Effect of chemical mutagens on growth of Okra (Abelmoschus esculentus L. Moench)

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Abstract
The purpose of the present study was to evaluate the effects of mutagens i.e. Ethyl methanesulfonate (EMS) and Sodium azide (SA) on the morphological growth of Okra. Seeds of two Okra varieties Subzpari and Pahuja were first pre-soaked in distilled water and then treated with 0.0 (Control), 0.1, 0.2 and 0.3% Sodium azide (SA) and Ethyl methanesulfonate (EMS) separately. The results showed that the germination rate was significantly decreased with increased lethality (%) and mutation frequency (%) in M1 of both varieties with increasing concentrations of EMS and SA against control. The results showed that different concentrations of EMS and SA affected morphological attributes of both okra varieties as the increase of mutagen concentration showed variable results in M1 generation compared with control, but M2 generation showed useful improvement in various traits. Such as the M2 generation of Subspari from 0.1% SA treatment showed a decrease in flowering time while an increase in the number of fruits per plant, fruit size, seeds yield (g) per plant and 100 seeds weight.

Keywords: Ethyl methane sulfonate; Growth; Induced mutagenesis; Okra; Sodium azide

Introduction
Okra [Abelmoschus esculentus (L.) Moench, 2n = 2x=130] is one of the important vegetables belongs to Malvaceae family and known by many names in different regions of the world as, Gumbo in the USA, ladyfinger in England, Quingombo in Spanish, Huang Gu Kui in China, Bhindi in India and Pakistan, Dherosh in Bangladesh. Okra plant is native to northern Africa from Ethiopia and Sudan. It is summer and rainy season crop and is widely cultivated from tropics to subtropics. Okra (Abelmoschus esculentusL.) is probably an amphidiploid (allotetraploid) derived from Abelmoschus tuberculatus with 2n = 58 and Abelmoschus ficulneus (L.) Wight and Arn. ex Wight with 2n = 72 chromosomes. Another edible okra species Abelmoschus caillei (A. Chev.) is found in the Central and West Africa. There are strong indications that Abelmoschus caillei is amphidiploids with Abelmoschus esculentus being one of the parental species. The lowest chromosome number2n=56 reported in Abelmoschus angulosus whereas, the highest
chromosome number 2n=196 reported in *Abelmoschus manihot* var. Caillei [1,2]. Fresh Okra fruits are a good source of carbohydrates, proteins, oil, potassium, magnesium, calcium, iodine, vitamins A and C, etc [3,4]. The Okra crop has many uses; fresh green pods are eaten as a vegetable, immature pods are fried, steamed, canned or pickled, stem and mature fruits are used in the paper industry. It is very useful about antiulcer, anticancer and hypoglycemic activities [5].

Mostly conventional breeding methods are employed for the improvement of crops and natural variability in the germplasm. This takes a long time to select genetic variability within local crops [6, 7]. To overcome this situation, induced mutagenesis can be used as an alternative method to create variability in different quantitative traits. Mutations are sudden genetic changes in living cell DNA that is neither genetically isolated nor genetically recombined [8]. According to Drake and others [9], mutations occur in the eukaryotic genome as 10⁸ base pair per generation. Mutations can be induced artificially by using physical and chemical mutagens. Induced mutations have been used as an important tool for the improvement of certain traits in the existing germplasm. After Muller [10] discovered the mutagenic effects of X-rays on the fly, induced mutations were used for plant breeding. Ethyl Methanesulfonate is a carcinogenic organic compound, a popular and powerful mutagen that produces random point mutations in the genomes through nucleotide substitutions [11,12]. Therefore, okra is of high nutritional value with medical applications and economic importance hence needs to improve the production, quality and quantity of fruit. Taking into account the importance of okra, the present study was done with the objectives of evaluating the germination and growth attributes of M₁ and M₂ generations okra through the treatment of sodium azide (SA) and ethyl methane sulfonate (EMS) under different concentrations.

**Materials and methods**

Seeds of two varieties of Okra (Pahuja and Subzpari) were obtained from registered seed companies, Hyderabad, Pakistan. The experiment was performed during the year 2015-16 at IBGE, University of Sindh, Jamshoro, Pakistan.

**Treatment of Okra seeds**

For chemical mutagenesis, 0.1% (0.5g/50ml), 0.2% (1.0g/50ml) and 0.3% (1.5g/50ml) solutions of ethyl methane sulfonate (ACROS) and sodium azide (ACROS) were prepared separately. For each treatment, 250 dry, healthy and uniform sized seeds of both okra varieties; Pahuja and Subzpari were initially presoaked in 100 ml distilled water in the lab at room temperature for six hours to ensure complete hydration. After removing the excess water presoaked 60 seeds of each Pahuja and Subzpari varieties were submerged in 50 ml solutions containing sodium azide and ethyl methane sulfonate separately with shaking (70rpm) in the dark for 60 minutes at room temperature [13]. After chemical treatment, seeds were washed with tap water and were sown in pots (5 seeds/pot) containing 10kg of sand and clay (1:1 ratio). For control, untreated seeds were soaked in distilled water for seven hours. The experiment was repeated twice and the total 120 seeds were used for each treatment. For M₂ generation study, mature, dry and uniform seeds of each treatment and control obtained from M₁ generation were grown in pots under the same soil conditions as in M₁ generation. The experiment was performed two times, thus a total 120 seeds for each treatment in the M₂ generation study. The mutagenic effect of ethyl methanesulfonate (EMS) and sodium azide (SA) on M₁ and M₂ generations of both okra varieties was assessed. Various growth attributes including germination (%), shoot and root lengths, stem diameter, number of branches, flowering time, number of fruits,
fruit length, seeds per fruit and 100 seeds weight were observed. The germination of seeds was observed after every week for three weeks and the germination percentage was calculated after three weeks by the formula as follows:

\[
\text{Germination (\%)} = \frac{\text{Number of seeds germinated}}{\text{Total number of seeds sown}} \times 100
\]

The survival of plants was noted after 30 days of sowing the lethality percentage was calculated as:

\[
\text{Lethality (\%)} = \frac{\text{Number of plants died}}{\text{Total number of seeds germinated}} \times 100
\]

The mutation frequency of ethyl methane sulfonate and sodium azide was calculated according to the reported method \[14\] as:

\[
\text{Mutation frequency (\%)} = \frac{\text{Viable mutants observed (V)}}{\text{Total plants studied (P)}}
\]

The shoot length of each plant was measured after 30 days of sowing and expressed in cm while root length was measured after harvesting of fruits and expressed in cm. The shoot diameter was measured on maturity time while number of branches was counted at maturity. The fruit size was measured at maturity, number of fruit per plant, number of seeds per fruit and 100 seeds weight was determined.

The experiment was performed according to the Randomized Complete Block Design (RCBD) manner. The collected data were expressed as the mean of triplicates ± standard deviation (SD) which was calculated.

\[
S = \sqrt{\frac{1}{N-1} \sum_{i=1}^{N} (x_i - \bar{x})^2}
\]

S=sample standard deviation
N=the number of observations
\(x_i=\)the observed values of a sample item
\(\bar{x}=\)the mean value of the observations

**Results and discussion**
Germination of seeds is a key parameter used to evaluate the impact of the mutagen on plants. The inhibition of seed germination after the treatment of seeds with various mutagens is a very effective method to study the effects of mutagens on plants.

The germination of seeds was found to be the highest in control plants as compared to other treated plants during M₁ and M₂ generations which showed that mutagenic treatments inhibited the germination in both varieties. The results revealed that germination was significantly reduced as the dose of mutagen was increased. The lowest germination response was noted in M₁ plants of Subs pri when seeds were treated with 0.3% of EMS and SA as 45% and 40% respectively compared to control which showed 90% germination (Table 1). Similarly, both varieties Subs pri and Puhja showed sensitivity to higher doses of mutagens as 38% and 40% lethality observed in 0.3% of SA and EMS respectively in Subs pri and 40% and 47% lethality in 0.3% SA and EMS treated M₁ plants of Puhja. Similarly, the highest % of survived mutants was in 0.1% EMS and SA treated plants of Subspri and Puhja with 92% and 93% mutant frequency respectively (Table 1). On the other hand, seed germination response was good in all seeds obtained from M₁ plants which showed 59% to 75% germination response in M₂ plants compared with control 84-88% germination (Table 1). Similarly, M₂ plants showed less lethality and mutant frequency than M₁ plants. Seed germination is a significant trait for measuring the response of plants to mutagenic treatment \[15\]. As observed in the present study, Etheret et al.\[16\] also reported the promotion of biological parameters by low-dose SA and EMS. Contrary to our results, seed germination was reduced to all doses of sodium azide in Eruca sativa L \[17\] and in Helianthus annuum \[18\]. Delays or inhibition of physiological activities including enzymatic activity, hormonal imbalance and mitotic activity essential for germination of seeds have explained the reduction in germination caused by mutagenesis \[17, 19\]. The results revealed
that there is a linear association between increased doses of mutagens (SA and EMS) and reduced germination of seeds. This was also reported in black gram [20], wheat [21] and Okra [22]. The shoot length in two okra varieties was observed to exhibit marginal differences in mutagens treated and control. The effect of EMS on shoot length of Subz pari was noted as a significant increase in shoot length (17.2±2.1 to 19.3±1.9cm) was observed in 0.1 & 0.2% EMS treated M₁ and M₂ plants compared with control (15.5±1.2 cm). On the other hand, 0.3% SA treated M₁ and M₂ plants of Puhja showed a significant decrease in shoot length as 10.5±2.12 and 10.76±2.5cm respectively compared to control (15.0±1.5cm) as shown in table 2.

Table 1. Effect of EMS and SA treatments on germination of Subz Pari and Pahuja varieties of Okra after 30 days of sowing

| Treatments | Germination (%) | Lethality (%) | Mutation Frequency (%) |
|------------|----------------|--------------|------------------------|
|            | Subz Pari | Pahuja | Subz Pari | Pahuja | Subz Pari | Pahuja | Subz Pari | Pahuja |
| Control    | 90 | 88 | 83 | 84 | 2 | 1.89 | 2 | 0.99 | ND | ND | ND | ND |
| EMS (%)    | 0.1 | 80 | 85 | 73 | 78 | 7 | 1.94 | 6 | 2 | 92 | 98 | 93 | 97 |
|            | 0.2 | 58 | 78 | 66 | 65 | 15 | 2 | 20 | 2 | 84 | 97 | 80 | 97 |
|            | 0.3 | 45 | 72 | 53 | 59 | 38 | 6 | 40 | 5 | 61 | 93 | 59 | 94 |
| SA (%)     | 0.1 | 86 | 88 | 80 | 80 | 7 | 3 | 12 | 3 | 92 | 97 | 87 | 96 |
|            | 0.2 | 50 | 76 | 56 | 81 | 19 | 4 | 22 | 6 | 80 | 95 | 77 | 93 |
|            | 0.3 | 40 | 75 | 42 | 72 | 40 | 8 | 47 | 10 | 59 | 91 | 52 | 89 |

Table 2. Effect of EMS and SA treatments on the shoot and root lengths, number of branches and stem diameter of Subz Pari and Pahuja varieties of Okra after 30 days against control Mean±S.D)

| Treatments | M₁ generation | Subz Pari |
|------------|---------------|-----------|
|            | Shoot Length (cm) | Root Length (cm) | No of branches/plant | Stem diameter (cm) | Shoot Length (cm) | Root Length (cm) | No of branches/plant | Stem diameter (cm) |
| Control    | 14.8±1.5 | 19.6±0.9 | 10.0±1.9 | 2.52±0.5 | 15.8±1.2 | 20.4±0.57 | 11.4±0.5 | 2.7±0.4 |
| EMS (%)    | 0.1 | 17.1±1.9 | 22.1±1.7 | 10.2±1.7 | 3.23±0.3 | 19.3±1.9 | 23.5±0.7 | 12.5±1.2 | 2.5±0.5 |
|            | 0.2 | 13.6±1.1 | 20.7±1.2 | 10.6±1.7 | 3.1±0.2 | 17.4±2.1 | 22.5±0.5 | 11.6±1.4 | 2.8±0.7 |
|            | 0.3 | 13.2±2.1 | 18.2±2.4 | 10.5±2.1 | 2.4±0.7 | 15.8±2.8 | 20.5±2.12 | 10.3±1.6 | 2.3±0.3 |
| SA (%)     | 0.1 | 11.1±1.2 | 24.0±2.1 | 10.8±1.8 | 2.9±0.6 | 12.6±2.3 | 20.4±2.3 | 11.2±1.7 | 2.85±0.7 |
|            | 0.2 | 11.4±1.2 | 19.5±0.7 | 9.5±1.8 | 3.17±0.2 | 11.9±2.7 | 17.4±1.53 | 10.8±1.6 | 2.6±0.5 |
|            | 0.3 | 10.5±2.1 | 13.0±1.4 | 9.4±1.2 | 3.1±0.12 | 12.8±1.3 | 21.0±2.3 | 10.6±1.8 | 2.08±0.4 |

| Treatments | M₂ generation |
|------------|---------------|
|            | Shoot Length (cm) | Root Length (cm) | No of branches/plant | Stem diameter (cm) |
| Control    | 15.0±1.5 | 20.2±0.9 | 10.2±0.6 | 2.28±0.5 | 15.5±1.2 | 21.6±0.57 | 11.8±1.9 | 2.58±0.7 |
| EMS (%)    | 0.1 | 15.9±1.3 | 19.7±2.1 | 11.1±1.7 | 2.5±0.6 | 17.9±2.6 | 22.5±0.7 | 14.5±2.1 | 2.07±0.7 |
|            | 0.2 | 14.2±2.1 | 19.3±1.8 | 10.8±0.8 | 2.43±0.6 | 17.2±2.1 | 21.5±1.2 | 12.5±1.6 | 2.43±0.5 |
|            | 0.3 | 12.5±1.3 | 17.8±2.7 | 11.2±1.1 | 2.3±0.7 | 15.9±2.1 | 18.8±0.92 | 10.5±0.7 | 2.24±0.6 |
| SA (%)     | 0.1 | 13.5±1.2 | 21.5±1.4 | 10.5±0.7 | 2.69±0.4 | 13.6±1.1 | 9.67±2.08 | 13.3±0.7 | 2.53±0.4 |
|            | 0.2 | 11.8±0.6 | 18.2±0.7 | 12.0±1.6 | 2.5±0.6 | 14.0±1.4 | 20.3±2.4 | 11.1±1.8 | 2.28±0.5 |
|            | 0.3 | 10.7±2.5 | 17.8±1.4 | 10.3±0.6 | 2.5±0.5 | 12.8±2.7 | 16.67±2.1 | 10.6±1.2 | 2.31±0.4 |
The root length in two okra varieties of EMS-treated plants increased slightly in M₁ and M₂, while the SA concentration was a little different in both M₁ and M₂ generations (Table 2). As the concentration of SA and EMS increases, the inhibitory effect of the mutagen decreased the root length. The decrease in root and shoot length is the effect of chemical mutagens on physiological systems [23]. The reduction in length of shoots and roots are due to mutagenesis has previously been studied in other plants [24]. Stimulation of shoot and root length was noted in lower doses of SA and EMS. It has also been noted in tomato that the seedling height is stimulated by low concentrations of mutagens [25]. Growth reduction is associated with auxin destruction, ascorbic acid changes, physiological and biochemical interference [22].

EMS and SA also affected the number of branches in Okra varieties under study compared to control plants. The maximum number of branches was noted in the M₂ generation, with EMS being 0.1%, while the Puhja variety had a slightly different response in 0.1, 0.2 and 0.3% of EMS in M₁, while in the M₂ generation. The maximum number of branches was noted with 0.1% and 0.2% EMS. For both of the M₁ and M₂ generations, SA showed almost similar results compared to control (Table 2). These results are similar to those reported by Biswas et al. [26] where the control plants produced significantly fewer branches than the mutagen in Trigonella species. The results are shown in (Table 2), the maximum shoot diameter was recorded in 0.1% of EMS treated plants. In the case of SA observed maximum diameter in 0.1% SA treated plant of Puhja compared to control plants when treated with SA and EMS, sabz pari variety shows a reduced diameter stem.

The results showed in table 3 revealed that as the concentration of the mutagens (EMS and SA) increased, the number of days of first flowering decreased. The effect of a higher dose of mutagens on flowering was also reported in urdbean [27] and garlic [28]. The number of fruits/plants is shown in Table 3. The results showed that the maximum number of fruits (23.0±1.1 and 21.0±1.4) was developed from 0.2% and 0.1% of EMS exposure in Puhja compared with control (15.0±1.2). An increase in number of fruits was noted due to 0.1 and 0.2% levels of EMS and 0.1% SA. Similar results were also obtained in cowpea [29]. The promotion of biological parameters by EMS and low-dose gamma-rays was previously reported in Vicia faba L. and Cicer arietinum [16, 30].

The results show that the effects of the two mutagenesis treatments are different and the varieties are diverse and passed down from generation to generation. According to results (Table 3), all mutagenic exposures of EMS and SA reduced the fruit size in M₁ compared with control. This decrease in fruit size was significant in higher concentrations (0.2 and 0.3%) of EMS and SA. The minimum fruit length (8.3±1.9 cm) was observed in 0.3% SA treated Puhja. However, an increase in fruit length was observed in two treatments in M₂ generation treated with 0.1% mutagen. The maximum fruit size (12.5±1.1 cm) was observed in M₂ plants previously treated with 0.1% EMS compared with control (11.5±0.3 cm) in Subs pari. Benke [28] also reported changes in garlic plants treated with gamma rays. Our results are not in agreement with Goyal et al. [27] who reported a gradual increase in fruit size with increasing concentrations of DES and EMS in urdbean.

The result showed (Table 3) that the total number of seeds per fruit for each chemical treatment was variable by variety and generation. An increased seed yield (29.6±1.2 g) per plant was noted in 0.1% SA treated M₂ plants of Subs pari variety as against 25.8±0.4 g in control while the minimum seed yield (17.8±0.8 g) was observed in M₂ plants of Subs pari treated with 0.3% EMS. This variability in yield of Okra seeds per plant was also reported by other researchers [31, 32].
Table 3. Effect of EMS and SA treatments on different yield parameters of Subz Pari and Pahuja varieties of Okra (Mean±S.D)

| Treatments | Pahuja | M1 generation | Subz Pari | M1 generation |
|------------|--------|--------------|-----------|--------------|
|            | Days to flowering | No of fruits / plant | Fruit size/ length (cm) | Seeds yield / plant (g) | 100 Seeds weight (g) | Days to flowering | No of fruits / plant | Fruit size/ length (cm) | Seeds yield / plant (g) | 100 seeds weight (g) |
| Control    | 60     | 15.0±1.2     | 10.3±0.4 | 27.5±0.6 | 5.86 | 52     | 13.5±1.2 | 11.7±0.5 | 26.0±0.5 | 6.3 |
| EMS (%)    | 0.1    | 58           | 21.0±1.4 | 8.8±0.5 | 20.6±1.8 | 5.3 | 46     | 14.0±0.5 | 10.4±0.5 | 20.5±1.5 | 5.3 |
|            | 0.2    | 46           | 23.0±1.1 | 8.7±1.2 | 18.5±2.1 | 5.1 | 43     | 13.3±0.7 | 9.7±1.1 | 19.4±1.8 | 5.28 |
|            | 0.3    | 47           | 14.5±0.7 | 8.9±1.1 | 21.0±2.2 | 5.0 | 32     | 10.5±0.7 | 7.5±1.2 | 18.3±1.6 | 4.89 |
| SA (%)     | 0.1    | 57           | 16.5±2.1 | 8.6±0.5 | 24.3±1.2 | 5.4 | 54     | 16.0±1.1 | 11.4±0.6 | 27.3±1.2 | 5.9 |
|            | 0.2    | 52           | 10.0±1.2 | 9.5±1.2 | 28.2±1.5 | 5.0 | 48     | 13.5±0.7 | 9.7±1.6 | 22.2±1.5 | 5.16 |
|            | 0.3    | 49           | 9.7±1.5  | 8.3±1.9 | 19.1±2.2 | 4.9 | 42     | 12.2±2.1 | 8.5±1.1 | 18.3±1.6 | 5.01 |

EMS and SA treatments also affected the seed weight of both okra varieties. The 100 seeds of each treatment were randomly collected and weighed. The results showed that there was a slight decrease in 100 seeds weight on different concentrations of EMS and SA in the M1 generation against control (Table 3). But M2 showed different results then M1 generation as an increase in 100 seeds weight was noted in M2 generation against control. The maximum seeds weight (7.74 g) was observed in Subs pari fruits of M2 plants treated with 0.1% SA against control (6.25 g). In cowpea reported similar results observed by [29].Effector mutagens have been found to reduce the quantitative traits of soybeans [33]. The obtained variability in all levels of mutagenesis observed for various traits revealed a range of improvements [34].

**Conclusion**

Okra is an important multipurpose vegetable rich in carbohydrates, proteins, minerals, vitamins, etc. Unfortunately, very little work is being done on this crop in Pakistan. The results showed that different concentrations of EMS and SA affected morphological attributes of both okra varieties as the increase of mutagen concentration showed variable results regarding morphological attributes in M1 generation against control, but M2 generation showed useful improvement in various traits. Such as the M2 generation of Subs pari from 0.1% SA treatment showed a decrease in flowering time while an increase in the number of fruits per plant, fruit size, seeds yield (g) per plant and 100 seeds weight. It is suggested that M3 generation of Okra varieties may be cultivated on different locations and morphological attributes may be further evaluated.

**Authors’ contributions**

Conceived and designed the experiments: M Rafiq. Performed the experiments: N Noor-ul-Ain & MM Nizamani. Analyzed the data: AH Kaleri, J Gul & MM Nizamani. Contributed materials/ analysis/tools: M Rafiq &SHA Naqvi. Wrote the paper: MM Nizamani.

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