

# Morphological response and genetic variability of four species of chili pepper (*Capsicum* spp.) under infection of pepper yellow leaf curl virus

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**Abstract.** Sayekti TWDA, Syukur M, Hidayat SH, Maharijaya A. 2021. Morphological response and genetic variability of four species of chili pepper (*Capsicum* spp.) under infection of pepper yellow leaf curl virus. *Biodiversitas* 22: 4758-4765. Chili pepper has various types and species, but only five known species are commonly used and consumed. Most cultivated chili is susceptible to various plant diseases, one of which is Pepper yellow leaf curl disease (PYLCD) caused by Pepper yellow leaf curl virus (PYLCV) (*Begomovirus*, *Geminiviridae*). To control PYLCD, resistant variety assembly is required to prevent virus infection in cultivated plants. From this research, testing on four chili species is expected to provide information regarding the resistance and performance of chili peppers to conditions infected with PYLCV. This study was conducted at Dramaga Bogor, West Java, Indonesia in two experimental units: planting under virus-free conditions (as control) and virus-infected conditions. Each experimental unit was carried out using a single factor Randomized Complete Block Design (RCBD) with three replications. Twenty-nine genotypes of chili pepper were used consisted of four species, including *C. annuum*, *C. frutescens*, *C. chinense*, and *C. baccatum*. Of the 29 genotypes tested, thirteen genotypes in the resistant, nine genotypes in moderate resistant, two genotypes in moderate susceptible, three genotypes in the Susceptible, and two genotypes in the highly susceptible category. The heritability, genotypic coefficient of variance (GCOV) and phenotypic coefficient of variance (PCOV) value obtained from testing for all characters is high, ranging from 65.16-99.12%, 14.87-82.60%, and 15.77-84.45%, respectively. Most of the genotypes from *C. chinense* showed good resistance to PYLCV. In general, by considering the category of the resistance level and other characters such as productivity, 'Jolokia' (*C. chinense*), 'Anies' (*C. annuum*) and 'Bonita' (*C. frutescens*) can be ascertained as potential candidate sources of resistance to PYLCV.

**Keywords:** *Begomovirus*, chili pepper, heritability, morphology, resistance

## INTRODUCTION

The world community has placed chili pepper (*Capsicum* spp.) as one of the essential horticultural commodities. We can find the use of chili pepper in every aspect of human life. The World Vegetable Center (WVC)/The Asian Vegetable Development and Research Center (AVRDC) has stored 286 accessions of chili pepper in 1986. This number has increased in 30 years until now; the number of collections owned by WVC has reached 8264 accessions, including accessions from at least 100 countries worldwide (Jarret et al. 2019). In addition, The Vegetable Research and Development Center (TVRC), Kasetsart University, Thailand, is also known to have collected 2827 accessions of chili pepper from all over Thailand in 1989 (Mongkolporn and Taylor 2011). Most of these accessions have been characterized conventionally and using molecular markers for various characters (Munoz-Concha et al. 2020), such as agronomic characters, fruit quality, and resistance to virus attack (Nankar et al. 2020). Chili pepper has various types and species, but until now, only five known species are commonly used and consumed by the public, namely *Capsicum annuum*, *C. frutescens*, *C. baccatum*, *C. chinense*, and *C. pubescens*.

Most of the chili genotypes are susceptible to various diseases, one of which is Pepper yellow leaf curl disease (PYLCD) caused by Pepper yellow leaf curl virus (PYLCV) (*Begomovirus*, *Geminiviridae*) (Sulandari et al. 2007). Infection of PYLCV can be recognized from its symptoms; thickening of the leaf bones, curly leaves, yellowing, and stunting (Sulandari et al. 2006; Gaswanto et al. 2016). Transmission of PYLCV occurs through whitefly (*Bemisia tabaci*) as a vector. Control of this disease is more directed at suppressing the population of vector insects, but it has not been effective in preventing the spread of PYLCV. According to Barchenger et al. (2019), assembling resistant plants is an effective way to control the spread of PYLCD.

Creating disease-resistant varieties requires sources of resistance traits. Sources of resistance to several members of *Begomovirus* have been extensively explored in various crops such as tomatoes (Yan et al. 2018), legumes (Blair and Morales 2008), and cotton (Zaidi et al. 2019). Likewise, many explorations and studies have been related to the resistance traits of chili pepper to PYLCV. Several sources of resistance to PYLCV were reported in several chili genotypes of the *C. chinense* species, including 'Bhut Jolokia' (Adluri et al. 2017).

Although the source of resistance to PYLCV has been identified from several chili plants, there have not been many chili genotypes that are fully resistant to PYLCV. These phenomena are thought to be due to the high potential for mixed infection and recombination/pseudo-recombination of *Begomovirus* (Singh et al. 2016). Mixed infection of several types of *Begomovirus* in one plant can cause an increase in the level of symptoms that are more severe than a single virus infection, while recombination or pseudo-recombination events can trigger new viral strains (Garcia-Gano et al. 2006; Mohamed 2010). This condition can then cause the breakdown of resistance of chili plants against PYLCV (Singh et al. 2016; Rubio et al. 2020). This shows that exploration is still needed to obtain information on the source of chili resistance properties against PYLCV. Through characterization, resistance testing and genetic analysis involving four chili species, it is hoped that data and information can be obtained about the level of resistance and performance of chili peppers under PYLCV-infected conditions.

## MATERIALS AND METHODS

### Study area and genetic material

This study was conducted at Dramaga Bogor, West Java, Indonesia (192 m asl). Plants were grown in a greenhouse with an average temperature of 24.7-32.9 °C and relative humidity of 49.2-82.2%.

Twenty-nine genotypes of chili pepper were used, consisted of five species included *C. annuum*, *C. frutescens*, *C. chinense*, and *C. baccatum*. The genetic materials of chili pepper that were used in this experiment are shown in Table 1. This experiment was carried out using two experimental units: planting under virus-free conditions (as control) and virus-infected conditions. Each unit was arranged in a randomized complete block design (RCBD) with three replications. In each trial unit, ten plants were used for biological replication. Transmission of PYLCV was done using The Whitefly (*Bemisia tabaci*) as a vector.

### Procedures

#### Vector preparation

Whitefly (*Bemisia tabaci*) was used as the vector for PYLCV transmission. The vectors were taken from chili pepper and eggplant gardens around the research site. Before being used for inoculation, the imago of whitefly from the field was maintained on virus-free cotton plants for a month to reproduce. Virus-free whitefly offsprings produced were used in the PYLCV transmission process.

#### Inoculation

Seeds of twenty-nine genotypes of chili pepper were sown in the seedling trays for two weeks or until the seedling reached the two-leaf phase. The well-growth seedling was transplanted to individual pots with a diameter of 20 cm and a height of 15 cm. Inoculation was done ten days after transplanting. Virus transmission was done using an individual inoculation procedure (Ganefianti

2010). In this experiment, we allowed 24 h of acquisition access feeding (AAF) followed by 48 h of inoculation access feeding (IAF). After the inoculation process was done, then the whitefly was eradicated. To ensure the plant was infected, plants were inoculated with ten whiteflies per plant.

Plants were observed for symptom development every day from one to sixty days post-inoculation. The symptom was scored and classified using a severity scale, 0 corresponds to healthy plant (no symptom); 1 for yellowing symptom; 2 for yellowing and curling symptoms; 3 for yellowing, curling, and cupping symptoms; and 4 for yellowing, curling, cupping and stunting symptoms. The score was then used to determine the severity of the disease of each genotype using the following formula:

$$DSI = [(\sum (n_i \times z_i)) / (N \times Z)] \times 100\%$$

Where: DSI: Disease Severity Index;  $n_i$ : class frequency;  $z_i$ : assessment class score; N: number of plants observed; Z: maximum disease index. Plant resistance was categorized into five categories: Resistant ( $0 \leq DSI \leq 10\%$ ), Moderately Resistant ( $10 < DSI \leq 20\%$ ), Moderately Susceptible ( $20 < DSI \leq 30\%$ ), Susceptible ( $30 < DSI \leq 50\%$ ) and Highly Susceptible ( $DSI > 50\%$ ). In addition to observing the symptoms of TYLCV attack, we also observed morphological, agronomic, and yield components of the chili pepper genotypes.

**Table 1.** The genetic material of 30 chili pepper

Genotype	Species
Yuni	<i>C. annuum</i>
IPBC12	<i>C. annuum</i>
IPBC15	<i>C. annuum</i>
Ungara	<i>C. annuum</i>
Anies	<i>C. annuum</i>
Seloka	<i>C. annuum</i>
SSP	<i>C. annuum</i>
IPBC5	<i>C. annuum</i>
IPBC3	<i>C. annuum</i>
F5-012328-6-2-1	<i>C. annuum</i>
F5-012328-1AB-2-1	<i>C. annuum</i>
F5-012328-6-2-2	<i>C. annuum</i>
F4-012328-1AB-3	<i>C. annuum</i>
F4-012328-3-3	<i>C. annuum</i>
F6-074	<i>C. annuum</i>
F1 Baja	<i>C. annuum</i>
Caman	<i>C. annuum</i>
Adelina	<i>C. annuum</i>
IPBC333	<i>C. frutescens</i>
IPBC290	<i>C. frutescens</i>
Bonita	<i>C. frutescens</i>
Taruna	<i>C. frutescens</i>
Cakra Putih	<i>C. frutescens</i>
Red Habanero	<i>C. chinense</i>
Jolokia	<i>C. chinense</i>
Fatali	<i>C. chinense</i>
Red Chupetinho	<i>C. chinense</i>
Bishop Crown	<i>C. chinense</i>
Lemon Drop	<i>C. baccatum</i>

### Data analysis

For qualitative characters, data were presented descriptively, and cluster analysis was performed. Quantitative character's data were subjected to analysis of variance, estimation of genetic parameters, and heritability estimates. The analysis of variance was conducted following the procedure of Steel and Torrie (1981). Estimation of genetic parameters and heritability was done by following methods of Lush (1949) and Johnson et al. (1955) (Table 2):

$$\begin{aligned}\sigma^2_G &= (M2-M3) / r \\ \sigma^2_e &= M3 \\ \sigma^2_P &= \sigma^2_G + \sigma^2_e \\ h^2_{bs} &= (\sigma^2_G / \sigma^2_P) \times 100\%\end{aligned}$$

Criteria of heritability (%) (Whirter 1979) are: low ( $0 < X < 20\%$ ), moderate ( $20 \leq X < 50\%$ ) and high ( $X \geq 50$ ).

## RESULTS AND DISCUSSION

### Qualitative character performance

Seventeen qualitative characters were observed, which showed a polymorphic pattern among genotypes. These characters included seedlings, stems, leaves, flowers, and fruit characters. Seedling characters were hypocotyl color consisted of green (23 genotypes), white (5 genotypes), and purple (1 genotype). Cotyledonous leaf shape comprised of lanceolate (18 genotypes), deltoid (2 genotypes), elong-deltoid (7 genotypes), and ovate (2 genotypes). Cotyledonous leaf color ranged from green to yellowish-green. Leaf shape fell into lanceolate (7 genotypes), deltoid (7 genotypes), and ovate (15 genotypes).

The flower position consisted of erect (15 genotypes), intermediate (5 genotypes), and pendant (9 genotypes). Fruit position comprised of erect (11 genotypes), intermediate (5 genotypes), and pendant (13 genotypes) while corolla color was white (21 genotypes), greenish-white (6 genotypes), and purple (2 genotypes). Sepals color fell into green (27 genotypes) and purple (2 genotypes) while anther color was categorized into gray (11 genotypes), greyish-green (1 genotype), purple (3 genotypes), green (7 genotypes), purplish-grey (1 genotype), purplish-white (4 genotypes), and yellow (2 genotypes). Number of Corolla: 5-crowns (11 genotypes), 6-crowns (11 genotypes), 5/6-crowns (6 genotypes), and 7-crowns (1 genotype).

Fruit shape was classified into elongate (14 genotypes), triangular (4 genotypes), campanulate (5 genotypes), and blocky (6 genotypes). Fruit shape at pedicle attachment included acute (1 genotype), obtuse (15 genotypes), truncate (7 genotypes), cordate (1 genotype), and lobate (5 genotypes). Fruit shape at blossom end consists of pointed

(19 genotypes), blunt (3 genotypes), sunken (5 genotypes), and sunken and pointed (2 genotypes). Fruit cross-sectional corrugation fell into slightly corrugated (24 genotypes), intermediate (1 genotype), and corrugated (4 genotypes). The fruit surface was smooth (20 genotypes), semi-wrinkled (8 genotypes), and wrinkled (1 genotype).

Cluster analysis was performed involving 17 polymorphic characters and they are presented in Figure 1. Cluster analysis showed that the 29 genotypes used were generally divided into two main clusters and several sub clusters. Partition of the clusters based on the similarity level of 0.625 produced five different clusters. Cluster I consisted of 22 genotypes from 3 different species, i.e., *C. annuum* ('Yuni', 'SSP', 'IPBC3', 'F6-074', 'IPBC12', 'Anies', 'IPBC5', 'F1 Baja', 'Caman', 'Seloka', 'Adelina', 'IPBC15', 'F5-012328-6-2-2', 'F5-012328-1AB-2-1', 'F5-012328-6-2-1', 'F4-012328-1AB-3', 'F4-012328-3-3', and 'IPBC333'), and *C. frutescens* ('IPBC333', 'IPBC290', 'Bonita', and 'Taruna'). Cluster II consisted of four genotypes from *C. chinense* species ('Fatali', 'Jolokia', 'Red Chupetinho', and 'Red Habanero'). Cluster III, Cluster IV, and Cluster V each consisted of one genotype, namely, 'Ungara' (*C. annuum*), 'Bishop Crown' (*C. chinense*), and 'Lemon Drop' (*C. baccatum*), respectively.

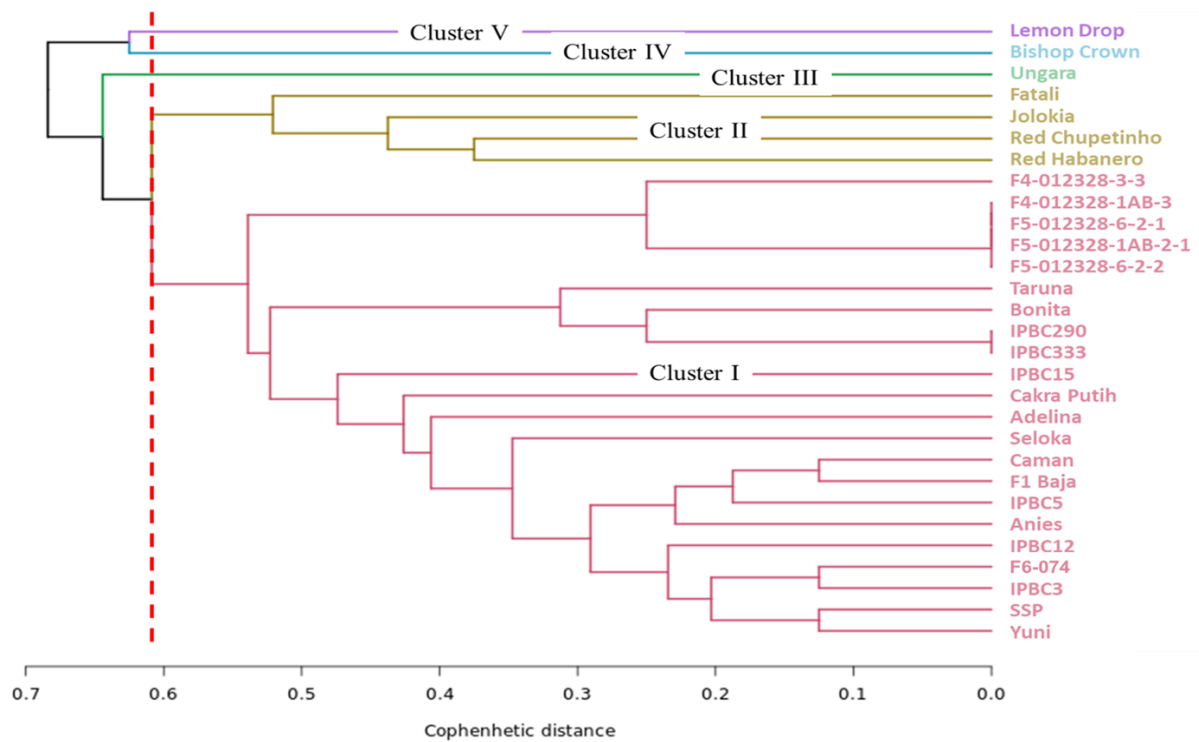
### Genetic variability

Analysis of variance revealed a significant genotype effect on all observed variables (Table 3). This result shows that there is a diversity of characters among the tested genotypes. The wide range of each character supports this claim. The mean value of the leaf length ranged from 5.00 cm ('Yuni') to 15.96 cm ('White Chakra'). The average leaf width ranged from 1.94 cm ('Yuni') to 9.34 cm ('Bishop's Crown'). The stem diameter ranged from 0.46 cm ('Caman') to 0.88 cm ('IPBC290' and 'Fatali'). The mean plant height ranged from 45.26 cm ('Ungara') to 126.67 cm ('IPBC290'). The fruit length ranged from 1.89 cm ('Ungara') to 16.15 cm ('Caman'). The diameter of the test genotype fruit ranged from 0.78 cm ('Yuni') to 3.92 cm ('Bishop Crown'). The observed fruit weights ranged from 1.30 g ('Red Chupetinho') to 15.36 g ('F4-012328-1AB-3'). The number of fruits per plant of the tested genotype under PYLCV-infected conditions ranged from 0 ('Adelina') to 42.6 ('Red Chupetinho'). The yield per plant of the tested genotype varied from 0 g ('Adelina') to 69.98 g ('Anies'). Disease severity index varied from 0% ('Adelina', 'Red Habanero', 'Red Chupetinho') to 75.0% ('Lemon Drop'). The coefficient of variance (CV) is the ratio of the standard deviation to the mean of a variable (Pena-Yam et al. 2019). The coefficient of variation from the test results shows a low value of less than 20% (Gomez and Gomez 1987) on all variables except for the disease severity index (25.51%), which indicates that the test results are valid.

**Table 2.** F-Ratios was used to test the effects for randomized complete block experiments

Sources of variation	Mean square	Expected mean squares (fixed model)
Replication (Rep)	M1	-
Genotype (Geno)	M2	$\sigma^2_e + r \sigma^2_G$
Error	M3	$\sigma^2_e$

Note:  $\sigma^2_e$ : environmental variance;  $\sigma^2_G$ : genotypic variance



**Figure 1.** Dendrogram showing the relationships among chili pepper genotypes using 17 qualitative traits

The results of the analysis of variability and heritability are presented in Table 4. The value of genotypic variance is greater than twice the standard deviation value ( $\sigma^2_G > 2 \sigma (\sigma^2_G)$ ). This result shows that the genetic diversity of each character is quite wide. The value of the observed phenotypic variance is greater than the genotypic variance, so it can be seen that the diversity of chili plant characters is not only influenced by genetic factors but is also influenced by the environment. To estimate the proportion of the influence of genetic factors on plant phenotype characters, an approach that can be taken is to estimate the heritability value of the related characters. The heritability value obtained from testing for all characters is high, ranging from 65.16 to 99.12%. Heritability value indicates the proportion of genetic factors in influencing the performance of a character (phenotype). The high

heritability value can confirm that the genetic diversity most influence the observed phenotype variation in each tested genotype.

The genotypic coefficient of variance (GCOV) and phenotypic coefficient of variance (PCOV) are presented in Table 5. The results of the analysis showed that PCOV values were higher than GCOV in all characters, indicating that the diversity of characters in plants is influenced by both genetic and environmental factors (Terfa and Gurmu 2020). All observed characters had high GCOV values (>20%) except for stem diameter, which was in the moderate category (15.77%). GCOV and PCOV values are grouped into three categories: low (<10%), moderate (10-20%), and high (>20%) (Terfa and Gurmu 2020; Hamidou et al. 2018).

**Table 3.** Estimation of the mean square for different parameters of *Begomovirus* infected chili pepper

Character	Range	Mean Square		CV (%)
		Rep	Geno	
Leaf length (cm)	5.00-15.96	0.0097 <sup>ns</sup>	24.1346 <sup>**</sup>	3.36
Leaf width (cm)	1.96-9.34	0.0147 <sup>ns</sup>	12.4521 <sup>**</sup>	3.41
Stem diameter (cm)	0.46-0.88	0.0015 <sup>ns</sup>	0.0351 <sup>**</sup>	5.39
Plant height (cm)	45.26-126.67	13.3544 <sup>ns</sup>	1853.5198 <sup>**</sup>	4.65
Fruit length (cm)	1.89-16.15	0.589 <sup>ns</sup>	40.9159 <sup>**</sup>	14.47
Fruit diameter (cm)	0.78-3.92	0.0258 <sup>ns</sup>	3.292 <sup>**</sup>	11.35
Fruit weight (g)	1.30-15.36	0.1513 <sup>ns</sup>	53.9417 <sup>**</sup>	9.04
Number of fruit per plant	0.00-42.60	1.2835 <sup>ns</sup>	268.0011 <sup>**</sup>	17.59
Yield per plant (g)	0.00-68.98	15.0792 <sup>ns</sup>	594.7606 <sup>**</sup>	15.91
Disease severity (%) <sup>T</sup>	0.00-75.00	0.0218 <sup>ns</sup>	0.0767 <sup>**</sup>	25.51

Note: Geno: Genotype; Rep: Replication; CV: Coefficient of Variance <sup>\*\*</sup> significant at level of 1%; 5%; <sup>ns</sup>: not significant; <sup>T</sup> = transformed data is used

**Table 4.** Genetic variability and heritability of *Begomovirus* infected chili pepper

Character	$\sigma^2_G$	$2\sigma(\sigma^2_G)$	Criteria of $\sigma^2_G$	$\sigma^2_P$	$h^2_{bs}$ (%)	Criteria of $h^2_{bs}$
Leaf length	7.99	7.08	Wide	8.14	98.19	High
Leaf width	4.13	3.65	Wide	4.18	99.12	High
Stem diameter	0.01	0.01	Wide	0.01	88.91	High
Plant height	613.08	543.63	Wide	627.35	97.72	High
Fruit length	13.37	12.00	Wide	14.16	94.39	High
Fruit diameter	1.08	0.97	Wide	1.13	96.03	High
Fruit weight	17.82	15.82	Wide	18.30	97.40	High
Number of fruit per plant	88.00	78.61	Wide	91.99	95.66	High
Yield per plant	187.83	174.56	Wide	219.09	85.73	High
Disease severity	0.02	0.01	Wide	0.03	65.16	High

Note:  $\sigma^2_P$ : phenotypic variance;  $\sigma^2_G$ : genotype variance;  $\sigma(\sigma^2_G)$ : stdev of genotypic variance;  $h^2_{bs}$ : broad sense heritability

**Table 5.** Estimated genotypic and phenotypic coefficient of variation of *Begomovirus* infected chili pepper

Character	GCOV (%)	Criteria of GCOV	PCOV (%)	Criteria of PCOV
Leaf Length	24.84	High	25.06	High
Leaf Width	36.12	High	36.28	High
Stem Diameter	14.87	Moderate	15.77	Moderate
Plant Height	30.51	High	30.86	High
Fruit Length	59.41	High	61.15	High
Fruit Diameter	55.85	High	57.00	High
Fruit Weight	80.16	High	81.22	High
Number of Fruit per Plant	82.60	High	84.45	High
Yield per Plant	39.02	High	42.14	High
Disease Severity	34.91	High	43.24	High

Note: PCOV: phenotypic coefficient of variance; GCOV: genotypic coefficient of variance

### Plant resistance level

The incubation period and disease severity of the 29 genotypes tested are shown in Table 6. The results showed a diversity of resistance levels among the tested genotypes. The PYLCV resistance level of chili peppers were divided into five categories, i.e., resistant, moderately resistant, moderately susceptible, susceptible, and highly susceptible.

The incubation period of the disease in each genotype varied from 5 to 45 DAI. The genotypes 'Adelina', 'Red Habanero', and 'Red Chupetinho' showed no symptoms of PYLCD during the observation period. Of the 29 genotypes tested, thirteen genotypes were classified into the Resistant category ('Adelina', 'Red Habanero', 'Red Chupetinho', 'F6-074', 'Jolokia', 'Fatali', 'F5-012328-6-2-1', 'Anies', 'F4-012328-1AB-3', 'F5-012328-1AB-2-1', 'Camán', 'IPBC3', and 'IPBC290'), nine genotypes were categorized as moderately resistant ('F1 Baja', 'Seloka', 'Bonita', 'IPBC12', 'F4-012328-3-3', 'Ungara', 'F5-012328-6-2-2', 'IPBC15', and 'Cakra Putih'), two genotypes were classified in moderately susceptible category ('Taruna' and 'IPBC333'), three genotypes were in the Susceptible category ('SSP', 'Bishop Crown', and 'IPBC5') and two genotypes were in the highly susceptible category ('Yuni' and 'Lemon Drop').

The yield reduction in resistant genotypes, namely 'Adelina' (*C. annuum*) (100%), 'Red Habanero' (55.80%), and 'Red Chupetinho' (47.63%), was relatively high. The Jolokia and Bishop Crown (*C. chinense*) genotypes showed a minor decrease in yield per plant, which only reached

20.87% and 21.81% respectively.

### Discussion

In assembling disease-resistant plants, apart from focusing on plant resistance, other traits that are not directly related also need to be developed to support increased yields and product quality. Sources of diversity are increasingly being explored, not only limited to the same species or commodity but also from other commodities (Beam and Ascencio-Ibanez 2020). Evaluation of various species becomes important to be carried out to obtain good sources of morphological, agronomic, and resistance character.

Evaluation of morphological and agronomic characters, in this study, was carried out on plants that had been inoculated with PYLCV. Qualitative characters were observed on normal plant organs that did not show symptoms of PYLCD. Qualitative characters are generally controlled by major genes and only slightly influenced by the environment (Seprico 2020), so viral infection is thought not to change these qualitative characters much. For the quantitative characters, plant characters that appear in the phenotype result from the influence of plant genetics and the environment. To determine the proportion of genetic influence in the diversity of characters, it can be seen through the heritability values of each observed character. High heritability values were observed in all the characters studied in this experiment. Heritability is the degree of genetic control associated with some important

traits (Usman et al. 2014) that indicates how much of the genetic variability has a genetic origin and gives the necessary information for the genetic selection process (Usman et al. 2017). This high value shows that most of the diversity that exists in the population is influenced by genetic factors.

In addition, in this study, genotypes from four species were used. These species differences led to a wide morphological diversity among the test genotypes. This diversity is shown in the heritability value. Compared to other characters, the disease severity character has a lower heritability value (65.16%). This is presumably because environmental influences have more roles in influencing the disease severity compared to other characters. According to (Sing et al. 2016), the disease severity in plants does not only depend on genetic factors related to plant resistance. Environmental factors, the interaction of insect vectors with the host, and viral recombination can significantly change plant resistance and affect the development of disease symptoms (Rubio et al. 2020). This assumption is supported by a high CV value, which indicates that factors other than genetics significantly affect disease severity.

The difference between PCOV and GCOV shows how much the environment influences the observed characters. In this study, the difference between PCOV and GCOV was small in all characters, which indicates that the characters were slightly influenced by environmental factors, except for the disease severity character, which has a large difference between PCOV and GCOV. Since PCOV estimates the genotypes and environment effects, the higher difference between GCOV and PCOV indicates a significant contribution of the environment through the interaction between genotypes and environment. This experiment cannot observe these interactions. The highest GCOV and PCOV values were obtained on fruit weight and the number of fruits per plant. The same results were also obtained in a study conducted by Paul et al. (2017) in Linseed plant (*Linum usitatissimum* L.), Adhikari et al. (2018) in rice (*Oryza sativa*), and Pena-Yam et al. (2019) and Ajith and Manju (2006) in chili, thus supporting the results of this study. The high GCOV supported by high heritability indicates that the diversity of characters in the population was high, and the selection of these characters should be possible (Hamidou et al. 2018).

**Table 1.** Resistance level and productivity of 29 chili pepper infected by Pepper Yellow Leaf Curl Virus

Genotype	Incubation period (DAI)	Disease severity index (%)	Yield per plant	Percentage reduction of yield per plant	Category
Adelina	-	0.00	0.00 <sup>p</sup>	100.00 <sup>a</sup>	Resistant
Red Habanero	-	0.00	34.45 <sup>f-k</sup>	55.80 <sup>g-k</sup>	Resistant
Red Chupetinho	-	0.00	48.09 <sup>cde</sup>	47.63 <sup>i-l</sup>	Resistant
F6-074	22-26	3.33	48.85 <sup>cde</sup>	60.18 <sup>e-j</sup>	Resistant
Jolokia	38	3.33	23.81 <sup>k-n</sup>	20.87 <sup>m</sup>	Resistant
Fatali	35	3.33	18.93 <sup>mn</sup>	34.80 <sup>lm</sup>	Resistant
F5-012328-6-2-1	39	5.83	34.17 <sup>f-k</sup>	51.78 <sup>h-k</sup>	Resistant
Anies	16	6.67	68.98 <sup>a</sup>	52.46 <sup>h-k</sup>	Resistant
F4-012328-1AB-3	28	7.50	52.87 <sup>bcd</sup>	46.80 <sup>i-l</sup>	Resistant
F5-012328-1AB-2-1	45	7.50	43.77 <sup>def</sup>	63.73 <sup>d-i</sup>	Resistant
Caman	16-21	8.33	30.75 <sup>h-l</sup>	62.04 <sup>d-i</sup>	Resistant
IPBC3	12	10.00	29.16 <sup>j-m</sup>	75.94 <sup>b-e</sup>	Resistant
IPBC290	15	10.00	28.95 <sup>j-m</sup>	64.55 <sup>c-h</sup>	Resistant
F1 Baja	12-13	10.83	55.47 <sup>bc</sup>	53.92 <sup>h-k</sup>	Moderately resistant
Seloka	16-18	11.67	42.57 <sup>efg</sup>	77.43 <sup>b</sup>	Moderately resistant
Bonita	15	13.33	41.38 <sup>efg</sup>	46.93 <sup>ijkl</sup>	Moderately resistant
IPBC12	12-19	15.00	35.59 <sup>f-j</sup>	54.92 <sup>g-k</sup>	Moderately resistant
F4-012328-3-3	24	15.00	18.57 <sup>mn</sup>	71.69 <sup>b-g</sup>	Moderately resistant
Ungara	22-25	16.67	30.49 <sup>h-l</sup>	31.71 <sup>lm</sup>	Moderately resistant
F5-012328-6-2-2	25-27	16.67	39.25 <sup>e-i</sup>	42.26 <sup>kl</sup>	Moderately resistant
IPBC15	15-18	18.33	17.59 <sup>n</sup>	80.24 <sup>bc</sup>	Moderately resistant
Cakra Putih	12	20.00	26.06 <sup>f-n</sup>	57.76 <sup>f-k</sup>	Moderately resistant
Taruna	12	20.83	20.77 <sup>lmn</sup>	62.62 <sup>d-j</sup>	Moderately susceptible
IPBC333	12	25.00	25.04 <sup>i-n</sup>	66.53 <sup>c-h</sup>	Moderately susceptible
SSP	26-29	30.83	28.70 <sup>j-m</sup>	70.78 <sup>c-g</sup>	Susceptible
Bishop Crown	26	31.67	40.38 <sup>e-h</sup>	21.81 <sup>m</sup>	Susceptible
IPBC5	25	44.17	59.35 <sup>b</sup>	71.58 <sup>b-g</sup>	Susceptible
Yuni	5-8	51.67	32.34 <sup>g-k</sup>	73.41 <sup>b-f</sup>	Highly susceptible
Lemon Drop	41	75.00	6.95 <sup>o</sup>	87.97 <sup>ab</sup>	Highly susceptible

Note: Number followed by the same letter in the same column were not significantly different to DMRT 5% level; DAI: Day After Inoculation; -: no symptoms appear until the end of the observation period

The resistance characters observed in this study included the incubation period and the disease's severity. The incubation period (the time between inoculation and symptom expression) and latent period (the time between inoculation and infectiousness of the house) have not been widely documented. If disease control is based on visual symptoms, the incubation period is important because it is related to the time of symptom appearance (Rimbaud et al. 2015). Control can be done quickly when the virus incubation period in cultivated plants is quickly seen. On the other hand, visual symptoms tend to be attractive to vectors. Viruses that cause visual symptoms can cause changes in the production and emission of volatile organic compounds or changes in nutrients and defense chemistry, which are important cues for insect vectors and affect vector feeding (Mauck et al. 2012). That is how the incubation period is thought to have a role in disease spread in the field.

Three genotypes ('Adelina', 'Red Habanero', and 'Red Chupetinho') showed no PYLCD symptoms until the end of the observation period. Plants may not show symptoms after being infected with the virus if they have complete resistance or partial resistance to prevent the development of the virus in plant tissues (Rubio et al. 2020). Fruit weight per plant in conditions infected with *Begomovirus* generally decreased compared to control (virus-free plants). 'Adelina' has no fruit until the end of the observation period due to flower loss and fruit formation failed to occur. This phenomenon is thought to be one of the consequences of PYLCD attacks, where Rusli et al. (1999) reported that one of the symptoms of PYLCD is flower loss. In addition, according to Harth et al. (2016), pollen reduction will occur in virus-infected plants so that fertilization does not happen, and no fruit is formed.

'Jolokia' is included in resistant plants with a long incubation period (38 DAI) and low DSI (3.33%). These results align with Adluri et al. (2017) who stated that the 'Bhut Jolokia' (*C. chinense*) genotype was thought to have good resistance to PYLCV. The Bishop Crown genotype was categorized as susceptible but under PYLCV-infected conditions, its yield reduction per plant was relatively low. Four of the five chili genotypes of the tested *C. chinense* species ('Red Habanero,' 'Red Chupetinho', 'Jolokia' and 'Fatali') had good resistance to PYLCV. These *C. chinense* species genotypes can be used as a source of resistance against PYLCV.

The genotype 'IPBC12' was included in the moderately resistant category, which in previous studies (Ganefianti 2010; Ayu 2019), 'IPBC12' included in the resistant category. This change in the resistance levels can be caused by several factors including changes in plant genetics (López-Fabuel et al. 2013) and differences in the infecting virus strains (Miras et al. 2014). The genotype 'F1 Baja' (*C. annum*) was found to be moderately resistant in the present study, which is in line with its varietal description in the Decree of the Ministry of Agriculture Number 014/kpts/SR.120/D.2.7/2/2016. Out of 29 chili genotypes evaluated, two genotypes from the *C. annum* species show a better resistance level than the 'F1 Baja', i.e., 'F6-074' and 'Anies'. In addition, 'Bonita' (*C. frutescens*) can also

be considered to have better resistance than the other genotype from *C. frutescens*, and it has a lower yield reduction per plant. These three genotypes also can be used as sources of chili pepper resistance against PYLCD.

This study revealed a wide genetic diversity among the tested genotypes in almost all the observed variables, thus enabling the breeders to select genotypes with desirable characters from the existing population. Most of the genotypes from *C. chinense* have good resistance and can be developed as a source of chili pepper resistance against PYLCV. The genotypes 'Red Habanero', 'Red Chupetinho', 'Jolokia', and 'Fatali' were resistant to PYLCV while the 'Bishop Crown' was classified susceptible but still be able to produce well. In general, by considering the resistance level and other characters such as productivity, 'Jolokia' (*C. chinense*), 'Anies' (*C. annum*) and 'Bonita' (*C. frutescens*) genotypes can be recommended as candidate sources of good resistance to PYLCV.

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