Cotton leaf curl virus disease in India- Integrated management strategies and the way ahead

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Cotton leaf curl virus disease (CLCuD)

- •First observed in 1989 on G barbadense in IARI,New Delhi
- •Appeared in patches around Sriganganagar and Punjab adjoining to international border with Pakistan in 1993 and spread to entire north zone(Around 15 lakh ha) in a span of 4-5 years
- •The problem is increasing due to development of frequent recombinants





Occurrence of cotton leaf curl virus disease in northern zone of India













Integrated management strategies

Date of sowing trials/recommendations

 Screening of released BG II hybrids against CLCuD/recommendations

Weed identification/eradication program



Date of sowing trials (2014-15)

Four experiments were conducted at each of the five locations

(Hisar, Sirsa, Sriganganagar, Abohar & Faridkot)

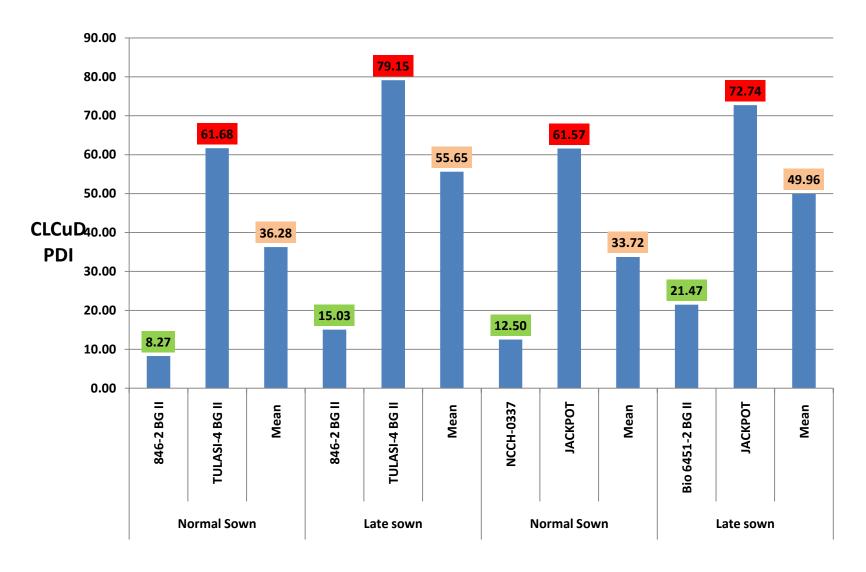
Pre-release hybrids –Normal date of sowing Pre-release hybrids –Late sown Released hybrids –Normal date of sowing Released hybrids –Late sown



Released hybrids	100
Pre-release hybrids	50
Number of rows per entry	2
Row length	5.4 metres (10 dibbles per row)
Number of plants	20 per replication
Number of replications	2
Spacing	67.5 x 60 cm
Experimental Design	Released hybrids:10x10 lattice; Pre release hybrids: RBD
Susceptible check	Standard susceptible check planted after every four lines of the test entry hybrids

- Sowing dates for normal sown crop: 14-24 May, 2014
- Sowing dates for late sown crop: 4-7 June, 2014

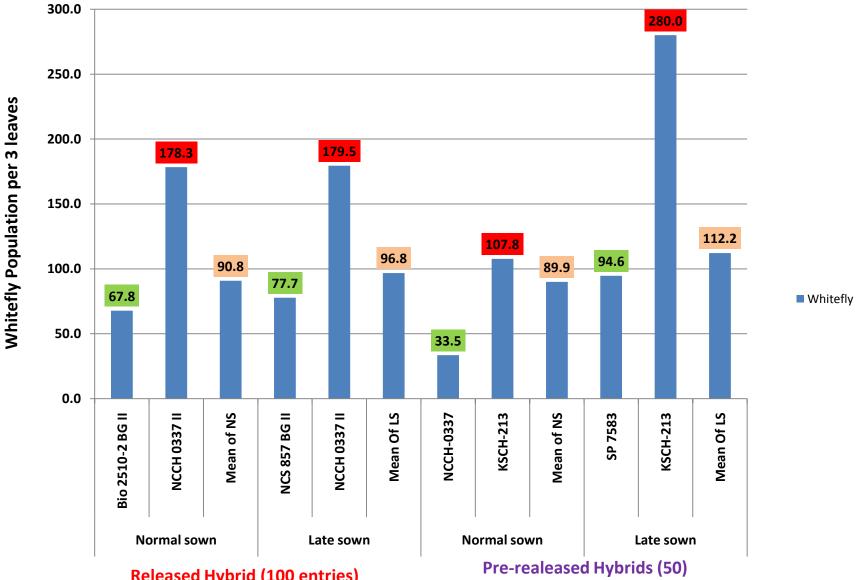




Released Hybrid (100 entries)

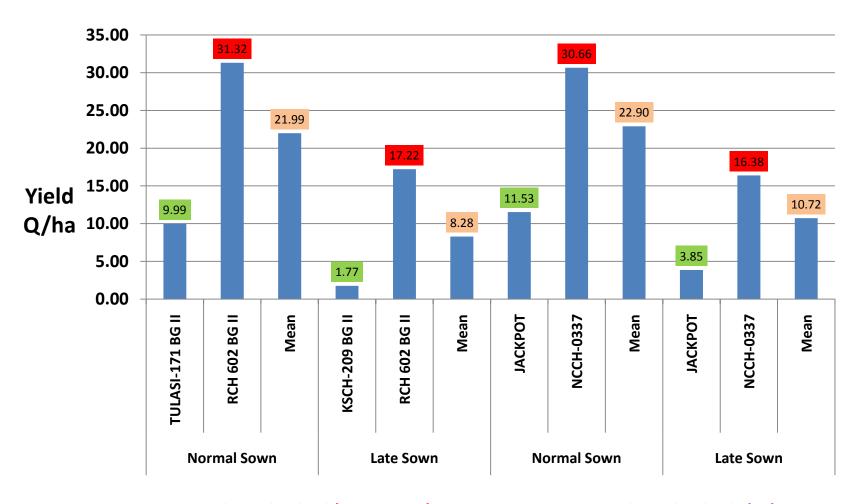
Pre-realeased Hybrids (50)





Released Hybrid (100 entries)





Released Hybrid (100 entries)

Pre-realeased Hybrids (50)



Multi location trials for shortlisting of CLCuD tolerant hybrids

 The released BG II hybrids for north zone are screened (2014-15 onwards) under replicated trials at five locations(Rajasthan-Sriganganagar, Punjab-Bhatinda & Faridkot, Haryana-Sirsa & Hisar) with susceptible check as infector rows after every four rows of test entries.

 The shortlisted tolerant hybrids only are advocated for marketing of seeds to the farmers.



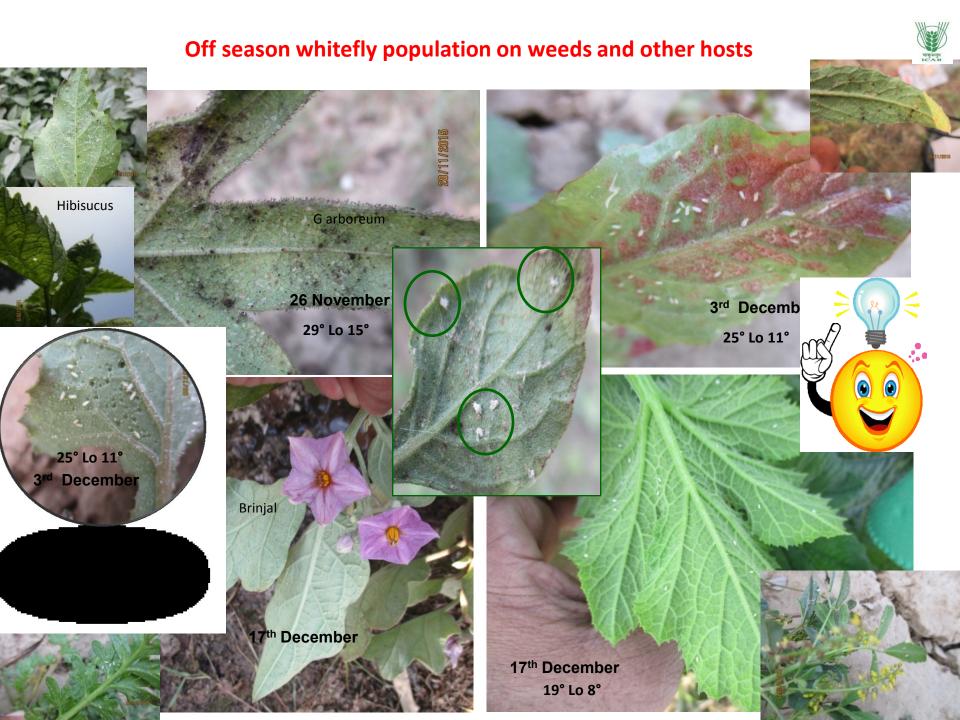


Chakrabarty, P.K, Sable, S., Monga, D. and Mayee, C. D. (2005). Polymerase chain reaction-based detection of Xanthomonas axonopodis pv malvacearum and cotton leaf curl virus. Indian J. Agric. Sci. 75: 524-27.



Detection of cotton leaf curl virus (CLCuV) in weeds showed that *Convulvus arvensis* (Hirankhuri), *Spinacea sp.* (Jungli Palak), *Solanum nigrum* (Blackberry nightshade), *Lantana camara* (*Raimuniya*), *Chenopodium album* (Bathua) were among naturally collected weeds which were detected positive by PCR amplification. *Achyrenthus aspera* (Puthkanda), *Digeria avensis* (Tandla), *Croton sprucifera* (Jungli mirch) and *Xanthium strumarium* (Gutpatna) were among artificially inoculated weeds which were found CLCuV positive by developing CLCuD type symptoms and by PCR amplification. Transmission of CLCuV from cotton to weeds and vice- versa was also successful.

Alok Kumar, D. Monga and Kishore Chand Kumhar (2017) Screening of cotton germplasm against leaf curl virus disease (CLCuD) and role of weeds for its development. J. Cotton res. Dev. 31 (1)87-96



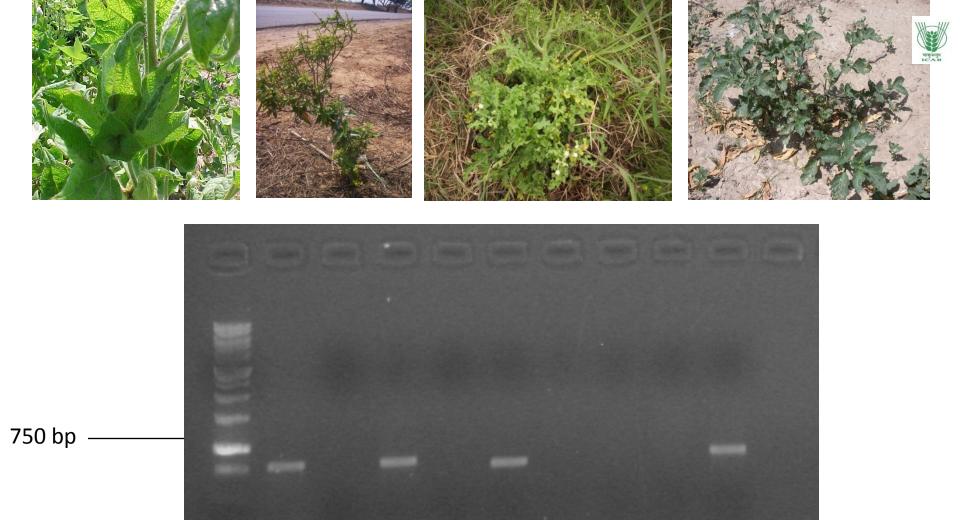


Figure: Amplification of CLCuV genomic components from the total DNA of weed plants with CP primers. Lane 1: Infected cotton; Lane 2: Healthy cotton; Lane 3: *Croton sparciflorus*; Lane 5: *Parthenium hysterophorus*; Lane 9: *Solanum americanum*; Lane 4, 6, 7 & 8: weeds showing no amplification.

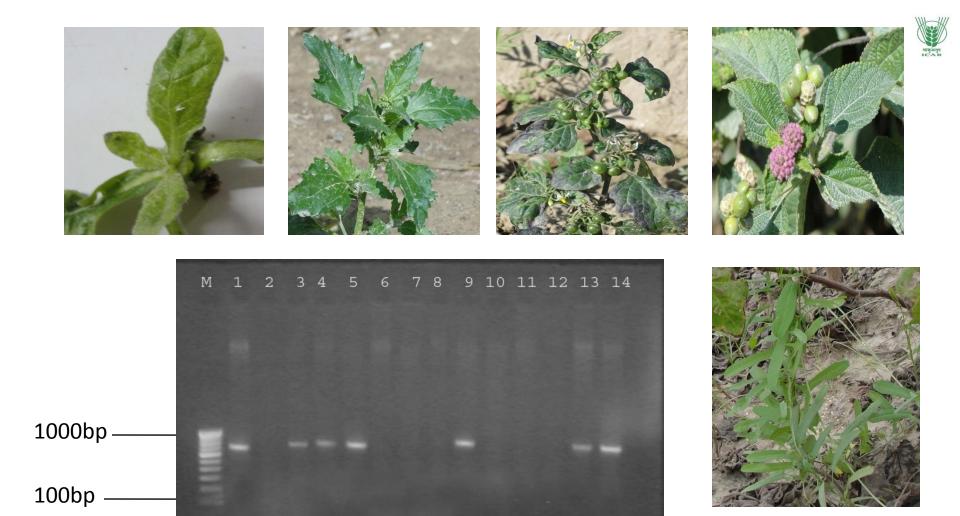
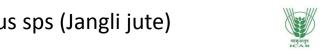


Figure: PCR amplification of weeds DNA collected from CICR RS Sirsa. Lane M-100 bp molecular marker Lane1. Spinacea spp Lane2. Chenopodium murate Lane3. Chenopodium album Lane4. Solanum nigrum Lane5. Lantana camaraLane6 Aerua sativa Lane7. Ageratum Lane8. Parthenium spp Lane9. Convolvulus arvensis Lane10 Solanum tubersum Lane11. Hibiscus rosa sinensis. Lane12. Chenopodium album Lane13. Check Lane14.check

Ageratum sps (Goat weed)



Chorchorus sps (Jangli jute)







Helianthus spp(Jungali suraj mukhi)



Tribulus terrestris (Bhakari)

CLCuV inoculum shifting from weeds to cotton











Weed eradication campaigns

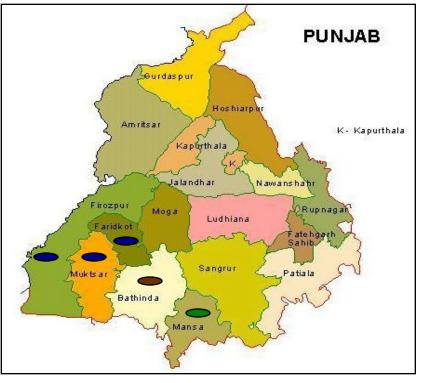
 Based on our trials regarding weeds serving as source of CLCuV inoculum, the state department officials conducted weed eradication programs starting from February onwards in a big way with active participation of farmers.

District wise incidence of CLCuD in Punjab



- > 50 % (V. Severe)
- 30.1-50 % (Severe)
- 20.1-30 % (Moderate)
- _____ 10.1 20 % (Low)
- ●0.1- 10 %(Very Low)
- 0.0 (Nil)





2015-16 2016-17

District wise incidence of CLCuD in Haryana



- > 50 % (V. Severe)
- 30.1-50 % (Severe)
- 20.1-30 % (Moderate)
- _____ 10.1 20 % (Low)
- ●0.1- 10 %(Very Low)
- 0.0 (Nil)





District wise incidence of CLCuD in Rajasthan



- > 50 % (V. Severe)
- 30.1-50 % (Severe)
- 20.1-30 % (Moderate)
- _____ 10.1 20 % (Low)
- 0.1- 10 %(Very Low)
- 0.0 (Nil)





2015-16 2016-17



The Way Ahead

CLCuD resistant lines identification/ Pyramiding resistance/Introgression of resistance

- In India, from among 5000 *G. hirsutum* lines during 2013-14 and 2128 lines during 2014-15 screened against CLCuD, none of the line was found to be resistant/ immune.
- However, lines identified as tolerant to disease are being used in pyramiding resistance against this disease.
- Recently two G hirsutum lines GVS8 & GVS 9 received from USA have shown immune reaction towards CLCuD and are being used for transfer of resistant sources.
- Introgression of CLCuD resistance from G arboreum is being attempted.

Gene pyramiding for CLCuD

Tolerant Sources: Bhiyani 251, MR 786, CSH 27, CSH 46 and CSH 538

Technical Programme

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1st    Year : F1s-2015
2nd    Year : F1s x F1s-2016
3rd    Year : (F1s x F1s) x (F1s x F1s)-2017
4th    Year : {(F1s x F1s) x (F1s x F1s)} x {(F1s x F1s) x (F1s x F1s)}
5th    Year : [{(F1s x F1s) x (F1s x F1s)} x {(F1s x F1s) x (F1s x F1s)}] x
[{(F1s x F1s) x (F1s x F1s)} x {(F1s x F1s)}]
so on .....
Selfing .... Selection of CLCuD resistant/tolerant plant
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Lines showing resistant reaction against CLCuD









Crossing program to introgress resistance to elite cultivars

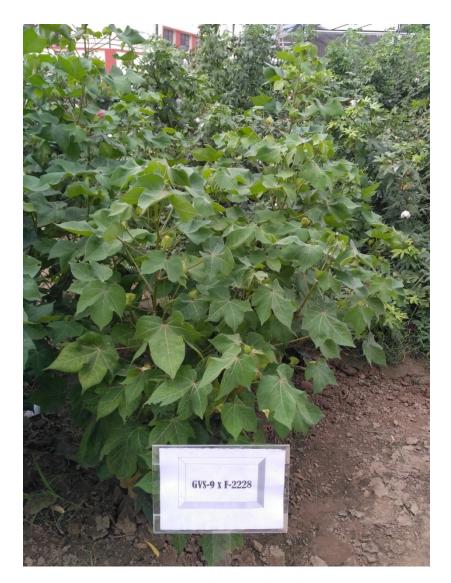






F1s showing resistant reaction against CLCuD





Introgression of CLCuD resistance from G. arboreum

Interspecific crosses:

• CSH 3129 (G. hirsutum) x CISA 310 (G. arboreum)-(2015) and subsequent crossing-(2016) using CSH 3129 (G. hirsutum) as recurrent parent.

F1 (CSH 3129 x CISA 310) x CSH 3129-(2017)

so on.... Identification of plants having CLCuD resistance.

Introgression using colchicine treated *G. arboreum*:

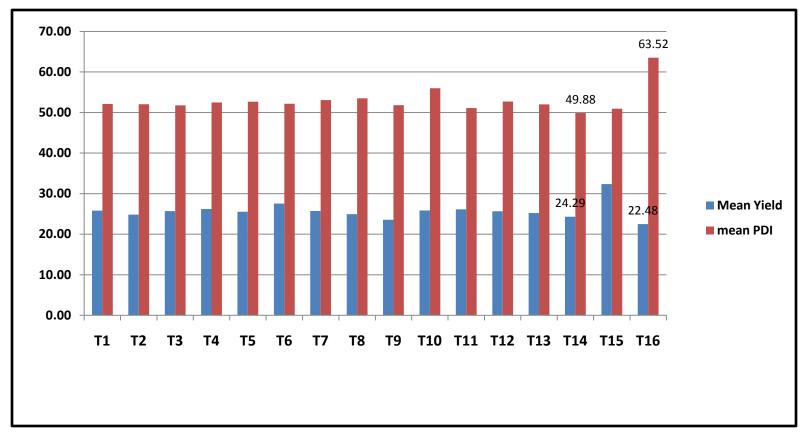
• CSH 3129 (G. hirsutum) x CISA 310 (G. arboreum) colchicine —(2016)treated and subsequent crossing using CSH 3129 (G. hirsutum) as recurrent parent.

F1 (CSH 3129 x CISA 310) x CSH 3129-(2017)

so on.... Identification of plants having CLCuD resistance.

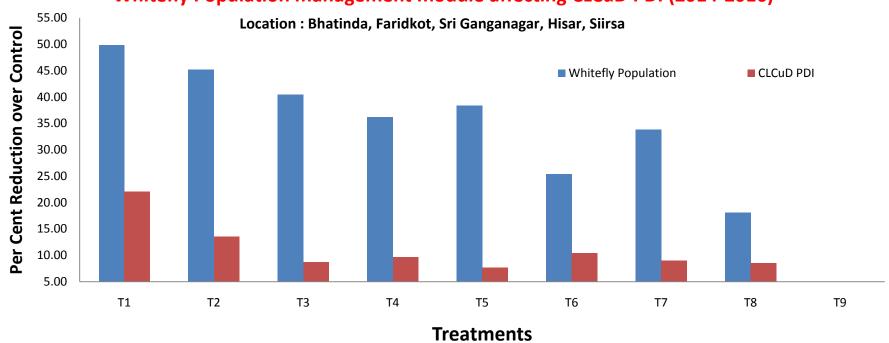
Path 3 e: Innovative interventions for the management of CLCuD in North Zone (Mean of two years 2015-16 and 2016-17)





Treatments		
T-1: Butter Milk @ 5%	T-9: Lachesis 30 @ 0.1%	
T-2: Cow Urine @ 6.6%	T-10: Digitalis 30 @ 0.1%	
T-3: Neem Oil @ 1%	T-11: Apis Mallifera 30 @ 0.1%	
T-4: Mustard Oil @ 3%	T-12: Bryonia 30 @ 0.1%	
T-5: Calcium Nitrate @ 0.5%	T-13: Natrum Mure 30 @ 0.1%	
T-6: Cow Urine + Calcium Nitrate	T-14: Polo @ 0.1% @ 0.1%	
T-7: Cow Urine + Butter Milk	T-15: Salicyclic Acid@200ppm (single spray)	
T-8: Butter Milk+ Calcium Nitrate	T-16: Control (Unsprayed)	

Whitefly Population management module affecting CLCuD PDI (2014-2016)



	Treatments		
T1	Spray of nimbicidine, V.lecanii, M.anisopliae, Difenthiurom, Hort.Min.Oil, Triazophos&Spiromesifen		
T2	Spray of nimbicidine, 20 DAS		
Т3	Spray of V.lecanii, 45 DAS		
T4	Spray of M.anisopliae, 60 DAS		
T5	Spray of Difenthiurom, 75 DAS		
Т6	Spray of Horticultural Mineral Oil (Hort.Min.Oil), 90 DAS		
T7	Spray of Triazophos, 105 DAS		
T8	Spray of Spiromesifen, 120 DAS		
Т9	Control		



Bioassay of selected EPFs and recommended chemical pesticides against whitefly nymphs under polyhouse condition

