A Simplified Procedure for Glucosinolates Quantification

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Introduction

The challenge in plant protection programs is to seek management alternative that mitigate environmental degradation while maintaining agricultural productivity and profitability.

Materials of natural origin are biodegradable and could serve as environmentally safe pesticide.

Glucosinolates (GSLs) is a group of naturally occurring thioglucosides in Cruciferous vegetables that can be used as biofumigants.
- **Biofumigation** is the incorporation of plant residues that release pesticidal compounds into the soil as they decompose. Cruciferous crops (e.g. mustard, cabbage, turnip, canola) can be used as biofumigants because they contain GSL, long recognized as powerful natural pesticides.

- GSL-containing plants inhibit weed seed germination and some soil pathogens. They can be used as cover crops to reduce dependence on synthetic pesticides.

- *Sclerotinia sclerotiorum* & *Phytophthora capsici* are two broad-spectrum soil-borne diseases that cause severe damage to vegetable production.
Sclerotinia rot of carrot caused by Sclerotinia sclerotiorum
Stem Lesion in pepper caused by *Phytophthora capsici*
Background

- *Sclerotinia sclerotiorum* is found worldwide on a wide range of host plants: 64 plant families - 225 genera and 361 species.

- The infected areas have a bleached appearance.

- >120 different GSL have been described - most have cancer preventing activities in human diet.

- On the contrary, GSLs develop goiter in animal food.

- GSLs by themselves are not biologically active. The volatile hydrolysis products isothiocyanates are toxic to some soil-borne plant pathogens.
- GSLs and their hydrolysis products are responsible for the sharp or biting taste of condiments (horseradish or mustard) and contribute to the characteristic flavors of plants whose leaves (brussels sprouts, cabbage), floral buds (broccoli, cauliflower), stems (Kohlrabi) or roots (radish, turnips) are consumed by humans.
The present study is a continuation of our previous work on natural products for pest control.

Objectives
1- To develop and validate a simplified procedure for quantification glucosinolates in *Brassica* accessions.

2- To identify *Brassica* species and/or accessions with high levels of glucosinolates for future research on fungicidal efficiency of their crude extracts.
Ten accessions that demonstrated relative cold tolerance, rapid maturity, and superior biomass production were selected from 48 accessions of the National Germplasm Repository:

<table>
<thead>
<tr>
<th>Accession</th>
<th>Species</th>
<th>Location</th>
</tr>
</thead>
<tbody>
<tr>
<td>Ames 8660</td>
<td><em>Brassica juncea</em></td>
<td>Greenhouse</td>
</tr>
<tr>
<td>Ames 8674</td>
<td><em>Brassica juncea</em></td>
<td>High Tunnels</td>
</tr>
<tr>
<td>Ames 8709</td>
<td><em>Brassica juncea</em></td>
<td>Field</td>
</tr>
<tr>
<td>Ames 8887</td>
<td><em>Brassica juncea</em></td>
<td>Indian mustard</td>
</tr>
<tr>
<td>PI-120923</td>
<td><em>Brassica juncea</em></td>
<td></td>
</tr>
<tr>
<td>PI-603015</td>
<td><em>Brassica juncea</em></td>
<td></td>
</tr>
<tr>
<td>Pacific gold</td>
<td><em>Brassica juncea</em></td>
<td></td>
</tr>
<tr>
<td>PI-169083</td>
<td><em>Brassica napus</em></td>
<td>Oil seed rape</td>
</tr>
<tr>
<td>PI-633215</td>
<td><em>Eruca sativa</em></td>
<td>Arugula</td>
</tr>
<tr>
<td>Ida gold</td>
<td><em>Brassica campestris</em></td>
<td>Field mustard</td>
</tr>
</tbody>
</table>
Analysis of Glucosinolates

- Extraction using boiling methanol
- Clean-up using celite column
- Separation by anion-exchange resin “Sephadex”
- Hydrolysis with Thioglucosidase
- Quantification
\[ K\overset{-}{O}_3S - O - N = C - S \]

- \( R = 2\)-hydroxy-3-butenyl; \( \text{CH}_2=\text{CHCHOHCH}_2^- \); Progoitrin
- \( R = 2\)-phenylethyl; \( \text{C}_6\text{H}_5\text{CH}_2\text{CH}_2^- \); Gluconasturtiin
- \( R = 3\)-indolylmethyl; \[ \text{N} \]
  \[ \text{H} \]; Glucobrassicin
- GC analysis is not amendable to some GSLs containing polar side chains (Lee et al. 2006).

- Total sulfur determination by wet digestion and ICP analysis (Bloem et al. 2005).

$$\text{Sulfatase} \uparrow \text{Thioglycosidase} \uparrow$$

\[ \text{K} \text{O}_3 \text{S} - \text{O} - \text{N} = \text{C} \text{S} \]

OH

OH

OH
- Separation was accomplished by adsorption on DEAE- Sephadex A-25 (2-[diethylamino] ethyl ether) ion exchange resin of 40-125 µm bead size.

- DEAE was pre-swelled overnight in 2M ammonium bicarbonate.

- The column (35 cm x 3 cm) was eluted with aqueous ammonium acetate (0.1M, 0.5M, 1M and 2M), one liter of each.

- Fractions containing GSLs were then combined, evaporated to dryness under reduced pressure and quantified.
Glucosinolates Separation

Sephadex

\[
\begin{align*}
\text{H} & \quad \text{N} & \quad \text{H} \\
\text{H} & \quad \text{H}
\end{align*}
\]

\[
\begin{align*}
\text{OH} & \quad \text{OH} \\
\text{NH}_4 & \quad \text{K}_3 \text{O}_3 \text{S} & \quad \text{O} & \quad \text{N} & \quad \text{C} \\
\text{S} & \quad \text{O} & \quad \text{S}\end{align*}
\]

\[\text{R}\]

Thioglucosidase
Sinigrin = volatile mustard oil + myrosinase, 5 mg/mL were incubated at 37 °C for 4 hrs.

\[ y = 0.0008x \]

\[ R^2 = 0.9493 \]
Conclusion

Environmental stress on growing plants can increase the concentration of GSLs per gm fresh tissue.

*Brassica Juncea* accession “Pacific gold” & *Brassica napus* accession PI-169083 are the most promising biofumigation cover crops among those tested.
THANK YOU