

Research developments in begomovirus in legumes : Past chievements, Present scenario and Future thrust areas

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Abstract - *Begomovirus* – A well recognized genus of plant viruses classified under Geminiviridae family of Group II (single stranded circular DNA) is known to cause virulence in a large range of hosts including tomato, okra, cotton, cassava, bittergourd, chilli, croton, cucumber, eggplant,

jatropha, mentha, mesta, papaya, potato, tobacco, legumes and many more. It is believed that organism has caused major economic losses on crops of all types. However, owing to today's demand of increased production, land area and productivity of pulses for future food and nutritional security, this paper will summarize collectively the research study and experiments conducted on legumes till date, present research work being conducted and future thrust areas on *begomoviruses* w.r.t. legumes like common bean (Phaseolus vulgaris), mung bean (Vigna radiata), urd bean (Vigna mungo), pigeon pea (Cajanus cajan), mothbean (Vigna aconitifolia), cowpea (Vigna ungiculata), velvetbean (Mucuna pruriens), frenchbean (Phaseolus vulgaris), soybean (Glysine max), long yard bean (Vigna sesquipedalis) etc. Prime focus areas of this review will include molecular characterization, DNA replication, phylogenetic analysis and infectivity patterns on host legumes.

Key Words: Begomovirus, legumes, *Bemisia tabaci,* yellow mosaic disease, bean common mosaic virus

1. Introduction-

Among all the diseases of plants, viral diseases are the most complex and the least known. Intensive research in the recent past has led to showcase some facts with clear understanding. This review will deal with morphology, taxonomy, genome, population genomics, gene expression, economic impact, host range, transmission, molecular characterization, antigenicity, infectivity and management of begomoviruses that forms largest group of disease causing viruses of plants and specifically of legumes.

2. Morphology-

Virions of geminivirus sub group III are germinate .i.e. twinned with two incomplete icosahedra. They are nonenveloped in nature and have a common dimension of 30-38 nm * 18-28 nm (length*diameter) with 22 pentameric capsomers per nucleocaspid and 110 identical protein

subunits (Galvez & Castano, 1976; Goodman *et al.,* 1977;Qazi et. al.,2007).

3. Taxonomy-

Mungbean yellow mosaic virus, Mungbean yellow mosaic India virus, Horsegram yellow mosaic virus and Dolichos

yellow mosaic virus are bipartite begomovirus that are responsible for causing yellow mosaic diseases in legumes across southern Asia. MYMIV is a bipartite isolate of Begomovirus that is more widespread in tropical and subtropical climates (Chakraborty et al., 2003; Usharani et al., 2005; John et al., 2008;Fazeli et al., 2009; Haq et al., 2011a, 2011b).

4. Genome-

Genome of begomovirus is bipartite (Honda and Ikegami 1986; Vanitharani et al.1996; Mandal et al. 1997; Karthikeyan et al. 2004).Both the virions and complementary sense standards have coding regions, positive and negative respectively. Double standard intermediates are responsible for replication of the genome by rolling circle replication where component B is dependent on component A for replication with a common region consisting of two divergent promoters differentially regulating temporal expression of viral genes (SIB 2008).

5. Population genomics-

Experiments on global scale population structure of begomovirus concluded seven major sub populations that can be further divided into 34 entirely different yet genetically close minor sub populations(Prasanna et. al.,2010).

6. Gene expression-

Common region is responsible for bidirectional transcription in *begomovirus* while protein expression takes place at subgenomic RNA level. Bipartite *begomovirus* majorly a disease causing mesobiotic pathogen in legumes has two components- Component A encodes six proteins CP(on vsense) & Rep, TrAP/AL2, REn, AC4, AV2(on c-sense) and component B encodes 2 proteins BV1(on v-sense) & BC1(on c-sense) both involved in movement (Sunter and Bisaro 1992, Bisaro 2006, Sunter and Bisaro 1992, Noueiry et al. 1994, Fontes et al.1994, Laufs et al. 1995, Sanderfoot and Lazarowitz 1996, Hanley-Bowdoin et al. 2000, Arguello-Astorga et al. 2004, Bisaro 2006, SIB 2008, Hanley-Bowdoin et al. 2013, ICTV 2017).

7. Economic Impact-

After first report of yellow mosaic disease on major pulse group,its economic impact was expressed by various scientists.Yellow mosaic disease has a potential to cause 85-100% yield loss in black gram and mungbean (Nene,1973).Yellow mosaic disease accounted for a total loss of 105,000 metric tonnes in soybean alone(Wrather et. al.,1997).Total yield losses in blackgram,soybean and mungbean collectively are expected to be around \$ 300 million per year(Verma and Malathi,2003).

8. Host Range-

Begomovirus are reported to hold a wider host range but in general it serves on dicotyledonous plant.Specific major legume genera includes Phaseolus, Vigna, Macroptilium, Calopogonium(Bird *et al.*, 1972; Bird, Sanchez & Vakili, 1973; Meiners *et al.*, 1973).Rarely infected legume genera are *Cassia, Cajanus, Glycine, Rhynchosia* and *Phaseolus* (J. Bird).Intrinsic research work led to identify Limabean or lablab bean(Capoor and Verma,1948),Mungbean,urdbean and cowpea (Nariyani,1960;Nene,1973), soybean (Suteri,1974,Fernandes et al. 2009),horsegram (Muniyappa *et al.*, 1975),Frenchbean (Singh, 1979), wild bush bean or quail bean (*Macroptillium lathyroides*) (Lima et al. 2013).

9. Transmission-

Original transmission was reported by whitefly Bemisia tabaci (Hemiptera: Alevrodidae) in a persistent, nonpropagative manner across the globe(Costa,1965). Bemisia tabaci race sidae is known to spread the virus(Bird et. al.,1972).Both males and females are capable to act as a vector of the virus and no evidence of larvae acting as vectors has been found (Bird et. al., 1973). Infection is possible in the hosts within six days of acquisition and inoculation(Bird, 1973). However longer feeding periods are needed to spread the disease effectively.16-21 days approximately is the period to act as potential vectors after acquisition period of 10-15 minutes (Gamez, 1971; Bird et al., 1973). All whiteflies sub types B. tabaci MEAM1 (Middle East-Asia Minor 1) (Biotype B), *B. tabaci* NW (New World) (Biotype A), B. tabaci NW2 (New World 2), and B. tabaci MED (Mediterranean; Biotype Q) (Barbosa et al. 2015) are responsible for transmission of virus except biotype-b which acts as a vector only on monocots. Circulative transmission is known not to occur but replication of the virus inside the vector is questionable (Rosen et. al., 2015).

10. Molecular characterization-

Molecular characterization and infectivity test led to the conclusion that viral isolates obtained from Indian cowpea were 94-98% similar in its genomic sequence of DNA-A and

DNA-B and shared a close relationship with the other isolates obtained so far, thus proving the Koch's postulates for *begomovirus* association with mungbean yellow mosaic disease-India in mungbean and cowpea(Singh et. al.,2011).

11. Antigenicity

All the begomoviruses were put to test and serological tests evaluated them to be closely related. Harrison and Robinson, 1999 used monoclonal antibodies to group various begomoviruses geographically on the basis of shared epitopes.

12. Infectivity-

YMD's first incidence was marked in western and northern India in lima bean (Capoor and Verma, 1948). Nariyani, 1960 was the first to report vellow mosaic disease in mungbean in Indain subcontinent. Infection in legumes are caused by bean golden mosaic virus which is a type strain of begomovirus. It includes Mungbean yellow mosaic India virus, Mungbean yellow mosaic virus, Dolichos yellow mosaic virus which have bipartite genome and produce mosaic and leaf yellowing as a the major symptoms in legumes (Varma and Malathi 2003 ; Balaji et al. 2004; Girish et al. 2005).Furthermore, Mungbean yellow mosaic India virus (MYMIV) and Mungbean yellow mosaic virus (MYMV) were reported to be more virulent, having more hosts and abundant(Fauguet and Stanley,2003) and the other two strains Dolichos yellow mosaic virus and Horsegram yellow mosaic virus are specific and rare(Maruthi et. al.,2005). Molecular characterization proved begomovirus subgroup-III isolates were responsible to cause mild mosaic infection on Vigna mungo var. Sylvestris L.(Naimuddin et. al., 2011).

13. Management-

For the management of *begomovirus* it is important to lower down the inoculum and and the vector population.

Cutting and destruction of infected portion of the plant in case of low or moderate infection and uprooting of the plant in case of severe infection is profitable.

Seed treatment with imidachlorpid @ 5ml/kg seed with two sprays of imidachlorpid @ 0.5 ml/l at 25 and 40 DAS were found to be effective against the infection (Jayappa et. al.,2017). Another effective method is two sprays of neemazal @ 3ml/l after seed treatment with imidachlorpid @ 5ml/kg seed (Jayappa et. al.,2017). Predators offer a better possibility of vector control. These predators include lacewings, bigeyed bugs, and minute pirate bugs. Lady beetles including *Clitostethus arcuatus* (on ash whitefly), the Asian multicolored lady beetle *Harmonia axyridis* and scale predators, such as *Scymnus* or *Chilocorus* species feed on whiteflies (UCIPM).

14. Conclusion and Future Prospects-

Begomovirus with their quantity, infectivity and strain differentiations have always attaracted us to conduct exemplary research on them. The wide host range they share, the virulence they have developed and their close association with vector whitefly has brought them to immediate research action and work at present. Also as it fears nutritional security in vegetables like okra, brinjal, tomato, chili, cassava and food security through legumes and cereals, it has come up as an important and immediate research interest. Disease symptoms and particle morphology suggest that bean golden mosaic viruses(subgroup-III of begomovirus) from Puerto Rico, El Salvador, Colombia, Guatamala, and probably Brazil are the same virus, although possible strain relationships are not worked out. Causal agents of similar diseases in other tropical areas have not been characterized. Thus, future research should be emphasized to workout strain relationships and solve complexity of gene order of *begomovirus*. Another challenge is to mark the common gene pool, if any, of begomovirus to understand population distribution and gene flow to check increasing levels of virulent strains. The disease causing ability of the begomovirus has increased due to evolution of more virulent strains and is likely to increase due to uncontrollable spread of whitefly population, tropical climate and widespread cultivation of legumes. Hence, Integrated approach towards the solution must be adopted to control the same. B. tabaci MEAM1 (Middle East-Asia Minor 1) (Biotype B) is found to violate rule boundaries of begomovirus and is found to transmit virus on monocots like maize which can later spread to legumes owing new threat to food security of future. Thus, thrust should be laid to limit strain of *begomovirus* on monocots. Extreme research work is also demanded in the field of protein mapping technology to mark both protein in a bipartite virus and understand their structure to break gene bond which would eventually control the disease spread. A new approach .i.e. application of new generation sequencing (NGS) technology for better diagnosis by metagenomic analysis and deep sequencing is yet to be carried out in *begomovirus*. This will allow us to develop better understanding of viruses and their associations and control them to achieve food security.

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