

DNA Damage Sensitivity Assays with *Arabidopsis* Seedlings

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[Abstract] We describe fast and reproducible sensitivity assays to quantify the response of *Arabidopsis* seedlings of different genotypes to a wide range of DNA damaging agents. We apply (1) γ -irradiation, which produces DNA breaks, (2) bleocin, a radiomimetic drug, (3) mitomycin C, a DNA intrastrand cross-linker, (4) hydroxyurea, an inhibitor of DNA synthesis and (5) UV-C, which causes mainly photoproducts. The “true leaf assay” and the “UV resistance assay” are based on easily determined phenotypes as readouts. Using a set of diverse damaging agents combined with different readouts allows establishing relative sensitivity/resistance compared to a reference line, e.g. wild type, determining the most effective type of induced damage and the potential repair pathway affected.

Materials and Reagents

1. *Arabidopsis thaliana* seeds [Wild type and T-DNA insertion lines of *arp6-3*, *swc6-1* and sensitive *ku70-2* (Rosa *et al.*, 2013) are used as examples. All mutants are in the Columbia-0 background.]
2. Sodium hypochlorite (Sigma-Aldrich, catalog number: 425044)
3. Tween-80 (Sigma-Aldrich, catalog number: 4780)
4. Sterile H₂O
5. Solid growth medium
6. Liquid plant growth medium (same as solid growth media but without agar)
7. Hydroxyurea (Sigma-Aldrich, catalog number: H8627)
8. 70% ethanol
9. Seed sterilization solution (see Recipes)
10. Bleocin (commercial name for bleomycin) (EMD Millipore, catalog number: 203408) (see Recipes)
11. Mitomycin C (Duchefa Biochemie BV, catalog number: M0133) (see Recipes)

Equipment

1. Sterile hood, preferably a biological safety cabinet to avoid exposure to the genotoxins
2. Bench top block shaker (e.g. Eppendorf, Thermomixer®)
3. UV crosslinker (254-nm UV light bulbs, 15 watts each) (Stratagene, model: Stratalinker 2400)
4. Gamma-irradiation source (Nordion, model: Co-60- Gamma-cell 220)
5. Forceps
6. Box or aluminum foil
7. Petri dishes for plant culture with solid growth medium (see Notes) (round 200 x 15 mm and 55 x 15 mm, and square 100 x 100 x 15 mm)
8. 1.5 ml Eppendorf tubes
9. Plant growth facilities with 16-h-light/8-h-dark cycles, at 21 °C

Procedure

A. Seed sterilization

1. Wash the seeds in a 1.5 ml Eppendorf tube with 1 ml of the sterilization solution, shake in a block shaker at 300 rpm for 6 min at room temperature. Allow seeds to set in the bottom of the Eppendorf and remove supernatant. Wash with 1 ml of sterile water, 5 times for 5 min in the shaker. After the final wash, remove supernatant and dry overnight in the closed sterile hood, leaving the tubes open.

Note: Sterilize approximately 350 seeds per genotype and treatment. All subsequent steps involving seed or plantlet manipulation should be carried in the sterile hood.

B. True leaf assay

1. Plate seeds in solid growth medium (\approx 75 to 100 per plate) using flame-sterilized and cooled forceps. Stratify in the dark (in a box or wrapped in aluminum foil) at 4 °C for 2 to 4 days and transfer to the growth chambers with illumination.

Note: Any other sterile plating techniques are suitable. Each plate makes one biological replicate; for each data point at least 3 biological replicates should be analyzed.

2. Treatments

- a. Gamma-irradiation: Treat 4-day-old-seedlings with a pulse of 100 Gray (Gy) with the Co-60- gamma source, with a dose of 27 to 34 Gy/min. Return plates to standard growth conditions to recover for 6 days until analysis.

Note: 4-day-old = 4 days after transfer to the growth chambers.

- b. Bleocin and mitomycin C: Using flame-sterilized and cooled forceps, transfer 4-day-old seedlings to small petri dishes containing 10 ml of liquid growth medium, either

without (mock) or with a drug (1 $\mu\text{g/ml}$ of bleocin or 10 $\mu\text{g/ml}$ of mitomycin C). After 5 days of incubation (with seedlings floating in liquid, but no shaking) in the illuminated growth chamber (9-day-old seedlings), remove the medium, wash extensively by flooding the plate 5 times with 20 ml of liquid media. Transfer seedlings to solid plates using flame-sterilized and cooled forceps. Allow seedlings to recover for 24 h before analysis.

Note: Seedlings should be carefully picked and handled, avoiding damage by the forceps. For examples of results see Figure 1A-C.

- c. Hydroxyurea: Plate seeds directly on plates, without or with 1 mM of the chemical, and grow at standard conditions until analysis at day 10.

3. Analysis

At day 10, photograph the plates for reference, and score the “percentage of plants with true leaves”.

Notes:

- a. *Upon undisturbed development, 10-day-old Arabidopsis seedlings possess one pair of true leaves.*
- b. *DNA damage causes the cells in the apical meristem to stop dividing, so that no true leaves are developed (Figure 1A).*
- c. *For the quantitative assay, calculate the percentage of plants with true leaves for each treated population in relation to the number of plants with true leaves after mock treatment. This calculation separates the effect of the DNA-damaging treatment from other factors, like vigor of the mutants or environmental factors.*
- d. *Only paired and normally shaped leaves should be scored; single, or small and narrow unexpanded leaves are not considered as true leaves.*

C. UV damage assay:

1. Plate seeds in solid growth medium (≈ 75 to 100 per plate) using flame-sterilized and cooled forceps. Stratify in the dark (in a box or wrapped in aluminum foil) at 4 °C for 2 to 4 days and transfer to the growth chambers with illumination.

Notes: Each plate makes one biological replicate; for each data point at least 3 biological replicates should be analyzed.

2. Treat seedlings at day 4 of growth, with 3 kJ/m^2 UV-C in a Stratalinker. Grow under standard conditions until day 12 and score the percentage of resistant plants (Figure 1D). Sensitive plants have phenotypes such as reduced global size, chlorotic cotyledons and senescence, whilst resistant plants have a normal size range and remain green.

Note: Divide the value in treated populations by the same parameter in the mock treatment, and average the biological replicates.

Before the treatment sterilize the interior of the Stratalinker with 70% ethanol and run it for 3 min with the maximum dose of UV-C. Then place plates inside and remove the lid, keeping it inside the Stratalinker. Quickly close the door, to avoid contamination and run the desired dosage of UV-C. When the treatment is finished close the plates still inside of the Stratalinker and return them to normal growth conditions.

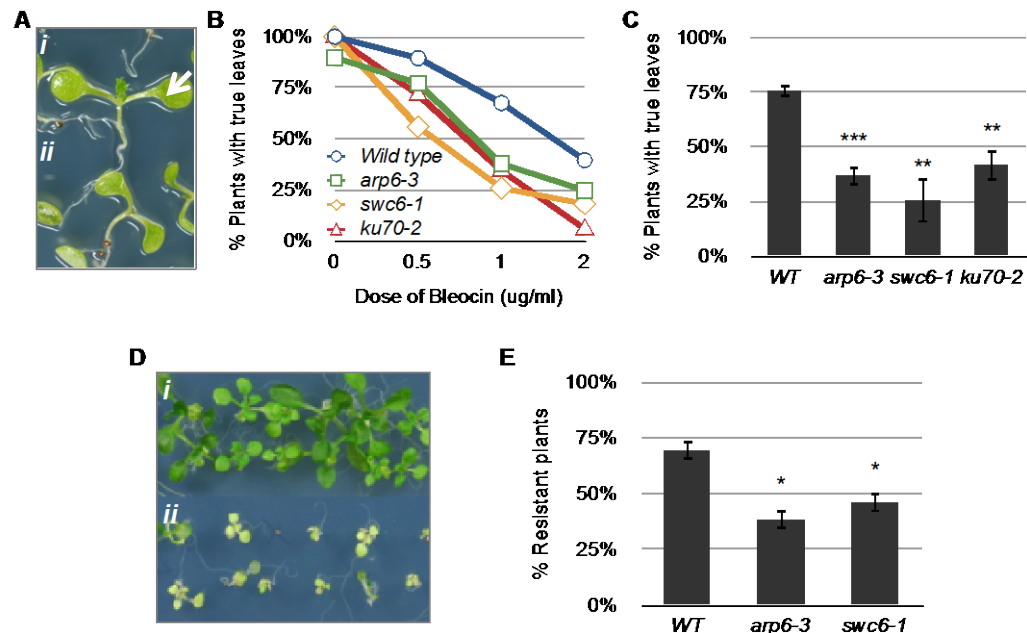


Figure 1. Examples of results. (A) Phenotypes of 10-day-old bleocin-treated seedlings with (*i*, arrow) and without (*ii*) true leaves. (B) True leaf assay showing the response of wild type, *arp6-3*, *swc6-1* and *ku70-2* to different doses of bleocin. Each plotted value corresponds to the scoring of one experiment with 60-80 seedlings. (C) True leaf assay with seedlings treated with 1 µg/ml of bleocin. The percentage of 10-d-old treated plants with true leaves was calculated in relation to mock samples. Bars depict the averages between 3 to 5 experiments with 75 to 100 seedlings per treatment. (D) Phenotypes of 12-day-old UV-C-treated resistant (*i*) and sensitive (*ii*) seedlings. (E) UV resistance assay of seedlings treated with 3 kJ/m² of UV-C. Bars depict the averages between 4 experiments with 75 to 100 seedlings per treatment. Asterisks indicate the significance between treated and non-treated samples according to P values from unpaired *t* tests: ****P* < 0.001 and **0.001 < *P* < 0.01.

Notes

1. To better characterize the response of the different genotypes to the DNA damaging agents, generate response curves with at least 3 different concentrations of the

- genotoxins (example in Figure 1B): 50 to 200 Gy of gamma-irradiation, 0.5 to 2 mg/ml of bleocin, 5 to 20 of mitomycin C, 0.5 to 2 of hydroxyhurea and 1.5 to 6 kJ/m² of UV-C.
2. Always keep the plates belonging to one experiment (mock + treatment) growing side-by-side. For example, take plates for mock and gamma-irradiation treatment out of the growth chamber at the same time.
 3. In this experiment we used Germination Media (GM) described in Reference 2 (detailed protocol available at <http://www.gmi.oeaw.ac.at/research-groups/ortrun-mittelsten-scheid/research-groups/ortrun-mittelsten-scheid/gm-medium-protocol>). We have not tested the use of regular MS plates but have no reason to believe the media will be a determining factor as long as the plates used in each experiment come from the same batch of media.

Recipes

1. Seed sterilization solution
5% sodium hypochlorite and 0.05% Tween-80
Mix 500 µl of sodium hypochlorite and 5 µl of Tween-80 in 10 ml of sterile H₂O
2. Bleocin (working solution: 1 µg /ml; stock: 10 mg/ml)
Dissolve 100 mg of Bleocin in 10 ml of H₂O
Make 0.2 ml aliquots and keep at -20 °C until use
Avoid thawing/freezing cycles
Dilute to the desired concentration in liquid media, using disposable 50 ml Falcon tubes
3. Mitomycin C (working solution: 10 µg/ml; stock: 0.5 mg/ml)
Dissolve 2 mg of Mitomycin C in 4 ml of H₂O immediately before use
Dilute to the desired concentration in liquid media, using disposable 50 ml Falcon tubes

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References

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