

Screening *Solanum* germplasm (Tomato and Brinjal) and identify rootstocks for resistance to Fusarium wilt

Comment [V1]: It will be better if you concise your topic.
Eg. Fusarium Wilt Management in Tomato and Brinjal through Screening Techniques

ABSTRACT:

The present study was carried out in Rabi, 2021 at the PG Research Block, Department of Vegetable Science, College of Horticulture, Sri Konda Laxman Telangana State Horticultural University, Rajendranagar, Hyderabad. In the present study, 25 genotypes of tomato and brinjal were studied for Fusarium wilt resistance at morphological level. Scoring of Fusarium wilt infection severity based on morphological symptoms revealed five groups viz., asymptomatic/no chlorosis in one cultivar (*Solanum torvum*) slight chlorosis of leaves in two genotypes (Arka Keshav and Surya), Moderate chlorosis with wilting or stunting of the plant in ten genotypes (EC-620509, EC-615055, EC-620378, EC-620389, EC-620394, EC-620422, EC-631369, AVTO-9803, EC-620428 and EC-631379), severe chlorosis with wilting and stunting of the plant in four genotypes (EC-620503, LA-1589, Marutham and PKM-1) and plant death was observed in eight genotypes (EC-620441, EC-620452, LA -0490, Money maker, Pusa rohini, Pusa sheethal, including susceptible checks (Pusa Ruby and Arka Vikas).

Comment [V2]: Please specify the objectives

Keywords: Tomato, Brinjal, Fusarium wilt, Genotypes.

Comment [V3]: Please includes conclusion in one sentence

1. INTRODUCTION:

Fusarium oxysporum is major soil borne fungal pathogens of both greenhouse and field grown tomatoes in the warm vegetable growing areas of the world (Rosewich *et al.*, 1999). *Fusarium oxysporum* penetrates the roots mainly through wounds and proceeds into and throughout the vascular system, leading to functional collapse, systemic wilting and often the death of the infected plant. *Fusarium oxysporum* f. sp. *lycopersici* (FOL) causes disease only in plants of the genus *lycopersicon* (Rowe, 1980) and inhabits most tomato growing regions worldwide, causing tomato production yield losses (Staniazsek *et al.*, 2007). This fungus responsible for vascular wilt disease in tomato and infects the vascular system of roots, inhibiting water transport, which in turn results in rapid plant death (Malhotra *et al.*, 1993; McGrath *et al.*, 1987). The first symptom of fusarium wilt in gardens and fields is usually the golden yellowing of a single leaflet or shoot, or a slight wilting and drooping of the lower leaves on a single stem. Yellowed and wilted leaflets drop early. Affected plants turn to bright yellow, wilt, dry up, and usually die before maturity. Three physiologic races of

FOL, named 1, 2 and 3 in order of their discovery (Booth 1971; Grattidge and Obrien,1982) are traditionally distinguished by each having a specific pathogenicity to tomato cultivars.

Comment [V4]: Insert management techniques and focus on screening techniques.

2. MATERIALS AND METHODS:

The experimental material comprised of 15 germplasm lines, 9 released varieties and 1 wild species (Table 1) which were obtained from NBPGR, Regional Station, Hyderabad; IARI, New Delhi; IIHR, Bengaluru; TNAU, Periyakulam; UC DAVIS, California, USA; TNAU, Coimbatore; KAU, Thrissur, and COE, Jeedimetla. All the cultivars were evaluated systematically during the research period. The experiment was laid out with Twenty-five genotypes of tomato and brinjal in Randomized Block Design (RBD) with three replications during Rabi, 2021 at PG Students Research Block of the College of Horticulture, Rajendranagar, Hyderabad. *Fusarium oxysporum* f. sp. *lycopersici* infected sample collected from Noble seeds private limited, Bengaluru. Further sub cultured on PDA media and incubated on BOD incubator at 24°C with 12h light and 12h dark. One week (7 days) old *F. oxysporum* f. sp. *lycopersici* cultures, grown on PDA plates, were flooded with about 10ml of sterile distilled water and the conidia were dislodged with a cell spreader, filtered through cheese cloth, and counted with a haemocytometer. The concentration was adjusted to 10 conidia per ml.

Comment [V5]: What may be the reason to choose these varieties. Please include the reason as well.

Comment [V6]: Write the crops selected, parts of crop, and how you know the it was infected fusarium.

Comment [V7]: Isolation process ?

Comment [V8]: Cite the method

Comment [V9]: Please cite the reference and make clear is this concentration is enough for causing disease.

Comment [V10]: Please make clear is it micro or macro conidia.

Table 1: List of genotypes used for disease screening along with their sources

Comment [V11]:

Comment [V12]: Please specify the crop: is it the variety of tomato or brinjal

S.NO	Name of the Variety/Accession	Source
1	EC-620509	NBPGR, Hyderabad
2	EC-615055	NBPGR, Hyderabad
3	EC-620428	NBPGR, Hyderabad
4	EC-620378	NBPGR, Hyderabad
5	EC-620389	NBPGR, Hyderabad
6	EC-620394	NBPGR, Hyderabad
7	EC-620422	NBPGR, Hyderabad
8	EC-631369	NBPGR, Hyderabad
9	EC-620503	NBPGR, Hyderabad

10	AVTO-9803	NBPGR, Hyderabad
11	EC-631379	NBPGR, Hyderabad
12	EC-620441	NBPGR, Hyderabad
13	EC-620452	NBPGR, Hyderabad
14	LA-1589	UC, DAVIS, California, USA
15	LA -0490	UC, DAVIS, California, USA
16	Money maker	UC, DAVIS, California, USA
17	Marutham	TNAU, Coimbatore
18	Pusa rohini	IARI, New Delhi
19	Pusa sheethal	IARI, New Delhi
20	Pusa ruby	IARI, New Delhi
21	PKM-1	TNAU, Periyakulam,
22	Arka Vikas	IIHR, Bengaluru
23	Arka Keshav	IIHR, Bengaluru
24	Surya	KAU, Thrissur
25	<i>Solanum torvum</i>	COE, Jeedimetla

EC: Exotic collection

For obtaining large amounts of the spore suspension, the fungus was grown on maize (solid substrate), 50g of maize was added to 250ml conical flasks to which 60% of distilled water was added, autoclaved, inoculated with 1ml spore suspension of *F. oxysporum* and incubated for 7 days. Sterile distilled water was added to these cultures and the spore suspension was adjusted to 10 conidia per ml. The spore suspension was used for further pathogenicity assay. Solid substrate served as a better medium for obtaining large amount of spore suspension (Nirmala devi, 2012). Twenty-five genotypes were sown in pro trays filled with cocopeat and watered regularly. Twenty days old healthy seedlings were selected and used for further pathogenicity test. Twenty days old seedlings were inoculated by standard root dip method. Conidia of all the isolates were recovered from one week old cultures. Seedlings were removed from the portrays, shaken to remove the adhering particles and washed carefully under tap water. The roots were trimmed with a sterile scissor and were submerged in the

Comment [V13]: Please cite the reference

Comment [V14]: What is the initial concentration and how you made it final concentration? Make it clear

Comment [V15]: How ?

Comment [V16]: Please specify the media you have used.

Comment [V17]: ?

Comment [V18]: Why you have used this method plz cite the reference?

Comment [V19]: Type of conidia and how many isolates you have used in your experiment? Make it clear.

conidial suspension for 30 minutes. The inoculated seedlings were transplanted to mini polybags, 15cm diameter, surface sterilized with 0.1% mercuric chloride containing soil and sand 1:1 ratio.

Comment [V20]: Why?

Procedure for Screening Fusarium wilt:

- Two kgs of sorghum grains were soaked in water overnight in flasks.
- The excess water which is left after maximum absorption by different grain substrate was drained off and the flasks containing soaked grains were plugged and autoclaved at 15 psi pressure for 30 minutes.
- The substrate in flasks was inoculated with actively growing 5 mm mycelial disc of the pathogen under aseptic conditions and inoculated flasks were incubated at $25\pm 1^{\circ}\text{C}$ in BOD incubator.
- *Fusarium oxysporum f. sp. lycopersici* mass multiplied on sterilize sorghum grains, each @ 5g/kg of soil were added to the pots/poly bags.
- One plant per pot/polybag was maintained.
- The data was recorded at 30 days after inoculation (DAI) and per cent disease incidence was calculated.
- The reaction of each genotype was categorized based on per cent disease incidence.
- The disease reaction was calculated as per the scale suggested by Morid *et al.* (2012).

Comment [V21]: Don't explain in bullet, plz write in paragraph form

Comment [V22]: How you confirmed that the culture is *F. oxysporum f. sp. lycopersici*

Comment [V23]: Make it clear

a. Data

on percentage of fusarium wilt incidence for twenty-five genotypes were recorded and presented as per the disease scale given by Morid *et al.* (2012)

Comment [V24]: I think it is just visual observation and doesnot make clear it may be due to fusarium species, so this disease scale doenot give any scientific evidence.

- 0- No symptoms
- 1- Slight chlorosis, wilting or stunting of the plant.
- 2- Moderate chlorosis, wilting or stunting of the plant.
- 3- Severe chlorosis, wilting or stunting of the plant.

4- Death of the plant.

b. Per cent disease incidence

Per cent disease incidence was calculated by using following formula,

$$\text{Per cent disease incidence} = \frac{\text{Number of plants infected}}{\text{Total number of plants planted}} \times 100$$

Comment [V25]: It would be better to use disease severity for screening rather than disease incidence percentage.

3. RESULTS AND DISCUSSION:

The percent disease incidence in genotypes was ranged from 0% to 100% (Table 2). Lowest per cent of disease incidence (0.00%) was observed in *Solanum torvum*. Highest per cent of disease incidence was found in genotypes viz., Pusa sheethal (100%), EC-620452(100%), LA-0490 (91.66%), EC-620441 (91.66%), Money maker (83.33%), Pusa rohini (83.33%) including susceptible checks (Pusa Ruby 100% and Arka Vikas83.33%). Arka Keshav and Surya recorded 8.33%, 16.66% disease incidence respectively. Genotypes viz., EC-620509, EC-615055, EC-620378, EC-620389, EC-620394, EC-620422, EC-631369, AVTO-9803, EC-620428 and EC-631379 recorded 50.00%, 33.33%, 33.33%, 41.66%, 50.00%, 41.66%, 33.33%, 50.00%, 50.00% and 41.66% disease incidence respectively. EC-620503, LA-1589, Marutham and PKM-1 recorded 58.33% disease incidence respectively.

Out of 25 genotypes, fusarium wilt incidence was reported ranging from 0 to 4 scores according to Morid *et. al* (2012). Among these *Solanum torvum* is highly resistant (HR) to fusarium wilt with “0” score. Arka Keshav and Surya were resistant (R) to fusarium wilt with 0.33 and 0.67 score respectively (Table 3). Genotypes viz., EC-620509, EC-615055, EC-620378, EC-620389, EC-620394, EC-620422, EC-631369, AVTO-9803, EC-620428 and EC-631379 were moderately resistant (MR) to fusarium wilt with scores *i.e.* 2.00, 1.33, 1.33,1.67, 2.00, 1.67, 1.33, 2.00, 2.00 and 1.67 respectively. EC-620503, LA-1589, Marutham and PKM-1 recorded 2.33 score were moderately susceptible (MS) to fusarium wilt. Pusa sheethal, EC-620452, LA -0490, EC-620441, Money maker, Pusa rohini including susceptible checks Pusa Ruby and Arka Vikas were susceptible (S) and highly susceptible (HS) to fusarium wilt recorded score viz., 4.00, 4.00, 3.67, 3.67, 3.33, 3.33, 4.00 and 3.33respectively.

These results were in uniformity with the findings of Chaudhary and sharma (2000), Gousset *et al.* (2005), Mahmoud *et al.* (2006), Ahmadvand *et al.* (2010), Bahattin *et al.*

(2010), Akansha Pandey and Sanjeev Dubey (2017), Antonio *et al.* (2017), Biswas and Ghosh (2018), Latifah *et al.* (2018) and Sushma *et al.* (2022).

Table 2: Scores and percent incidence of fusarium wilt in 25 genotypes

S.NO	Name of the genotypes	Score	Percent incidence (%)
1	EC-620509	2.00	50.00
2	EC-615055	1.33	33.33
3	EC-620428	2.00	50.00
4	EC-620378	1.33	33.33
5	EC-620389	1.67	41.66
6	EC-620394	2.00	50.00
7	EC-620422	1.67	41.66
8	EC-631369	1.33	33.33
9	EC-620503	2.33	58.33
10	AVTO-9803	2.00	50.00
11	EC-631379	1.67	41.66
12	EC-620441	3.67	91.66
13	EC-620452	4.00	100.00
14	LA-1589	2.33	58.33
15	LA -0490	3.67	91.66
16	Money maker	3.33	83.33
17	Marutham	2.33	58.33
18	Pusa rohini	3.33	83.33
19	Pusa sheethal	4.00	100
20	Pusa ruby	4.00	100
21	PKM-1	2.33	58.33
22	Arka Vikas	3.33	83.33
23	Arka Keshav	0.33	8.33
24	Surya	0.67	16.66

25	<i>Solanum torvum</i>	0.00	0.00
		S.Em±	5.75
		SD	28.76

Table 3: Disease reaction of genotypes for fusarium wilt incidence

S. No	Reaction	Score	Numberofgenotypes	Genotypes
1.	Highly resistant (HR)	0	1	<i>Solanum torvum</i>
2.	Resistant (R)	1 (0-1)	2	Arka Keshav, Surya
3.	Moderatelyresistant (MR)	2 (1-2)	10	EC-620509, EC-615055,EC-620378, EC-620389,EC-620394, EC-620422, EC-631369, AVTO-9803, EC-620428, EC-631379
4.	Moderatelysusceptible(MS)	3 (2-3)	4	EC-620503, LA-1589,Marutham, PKM-1
5.	Susceptible (S)and Highly Susceptible (HS)	4 (3-4)	8	EC-620441, EC-620452,LA -0490, Money maker, Pusa rohini, Pusa sheethal,Pusa ruby, Arka Vikas

4. CONCLUSION:

In the present investigation we concluded that *Solanum torvum* is highly resistant (HR) to fusarium wilt with “0” score. Arka Keshav and Surya were resistant (R) to fusarium wilt with 0.33 and 0.67 score respectively.

Comment [V26]: I think conclusion should be overall along with recommendation

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Comment [V27]: Which system for citation has been used and make uniform.

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