

University of Helsinki



Faculty of Agriculture and Forestry



Disease management, especially viruses in potato and sweetpotato

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Most important food crops:

Maize	840 milj. tn
Rice	696
Wheat	654
Potato	324
Cassava	230
Barley	123
Sweetpotato	108

FAOSTAT 2010



Over 1500 viruses detected in plants so far .

Potato

Sweetpotato

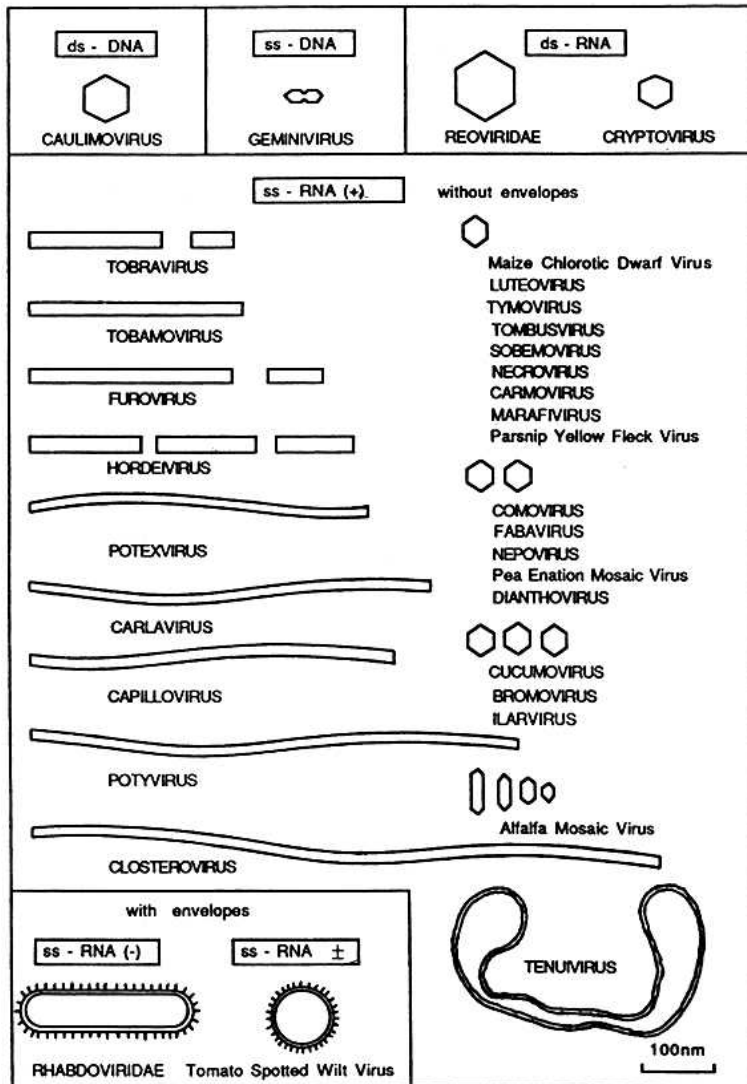


Figure 5.1 The families and groups of viruses infecting plants. Outline diagrams are drawn approximately to scale. (Courtesy R. I. B. Francki.) One additional group, the commelina yellow mottle virus group with dsDNA and bacilliform particles, was approved by the I.C.T.V. in August 1990.

Alfalfa mosaic virus (AMV), *Alfamovirus*, *Bromoviridae*
 Andean potato latent virus (APLV), *Tymovirus*, *Tymoviridae*
 Andean potato mottle virus (APMV), *Comovirus*, *Comoviridae*
 Arracacha virus B (AVB), tentative *Cheravirus*, *Sequiviridae*
 Beet curly top virus (BCTV), *Curtovirus*, *Geminiviridae*
 Cucumber mosaic virus (CMV), *Cucumovirus*, *Bromoviridae*
 Eggplant mottled dwarf virus (EMDV), *Nucleorhabdovirus*, *Rhabdoviridae*
 Potato aucuba mosaic virus (PAMV), *Potexvirus*, *Flexiviridae*
 Potato black ringspot virus (PBRV), *Nepovirus*, *Comoviridae*
 Potato deforming mosaic virus = Tomato yellow vein streak virus (ToYVSV), *Begomovirus*, *Geminiviridae*^d
 Potato latent virus (PotLV), *Carlavirus*, *Flexiviridae*
 Potato leafroll virus (PLRV), *Polerovirus*, *Luteoviridae*
 Potato mop-top virus (PMTV), *Pomovirus*, -
 Potato rough dwarf virus = Potato virus P, tentative *Carlavirus* ^e
 Potato virus A (PVA), *Potyvirus*, *Potyviridae*
 Potato virus M (PVM), *Carlavirus*, *Flexiviridae*
 Potato virus S (PVS), *Carlavirus*, *Flexiviridae*
 Potato virus T (PVT), *Trichovirus*, *Flexiviridae*
 Potato virus U (PVU), *Nepovirus*, *Comoviridae*
 Potato virus V (PVV), *Potyvirus*, *Potyviridae*
 Potato virus X (PVX), *Potexvirus*, *Flexiviridae*
 Potato virus Y (PVY), *Potyvirus*, *Potyviridae*
 Potato yellow dwarf virus (PYDV) *Nucleorhabdovirus*, *Rhabdoviridae*
 Potato yellow mosaic virus (PYMV), *Begomovirus*, *Geminiviridae*
 Potato yellow vein virus (PYVV), tentative *Crimivirus*, *Closteroviridae*
 Potato yellowing virus (PYV), tentative *Alfamovirus*^f
 Solanum apical leaf curl virus (SALCV), tentative *Begomovirus*
 Sowbane mosaic virus (SoMV), *Sobemovirus*, -
 Tobacco mosaic virus (TMV), *Tobamovirus*, -
 Tobacco necrosis virus (TNV), *Necrovirus*, *Tombusviridae*
 Tobacco rattle virus (TRV), *Tobravirus*, -
 Tobacco ringspot virus (TRSV), *Nepovirus*, *Comoviridae*
 Tobacco streak virus (TSV), *Ilarvirus*, *Bromoviridae*
 Tomato black ring virus (TBRV), *Nepovirus*, *Comoviridae*
 Tomato mosaic virus (ToMV), *Tobamovirus*, -
 Tomato mottle Taino virus (ToMoTV), *Begomovirus*, *Geminiviridae*
 Tomato spotted wilt virus (TSWV), *Tospovirus*, *Bunyaviridae*

CMV
 ICLCV
 IYVV
 SPPV
 SPCV
 SPCFV
 SPCSV
 SPFMV
 SPGVaV
 SPLV
 SPLCV
 SPLCCaV
 SPLCV-CN
 SPLCGV
 SPLCLaV
 SPLCESV
 SPLCSCV
 SPLCUV
 SPLSV
 SPMMV
 SPMSV
 SPMaV
 SPSMV-1
 SPVMV
 SPV2
 SPVC
 SPVG
 SPVCV
 SPYDV



Incidence of PLRV, PVY, PVX and PVA in seed potatoes sold at rural markets in Kenya, September 2006

Gildemacher et al. 2009. Am. J. Pot Res 86:373–382

Virus incidence levels (%)

Market	District	PLRV	PVY	PVX	PVA	Virus free	Multiple infections
Elburgon	Nakuru	29	83	39	10	8	50
Kagio	Kirinyaga	68	91	83	56	0	96
Karatina	Nyeri	91	78	83	28	1	93
Kihingo	Laikipia	71	48	100	9	0	79
Mau Narok	Nakuru	61	83	30	15	9	68
Meru	Meru Central	91	58	70	40	1	84
Molo	Nakuru	49	70	64	14	6	66
Murang'a	Muranga	95	100	64	64	0	100
Nanyuki	Meru Central	96	100	55	65	0	98
Naru Moru	Nyeri	63	29	46	65	6	75
North Kinangop	Nyandarua	99	98	34	78	0	98
South Kinangop	Nyandarua	74	94	23	65	3	83
Grand mean		74	77	57	42	3	82



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of Helsinki



SEED POTATO DEVELOPMENT PROJECT IN TANZANIA

(2012-2014)

1. Healthy planting materials established
2. Seed potato production started
3. Seed potato certification started
4. Training for the aforementioned activities
(14 Tanzanian specialists: 6 women and 8 men)

Funding: Ministry for Foreign Affairs of Finland and the Tanzanian government



http://www.placesonline.com/maps_ok/tanzania.jpg



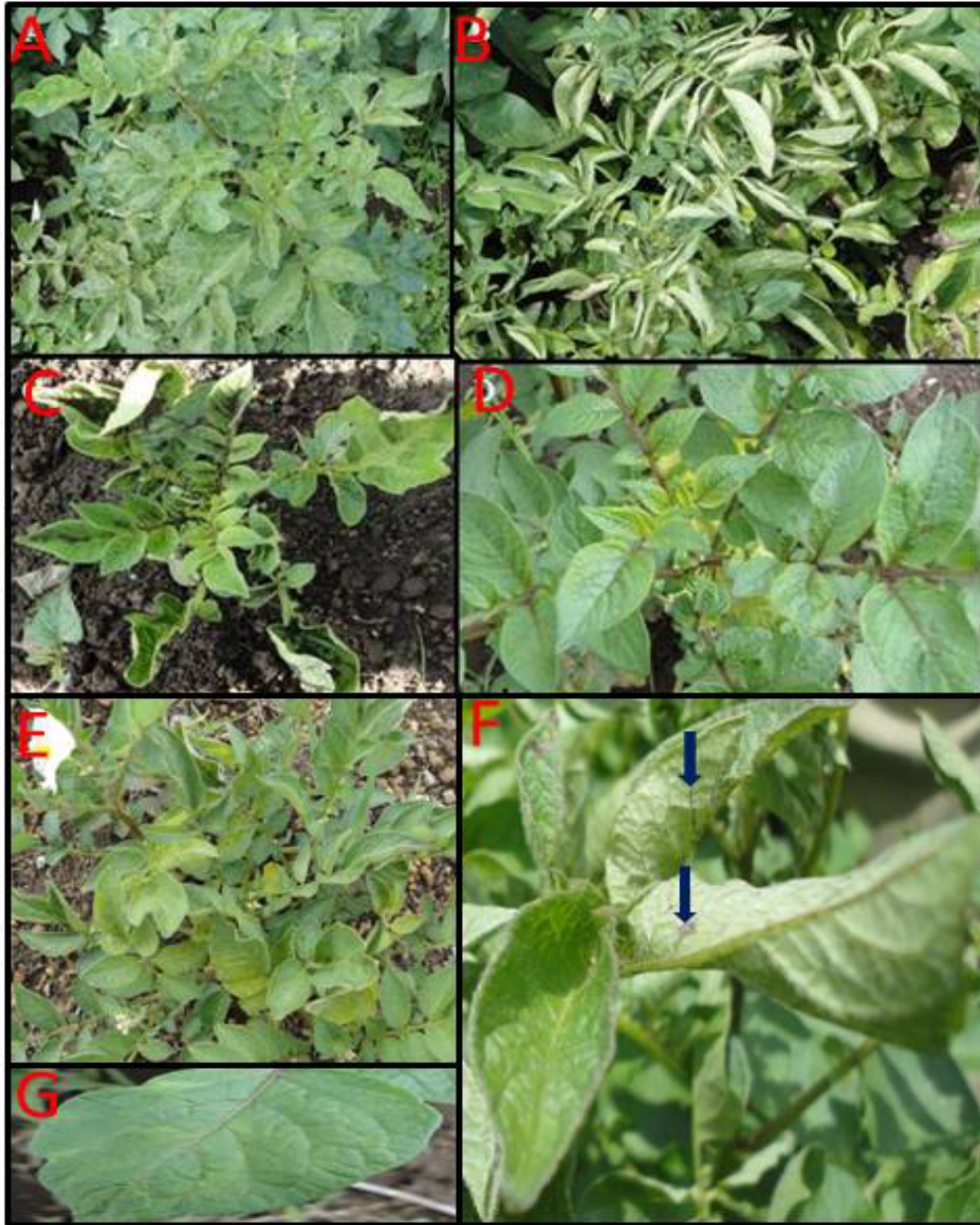
Mbeya, Tanzania



Agricultural Research Institute Uyole
Mbeya, Tanzania



© Evangelista Chiunga



Viruses occurring in potatoes in Mbeya region, Tanzania

Evangelista Chiunga

ARI-Uyole

ARI-Mikocheni

University of Helsinki

Leaf samples collected from 13 potato farmers' fields in October-December 2011

Table 2. DAS- ELISA results from collected potato leaf samples

Locality	n	PVS	PVY	PVX	PVA	PLRV	PVM
Umalila	39	19	0	6	0	2	0
Kawetele	39	30	4	0	0	25	1
Kikondo	39	15	0	6	7	21	3
Mwakareli	39	10	0	14	2	14	1
Bonde la uyole	39	30	2	2	0	16	5
ARI- Uyole	24	16	0	3	0	7	1
Σ	219	120	6	31	9	85	11
		56%	3%	14%	4%	39%	5%

n: Total number of potato leaves sampled per locality

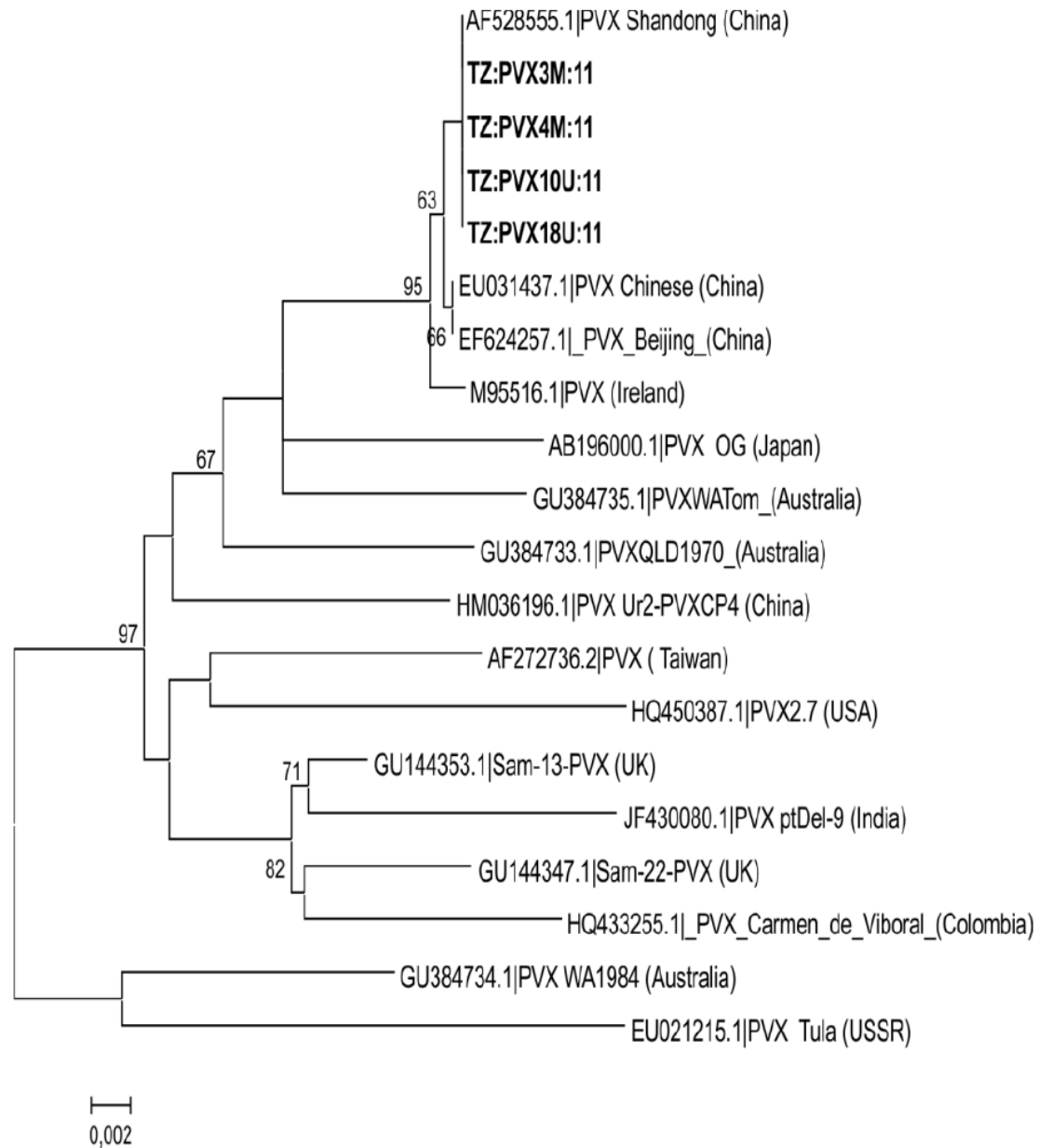
Σ : Total number of potato leaves detected positive for each virus

Evangelista Chiunga
Master's thesis
University of Helsinki

Table 3. Mixed virus infections in potato plants

	S + LR	S+X	S + X	S + Y	S + A	S + M	S + X + LR	S + M + LR	S+M+LR+A	S + X + LR + A	X + LR	X + LR + A	X + M + LR	Y + LR	M + LR	M + LR + A	LR + A
Locality																	
Umalila	1	1	2	0	0	0	0	0	0	0	1	0	0	0	0	0	1
Kawetele	16	0	0	2	0	0	0	0	0	0	0	0	0	2	1	0	0
Kikondo	9	0	0	0	1	0	1	0	0	1	0	1	1	1	0	1	0
Mwakareli	1	2	0	0	0	0	2	0	1	0	1	0	0	0	0	0	0
Uyole	5	0	0	2	0	1	0	4	0	0	2	0	0	0	0	0	0
ARI-Uyole	6	2	0	0	0	0	0	0	0	0	1	0	0	0	0	0	0
Σ	38	5	2	4	1	1	3	4	1	1	5	1	1	3	1	1	1

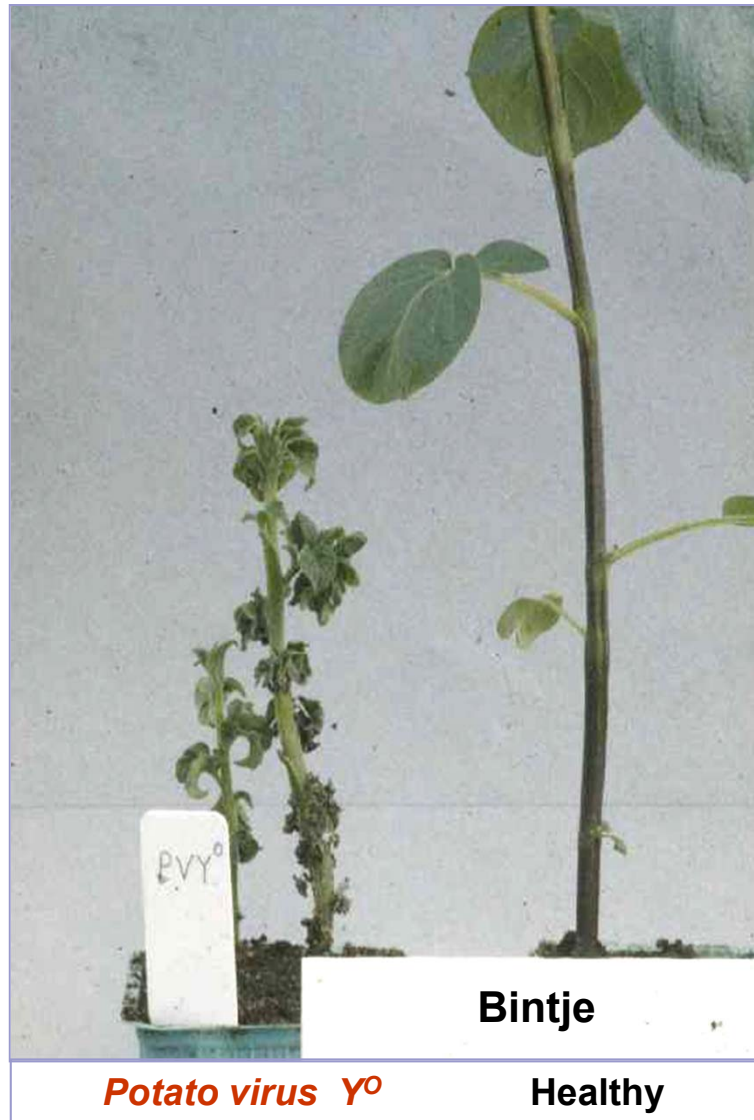
International trade of potatoes distributes viruses



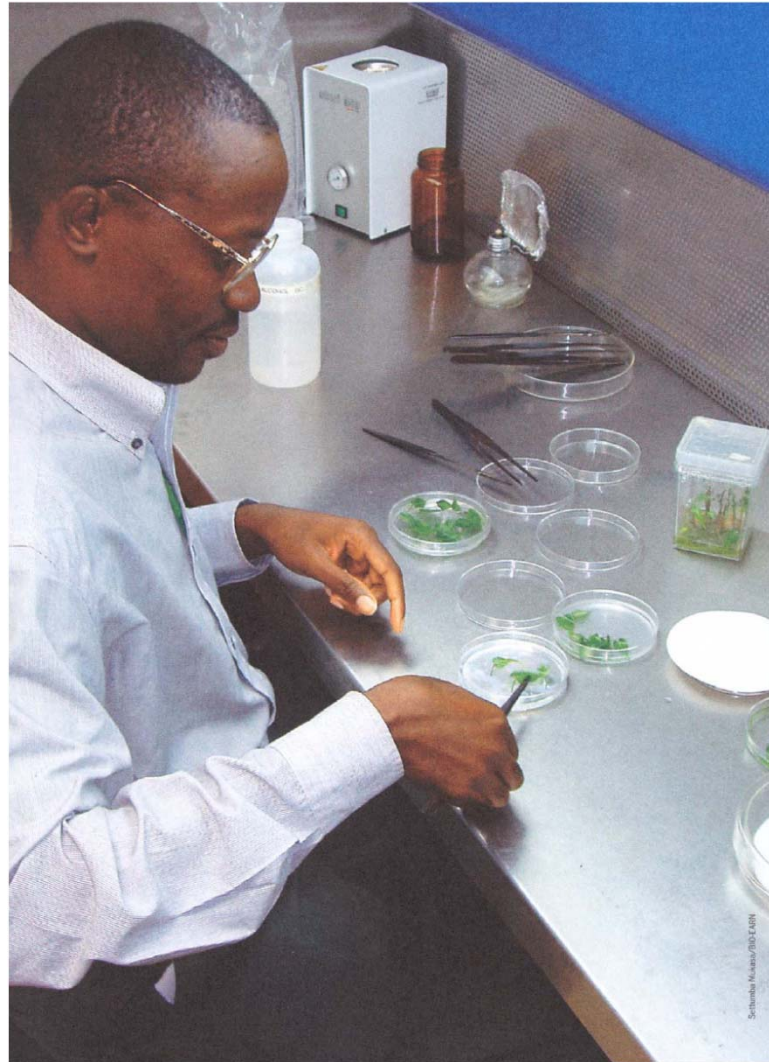
Control of virus diseases

1. Production and availability of healthy planting materials

Growth reduction caused by *Potato virus Y* strain O (PVY^O) when the plant grows from an infected seed tuber.



Production of healthy plants using tissue culture techniques



Dr. Settumba Mukasa / BIO-EARN / Makerere University, Uganda

Jari Valkonen

Production of pathogen-free clones of potato cultivars



**Regenerating
meristem**



**Virus-free shoots
propagated *in vitro***

Jari Valkonen

Photos: Jenni Kesulahti

Finnish Seed Potato Center, Tyrnävä



Finnish Seed Potato Center SPK, Tyrnävä



Finnish Food Safety Authority EVIRA, Loimaa



Control of virus diseases

**Production and availability of healthy planting materials:
practices that can be applied at the farm level**

- **Crop rotation (voluntary plants can be sources of viruses)**
- **”Select-the-Best” -practice introduced by International Potato Center (CIP): farmers mark disease-free potato plants in the field and use the tubers of them as seed.**
- **Symptomless sweetpotato vines used as planting materials**

Subsistence sweetpotato cultivation and piece-meal harvesting, Uganda



Jari Valkonen

Control of virus diseases

**Production and availability of healthy planting materials:
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- **”Select-the-Best” -practice introduced by International Potato Center (CIP): farmers mark disease-free potato plants in the field and use the tubers of them as seed.**
- **Symptomless sweetpotato vines used as planting materials**

Control of virus diseases

1. Production and availability of healthy planting materials

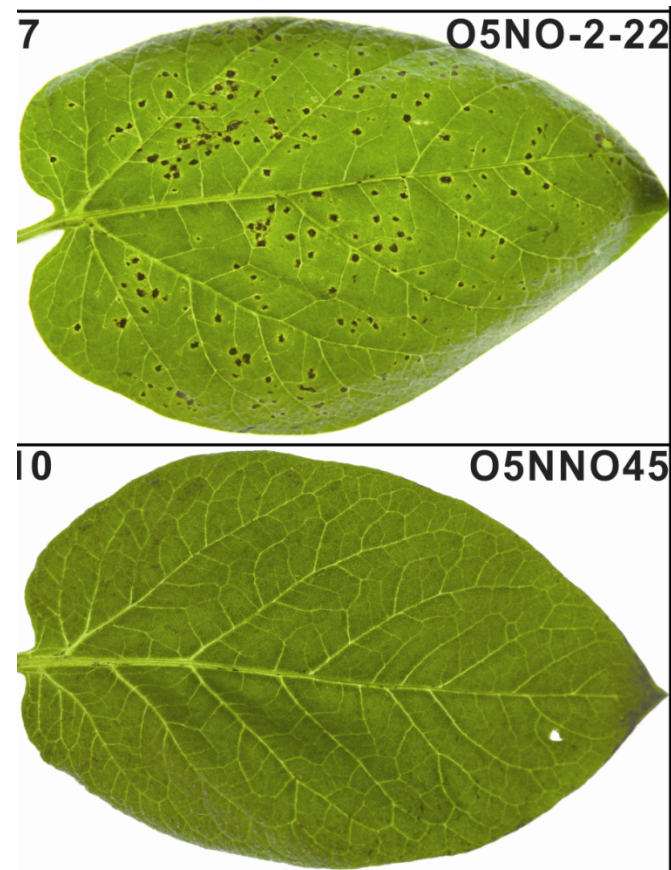
2. Use of virus-resistant cultivars

Gene-for-gene resistance

Hypersensitive, strain-specific resistance to *Potato virus Y* (PVY) in potato plants

**HR to PVY strain O is controlled by the gene *Ny* in potato:
Necrotic local lesions on the inoculated leaf.**

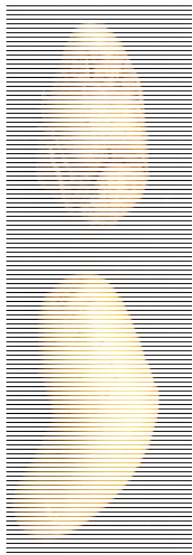
***Ny* does not recognize PVY strain N:
the leaf becomes infected and no necrotic lesions appear**



Strain N of PVY (PVY^N) overcomes the resistance conferred by *Ny*



cv. Nicola



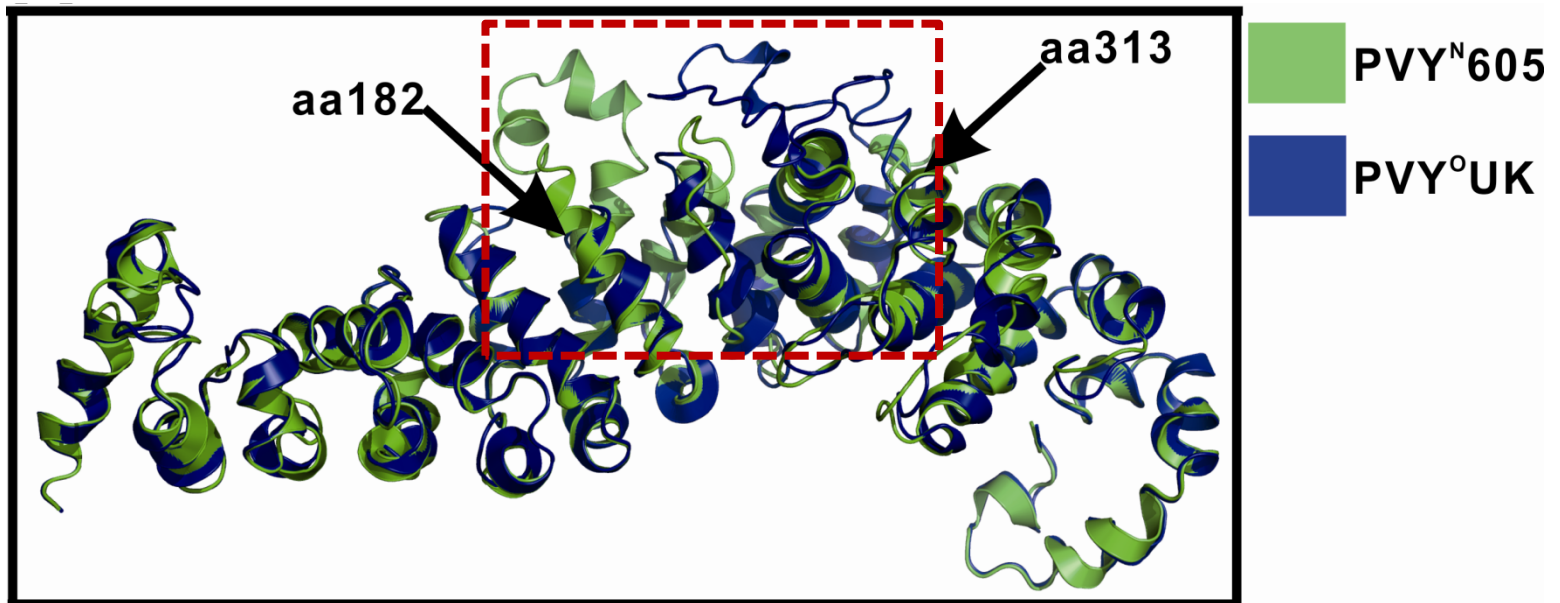
cv. Annabelle

PVY^N-NTN occurs in all potato growing areas, including Latin America, USA, Canada, Europe, Africa and Asia.

In some cultivars and under certain growth conditions, necrotic symptoms develop in tubers.

(Photos Y. Tian & J. Valkonen)

Specific recognition of PVY strain O by gene *Ny* in potato can be explained by predicted structural differences between strains O and N



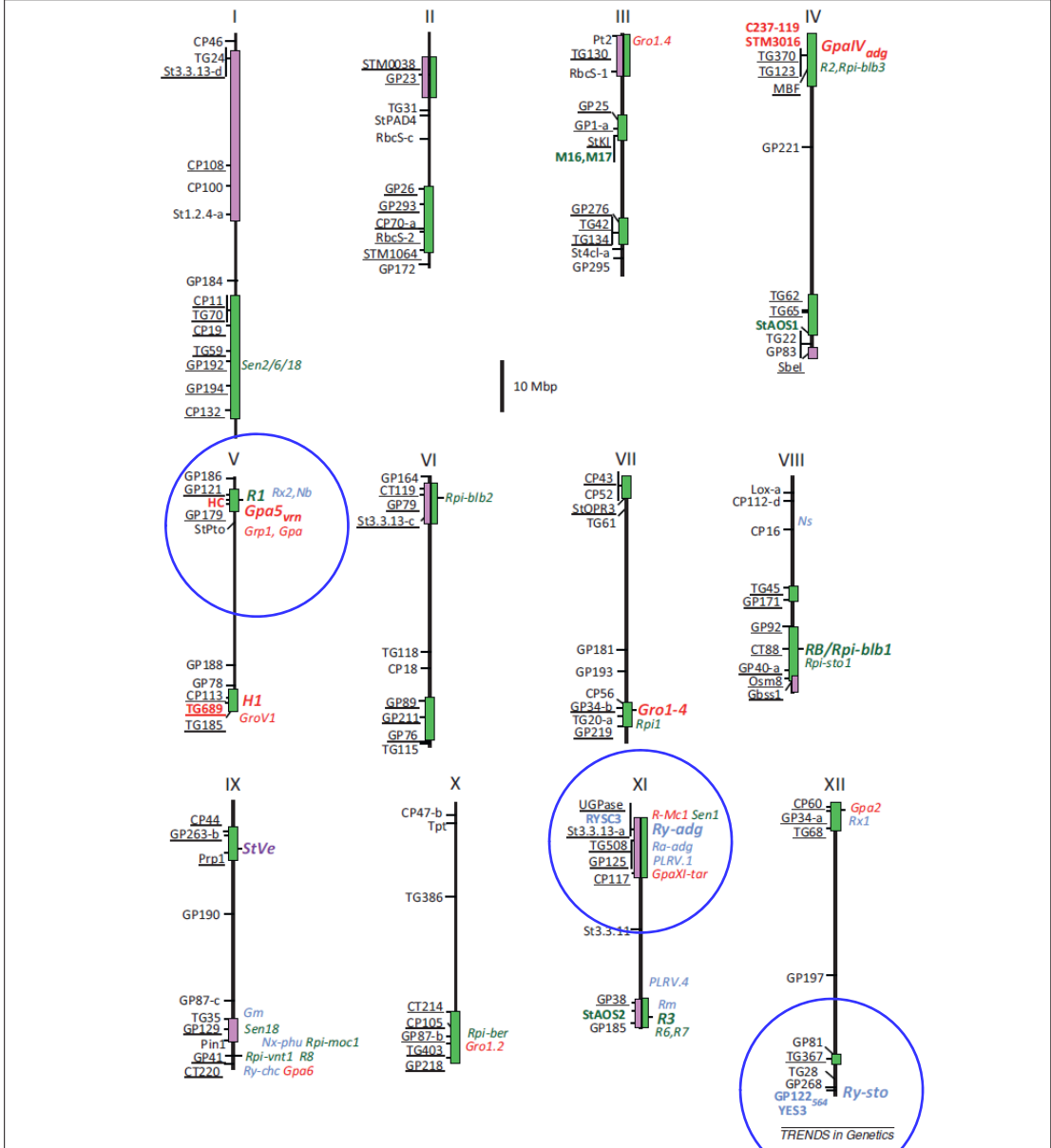
3D modeling using I-TASSER

Tian & Valkonen 2013, MPMI 26: 297–305

Virus resistance genes mapped to potato chromosomes

Extreme resistance (ER)
 (virus multiplication inhibited)

- No visible response
- Effective against all strains

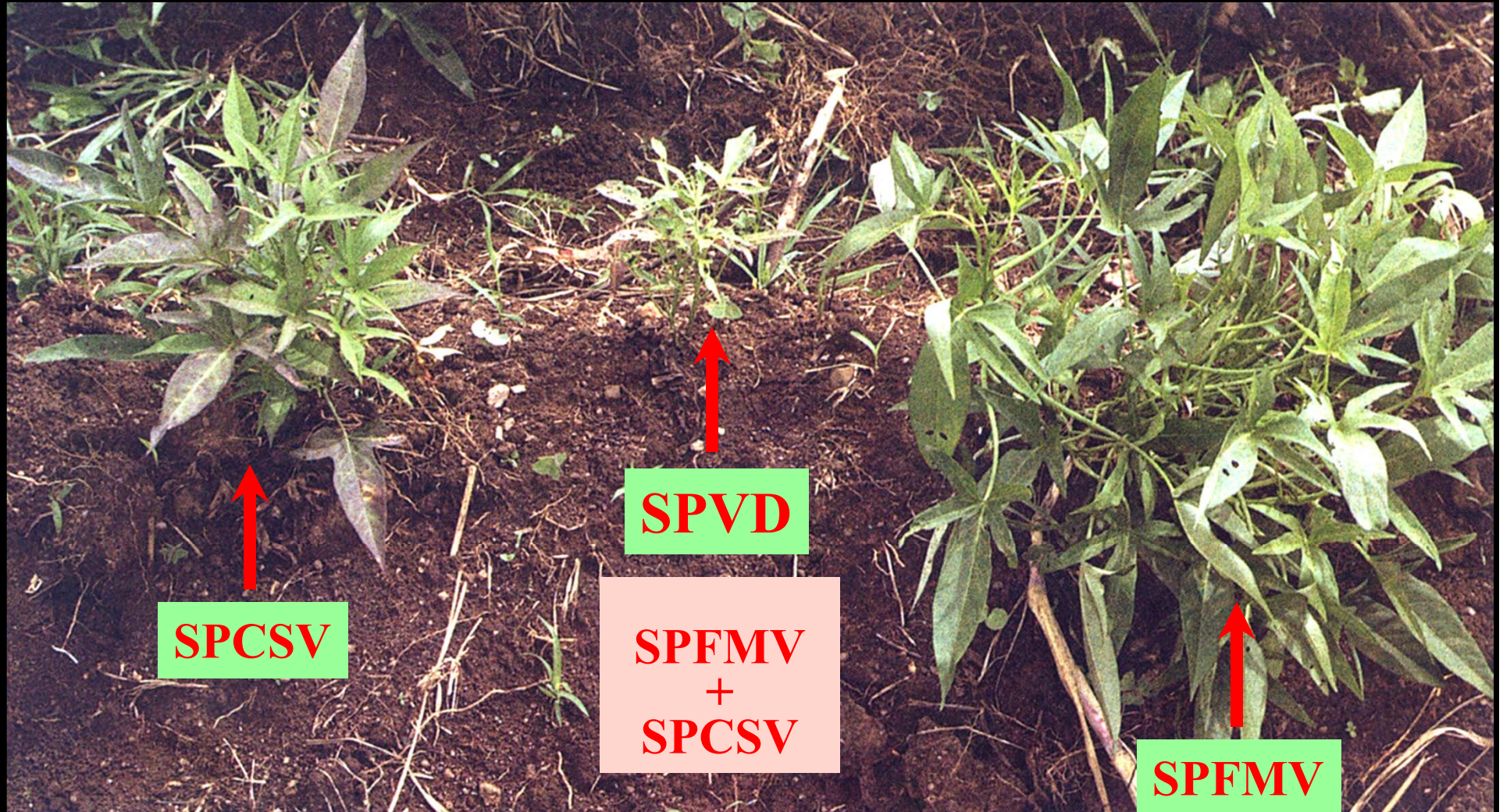


Control of virus diseases

2. Use of virus-resistant cultivars

- Durability of resistance in mixed virus infections?
Which viruses should be targeted by resistance?**

Viral synergism
Sweet Potato Virus Disease (SPVD)
- the main disease of sweetpotatoes



Karyeija et al. 2000. Virology 269: 26-36.
Gibson et al. 2004. Virus Res. 100: 115-122.

R.W.Gibson

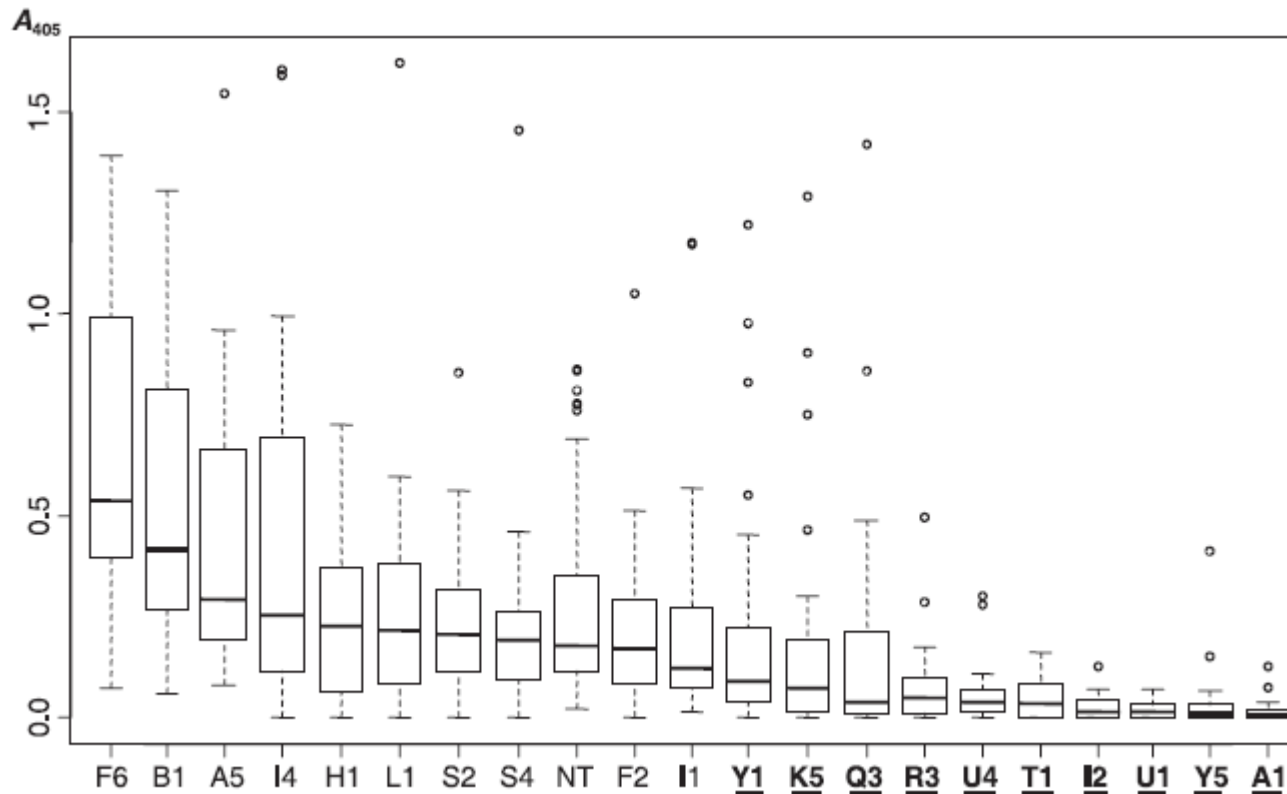


R.W.Gibson



**SPFMV
+
SPCSV
= SPVD**

Virus-derived transgenic resistance to SPCSV: Titers of SPCSV in transgenic sweetpotato plants



The resistance to SPCSV broke down following infection with SPFMV, resulting in development of SPVD

KREUZE et al. (2008) MOLECULAR PLANT PATHOLOGY 9: 589–598



Control of virus diseases

2. Use of virus-resistant cultivars

- Durability of resistance in mixed virus infections?
Which viruses should be targeted by resistance?

Transgenic virus-specific resistance based on RNA silencing is vulnerable to break-down in mixed virus infections.

Other sources of resistance ?

NASPOT varieties bred by Dr Mwanga in Uganda ?




Control of virus diseases

- 1. Production and availability of healthy planting materials**
- 2. Use of virus-resistant cultivars**
- 3. Comprehensive knowledge on viruses infecting the crop and prevailing in the cultivation environment**



**Generic methods needed for detection
of new viruses and virus strains**

**Detection of alien viruses and viroids
in plants by siRNA deep-sequencing**



RNA silencing mechanism is activated by double-stranded RNA (dsRNA)

Viruses form dsRNA

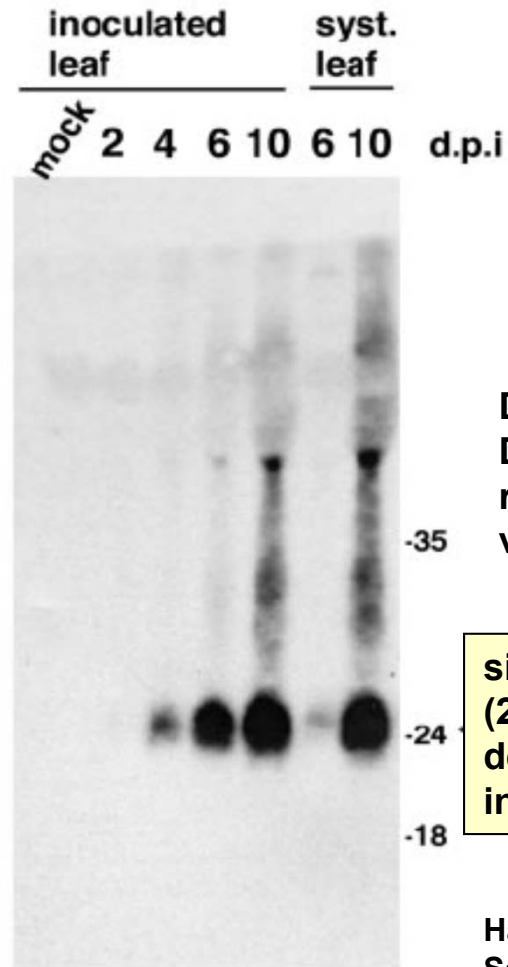
- replicative forms of RNA viruses
- ds secondary structures of single-stranded RNA virus genomes
- ds secondary structures of RNA transcripts produced by DNA viruses

RNA silencing is an antiviral defence mechanism:

**Viruses are always detected by the
RNA silencing defence mechanism in infected plants!**

Viral RNA is detected and cleaved to small RNA by the RNA silencing mechanism

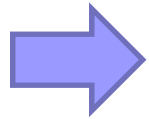
Northern blot,
virus-specific probe



Dicer-like (DCL) enzymes
DCL2 and DCL4 are
responsible for detecting
viral RNA in plants

siRNA
(21, 22 and 24 nt)
derived from Potato virus X
in an infected plant

Hamilton & Baulcombe, 1999
Science 286:950-952

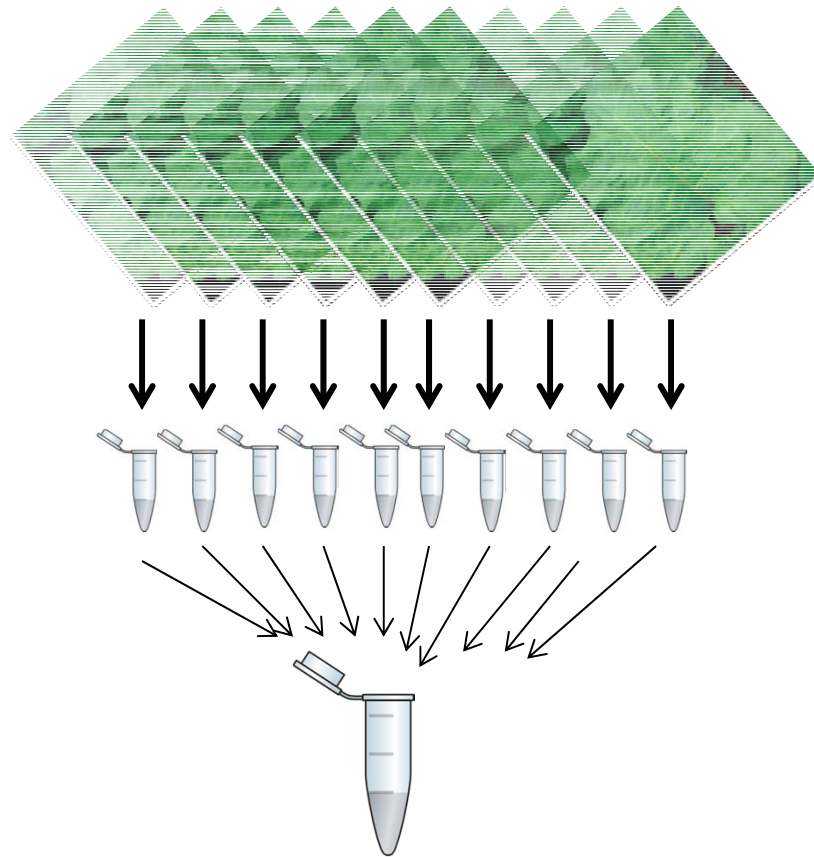


The plant detects the virus – we analyze the end products resulting from the defence response

siRNA can be isolated, sequenced, and viruses detected

Kreuze, J.F., Perez, A., Untiveros, M., Quispe, D., Fuentes, S., et al. (2009) Complete viral genome sequence and discovery of novel viruses by deep sequencing of small RNAs: a generic method for diagnosis, discovery and sequencing of viruses. *Virology* 388: 1–7.

Procedure



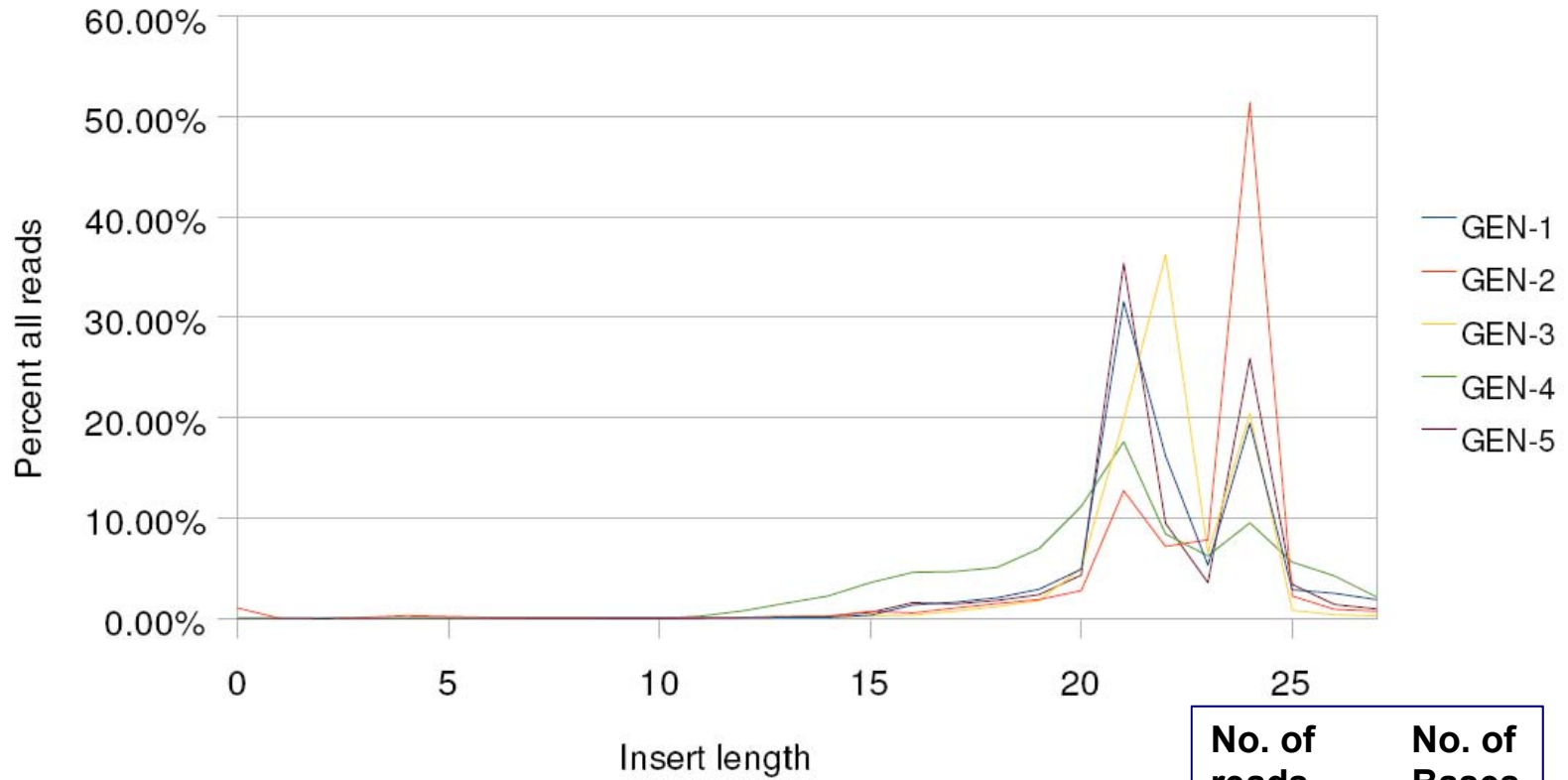
**1. Collect samples
(store in freezer)**

**2. Extract total RNA, check
quality, measure concentration,
store in freezer**

**3. Pool similar amounts of
RNA from different samples
(e.g., 50 plants)**

**The pool of RNA (5-10 μg) will be
sent to a sequencing laboratory**

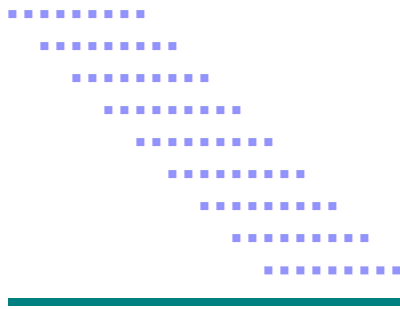
Size classes of small RNA reads



	No. of reads	No. of Bases
GEN-1	1'508'504	48'272'128
GEN-2	4'749'857	151'995'424
GEN-3	4'155'175	132'965'600
GEN-4	1'728'098	55'299'136
GEN-5	1'218'575	38'994'400

Data analysis ja utilization - A

5. Original RNA molecules or parts thereof are reconstructed from overlapping siRNA sequences:



6. Reconstructed RNA molecules are identified by comparison with sequences in public databases



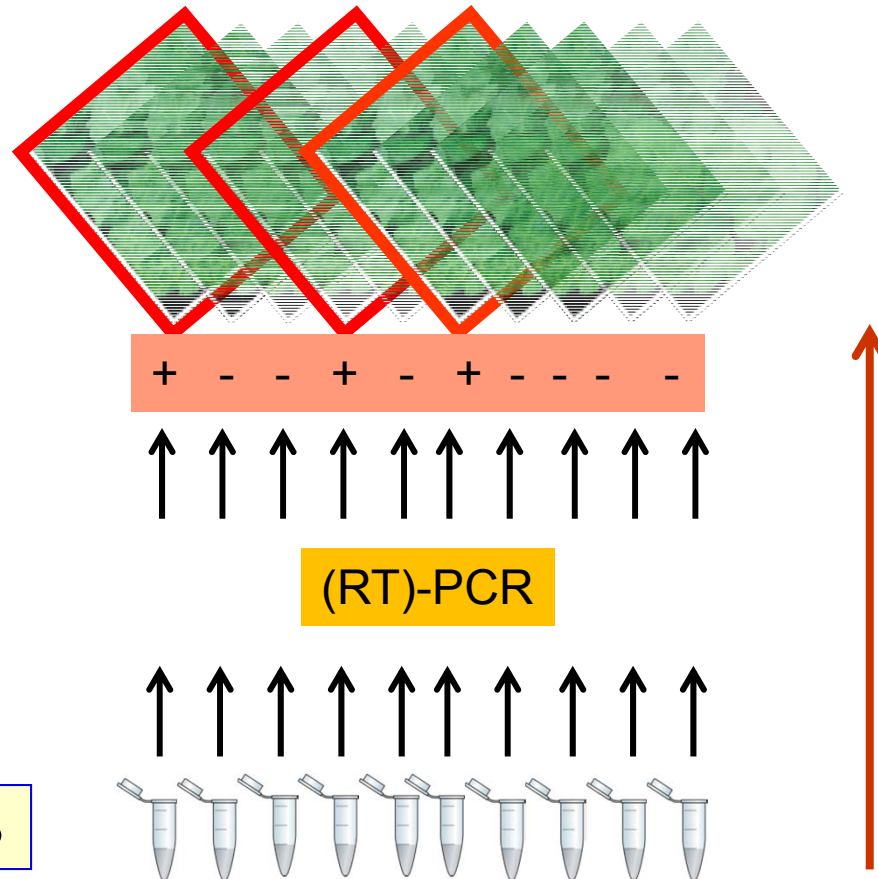
7. VIRUSES DETECTED!



8. Are the detected viruses new strains, or new viruses? Analyses can be continued by mapping siRNA reads to the genomes of viruses found in the sample (or to the closest-related viruses).

Data analysis using bioinformatics may be all what is needed

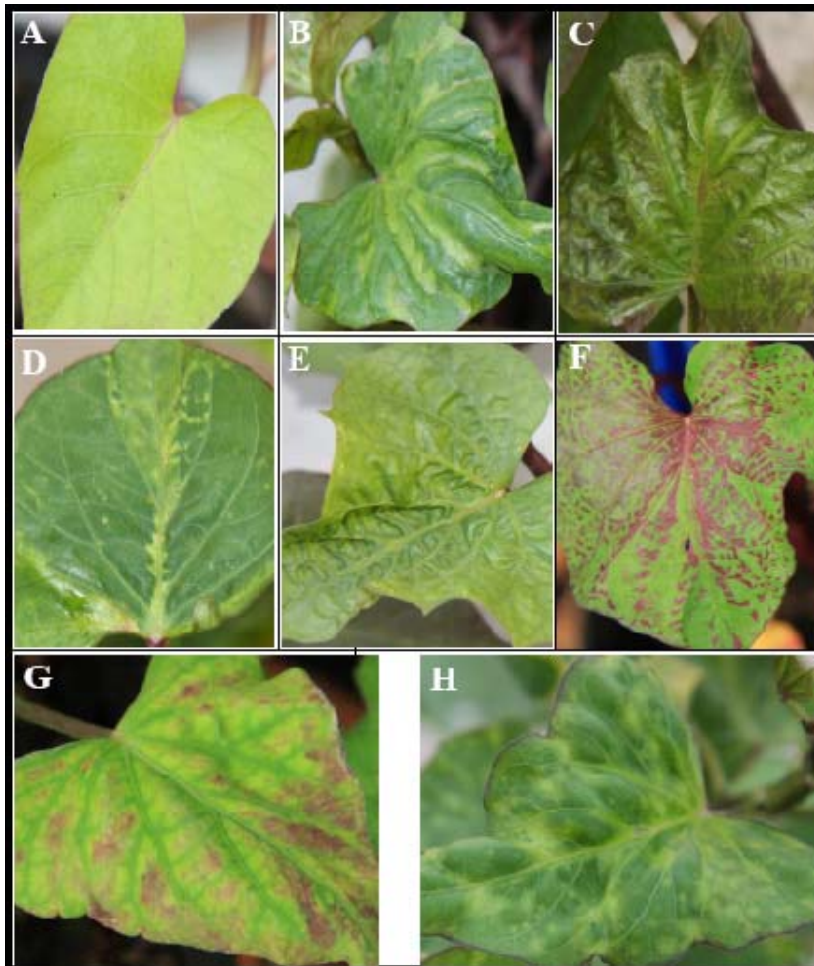
Identification of infected plants if needed



Data analysis ja utilization - B

9. PCR primer pairs designed based on the reconstructed viral genomes; original samples (RNA stored in freezer) tested by RT-PCR to detect the infected plants

SPPV-A	<i>Sweet potato pakakuy virus, strain A</i>	DNA, pararetro
SPPV-B	<i>Sweet potato pakakuy virus, strain A</i>	DNA, pararetro
SPLCGV	<i>Sweet potato leaf curl Georgia virus</i>	DNA, gemini
SPCSV-WA	<i>Sweet potato chlorotic stunt virus, strain WA</i>	RNA, crini
SPFMV	<i>Sweet potato feathery mottle virus</i>	RNA, poty
SPVC	<i>Sweet potato virus C</i>	RNA, poty



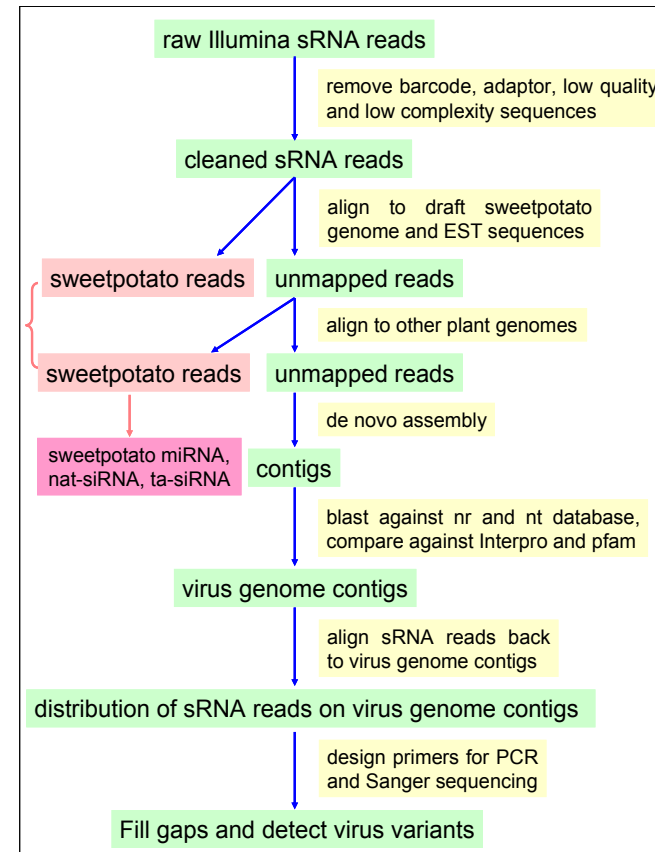
Kashif, Pietilä, Artola, Jones, Tugume, Mäkinen and Valkonen:

Detection of viruses in sweetpotatoes from Honduras and Guatemala augmented by deep-sequencing of small-RNAs.

Plant Disease 96:1430-1437 (2012)

Going to scale: the pan African sweetpotato virome

- n ~2000 samples and bioinformatics pipeline for virus identification using siRNA assembly and genome subtraction



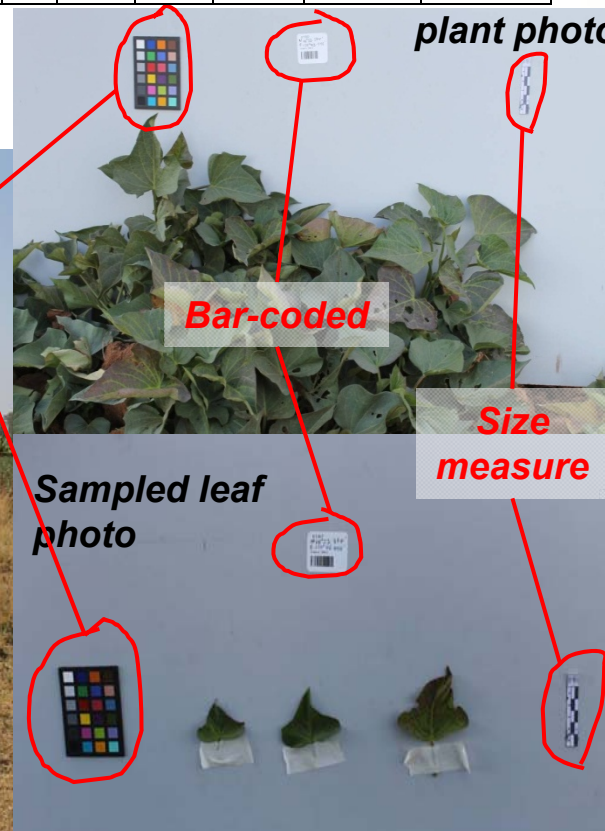
Recorded data

Sample number	Date	Region	District	Locality	Field number	GPS			Crop age (months)	Field size	Photo number field	Photo number plant	Photo number leaf	Intercrop	Cultivar/species	Notes
						Longitude	Latitude	Altitude (m)								
ET1	23/01/2012	SNNRP	Arba Minch Zuria	Zigiti Baqole	1	N06 04' 552"	E 037 27' 450"	2455	4	0.01	100-0304	100-0305	100-0306	Enset + kale	Local (Ambo)	
ET2	23/01/2012	SNNRP	Arba Minch Zuria	Zigiti Baqole	1	N06 04' 552"	E 037 27' 450"	2455	4	0.01	100-0304	100-0309	100-0310	Enset + kale	Local (Ambo)	
ET3	23/01/2012	SNNRP	Arba Minch Zuria	Zigiti Baqole	1	N06 04' 552"	E 037 27' 450"	2455	4	0.01	100-0304	100-0311	100-0312	Enset + kale	Local (Ambo)	
ET4	23/01/2012	SNNRP	Arba Minch Zuria	Zigiti Baqole	2	N06 03' 652"	E 037 27' 728"	2365	3.5	0.25	100-0313	100-0315	100-0317	Potato volunteers	Ambo	
ET5	23/01/2012	SNNRP	Arba Minch Zuria	Zigiti Baqole	2	N06 03' 652"	E 037 27' 728"	2365	3.5	0.25	100-0313	100-0318	100-0319	Potato volunteers	Tula	

Recorded data



Standard color chart for image correction



Mozambique

sample		Virus identified											
#	region	SPMM											new
		SPFMV	SPVC	SPVG	SPV2	V	SPCSV	SPCFV	SPLCV	SPPV	SPSMV		
1	Angonia											X	
2	Angonia	X										X	
3	Angonia	X	X									X	
4	Angonia	X	X		X	X	X		X	X		X	
5	Angonia	X	X	X			X		X	X		X	p
7	Angonia	X					X		X	X		X	
8	Angonia	XX	XX	X								X	p
10	Angonia	X							X	X		X	
14	Gurue												
15	Gurue	XX	XX		X	XX	X				X	X	
17	Gurue	X	X		X	XX	X		X	X		X	
18	Gurue	XX	X	X	X	XX					X	X	p
19	Gurue	X					X				X	X	
20	Gurue	XX	X	X							X	X	p
21	Gurue	X											
22	Gurue	X		X			X		X			X	
24	Gurue										X		
25	Gurue	XX	X		X	XX	X				X	X	
27	Gurue	X	X		X	X	X				X	X	
28	Maputo	XX							X	X			
31	Maputo	XX			X	XX	X				X		p
32	Maputo	XX	X	X					X	X	X		p
33	Maputo	X	X		X	X			X	X			
34	Maputo	X	XX			X	X		X	X			
35	Maputo	X					X				X		
36	Maputo	XX	XX	X		XX	XX				X		
37	Maputo	X	X	X					X	X			
39	Maputo	X	X				X				X		
41	Chockwe	XX	X	X		XX	X		X	X	X		
42	Chockwe	X	X	X		X	X		X	X			
44	Chockwe	XX	X	X		X		XX	X	X	X		
46	Chockwe	XX	X	X	X	XX	X		X	X	X		p,a
49	Chockwe	X				XX	X		X	X			
51	Chockwe	X	X	X		XX	X		X	X			p,a
55	Chockwe	XX				X	X		X	X			
66	TANZANIA	X					X		X	X			

Ghana & Burkina Faso

Virus identified

sample #	region	SPFMV	SPVC	SPCSV	SPCFV	SPLCV	SPVCV	SPPV	SPSMV	new
116	Baugonia					X		X		
23a	Bompro	X					X	X		
19a	Dompase	XX	XX	X		X				
30a	Dzogodze	XX				X		X		
104	Ejura	X				X		X	X	
102	Ejura	XX	X			X				
103	Ejura	X	XX	X				X		
105	Ejura	X		X		X		X	X	
11b	Esukyeano	X		X	XX	X		X		
12a	Esukyeano	X			X			X	X	
152b	Fumesua	X		X		X		X	X	
156a	Fumesua	X	X	X	X	X		X		
27a	Gomakarde	XX	X					X		
26a	Gomakarde	XX	X					X		
25a	Gomakarde	X		X		X		X		
133	Kamboinse	X		X		X		X	X	
137	Kamboinse	X		X		X				
140	Kamboinse	X		X		X		X	X	
135	Kamboinse	X		X				X		
13a	Komenda	XX	XX	X	X			X		
14a	Komenda	X	X					X		
15a	Komenda	XX			XX			X		
39a	Kporkuve	X		X		X		X		
6a	Krobo Kwamu	XX	XX		XX			X		
7a	Krobo Kwamu	X	X		X	X		X		
8a	Krobo Kwamu	X	X		X	X		X		
9a	Krobo Kwamu	X	X		X	X		X		
4a	Krobo Kwamu	X		X	XX	X		X		
32a	Kudzordzi Korpe	XX	XX			X		X		
31a	Kudzordzi Korpe	X				X		X		
40b	Lume			X		X		X		
41a	Lume					X		X		
43a	Lume	X				X		X		
119	Manchoro					X		X		
122	Manchoro					X			X	
123	Manchoro					X		X	X	
124	Nimbasinia					X		X		
126	Nimbasinia	XX			X	X		X		
113	Nyangua					X		X	X	
111	Nyangua	XX	X		X	X		X		
130	Tekuru					X		X		
34a	Vume	XX	XX	X		X		X		
36a	Vume	X						X		
35a	Vume	X				X		X		

→ Cotton leaf curl Gezira
alphasatellite – Burkina
Faso

→ 'Nodavirales superfamily'
s



Control of virus diseases

- 1. Production and availability of healthy planting materials**
- 2. Use of virus-resistant cultivars**



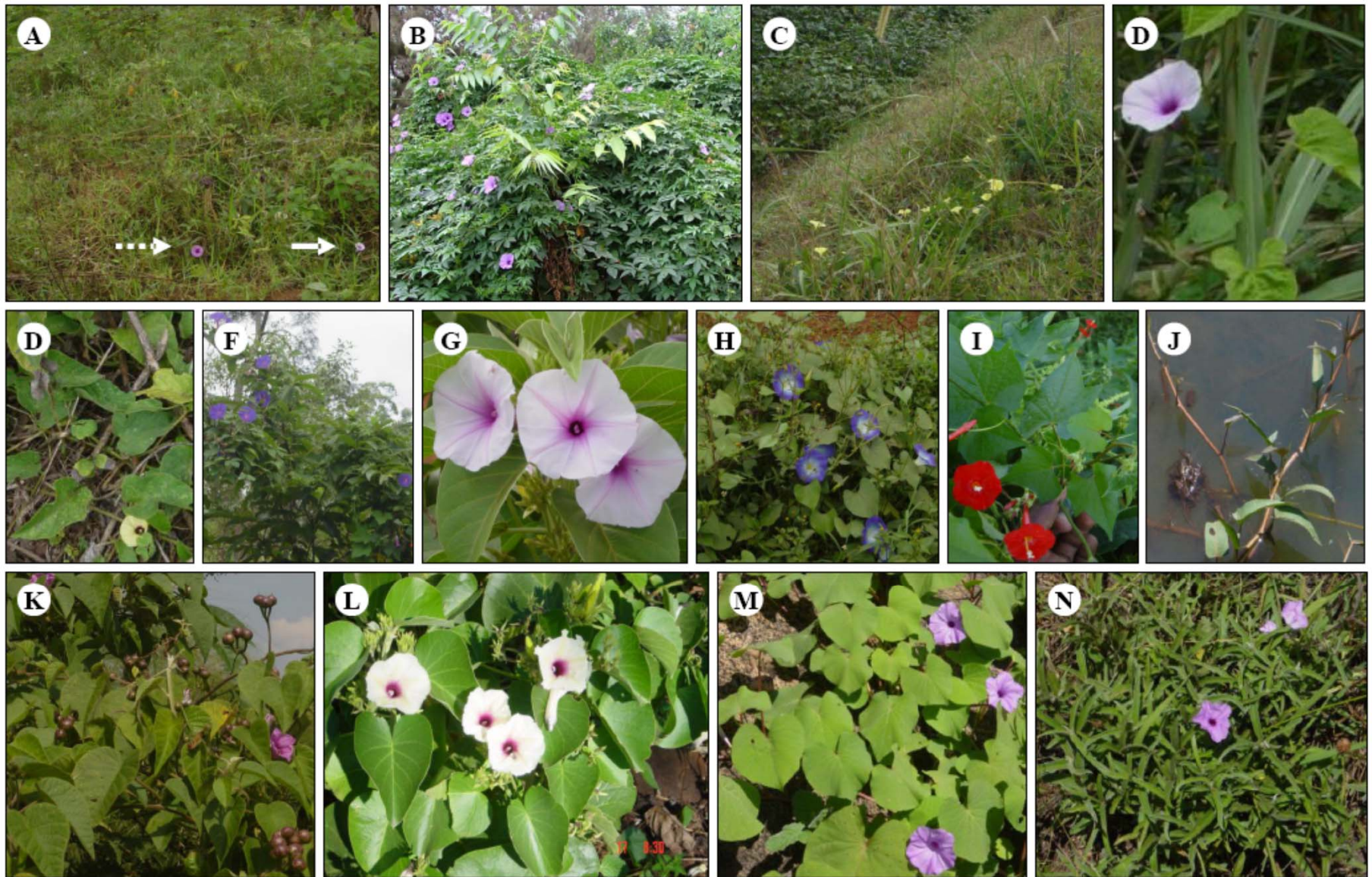
Control of virus diseases

- 1. Production and availability of healthy planting materials**
- 2. Use of virus-resistant cultivars**
- 3. Comprehensive knowledge on viruses infecting the crop and prevailing in the cultivation environment**

•Viruses infecting wild plants?



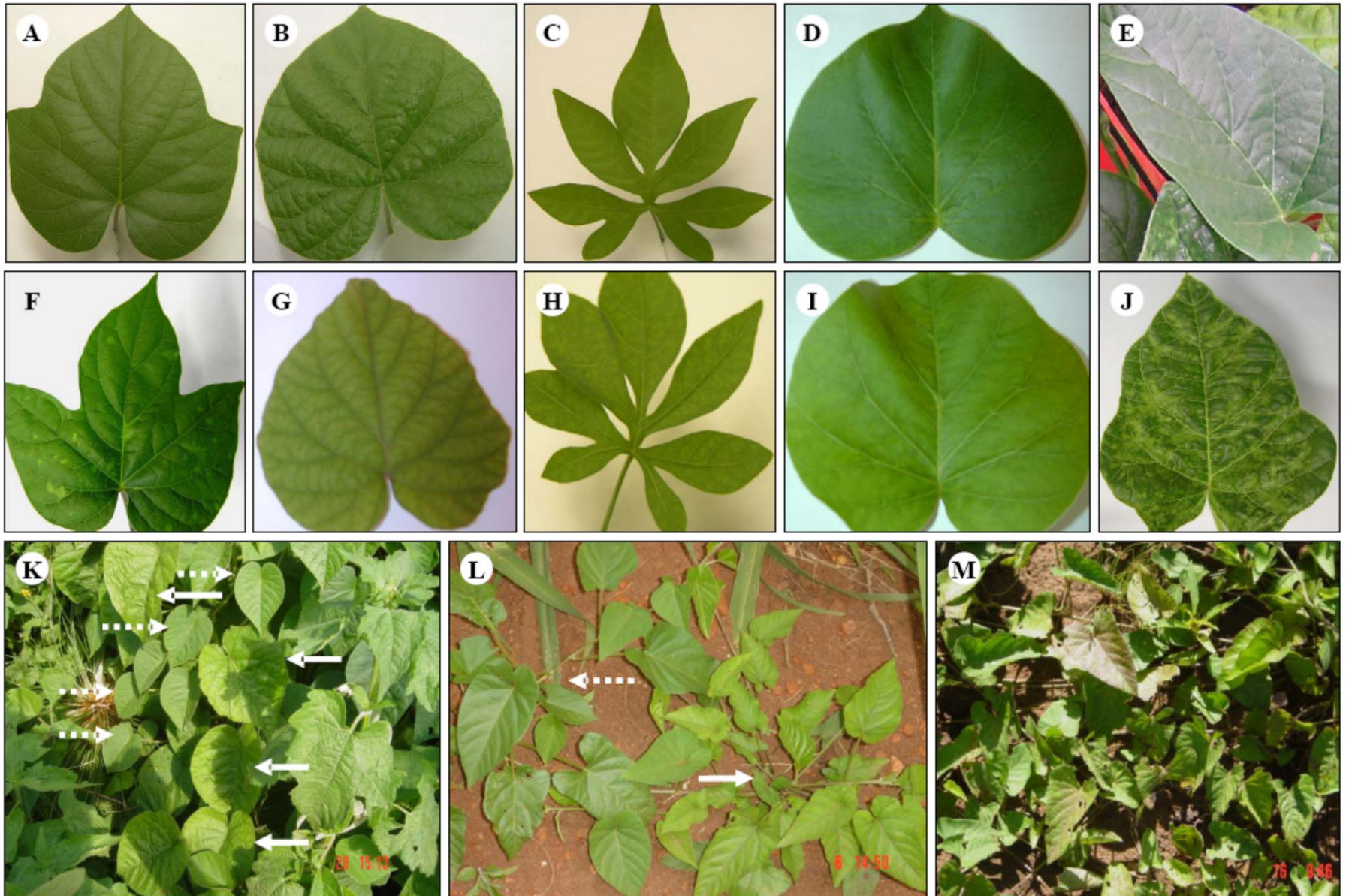
***Ipomoea aquatica* just fished from the edge of a swamp by Arthur Tugume
in Katakwi district, Uganda, in April 2007**



Wild species of *Ipomoea* in Uganda

Tugume et al. 2008.
Phytopathology 98: 640-652.

Various types of virus-like symptoms in wild *Ipomoea* species in Uganda



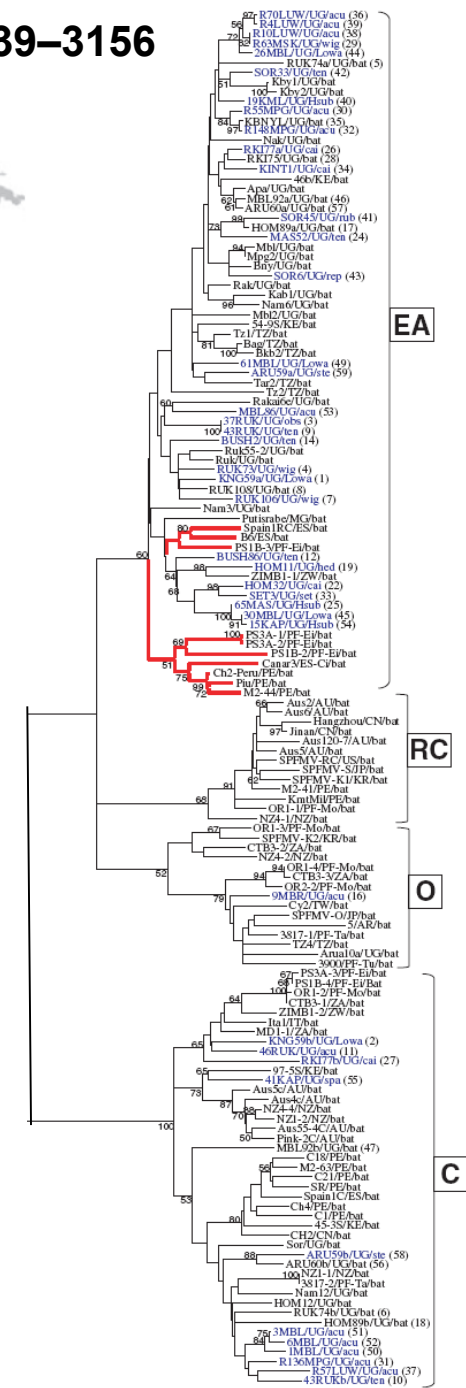
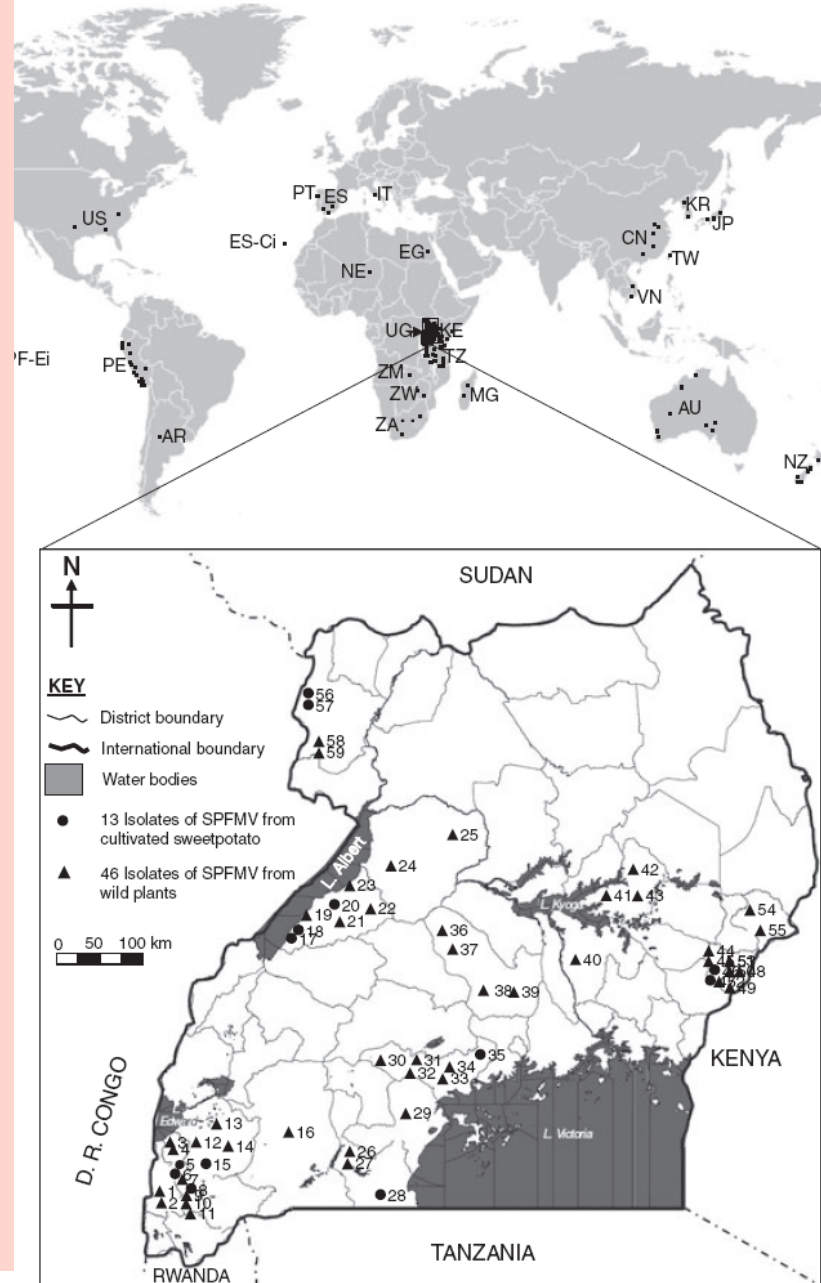
Survey in Uganda:

In all, 2,864 wild plants were observed for virus-like symptoms and tested for SPFMV in 2004 and 2007.

22 *Ipomoea* species, *Hewittia sublobata*, and *Lepistemon owariensis* were infected with SPFMV.

19 species were new hosts for SPFMV.

Tugume, Mukasa & Valkonen 2008. Phytopathology 98: 640-652.



Conclusions

- 1. Production of healthy planting materials is in progress but does not yet reach all farmers**
- 2. Virus-resistant varieties of potato are needed, and while especially landrace sweetpotatoes show resistance to many viruses, resistance to mixed virus infections needs to be identified**
- 3. New methods will allow building comprehensive knowledge on viruses infecting the current crops and prevailing in the cultivation environment, especially in wild plant species, which informs strategies aiming to prevent infection of crops with viruses.**

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Potato Project Team at ARI-Uyole and TOSCI, Tanzania



Ministry for Foreign Affairs of Finland

Academy of Finland

A Decade of Bioscience
Development in Eastern Africa:

The BIO-EARN Programme 1999–2010



TABLE 4. Number of plants that were expressing virus-like symptoms, those that were symptomless, and those that tested positive for *Sweet potato feathery mottle virus* (SPFMV) in nitrocellulose membrane enzyme-linked immunosorbent assay (NCM-ELISA) from each plant species observed and tested in 2004 and 2007^a

Plant species	Total no. of plants	With virus-like diseases		Symptomless plants		Overall
		Total (%) ^b	SPFMV+ ^c	Total (%) ^d	SPFMV+ (%) ^e	SPFMV+ (%)
2004						
<i>Ipomoea acuminata</i>	55	15 (27.3)	60	40 (73.7)	3 (7.5)	12 (21.8)
<i>I. aquatica</i>	12	4 (33.0)	100	8 (67.0)	3 (37.5)	7 (58.3)
<i>I. blepharophylla</i>	14	4 (29.0)	75	10 (71.0)	7 (70.0)	10 (71.4)
<i>I. cairica</i>	99	9 (9.1)	88.9	90 (90.9)	29 (32.2)	37 (37.4)
<i>I. cordofana</i>	10	0 (0.0)	0	10 (100.0)	1 (10.0)	1 (10.0)
<i>I. crepidiformis</i>	18	4 (22.0)	75	14 (78.0)	4 (28.6)	7 (38.9)
<i>I. eriocarpa</i>	134	5 (4.0)	0	129 (96.0)	1 (0.8)	1 (0.7)
<i>I. grantii</i>	9	0 (0.0)	0	9 (100.0)	1 (11.1)	1 (11.1)
<i>I. hederifolia</i>	13	5 (38.0)	60	8 (62.0)	1 (12.5)	4 (30.7)
<i>I. hildebrandtii</i>	11	1 (9.1)	100	10 (90.9)	2 (20.0)	3 (27.3)
<i>I. involucrata</i>	78	3 (4.0)	33.3	75 (96.0)	1 (1.3)	2 (2.6)
<i>I. obscura</i>	79	10 (12.7)	100	69 (87.3)	2 (2.9)	12 (15.2)
<i>I. polymorpha</i>	10	0 (0.0)	0	10 (100.0)	2 (20.0)	2 (20.0)
<i>I. purpurea</i>	10	2 (20.0)	100	8 (80.0)	3 (37.5)	5 (50.0)
<i>I. repens</i>	24	7 (29.0)	42.8	17 (71.0)	0 (0.0)	3 (12.5)
<i>I. rubens</i>	27	3 (11.0)	66.7	24 (89.0)	8 (33.3)	10 (37.0)
<i>I. sinensis</i>	143	45 (31.5)	88.9	98 (68.5)	10 (10.2)	50 (35.0)
<i>I. spathulata</i>	15	1 (7.0)	100	14 (93.0)	6 (42.8)	7 (46.7)
<i>I. stenobasis</i>	13	6 (46.0)	100	7 (54.0)	1 (14.3)	7 (53.8)
<i>I. tenuirostris</i>	230	23 (10.0)	74	207 (90.0)	2 (0.9)	19 (8.3)
<i>I. velutipes</i>	10	1 (10.0)	100	9 (90.0)	2 (22.2)	3 (30.0)
<i>I. wightii</i>	62	10 (16.1)	90	52 (83.9)	4 (7.7)	13 (21.0)
<i>Hewittia sublobata</i>	420	67 (16.0)	91	353 (84.0)	22 (6.2)	83 (19.8)
<i>Lepistemon owariensis</i>	44	5 (11.4)	80	39 (88.6)	21 (53.8)	25 (56.8)
2007						
<i>I. acuminata</i>	102	20 (20.0)	80	82 (80.0)	14 (17.1)	30 (29.0)
<i>I. aquatica</i>	12	3 (25.0)	33.3	9 (75.0)	2 (22.1)	3 (25.0)
<i>I. blepharophylla</i>	–	–	–	–	–	–
<i>I. cairica</i>	121	6 (5.0)	66.7	115 (95.0)	10 (8.7)	14 (11.5)
<i>I. cordofana</i>	–	–	–	–	–	–
<i>I. crepidiformis</i>	12	0 (0.0)	0	12 (100.0)	1 (8.3)	1 (8.3)
<i>I. eriocarpa</i>	–	–	–	–	–	–
<i>I. grantii</i>	–	–	–	–	–	–
<i>I. hederifolia</i>	34	8 (24.0)	75	26 (76.0)	1 (3.8)	7 (20.5)
<i>I. hildebrandtii</i>	6	1 (17.0)	100	5 (83.0)	0 (0.0)	1 (16.7)
<i>I. involucrata</i>	–	–	–	–	–	–
<i>I. obscura</i>	75	10 (13.0)	50	65 (87.0)	4 (4.6)	8 (10.7)
<i>I. polymorpha</i>	–	–	–	–	–	–
<i>I. purpurea</i>	14	1 (7.1)	100	13 (92.9)	0 (0.0)	1 (7.1)
<i>I. repens</i>	3	1 (33.0)	100	2 (67.0)	0 (0.0)	1 (33.3)
<i>I. rubens</i>	10	2 (20.0)	50	8 (80.0)	0 (0.0)	1 (10.0)
<i>I. sinensis</i>	231	31 (13.0)	67.8	200 (87.0)	7 (3.5)	28 (12.1)
<i>I. spathulata</i>	27	2 (7.4)	100	25 (92.6)	2 (8.0)	4 (14.8)
<i>I. stenobasis</i>	10	1 (10.0)	100	9 (90.0)	0 (0.0)	1 (10.0)
<i>I. tenuirostris</i>	165	19 (12.0)	73.6	146 (88.0)	8 (5.5)	22 (13.3)
<i>I. velutipes</i>	–	–	–	–	–	–
<i>I. wightii</i>	51	6 (12.0)	50	45 (88.0)	1 (2.2)	4 (7.8)
<i>H. sublobata</i>	267	22 (8.2)	63.6	245 (91.8)	4 (1.6)	18 (6.7)
<i>L. owariensis</i>	53	4 (8.0)	75	49 (92.0)	5 (10.2)	8 (15.1)

^a Dash (–) indicates that no observations and NCM-ELISA tests were conducted for a given species in the year 2007.

^b Numbers in parenthesis represent the proportion of plants with virus-like symptoms as a percentage of total number of plants observed in a given year.

^c Percentage and number of symptomatic plants that consistently and unambiguously showed a positive reaction in NCM-ELISA for SPFMV among the total number of plants with virus-like disease symptoms in a given year.

^d Numbers in parenthesis represent the proportion of plants without virus-like symptoms as a percentage of total number of plants observed in a given year.

^e Percentage and number of symptomatic plants that consistently and unambiguously showed a positive reaction in NCM-ELISA for SPFMV among the total number of plants without virus-like disease symptoms in a given year. Numbers in parenthesis represent the percentage of symptomless plants infected with SPFMV.

Tugume, Mukasa & Valkonen 2008. Phytopathology 98: 640-652.

Natural wild hosts of Sweet potato feathery mottle virus show spatial differences in virus incidence and virus-like diseases in Uganda.

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Seed Potato Development Project in Tanzania

1. Strengthened capacity for potato pathogen testing and cleaning
2. Capacity for pre-basic potato production improved
3. Increased capacity of certified potato production in Tanzania
4. Field management practices improved

Tanzania Official Seeds Certification Institute (TOSCI).

TOSCI's capacity to inspect and certify seed will be enhanced in the project through training in Quality Declared Seed protocol for potato.

- Increased number of private and community based seed potato producers;
- Increase in certified seed production (has);
- Number of TOSCI inspectors trained;
- Testing capacity of TOSCI (samples per season); and
- Number of farmers benefiting from clean seed.

All field multiplied seeds will be certified by TOSCI using the FAO QDS standard and/or other Tanzanian seed standards that may be in force. Prior to the certification, there need to be capacity building activities for TOSCI staff.