegetable fruit crop in the Nigerian Savanna, and its average consumption is about 18% daily [5]. It is a highly perishable crop and losses account for as high as about 50% between the points of production and consumption [6]. The fruit is highly perishable due to high moisture content and susceptibility to plant pathogens both in the field and during postharvest handling. The post harvest losses in fresh tomato account for about 25.80% and these losses vary from time to time, season to season and even from one Region to another depending on the interaction of a susceptible host, a virulent pathogen and favourable environmental conditions [7,8]. In all parts of the world, bacteria and fungi have been found to be the most frequently occurring rot causing pathogens of tomato fruits [9]. The most commonly isolated and identified fruit rot fungi pathogens are *Fusarium oxysporum*, *Alternaria solani* and *Aspergillus niger* [10,11,12]. Other pathogens equally responsible for fruit rot disease in tomato include the following fungi, *A. flavus*, *Alternaria alternata*, *Botrytis cinerea*, *Curvularia* spp, *Fusarium moniliforme*, *Geotrichum* spp, *Mucor* spp, *Penicillium* spp, *Phytophthora* spp and *Rhizopus stolonifer*; and bacteria such as *Erwinia* spp [13,5,14]. The aim of this study was to isolate and identify the fungi which cause field rot disease of tomato fruits in Tarka Local Government Area, Benue State, Nigeria and also to determine the susceptibility of the fruits to the isolated pathogens.

## 2. MATERIALS AND METHODS

### 2.1 Study Area

The isolation and identification of pathogens which cause fruit rot disease of tomato were carried out at the Plant Pathology Laboratory, Department of Crop and Environmental Protection, Federal University of Agriculture, Makurdi, Benue State in 2015 and 2016.

### 2.2 Collection of Infected Tomato Samples

Infected fruits with different degrees of rot symptoms were randomly collected from tomato farms at Tarka Local Government Area and packaged as reported by Sani and Gwa [11].

### 2.3 Sterilization of Samples

Glass wares were washed in running tap water and sterilized in an oven at 120°C for 30 minutes. Infected samples were washed with tap water, cut with a sharp sterilized blade into small pieces approximately 2x2 mm<sup>2</sup> in diameter at the interphase of healthy and infected tissues as reported by Lum and Takor [15]. Samples were further sterilized in 5% Sodium hypochlorite solution for about 20 seconds [11]. The small pieces were rinsed in four successive changes of sterile water in order to remove the chlorox chemical on them and were blotted on sterile paper for about 10 minutes before inoculation of the tissues.

### 2.4 Inoculation of Plant Tissues

Sterile Potato Dextrose Agar (PDA) (20 ml) was poured in 90 mm Petri dishes and left to solidify before inoculation. The medium was amended with 0.16 g of powdered streptomycin sulphate to suppress bacteria growth. Four pieces of the infected tissues were aseptically transferred to the Petri dishes containing the solidified PDA and incubated at 30±5° C for five days. Plates were regularly monitored and growth colonies were examined to determine the frequency of occurrence of each of the pathogens identified.

## 2.5 Determination of Frequency

The frequency of occurrence of the isolated pathogens was calculated by counting the number of times each isolate occurred out of the total number of isolates. Colonies that grew on the plates were examined and aseptically sub-cultured in order to obtain pure cultures of the various isolates.

## 2.6 Identification of Pathogens

Identification of the pure cultures was done by preparing slides of fungal isolates from the different pure cultures and examining them with the aid of a compound microscope. Morphological and cultural characteristics of the isolated fungi were compared with identification guides to establish the identity of the isolated fungi [16].

## 2.7 Pathogenicity Test of Isolated Fungi

Pathogenicity tests were carried out using the method of Sani and Gwa [11] to ascertain the ability of the various isolates to cause rotting on apparently healthy looking tomato fruits free of disease symptoms and physical injuries. The fruits were washed in three successive changes of running tap water and 5% sodium hypochlorite solution was used to sterilize them for about 2 minutes to remove surface contaminants. Cylindrical discs about 5 mm in diameter were made on the ripe and healthy looking tomato fruits using a sterilized cork borer. Mycelial discs (4 mm in diameter) from five days old cultures of *Aspergillus flavus*, *A. niger*, *Alternaria solani*, *Phytophthora infestans*, *Oidium neolycopecisci*, *Sclerotium rolfsii*, *Fusarium oxysporum*, *Verticillium* spp and *Colletotrichum* spp were aseptically obtained and inserted separately into

the holes created. The holes were covered with Petroleum jelly to prevent contamination by other microbes. Tomato fruits treated with sterile PDA instead of the inocula of the various isolates served as the control. The tested isolates and the control were replicated three times. A total of 33 ripe healthy tomato fruits were used for the pathogenicity test. The treatments were completely randomized and the inoculated tomato fruits were incubated at ambient room temperature ( $30\pm 5^{\circ}\text{C}$ ) under sterile conditions for 5 days for growth of the fungi. Symptoms of rots obtained from fruits artificially inoculated with the fungal isolates were compared with those already observed when infected on the field. The artificially infected fruits were re-isolated aseptically, and cultured on PDA plates; proper morphological and cultural examination and comparisons were made with those infected on the field [17].

## 2.8 Statistical Analysis

Data collected were analyzed by one-way analysis of variance (ANOVA) according to Gomez and Gomez [18]. The two-tailed paired Student's t-test was used for comparing the mean frequency of occurrence of fungal isolates for the two years at 5% level of probability.

## 3. RESULTS

In this study, a total of nine different fungi namely *Aspergillus flavus*, *A. niger*, *Alternaria solani*, *Phytophthora infestans*, *Oidium neolycopecisci*, *Sclerotium rolfsii*, *Fusarium oxysporum*, *Verticillium* spp and *Colletotrichum* spp were isolated as pathogens causing tomato fruit rot disease on the field between July and December, in 2015 and 2016 (Fig. 1a–e).

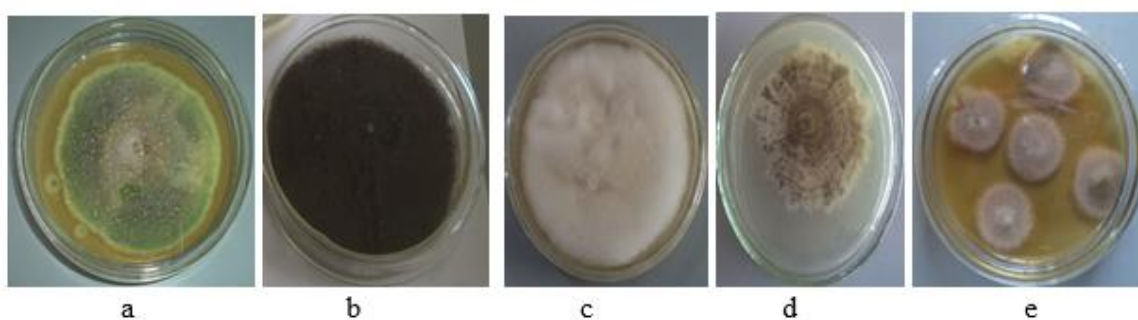


Fig. 1. Colonies of some Fungi on Potato Dextrose Agar (a) *Aspergillus flavus*; (b) *A. niger*; (c) *Fusarium oxysporum*; (d) *Colletotrichum* spp; (e) *Alternaria solani*

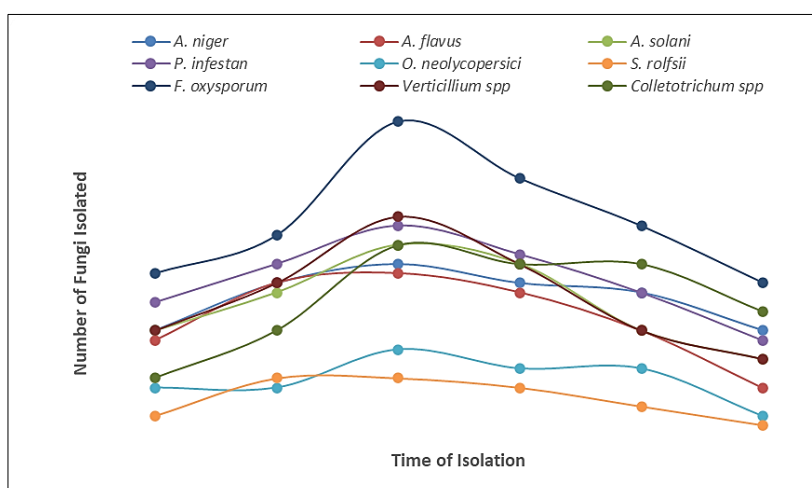


Fig. 2. Number of fungi isolated on tomato fruits from July to December, 2015

Table 1. Mean variation of fungi isolated on tomato fruits in 2015 and 2016 cropping seasons

Fungi Isolated	Year		Df	T-Value	P-Value
	2015	2016			
<i>Aspergillus niger</i>	8.83±0.65	6.67±0.56	9	2.52	0.03*
<i>Aspergillus flavus</i>	7.17±0.79	6.17±1.10	9	0.75	0.47
<i>Alternaria solani</i>	8.83±0.95	6.00±0.97	9	2.10	0.06
<i>Phytophthora infestans</i>	10.00±0.86	7.33±0.88	9	2.17	0.05*
<i>Oidium neolycoopersici</i>	4.33±0.56	2.50±0.43	9	2.61	0.02*
<i>Sclerotium rolfsii</i>	3.17±0.65	1.66±0.33	7	2.04	0.08
<i>Fusarium oxysporum</i>	14.33±1.70	9.33±0.95	7	2.60	0.03*
<i>Verticillium spp</i>	9.00±1.20	6.50±1.10	9	1.54	0.15
<i>Colletotrichum spp</i>	8.50±1.10	6.83±1.20	9	1.06	0.31

\*indicates statistical significance at 95% CL

Table2. Pathogenicity test of fungal isolates in artificially inoculated fruits of UC 82B variety of tomato

Isolates	Rotting on tomato fruits		
	Days to rotting	Inoculated fruits	Uninoculated fruits (Control)
<i>Aspergillus niger</i>	3	++	-
<i>Alternaria solani</i>	4	++	-
<i>Aspergillus flavus</i>	3	++	-
<i>Verticillium spp</i>	4	++	-
<i>Oidium neolycoopersici</i>	4	+	-
<i>Phytophthora infestans</i>	4	++	-
<i>Colletotrichum spp</i>	4	++	-
<i>Fusarium oxysporum</i>	2	+++	-
<i>Sclerotium rolfsii,</i>	5	+	-

+ = slight rotting; ++ = moderate rotting; +++ = severe rotting; - = no rotting

The number of fungi isolated showed that *F. oxysporum* occurred most frequently while *S. rolfsii* was consistently the least from July to December, 2015 (Fig. 2). The results further indicated that for each fungus, the least number was obtained in July and December while the peak number was recorded in September, 2015.

The mean number of rot causing fungi isolated on tomato fruits from July to December in the 2015 and 2016 cropping seasons showed that *S. rolfsii* was the least while *F. oxysporum* was the highest (Table 1). The number of fungi isolated was more in the 2015 cropping season than in 2016. The mean number of *A. niger*, *P. infestans*,

*O. neolycopersici* and *F. oxysporum* differed significantly ( $P \leq 0.05$ ) in both years. The rest of the fungi did not vary significantly in both years.

Results of the pathogenicity test indicated that all the nine isolates were able to elicit rotting in the tomato fruits (Table 2). The results however, revealed that *F. oxysporum* was more aggressive than the other pathogens and caused severe rotting of the fruits. The results further showed that *Oidium neolycopersici* and *S. rolfsii* caused slight rotting of the fruits while the rest of the isolates produced moderate rotting. There was no rotting observed in the control treatment (tomato fruits uninoculated with the pathogenic isolates).

#### 4. DISCUSSION

The results of this study showed that *Aspergillus niger*, *A. flavus*, *Alternaria solani*, *Phytophthora infestans*, *Oidium neolycopersici*, *Sclerotium rolfsii*, *Fusarium oxysporum*, *Verticillium* spp and *Colletotrichum* spp are pathogens which cause tomato fruit rot in Tarka, Benue State. Similar results were obtained by Ibrahim et al. [19], Matthew [20], Laila et al. [21], Yusuf et al. [5], Nizamani et al. [14] who indicated that fungi are major rot causing pathogens of tomato fruits. In another study, Sani and Gwa [11] isolated *A. flavus*, *A. niger*, *F. oxysporum*, *F. moniliforme* and *Rhizoctonia solani* from rotted tomato fruits in Dutsin-Ma, Nigeria and found them pathogenic to healthy ones. Similarly, Onuorah and Orji [13] isolated *A. niger*, *Rhizopus stolonifer*, *F. oxysporum*, *Saccharomyces cerevisiae*, *Alternaria alternata*, *Penicillium digitatum* and *Geotrichum candidum* from rotted tomato fruits in Awka, Nigeria and found *A. niger* with the highest percentage of rots. Yusuf et al. [5] isolated *Aspergillus* spp, *Fusarium* spp, *Penicillium* spp and *Rhizopus* spp from infected tomato in Anyigba, Kogi State and found *Aspergillus* spp with the highest frequency (38.89%) and *Fusarium* spp with the least (5.56%), contrary to these results which indicated that *F. oxysporum* was the most frequently isolated fungus in both cropping seasons (mean number=14.33 in 2015 and 9.33 in 2016) while *Sclerotium rolfsii* was the least (mean=3.17 in 2017; 1.60 in 2016).

Similarly, Abdulkadir et al. [12] isolated *F. oxysporum* from tomato fruits in Makurdi, and found the fungus responsible for *Fusarium* wilt disease of tomato. In another study carried out in Ethiopia, Lemma et al. [22] isolated *Alternaria* spp, *Fusarium* spp, *Rhizopus* spp, *Penicillium* spp and *Erwinia carotovora* from infected tomato

samples. In related studies, Mugao and Birgen [23] isolated *Erwinia* spp, *Botrytis* spp, *Alternaria* spp, *Geotrichum* spp and *Rhizopus* spp from infected tomato fruits in Mwea, Kenya and found healthy tomato fruits susceptible to them. Nizamani et al. [14] isolated *Erwinia* spp, *Botrytis* spp, *Alternaria* spp, *Geotrichum* spp and *Rhizopus* spp from rotted tomato fruits and observed that *A. solani* was the main cause of post-harvest tomato fruit rot in Tandojam, Pakistan, contrary to the results obtained in this study. Similar findings were reported by Sajad et al. [8] and Cristina et al. [24] that the major pathogens which caused tomato fruit rots were *Alternaria* sp., *Fusarium* sp., *G. candidum* and *R. stolonifer*.

This study revealed that fungal infection was more in September of both years than in any other month. This is probably due to favourable environmental conditions such as high rainfall, relative humidity and soil moisture which enhanced spore formation and dissemination of pathogens at the expense of the tomato plants which were susceptible.

#### 5. CONCLUSION

This study showed that tomato fruits were susceptible to fungal pathogens such as *Aspergillus niger*, *A. flavus*, *Alternaria solani*, *Phytophthora infestans*, *Oidium neolycopersici*, *Sclerotium rolfsii*, *Fusarium oxysporum*, *Verticillium* spp and *Colletotrichum* spp. *Sclerotium rolfsii* had the least frequency while the most devastating pathogen was *F. oxysporum*. The results further revealed that in both years, the number of fungi isolated was least in July and December and highest in September. It is therefore, recommended that appropriate measures be taken during the growing, harvesting and postharvest handling of tomato to mitigate diseases and increase the output for sustainable crop production and to ensure food security for the global teeming populations.

#### COMPETING INTERESTS

Authors have declared that no competing interests exist.

#### REFERENCES

1. Wani AH. An overview of the fungal rot of tomato. *Mycopathology*. 2011;9:33–38.
2. FAO. FAOSTAT statistical database. [Rome]: FAO ; 2020.

- Available:<http://www.fao.org/faostat/en/#data/QC/visualize>. Retrieved on 12/05/2021.
3. Onuorah S, Obika I, Okafo, U. Filamentous fungi associated with the spoilage of post-harvest sweet orange fruits (*Citrus sinensis*) sold in Awka major markets, Nigeria. *Bioengineering and Bioscience*. 2015;3(3):44–49.
  4. Wogu MD, Ofuase O. Microorganisms responsible for the spoilage of tomato fruits, *Lycopersicon esculentum* sold in markets in Benin City, Southern Nigeria. *School of Academics and Journal of Biosciences*. 2014;2(7):459–466.
  5. Yusuf L, Agieni GA, Olorunmowaju Al. Isolation and identification of fungi associated with tomato (*Lycopersicon esculentum* M.) rot. *Sumerianz Journal of Agriculture and Veterinary*. 2020;3(5):54-56.
  6. Mbuk EM, Bassey NE, Udoh ES, Udoh EJ. Factors influencing postharvest loss of tomato in urban market in Uyo, Nigeria. *Nigerian Journal of Agriculture, Food and Environment*. 2011;7:40-46.
  7. Mujib UR, Naushad K, Inayatullah J. Post-harvest losses of tomato crop. *Sarhad Journal of Agriculture*. 2007;23:1279-1284.
  8. Sajad AM, Jamaluddin, Abid HQ. Fungi associated with the spoilage of post harvest tomato fruits in different markets of Jabalpur, Madhya-Pradesh, India. *International Journal of Current Research Review*. 2017;9:12-16.
  9. Obetta SE Nwakonobi TU, Adikwu OA. Microbial effects on selected stored fruits and vegetables under ambient conditions in Makurdi, Benue State, Nigeria. *Research Journal of Applied Sciences, Engineering and Technology*. 2011;3: 393–398.
  10. Nowicki M, Nowakowska M, Niezgodna A, Kozik EU. *Alternaria* black spot of crucifers: Symptoms, importance of disease and perspectives of resistance breeding vegetable crops. *Research Bulletin*. 2012;76:5-19.
  11. Sani S, Gwa VI. Fungicidal effect of *Azadirachta indica* and *Zingiber officinale* extracts in the control of *Fusarium oxysporum* and *Rhizoctonia solani* on tomato (*Solanum lycopersicum*) fruits. *Innovative Techniques in Agriculture*. 2018;2(4):439–448.
  12. Abdulkadir HK, Ekefan EJ, Gwa VI. Antagonistic potential of *Trichoderma harzianum* against *F. oxysporum* f. sp. *lycopersici* isolates causing *Fusarium* wilt disease of tomato (*Solanum lycopersicum* L.). *FUDMA Journal of Agriculture and Agricultural Technology*. 2023;9(1): 143–149.
  13. Onuorah S, Orji MU. Fungi associated with the spoilage of post-harvest tomato fruits sold in major markets in Awka, Nigeria. *Universal Journal of Microbiology Research*. 2015;3(2):11–16.
  14. Nizamani S, Khaskheli AA, Jiskani AM, Khaskheli SA, Khaskheli AJ, Poussio GB, Jamro H, Khaskheli MI. Isolation and identification of the fungi causing tomato fruit rot disease in the vicinity of Tandojam, Sindh. *Agricultural Science Digest*. 2021; 41:186–190.
  15. Lum AF, Takor MC. Taro leaf blight: disease assessment, farmers' knowledge and management potential of goatweed extract in South West Cameroon. *Journal of Agriculture and Crops*. 2021;7(4):159–166.
  16. Agrios G. *Plant Pathology*. 5th ed. Elsevier Academic Press, London; 2005.
  17. Gwa VI, Richard IB. Susceptibility of white yam (*Dioscorea rotundata* Poir) tuber to rot fungi and control with extracts of *Zingiber officinale* Rosc. *Azadirachta indica* A. Juss. and *Piper guineense* Schumach. *Journal of Plant Pathology and Microbiology*. 2018; 9:9.
  18. Gomez KA, Gomez AA. *Statistical procedures for agricultural research*. 2nd ed. John Wiley and Sons; 1984
  19. Ibrahim AD, Musa K, Sani A, Aliero AA, Yusuf BS. Microorganisms associated with the production of volatile compounds in spoilt tomatoes. *Research in Biotechnology*. 2011;2(2):82–89.
  20. Matthew T. Post-harvest microbial deterioration of tomato (*Lycopersicon esculentum*) fruits. *Report and Opinion*. 2011;3(4):52–57.
  21. Laila N, Sajib P, Mahmud MR. Investigation of potential biological control of *Fusarium oxysporum* f. sp. *lycopersici* by plant extracts, antagonistic sp. and chemical elicitors in vitro. *Fungal Genomics and Biology*. 2018;8(1). DOI:10.4172/2165-8056.1000155
  22. Lemma Z, Dawit W, Negari M, Chaka A, Selvaraj T, Gebresenbet G. Identification of post-harvest rotting microorganisms from tomato fruits (*Solanum esculentum* Mill.). *Journal of Stored Products*

- and Postharvest Research. 2014;5(3): 14–19.
23. Mugao LG, Birgen JK. Pathogens associated with tomato post-harvest losses in Mwea, Kenya. *International Journal of Multidisciplinary Research and Growth Evaluation*. 2021;2(2):225–232.
24. Cristina RA, Perez JJ, Guillermo HML, Bernado MA, Omar RE. Control of phytopathogenic microorganisms of post-harvest in tomato (*Lycopersicon esculentum* Mill.) with the use of citrus extract. *Journal of Plant Science and Phytopathology*. 2018;2:37–43.

---

© 2023 Gwa and Lum; This is an Open Access article distributed under the terms of the Creative Commons Attribution License (<http://creativecommons.org/licenses/by/4.0>), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

*Peer-review history:*

*The peer review history for this paper can be accessed here:*

<https://www.sdiarticle5.com/review-history/109818>