Title: Cassava brown streak disease: historical timeline, current knowledge and future prospects

Key words:

CBSD; CBSV; UCBSV; CMD; Food Security; Virology

Authors:

Katie R. Tomlinson (1); Andy M. Bailey (1); Titus Alicai (2); Sue Seal (3);
Gary D. Foster (1)*

(1) School of Biological Sciences, University of Bristol, 24 Tyndall Avenue,
Bristol, BS8 1TQ, UK

(2) National Crops Resources Research Institute, P.O. Box 7084,
Kampala, Uganda

(3) Natural Resources Institute, University of Greenwich, Central Avenue,
Chatham Maritime, Kent ME4 4TB, UK

*Corresponding author email: Gary.Foster@bristol.ac.uk, Tel: 0117 3941178

Word count:

Abstract: 197

Review: 6716

Acknowledgements: 27

Figure Legends: 1 = 40; 2 = 9; 3 = 20; 4 = 6; 5 = 658

Suggested running title:

CBSD: timeline, current knowledge and future prospects
Abstract

Cassava is the second most important staple food crop in terms of per capita calories consumed in Africa and holds potential for climate change adaptation. Unfortunately, productivity in East and Central Africa is severely constrained by two viral diseases: cassava mosaic disease (CMD) and cassava brown streak disease (CBSD). CBSD was first reported in 1936 from northeast Tanzania. For approximately seventy years CBSD was restricted to coastal East Africa and so had a relatively low impact on food security compared to CMD. However, at the turn of the 21st century CBSD re-emerged further inland, in areas around Lake Victoria and it has since spread through many East and Central African countries, causing high yield losses and jeopardising the food security of subsistence farmers. This recent re-emergence has attracted intense scientific interest, with studies shedding light on CBSD viral epidemiology, sequence diversity, host interactions and potential sources of resistance within the cassava genome. This review reflects on 80 years of CBSD research history (1936 – 2016) with a timeline of key events. We provide insights into current CBSD knowledge, management efforts and future prospects for improved understanding needed to underpin effective control and mitigation of impacts on food security.
Introduction

Cassava (*Manihot esculenta* Crantz, family *Euphorbiaceae*) produces carbohydrate rich storage roots, which are a staple food crop for approximately 800 million people worldwide (FAO 2013). In Africa, cassava is the second most important food staple in terms of *per capita* calories consumed (Nweke 2004). Storage roots are used as a fresh carbohydrate source and can also be processed into flour, which may be consumed by the grower’s family, sold in local markets or used to produce several industrial food products (Hillocks & Thresh 2002). Subsistence farmers rely on cassava for a vital energy source, as it can be planted and harvested throughout the year, tolerates periods of unpredictable droughts and grows on marginal soils (Hillocks & Thresh 2002). Recent modeling has suggested that cassava may be highly resilient to future climate change and could provide Africa with adaptation opportunities, which are not offered by other staple food crops (Jarvis et al. 2012).

Cassava was introduced into Africa from Brazil by Portuguese traders in the 16th century and subsequently integrated into local agriculture in countries across the continent (Jones 1959). Africa produces over half of global cassava (57%) (Bennett 2015), however the continent’s average fresh yield (9.9 t ha\(^{-1}\)) lags behind potential yields (15 – 40 t ha\(^{-1}\)) achieved under experimental conditions (Fermont et al. 2009). There are many reasons behind reduced yields, including restricted access to labour, poor soil quality and premature harvesting (Fermont et al. 2009). Productivity in East and Central Africa is significantly constrained by two viral diseases: cassava mosaic disease (CMD) and cassava brown
streak disease (CBSD), which together are estimated to cause annual losses worth US$1 billion (IITA 2014a) and adversely affect food security in the entire region (Patil et al. 2015).

In this paper, we review CBSD research history, highlighting key events in a timeline (Fig. 1) and provide future prospects for further understanding and effective control. The review is split into two phases according to the geographical distribution of CBSD. Phase one covers the small number (n=65) of reports published between 1936 and the early 1990s when CBSD was reported to be restricted to low altitude areas (<1000 meters above sea level (masl)) along coastal East Africa and lake shore districts of Malawi (Legg et al. 2011). Phase two examines CBSD re-emergence after the mid 1990s, when CBSD spread across East and Central Africa (Legg et al. 2011). We review the corresponding increased number (n=277) of reports on CBSD geographical expansion, viral molecular characterization, host interactions, diagnostic techniques and control efforts.

We offer insights into what can be learnt from CBSD history, in particular the need for application of knowledge to protect against and predict multiple biotic threats to staple food crops through improved understanding of CBSD epidemiology, diagnostics, surveillance and predictive modeling. This calls for effective international scientific collaborations across multiple areas of expertise and the rapid application of research and technologies to solve problems affecting farmers.
Initial emergence and symptom description (1930s – early 1990s)

The first report of CBSD from northeast Tanzania (then called Tanganyika) describe distinctive foliar symptoms on lower mature cassava leaves and rot of storage roots (Storey 1936). Nichols (1950) later reported that symptoms could be expressed on all parts of the plant and include storage root necrosis (Fig. 3A), radial root constrictions (Fig. 3B), foliar chlorosis (Fig. 3C) and occasionally brown streaks or lesions on stems (Fig. 3D).

It was noted that two main types of foliar symptoms exist: (1) feathery chlorosis along secondary vein margins, which eventually coalesce to form blotches, (2) chlorotic mottling with no veinal association (Nichols 1950). These distinctive symptoms lack the leaf distortion observed in CMD infected cassava plants. CBSD symptoms are variable in terms of severity, onset of symptom expression and parts of the plant affected, depending on the viral strain, cassava cultivar, environmental conditions and the age of the plant when infected (Nichols 1950). This variability makes diagnosis difficult for farmers (Nichols 1950) and can result in farmers being unaware that their crop is affected until they harvest storage roots (Legg & Kanju 2015). The difficulty in diagnosing CBSD has meant that infected stems have been transported to areas where CBSD has previously been absent and used for planting material. Symptom variability has also hampered epidemiology studies, as the disease can go unnoticed in an area for long periods.

The surveying and symptom scoring of infected plants across different geographical areas revealed that most plants with foliar symptoms usually also develop root necrosis (Hillocks et al. 2001). In the most
sensitive cultivars, CBSD reduces root weight by up to 70% with necrosis developing at six months post planting (Hillocks et al. 2001). Whereas, it was reported that the local Tanzanian cultivar "Nachinyaya" did not develop root necrosis and so was relatively tolerant to CBSD (Hillocks et al. 1996).

**Early geographical distribution**

Storey (1939) reported that CBSD was widespread in coastal Tanzania and by 1950, the disease was endemic across coastal areas of East Africa from northeast Kenya, Tanzania to northern Mozambique at altitudes below 1000 masl (Nichols 1950). The disease was reported in Uganda in 1945 and may have been introduced through infected cuttings sent from the Amani research station in Tanzania (Nichols 1950; Jameson 1964). Strict roguing of infected plants, replacement with non-infected planting material and quarantine appear to have prevented spread of CBSD in Uganda at this time (Nichols 1950). Significantly, a lack of plant-to-plant vector transmission at higher altitudes was reported (Nichols 1950; Jennings 1960).

**Causal agent characterization**

Storey (1936) suspected a viral causal agent, as CBSD was successfully transmitted through grafting stem cuttings. Subsequently, Lister (1959) mechanically transmitted CBSD to indicator hosts, including *Petunia hybrida*, *Datura stramonium*, *Nicotiana tabacum*, *N. rustica* and *N. glutinosa*, which produce a range of symptoms depending on the sensitivity of the host and viral variant. In 1976, sap transmission of CBSD from infected cassava material to *N. clevelandii* produced two
distinct symptom types, which suggested that two viral variants may be responsible for CBSD (Bock & Guthrie 1976).

Virus particles were identified by electron microscopy analysis of CBSD infected *N. debneyi* (Bock 1994). The infected samples contained 650nm filamentous particles with a similar morphology to viruses within the *Carlavirus* genus (Bock 1994). However, pinwheel inclusions, typical of *Potyviridae* were identified in *N. benthamiana* (Lennon *et al.* 1985). Pinwheel inclusions were subsequently found through more thorough electron microscopy of CBSD infected cassava samples, albeit at low concentrations (Were *et al.* 2004).

The *Potyviridae* sequence identity was finally confirmed in 2001 through reverse transcription PCR (RT-PCR) on CBSD infected *N. benthamiana* samples (Monger *et al.* 2001a). When the RT-PCR product was sequenced, it aligned most closely to the coat protein sequence of *Sweet potato mild mottle virus* (SPMMV, genus *Ipomovirus*, family *Potyviridae*) (Monger *et al.* 2001a). The same RT-PCR technique was used to detect CBSV in symptomless cassava leaves, highlighting the sensitivity of the RT-PCR technique (Monger *et al.* 2001a).

**Early control efforts**

In the 1930s, a cassava breeding programme was launched in Tanzania, which included breeding for CBSD and CMD resistance at the Amani research station (Jennings 1957; Nichols 1946). Early breeding to develop virus resistant cultivars involved crossing cultivated cassava with wild relatives, including *M. glaziovii*, *M. dichotoma*, *M. catingea*, *M. saxicola* and *M. melanobasis*, which are believed to have higher levels of
CBSD resistance (Jennings 1957; Kawuki et al. 2016). The breeding programme produced the *M. esculenta-M. glaziovii* hybrid known as "Namikonga" in Tanzania or "Kaleso" in Kenya, which for many years offered relatively high levels of CBSD tolerance (Hillocks & Jennings 2003; Kaweesi et al. 2014). However, "Namikonga" was not widely distributed to farmers, which may be because of its susceptibility to CMD (Hillocks & Jennings 2003; Kawuki et al. 2016).

**Initial vector transmission studies**

Until relatively recently, very little was known about vector transmission of CBSVs. It had been noted that CBSD outbreaks tended to coincide with increases in whitefly populations (Storey 1939; Hillocks & Jennings 2003). However initial attempts to transmit CBSV with whitefly (*B. tabaci*) or aphid (*Myzus persicae*) were unsuccessful (Bock 1994).

**Geographical distribution in the early 1990s**

In the early 1990s, there were reports of high CBSD incidences in areas of Tanzania, Mozambique and Malawi (Hillocks & Jennings 2003). Surveys revealed CBSD incidences reaching 36% - 50% in cassava fields along coastal areas of Tanzania (Legg & Raya 1998; Hillocks et al. 1999). Similarly, CBSD incidences in Malawi reached 75% in many fields surrounding Lake Malawi and nearly all plants inspected in northern coastal areas of Mozambique were expressing CBSD symptoms (Hillocks et al. 2002; Hillocks & Jennings 2003). In a control effort, virus-free CBSD tolerant cultivars were distributed to farmers in Mozambique who depended heavily on CBSD sensitive cassava cultivars for food security (Hillocks & Jennings 2003). CBSD was also re-discovered in
Uganda in 1994 at a site near Entebbe (Thresh et al. 1994). This led researchers to call for concerted efforts to understand CBSD through improved surveillance (Hillocks & Jennings 2003).

**Reflections on initial emergence (1930s – early 1990s)**

Despite CBSD being endemic across coastal East Africa during this period, relatively little work was done to understand and control CBSD. This is reflected by the low number of scientific papers, reports or reviews which feature CBSD published between 1936 and early 1990s (n=65) (Fig. 2). The slight increase in references to CBSD in the 1970s is due to a small number of reports (n= 27) on the threat posed by CBSD. In hindsight, these reports should have served as a warning to take control actions, which may have prevented the later expansion of CBSD across the region.

There was a general lack of scientific interest in CBSD at this time due to many factors, including the restricted occurrence of CBSD to low altitude areas along coastal eastern Africa and the devastating impacts of CMD on food security. During this period, CMD was a greater priority due to its prevalence across all cassava-growing areas of Africa, resulting in famines, higher economic losses and forcing many farmers to abandon the crop (Thresh et al. 1994; Thresh & Cooter 2005; Alabi et al. 2011). To help control the disease, CMD resistant cultivars were distributed to areas severely affected (Legg & Thresh 2000). Unfortunately these cultivars show varying levels of CBSD susceptibility (Legg et al. 2006). It is not known whether the deployment of these cultivars has contributed to the increased distribution of CBSD in the field.
Re-emergence and expansion across East and Central Africa (mid 1990s – 2016)

In 2004, the apparent restriction of CBSD to coastal lowlands changed with the re-emergence of CBSD at altitudes above 1000 masl (Alicai et al. 2007). Infections of cassava plants showing CBSD symptoms at higher altitudes in Uganda were confirmed by RT-PCR. Coat protein sequences aligned to CBSV isolates from Mozambique and Tanzania with sequence identities from 77.0 to 82.9% (Alicai et al. 2007). It is not known whether CBSD had been re-introduced to Uganda through infected cuttings or whether the disease had existed at a low level since it was first introduced in the 1940s (Alicai et al. 2007). Shortly after this first report, overall incidence of CBSD in Uganda increased from 12% in 2008 to 27% in 2011 (T. Alicai personal communication) and similar increases were reported in Tanzania and Kenya (Mware et al. 2009; Ntawuruhunga & Legg 2007; Legg et al. 2011). There have since been CBSD reports from Burundi (Bigirimana et al. 2011), Rwanda (FAO 2011), eastern Democratic Republic of Congo (Mulimbi et al. 2012), South Sudan (T. Alicai personal communication) and Mayotte Island (Roux-Cuvelier et al. 2014).

It is difficult to obtain a truly accurate estimation of the economic damage caused by CBSD, however an overall loss of US$750 million a year is estimated across Kenya, Tanzania, Uganda, and Malawi (Hillocks & Maruthi 2015). CBSD is now one of the leading causes of cassava losses in East Africa (Pennisi 2010) and its on-going spread threatens the major cassava growing areas of Central and West Africa (Legg et al. 2014).
The dramatic increase in the impact of CBSD on food security is reflected in the increase of papers, reports and reviews, which refer to CBSD published from the mid 1990s to 2016 (n= 277) (Fig. 2). The expansion of the CBSD epidemic across the Great Lakes region of East and Central Africa has necessitated the rapid development and implementation of effective control strategies. Several important projects were initiated following CBSD re-emergence, which aimed to develop research, extension and policy capacity in the countries affected. Key targets have been to breed or genetically engineer resistant cultivars, provide certified virus-clean planting material and improve viral surveillance and diagnosis (Legg et al. 2014).

**Recent local and regional CBSD epidemiology**

The reasons behind the sudden increase in CBSD incidence and geographical range remain poorly understood. Studies have shown CBSD spread and development is enhanced by high disease pressure, use of susceptible genotypes and high whitefly numbers (Katono et al. 2015). CBSD is dispersed locally and over long distances through the trade transportation of infected planting material, whereas whiteflies are only able disperse and amplify CBSD locally (McQuaid et al. 2017).

The ability of *B. tabaci* to transmit CBSV from infected to healthy plants was confirmed under quarantine insectary and glasshouse conditions in Maruthi et al. (2005). It has since been shown that CBSD viruses are transmitted semi-persistently, with whiteflies acquiring viruses in 5 – 10 minutes, retaining them for up to 48 hours, and transmitting them over relatively short distances of less than 17 meters in a cropping season.
(Maruthi et al. 2016). CBSD outbreaks occur from 3 to 12 years after increases in whitefly numbers (Legg et al. 2011). Critically, one of the primary cause for increases of both CMD and CBSD in the African Great Lakes region appears to be super-abundant numbers of whiteflies (Fig. 4), which are able to thrive at altitudes above 1000 masl (Alicai et al. 2007; Jeremiah et al. 2015).

Survey data has revealed that the transportation of infected material to areas where CBSD was previously absent has enabled the disease to spread from independent hot spots (Legg et al. 2011). This is because cassava stems used for vegetative planting material are exchanged by farmers across localities and transported over long distances. One report concluded that plants can also be infected through the use of contaminated cutting tools, which could contribute to in-field spread (Rwegasira & Chrissie 2015), however a similar study showed that such practices did not result in transmission of CBSVs (Maruthi et al. 2016).

CBSD viruses are found only in Africa and so it appears that these viruses have evolved within in East Africa on an unknown species and subsequently jumped host into cassava in a new encounter situation (Monger et al. 2010). Therefore there may be other hosts for CBSVs, which could serve as a viral inoculum sources in the field (Monger et al. 2010). CBSV has been detected in the wild perennial species M. glaziovii (Ogwok et al. 2014); the importance of this to CBSD epidemiology is not currently known.
Molecular characterisation of unusual CBSD viral genome features

CBSVs belong to the *Ipomovirus* genus of the *Potyviridae* family (Monger *et al.* 2001a). *Ipomoviruses* have positive sense, single-stranded genomes, which are translated as large polyproteins and autocatalytically cleaved by virus-encoded proteases into ten mature proteins with an additional P3N-PIPO protein produced through ribosomal frame-shifting (Valli *et al.* 2015). The genome organisation of CBSVs is shown in Fig. 5.

Molecular characterization of coat protein sequences has revealed that there are at least two genetically distinct species: *Cassava brown streak virus* (CBSV) and *Ugandan Cassava brown streak virus* (UCBSV) (Monger *et al.* 2001b; Winter *et al.* 2010), which typically have 76–78% nucleotide and 87–90% amino acid identity (Mbanzibwa *et al.* 2009b).

Genome analysis has revealed that CBSVs share unusual features (Mbanzibwa *et al.* 2009a; Monger *et al.* 2010). Firstly, CBSVs lack the multi-functional helper-component protease protein (HCPro), which possesses silencing suppressor, vector transmission and long distance movement *in planta* activities in *Potyviridae* viruses (Valli *et al.* 2015). The HCPro protein is found in all other known *Potyviridae* viruses, except for *Squash vein yellowing virus* (SqVYV) and *Cucumber vein yellowing virus* (CVYV) (Mbanzibwa *et al.* 2009a). In CBSVs, HCPro appears to have been replaced by silencing suppressor activity of the P1 serine proteinase (Mbanzibwa *et al.* 2009a). CBSV and UCBSV P1 proteins are most closely related to P1 of SPMMV and P1b of SqVYV and CVYV, which are related to the tritomovirus P1 proteins (Mbanzibwa *et al.* 2009a). The CBSV and UCBSV P1 proteins both contain zinc finger and LXKA motifs (Mbanzibwa *et al.* 2009a). The zinc finger and LXKA motifs
in CVYV P1b are essential for silencing suppressor activity through the binding and sequestering of siRNA required for silencing (Valli et al. 2008). It is therefore likely that the same motifs are responsible for the silencing suppressor activities of CBSV and UCBSV P1 proteins (Mbanzibwa et al. 2009a).

CBSV and UCBSV also encode novel Ham1 proteins with conserved Maf/Ham1 motifs (Mbanzibwa et al. 2009a). Proteins with Maf/Ham1 domains are found across prokaryotic and eukaryotic organisms and have nucleoside triphosphate pyrophosphatase activities, which reduce mutation rates by preventing the incorporation of non-canonical nucleotides into RNA and DNA (Galperin et al. 2006). The functions of CBSV and UCBSV Ham1 proteins are yet to be elucidated but they are likely to provide essential functions in the lifecycles of CBSVs. For instance, Ham1 proteins may reduce mutation rates under oxidative stress conditions in mature cassava leaves where CBSV viruses are found at the highest concentrations within the plant (Ogwok et al. 2014).

*Euphorbia ring spot virus* (EuRV, genus *Potyvirus*, family *Potyviridae*) also encodes a Ham1 protein with an uncharacterized function (Knierim et al. 2016). EuRV, CBSV and UCBSV are part of a small number of viruses which are able to infect plants in the *Euphorbiaceae* family and so perhaps Ham1 proteins are a euphorbia host adaptation (Monger et al. 2010).

**Differences between CBSVs infections and genome sequences**

In 2010, CBSV was found in infected cassava samples from Mozambique and Tanzania and UCBSV in Kenya, Uganda, Malawi and north-western Tanzania (Winter et al. 2010). However, recent
phylogenetic analysis of whole genome sequences has revealed that the viral species are not limited to agro-ecological zones and that there may be three separate species within the UCBSV clade (Ndunguru et al. 2015).

CBSV and UCBSV produce distinctly different symptoms on cassava and indicator hosts. CBSV causes more severe root necrosis and feathery chlorosis along vein margins, which develops into chlorotic blotches, whereas UCBSV causes circular chlorotic blotches between veins in cassava (Mohammed et al. 2012; Winter et al. 2010; Nichols 1950). CBSV tends to accumulate to higher titers than UCBSV in cassava (Kaweesi et al. 2014) and indicator plants (Mohammed et al. 2012; Ogwok et al. 2014).

Sequence differences between CBSV and UCBSV genomes should explain differences in symptom severities, viral loads and host-interactions observed between the two viral species. Key areas of CBSV and UCBSV genomes show relatively high levels of divergence, including the P1 and Ham1 regions, with only 59% and 47% amino acid identities respectively (Winter et al. 2010). One suggestion for the low level of Ham1 sequence similarity is that Ham1 genes may have been acquired separately by CBSV and UCBSV from a eukaryotic host (Monger et al. 2010). Alternatively CBSV and UCBSV Ham1 sequences may be derived from a common ancestor, which have diverged due to differential selection pressures on the genome sequences of the two species (Monger et al. 2010).


Evolution of CBSVs

Statistical analysis of CBSV and UCBSV genomes using the Empirical Bayes approach has predicted amino acid sites in UCBSV and CBSV coat protein and UCBSV Ham1 sequences, which appear to have been under positive selection (Mbanzibwa et al. 2011a). It is possible that positive selection at these different amino acid positions may be enabling adaptive evolution of the two viral species (Mbanzibwa et al. 2011a). Recent whole genome sequence analysis has revealed that there is a higher diversity of CBSV isolates compared to UCBSV (Alicai et al. 2016). This diversity may be enabling CBSV to rapidly adapt to overcome host resistance mechanisms, which breeders have been selecting for (Alicai et al. 2016).

Whole genome analysis has also identified putative homologous recombination sites within the genomes of CBSV and UCBSV isolates (Ndunguru et al. 2015). To date there has been no evidence for recombination between CBSV and UCBSV isolates (Mbanzibwa et al. 2011a; Ndunguru et al. 2015). However, analysis of more CBSV and UCBSV genome sequences should provide insights into the importance of recombination in CBSD viral evolution.

Potential for interactions between CBSVs

There is potential for CBSV and UCBSV isolates to interact as RT-PCR has revealed that mixed infections are common, making up 34% - 50% of tested infections in Kenya (Kathurima et al. 2016), Tanzania (Mbanzibwa et al. 2011b) and Uganda (Ogwok et al. 2014). The potential interactions between the two viral species are not currently understood. Two of the CMD causal viruses: African cassava mosaic virus (ACMV) and East
African cassava mosaic virus (EACMV) were shown to interact synergistically, leading to increased viral titers (Vanitharani et al. 2004). It is therefore possible that similar synergistic interactions occur between CBSD viral isolates.

Breeding for CBSD resistance

To date, there is no cassava cultivar with a high level of CBSD resistance available to farmers (Abaca et al. 2013). Breeding cassava is notoriously difficult due to high heterozygosity and a challenging cross-pollination process (Ceballos et al. 2012). Breeding is further complicated by cultivars showing variation in CBSD resistance across different environments, which necessitates the testing of cultivars in different agro-ecological zones to ensure their resistance is stable (Tumuhimbise et al. 2014).

Breeders and farmers across Tanzania, Kenya, Uganda and Malawi have been selecting cultivars which strongly express foliar symptoms but develop low levels of storage root necrosis (Hillocks et al. 2016). Twenty five best bet clones from five countries across East and Southern African were selected, virus-cleaned, shared and regionally evaluated across diverse environments for sources of CBSD and CMD resistance under the 5CP project (IITA 2014b). Breeding efforts also include a seven-year evaluation process of Tanzanian and Ugandan germplasm, whereby extensive intra-specific hybridizations have generated tolerant clones which develop relatively low levels of root necrosis of 12% compared to >80% in sensitive cultivars (Kawuki et al. 2016).
Although tolerant cultivars develop reduced symptoms, they remain susceptible to CBSD viruses and thereby their adoption does not remove viral inocula from the field. Therefore considerable efforts have been made to screen and breed cassava cultivars for CBSD resistance, which are able to restrict CBSD viral replication and/or movement. Promisingly, protoplast studies have recently shown that the elite breeding line KBH2006/18 line can inhibit CBSD viral replication, which offers exciting opportunities to characterize resistance and resistance-breaking viral virulence factors (Anjanappa et al. 2016).

**Responses of different cassava cultivars to CBSD**

Cassava cultivars respond very differently to infection by CBSVs; they produce a range of symptoms and are associated with varying viral loads at different time points of infection (Kaweesi et al. 2014). Sensitive cultivars show severe shoot and root symptoms, whereas cultivars with higher tolerance tend to express foliar symptoms but usually lack or exhibit mild root necrosis (Hillocks & Jennings 2003). Cultivars such as NASE 3 show high levels of resistance to UCBSV infection but remain susceptible to CBSV (Ogwok et al. 2016). It has been shown that cultivars, such as “Namikonga” support lower viral titers than susceptible cultivars, such as Albert (Maruthi et al. 2014). However, symptom severity is not always correlated with viral load, as the cultivar NASE 1 supports a relatively high viral load but produces no foliar or root necrosis symptoms, whilst the cultivar NASE 14 supports a low viral load but expresses severe root necrosis (Kaweesi et al. 2014).
This disparity between viral titers and symptom development has necessitated the use of viral load quantification during breeding to identify and select cultivars, which support low CBSD viral titers. Until recently, quantification of CBSD viruses in cassava has been based on quantitative RT-PCR, which measures the abundance of viral transcripts relative to the abundance of plant reference gene transcripts (Kaweesi et al. 2014; Moreno et al. 2011; Abarshi et al. 2012; Ogwok et al. 2014). However, the expression of plant reference genes can vary in different plant tissues, under varying developmental and environmental conditions (Brunner & Yakovlev 2004) and during viral infection (Liu et al. 2012). To overcome this, Shirima et al. (2017) have recently adapted the qRT-PCR technique to enable absolute quantification of CBSV mRNA without normalization to plant reference genes. The higher levels of accuracy offered by this technique should be valuable in breeding efforts to generate cassava cultivars, which support very low CBSD viral loads.

Identification of CBSD tolerance markers in cassava genomes

Despite the importance of cassava in developing countries it has received relatively little scientific attention when compared to maize, rice and wheat (Varshney et al. 2012). Genomic studies of cassava are now enabling the identification of genetic markers associated with tolerance within the genomes of tolerant cultivars. In 2009 the first cassava genome assembly and annotation was publicly released (Prochnik et al. 2012). Since then a large linkage map has been built using simple sequence repeats (SSR) and single nucleotide polymorphisms (SNPs) to identify quantitative trait markers associated with CBSD tolerance across diverse African farmer-preferred cultivars (Prochnik et al. 2012; Patil et
This has revealed a number of putative CBSD tolerance alleles across different genetic clusters in different cassava genotypes (Abaca et al. 2013; Ferguson 2015). If validated these alleles will be useful as markers in marker-assisted breeding and combined into cultivars for effective and durable CBSD tolerance (Pariyo et al. 2013).

**Transcriptional responses to CBSD viruses in different cassava cultivars**

To date very little is known about the function of these putative CBSD tolerance alleles. RNA-sequencing analysis of transcripts, which are overexpressed during CBSD infection of the tolerant cultivar "Namikonga" has implicated NAC transcription factors, as well as genes involved with jasmonic acid hormone signaling and the biosynthesis of phenylpropanoid, terpenoid and steroid secondary metabolites (Maruthi et al. 2014). In other plants, jasmonic acid and secondary metabolites are linked to abiotic and biotic stress responses (Wasternack & Hause 2013; Petrussa et al. 2013; Izbiańska et al. 2014).

Transcriptional studies are also helping to gain understanding into the mechanisms behind these different interactions between cassava cultivars and different CBSD viruses. Ogwok et al. (2016) recently demonstrated that Dicer like proteins (DCL): 2 and 4 and Argonaute (AGO) 2 are differentially expressed during CBSV and UCBSV infections in different cassava cultivars. DCL and AGO proteins are integral to the plant antiviral defense mechanism of silencing viral RNA (Llave 2010). Further studies are required to gain a fuller understanding how genes involved with host silencing of viral RNA are
differentially expressed in different cultivars in response to different CBSVs.

Transcriptome analysis has also revealed that beta-1,3-glucanase, which is involved with callose degradation at plasmodesmata is up-regulated during CBSD infection of the susceptible cultivar 60444 but not in the elite breeding line KBH2006/18, which shows relatively high levels of CBSD resistance (Anjanappa et al. 2017). The degradation of callose at plasmodesmata has previously been shown to promote viral movement (Zavaliev et al. 2011). Anjanappa et al. (2017) suggest that enhanced callose degradation at plasmodesmata during CBSD infection of 60444 may promote viral movement, whereas the higher amount of callose present at plasmodesmata during KBH2006/18 infection is sufficient to limit systemic viral movement and thereby restrict infection.

CBSD resistance through genetic transformation

There are promising attempts to introduce CBSD resistance into cassava through genetic engineering. The mechanism utilized involves the transgenic expression of inverted repeat CBSD viral sequences to trigger post-transcriptional gene silencing (PTGS) of the corresponding sequences during infection and hence confer viral resistance to the plant (Patil et al. 2011). The approach was successful in N. benthamiana; transgenic expression of UCBSV coat protein hairpin constructs resulted in high levels of resistance to six diverse CBSV and UCBSV isolates (Patil et al. 2011). The same construct was expressed in cassava and conferred resistance to CBSV and UCBSV under field conditions with high disease pressure (Yadav et al. 2011; Ogwok et al. 2012). Vegetative
stem cuttings taken from transgenic plants retained CBSD resistance, enabling their use in vegetative propagation (Odipio et al. 2013).

To ensure that transgenic plants are resistant to both CBSV and UCBSV viruses, the cultivar TME 204 was transformed with a construct (p5001) containing fused tandem repeat coat protein sequences from both CBSV and UCBSV to produce the transgenic line: TME 204 p5001 (Beyene et al. 2017). This transgenic line was resistant to CBSD when graft challenged (Beyene et al. 2017) and grown within confined field trials across different agro-ecological locations in Uganda and Kenya, where plants were exposed to a range of both CBSV and UCBSV isolates over multiple vegetative propagation cycles (Wagaba et al. 2017).

It is vitally important that improved cultivars are resistant to both CBSD and CMD. Transgenic CBSD resistance was conferred to cultivars TME 7 and TME 204, which are naturally CMD resistant due to the presence of the single dominant CMD2 resistance locus (Vanderschuren et al. 2012; Beyene et al. 2016). Critically however, these TME cultivars lost their CMD2 resistance through an unknown mechanism during somatic embryogenesis (Beyene et al. 2016). Work is ongoing to cross the CBSD resistant transgenic line TME 204 p5001 with a wild type CMD2 type cultivar to combine durable CBSD and CMD resistance into a single cultivar (Beyene et al. 2017).

Once biosafety issues have been addressed, the potential benefits of GM cassava to smallholder farmers are substantial. It was estimated that net value for the release of CBSD resistant cultivars would be US$436 million for western Kenya and US$790 million for Uganda over 35 year period starting in 2025 (Taylor et al. 2016). The Virus Resistant Cassava for Africa (VIRCA Plus) project is working to deliver CMD and CBSD
resistant cassava cultivars to smallholder farmers in Uganda and Kenya and so improve their livelihoods and food security (Taylor et al. 2016).

**Distribution of certified virus-clean planting material**

The lack of cultivars highly resistant to CBSD makes the existence of clean seed system critical for effective management of CBSD. Clean cassava seed systems are non-existent in most eastern Africa countries where CBSD is a problem. The Great Lakes Cassava Initiative was launched in 2008 with an overall goal to distribute certified virus-clean CBSD tolerant cultivars to 1.15 million farmers across six East and Central African countries over a four year period (Catholic Relief Services 2010). As tolerant cultivars still retain viruses within their stems, planting material must be subjected to a cleaning process and highly sensitive diagnostic testing before it can be multiplied and supplied to farmers. This should reduce disease pressure in affected areas, as at least initially the majority of crops will be disease-free (Mwangangi 2014). The production of certified virus-clean cassava germplasm is particularly important during the transportation of vegetative planting material due to the risks CBSD poses to cassava growing areas which are currently unaffected (Legg et al. 2011). The cleaning process involves culturing meristem tissue *in vitro*, and subjecting it to thermo and/or chemotherapy, which inactivates viruses and prevents viral replication or movement within tissues.

Mathematical modeling has shown that in order for the clean seed system to be sustainable, multiplication sites should only be set up in areas with low-disease pressure and low vector population density (McQuaid et al. 2015). Modeling has also shown that to reduce CBSD
dispersal and increase cassava yields, virus-free planting material should be distributed to a number of different growers across a widespread area with restricted trade (McQuaid et al. 2017). Once certified virus-clean material has been distributed, farmers must also be thoroughly trained in the identification of disease symptoms to enable sufficient roguing to further reduce CBSD spread (McQuaid et al. 2015; Legg et al. 2017). Cassava clean seed system projects have recently been piloted in Uganda, Tanzania and Nigeria. It is hoped that similar systems will be established and effective across other African countries, including Kenya (CSS Report 2016).

**CBSVs diagnostics**

As many CBSD infected plants remain symptomless, highly sensitive diagnostic techniques are required in the production and transportation of material (Abarshi et al. 2010). There have been several important advancements in cassava disease diagnostic techniques, including the optimization of RT-PCR to enable reliable simultaneous detection of CBSV and UCBSV (Mbanzibwa et al. 2011b), as well as cassava mosaic begomoviruses in a single multiplex RT-PCR reaction (Abarshi et al. 2012b). Next generation high through-put sequencing (NGS) has been used to screen large numbers of plants for the presence of CBSVs to ensure it is virus-free before dissemination as planting material. Adams et al. (2013) demonstrated that with NGS it was possible to detect 1% infected plants out of a total of 300 plants with 95% probability. Although useful tools, to date many of these techniques are relatively resource intensive and so it is vitally important that affordable diagnostic tools are available in African countries to enable sensitive CBSD detection locally,
even in cassava fields. One promising technique is reverse transcription loop-mediated isothermal amplification (RT-LAMP), which is able to detect and differentiate the presence of CBSV and UCBSV viruses with lower consumables, resources and instrument costs than RT-PCR (Tomlinson et al. 2013).

Conclusions
In the past 20 years, CBSD has become a major cause of food insecurity across East and Central Africa and only since its recent geographic expansion has the disease received the scientific interest it deserves. Once the CBSD pandemic unraveled, it was largely too late to restrict the disease to limited outbreak areas. Lessons must be learnt from this to prevent similar disease outbreaks in the future. Critically, that scientific interest should be applied to predicting and preventing future outbreaks before they are able to emerge and cause devastating yield losses across large areas.

In terms of understanding CBSD, recent studies have begun to show that CBSVs are diverse and that CBSV has a high evolutionary capacity (Alicai et al. 2016). Many control efforts are being aided by advancing molecular techniques, including marker-assisted breeding, development of genetically modified resistant lines, provision of certified virus-clean planting material and the use of sensitive diagnostics. Despite this progress, there are still many areas of CBSD biology and epidemiology, which remain poorly understood and offer opportunities to further understand and control the disease.
Future prospects

Understand key drivers in CBSD epidemiology

Relatively little is known about the complex interactions between viral variants, vectors, cassava cultivars and environmental conditions and how they may be influencing the spread of CBSD. Therefore CBSD incidence, prevalence and whitefly populations in farmers' fields need to be regularly monitored in major cassava producing areas to track periodic changes in the general status of the disease in affected countries and those at risk. Where control interventions are deployed they should be evaluated for their impact in controlling CBSD. Availability of the above information is required in development of predictive models that will provide an evidence base for disease control decisions and resource allocation. The effectiveness of CBSD control strategies also heavily depends on the level of farmer engagement and awareness. In Uganda, extension work includes efforts to raise farmer awareness of CBSD and deliver information on its management (Kumakech et al. 2013).

Gain insights into viral populations

We currently know very little about viral populations within wild hosts, which may serve as important sources of viral inoculum and enable the evolution of CBSD and other emerging viral diseases. Next generation deep sequencing can be used to detect viral populations of which very little sequence information is known (Prabha et al. 2013). It would be fascinating to apply this to cassava and characterize viral populations within CBSD infected cassava and wild hosts surrounding cassava crops.
This could shed light on viral evolution and the contribution of wild hosts in epidemiology. It may also help to identify potential unknown viral diseases, against which preemptive control could be taken in anticipation of emerging diseases (Newbery et al. 2016).

**Measures to restrict CBSD spread into unaffected areas**

To date, CBSD viruses are only found in East and Central Africa. However, CBSD distribution could increase should infected material be transported to other cassava growing areas of Africa, Latin America and Asia, which would result in huge economic losses and food insecurity (J. Legg et al. 2014b). Therefore movement of cassava material from CBSD affected countries should be subject to strict quarantine measures to ensure that planting material is virus-free before transportation. Such measures will facilitate movement of superior cultivars for production or breeding purposes.

**Utilize diverse cultivars for genomic resources**

It is important to continue to maintain and investigate diverse cassava germplasm from across Africa and Latin America and their wild relatives for potential sources of disease resistance and other beneficial agronomic traits (Turyagyenda et al. 2012). This will enable farmers to adapt to changing environmental, socio-cultural and market conditions (Pautasso et al. 2013).

**Surveillance of viral diseases**

To target control efforts, it is vitally important to accurately survey viral disease distribution. The IITA has recently launched the Cassava
Disease Surveillance platform in Nigeria, which offers opportunities for cassava breeders and extension workers to upload images of plants suspected to be infected with CBSD and other diseases. The images are analysed by a team of experts to enable rapid diagnosis and coordination of emergency control responses (IITA 2016). Similarly the West African Virus Epidemiology project launched in 2015 aims to use field surveys to gain a clear understanding of the viruses which affect cassava in West Africa to predict viral emergence and inform policy decisions. Structured surveys under the Cassava Virus Diagnostics project in eastern and southern Africa are tracking area-wide changes in cassava viral diseases over time. This will provide the basis for disease control intervention decision-making and impact assessment.

**Predicted affects of climate change on cassava production**

Cassava demonstrates relatively high levels of resilience to temperature and rainfall fluctuations predicted in climate change models (El-Sharkawy 2004). A model based on temperature and rainfall projections across Africa has predicted that compared to other staple food crops, overall cassava is the least likely to be adversely affected by climate change (Jarvis *et al.* 2012). This makes cassava an attractive food security crop for climate change adaptation in Africa. However, climate change is also predicted to affect the distribution and abundance of cassava pests and diseases, including *B. tabaci* (Jarvis *et al.* 2012). Recent ecological niche modeling has predicted that with climate change, the potential distribution of CBSD and CMD carrying *B. tabaci* will spread over West, Central and the south-western coast of Africa where cassava production is high and CBSD is currently absent (Herrera Campo *et al.* 2011). Therefore
monitoring and controlling *B. tabaci* populations is a major priority. The deep sequencing technique could be extended to *B. tabaci*, enabling the mapping of the most active and abundant viral species carried by *B. tabaci* populations across different agricultural regions (Ng *et al.* 2011).

**Understanding CBSVs infection mechanisms and virulence determinants**

Despite the increasing number of sequenced, CBSVs genomes, there is very little is known about virulence determinants within CBSV and UCBSV genomes responsible for key functions during infection and their effect on disease symptomatology. To date, only the silencing suppression activity of the UCBSV P1 protein has been characterized (Mbanzibwa *et al.* 2009a). The construction of infectious clones will enable the targeted mutagenesis of key viral sequences to identify the functions of viral proteins and the host proteins they interact with, which should serve as potential targets to restrict viral infection. Current work to develop and manipulate CBSVs infectious clones is ongoing at various institutions.

**Collaborative sharing of information and resources**

There are many opportunities to exploit recent progress made in understanding CBSD through progress in cassava, viral and vector research. There is a need for this research to be integrated into a central, easily accessible platform (Ayling *et al.* 2012). This will require experts across diverse backgrounds and countries openly communicate, engage, share data and collaborate through networks such the Global Cassava Partnership for the 21st Century. Such partnerships should help to
generate solutions to controlling CBSD and enable cassava to fulfill its potential of feeding billions of people by 2050 (Legg et al. 2014).

**Acknowledgements**

This work was made possible by the South West Training Partnership, funded by Biotechnology and Biological Sciences Research Council, UK.

All photos were taken by Katie Tomlinson.

**Conflict of interest**

The authors declare that they have no conflicts of interest with the contents of this article.
References

Abaca, A. et al., 2013. Genetic relationships of cassava genotypes that are susceptible or tolerant to Cassava brown streak disease in Uganda. *Journal of Agricultural Science*, 5(7).


Alicai, T. et al., 2016. Characterization by next generation sequencing reveals the molecular mechanisms driving the faster evolutionary rate of Cassava brown streak virus compared with Ugandan cassava brown streak virus. *Scientific Reports*, 6, p.36164.


Clean Seed System Report, 2016. Commercializing clean cassava planting material delivery system in Uganda. The second annual review and planning meeting held at the National Crops Resources Research Institute: 28th - 30th June 2016.


utilization, Wallingford: CABI.


IITA, 2014b. IITA Report. 5CP Updates: Highlights of the progress of the new cassava varieties and clean seed to combat CBSD and CMD project (5CP).


Mbanzibwa, D.R. *et al.*, 2011b. Simultaneous virus-specific detection of the two Cassava brown streak-associated viruses by RT-PCR reveals wide


Mwangangi, M.M., 2014. Evaluation of meristem tip culture, chemotherapy and thermotherapy on the reduction of *Cassava brown streak virus* in infected cassava (Guzo variety) cuttings. *Master of Science in Biotechnolog thesis submitted to Tomo Kenyatta University of Agriculture and Technology*.


Newbery, F., Qi, A. & Fitt, B.D., 2016. Modelling impacts of climate change on arable crop diseases: progress, challenges and applications. *Current*


Ogwok, E. et al., 2016. Comparative analysis of virus-derived small RNAs within cassava (Manihot esculenta Crantz) infected with Cassava brown streak viruses. Virus Research, 215, pp.1–11.


9498.


Figure 1: Key events in CBSD geographical distribution and research history (1936 – 2016).

Figure 2: Cumulative number of scientific papers, reports or reviews which refer to CBSD, published in each decade between 1936 and 2016; dramatic increase in publications from the mid 1990s following the re-emergence of CBSD (accessed on Google scholar in December 2016).

Figure 3: CBSD storage root necrosis (A), radial root constrictions (B), foliar chlorosis (C) and brown streaks or lesions on stems (D).

Figure 4: Super-abundant whiteflies on cassava in Uganda.

Figure 5: CBSVs genomes encode a large polyprotein which is auto-catalytically cleaved at specific cleavage sequences by virus-encoded proteases into ten mature proteins and an additional P3N-PIPO protein is produced through a +2 ribosomal frameshift in the P3 region (Valli et al. 2015). CBSVs proteins are: P1
Proof

39

= serine protease/silencing suppressor, P3 = third protein, PIPO = pretty interesting Potyviridae ORF, 6K1 and 6K2 = 6-kDa proteins, CI = cylindrical inclusion protein, Vpg = viral genome-linked protein, NIa-Pro = main viral protease, NIb = viral RNA dependent RNA polymerase, Ham1 = putative pyrophosphatase, CP = coat protein. Note unusual features: presence of single P1 protein, absence of HC-Pro and presence of novel Ham1 protein (Mbanzibwa et al. 2009a).
Figure 1: Key events in CBSD geographical distribution and research history (1936 – 2016).

254x179mm (72 x 72 DPI)
Figure 2: Cumulative number of scientific papers, reports or reviews which refer to CBSD, published in each decade between 1936 and 2016; dramatic increase in publications from the mid 1990s following the re-emergence of CBSD (accessed on Google scholar in December 2016).
Figure 3: CBSD storage root necrosis (A), radial root constrictions (B), foliar chlorosis (C) and brown streaks or lesions on stems (D).

167x129mm (72 x 72 DPI)
Figure 4: Super-abundant whiteflies on cassava in Uganda.

114x88mm (72 x 72 DPI)
Figure 5: CBSVs genomes encode a large polyprotein which is auto-catalytically cleaved at specific cleavage sequences by virus-encoded proteases into ten mature proteins and an additional P3N-PIPO protein is produced through a +2 ribosomal frameshift in the P3 region (Valli et al. 2015). CBSVs proteins are: P1 = serine protease/silencing suppressor, P3 = third protein, PIPO = pretty interesting Potyviridae ORF, 6K1 and 6K2 = 6-kDa proteins, CI = cylindrical inclusion protein, Vpg = viral genome-linked protein, NIa-Pro = main viral protease, NIb = viral RNA dependent RNA polymerase, Ham1 = putative pyrophosphatase, CP = coat protein. Note unusual features: presence of single P1 protein, absence of HC-Pro and presence of novel Ham1 protein (Mbanzibwa et al. 2009a).