Proteomic Analysis of Cassava Mosaic Virus (CMV) Responsive Proteins in Cassava Leaf

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Proteomics is becoming an increasingly important tool for the study of many different aspects of plant functions, such as investigating the molecular processes underlying host-pathogen interaction, plant physiology, development and differentiation. Cassava mosaic disease (CMD), caused by cassava mosaic virus (CMV), is the most serious disease in cassava. However, the molecular mechanisms underlying CMD in cassava during CMV infection is not yet clearly understood. The current study determined and identifies the differentially expressed proteins from cassava leaves during the infection of CMV viz., Indian Cassava mosaic virus (ICMV) and Sri Lankan Cassava Mosaic Virus (SLCMV). 2D gel electrophoresis was used to identify the cassava responsive proteins during the virus infection and the differentially expressed proteins were analysed by matrix-assisted laser desorption/ionization-time-of-flight (MALDI-TOF) mass spectrometry. There are 19 proteins were differentially expressed in cassava leaves by CMV infection. Among them 18 were giving good spectra by MALDI-TOF mass spectrometry. Analysis of Peptide Mass Fingerprint (PMF) data of these 18 proteins revealed the identity of the differentially expressed proteins, which suggest their importance and relevance on plant growth and development, and defence. This work paves the way towards a comprehensive analysis of CMV infection of cassava. Identification of the differentially expressed proteins by their sequence homology to known proteins suggests a possible direct or indirect role on plant defence during CMV infection. This study revealed the differentially expressed proteins, expressed during interaction between cassava and CMV that might play important roles either in viral pathogenesis or resistance.
**Introduction**

Cassava (*Manihot esculenta* Crantz) is a food security perennial crop of the Euphorbiaceae family, which originated in South America and reached Africa and Asia during the 16th and 17th centuries. Currently, cassava is extensively cultivated as an annual crop in tropical and subtropical regions for their rich source of carbohydrates (85%) and protein (1-2%) for human food in the world and the world’s sixth food crop for more than 800 million people (Liu *et al.*, 2011; Howeler *et al.*, 2013). It has a high growth rate under optimal conditions and the tuberous roots as well as the leaves are used as human food, animal feed and industrial products (Mann, 1997; El-Sharkawy, 2004; Sheffield *et al.*, 2006; Gbadegesin *et al.*, 2008). Although cassava roots and leaves combine high energy, protein and high levels of some vitamins, minerals and dietary fibre (Prathibha, 1995; Bradbury, 1998) and high productivity under drought and poor soil conditions, it is highly susceptible to various diseases, post-harvest physiological deterioration (Gegios *et al.*, 2010; Stephenson *et al.*, 2010; Vanerschuren *et al.*, 2014; Patil *et al.*, 2015; Urrota *et al.*, 2016). Cassava improvement programs are focused on addressing these constraints but are hindered by their high heterozygosity, difficulty in synchronizing flowering, low seed production and a poor understanding of the physiology of this plant (Ceballos *et al.*, 2004).

The yield of cassava can be reduced to up to 100% due to cassava mosaic disease (CMD), which is caused by various isolates of cassava mosaic geminiviruses (CMGs). In India, two CMGs namely *Indian Cassava mosaic virus* (ICMV) and *Sri Lankan Cassava Mosaic Virus* (SLCMV) are the causal agent of CMD in cassava. Due to its less importance, the research to improve cassava has lagged behind that of other crops such as rice, wheat, maize, and potatoes. Therefore, only relatively minor increases in cassava’s productivity have been obtained.

Analysis of proteins expressed in cassava leaf tissues will provide a better understanding of the constitutive differences controlling the plant’s growth, development, and defences during CMV infection. Furthermore, the recent molecular biological techniques of differential expression of genes or proteins during plant-pathogen interaction can be used as a powerful tool to dissect the molecular mechanism underlying the susceptibility of different cassava cultivars to CMV infection. In recent years, differential expression of eukaryotic proteins has been employed as a research approach in many laboratories to detect proteins that change in response to pathogen ingress. The main advantage of this technique is that it permits the simultaneous identification of up and down regulated proteins and may serve as genetic and diagnostic markers, as well as providing insights into the underlying mechanisms of disease incidence. Most of the previous studies focused on the effects of environmental factors and physiology of cassava in relationship to crop yield (extensively reviewed in El-Sharkawy, 2004). At the molecular level, there are few reports on genes and proteins that may play important roles in controlling cassava storage root formation and yield. Yet, there has not been any report in the literature on molecular and biochemical investigation of leaf genes or proteins of cassava during CMV infection.

Particularly, to date, there have been very few papers in the literature about proteomic analysis on storage root (*Souza et al.*, 1998; Cabral and Carvalho, 2001; De Souza *et al.*, 2002; Shewry, 2003; Sheffield *et al.*, 2006), somatic embryos, plantlets and tuberous root (*Li et al.*, 2010). However, a proteome analysis of cassava leaf during growth and
development was reported earlier by Mitprasat et al., (2011).

This study represents the proteomic analysis of cassava leaves during CMV infection. To further fulfill the lacking knowledge in the literature, leaf proteins that were differentially expressed during CMV infection of cassava were examined using a proteomic approach. Two-dimensional (2-D) gel electrophoresis combined with mass spectrometry revealed a number of candidate proteins that are differentially expressed between CMV-infected and non-infected healthy cassava leaves.

Materials and Methods

Genetic materials

Cassava cultivar H226 was obtained from the germplasm pool, Tapioca and Castor Research Station (TCRS), Yethapur, Tamil Nadu Agricultural University (TNAU). The healthy (meristem-derived virus free) and CMV infected (artificially inoculated meristem-derived) cassava leaves were used as the protein source in this study.

Cassava meristem culture

The virus-free healthy cassava plants were developed through apical meristem culture at Faculty of Horticulture, TNAU as described previously (Raghu et al., 2011). All the meristem-derived plants were fertilized with Hoagland solution (Hoagland and Arnon 1950) and the absence of CMV was detected by Polymerase chain reaction (PCR) using CMGs degenerate primers (Forward: 5’- TGT GAC CTT GAT TGG CAC CTG-3’; Reverse: 5’-CTC GAC GAG TGG TTT CAC GA-3’ for ICMV and Forward: 5’-TAG CTG CCC TGT GTT GGA C-3’; Reverse: 5’-TGA GAA ACC CAC GAT TCA GA-3’ for SLCMV). Reaction conditions were essentially those of Sambrook et al., (1989). PCR parameters were 94°C for 2 min then 40 cycles of 1 min at 94°C, 1 min at 63°C and 1 min at 72°C, followed by the final extension of 10 min at 72°C.

Proteomic analysis

Sampling

Three biological replicates of leaf tissues were collected from healthy (control) and CMV infected cassava plants (Figure 1) and immediately transferred into liquid nitrogen (LN₂) and stored at -80°C until further use.

Protein extraction

Three biological replicates of leaf tissues were ground finely in a mortar cooled with Liquid Nitrogen and suspended in 10% (w/v) trichloroacetic acid in acetone with 0.07% (w/v) dithiothreitol (DTT) at -20°C for 1 h, followed by centrifugation for 15 min at
35,000 g. The pellets were washed with ice-cold acetone containing 0.07% DTT, incubated at -20°C for 1 h, and centrifuged again at 4°C. This step was repeated thrice and the final pellets were lyophilized. The powder was then solubilized in lysis buffer (at 37°C) and the protein content was determined by the Bradford method (Bradford, 1976; Salekdeh et al., 2002; Jagadish et al., 2010).

**2D gel**

Equal amounts of protein (150 µg) from healthy and CMV infected samples were separated by Two-dimensional polyacrylamide gel electrophoresis (2D-PAGE), as described by Yan, et al., (2005). In the first dimension, IPG strips (BioRad Laboratories, USA) of 17-cm length and pH 4–7 were used. Electrophoresis was carried out at 400 V for 1 h, followed by 1000 V for 1 h and 2950 V for 24 h. After IEF, the proteins were separated by SDS-PAGE in the second dimension using 13% polyacrylamide gels (Salekdeh et al., 2002). The gels were stained by silver staining method (Blum et al., 1987). For each biological replicate, one set of gels with high resolution, run at different times, was selected for further analysis. The relative abundance of protein spots was quantified with Melanie III (GeneBio, Geneva, Switzerland), after silver staining the gels, and scanned with a densitometer (GS-700, Bio-Rad).

**Matrix-assisted laser desorption/ionization-time of flight mass spectrometry (MALDI-TOF MS) and database searching**

Selected spots were excised from preparative gels (stained with AgNO3) (Salekdeh et al., 2002) and extracted by an addition of 10µl of the extraction buffer, followed by an addition of 10-15 µl of acetonitrile. Pooled extracts were dried in a lyophilizer (SFDSN06, Samwon Freezing Engineering Co., Busan) and the extracts were re-dissolved in 1µl of extraction buffer and 1µl of matrix solution (α-acyano-4-hydroxycinnamic acid, HCCA) and targeted onto a MALDI-TOF plate. After drying the samples completely onto the targeting plate, MALDI-TOF/MS was conducted using a Voyager-DE STR mass spectrometer (Applied Biosystems, Franklin Lakes, NJ, USA) equipped with delay ion extraction. Mass spectra were obtained over a mass range of 808-2705 Da. For identification of proteins, the peptide mass fingerprinting data were used to search against the Swissprot database using the Mascot program (http://www.matrixscience.com). The following parameters were used for database searches: taxonomy, viridiplantae, cleavage specificity, trypsin with one missed cleavage allowed; peptide tolerance of 100 ppm for the fragment ions; and allowed modifications, Cys Carbamidomethyl (fixed), and oxidation of Met (variable).

**Results and Discussion**

**CMV responsive protein profiling in cassava leaves by 2D-gel analysis**

Comparison of mRNA or proteins isolated from target tissues of infected and healthy (control) (Figure 1) plants can provide information on the biochemical and molecular changes associated with CMV infection of cassava cv.H226. Thus, a proteomic analysis could be a powerful approach to identify responsive proteins associated to a biotic stress, such as pathogen infection. In this study, we adopted a proteomic strategy using 2-D gel electrophoresis to understand the molecular changes in cassava leaves infected by CMV versus that healthy cassava cv. H226.

CMV was artificially inoculated in healthy (meristem-derived) plants of cassava cv.H226 by whiteflies (Bemisia tabaci) and leaf tissues
were collected at after 21 days post inoculation for the proteomic analysis. Proteins were extracted from leaves using the TCA precipitation method and separated by 2-D gel electrophoresis as previously described by Yan et al., (2005). Silver staining of cassava leaf proteins separated by 2-D gel electrophoresis allowed the detection of around 300-350 spots (Figure 2). Comparison of 2-D gel electrophoretic pattern of leaf proteins between infected and healthy cassava leaves revealed the differential expression of nineteen protein spots (Table 1). Among the 19 protein spots that were differentially expressed, 11 (58%) were found to be up-regulated (Figure 3a) and 8 (42%) were found to be down-regulated by CMV infection of cassava cv. H226.

Analysis of differentially expressed proteins of cassava during CMV infection

Among the 19 differentially expressed protein spots, only 18 protein spots resulted in good spectra by MALDI-TOF while spot #1 did not. PMF data analysis of the eighteen protein spots derived by MALDI-TOF mass spectrometry using MASCOT search algorithm showed homology to ribosomal protein 4, chaperone protein DNAj, putative cytochrome c oxidase subunit II PS17, ATP binding cassette transporter, maturase K, oxygen-evolving enhancer protein 1, ascorbate peroxidase APX2, ATP synthase beta subunit, protein kinase-coding resistance protein, 2-oxoglutarate-Fe(II)-dependent oxygenase domain containing protein, component of cytosolic 80S ribosome, 40S small subunit and NADP-dependent sorbitol-6-phosphate dehydrogenase.

Protein spot #17 and #6 have the higher (6.730) and lower (0.203) abundance ratio, respectively. The average spot abundance ratio of down regulated proteins was 0.482 and that of up-regulated proteins was 3.556. Because of limited genome information of cassava in database, only 4 differentially-expressed protein spots (#10, #13, #15 and #16) were identified by the peptide mass fingerprint analysis. The differential expression of proteins in spots #10, #13, #15 and #16 was significant, while the remaining proteins were found to be marginally significant (Table 1). We attribute the somehow low number to differentially expressed proteins that have significant Mascot scores to the limited genome information of cassava in the database.

The proteomic analysis conducted here showed that the differentially expressed proteins identified which are either up or down regulated during CMV infection of cassava cv. H226 may play important roles related to plant growth, development and the defense against virus infection.

The present study is the second report on proteomics of cassava and the first one studying the cassava leaf proteome during CMV infection. Our aim was to identify major leaf proteins that exhibit differential expression pattern during CMD, which is caused by CMV in cassava plants. Biotic and abiotic stress results in alterations of plant homeostasis, including reduced photosynthetic rate and ionic imbalance. Induction of disease tolerance in plants involves a complex network of signal perception, amplification, and transduction which might include protein phosphorylation cascades, ion fluxes, oxidative stress and generation of secondary signals and activation of various genes involved in disease tolerance. Gene and protein expression profiling has become an important tool to investigate how an organism responds to environmental changes. Yet, there has not been any report in the literature on molecular and biochemical investigation of leaf genes or proteins involve in CMD of cassava.
Particularly, to date, there have been very few papers in the literature about proteomic analysis of cassava (Cabral and Carvalho 2001; Sheffield et al., 2006; Li et al., 2010; Mitprasat et al., 2011). To further fulfil the lacking information in the literature, leaf proteins that are differentially expressed during CMD in cassava were examined in this study using a proteomic approach.

Recent whole genome expression profiling techniques such as microarrays and proteomics have been used to dissect the molecular mechanism(s) leading to the development of a phenotype. Differential expression of genes or proteins in storage root (Cabral and Carvalho, 2001; De Souza et al., 2002; Sheffield et al., 2006), somatic embryos, plantlets and tuberous root (Li et al., 2010) and leaf (Mitprasat et al., 2011) during growth and development of cassava has been reported earlier. In this study, we adopted a proteomics strategy to understand the molecular changes in leaves of healthy and infected cassava (cv.H226) plants.

Silver staining of the cassava leaf proteins separated by 2-D gel electrophoresis allowed the detection of around 300–350 spots. Comparison of 2-D gel electrophoretic pattern of leaves proteins between healthy and infected plants revealed the differential expression of 19 protein spots (Table 1). Among the 19 differentially expressed proteins, eleven protein spots (#3, #4, #5, #7, #10, #13, #14, #16, #17, #18 and #19) were found to be up-regulated and eight protein spots (#1, #2, #6, #8, #9, #11, #12, and #15) were found to be down-regulated in cassava plants cv.H226 during CMV infection. Analysis of PMF data coupled with MASCOT searches allowed the identification of eighteen proteins showing significant or marginally significant homology to known proteins. Most likely the fact that only 4 differentially expressed proteins had significant matches while the other 14 were marginally significant is due to at least in part to the limited genome information of cassava on the NCBI database. Cabral and Carvalho (2001) and De Souza et al., (2002) reported similar findings about proteins associated with storage root formation in cassava. Mitprasat et al., (2011) reported that around 39 spots, which were successfully identified by ion trap LC–MS/MS, were significantly altered (P=0.05) during week 4 to 8 of growth in cassava leaf proteomic analysis during plant development, from planting of stem cutting to storage root formation.

**Translational control**

The ribosome is a two-subunit ribonucleoprotein complex that catalyzes the peptidyl transferase reaction of polypeptide synthesis, an absolute requirement. Chang et al., (2005) characterized 251 Evolutionarily Conserved and Variable Proteins of cytosolic 80S and 40S ribosomes in Arabidopsis. The present study revealed that the up regulation of component of cytosolic 80S ribosome and 40S small subunit shows induced protein synthesis. This may be due to the expression of viral and plant proteins which are involved in host pathogen interaction.

**Metabolism related proteins**

The group of differentially expressed proteins which are involved in primary metabolism can also provide substrate for the synthesis of secondary metabolites. The 6 proteins identified from this group comprised: NADP-dependent sorbitol-6-phosphate dehydrogenase (spot #19), 2-oxoglutarate-Fe(II)-dependent oxygenase domain containing protein (spot #17), cytochrome c oxidase (spot #6), ATP synthase beta subunit (spot #15), maturase K (spot #9) and oxygen-evolving enhancer protein 1 (spot #12).
Table 1: Abundance ratio and identity of induced proteins among cassava mosaic virus (CaMV) infected cassava leaves

| Spt ID | Up/Down regulation | Experimental pI value | Experimental Mass | Abundance ratio | Coverage | Mowse score | Theoretical pI | MW | Accession No | Putative Function |
|--------|---------------------|-----------------------|-------------------|-----------------|----------|-------------|----------------|----|--------------|------------------|
| #2     | Down                | 4.65                  | 12037             | 0.274           | 16       | 64          | 06.13          | 8300| XP002322858 | Predicted protein (Populus trichocarpha) |
| #3     | Up                  | 4.6                   | 15116             | 3.960           | 38       | 68          | 10.16          | 2197| AAG52804     | Ribosomal protein 4 (Leptobryum stellatum) |
| #4     | Up                  | 4.7                   | 15707             | 3.790           | 46       | 53          | 09.69          | 8150| XP002535156 | Chaperone protein DNAj, putative (Ricinus communis) |
| #5     | Up                  | 5.4                   | 17007             | 2.451           | 83       | 59          | 09.94          | 6615| CAN63043     | Hypothetical protein (Vitis vinifera) |
| #6     | Down                | 5.86                  | 14503             | 0.203           | 100      | 43          | 09.62          | 1707| P84733       | Putative cytochrome c oxidase subunit II PS17 (Pinus strobus) |
| #7     | Up                  | 6.2                   | 14514             | 4.070           | 33       | 67          | 9.53           | 3011| XP002331812 | Predicted protein (Populus trichocarpha) |
| #8     | Down                | 6.48                  | 18301             | 0.290           | 35       | 60          | 9.11           | 1106| XP002969857 | ATP binding cassette transporter (Selaginella moellendorfii) |
| #9     | Down                | 4.2                   | 31021             | 0.620           | 23       | 62          | 9.65           | 3240| AEK35190     | maturase K, partial (chloroplast) (Datura stramonium) |
| #10    | Up                  | 4.72                  | 20402             | 2.513           | 24       | 78          | 5.43           | 3307| XP002951214 | Hypothetical protein (Volvox carteri f. nagariensis) |
| #11    | Down                | 5.2                   | 25106             | 0.590           | 60       | 55          | 10.10          | 5990| XP002538199 | Conserved Hypothetical protein (Ricinus communis) |
| #12    | Down                | 5.34                  | 25051             | 0.783           | 52       | 40          | 5.36           | 1066| P84989       | Oxygen-evolving enhancer protein 1 (chloroplast) (Populus euphratica) |
| #13    | Up                  | 5.72                  | 34003             | 2.240           | 46       | 93          | 5.73           | 1730| AAX84679     | Ascorbate peroxidase APX2 (Manihot esculenta) |
| #14    | Up                  | 5.86                  | 34001             | 4.554           | 29       | 56          | 5.96           | 2544| XP002985124 | Hypothetical protein (Selaginella moellendorfii) |
| #15    | Down                | 5.4                   | 64010             | 0.590           | 55       | 101         | 5.03           | 3671| CAJ80585     | ATP synthase beta subunit (Physalis aequata) |
| #16    | Up                  | 6.07                  | 60101             | 3.146           | 39       | 71          | 8.48           | 1618| ACO25596     | Protein kinase-coding resistance protein (Nicotiana repanda) |
| #17    | Up                  | 6.66                  | 40008             | 6.730           | 19       | 61          | 5.89           | 3740| NP190532     | 2-oxoglutarate-Fe(II)-dependent oxygenase domain containing protein (Arabidopsis thaliana) |
| #18    | Up                  | 5.94                  | 61017             | 2.580           | 38       | 62          | 10.18          | 2976| XP002954017 | Component of cytosolic 80S ribosome and 40S small subunit (Volvox carteri f. nagariensis) |
| #19    | Up                  | 6.85                  | 38012             | 3.094           | 30       | 64          | 9.16           | 2852| AAM77729     | NADP-dependent sorbitol-6-phosphate dehydrogenase (Prunus emarginata) |

Spot ID, Experimental and theoretical pI, and MW correspond to the protein spot numbers indicated in Fig. 3. Proteins were identified by using the peptide masses from MALDI-TOF analysis, followed by data base search. Corresponding accession numbers for the identified proteins were obtained from NCBI (www.ncbi.nlm.nih.gov/)
**Fig. 1** Healthy and Cassava mosaic virus (CaMV) infected cassava plants. a) Healthy uninfected shoots without mosaic symptoms; b) CaMV infected shoots showing pronounced mosaic pattern with narrow, severely twisted and distorted leaves.

**Fig. 2** Two-dimensional gel electrophoresis analysis of total proteins extracted from the leaf tissue of cassava cv.H226 under control conditions. In the first dimension (IEF), 150 µg of protein was loaded on a 17-cm IPG strip with a linear gradient of pH 4–7. In the second dimension, 13% SDS-PAGE gels were used with molecular weight (Mr) standards. Proteins were visualized by silver staining. The arrows indicate 19 proteins that showed up and down regulation and significantly under healthy and infected conditions.
**Fig. 3** Magnified view of differentially expressed protein spots induced by cassava mosaic virus (CaMV) infection in cassava cv.H226 in a 2D gel electrophoresis. Up-regulated (A) and down-regulated (B) proteins are shown.
Sorbitol is known to be the primary transport product of photosynthesis. NADP-dependent sorbitol-6-phosphate dehydrogenase (spot #19) is a key enzyme in sorbitol biosynthesis, where it catalyzes the NADPH-dependent reduction of glucose-6-phosphate to sorbitol-6-phosphate (Herrera et al., 2010; Zhu et al., 2011).

Ferrous iron dependent oxygenases are a superfamily of enzymes that catalyse a wide range of reactions including hydroxylation, desaturation and oxidative ring closures. All the previous research studied 2-oxoglutarate (2OG) -Fe(II)-dependent oxygenase domain containing protein (spot #17) have an absolute requirement for Fe (II) and catalyse a variety of two-electron oxidations, including hydroxylation, desaturation and oxidative ring closure reactions (Prescott, 1993; Prescott and John, 1996). In almost all cases, the oxidation of the ‘prime’ substrate is coupled to the conversion of 2OG into succinate and CO2. One of the oxygens of the dioxygen molecule is incorporated into succinate. In the case of desaturation reactions, the other dioxygen-derived oxygen is presumably converted to water. In hydroxylation reactions, the partial incorporation of oxygen from dioxygen into the alcohol product occurs with significant levels of exchange of oxygen from water being observed (Baldwin et al., 1993; Lloyd et al., 1999).

The changing demands for energy and biosynthetic intermediates during plant growth and development are accommodated to a large extent by changes in the number and activity of mitochondria. Mitochondrial oxidative phosphorylation (OX-PHOS) in most eukaryotes is based on the sequential operation of five protein complexes termed complex I (NADH dehydrogenase), complex II (succinate dehydrogenase), complex III (cytochrome c reductase), complex IV (cytochrome c oxidase) and complex V (ATP synthase complex). The Cyt c oxidase complex is the terminal electron acceptor of the mitochondrial inner membrane respiratory chain. Evidence for possible activities of the cytochrome-linked systems in plants has been seen in: (a) oxidative systems in nonphotosynthetic tissues; (b) respiratory mechanisms in dark reactions in photosynthetic cells; and (c) light induced reactions (Smith, 1958).

Respiratory oxidative phosphorylation represents a central functionality in plant metabolism (Vanlerberghe and McIntosh, 1997; Plaxton and Podesta, 2006). Respiration rates, maximum activity of cytochrome c oxidase (protein spot #6), and active mitochondrial number consistently decreased in plants infected with CMV. Plant growth during CMV infection also reduced cytochrome pathway activity and total mitochondrial ATP production.

ATP is a ubiquitous, energy-rich molecule of fundamental importance in living organisms. It is a key substrate and vital cofactor in many biochemical reactions and is thus conserved by all cells. Chivasa et al., (2011) identified subunits of the vacuolar and chloroplastic ATP synthase proteins as responsive to fumonisin B1 (FB1), with the great majority belonging to the mitochondrial F_{1}F_{0}-ATP synthase machinery. In this study, the ATP synthase beta subunit (protein spot #15) was down regulated that show that many biochemical reactions are affected during host-pathogen interaction between cassava and CMV. Similarly, Nwugo et al., 2013 also found that ATP synthase beta subunit got down regulated during Candidatus Liberibacter asiaticus infection in grapefruit plants. matK is an invaluable gene present in chloroplast-encoded group II intron maturase (Muller et al., 2006; Barthet et al., 2007). RNA editing mechanisms previously reported in matK (Vogel et al., 1997; Tillich
et al., 2005) may correct the reading frame in species with frame shift indels and premature stop codons and restore the codon identities needed to form the proper amino acids for function. Further, genome studies of the holoparasite Epifagus virginiana (Ems et al., 1995) and Adiantum capillus-veneris (Wolf et al., 2003) support that matK has a function in the plant. This putative enzyme critically impacts all chloroplast function including photosynthesis. Several chloroplast genes have light-induced expression (Klein and Mullet, 1990; Klein, 1991; Baumgartner et al., 1993). These genes are involved in two major activities of the chloroplast: photosynthesis and chloroplast development (Klein, 1991). Chloroplast development requires turning on protein translation in this organelle and increases the expression of all RNAs and proteins related to the translation machinery (Baumgartner et al., 1993). Maturase is needed for processing introns in order to generate the needed proteins and tRNAs for photosynthesis and/or the chloroplast translation machinery. However, maturase K (protein spot #9), which is involved directly or indirectly in the regulation of plant development, was found to be down regulated during CMV infection in cassava in our study.

The present study showed the down regulation of oxygen-evolving enhancer protein 1 (protein spot #12), which is involved in photosynthesis of Cassava. Similarly Nwugo et al., (2013) also observed that a down regulation of Oxygen-evolving enhancer (OEE) proteins during Candidatus Liberibacter asiaticus’ (Las) infection which causes Huanglongbing (HLB) disease in grapefruit (Citrus paradisi). OEE proteins 1 and 2 are subunits of the oxygen-evolving system of PSII and are involved in stabilizing the Mn cluster (Pushkar et a., 2008). HLB-affected trees generally show leaf yellowing (chlorosis) which is likely due to a reduction in chlorophyll biosynthesis (Sagaram et al., 2009; Liao et al., 2012) and Mg is important in chlorophyll biosynthesis. Thus, a virus-mediated reduction of the Mg content together with a reduction in Fe content of leaves of cassava plants could play a role in CMD-associated chlorosis.

**Molecular chaperones**

Virus proliferation depends on the successful recruitment of host cellular components for their own replication, protein synthesis and virion assembly. In the course of virus particle production, a large number of proteins are synthesized in a relatively short time, whereby protein folding can become a limiting step. Most viruses therefore need cellular chaperones during their life cycle. In addition to their own protein folding problems, viruses need to usurp or divert cellular resources, including host factors, away from their normal function (Witham and Wang, 2004) and interfere with cellular processes such as signal transduction, cell cycle regulation and induction of apoptosis in order to create a favourable environment for their proliferation and to avoid premature cell death (Mayer, 2005; Scholth of, 2005).

Chaperones are involved in the control of these cellular processes and some viruses reprogram their host cell by interacting with them.

Molecular chaperones are thought to be involved as there are molecules present in anucleate sieve elements (SE) sap that are larger than the size exclusion limit (SEL) of plasmodesmata (PD) and movement can be bidirectional (Golecki et al., 1999; Oparka, 2004). It is thought that some proteins partially unfold and bind to another molecule that assists passage through the PD (Lucas, 1999; Ding et al., 2003). On the SE side, chaperone molecules would be required for the correct re-formation of the protein.
Molecular chaperones may function by binding specifically to interactive protein surfaces exposed transiently during a cellular pro-cess, preventing them from undergoing incorrect interac-tions that might produce non-functional structures (Ellis, 1990).

Hsp70 chaperones, as central components of the cellular chaperone network, are frequently recruited by viruses. The chaperone function of Hsp 70 proteins in these events is regulated by members of the DnaJ-like protein (protein spot #4) family, which occurs through direct interaction of different Hsp70 and DnaJ-like protein pairs that appear to be specifically adapted to each other. Bargen et al., (2001) identified DnaJ-like protein as a non-structural protein encoded by the mRNA segment (NSm) of tomato spotted wilt virus (TSWV) interacting host proteins, implying an involvement of molecular chaperones during systemic spread of the virus by cell-to-cell movement of nucleocapsid through modified plasmodesmata (PD) in tobacco and Arabidopsis.

**Signalling and disease resistance**

When plants are exposed to stressful environmental conditions, the production of Reactive Oxygen Species (ROS) such as O$_2^-$, OH•, and H$_2$O$_2$ increases and can cause significant damage to the cell components such as DNA, proteins and lipids (Thakur and Sohal, 2013). It is known that an excess of free oxygen radicals leads to programmed cell death (PCD) (Pellinen et al., 2002; Vranova et al., 2002). However, ROS are also utilized in various metabolic processes such as formation of lignin in the cell wall (Inze and Montagu, 1995), leaf and flower abscission, cell senescence, ripening of fruit and flowering (Mehlhorn et al., 1996). For the protection from oxidative damage, plant cells contain both oxygen radical detoxifying enzymes such as catalase, peroxidase, and superoxide dismutase, and nonenzymatic antioxidants such as ascorbate peroxidase and glutathione-S-transferase (Pnueli et al., 2003). These enzymes play a crucial role in the protection of plant cells from oxidative damage at the sites of enhanced ROS generation (Kuniak and Sklodowska, 2001). The cooperative function of these antioxidants plays an important role in scavenging ROS and maintaining the physiological redox status of organisms (Cho and Seo, 2005).

Antioxidant defenses, which can detoxify ROS, are present in plants (Mittler, 2002; Apel and Hirt, 2004; Foyer and Noctor, 2005). A major hydrogen peroxide detoxifying system in plant cells is the ascorbate-glutathione cycle, in which, ascorbate peroxidase (APX) enzymes play a key role catalyzing the conversion of H$_2$O$_2$ into H$_2$O, using ascorbate as a specific electron donor (Dąbrowska et al., 2007). Different APX isoforms are present in distinct subcellular compartments, such as chloroplasts, mitochondria, peroxisome, and cytosol (Caverzan et al., 2012). The APX responses are directly involved in the protection of plant cells against adverse environmental conditions. In the present study, the cell damage due to the CaMV infection was reduced by enhanced production of ascorbate peroxidase (protein spot #13) in the leaf tissues.

Constant exposure to pathogen attack during their long evolutionary history of host plants has resulted in plant-pathogen coevolution. Interactions between plant pathogens and their host plants involve specific recognition and subsequent activation of a cascade of plant defense responses. Plant resistance gene (R-gene) plays an important role in plant-pathogen recognition (Bendahmane, 2002). The protein encoded by the majority of the disease resistance genes present several highly conserved domains: nucleotide binding
site (NBS), leucine-rich repeat (LRR), toll/interleukin receptor (TIR) domain, protein kinase (PK) domain etc. (Jones et al., 2001; Xiao et al., 2006). Intercellular signalling protein kinases that play a signalling role in the regulation of cellular energy metabolism. Their activity largely depends upon the concentration of cellular AMP which is increased under conditions of low energy or metabolic stress. Gao et al., (2010) expressed 73 resistant gene analogs (RGAs) of the protein kinase (PK) class in tobacco with challenged inoculation with Tobacco mosaic virus (TMV) or the tobacco black shank pathogen (Phytophthora parasitica var. nicotianae). The expression of two RGAs of the PK class was induced by P. parasitica var. nicotianae. Infection by either TMV or P. parasitica var. nicotianae enhanced the expression of protein kinase genes coding resistance proteins. The present study shows that the up regulation of protein kinase-coding resistance protein (protein spot #16) by CaMV should provide valuable information for cloning related resistance genes in cassava.

Previous reports have shown that the Arabidopsis ABC transporters (spot #8) AtABCG36, AtABCG40, and NpPDR1 are involved in plant defense responses (Lipka et al., 2005; Kobae et al., 2006; Stein et al., 2006; Clay et al., 2009; Badri et al., 2012). For example, Badri et al., (2012) demonstrated the involvement of seven root-expressed ATP-binding cassette (ABC) transporters (Atabcg36, Atabcg37, Atabcc5, Atabcf1, Atabcf3, Atnap5 and Atath10) in higher expression of defense genes by secreting phytoalexin in Arabidopsis thaliana after pathogen inoculation. Atabcg37 and Atabcc5 secreted higher levels of the phytoalexin camalexin, and Atabcg36 secreted higher levels of organic acids, specifically salicylic acid (SA).

This extensive study effectively provides a basis for further functional characterization of differentially expressed leaf proteins, which can help understand how biochemical processes in cassava leaves may be involved in cassava mosaic disease and dissect the molecular basis of host-pathogen interaction between cassava and cassava mosaic virus.

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Compliance with Ethical Standards

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