# **Tackling Acrylamide in Canadian Wheat Bakery Products**

Ali Salimi Khorshidi<sup>1\*</sup>, Pham A. Tuan<sup>2</sup>, Belay T. Ayele<sup>2</sup>, Martin G. Scanlon<sup>1</sup> <sup>1</sup>Department of Food and Human Nutritional Sciences, University of Manitoba <sup>2</sup> Department of Plant Science, University of Manitoba \*Corresponding author. E-mail: Ali.SalimiKhorshidi@umanitoba.ca

Introduction	Results				
• Monitored by WHO & European Commission	<b>Table 1.</b> Expression levels of genes in studiedcultivars at 14 and 21 dpa				
• Benchmark levels set	Cultivar	dpa	Relative Expression		
y Caused by wheat asparagine (ASN)	Cultivar	upa	TaASN1	TaASP2	TaGSr1
	Brandon	14	7.02	0.59	1.16
e editing to reduce asparagine dentifying genes related to ASN		21	26.1	0.06	0.27
	BW5011	14	4.79	1.20	1.90
Knocking down suspect genes		21	19.8	0.11	0.58
Develop low asparagine cultivars	Glenn	14	4.83	0.91	0.59
Objective		21	19.4	0.12	0.48
	Prosper	14	1.25	0.86	0.58
derstand which genes are involved		21	6.89	0.29	0.42
umulation of ASN in Canadian					
ercial wheat cultivars					

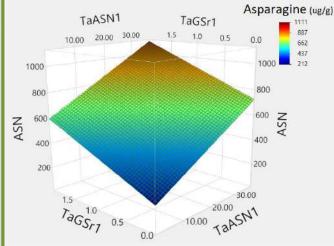


Figure 1. Relationship between TaASN1, TaGSr1 and wheat ASN concentration

- Significant differences in the expression levels among cultivars (Table 1)
- No effect of location on expression levels of almost all genes
- Significant change in expression levels of genes from 14 dpa to 21 dpa (Table 1)
- Strong correlations between ASN and expression levels of asparagine synthetase and glutamine synthetase genes (Figure 1)

### **Highlights**

- Significant differences in expression levels among Canadian varieties for asparagine and glutamine synthetase genes
- Genes targeted for development of low ASN cultivars through gene-editing identified

## Reference

Curtis et al. (2020). Contrasting gene expression patterns in grain of high and low asparagine wheat genotypes in response to sulphur supply. Ann Appl Biol. 2020;1–17.



Canada, Swift Current, SK.

## Gene

- 1. Id
- 2. Kr

Acry

Carc

Mainly

3. D

To und in accu comme

## **Materials & Methods**

Four commercial wheat cultivars (AAC Brandon, BW5011, Glenn, Prosper) were grown at two MB locations (2020) under 90 lb of Nitrogen per acre.

Whole wheat flour of mature seeds was analyzed for ASN level using an enzymatic kit and UV-Vis Spectrometry.

Seeds at 14 & 21 days post anthesis (dpa) were used for gene expression studies. Expression patterns of 10 genes from ASN synthetase, glutamine synthetase & asparaginase gene families were studied using RT-qPCR.