

Digital Receipt

This receipt acknowledges that Turnitin received your paper. Below you will find the receipt information regarding your submission.

The first page of your submissions is displayed below.

Submission author: Ni Putu Pandawani

Assignment title: Article

Submission title: Distribution and molecular characterization of Squash mosa...

File name: acterization_of_Squash_mosaic_virus_on_cucumber_in_-MAR...

File size: 428.52K

Page count: 7

Word count: 3,777

Character count: 19,445

Submission date: 01-Apr-2022 05:37PM (UTC+0700)

Submission ID: 1798773431



Distribution and molecular characterization of Squash mosaic virus on cucumber in Gianyar, Bali

by Ni Putu Pandawani

Submission date: 01-Apr-2022 05:37PM (UTC+0700)

Submission ID: 1798773431

File name: acterization_of_Squash_mosaic_virus_on_cucumber_in_-MARET_22.pdf (428.52K)

Word count: 3777

Character count: 19445

ISSN: 1411-7525 E-ISSN: 2461-0399 DOI: 10.23960/j.hptt.12248-54

RESEARCH PAPER

13

Distribution and molecular characterization of Squash mosaic virus on cucumber in Gianyar, Bali

Listihani¹, Ni Putu Pandawani¹, Tri Asmira Damayanti², Mimi Sutrawati³, Dewa Gede Wiryangga Selangga⁴, Ketut Ayu Yuliadhi⁵, Trisna Agung Phabiola⁵, & Gusti Ngurah Alit Susanta Wirya⁵

Manuscript received: 17 December 2021. Revision accepted: 4 March 2022. Availabe online: 24 March 2022.

ABSTRACT

Squash mosaic virus (SqMV) has been reported to infect cucumber plants in Java and cause a decrease in fruit quality and quantity. Until now, there is no information regarding Sqmv infection in Cuc vitaceae in Bali. In a preliminary research conducted during 2020, we found mosaic symptoms in Gianyar, Bali. SqMV was inferred to be the possible cause of the virus-disease-like symptoms. The study aimed to determine the precence of SqMV in Gianyar, Bali and to obtain information related to its molecular character. This research method includes surveys, field observations, virus identification by RT-PCR, and DNA analysis. Severe disease incidence caused by SqMV was observed on cucumber cultivation in Gianyar that was in the range of 5.81–66.87%. PCR using specific primer for coat protein (CP) gene of SqMV was successfully amplified the DNA fragments of ± 582 bp on samples from Payangan, Tegallalang, Ubud, Sukawati, Blahbatuh, and Gianyar districts, except for samples from Tampaksiring. This indicates that the presence of SqMV is widespread in Java and Gianyar. The SqMV isolate from Bali had the highest nucleotide homology at 91.9–93.4% and amino acids 94.0–94.5 and was closely related to the Brazil isolate (KT923125) and had loger isolate homology than other countries (China, Trinidad, Spain, Japan, Arizona, Cekoslowakia, and Australia). It proved that those mosaic symptoms on cucumber is associated with SqMV infection. SqMV

INTRODUCTION

Squash mosaic virus (SqMV) in 34 tion causes greenish-yellow mosaic symptoms on cucumber leaves. The leaves become malformations with narrow leaf size and stunted plants. Cucumber plants infected with SqMV showed shrinkage of the fruit base at fruit malformations. The infected fruit can change in taste and nutritional content, seed germination, fruit weight, and the amount of fruit production (Lestari &

24 responding author:

Gusti Ngurah Alit Susanta Wirya (susantawirya@unud. ac.id)

Study Program of Agrotechnology, Faculty of Agriculture and Business, University of Mahasaraswati, Jl. Kamboja No.11A, Der 10 ar, Bali, Denpasar, Indonesia 80233

²Department of Plant Protection, Faculty of Agriculture, IPB University, Jl. Raya Dramaga, Bogor, West Java, Indonesia 16680

³Faculty of Agriculture, Bengkulu University, Jl. W.R Supratman, Kandang Limun, Bengkulu 38371A

⁴Faculty of Agriculture, Warmadewa University, Jl. Terompong No. 24, Tanjung Bungkak, Denpasar, Bali 80239 ⁵Stuc 20 rogram of Agroecotechnology, Faculty of Agriculture, Udayana University, Jl. P.B. Sudirman, Denpasar, Bali 80234, Indonesia

Nurhayati, 2014). According to A 35t al. (2012), *Cucumis sativus*, *C. melo, Cucurbita pepo, C. moschata*, and *C. maxima* plants infected with SqMV showed mosaic symptoms. In advanced symptoms, SqMV causes a decres in fruit production and malformations.

SqMV is a seed-borne virus in cucurbit plants and can spread during planting and at harvest after primary infection of infected seeds (Şevik & Toksöz, 2008). SqMV can also be transmitted mechanically or via inject vectors (Maina et al., 2017). SqMV infection in the field occurs with the help of vector insects, namely beetles from the Chrysomelidae family (Acalymma thiemei, Diabrotica sp., and Aulacophora similis) and Coccinelidae (Epilachna chrysomelina) (Ali etgl., 2012).

Regulation of the minister of agriculture of the republic of Indonesia number 25 year 2020 concerning types of quarantined plant destruction organisms (OPTK) stipulates the status of SqMV as OPTK category A2 group 1 which is still limited in West Java and it's that cannot released from carrier media (Kepmentan, 2020). Lestari & Nurhayati (2014) reported that SqMV had infected 100% of cucumber seeds. SqMV has been detected in five cucumber varieties in Indonesia and has been detected in Java (Purba et al., 2017; Listihani, 2019). Several factors that can trigger an

increase in disease severity in an area are changes in climatic conditions, plant cultivation techniques, and traffic of plant material betweer 12 gions (Ahanger et al., 2013; Hanssen et al., 2010). Tobacco mosaic virus (TMV) increases its multiplication when temperature reaches 34–36 °C (Ahanger et al., 2013). Given the nature of SqMV, it is possible that the source of SqMV infection in Indonesia comes from imported vegetable seeds.

Until now, there have no re 32 ts of SqMV infection in cucumber plants in Bali. Therefore, this study aimed to determine the presence of SqMV in cucumber plants in Bali. This information is very important because there is no information regarding the detection and presence of SqMV in Bali.

MATERIALS AND METHODS

Collection and Disease Incidence. The survey and sampling of cucumber cultivations were carried out in Gianyar Regency, Bali Province (Payangan, Tegallalang, Tampaksiring, Ubud, Sukawati, Blahbatuh, and Gianyar Districts). Sampling was carried out by purpose sampling method, as many as 10 symptomatic samples were taken from each location. The total samples taken were 90 samples, then used as material for virus detection. Calculation of disease incidence by random sampling with each location as many as 50 plants. The disease incidence (DI) in the field is calculated by:

 $DI = \frac{The \ number \ of \ symptomatic \ samples}{Total \ samples \ in \ the \ field} \times 100\%$

tal RNA Extraction from Symptomatic Leaves and RT-PCR Amplification. Total RNA was extracted from samptomatic plant leaf tissue using the CTAB method. Total viral RNA was isolated from infected leaf with a procedure described by Doyle & Doyle (1990). Fresh tissue (0.1 g) grinded with liquid nitrogen and added 500 μL of 10% CTAB buffer (cetyl-trimethyl-ammonium bromide, 0.1 M Tris-HCl pH 8, 0.05 M EDTA, 0.5 36 NaCl, 1% β-mercaptoethanol). Then 7 transferred to 1.5 mL micro tubes and incubated in a water bath at 65 °C for 30 min, then the micro tubes were inverted every 10 min. After 30 min the micro tube containing the mixture was taken from a water bath and allowed to stand for 2 min at room temperature, then added 500 µL of the hloroform: Isoamyl alcohol mixture with a ratio of $\overline{24}$: 1 (v:v). The mixture was vortexed for 5 min until well mixed, then centrifuged (Microlitre Centrifuge Z 207M, Hermle, Germany) at a speed of 14,000 rpm for 15 min. A total of 450 µL of supernatant was taken and transferred into a new micro tube, then 3 M of sodium acetate was added. The mixture was vortexed and incubated at -80 °C for 2 hours or -20 °C 12 one night. After incubation, the nucleic acid mixture was centrifuged at 12,000 rpm for 13 min to precipitate the nucleic acids. The nucleic acid pellets were washed by 500 µL of 70% ethanol, then centrifuged again at 8000 rpm for 5 min, pellets were dried. After drying, the pellets containing total RNA were dissolved in 50 to 100 µL of TE buffer (pH 8) (10mM Tris-HCl, 8 mM EDTA) and it was stored at -20 °C until ready for use.

The composition of the reverse transcription (RT) PCR mixture consisted of 1 μL oligo dNTP 10 mM, 2 μL total RNA, ar 26.75 μL dH2O. All reagents were vortex gently and incubated at 65 °C for 5 min in water bath, then immediately cooled 17 ice. Next to the reactants were added 2 μL of RT buffer, 1 μL dNTP 18 mM, 0.5 μL DTT 50 mM, 0.5 μL RNAse inhibitor (RiboLock RNase Inhibitor 20 15 ts/μL), 0.25 μL MmuLV (Revertaid 200 units/μL) (Thermo Fisher Scientific, Waltham, MA, USA) 11 a total volume of 10 μL. The reverse transcription reaction was carried out at 42 °C for 60 min followed by 70 °C for 10 min in order to deactivate the enzyme in the PCR machine. The cDNA product can then be used as a template for amplification.

Amplification of cDNA total using machine PCR thermal cycler (MultiGene OptiMax Thermal Cycler 230V Model, Labnet, USA). The primer pair used to amplify SqMV was SqMV RNA2-f1 (5'-GGTG-CAGCAGCTTGGAACTTATAATCCAATTTGG-3')/SqMV RNA2-r1 (5'-TGGGAAAGAAGCCACAA-CAAAACCC-3'), with target amplicon sizes ± 12 bp (Chinnaraja et al., 2015). The composition of the amplification reaction for a total volume of 25 μL was 12.5 μL Go Taq green 2x (Thermo scientific, USA), 1 μL 10 μM reverse primer, 1 μL 10 μM forward primer, 9.5 μL nuclease-free water, and 1 μL cDNA.

DNA was electrophoresis on 1% agarose gel [0.3 g of agarose dissolved in 0.5x TBE buffer (45 mM Tris-borat, 1 mM EDTA) up to 30 mL]. The agarose gel solution was cooled to 50 °C for 15 min, then fluor-oVuo TM nucleic acid dye (Smobio, Taiwan) was added. Electrophoresis was carried out at a voltage of 100 V for 20 min. The results of the electrophoresis were then visualized under ultraviolet transilluminator and documented with a digital camera.

Identification of SqMV Based on DNA Sequence Analysis. The amplified DNA fragments were sent to 1st Base Malaysia for the sequencing. The sequence

Vol. 22, No. 1, 2022: 48-54

was analyzed using BLAST (Basic Local Alignment Search Tool) program (https://blast.ncbi.nlm.nih.gov/Blast.cgi) aims to see the level of similarity to the nucleotide sequences of related species found in Gen-Bank. Matrix homo 31y identities were analyzed using BioEdit software 7.013 for Windows 95/98/NT/2000/XP (Hall, 2007) and phylogenetic trees were analyzed using MEGA v6.0 software (Tamura et al., 2013) using bootstrap 1000 times replication.

RESULTS AND DISCUSSION

Disease Symptoms on Cucumber Plants. During field surveys in several cucumber cultivation areas in Gianyar Regency, we found varying virus-like symptoms on cucumbers such as mosaic, yellowing, vein banding, vein clearing, and leaf malformation, and mottle (Table 1). Mosaic symptoms were found in all districts, except in Tampaksiring. Mosaic symptoms observed in the field were seen on young leaves (Fig-

ure 1A) and on fruit showed malformations accompanied by mosaic symptoms (Figure 1B). Symptoms like these are typical symptoms of Squay infection in Cucurbitaceae plants (Aulia, 2004; Ali et al., 2012; Firmansyah et al., 2017).

Diseases Incidence of Virus like Symptoms. The incidence of disease in the field ranged from 5.81% to 66.87% (Table 1). The disease incidence was more than 50% in Ubud and Blahbatuh Districts, while the incidence of disease is less than 10% in Sukawati District. Good sanitation practices and vectors of rare insects on cucumber plantations in Sukawati lead to low incidence. According to Aulia (2004) showed that the percentage of SqMV infection in Cucurbitaceae plants in Bogor was the highest pared to other viruses (CMV, PRSV, ZYMV). Ali et al. (2012) reported that the percentage of Cucurbitaceae plants infected with SqMV was only 3.8%. Meanwhile, Şevik & Toksöz (2008) reported that SqMV infection in pumpkins (C.

Table 1. Symptoms and incidence of disease on cucumber plants in Gianyar Regency, Bali Province

Districs	Plant age (DAP)*	Disease symptoms	Disease incidence (%)
Payangan	33-40	Mosaic, yellowing, leaf malformation and mottle	33.97 (123/362)
Tegallalang	24-42	Mosaic, vein banding, yellowing, leaf malformation	37.83 (213/563)
Tampaksiring	28 - 37	Vein banding, yellowing, leaf malformation	19.11 (134/701)
Ubud	35-42	Mosaic, yellowing, vein banding, and mottle	66.87 (523/782)
Sukawati	20-33	Mosaic and yellowing	5.81 (42/739)
Blahbatuh	30-43	Mosaic, vein clearing, vein banding, and mottle	63.80 (520/815)
Gianyar	33–46	Mosaic, yellowing, vein banding, vein clearing, and leaf malformation	31.67 (134/428)

^{*=} Days after planting (DAP).

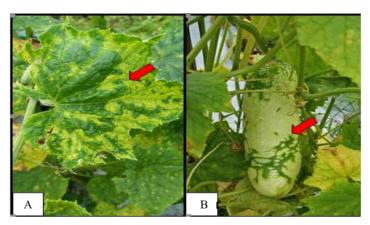


Figure 1. Symptoms of SqMV infection on cucumber in Bali; (A) Mosaic and curly on leaf; (B) Mosaic and malformation on fruit.

moschata and C. maxima) was 21.11% and in zucchini as much as 20%, while Dikova & Hristova (2002) stated that seed-borne SqMV in Cucurbitaceae plants reached 91%. The spread of SqMV in the field is due to the inoculum sources that are continuously available in the field due to the unavailability of virus-free seeds.

Virus Identification by RT-PCR, Sequencing and Sequence Analysis. Specific DNA band SqMV measuring ±582 bp was successfully amplified in several samples showing mosaic symptoms from Payangan, Tegallalang, Ubud, Sukawati, Blahbatuh, and Gianyar Districts, except for samples from Tampaksiring (Figure 2). No mosaic symptoms were found on cucumbers in Tampaksiring and no SqMV was detected by RT-PCR, it was suspected that there were other viruses that infect cucumber plants. Chinnaraja et al. (2015) succeeded in detecting SqMV from watermelon and pumpkin plant samples in Trinidad with an amplicon size of ±582 bp using the same primer.

The homology of SqMV nucleotides and amino acids between isolates in other countries was 87.2-96.5% and 89.8-96.1%, respectively. These data indicated that the samples from Bali which were 25 gned with the sequence from GenBank were SqMV isolates. The nucleotide sequence similarity of the CP SqMV gene ranged from 81.1–99.6% for the same virus species (Haudenshield & Palukaitis, 1998). The SqMV isolate from Bali had the highest nucleotide homology at 91.9-93.4% and amino acids 94.0-94.5% and was closely related to the Brazil isolate (KT923125) and had lower isolate homology than other countries. The relationship between isolates based on phylogenetic analysis showed that SqMV isolates from Bali were in the same group as SqMV watermelon isolates from Brazil and separated from SqMV isolates from other countries (China, Trinidad, Spain, Japan, Arizona, Cekoslowakia, and Australia) (Figure 3). In this study, the differences in these groups are not due to geographic location and hosts (watermelon, squash, crookneck pumpkin).

SqMV is an exotic virus because it includes quarantine plant pest organisms (OPTK) category A2 group 1. Prior to 2015, SqMV was included in quarantine plant pest organism category A1 (OPTK A1) whose presence had not been reported in Indonesia. In 2015 until now, SqMV was reported in Indonesia on cucumber plants in Bogor (Kementan, 2020), so that in 2015, SqMV changed its status from OPTK A1 to OPTK A2, namely OPTK that already exists in Indonesia, but its existence is still limited. in some areas only.

SqMV may have infected Cucurbitaceae plants in Indonesia for a long time, but not many have reported its molecular character. SqMV has been reported to infect cucumber plants in Bogor based on serological detection (Firmansyah et al., 2017). In addition, the molecular character of SqMV isolates from Java has been reported by Listihani (2019). The presence of SqMV infection in cucumber plants in Gianyar in this study indicated that the distribution was increasing and it was strictly find the field.

This is the first report of SqMV infection in cucumber plants in Bali. The absence of an appropriate detection method and clean sanitation in the field, the present of insect vectors all year round in the field, as well as the use of non-virus-free seeds are the causes of the rapid spread of SqMV distribution in the field (Firmansyah et al., 2017). Several factors that can trigger the emergence of new diseases in an area are plant cultivation techniques and traffic of plant material between regions well as changes in climatic conditions (Jones, 2016). Nancarrow et al. (2014), studied the effects of elevated (10–21 °C, night/day) or ambient (5–16 °C, night/day) temperature winter growing season regimes on wheat plants infected with BYDV. Infected

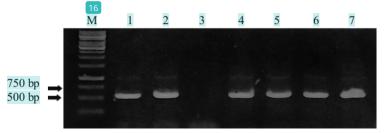


Figure 2. Visualization of DNA amplification of SqMV with primer pair of SqMV RNA2-f1/RNA2-f18 n 1% agarose gel. (M) DNA marker (1 kb ladder) (Thermo Scientific), (1 to 7) DNA samples from: (1) Payangan; (2) Tegallalang; (3) Tampaksiring; (4) Ubud; (5) Sukawati; (6) Blahbatuh; (7) Gianyar.

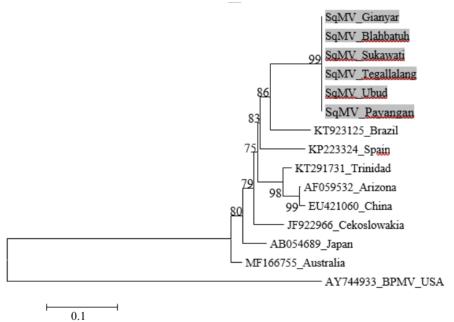


Figure 3. Phylogeny tree of nucleotide sequences of CP SqMV Bali Isolates using MEGA 6.0 (Neighbour Joining with bootstrap 1000x) with SqMV from other countries that deposited in GenBank. *Bean pod mottle virus* (BPMV) from America is used as out groups. The highlight mark is a SqMV isolates from Bali.

plants grown under elevated temperature were larger, developed virus symptoms earlier, and had higher virus titers than plants grown at ambient temperature. SqMV can infect cucumber seeds up to 100% (Firmansyah et al., 2017). This does not rule out the possibility that the source of SqMV infection in Indonesia comes from imported seeds.

CONCLUSION

Variations in symptoms were found in all sampling locations, but mosaic symptoms were not found in Tampaksiring. based on RT-PCR, plants showing mosaic symptoms were positive for SqMV. Therefore, cucumber plantings at all locations in Gianyar were positive for SqMV infection, except Tampaksiring. This study is the first report of SqMV infection in Bali. The existence of SqMV in Gianyar, Bali shows the increasing number of sources of inoculum in the field. Therefore, it is necessary to prevent the distribution of seeds and seedlings to other areas that have not been infected with SqMV.

ACKNOWLEDGMENTS

We would like to thank the Head of the Plant Disease Laboratory, Udayana University for providing a tool for molecular detection.

FUNDING

Research with independent funding sources.

AUTHORS' CONTRIBUTIONS

I, NPP, DGWS, KAY, TAP, and GNASW carried out the survey and sampling including observation of disease symptoms on cucumber cultivations in Gianyar, Bali. TAD and MS supported the research materials. I, DGWS, KAY, and GNAWS performed molecular work and data analysis. I, DGWS, NPP, KAY, TAP, GNAWS, TAD, and MS papared the manuscript. The author provides feedback and comments on the flow of research, data analysis and interpretation as well as shape of the manuscript. All the authors have read and approved the final manuscript.

COMPETING INTEREST

Authors declare no competing interest such as financial or non-financial interests, professional or personal relationships that are directly or indirectly connected to the work submitted for publication.

REFERENCES

- Ahanger RA, Bhat HA, Bhat TA, Ganie SA, Lone AA, Wani IA, Ganai SA, Haq S, Khan OA, Junaid JM, & Bhat TA. 2013. Impact of climate change on plant diseases. *Int. J. Modern Plant & Anim Sci.* 1(3): 105–115.
- Ali A, Mohammad O, & Khattab A. 2012. Distribution of viruses infecting cucurbit crops and isolation of potential new virus-like sequences from weeds in Oklahoma. *Plant Dis.* 96(2): 243–248. https://doi.org/10.1094/PDIS-05-11-0419
- Aulia R. 2004. Inventarisasi dan deteksi virus penyebab mosaik pada Famili Cucurbitaceae di Kotamadya Bogor, Pasir Muncang dan Cibodas [Inventory and detection of virus causing mosaic in the Cucurbitaceae Family in the Bogor, Pasir Muncang and Cibodas]. Skripsi. IPB University. Bogor.
- Chinnaraja C, Ramkissoon A, Rajendran R, Tony ST, Ramsubhag A, & Jayaraj J. 2015. First report of Zucchini yellow mosaic virus and Squash mosaic virus infecting cucurbits in Trinidad. Plant Dis. 100(4): 866. https://doi.org/10.1094/PDIS-09-15-0988-PDN
- Dikova B & Hristova D. 2005. Detection of Squash mosaic virus (SqMV), Zucchini yellow mosaic virus (ZYMV) and Cucumber mosaic virus (CMV) in cucurbit seeds. Bulg. J. Agric. Sci. 11(1): 49-59.
- Doyle JJ & Doyle JL. 1990. Isolation of plant DNA from fresh tissue. *Focus*. 12(1): 13–15.
- Firmansyah D, Widodo, & Hidayat SH. 2017. Chitosan and plant growth promoting rhizobacteria application to control *Squash mosaic virus* on cucumber plants. *Asian J. Plant Pathol.* 11: 148–155. https://doi.org/10.3923/ajppaj.2017.148.155
- Hall T. 2007. BioEdit version 7.0.9. Carlsbad: Computer Program and documentation, Ibis Biosciences. http://www.mbio.ncsu.edu/BioEdit/bioedit.html

- Hanssen IM, Lapidot M, & Thomma BPHJ. 2010. Emerging viral diseases of tomato crops. Mol. Plant Microbe Interac. 23(5): 539–548. https://doi.org/10.1094/MPMI-23-5-0539
- Haudenshield JS & Palukaitis P. 1998. Diversity among isolates of *Squash mosaic virus*. *J. Gen. Virol.* 79(10): 2331–2341. https://doi.org/10.1099/0022-1317-79-10-2331
- Jones RAC. 2016. Future scenarios for plant virus pathogens as climate change progresses. Adv. Virus Res. 95: 87–147. https://doi.org/10.1016/ bs.aivir.2016.02.004
- [Kementan] Kementrian Pertanian Republik Indonesia. 2020. Peraturan Menteri Pertanian Republik Indonesia nomor 25 tahun 2020 tentang jenis organisme pengganggu tumbuhan karantina [Regulation of the Minister of Agriculture of the Republic of Indonesia number 25 year 2020 about types of quarantine plant disordering organisms]. Kementerian Pertanian Republik Indonesia. Jakarta.
- Lestari SM & Nurhayati E. 2014. Efisiensi tular benih *Squash mosaic virus* pada Cucurbitaceae [Seed transmission efficiency of *Squash mosaic virus* on cucurbitaceae]. *J. Fitopatol. Indones.* 10(3): 81–86. https://doi.org/10.14692/jfi.10.3.81
- Listihani. 2019. Karakterisasi dan epidemic virus pada tanaman mentimun di Pulau Jawa [Characterization and epidemics of virus on cucumber in Java Island]. *Dissertation*. IPB University. Bogor.
- Maina S, Edwards OR, & Jones RAC. 2017. Two complete genome sequences of *Squash mosaic virus* from 20-year-old cucurbit leaf samples from Australia. *Genome Announc*. 5(32): e00778-17. https://doi.org/10.1128/genomeA.00778-17
- Nancarrow N, Constable FE, Finlay KJ, Freeman AJ, Rodoni BC, Trebicki P, Vassiliadis S, Yen AL, & Luck JE. 2014. The effect of elevated temperature on *Barley yellow dwarf virus*-PAV in wheat. *Virus Res.* 186: 97–103. https://doi.org/10.1016/j.virusres.2013.12.023
- Purba ER, Lestari SM, Nurhaelena Y, & Hidayat SH. 2017. Deteksi Squash mosaic virus pada lima varietas mentimun (Cucumis sativus L.) [Squash mosaic virus detection on five cucumber (Cucumis sativus L.) Varieties]. J. Hort. Indonesia. 8(2): 104–110. https://doi.org/10.29244/ jhi.8.2.104-110

Şevik MA & Toksöz Y. 2008. Occurrence of *Squash mosaic virus* (SqMV) infecting pumpkin and squash growing in Samsun, Turkey. *J. Turk. Phytopath.* 37(1–3): 15–25.

Tamura K, Stecher G, Peterson D, Filipski A, & Ku-

mar S. 2013. MEGA6: Molecular evolutionary genetics analysis version 6.0. *Mol. Biol. Evol.* 30(12): 2725–2729. https://doi.org/10.1093/molbev/mst197

Distribution and molecular characterization of Squash mosaic virus on cucumber in Gianyar, Bali

ORIGINALITY REPORT SIMILARITY INDEX **INTERNET SOURCES PUBLICATIONS** STUDENT PAPERS **PRIMARY SOURCES** journal.ugm.ac.id Internet Source Mimi Sutrawati, Parwito Parwito, Priyatiningsih Priyatiningsih, Agustin Zarkani et al. "FIRST REPORT OF Begomovirus INFECTION ON PAPAYA IN BENGKULU, INDONESIA", Jurnal Hama dan Penyakit Tumbuhan Tropika, 2021 Publication docplayer.net 2% Internet Source Yashanti Berlinda Paradisa, Sri Sulandari, Sedyo Hartono, Susamto Somowiyarjo et al. "RECOMBINANT ANTIBODY PRODUCTION BY **CLONING OF Pepper yellow leaf curl** Indonesia virus (PepYLCIV) COAT PROTEIN GENE", Jurnal Hama dan Penyakit Tumbuhan Tropika, 2022 Publication

Penyakit Tumbuhan Tropika, 2020

Publication

11	journals.plos.org Internet Source	1%
12	Dewa Gede Wiryangga Selangga, Listihani Listihani. "MOLECULAR IDENTIFICATION OF Pepper yellow leaf curl Indonesia virus ON CHILI PEPPER IN NUSA PENIDA ISLAND", JURNAL HAMA DAN PENYAKIT TUMBUHAN TROPIKA, 2021 Publication	<1%
13	K. Subramanya Sastry, Bikash Mandal, John Hammond, S. W. Scott, R. W. Briddon. "Encyclopedia of Plant Viruses and Viroids", Springer Science and Business Media LLC, 2019 Publication	<1%
14	hdl.handle.net Internet Source	<1%
15	link.springer.com Internet Source	<1%
16	opoposos into odultr	
10	openaccess.iyte.edu.tr Internet Source	<1%
17		<1 % <1 %

Gerritsen, J., S. Fuentes, W. Grievink, L. van Niftrik, B. J. Tindall, H. M. Timmerman, G. T. Rijkers, and H. Smidt. "Characterization of Romboutsia ilealis gen. nov., sp. nov., isolated from the gastro-intestinal tract of a rat, and proposal for the reclassification of five closely related members of the genus Clostridium into the genera Romboutsia gen. nov., Intestinibacter gen. nov., Terrisporobacter gen. nov. and Asaccharospora gen. nov.", INTERNATIONAL JOURNAL OF SYSTEMATIC AND EVOLUTIONARY MICROBIOLOGY, 2014.

Publication

20

I M Merdana, I K Suada, N L Watiniasih, I W Sudira, L M Sudimartini, N L S Ani, K Budiasa. "Improving the quality of wastewater in traditional chicken slaughterhouses using moringa seed powder", IOP Conference Series: Earth and Environmental Science, 2021

<1%

Publication

dergipark.ulakbim.gov.tr

<1%

22

Esam K.F. Elbeshehy, Ehab M.R. Metwali, Omar A. Almaghrabi. "Antiviral activity of Thuja orientalis extracts against watermelon mosaic virus (WMV) on Citrullus lanatus", Saudi Journal of Biological Sciences, 2015

<1%

23	bmcresnotes.biomedcentral.com Internet Source	<1%
24	s3ilmupertanian-pps.unud.ac.id Internet Source	<1%
25	worldwidescience.org Internet Source	<1%
26	A. A. Fajinmi. " Agro-ecological incidence and severity of , genus , family Potyviridae, on cultivated pepper (L.) in Nigeria ", Archives Of Phytopathology And Plant Protection, 2011	<1%
27	Khalid Amari, Caiping Huang, Manfred Heinlein. "Potential Impact of Global Warming on Virus Propagation in Infected Plants and Agricultural Productivity", Frontiers in Plant Science, 2021 Publication	<1%
28	Liming Liu, Kunlun Xie, Abra Rosaline Tsekpuia, Bin Peng, Mei Liu, Qinsheng Gu. "Construction and biological characterization of an Agrobacterium-mediated infectious cDNA of squash mosaic virus", Virus Research, 2019 Publication	<1%
29	Soad Hassan Taha, Ibrahim Mohamed El- Sherbin, Aida Soliman Salem, Mahmoud	<1%

Abdel-Hami, Ali Hussein Hamed, Ghada Abady

Ahmed. "Antiviral Activity of Curcumin Loaded Milk Proteins Nanoparticles on Potato Virus Y", Pakistan Journal of Biological Sciences, 2019

Publication

- apsjournals.apsnet.org <1% 30 Internet Source www.galizalivre.org Internet Source www.walshmedicalmedia.com 32 Internet Source R.W.S. AGUIAR, G.B. ALVES, A.P. QUEIROZ, I.R. 33 NASCIMENTO, M.F. LIMA. "Evaluation of Weeds as Virus Reservoirs in Watermelon Crops", Planta Daninha, 2018 Publication Deni Firmansyah, Widodo ., Sri Hendrastuti <1% 34 Hidayat. "Chitosan and Plant Growth Promoting Rhizobacteria Application to Control Squash mosaic virus on Cucumber Plants", Asian Journal of Plant Pathology, 2017 **Publication** Lebeda, Ale_, M Widrlechner, J Staub, H Ezura, <1%
 - Lebeda, Ale_, M Widrlechner, J Staub, H Ezura, J Zalapa, and E Kristkova. "Cucurbits (Cucurbitaceae; Cucumis spp., Cucurbita spp., Citrullus spp.)", Genetic Resources

Chromosome Engineering & Crop Improvement, 2006.

Publication



Tzvi Tzfira, Vitaly Citovsky. "Chapter 21 Probing Interactions Between Plant Virus Movement Proteins and Nucleic Acids", Springer Science and Business Media LLC, 2008

<1%

Publication

Exclude quotes

On

Exclude matches

Off

Exclude bibliography